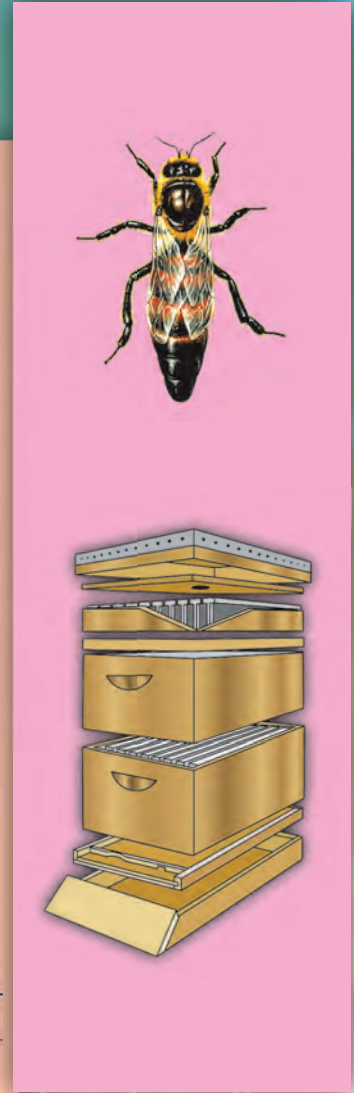
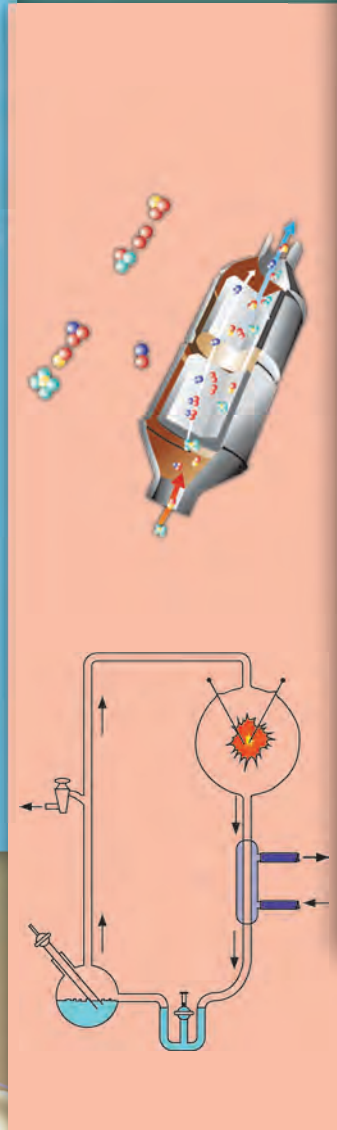
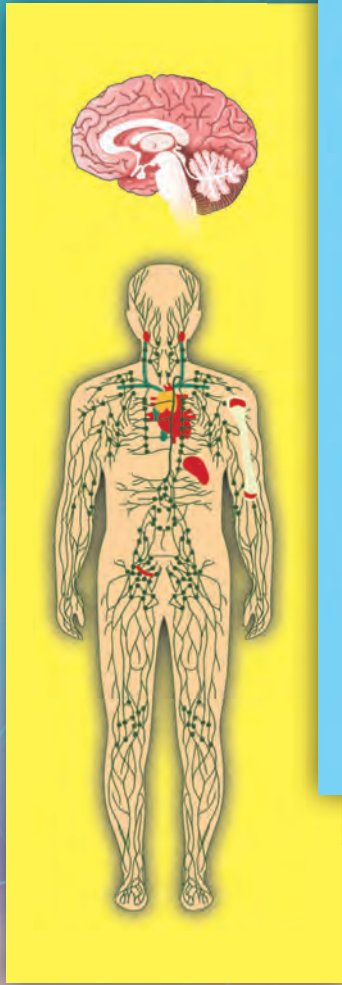




BIOLOGY

Standard XII



The Constitution of India

Chapter IV A

Fundamental Duties

ARTICLE 51A

Fundamental Duties- It shall be the duty of every citizen of India-

- (a) to abide by the Constitution and respect its ideals and institutions, the National Flag and the National Anthem;
- (b) to cherish and follow the noble ideals which inspired our national struggle for freedom;
- (c) to uphold and protect the sovereignty, unity and integrity of India;
- (d) to defend the country and render national service when called upon to do so;
- (e) to promote harmony and the spirit of common brotherhood amongst all the people of India transcending religious, linguistic and regional or sectional diversities, to renounce practices derogatory to the dignity of women;
- (f) to value and preserve the rich heritage of our composite culture;
- (g) to protect and improve the natural environment including forests, lakes, rivers and wild life and to have compassion for living creatures;
- (h) to develop the scientific temper, humanism and the spirit of inquiry and reform;
- (i) to safeguard public property and to abjure violence;
- (j) to strive towards excellence in all spheres of individual and collective activity so that the nation constantly rises to higher levels of endeavour and achievement;
- (k) who is a parent or guardian to provide opportunities for education to his child or, as the case may be, ward between the age of six and fourteen years.

The Coordination Committee formed by GR No. Abhyas - 2116/(Pra.Kra.43/16) SD - 4 Dated 25.4.2016 has given approval to prescribe this textbook in its meeting held on 30.01.2020 and it has been decided to implement it from academic year 2020-21.

BIOLOGY

STANDARD TWELVE



Download DIKSHA App on your smartphone. If you scan the Q.R.Code on this page of your textbook, you will be able to access full text and the audio-visual study material relevant to each lesson provided as teaching and learning aids.



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The Constitution of India

Preamble

WE, THE PEOPLE OF INDIA, having solemnly resolved to constitute India into a SOVEREIGN SOCIALIST SECULAR DEMOCRATIC REPUBLIC and to secure to all its citizens:

JUSTICE, social, economic and political;

LIBERTY of thought, expression, belief, faith and worship;

EQUALITY of status and of opportunity; and to promote among them all

FRATERNITY assuring the dignity of the individual and the unity and integrity of the Nation;

IN OUR CONSTITUENT ASSEMBLY this twenty-sixth day of November, 1949, do HEREBY ADOPT, ENACT AND GIVE TO OURSELVES THIS CONSTITUTION.

NATIONAL ANTHEM

Jana-gana-mana-adhināyaka jaya hē
Bhārata-bhāgya-vidhātā,

Panjāba-Sindhu-Gujarāta-Marāthā
Drāvida-Utkala-Banga

Vindhya-Himāchala-Yamunā-Gangā
uchchala-jaladhi-taranga

Tava subha nāmē jāgē, tava subha āsisa māgē,
gāhē tava jaya-gāthā,

Jana-gana-mangala-dāyaka jaya hē
Bhārata-bhāgya-vidhātā,

Jaya hē, Jaya hē, Jaya hē,
Jaya jaya jaya, jaya hē.

PLEDGE

India is my country. All Indians
are my brothers and sisters.

I love my country, and I am proud
of its rich and varied heritage. I shall
always strive to be worthy of it.

I shall give my parents, teachers
and all elders respect, and treat
everyone with courtesy.

To my country and my people,
I pledge my devotion. In their
well-being and prosperity alone lies
my happiness.

Preface

Dear Students,

We welcome you all to Std. XII. Now you are familiar to the subject of Biology as a separate discipline in standard XI. You have already been acquainted with many concepts of Biological Sciences from Standard six onwards, especially in the subject of General Science up to standard Eight and Science and Technology for standard Nine and Ten.

This textbook aims to create awareness about the biological sciences specially Botany, Zoology and allied aspects of biological sciences. The National Curriculum Framework (NCF) was formulated in 2005, followed by the State Curriculum Framework (SCF) in 2010. Based on the given these two frameworks, reconstruction of the curriculum and preparation of a revised syllabus has been undertaken which will be introduced from the academic year 2019-20. The textbook incorporating the revised syllabus has been prepared and designed by the Maharashtra State Bureau of Textbook Production and Curriculum Research, (Balbharati), Pune.

The subject biology intends to give students understanding, and appreciation of the vast diversity of living beings, their special adaptations to their environments and evolutionary relationships. No compromise is made in any manner over the use of language in the Biology context, but at the same time, the textbook is presented in a simple lucid language. In addition, relevant diagrams, graphs, tables used in the textbook will bring about more clarity in the understanding of various terminologies and biological concepts. All the illustrations are in colour form. This will surely enable students to understand various concepts of botany and zoology thoroughly and correlate this with their day-to-day practical life. The new syllabus focuses on the conceptual principles of overall life processes, its understanding, and application in day-to-day life and ability to solve different upcoming problems and issues like inheritance and its significance, conservation; different diseases and remedies, the application of technology, etc. The general teaching-learning objectives of the revised syllabus are further determined based on the 'principle of constructivism' i.e. self-learning.

The curriculum and syllabus confirms to the maxims of teaching such as moving from concrete to abstract, known to unknown and from part to whole. For the first time, in the syllabus of biology various independent activities have been introduced. These activities will not only help to understand the content knowledge but also provide scope for gaining relevant and additional application based knowledge on your own efforts. Q. R. Code have been introduced for gaining the additional information, abstracts of chapters and practice questions/ activities.

The efforts taken to prepare the textbook will not only enrich the meaningful learning experience of the students, but also benefit other stakeholders such as teachers, parents as well as those aspiring candidates preparing for the competitive examinations.

We look forward to a positive response from the teachers and students.
Our best wishes to all!



(Vivek Gosavi)

Director

Pune

Date : 21 February 2020

Bharatiya Saur : 2 Phalguna 1941

Maharashtra State Bureau of Textbook
Production and Curriculum Research, Pune 4

- For Teachers -

Dear Teachers,

We are happy to introduce the revised textbook of Biology for Std XII in continuation of Std XI. This book is a sincere attempt to follow the maxims of teaching as well as develop a 'constructive' approach to enhance the quality of learning and teaching as well. The present day education demands for more activity based, experimental and innovative learning opportunities is the need of the hour. The present curriculum has been restructured so as to bridge the credibility gap that exists between what is being taught and what students learn from the experiences in the outside world. Guidelines provided below will help to enrich the teaching-learning process to achieve the desired learning outcomes.

- To begin with, get familiar with the textbook.
- Always teach with proper planning.
- The present book has been prepared for constructive and activity-based teaching.
- Teachers must skillfully plan and organize the activities provided in each chapter to develop interest as well as to stimulate the thought process among the students.
- Use teaching aids as required for the proper understanding of the subject.
- Use demonstration, discussion method for teaching.
- Follow the order of the chapters strictly as listed in the contents because the units are introduced in a graded manner to facilitate knowledge building.
- Facilitate peer learning as much as possible by reorganizing the class structure frequently.
- Teaching-learning interactions, processes and participations of all students are very essential and so is your active guidance.
- Ask questions based on previous knowledge.
- Do not use the boxes titled 'Do you know?' for evaluation. However, teachers must ensure that students read this extra information.
- Information provided in boxes with the title 'Can You Tell', 'Always Remember' should be considered for evaluation.
- Exercise is given at the end of lesson. In exercise different type of questions/ activities are given.
- Exercises provided after each unit are prepared using different learning parameters like observation, co-relation, critical thinking, analytical reasoning etc.
- Evaluation pattern should be based on the above mentioned parameters. Equal weightage should be assigned to all the topics. Use different combinations of questions. Stereotype questions should be avoided.
- 'Can You Recall' is the first main starting point of lesson which helps for the introduction of topic. This will also helpful for students regarding understanding the content of lesson.
- 'Internet My Friend' is given for collecting extra important information related to topic.
- 'Use Your Brain Power' is used for the application level questions in different lessons.
- 'Do Your Self', 'Find Out', 'Observe and Discuss' and 'Try This' are used for activity based learning.
- 'Know the Scientist' is used for the information of different scientist related to concepts in lesson.
- 'Activity' is used in lesson and exercise for better understanding and application of the content which studied.
- Teacher should use their freedom to acquaint the students with flora and fauna of given region.
- Remember that mathematical and statistical tools are also important to understand biology
- List of abbreviations are provided towards the end of the textbook for further clarification.
- Use Q. R. Code given in the textbook.

Best wishes for a wonderful teaching experience and fruitful welcome!

Competency Statements

Standard XII

Unit	After studying content in the textbook student will....
Unit 1 : Reproduction	<ol style="list-style-type: none">1. Know the significance of reproduction in life of species.2. Explain the difference between asexual and sexual reproduction in plants and animals.3. Recognize the importance of asexual and sexual reproduction in plants and animals.4. Compare and analyze different modes of asexual reproduction.5. Know the reduction in the size of gametophytic generation.6. Know the different adaptations in the flowers depending upon the agency to accomplish pollination.7. Describes mechanism of sexual reproduction.8. Recognize, analyze and compare structural similarities, differences and progressive evolutionary changes in reproduction in lower and higher plants and animals.9. Explain embryo development both in plants and animals.
Unit 2 : Genetics and Evolution	<ol style="list-style-type: none">1. Explain the mechanism of inheritance and variation.2. Elaborate the role of chromosome, its molecular basis of heredity.3. Explain the laws of inheritance and further elaborate the reasons of variation.4. Describe the basis of origin of life, geological time scale, evidences.5. Explain, describe and compare different theories of evolution.6. Explain the structure and functions of genetic material.7. Use of genetics in studying patterns of sex determination in honey bees, birds and human beings mentioning different genetic disorders.8. Explain inheritance of sex linked characters in humans.9. Define concept of genomics, applications of genetic engineering and gene regulation.10. Explain chromosomal theory of inheritance, linkage and crossing over.11. Understand evidences for DNA as genetic material and genetic code.
Unit 3 : Physiology	<ol style="list-style-type: none">1. Explain the scientific reasons behind various physiological activities based on relationship.2. Understand the relationship between chemical reactions, structural organization involved and its impact on organism.3. Analyze and explain the experimental setup.4. Draw diagrams and give comments on findings and observations.5. Describe the contribution of different workers or scientists and its significance.6. Understand and explain role of physiology in biology.7. Explain and draw mechanisms of different physiological processes.8. Explain importance, source and methods of absorption of water, water as 'elixir of life'.9. Explain loss of excess water, significance of transpiration, transpiration as 'necessary evil'.10. Define growth, types of growth, phases of growth, growth curves, growth rates.11. Explain minerals, their role, sources and methods of absorption.12. Differentiate respiration and breathing.13. Explain circulatory system.

Unit 4 : Applied Biology	<ol style="list-style-type: none"> 1. Explains correlation between diseases and health. 2. Identify and elaborate various types and effects of Addications. 3. Elaborate the role of microbes in food production. 4. Describes, compares, reviews different techniques developed for betterment of life. 5. Understand applications of technology used to overcome problems in daily life. 6. Suggest remedial measures for improvement of social health. 7. Describe and suggest career opportunities in the fields of dairy, poultry and other field. 8. Explain role of microbes in upcoming fields as Biocontrol agents, Sewage treatment, Nanotechnology. 9. Elaborate the need of bio technology.
Unit 5 : Ecology and Environment	<ol style="list-style-type: none"> 1. Explains the correlation, interaction and effect of environment on organisms. 2. Understand and explain the relationship in ecosystem, role of energy flow. 3. Analyze, understand and explain environmental issues and their impact. 4. Contribute, plan and implement programs about conservation of environment. 5. Use information gathered to save biodiversity, find remedies to solve environmental issues.

Contents

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Can you recall?

1. How do plants reproduce without seeds ?
2. How does vegetative propagation occur in nature ?

Reproduction is the production of young ones like parents. Reproduction is an essential process as it leads to continuation of species as well as to maintain the continuity of life. Each organism has its own particular method of reproduction. All these methods generally fall into two categories :

- i. Asexual reproduction
- ii. Sexual reproduction.

1.1 Asexual Reproduction :

Asexual reproduction does not involve fusion of two compatible gametes or sex cells. It is the process resulting in the production of genetically identical progeny from a single organism and inherits the genes of the parent. Such morphologically and genetically identical individuals are called **clones**. Organisms choose to reproduce asexually by different modes or ways:

- i. **Fragmentation** : Multicellular organisms can break into fragments due to one or the other reasons. e.g. *Spirogyra*. These fragments grow into new individuals.

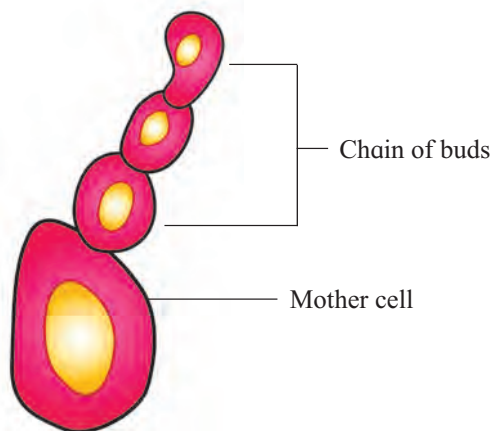


Fig. 1.1 : Budding in Yeast

- ii. **Budding** : It is the most common method of asexual reproduction in unicellular *Protosiphon* and yeast. Usually it takes place during favourable conditions by producing one or more outgrowths (buds). These buds on separation develop into new individual.
- iii. **Spore formation** : In *Chlamydomonas* asexual reproduction occurs by flagellated, motile zoospores which can grow independently into new individuals.

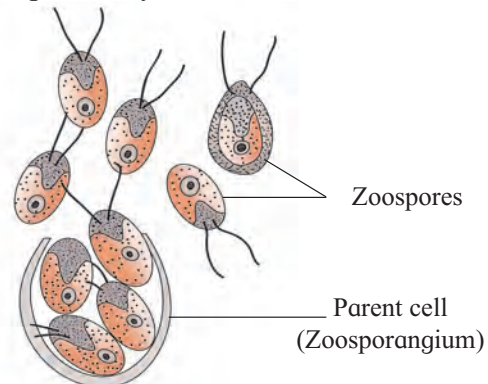


Fig. 1.2 : Zoospores in *Chlamydomonas*

Other methods of asexual reproduction include - **Binary fission** which occurs in *Chlorella*, *Diatoms* and *Chlamydomonas*; **Conidia formation** in *Penicillium* and **Gemma formation** as in *Marchantia*.



Activity :

Sprinkle a small spoonful of yeast pellets/powder over warm water and then add sugar. Cover it and wait for 10 minutes. Yeast becomes bubbly over the water proving that it is still active.



Can you recall?

The capacity to reproduce by vegetative propagation :

- Root - Sweet potato, *Asparagus*, *Dahlia*.
- Leaf - *Bryophyllum*, *Kalanchoe*, *Begonia*, etc.
- Stem - rhizome (turmeric), tubers (potato), bulbs (onion), etc.
- How does vegetative propagation occur in nature?

Vegetative Reproduction :

Plants reproduce asexually through their vegetative parts. Hence, the new plants formed are genetically identical to their parents.

There are also few methods which would not occur naturally in the plants. Agriculture and horticulture exploit vegetative reproduction in order to multiply fresh stocks of plants. Artificial methods are used to propagate desired varieties according to human requirements. The various methods are as follows :

a. Cutting :

The small piece of any vegetative part of a plant having one or more buds is used for propagation viz. Stem cutting - e.g. Rose, *Bougainvillea*; leaf cutting - e.g. *Sansvieria*; root cutting e.g. Blackberry.

b. Grafting :

Here parts of two plants are joined in such a way that they grow as one plant. In this method, part of the stem containing more than one bud (**Scion**) is joined onto a rooted plant called **stock**, is called grafting. Whereas budding is also called **bud grafting** in which only one bud is joined on the stock, e.g. Apple, Pear, Rose, etc.

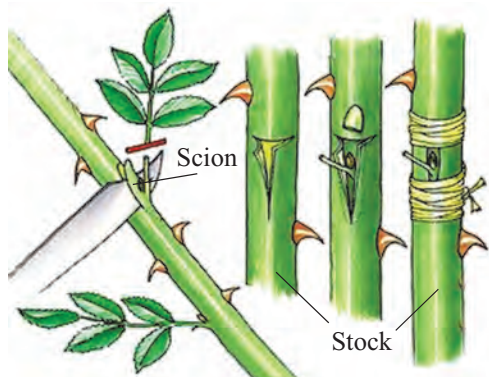


Fig. 1.3 : Grafting in Rose

c. **Tissue culture** : It is a method by which a small amount of plant tissue is



Do you know ?

Why does gardner choose to propagate plants asexually?

carefully grown to give many plantlets. Micropropagation method is also used now a days.

1.2 Sexual Reproduction :

It involves fusion of two compatible gametes or sex cells. All organisms reach to the maturity in their life before they can reproduce sexually. In plants, the end of juvenile or vegetative phase marks the beginning of the reproductive phase and can be seen easily in the higher plants at the time of flowering.

The flower is specialized reproductive structure of a plant in which sexual reproduction takes place. The function of flower is to produce haploid gametes and to ensure that fertilization will take place. Typical flower consists of four different whorls viz. calyx, corolla, androecium and gynoecium.

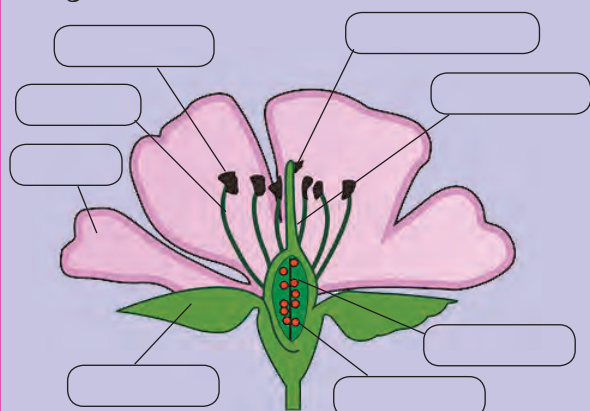
Sexual reproduction involves two major events viz. meiosis and fusion of gametes to form diploid zygote and the production of genetically dissimilar offsprings. Variations are useful from the point of view of the survival and the evolution of species, over the time.

Sexual reproduction is characterised by fusion of the male and female gametes (fertilization), the formation of zygote and embryogenesis. Sequential events that occur in sexual reproduction are grouped into three distinct stages viz, Pre-fertilization, Fertilization and the Post-fertilization.



Activity :

Label the parts of flower in the given diagram :





Always Remember

Diploid sporophyte is the predominant plant body in all angiosperms, where meiosis takes place to produce haploid spores that form gametophyte. Gametophytes are considerably reduced and develop within the flower. They produce gametes.

The male reproductive whorl of flower is called **androecium**. Individual member of androecium, is called **stamen**. Stamen consists of filament, connective and anther. having two anther lobes (theca).

Structure of Anther :

Anther is generally dithecous (having two lobes) and tetrasporangiate. Each lobe of anther contains two **pollen sacs**. In dithecous anther four pollen sacs are present. Therefore, it is **tetrasporangiate**. An immature stage of anther is represented by group of parenchymatous tissue surrounded by single layered epidermis. The heterogeneity (differentiation) arises when some hypodermal cells get transformed into **archesporial cells**.

T. S. of Anther :

The archesporial cell divides into an inner sporogenous cell and outer primary parietal cell. Sporogenous cell forms sporogenous tissue. Each cell of sporogenous tissue is capable of giving rise to a microspore tetrad. Parietal cell undergoes divisions to form anther wall layers. The wall of mature anther consists of four layers. **Epidermis** is the outermost protective layer made up of tabular (flattened) cells. **Endothecium** is sub-epidermal layer made up of radially elongated cells with fibrous thickenings. Inner to endothecium is **middle layer** made up of thin walled cells (1-2 layered), which may disintegrate in mature anther. **Tapetum** is the inner most nutritive layer of anther wall. It immediately encloses the sporogenous tissue (microspore mother cells).

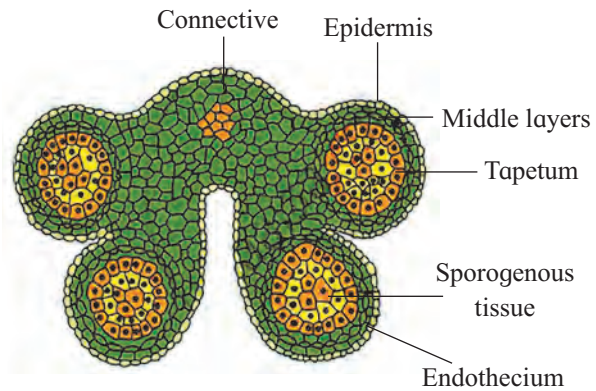


Fig. 1.4 : (a) T. S. of anther

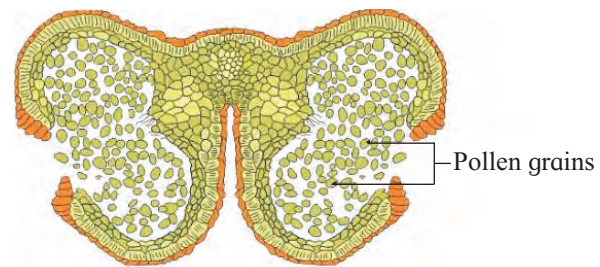


Fig. 1.4 : (b) Dehiscent anther

1.3 Microsporogenesis :

Each microspore mother cell divides meiotically to form tetrad of haploid microspores (**pollen grains**).

Structure of microspore :

Typical pollen grain is a non-motile, haploid, unicellular body with single nucleus. It is surrounded by a two layered wall called **sporoderm**. The outer layer **exine** is thick and made up of complex, non-biodegradable, substance called **sporopollenin**. It may be smooth or with a sculptured pattern (characteristic of the species). It is resistant to chemicals. At some places exine is very thin showing thin areas known as **germ-pores**. These are meant for the growth of emerging pollen tube during germination of pollen grain. The inner wall layer, **intine** consists of cellulose and pectin.



Find out

Why pollen grains can remain well preserved as fossil?



Always Remember

- Pollen viability (viability is the functional ability of pollen grain to germinate to develop male gametophyte) depends upon environmental conditions of temperature and humidity. It is 30 minutes in rice and wheat. But in some members of family Solanaceae, Rosaceae, Leguminosae, it lasts even for months.

Development of male gametophyte :

Pollen grain marks the beginning of male gametophyte. It undergoes first mitotic division to produce bigger, naked **vegetative cell** and small, thin walled **generative cell**. The vegetative cell is rich in food and having irregular shaped nucleus. The generative cell floats in the cytoplasm of vegetative cell.

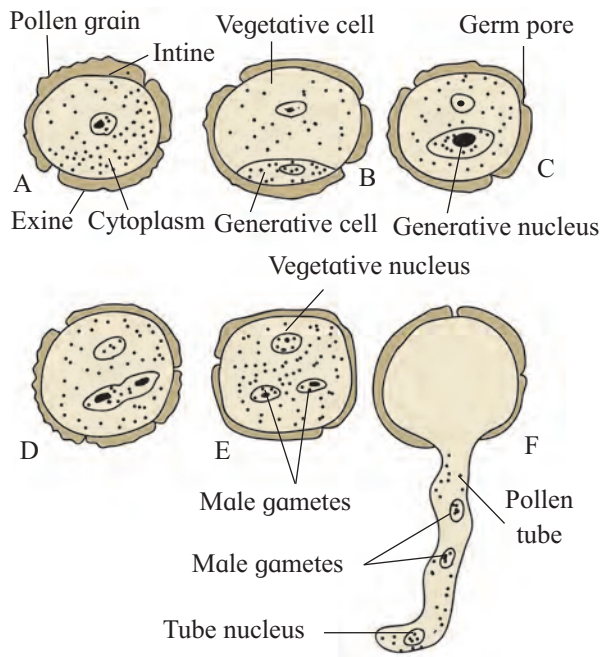


Fig. 1.5 : Development of male gametophyte

The second mitotic division is concerned with generative cell only and gives rise to two non-motile male gametes. The mitotic division of generative cell takes place either in pollen grain or in the pollen tube. The pollen grains are shed from the anther, at this two-celled stage in most of the angiosperms.

Female reproductive whorl of flower is **gynoecium** (Pistil). Individual member of gynoecium is called **carpel** (megasporophyll). A flower with many, free carpels is called **apocarpous** (e.g. *Michelia*). A **syncarpous** flower is one that has many carpels fused together (e.g. Brinjal). Typical carpel has three parts viz, ovary, style and stigma. The number of ovules in the ovary varies e.g. paddy, wheat and mango are **uniovulate** whereas tomato and lady's finger are **multiovulate**.

1.4 Structure of Anotropous ovule :

Each ovule develops inside the ovary and is attached to the **placenta** by a small stalk called **funiculus**. The place of attachment of funiculus with the main body of ovule, is called **hilum**. In angiosperms, the most common type of ovule is **anatropous** in which micropyle is directed downwards and is present adjacent to the funiculus (funicle). The ovule consists of central parenchymatous tissue, the **nucellus** which is surrounded usually by two protective coverings called **integuments** viz. Outer and an inner integument.

A narrow opening at the apex of the ovule is called **micropyle**. Chalaza is the base of ovule directly opposite to micropyle. **Embryo sac** (female gametophyte) is oval, multicellular structure embedded in the nucellus.

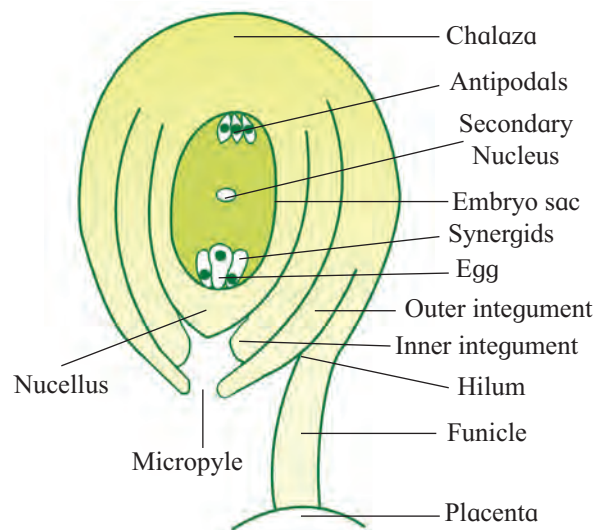


Fig. 1.6 : Anatropous Ovule

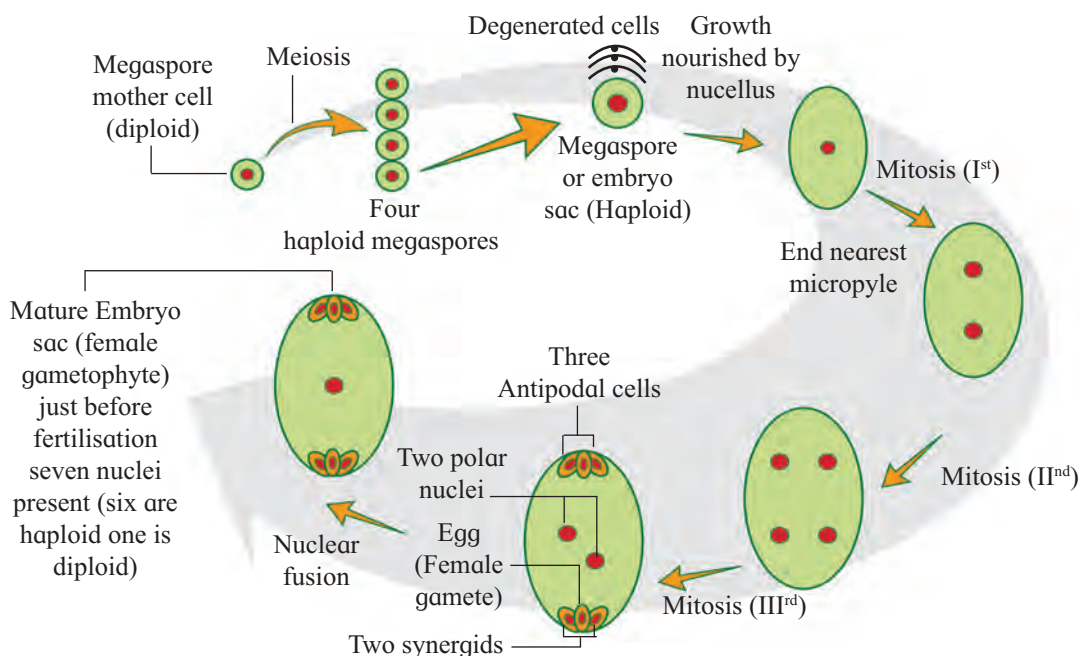


Fig. 1.7 : Development of female gametophyte

1.5 Megasporogenesis :

It is the process of formation of haploid megaspores from diploid **megaspore mother cell** (MMC). Megaspore mother cell becomes distinguished in the nucellus, more or less in the centre but towards micropylar end of ovule.

Development of female gametophyte :

Megaspore mother cell undergoes meiosis to form linear tetrad of haploid cells i.e. megaspores. Upper three megaspores abort and lowest one towards centre of nucellus remains functional. It acts as the first cell of female gametophyte. Generally one megaspore towards centre is functional megaspore. It is infact the first cell of female gametophyte. It undergoes three successive, free nuclear **mitotic** divisions. Thus, total eight nuclei are formed, four of which are located at each pole. One nucleus from each pole migrates towards the centre and are called **polar nuclei**. Three nuclei towards micropylar end constitute **egg apparatus**. It consists of large central, haploid egg cell and two supporting haploid synergid cells. Synergid shows hair like projections called **filiform apparatus**, which guide the pollen tube towards the egg.

Antipodal cells are group of three cells present at the chalazal end. The two haploid polar nuclei of large central cell fuse to form diploid **secondary nucleus** or **definitive nucleus**, just prior to fertilization. This seven-celled and eight nucleated structure is called an **embryo sac**. This method of embryo sac development from a single megaspore is described as **monosporic development**. In angiosperms, the development of female gametophyte is endosporous i.e. within the megaspore. Female gametophyte is colourless, endosporic and is concealed in the ovule enclosed by ovary.

1.6 Pollination :

Pollen grains being non motile, angiosperms have evolved the strategy to use abiotic agents (wind, water) and biotic agents (birds, insects, snails) to their flowers, feeding the visitors and exploiting their mobility for pollination and also seed dispersal. Pollen grains are non-motile and they are usually carried from flower to flower by means of external agents. Pollination is the transfer of pollen grains from anther to the stigma of the flower. It is the prerequisite for fertilization because both the male and female gametes are non-motile. Moreover gametes are produced at two different sites.

Self pollination is a type of pollination which occurs in a single flower or two flowers of a single plant. It results in inbreeding or selfing. In contrast cross pollination is the transfer of pollen grains from the anther of one flower to the stigma of another flower of different plants of same species. Pollination can be further divided into three types on the basis of source of pollination.

a. Autogamy (self pollination) :

It is a type of pollination in which bisexual flower is pollinated by its own pollen grains. Offsprings are genetically identical to their parents e.g. pea, *Clitoria*



Always Remember

- Flowers which use autogamy consist of several adaptations in the structure of a flower to facilitate this process. It occurs without external pollinating agents.
- When flower opens to expose its sex organs, it is called **Chasmogamous**.
- The contrivances (a condition that leads to) favour self pollination are- Bisexuality, Homogamy and Cleistogamy.
- Homogamy : When anther and stigma of a flower become mature at the same time, called **homogamy**.
- Some flowers are self pollinated even before the opening of flower. Such condition is called **cleistogamy**. Underground flowers in some plants which exhibit cleistogamy are never opened e.g. *Commelina benghalensis*.
- Plants like *Viola*, *Commelina* can produce both chasmogamous and cleistogamous flowers on the same plant.



Think about it

Why do some plants have both chasmogamous and cleistogamous flowers ?

b. Geitonogamy :

It is the transfer of pollen grain to a stigma of a different flower produced on the same plant. It is functionally similar to cross pollination as it involves pollinating agents, but it cannot bring about genetic variations and is only of ecological significance e.g. *Cucurbita maxima*. It is similar to autogamy as pollen grains come from same plant.

c. Xenogamy (cross polination/ out breeding) :

It is a type of cross pollination when pollen grains of one flower are deposited on the stigma of a flower of different plant belonging to same species, with the help of pollinating agency. It generates genetically varied offsprings.

Majority of flowering plants depend on the transfer of pollen grains. Virtually all seed plants need to be pollinated. Most of the food and fibre crops grown throughout the world, depend upon pollinators for reproduction.

The agents responsible for pollination have been grouped into two main categories :

A. Abiotic agents

B. Biotic agents

A. Abiotic Agents : These are non-living agents which include wind and water.

1. Pollination by wind (Anemophily) :

Most of the important crop plants are wind pollinated. These include wheat, rice, corn, rye, barley and oats. Palms are also wind pollinated.

Adaptations in anemophilous flowers :

- The flowers are small, inconspicuous, colourless, without nectar and fragrance (odour).
- The pollen grains are light in weight, dry and produced in large numbers to increase chances of pollination considering wastage of pollengrains.
- Stigma is feathery to trap pollens carried by wind currents.

- Stamens are exerted with long filaments and versatile anthers.
- Stamens and stigmas are exposed to air currents.

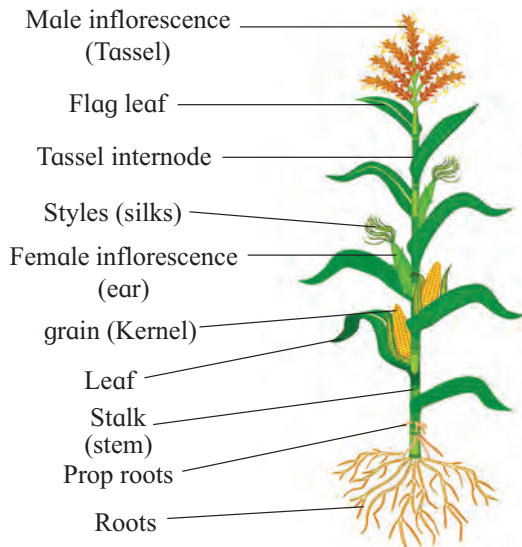


Fig. 1.8 : Pollination by wind (Maize)



Always Remember

The pollens of wind pollinated plants are most frequently associated with symptoms of hayfever among people those are sensitive to pollens. It is caused by hypersensitivity to pollen.

2. Pollination by water (Hydrophily) :

Found only in some 30 genera of aquatic monocots. E.g. *Vallisneria*, *Zostera*, *Ceratophyllum* etc.

Adaptations in hydrophilous flowers :

- Flowers are small and inconspicuous.
- Perianth and other floral parts are unwettable.
- Pollen grains are long and unwettable due to presence of mucilage.
- Nectar and fragrance are lacking in flowers.

Hydrophily is of two types -

Hypohydrophily : Pollination occurs below the surface of water. Here the pollen grains are heavier than water, sink down and caught by stigmas of female flowers, e.g. In *Zostera* (sea

grass) the pollen grains are long, ribbon like and without exine.

Epihydrophily : The pollen grains float on the water surface and reach the stigma of female flower. e.g. *Vallisneria* is a submerged dioecious, fresh water aquatic plant in which female flowers reach the water surface temporarily to ensure pollination and male flowers float on the surface of water.

- Specific gravity of pollen grains is equal to that of water. That is why they float on surface of water.
- Some aquatic plants are anemophilous e.g. *Potamogeton*, *Halogaris*, etc.
- Some aquatic plants are entomophilous e.g. Lotus, water hyacinth, waterlily, etc.

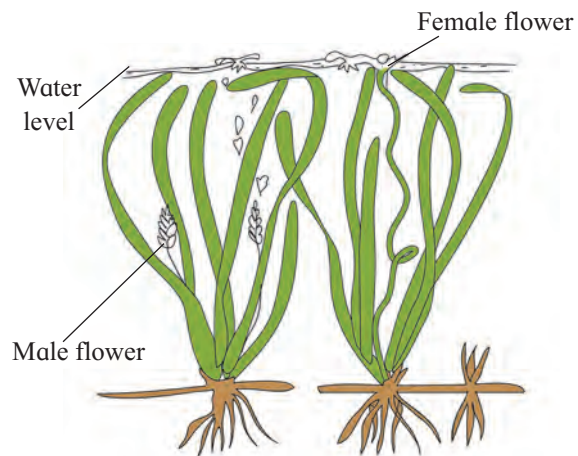


Fig. 1.9 : Male and female plants-Vallisneria

B. Biotic Agents : It includes living agents. About 80% of plants require the help of other living, moving creatures such as insects, birds, bats, snails to transfer their pollens from one flower to another. These also sustain our ecosystems and produce natural resources by helping plants to reproduce.

1. Pollination by insects (Entomophily) :

It occurs in Rose, Jasmine, *Cestrum*, etc.

Adaptations in entomophilous flowers :

- They are large, showy and often brightly coloured.
- The flowers produce sweet odour (smell) and have nectar glands.

- The stigma is rough due to presence of hair or is sticky due to mucilaginous secretion.
- The pollen grains are spiny and surrounded by a yellow sticky substance called pollen-kit.
- Some plants have special adaptations for the insect visitor to help in cross pollination, e.g. lever mechanism or turn-pipe mechanism in *Salvia*.



Do you know ?

In biotic pollination, plants are adapted to encourage the specific pollinators they need. They are said to have developed pollination contrivance. Plants and pollinators have co-evolved physical characteristics that make them to interact successfully. Such characteristics are considered pollination syndromes.



Fig. 1.10 : Lever mechanism in *Salvia*



Think about it

You may see bumblebee early in the year as they try to find a suitable place to establish a nest and rear a colony. If you find a bumblebee nest please leave it alone. Their nest lasts only for a season. Educate the world about the need to help the bees.

2. Pollination by birds (Ornithophily) :

Only a few types of birds are specialised for pollination. They usually have small size and long beaks e.g. Sun birds and humming birds. Some ornithophilous plants are *Bombax*, *Callistemon* (Bottle Brush), *Butea*, etc.



Fig. 1.11 : Ornithophily

Adaptations in ornithophilous flowers :

- Flowers are usually brightly coloured, large and showy.
- They secrete profuse, dilute nectar.
- Pollen grains are sticky and spiny.
- Flowers are generally without fragrance, as birds have poor sense of smell.

3. Pollination by Bats (Chiropterophily) :

Bats can transport pollens over long distance, some times several kilometers.

Adaptations in Chiropterophilous flowers :

- Flowers are dull coloured with strong fragrance.
- They secrete abundant nectar.
- Flowers produce large amount of edible pollen grains, e.g. *Anthocephalous* (kadamb tree), *Adansonia* (Baobab tree/ Gorakh chinch), *Kigelia* (Sausage tree).

1.7 Outbreeding devices (contrivances):

Many plants have mechanisms that discourage or prevent self pollination. To promote cross pollination and increase genetic diversity, plants have evolved a wide variety of sexual strategies. Genetic diversity is an essential factor for evolution by natural selection. Continued self pollination results in the inbreeding depression.

Thus, plants have developed many devices to encourage cross pollination. The examples of outbreeding devices are as follows:

Unisexuality :

In this case, the plant bears either male or female flowers. It is also called as **dioecism**. As flowers are unisexual, self pollination is

not possible. Plants may be monoecious, e.g. Maize or dioecious, e.g. Mulberry, Papaya.

Dichogamy :

In this device, anthers and stigmas mature at different times in a bisexual flower so as to prevent self pollination. It can be further divided into two types:

1. **Protandry** : In this type, androecium matures earlier than the gynoecium, e.g. in the disc florets of sunflower.
2. **Protogyny** : In this type, gynoecium matures earlier than the androecium, e.g. *Gloriosa*.

Prepotency :

Pollen grains of other flowers germinate rapidly over the stigma than the pollen grains from the same flower, e.g. Apple.

Heterostyly (heteromorphy):

In some plants like *Primula* (Primrose), there are two or three forms/ types of flowers in which stigmas and anthers are placed at different levels (heterostyly and heteroanthy). This prevents the pollens from reaching the stigma and pollinating it. In heteromorphic flowers, pollen grains produced from anther pollinate stigmas produced at the same level.

Herkogamy :

It is a mechanical device to prevent self pollination in a bisexual flower. In plants, natural physical barrier is present between two sex organs and avoid contact of pollen with stigma of same flower, e.g. *Calotropis*-pentangular stigma is positioned above the level of anthers (pollinia).

Self incompatibility (self sterility):

This is a genetic mechanism due to which the germination of pollen on stigma of the same flower is inhibited, e.g. Tobacco, *Thea*.



Do you know ?

In all breeding programmes, the plants are hand pollinated to ensure cross pollination between selected varieties. e.g. wheat, rice.

1.8 Pollen - Pistil Interaction :

It is the interaction of pollen grains with sporophytic tissue (stigma). It begins with pollination and ends with fertilization. All the events from the deposition of pollen grain on stigma to the entry of pollen tube in the ovule (synergid) are referred as pollen - pistil interaction. Pollination does not guarantee the transfer of right type of pollen, often wrong type also land on stigma. The pistil has the ability to recognise and accept the right or compatible pollen of the same species. Thus wrong type of pollen is discarded by pistil. Compatibility and incompatibility of the pollen-pistil is determined by special proteins. This process involves pollen recognition followed by promotion or inhibition of pollen.

The stigmatic surface of flower refuse other wrong type or incompatible pollen grains. A physiological mechanism operates to ensure that only intraspecific pollen germinate successfully. The compatible pollen absorbs water and nutrients from the surface of stigma, germinates and produces pollen tube. Its growth through the style is determined by specific chemicals. The stigmatic surface provides the essential prerequisites for a successful germination, which are absent in the pollen. The pollen tube is finally pushed through the ovule and reaches the embryo sac. The tip of the pollen tube enters in one of the synergids and then ruptures to release the contents. Due to pollen pistil interaction, intense competition develops even in the compatible pollen grains (gametes).

It also plays important role in sexual reproduction and seed formation. Pollen grain can also be induced to germinate in a synthetic medium. Sucrose induces pollen germination and tube growth *in vitro*. Addition of boric acid facilitates and accelerates pollen germination.

Artificial hybridization :

It is one of the major approaches used in the crop improvement. Only the desired

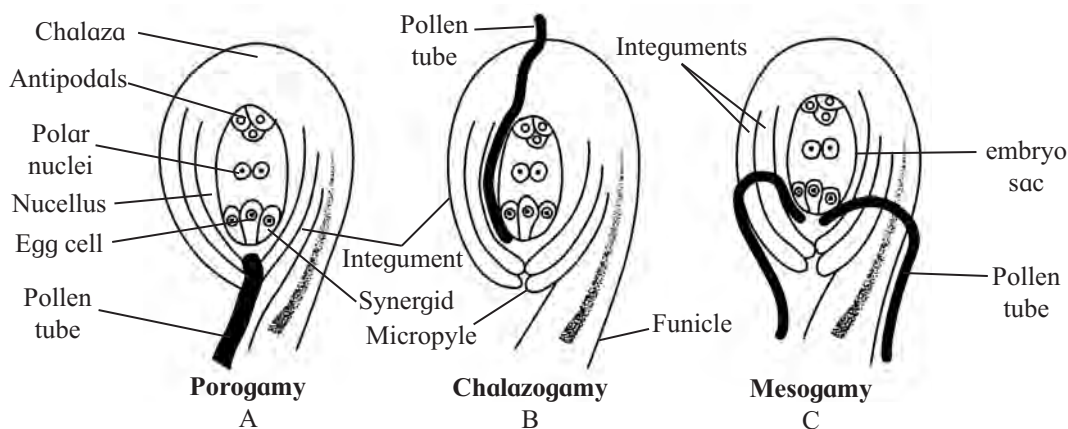


Fig. 1.12 : Entry of pollen tube into the ovule

pollen grains are hand pollinated and used for fertilization. This is accomplished through emasculation and bagging procedure.

1.9 Double Fertilization :

Double fertilization is a complex fertilization mechanism in flowering (angiospermic) plants. It was discovered by Nawaschin in the liliaceous plants like *Lilium* and *Fritillaria*.

After a pollen grain has reached the surface of the stigma, it germinates and forms a pollen tube, which penetrates the stigma, style, ovary chamber and then enters ovule. The growth of pollen tube is guided by the chemicals secreted by the synergids. It usually enters ovule through the micropyle. It is termed as **porogamy**. But in some cases, it is found to enter through chalaza, known as **chalazogamy** and in some plants by piercing the integuments, called **mesogamy**. Finally, it penetrates embryo sac of ovule at its micropylar end.

The pollen tube carrying male gametes penetrates in one of the synergids. Watery contents of synergid are absorbed by pollen tube which then ruptures and release the contents, including the two non-motile male gametes. As non motile male gametes are carried through hollow pollen tube, it is known as **siphonogamy** that ensures fertilization to take place. **Syngamy** and **triple fusion** are two events of sexual reproduction in angiospermic flowering plants. Syngamy is the fusion of

haploid male gamete with haploid female gamete (egg) to produce a **diploid zygote**, whereas in triple fusion, second haploid male gamete fuses with diploid secondary nucleus producing **primary endosperm nucleus (PEN)** that develops into **triploid endosperm**. The zygote develops into an embryo. Syngamy is a type of generative fertilization whereas triple fusion is a type of vegetative fertilization.

Here, both the male gametes participate and therefore, it is described as or called **double fertilization**.

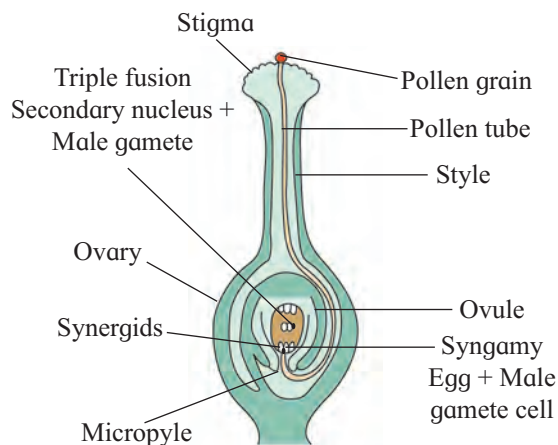


Fig. 1.13 : Double fertilization

Significance of Double Fertilization :

- It is a unique feature of angiosperms. It ensures that the parent plant invests a seed with a food store, only if the egg is fertilized.
- The diploid zygote develops into an embryo which consequently develops into a new plant.

- The triploid PEN develops into nutritive endosperm tissue.
- It restores the diploid condition by fusion of haploid male gamete with haploid female gamete (i.e. through syngamy).
- It also helps to avoid polyembryony.

1.10 Development of Endosperm :

The triploid primary endosperm nucleus repeatedly divides mitotically to form nutritive tissue, called **endosperm**. In post-fertilization changes within the ovule, the embryo and endosperm are seen to develop simultaneously.

The other cells of embryo sac disorganized sooner or later. The formation of triploid endosperm nucleus triggers cell division which leads to the formation of endosperm.

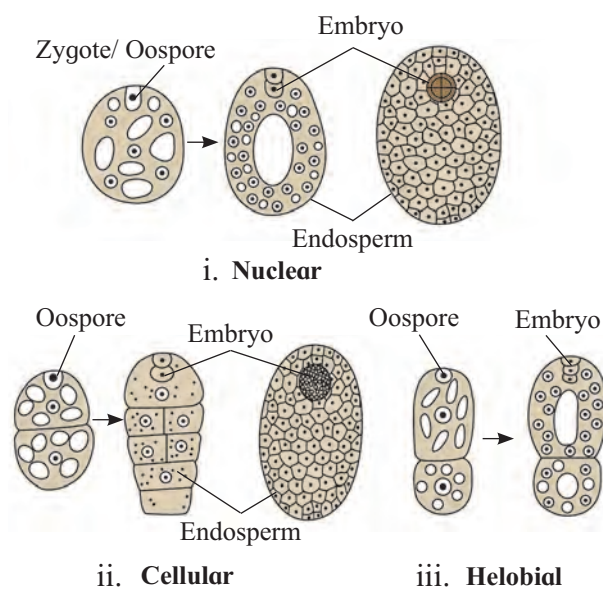


Fig. 1.14 : Types of Endosperm

There are three types of endosperms on the basis of mode of development. These are i. Nuclear type, ii. Cellular type, iii. Helobial type :

a. Nuclear Type :

It is the most common type found in 161 angiospermic families. Here, the primary endosperm nucleus repeatedly divides mitotically without wall formation to produce large number of free nuclei. A big central

vacuole appears in the centre of cell pushing the nuclei towards the periphery. Later, walls develop between the nuclei, hence multicellular endosperm is formed. But in several cases cell wall formation remains incomplete. e.g. wheat, sunflower and coconut. Coconut has multicellular endosperm in the outer part and free nuclear as well as vacuolated endosperm in the centre.

b. Cellular Type :

In some plants, division of triploid primary endospermic nucleus is immediately followed by wall formation. So that the endosperm is cellular right from the beginning. It is mostly observed in 72 families of dicots as in members - *Balsam*, *Petunia*, *Adoxa*, etc.



Use your brain power

What do you call the kernel that you eat in tender coconut ?

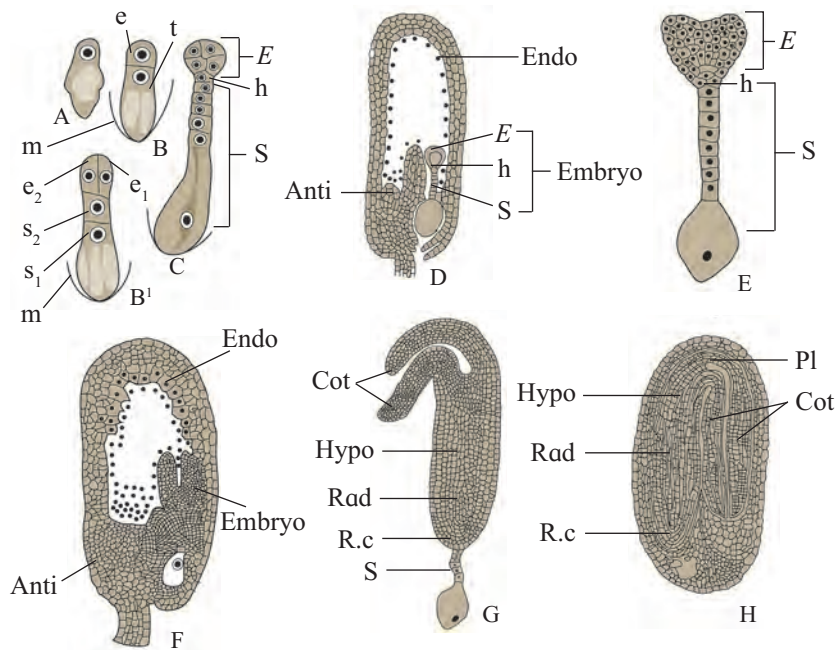
c. Helobial Type :

It occurs in the order Helobiales of monocotyledons. In this case, first division of primary endosperm nucleus is followed by a transverse wall, which divides the cell unequally. The smaller cell is called **chalazal cell** and larger cell is the **micropylar cell**. Then the nuclei in each cell divide by free nuclear divisions and then walls develop between nuclei in micropylar chamber. It is intermediate between cellular and nuclear type endosperm e.g. *Asphodelus*.

Mosaic Endosperm : Endosperm containing tissues of two different types is called mosaic endosperm. In plants like corn the endosperm contains patches of two different colours. It forms a sort of mosaic pattern.

1.11 Development of Embryo :

The process of development of zygote into an embryo is called **embryogenesis**. The embryo is developed at the micropylar end of embryo sac. The growth of embryo triggers only



A. Oospore. B. Two celled proembryo. e=embryonal initial; t=suspensor initial; m=Embryo sac membrane. B¹=4-celled I-shaped proembryo; e₁, e₂ are from embryonal initial; s₁, s₂ are from suspensor initial. C. Further development of embryo. S=Suspensor, h=Hypophysis; E=Embryonal mass D. L. S. of ovule Endo=Endosperm in free nuclear stage. Anti=Antipodal tissue. Embryo= Developing embryo E. Embryo showing further development of embryonic octants and hypophysis. F. L. S. of ovule. Endosperm becoming cellular. G. Embryo Cot=Cotyledons; Hypo=Hypocotyl; Rad=Radicule; R.c=Root-cap H. Mature seed. Pl=Plumule. Endosperm has been consumed almost completely.

Fig. 1.15 : Development of Dicot Embryo as in *Capsella*

after certain amount of endosperm is formed. After fertilization the embryonic development begins.

The zygote divides to form two-celled **proembryo**. The larger cell towards the micropyle is called basal or **suspensor initial** cell and smaller cell towards chalaza is called terminal or **embryonal initial** cell. The suspensor cell divides transversely in one plane to produce filamentous suspensor of 6-10 cells.

The first cell of the suspensor towards the micropylar end becomes swollen and function as a **haustorium**. The lowermost cell of suspensor is known as **hypophysis**. The suspensor helps in pushing the embryo in the endosperm. The embryonal initial undergoes three successive mitotic divisions to form octant. The planes of divisions are at right angles to each other. The lower tier of four cells of octant give rise to hypocotyl and radicle whereas four cells of

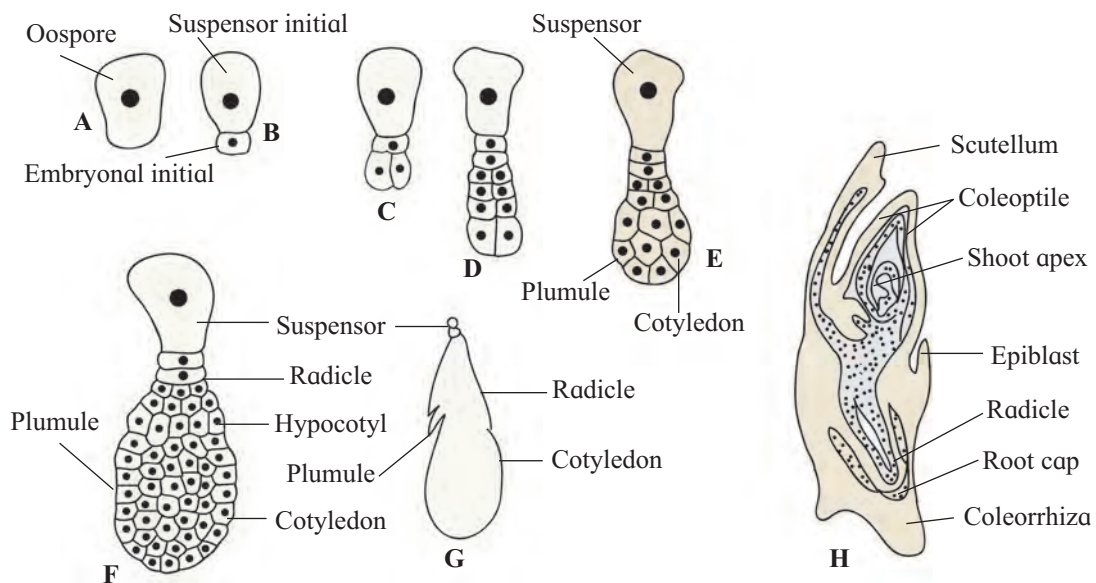


Fig. 1.16 : Development of Monocot (grass) Embryo

upper tier form the plumule and the one or two cotyledons. The hypophysis by further division gives rise to the part of radicle and root cap. Subsequently, the cells in the upper tier of octant divide in several planes so as to become heart shaped which then forms two lateral cotyledons and a terminal plumule. Further enlargement of hypocotyl and cotyledons result in a curvature of embryo and it appears horse-shoe shaped.

The embryo development is similar in both dicots and monocots up to the octant stage. The difference appears later. In monocot embryo, single cotyledon occupies terminal position and plumule is lateral. The single shield shaped cotyledon is called as **scutellum**. The protective sheath of plumule is called **coleoptile** and that of radicle is **coleorrhiza**. **Finally, ovule is transformed into the seed and ovary into the fruit.**

1.12 Seed and Fruit Development :

The goal of reproduction, in every living organisms including plants, is to create offsprings for the next generation. One of the ways that plants can produce offsprings is by forming (making) seeds. The flowers must be pollinated in order to produce seeds and fruit. Seed development is initiated by fertilization. The integuments of the fertilized ovule persist and get transformed into the seed coat of mature seed.

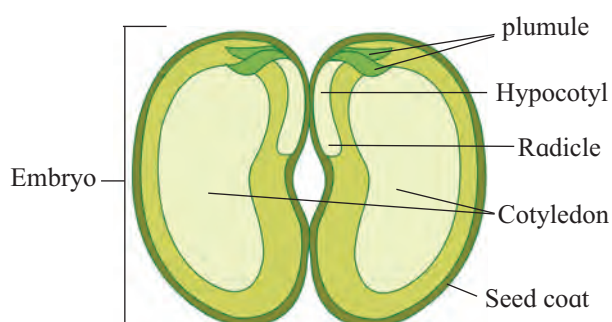


Fig. 1.17 : Bean seed (Dicot)

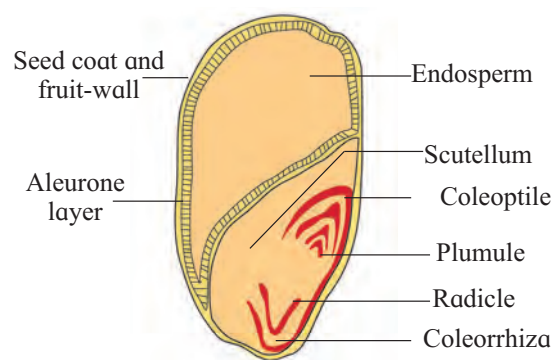


Fig. 1.18 : V. S. Maize grain (Monocot)

Seed sometimes consists of two distinct coverings, a typical outer seed coat, the **testa** and the inner thin, membranous **tegmen**.

The nucellus in the ovule may persist in some genera like black pepper and beet as a thin, papery layer, the **perisperm**. In some seeds, the food reserves in the endosperm are partially used up in the development of an embryo. Obviously, in such seeds the endosperm remains conspicuous and fills a greater part of the seed. Thus, the resultant seed is **endospermic** or **albuminous** e.g. Castor, Coconut, Maize, etc.

In other seeds, embryo absorbs food reserve from the endosperm completely during its developmental stages. Thus, endosperm disappears (disorganizes) in mature seeds. The resultant seed is **non-endospermic** or **ex-albuminous** e.g. Pea, bean, etc.

The cotyledons in some non-endospermic seeds act as a food storage and in others they are the first photosynthetic organs. Micropyle persists as a small pore in seed coat to allow the entry of water and oxygen during soaking.

Fruit development is triggered by hormones produced by developing seeds. As mentioned earlier, after fertilization the zygote is formed and the ovary begins to differentiate into the fruit and ovary wall develops into **pericarp**. Pericarp is basically three layered which get differentiated in the fleshy fruit like mango, coconut, etc.



Can you recall?

1. What are the parts of the fruit ?
2. What is the difference between true fruit and false fruit ?

Significance of seed and fruit formation :

- Fruits provide nourishment to the developing seeds.
- Fruits protect the seeds in immature condition.
- Seeds serve as important propagating organs (units) of plant.
- Seeds and fruits develop special devices for their dispersal and thus help in the distribution of the species.



Try This

Help to rebuild natural ecosystem. Mix seeds and potting soil together with dry clay. Mould the mixture with water into small balls and allow them to dry in sun. Throw the same at places suitable for germination.

Dormancy is a temporary state of metabolic arrest that facilitates the survival of organisms during adverse environmental conditions. Structural or physiological adaptive mechanism for survival is called **dormancy**. Mature and viable seeds will not germinate even in the presence of favourable conditions and they are dispersed at different places during dormancy. Viable seeds germinate only after completion of dormancy period.



Think about it

1. How long seeds stay viable/ healthy?
2. Can old seeds still grow?

Some examples of oldest mature seeds that have grown into viable plants are as follows :

- *Lupinus arcticus* - 10,000 years
- *Phoenix dactylifera* - 2000 years
- Some seeds are short lived, e.g. *Citrus*.
- Some tiny seeds are easy for dispersal. e.g. *Striga*, Orchids, *Orobancha*.

1.13 Apomixis :

It is phenomenon of formation of embryo(s) through asexual method of reproduction without formation of gametes and the act of fertilization. Alternatively, it is unusual sexual reproduction where there is no meiosis and syngamy. Embryo develops in the ovule and ovule develops to form seed .

In apomixis, when a gametophyte organ or cell produces embryo like structure without fertilization, it is termed as **apogamy**. Similarly when diploid sporophyte cell produces a diploid gametophyte without undergoing meiosis is called **apospory**, e.g. Orange, Mango.



Internet my friend

Collect information about seed mother Rahibai's story. How does she save over 80 varieties of native seeds?

The main categories of apomixis are:

a. Recurrent apomixis :

In this type, the embryo sac generally rise either from an archesporial cell or from some other part of the nucellus. In **diplospory**, the unreduced embryo sac is derived from the diploid megaspore mother cell e.g. *Taraxacum*. In apospory, the nucellar cells give rise to apomictic embryo sac.

b. Non-recurrent apomixis :

In this type, megaspore mother cell undergoes usual meiotic division and a haploid embryo sac is formed. Here, the embryo arises either from the egg by parthenogenesis or from some other haploid cells of gametophyte through apogamy. Plants produced by this method are generally sterile and do not reproduce sexually, e.g. *Nicotiana*.

c. Adventive Embryony :

In this type, embryos may develop from somatic nucellus or integuments along with normal **zygotic embryo**. It is common in Mango, Orange, Lemon, etc. It gives rise to a condition called **polyembryony**.

Genetically identical plants can be produced effectively and rapidly by apomixis.

1.14 Parthenocarpy :

This term is coined by Noll (1902). It is the condition in which fruit is developed without the process of fertilization. It occurs naturally in some varieties of Pineapple, Banana, Papaya, etc. In these plants, it seems that the placental tissue in the unfertilized ovary produces auxin IAA (Indole-3 Acetic Acid) which is responsible for enlargement of ovary into fruit. The fruit resembles the normally produced fruit but it is seedless.



Use your brain power

What do bananas and figs have in common ?

1.15 Polyembryony :

It is the development of more than one embryos, inside the seed and the condition is described as polyembryony. It was first noticed by **Leeuwenhoek** (1719) in the seeds of *Citrus* genus. It is the occurrence of more than one embryo in a seed which consequently results in the emergence of multiple seedlings. The additional embryos result from the differentiation and development of various maternal and zygotic tissues associated with the **ovule of seed**. Polyembryony may be **true** or **false** depending upon whether many embryos arise in the same embryo sac or in different embryo sacs in the same ovule. In **adventive polyembryony**, an embryo develop directly from the diploid cell of nucellus and integuments as in *Citrus*. In **cleavage polyembryony**, zygote proembryo sometimes divides (cleaves) into many parts or units. Each unit then develops into an embryo. Polyembryony increases the chances of survival of the new plants. Nucellar adventive polyembryony is of great significance in horticulture.



Think about it

Why are some seeds of *Citrus* referred to as polyembryonic ?



Do you know ?

1. Parthenogenesis is the development of embryo directly from egg cell or a male gamete. It is a kind of apogamy.
2. Agamospermy : Here plants produce seeds. But embryo, inside it, is produced without (omitting) meiosis and syngamy.
3. Parthenocarpy can be induced artificially by - spraying of gibberellins, delaying pollination, use of foreign pollens, etc.
4. Genetically uniform parental type seedlings are obtained from nucellar embryos.



Activity :

Prepare chart for natural vegetative propagation exhibited by flowering plants indicating the vegetative part/s and the different examples.

Organ	Part	Name of plant

Exercise

Q. 1 Multiple choice questions.

1. Insect pollinated flowers usually possess
 - a. Sticky pollens with rough surface
 - b. Large quantities of pollens
 - c. Dry pollens with smooth surface
 - d. Light coloured pollens
2. In ovule, meiosis occurs in
 - a. Integument
 - b. Nucellus
 - c. Megaspore
 - d. Megaspore mother cell
3. The ploidy level is NOT the same in
 - a. Integuments and nucellus
 - b. Root tip and shoot tip
 - c. Secondary nucleus and endosperm
 - d. Antipodals and synergids
4. Which of the following types require pollinator but result is genetically similar to autogamy?
 - a. Geitonogamy
 - b. Xenogamy
 - c. Apogamy
 - d. Cleistogamy
5. If diploid chromosome number in a flowering plant is 12, then which one of the following will have 6 chromosomes?
 - a. Endosperm
 - b. Leaf cells
 - c. Cotyledons
 - d. Synergids
6. In angiosperms, endosperm is formed by/ due to
 - a. Free nuclear divisions of megaspore
 - b. polar nuclei
 - c. polar nuclei and male gamete
 - d. synergids and male gamete
7. Point out the odd one
 - a. Nucellus
 - b. Embryo sac
 - c. Micropyle
 - d. Pollen grain

Q. 2 Very short answer type questions :

1. Name the part of gynoecium that determines the compatible nature of pollen grain.
2. How many haploid cells are present in a mature embryo sac ?
3. Even though each pollen grain has 2 male gametes, why at least 20 pollen grains are required to fertilize 20 ovules in a particular carpel?
4. Define megasporogenesis.
5. What is hydrophily ?
6. Name the layer which supplies nourishment to the developing pollen grains.
7. Define parthenocarpy.
8. Are pollination and fertilization necessary in apomixis ?
9. Name the parts of pistil which develop into fruits and seeds.
10. What is the function of filiform apparatus ?

Q. 3 Short Answer Questions :

1. How polyembryony can be commercially exploited ?
2. Pollination and seeds formation are very crucial for the fruit formation. Justify the statement.
3. Incompatibility is a natural barrier in the fusion of gametes. How will you explain this statement?
4. Describe three devices by which cross pollination is encouraged in Angiosperms by avoiding self pollination.

Q. 4 Long Answer Questions :

1. Describe the process of double fertilization.
2. Explain the stages involved in the maturation of microspore into male gametophyte.

3. Explain the development of dicot embryo.

4. Draw a labelled diagram of the L.S. of anatropous ovule and list the components of embryo sac and mention their fate after fertilization.

Q. 5 Fill in the blanks:

The collects the pollen grains.

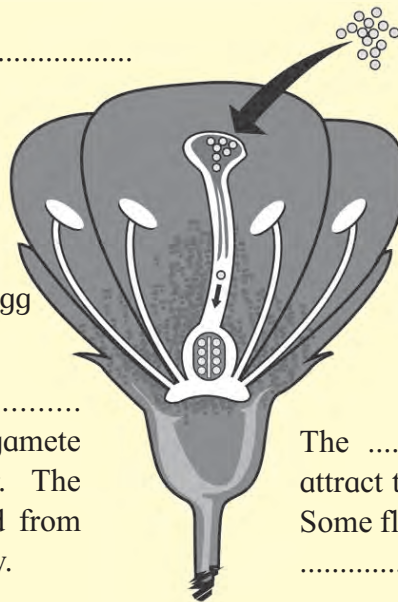
The male whorl, called the produces

The pollen grains represent the

The contains the egg or ovum.

..... takes place when one male gamete and the egg fuse together. The fertilised egg grows into seed from which the new plants can grow.

The is the base of the flower to which other floral parts are attached.



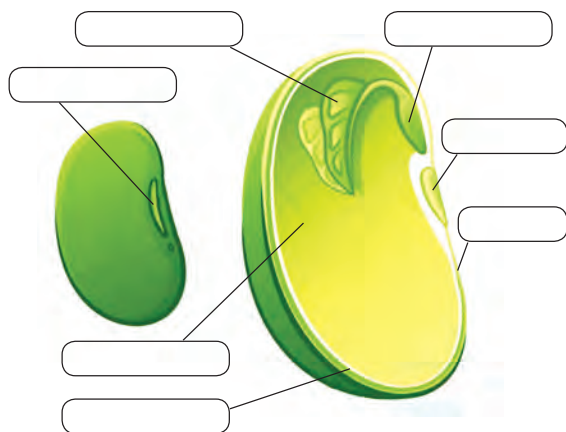
..... is the transfer of pollen grains from anther of the flower to the stigma of the same or a different flower

Once the pollen reaches the stigma, pollen tube traverses down the to the ovary where fertilisation occurs.

The are coloured to attract the insects that carry the pollen. Some flowers also produce or that attracts insects.

The whorl is green that protects the flower until it opens.

Q. 6 Label the parts of seed.



Q. 7 Match the column.

Column - I (Structure before seed formation.)		Column - II (Structure after seed formation.)	
A.	Funiculus	I.	Hilum
B.	Scar of Ovule	II.	Tegmen
C.	Zygote	III.	Testa
D.	Inner integument	IV.	Stalk of seed
		V.	Embryo

- a. A - V, B - I, C - II, D - IV
- b. A - III, B - IV, C - I, D - V
- c. A - IV, B - I, C - V, D - II
- d. A - IV, B - V, C - III, D - II

Project :

Natural vegetative propagation by leaves only in different vascular plants.



Can you recall?

1. Enlist the various life processes. Name the life process which is responsible for continuation of the human race.
2. What are the common methods of reproduction in the unicellular organisms like *Euglena*, *Amoeba* and *Paramecium*?
3. What type of asexual reproduction occurs in *Hydra*?
4. What are the different methods of reproduction in animals?

We know that reproduction is one of the major life processes of any living organism. It helps in maintaining the continuity of the species. **Reproduction** is defined as the biological process of formation of new life forms from pre-existing similar life. It thus becomes a vital process which enables the species to survive over a long period, even though the individuals or organisms live naturally for a limited period of time i.e. their life span. In this chapter, we will learn about the various methods of reproduction in animals the human reproductive system, gametogenesis and fertilization, early embryology, parturition and reproductive health.

Reproduction in animals occurs mainly by two methods i.e. asexual and sexual.

2.1 Asexual Reproduction in animals :

It is a common method among lower animals. It does not involve meiosis nor the gamete formation and fusion. The formation of progeny is by a single parent only and does not involve both the sexes, so it is called asexual reproduction. The progeny or daughter cells are genetically identical to the single parent and are also referred to as **clones**. The lower animals reproduce asexually by gemmule formation and budding.

Gemmule Formation:

Gemmule is an internal bud formed only in sponges. It has asexually produced mass or aggregation of dormant cells, the archaeocytes capable of developing into a new organism. The archaeocytes get coated by a thick resistant layer of secretion by amoebocytes. The gemmule is formed to overcome unfavourable conditions. On return of favourable conditions of water and temperature, the gemmules hatch and develop into a new individual. e.g. *Spongilla*.

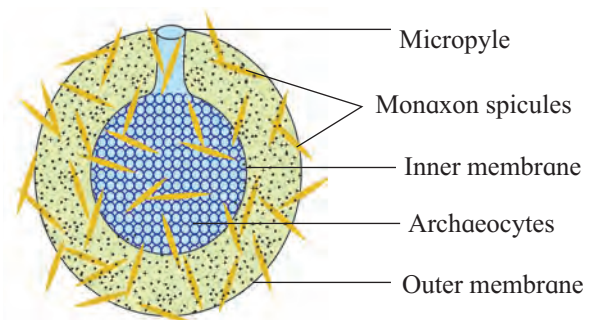


Fig. 2.1 : Gemmule

Budding:

It is a simple method of asexual reproduction normally occurring in favourable conditions. It is seen in a variety of animals like coelenterates (*Hydra* and corals) and in some colonial ascidians. In *Hydra*, a small outgrowth (bud) is produced towards the basal end of the body.

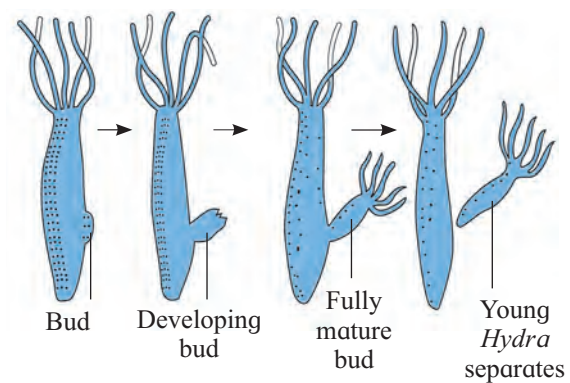
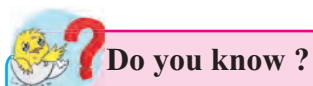


Fig. 2.2 : Budding in *Hydra*

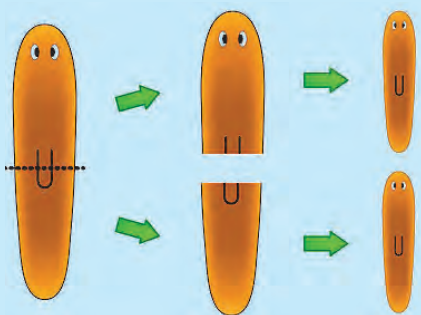
It develops as a bud which grows and forms tentacles and develops (get transformed) into a new individual. This process is called **budding**. The young *Hydra* gets detached from the parent and becomes an independent new organism.



Do you know ?

Regeneration:

A word which in biology refers to the process observed in all living organisms from the unicellular bacteria upto the most complex multicellular forms e.g. humans. By this process, the organism can fundamentally repair or regrow or restore its lost or damaged part. Though it involves asexual processes, it differs distinctly from reproduction e.g. a damaged *Hydra* can regenerate its lost part. Similarly *Planaria* if wounded, its cells become active and regenerate lost part or organ back to its original state. They can also reproduce asexually by fragmentation. Also, it is seen in planarians that the anterior end exerts a pull on the posterior end resulting in a constriction in the middle part and splitting into two pieces. Each piece grows into a new *Planaria*. i.e. two clones of the original have been formed.



2.2 Sexual reproduction in animals :

It is the process which involves the production of offspring by the formation and fusion of gametes. It is also called **amphimixis**. In animals, gamete formation primarily involves meiosis.

The sexually reproducing animals show two main phases in their life time. The earlier **juvenile phase** mainly represents physical growth phase starting from birth. The animals can not reproduce sexually in this phase. The later **Reproductive maturity phase** is attained usually after physical growth is almost over. It involves growth and activity of the sex organs. The animal can reproduce sexually in this phase. Both these periods (phases) are of variable duration in different animals. After attaining sexual maturity, the animal exhibits various events, namely pre-fertilization (gametogenesis and gamete transfer), fertilization (fusion of male and female gametes) and post fertilization events (formation of zygote and embryogenesis).

The sexually reproducing animals show various breeding patterns. Some like the goat, sheep, and donkey are **seasonal breeders** while humans and apes are **continuous breeders**. They can breed throughout the year.

Human Reproduction :

Humans are sexually reproducing animals. The process of reproduction involves various sequential steps such as gametogenesis, insemination, internal fertilization (i.e. fusion of male and female gametes), zygote formation and embryogenesis, gestation and parturition.

The gametes, sperms and eggs are produced by the primary sex organs, testis in male and ovary in female. Organs other than testis and ovary, are called secondary sex organs of the male and female. As male and female can be externally differentiated by certain specific features called **secondary sexual characters**, they are called **sexual dimorphic characters**. In males, presence of beard, moustache, hair on the chest, muscular body, enlarged larynx (Adam's apple) are secondary sexual characters while in females these characters are the developed breast, broader pelvis and high pitched voice.

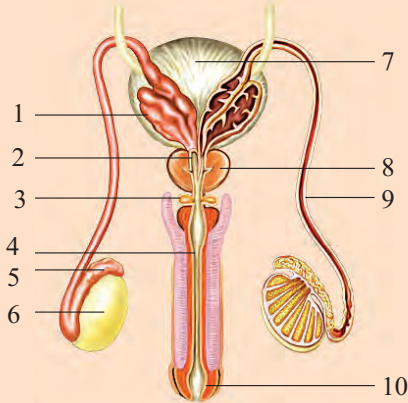
A. Male Reproductive System :

It consists of the primary male organ (gonad) called testes, the accessory ducts and glands which form internal and external genitalia.



Can you recall?

Label the given male reproductive system you have studied.



a. Testes:

A pair of testes, mesodermal in origin, are formed in the lower abdominal cavity. They are located in a pouch called **scrotum**. During early foetal life, the testes develop in abdominal cavity and later they descend into the scrotal sac through a passage called **inguinal canal**. Each testis is oval in shape, 4 to 5cm long, 2 to 3cm wide and 3cm thick.

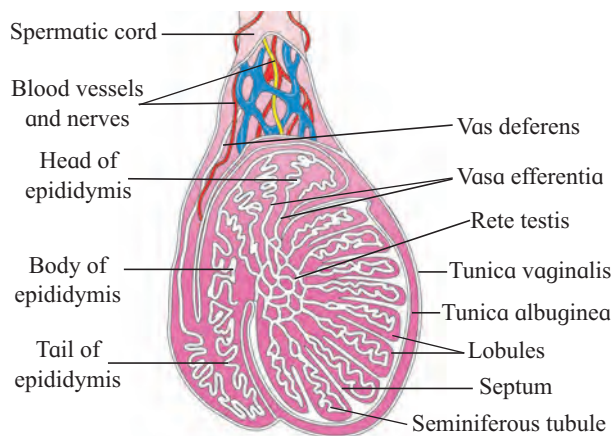


Fig. 2.3 : L. S. of testis

Histology of Testis:

The testis is externally covered by a collagenous connective tissue layer called **tunica albuginea**. Outer to it is an incomplete peritoneal covering called **tunica vaginalis**, and inner to it is **tunica vasculosa**, a thin membranous and vascular layer. Fibers from tunica albuginea divide each testis into about 200-300 testicular **lobules** (refer dig. 2.3 L. S. of testis). Each with 1-4 highly coiled seminiferous tubules. Each **seminiferous tubule** is internally lined by cuboidal germinal epithelial cells (spermatogonia) and few large pyramidal cells called **Sertoli** or **sustentacular cells**.

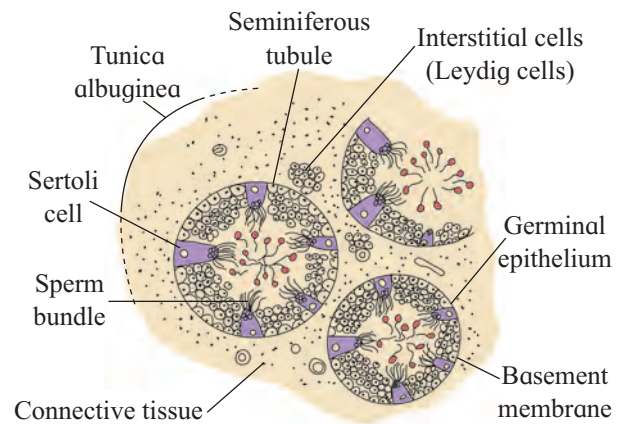


Fig. 2.4 : T. S. of Testis

The germinal epithelial cells undergo gametogenesis to form the **spermatozoa**. Sertoli cells provide nutrition to the developing sperms. Various stages of spermatogenesis can be seen in the seminiferous tubules. The inner most spermatogonial cell (2n), primary spermatocyte (2n), secondary spermatocyte (n), spermatids (n) and sperms (n). The **Interstitial** or **Leydig's cells** lie in between the seminiferous tubules. They secrete the male hormone **androgen** or **testosterone**.



Do you know ?

1. Presence of the peritoneal covering around the testis is an indication of its abdominal origin.
2. The testis are suspended in the scrotum by the spermatic cord.
3. Testosterone hormone stimulates the descent of testis and the fibro-muscular band called **gubernaculum** in the scrotum.
4. In some males a loop of the intestine may pass through the inguinal canal into the scrotum and cause a condition called **inguinal hernia**.

b. Accessory ducts:

The accessory ducts include rete testis, vasa efferentia, epididymis, vas deferens, ejaculatory duct and urethra. All the seminiferous tubules of the testis at the posterior surface form a network of tubules called **rete testis**. 12-20 fine tubules arising from rete testis are **vasa efferentia**. They carry the sperms from the testis and open into the **epididymis**. It is a long and highly coiled tube which is differentiated into an upper caput-, middle corpus- and lower cauda epididymis. The sperms undergo maturation in epididymis. Posteriorly, it leads into the **vas deferens** which travels upto the abdominal cavity and loops over the ureter to open into the urethra. Before doing so, it joins the duct of seminal vesicle to form the **ejaculatory duct**. The ejaculatory duct passes through the **prostate gland** and opens into the urethra. The **urethra** provides a common passage for the urine and semen and hence is also called **urinogenital duct**. In males the urethra is long and extends through the penis. It opens to the outside by an opening called the urethral meatus or **urethral orifice**. All the accessory ducts except urethra are present in pairs.

c. Glands:

The male accessory glands are as follows:

- **Seminal vesicles:** It is a pair of glands lying on the posterior side of urinary bladder. It secretes an alkaline **seminal fluid** which contains fructose, fibrinogen and prostaglandins. It contributes about 60% of the total volume of the semen. Fructose provides energy for sperm movement while fibrinogen coagulates the semen into a bolus for quick propulsion in the vagina. The prostaglandins stimulate reverse peristalsis in vagina and uterus aiding faster movement of sperms towards the egg in the female body.
- **Prostate gland:** It is a large and single gland made up of 20-30 lobes and is located underneath the urinary bladder. It surrounds the urethra and releases a milky white and slightly acidic prostatic fluid into the urethra. It forms about 30% of volume of semen. It contains citric acid, acid phosphatase and various other enzymes. The acid phosphatase protects the sperms from the acidic environment of vagina.



Activity :

Find the symptoms of prostate cancer.



Always Remember

Prostate cancer is cancer of the prostate gland. Men who are over 50 years of age and have a daily high consumption of fat, have an increased risk of prostate cancer.



Internet my friend

What is the role of prostaglandin?

- **Cowper's gland / Bulbourethral gland** : It is a small, pea sized and paired gland situated on either side of urethra. These

glands secrete an alkaline, viscous, mucous like fluid which acts as a lubricant during copulation.

Semen :

It is the viscous, alkaline and milky fluid (pH 7.2 to 7.7) ejaculated by the male reproductive system. Normally 2.5 to 4.0 ml of semen is given out during a single ejaculation and it contains about 400 million sperms. It contains secretion of the epididymis and the accessory glands for nourishing (fructose), neutralizing acidity (Ca^{++} , bicarbonates), activation for movement (prostaglandins).

d. External genitalia:

It includes the penis and the scrotum. The **penis** is the male copulatory organ. It is cylindrical and muscular with three bundles of erectile tissue- a pair of postero-lateral tissue called **corpora cavernosa** and a median **corpus spongiosum**. The swollen tip of the penis is called **glans penis**. It is covered by a loose fold of skin called **foreskin** or **prepuce**.

Scrotum :

It is a loose pouch of pigmented skin lying behind the penis and is divided into a right and left scrotal sac by a septum of tunica dartos made of smooth muscle fibres. The foetal testes are guided into and retained in the scrotum by a short fibro muscular band called **gubernaculum**. The testes remain suspended in scrotum by a spermatic chord. Failure of testis to descend into scrotum is called **cryptorchidism**. The failure also results in the sterility. The cremaster and dartos muscles of scrotum help in drawing testes close or away from the body. This helps in maintaining the temperature of the testis $2\text{-}3^{\circ}\text{C}$ lower than the normal body temperature, necessary for spermatogenesis.

B. Female Reproductive System:

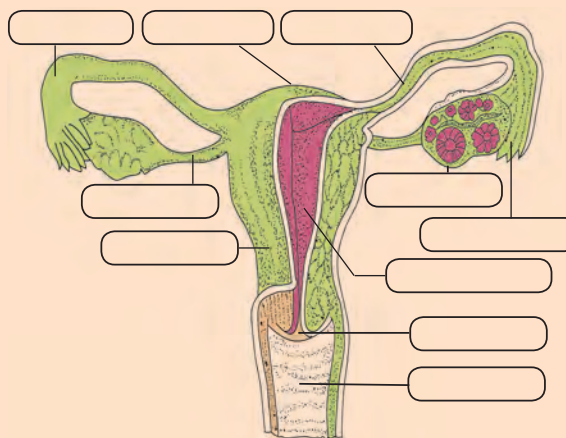
The female reproductive system consist of the following parts :

1. A pair of ovaries
2. A pair of oviducts
3. Uterus
4. Vagina
5. External genitalia (vulva)
6. A pair of vestibular glands
7. A pair of mammary glands



Can you recall?

Give labels to given female reproductive system:



1. Ovary : It is the primary female sex organ. Its main function is production of egg or ovum and the female reproductive hormones. It is solid, oval or almond shaped organ. It is 3.0 cm in length, 1.5 cm in breadth and 1.0 cm thick. It is located in the upper lateral part of the pelvis near the kidneys. Each ovary is held in position by ligaments by attaching it to the uterus and the abdominal wall. The largest of these is the broad ligament formed by a fold of peritoneum. It holds the ovary, oviduct and the uterus to the dorsal body wall. The ovarian ligament attaches ovary to the uterus. The ovary produces five hormones viz, estrogen, progesteron, relaxin, activin and inhibin.

Structure and development of the ovary :

Each ovary is a compact structure differentiated into a central part called medulla and the outer part called cortex. The cortex is covered externally by a layer of germinal

epithelium. The stroma or loose connective tissue of the medulla has blood vessels, lymph vessels, and nerve fibres. The outer cortex is more compact and granular. It shows large number of tiny masses of cells called ovarian follicles. These are collectively formed from the immature ova originating from cells of the dorsal endoderm of the yolk sac. The cells migrate to the gonadal ridge during embryonic development and divide mitotically. Now these cells are called **oogonia**. As the oogonia continue to grow in size they are surrounded by a layer of granulosa cells and form the rudiments of the ovarian follicles. The process of oogenesis starts much before the birth of the female baby and by the end of twelve weeks the ovary is fully formed. It has more than two million **primordial follicles** in it.

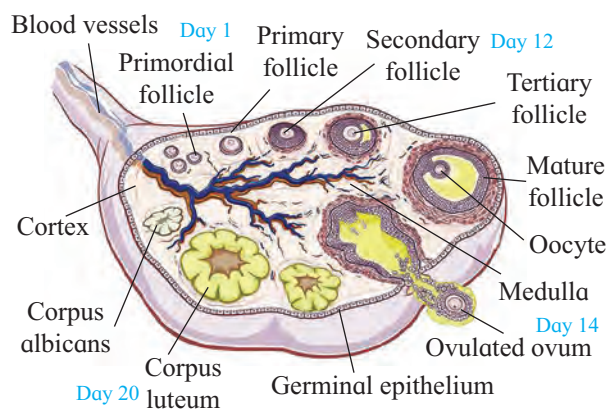


Fig. 2.5 : T.S. of ovary

The cells of **germinal epithelium** give rise to groups of oogonia projecting into the cortex in the form of cords called **egg tubes** of **Pfluger**. Each cord at its end has a round mass of oogonial cells called **egg nests**, from which the primordial ovarian follicles develop. Each primordial follicle has, at its center a large **primary oocyte** (2n) surrounded by a single layer of flat follicular cells. The primary oocyte starts with its meiotic division but gets arrested at meiosis I. Of the two million primordial follicles embedded in the foetal ovary only about one million remain at birth and only about 60,000 - 80,000 remain at the

time of **puberty**. The large scale destruction of the primordial follicles during growth is called **atresia**.

The development of the primordial follicles into mature or Graafian follicles restarts with the onset of puberty. During each menstrual cycle only one of the primordial follicle starts growing to form the **Graafian follicle**.

In each cycle, alternately one of the two ovaries produces the Graafian follicle.

The 1st menstrual cycle or **menarche** begins normally at about 13 years and Menopause i.e. stopping of the cycles happens at age 45 to 55 years. The period in between menarche and menopause is the reproductive age of the female and is approximately 32 years. In this time the female will be producing a maximum of about 416 eggs ($32 \times 13 = 416$ eggs).

Ovarian histology of a mature female :

In the histology of ovary, we have discussed the primary structure of ovary. The following discussion includes the changes seen in a mature ovary, primarily in the cortex. The different stages of development of the oocyte can be seen. These changes in the ovary are cyclic, occurring during each menstrual cycle and it involves maturation of the primordial follicles into primary, secondary and Graafian follicles. Each primary follicle has multilayered cuboidal follicular cells. The stroma cells add **theca** over the follicle. It now changes into a **secondary follicle**. There is growth of the oocyte and the granulosa cells increase in number. They start producing the hormone estrogen. The secondary follicle grows into the **Graafian follicle** by addition of more follicular cells. As this process of maturation of follicles takes place, they begin to move towards the surface of ovary. The Graafian follicle presses against the thin wall of the ovary giving it a blistered appearance. The egg is released from the Graafian follicle during **ovulation** and the remaining part of the follicle changes into a temporary endocrine gland called **corpus**

luteum. If fertilization does not take place the corpus luteum degenerates into a white scar called **corpus albicans**.



Use your brain power

In t. s. of ovary, can all the stages of follicles be seen simultaneously?

Structure of Graafian follicle :

Graafian follicle is a mature ovarian follicle. An eccentric secondary oocyte is surrounded by a non-cellular layer of **zona pellucida** secreted by the vitelline membrane of oocyte. The outermost protective and fibrous covering is called **theca externa**. Inner to it is cellular **theca interna**. It produces the hormone estrogen. Inner to the theca interna, the follicular cells form the **membrana granulosa**. From the membrana granulosa the cells differentiate into **discus proligerus** and the **corona radiata** cells. **Cumulus oophorus** is the term used for the oocyte and surrounding granulosa cells. A fluid filled cavity called **antrum** lies between the oocyte and the **membrana granulosa**. It is filled with a fluid called **liquor folliculi**.

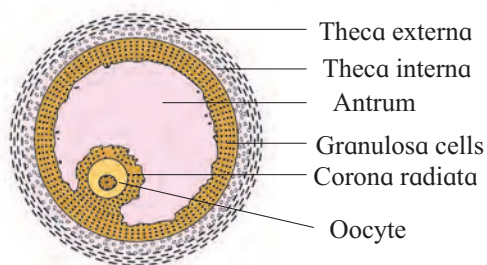


Fig. 2.6 : Graafian Follicle

2. Oviduct / Fallopian tube / Uterine tube:

These are a pair of muscular ducts lying horizontally over the peritoneal cavity. The proximal part of the tube lies close to the ovary, and distally it opens into the uterus. Each tube is 10 to 12 cm in length. It is internally lined by ciliated epithelium. It can be divided into three regions :

- a. **Infundibulum** : The proximal funnel like part with an opening called **ostium** surrounded by many finger like processes called **fimbriae** (of these at least one is long and connected to the ovary). The cilia and the movement of fimbriae help in driving the ovulated egg to the ostium.
 - b. **Ampulla** : It is the middle, long and straight part of the oviduct. Fertilization of the ovum takes place in this region.
 - c. **Isthmus / Cornua** : The distal narrow part of the duct opening into the uterus.
3. **Uterus** : It is commonly also called the womb. It is a hollow, muscular, pear shaped organ, located above and behind the urinary bladder. It is about 7.5 cm long, 5 cm broad and 2.5 cm thick. The uterus can be divided into three regions :
 - a. **Fundus** : It is the upper dome shaped part. Normally implantation of the embryo occurs in the fundus.
 - b. **Body** : It is the broad part of the uterus which gradually tapers downwards.
 - c. **Cervix** : It is the narrow neck about 2.5 cm in length. It extends into the vagina. Its passage has two openings : an **internal os** towards the body, and an **external os** towards the vagina.

Internally the uterine wall can be distinguished into three layers : Outermost **perimetrium**, middle thick muscular **myometrium**, made up of thick layer of smooth muscles. Vigorous contractions of these muscles cause **labour** during the **parturition** (child birth). The innermost layer called **endometrium** or mucosal membrane is made up of stratified epithelium. The thickness of this layer regularly undergoes changes in during the menstrual cycle. It is richly supplied with blood vessels and uterine glands. These provide nourishment to the developing foetus.



Do you know ?

Uterus cancer:

Most of the uterine cancers begin in the layer of cells that form the lining of endometrium of uterus.

Symptoms : Abnormal bleeding between periods, vaginal bleeding after menopause, an abnormal watery, blood-tinged discharge from vagina, pelvic pain.

Detection : It is diagnosed with Pap smear test, biopsy, Ultrasound.

Treatment : Chemotherapy, radiation, surgical removal of uterus (hysterectomy).

4. Vagina : It is a tubular, female copulatory organ, 7 to 9 cm in length. It lies between the cervix and the vestibule. The vaginal wall has an inner mucosal lining, the middle muscular layer and an outer adventitia layer. The mucosal epithelium is stratified and non-keratinised and stores glycogen. There are no glands but the cervical secretion of mucus is received in the vagina. The opening of the vagina into the vestibule is called **vaginal orifice**. This opening is covered partially by a fold of mucus membrane called **hymen**. The vagina acts as a passage for menstrual flow as well as birth canal during parturition.

5. External genitalia : The external genital organs of female include parts external to the vagina and are collectively called '**vulva**' (covering or wrapping), or **pudendum**. They include the following parts :

- Vestibule - It is a median vertical depression of vulva enclosing the urethral and vaginal opening.
- Labia minora - These are another pair of thin folds inner to the labia majora with which they merge posteriorly to form the fore chette while towards anterior end they converge into a hood-like covering around the clitoris.

- Clitoris - A small conical and sensitive projection lying at the anterior end of labia minora. It has a pair of erectile tissue - The corpora cavernosa and is homologous to the penis.
- Labia majora - These are a pair a fleshy folds of skin forming the boundary of vulva. They are homologous to the scrotum. They surround and protect the other parts of external genitalia and enclose the urethral and vaginal openings in the vestibule.
- Mons pubis - It is a fleshy elevation above the labia majora. The Mons pubis and outer part of labia majora show pubic hair.

6. Accessory glands / Vestibular glands / Bartholin's glands : It is a pair of glands homologous to the Bulbourethral or Cowper's glands of the male. They open into the vestibule and release a lubricating fluid.

Mammary glands :

Accessory organs of female reproductive system for production and release of milk after parturition. Development of the mammary gland occurs at puberty under the influence of estrogen and progesteron. Lactotropic hormone (LTH) or prolactin helps in development of lactiferous tubules during pregnancy.

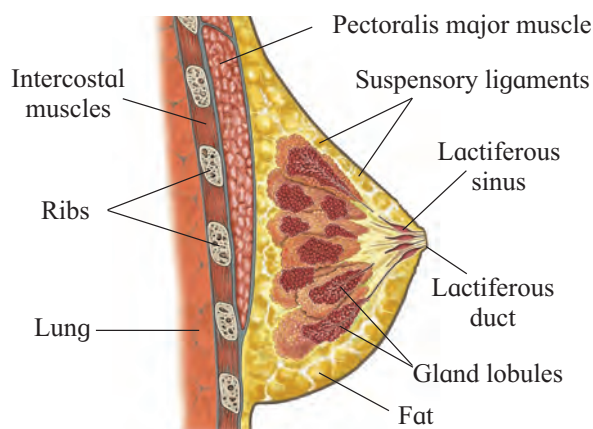


Fig. 2.7 : Section view of Mammary gland

The mammary glands are a pair of rounded structures present in the subcutaneous tissue of the anterior thorax in the pectoral region (from

2nd to 6th rib). These are modified sweat glands. Each mammary gland contains fatty connective tissue and numerous lactiferous ducts. The glandular tissue of each breast is divided into 15-20 irregularly shaped mammary lobes, each with alveolar glands and lactiferous duct. Alveolar glands secrete milk which is stored in the lumen of alveoli. The alveoli open into mammary tubules. The tubules of each lobe join to form a mammary duct. Many mammary ducts join to form a wider mammary ampulla, which is connected to lactiferous duct. These converge towards the nipple located near the tip of the breast. It is surrounded by a dark brown coloured and circular area of the skin called **areola**.



Do you know ?

Breast cancer :

Symptoms: First symptom of breast cancer is a lump in breast or underarm. Lump is painless. Swelling of all or part of breast. Skin irritation, Breast or nipple pain, nipple retraction, Redness, scaliness or thickening of nipple or breast skin, discharge, etc.

Detection: Mammogram (x-ray), ultrasound, MRI, Biopsy, Blood test.

Treatment: Radiation therapy, chemotherapy lumpectomy, Mammoplasty



Do you know ?

Weaning : Mother's milk is replaced gradually by solid food after some time. This process is called weaning.

Puberty / Sexual maturity in Males :

Puberty is the age at which the reproductive system becomes functional, sex organs begin to produce gametes and sex hormones. In males the onset of puberty occurs at age 12-15 years. Under the influence of testosterone, the secondary sexual characters appear. Thereafter it remains functional throughout the life.

Puberty / Sexual maturity in Females :

The reproductive system becomes functional at puberty. It is characterised by onset of menstrual cycle also called **menarche**, which usually occurs at age 10-14 years. However, unlike in the males, the mature females show cyclic changes in their reproductive system- the menstrual cycles. These cycles continue only upto **menopause**. This normally happens at age 45-50 years. The period from menarche to menopause is thus the reproductive age of the female. The female is unable to bear children (by natural method) after menopause. Menarche, menstrual cycles and menopause are controlled by gonadotropic hormones.

2.3 Menstrual cycle (Ovarian cycle):

Menstrual cycle is the characteristic feature of primates including human. It involves a series of cyclic changes in the ovary and the female reproductive tract, mainly in the uterus. These changes take place under the effect of gonadotropins and the ovarian hormones respectively. The cycles are repeated with a periodicity of approximately 28 days. The middle of each cycle is characterised by the release of an egg. This egg in every cycle comes alternately from one of the two ovaries. The cycle is divided into four phases.

a. Menstrual phase:

The beginning of each cycle is taken as the first day where menses or loss of blood (45-100ml) takes place and it lasts for approximately five days (average 3-7 days).

Endometrium of uterus breaks down under the effect of prostaglandins released due to decreased levels of progesterone and estrogen. Due to this blood, tissue fluid, mucus, endometrial lining and the unfertilized oocyte is discharged through vagina. The endometrial lining becomes very thin i.e. about 1 mm. The menstrual discharge continues for an average of 5 days, however this blood does not clot

due to presence of fibrinolysin. Menstrual phase occurs when an ovulated egg does not get fertilized and it is thereby shed out along with the menstium. It is thus called ‘funeral of unfertilized egg’.

During these five days, many primordial follicles develop into primary and few of them into secondary follicles under the effect of FSH.

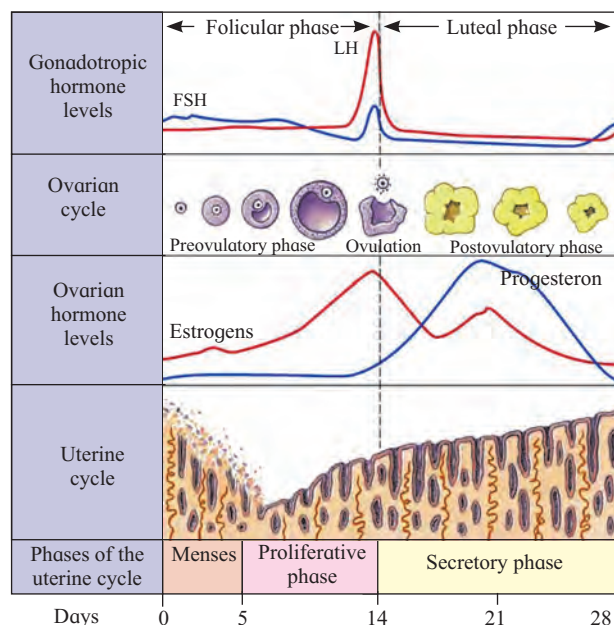


Fig. 2.8 : Hormones and the menstrual cycle.



Internet my friend

1. Enlist the examples of primates and non primate animals.
2. Collect information about female reproductive cycles differentiating both primates and non-primates.

b. Proliferative phase / Follicular phase / Post menstrual phase:

This phase is the duration between the end of menstruation and release of ovum (ovulation). Duration of this phase is more variable than other phases. Generally, it extends from 5th to 13th day of menstrual cycle.

A few (6 to 12) secondary follicles proceed to develop but usually one of them develops into a graafian follicle (mature follicle). The other secondary follicles degenerate. This process

of degeneration is called **atresia**. Developing secondary follicles secrete the hormone estrogen. The stimulation for proliferation of new follicles is influenced by GnRH which stimulates release of FSH.

Endometrium begins to regenerate under the effect of gradually increasing quantity of estrogens. Regeneration also involves formation of endothelial cells, endometrial or uterine glands and network of blood vessels. Thickness of endometrium reaches 3-5 mm.

c. Ovulatory phase:

It is the shortest phase of menstrual cycle. It involves rupturing of the mature graafian follicle and release of ovum (secondary oocyte) into the pelvic cavity; usually on 14th day of menstrual cycle. Rapid secretion of LH by positive feedback mechanism causes the mature follicle to rupture. Ovulation may be accompanied by mild or severe pains in lower abdomen.

d. Secretory phase / Luteal phase:

Duration of this phase is between the ovulation and beginning of the next menses. This phase is the longest phase. It lasts for 14 days; from 15th to 28th day of the cycle.

After release of secondary oocyte, remaining tissue of graafian follicle transforms into corpus luteum under the effect of LH. Corpus luteum begins to secrete progesteron and estrogens. The ovulated egg may get fertilized within 24 hours. However, in the absence of fertilization, corpus luteum can survive for only two weeks and then degenerate into a white scar called **corpus albicans**.

The corpus luteum releases progesteron, small amount of estrogens and inhibin. Under the influence of these hormones, the endometrial glands grow, become coiled and start uterine secretions. Endometrium becomes more vascularized and thickens up to 8-10 mm. Inhibin stops secretion of FSH. These changes are necessary for fertilization and subsequent implantation.

However, if the ovulated egg gets fertilized and the embryo is implanted, there is secretion of human chorionic gonadotropin (hCG), which extends the life of corpus luteum and stimulates its secretory activity. Presence of hCG in maternal blood and urine is an indicator of pregnancy. In absence of fertilization, next menstrual cycle begins.



Always Remember

Hygiene practices during menstruation :

- Keep the pubic area clean.
- Change the sanitary napkin every 4-5 hours.
- Maintaining personal hygiene during menstruation is important to reduce the risk of infection.
- Dispose used sanitary napkin properly.
- Using damp and dirty clothes or using a sanitary napkin for a longer time can act as a perfect environment for growth and multiplication of harmful bacteria and lead to infections.



Use your brain power

Why the menstruation is painful in some women?



Can you tell?

Can you tell the names of primates who show the presence of menstrual cycle?

2.4 Gametogenesis:

The gametogenesis is the process of formation of gametes in sexually reproducing animals. The male gamete is sperm and the female gamete is ovum or egg. The gametes are formed from primordial germ cells of gonads.

Spermatogenesis:

The process of formation of the male gamete (sperm) or spermatozoa from the germinal epithelium of testis is called

spermatogenesis. At the onset of puberty, the hypothalamus begins secretion of gonadotropin releasing hormone (GnRH). It initiates the significant increase in the secretion of follicle stimulating hormone (FSH) which induces spermatogenesis. Each seminiferous tubules is lined by a single layer of cuboidal epithelial cells called germinal epithelium.

The cells of germinal epithelium undergo spermatogenesis to produce sperms. Process of spermatogenesis involves three phases.

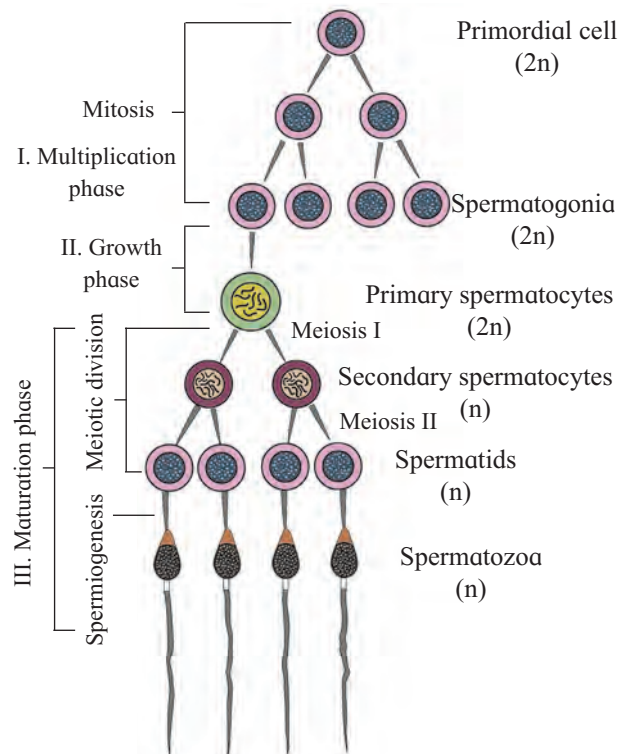


Fig. 2.9 : Spermatogenesis

I. Multiplication phase: The primordial germ cells (2n) of seminiferous tubules undergo repeated mitotic divisions to produce large number of spermatogonia (2n). Each spermatogonium is diploid and with 46 chromosomes.

II. Growth phase: Some of the spermatogonia stop dividing and grow in size to develop into primary spermatocytes (2n) due to accumulation of food.

III. Maturation phase: It involves meiotic or reduction division. The spermatocyte undergoes the first phase of meiotic division (meiosis I) leading to formation of two haploid cells called secondary spermatocytes (n), which are with 23 chromosomes each. The secondary spermatocyte undergoes second phase of meiotic division (meiosis II) to produce four haploid spermatids. The spermatid is non-motile and non-functional. It gets transformed into a functional spermatozoa by the process called **spermiogenesis**. During this process of change, the spermatids remain held to each other and to the sertoli cells by cytoplasmic bridges. The sperm heads remain attached to the sertoli cells and their tails hanging in the lumen of seminiferous tubule. During spermiogenesis, length of spermatid increases. Centrioles are rearranged as primary and distal centrioles. Mitochondria become spirally coiled and acrosome is formed from golgi complex.

Structure of sperm:

Sperm is the male gamete. It is a motile, microscopic elongated cell. It is divisible into three parts- head, middle piece and tail.

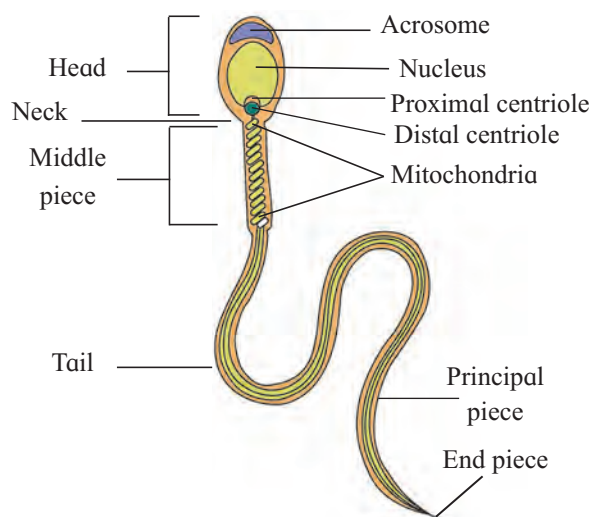


Fig. 2.10 : Structure of Sperm

Head: The sperm head is oval in shape and contains haploid nucleus. Above the nucleus, there is a cap like structure called **acrosome**. It is formed from the golgi body. Acrosome

contains hydrolytic enzymes; hyaluronidase and proteolytic enzymes like zona lysins and corona penetrating enzymes.

Neck: It is a very short region having two centrioles i.e. proximal centriole and distal centriole.

Middle piece: It has an axial filament surrounded by 10-14 spiral turns of mitochondria (nebenkern). It produces energy necessary for the movement of sperm.

Tail: It is a long, slender and tapering part containing cytoplasm and fine thread- axial filament. The axial filament arises from the distal centriole and travels through out the length of tail. It is partly surrounded by plasma membrane (main piece). The part without plasma membrane is called end piece.

Oogenesis:

It is process of formation of the haploid female gamete i.e. egg or ovum from the diploid germinal epithelium. It involves the process of meiosis (and mitosis). Like spermatogenesis, oogenesis process can be divided into three stages :

- I. Multiplication phase
- II. Growth phase
- III. Maturation phase

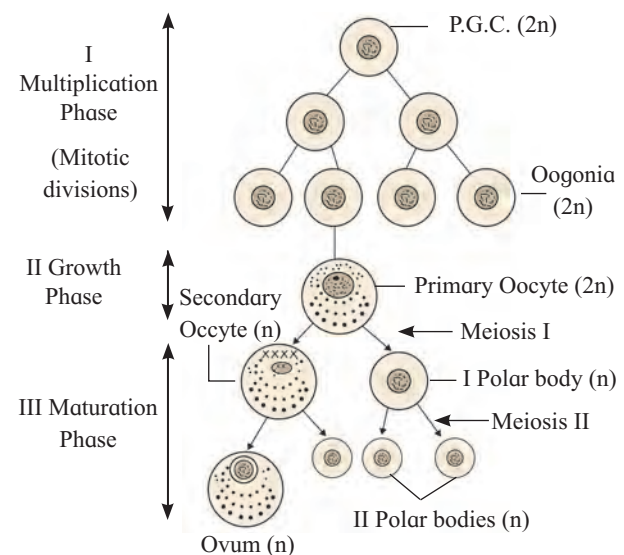


Fig. 2.11 : Oogenesis



Activity :

Prepare a chart of comparison between spermatogenesis and oogenesis.

I. Multiplication phase: In this stage, the primary germinal cells PGCs ($2n$) of ovary undergo repeated mitotic division to form millions of gamete mother cells or oogonial cells ($2n$). This process is completed in the embryonic stage of human females.

II. Growth phase: Some of the oogonia stop division and begin to increase in size and form the primary oocytes ($2n$). Cellular organelles like ER, golgi apparatus and mitochondria increase in number.

III. Maturation phase: Oogenesis takes place in the ovaries. The process is initiated prior to birth of the female baby. The primary oocytes ($2n$) enter the maturation phase which includes meiotic division (Meiosis I and Meiosis II). The diploid primary oocytes undergo meiosis I (reduction division) to form 2 haploid daughter cells. This division is peculiar in females as both the daughter cells are with haploid number of chromosomes i.e. 23 chromosomes. But due to unequal division of cytoplasm, of the 2 daughter cells produced, one is a large cell called secondary oocyte (n) and another is a small cell called 1st polar body (n). Normally the 1st polar body does not enter meiosis II. The secondary oocyte (n) proceeds meiosis II, only upto metaphase II. Its division is further stopped or arrested at this stage. The secondary oocyte is shed from the graafian follicle and ovary. The restart and completion of meiosis II will happen only with entry of the sperm. This last phase is usually completed in the ampulla of the fallopian tube at the time of fertilization. In this division also, the two unequal daughter cells are formed- the large cell is ovum (n) and the small cell is 2nd polar body (n). The ovum (n) so formed functions as the female gamete

and is ready for fertilization. (Completion of meiosis II and completion of fertilization go hand in hand. If the secondary oocyte does not receive the sperm / spermatozoa, it is shed off along with menstrum).

Structure of secondary oocyte:

In human, unfertilized egg when ovulated i.e. released from the ovary is actually the secondary oocyte. It is non-cleidoic (without shell) and microlecithal (yolk is present in very small quantity). It is approximately 0.1mm (100 microns) in size. It is rounded, nonmotile and haploid female gamete. The nucleus of the egg appears large and is called **germinal vesicle**. Typical nucleus or pronucleus is formed at the time of fertilization. The cytoplasm of egg is also called **ooplasm**. It is devoid of centrioles. The egg is surrounded by various coverings.

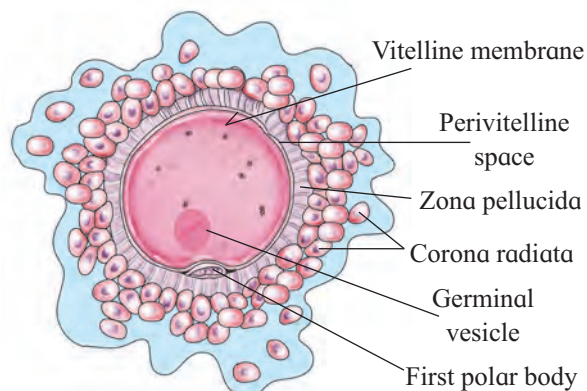


Fig 2.12 : Unfertilized egg/ Ovum

The egg membrane is called vitelline membrane. It secretes a non-cellular glycoproteinous membrane, **zona pellucida** on its outside. Adhering to the outer surface of zona pellucida are several radially elongated cells forming the **corona radiata**. These cells are derived from the innermost layer of granulosa cells. They are firmly held to the zona pellucida and to each other by hyaluronic acid (mucopolysaccharide). Between the vitelline membrane and the zona pellucida is a fluid filled perivitelline space. The first polar body lies in this space.

The egg shows polarity. The side having germinal vesicle and first polar body is called **animal pole** while the side opposite to it is called **vegetal pole**.

2.5 Fertilization / Syngamy:

Sexual reproduction primarily involves formation and fusion of gametes. Fertilization is the later process which involves fusion of the haploid male and female gametes resulting in the formation of a diploid zygote (2n). Like in other mammals, in humans the process of fertilization is internal and it usually takes place in the ampulla of the fallopian / uterine tube. The fertilized egg or zygote will develop into an embryo and this process occurs within the uterus.

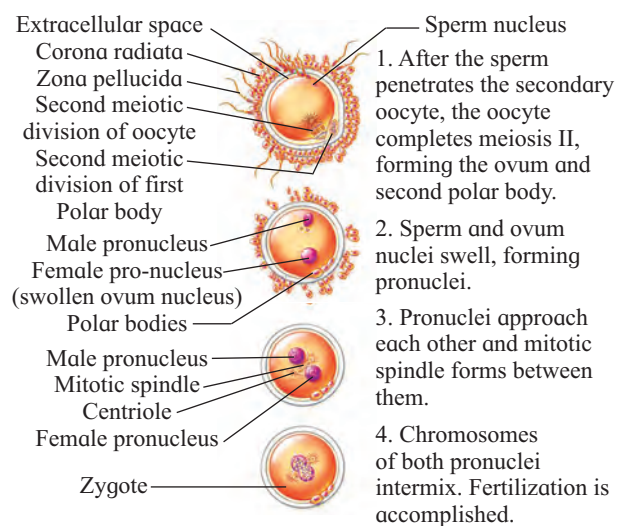


Fig. 2.13 : Process of Fertilization

Mechanism of fertilization :

Semen released during ejaculation has sperms and some secretions. The coagulated semen now undergoes liquification and sperms become active. The mechanism of fertilization is as follows :

a. Movement of sperm towards egg:

It involves capacitation of sperms reaching the vagina. Here as many as 50% are demotilised / broken / destroyed. Remaining sperms undergo capacitation. This process requires 5-6 hours. Acrosome membrane

becomes thin, Ca^{++} enters the sperm and their tails begin to show rapid whiplash movements.



Always Remember

Indian law under the Hindu marriage Act has defined the marriageable age of a boy and girl. As per this act, minimum age for boy must be 21 and for a girl must be 18 years, at the time of marriage.

As a result of capacitation, sperms become extra active and begin to start moving upwards from vagina to uterus and to the oviducts. The prostaglandins activate the sperms. The vestibular secretions of the female also enhance sperms motility. The sperms swim at an average speed of 1.5 to 3.0 mm/min.

Sperms reach upto the ampulla as a result of their own swimming and partly by contraction of uterus and fallopian tubes stimulated by oxytocin of female. After capacitation the sperms may reach ampulla within 5 minutes. Sperms can remain viable for 24-48 hours (Ovum for about 24 hours).

b. Entry of sperm into the egg : Out of 200 to 400 million sperms, only few hundred manage to reach the ampulla. Though many sperms reach the ampulla but only a single sperm fertilizes the ovum. A sperm after reaching the egg / ovum comes to lie against it. Its acrosome releases lysins : hyaluronidase and corona penetrating enzymes. They separate and dissolve the cells of corona radiata, so the sperm head passes through the zona pellucida of egg. The zona pellucida has fertilizin receptor proteins (ZP3, ZP2). The fertilizin binds to specific acid protein- antifertilizin of sperm. It brings about attraction of sperms to the egg to enhance fertilization. Fertilizin-antifertilizin interaction is species specific. Thus, the fertilizin-antifertilizin reaction is also called compatibility reaction.

Acrosome reaction : As the sperm head touches the zona pellucida in the animal pole region, its acrosome covering ruptures to release lytic enzymes, acrosin or zona lysin. They act on the zona pellucida at the point of contact. This causes egg reaction - A small fertilization cone / cone of reception is formed on the egg membrane. The sperm head comes in contact with this cone. It results in production of a weak wave of depolarisation. Plasma membrane of the both cells dissolve at the point of contact. The sperm nucleus and the centrioles enter the egg, while other parts remain outside.

As soon as the sperm head touches the vitelline membrane, a cortical reaction gets activated changing the vitelline membrane into a fertilization membrane by deactivating the sperm receptors of zona pellucida. A distinct perivitelline space is created around the fertilization membrane. This prevents any further entry of other sperms into the egg i.e. polyspermy is avoided.



Do you know ?

1. What would happen if the sperm fuses with the egg before it reaches the fallopian tube?
2. What is ectopic pregnancy? Can ectopic pregnancy continue upto full term?

c. Activation of ovum : The ovum before fertilization was at metaphase II stage. With a contact of sperm head to the vitelline membrane of egg, it gets activated to resume and complete its meiosis II. With this it gives out the second polar body. The germinal vesicle organizes into female pronucleus. At this stage, it is the true ovum or egg.

Fusion of egg and sperm : The coverings of male and female pronuclei degenerate allowing the chromosomal pairing. This results in the formation of a **synkaryon** by the

process called **syngamy** or **karyogamy**. The zygote is thus formed. The proximal centriole received from the sperm helps in formation of the synkaryon spindle and cleavage of cell into two blastomeres.



Internet my friend

Find out about extraembryonic membranes.

Significance of fertilization :

- Secondary oocyte completes the process of oogenesis and is transformed into a mature ovum (n).
- The diploid chromosome number is restored in the zygote by the process of syngamy.
- The ovum lacks the centrioles necessary for further divisions, are received from the sperm during fertilization.
- Fertilization involves fusion of male and female gametes from the two parents. It results in variations which are significant to evolution.
- Sex of the offspring is determined.



Always Remember

- Secondary oocyte (egg) is ovulated after LH surge at about the middle of menstrual cycle i.e. day 14.
- Egg (arrested at metaphase II) reaches the ampulla of uterine tube in 12-24 hours after ovulation. The cilia and the fimbriae of the fallopian tube help, direct the egg to ostium.
- During coitus/ intercourse semen is deposited into the vagina of the female. This process is called insemination.
- Human male during ejaculation gives out about 2-4ml of semen with an average count of 200-400 million sperms.

2.6 Embryonic development :

The zygote formed as a result of syngamy is activated to divide.

Cleavage :

It is the process of early mitotic division of the zygote into a hollow multicellular blastula. It does not involve the growth of the daughter cells. The cells formed by cleavage are called **blastomeres**.

Since, there is no growth phase between the cleavages, the size of blastomeres will be reduced with every successive cleavage. As the size reduces, the metabolic rate increases. Subsequent cleavages are thus faster than earlier one. This requires rapid replication of DNA and high consumption of oxygen.

Process of cleavage : In human, cleavage is holoblastic i.e. the whole zygote gets divided. The cleavage planes may be longitudinal or meridional and equatorial or horizontal. It is radial and indeterminate i.e. fate of each blastomere is not predetermined.



Internet my friend

What is meroblastic cleavage? In which organisms, is it observed?

The 1st cleavage in the zygote is meridional and occurs at about 30 hours after fertilization. It divides longitudinally into two blastomeres, one slightly larger than the other. The 2nd cleavage is also longitudinal but at the right angle to the 1st one and occurs after 30 hours of 1st cleavage. The 3rd cleavage is horizontal. After 3rd cleavage the embryo is in 8-cell stage. As the cleavages are going on the young embryo is gradually being pushed towards the uterus. By the end of 4th day after fertilization, embryo is a solid ball of 16-32 cells and externally looking like mulberry. This stage is thus called **morula**. The morula shows cells of two types : smaller, clearer cells towards the outer side and inner cell mass of larger cells. Cells are compactly

arranged. Till the formation of morula the zona pellucida is retained around the embryo and thus, there is no change in the overall size from zygote to morula. The morula reaches the isthmus and gains entry into the uterus by the end of day 4.

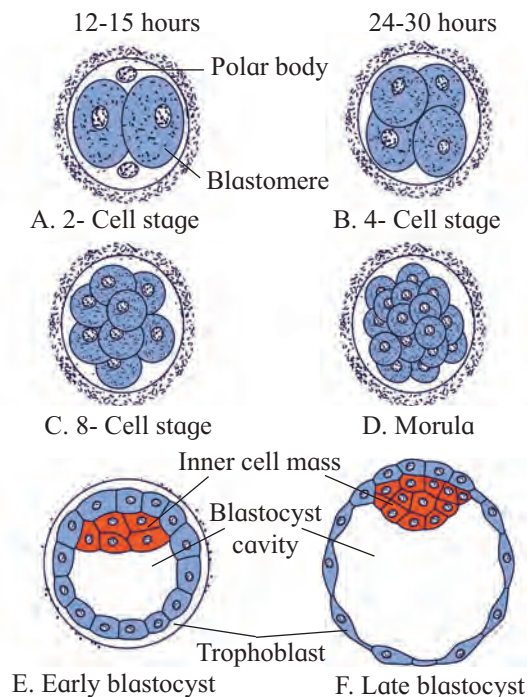


Fig. 2.14 : Process of cleavage and formation of Blastula

Blastulation :

Blastulation is the process of formation of the hollow and multicellular blastocyst. The embryo (blastocyst) that enters the uterus remains floating in uterine cavity for 2-4 days after its entry. i.e. till the end of 7th day after fertilization. The outer layer of cells seen in the morula now form the layer called **trophoblast**.

Cells from the trophoblast begin to absorb the glycogen rich uterine milk. The blastocyst doubles in size from 0.15 mm to 0.30 mm. With more fluid entering inside the blastocyst cavity is formed. These outer cells become flat and are called **trophoblast cells** (since they help only in absorbing nutrition for the developing embryo). The inner larger cells form **inner cell mass** or **embryoblast** (the embryo proper develops from the embryoblasts). These remain attached

to the trophoblasts on only one side. The trophoblast cells in contact with the embryonal knob are called **cells of Rauber**. At this stage, the blastocyst shows polarity. The side with inner cell mass is called the embryonal end and the side opposite to it is the abembryonal end. By the end of the 7th day the blastocyst is fully formed. It is now ready for implantation and gastrulation. The function of zona pellucida is to prevent the implantation of the embryo at an abnormal site. It does not expose the sticky and phagocytic trophoblast cells till it reaches the implantation site i.e. within the uterus, hence zona pellucida now ruptures.



Can you recall?

What do you mean by Monozygotic Dizygotic and Conjoined twins.

Implantation :

The blastocyst after its formation, gets implanted or embedded into the endometrium of the uterus. This process usually begins on day 7 after fertilization and by end of 10th day, the embryo is completely buried inside the endometrium. The embryo usually implants in the region of the fundus of uterus. In the process, the embryo attaches itself by its embryonic pole, close to the endometrium. The trophoblast cells of the animal pole have the power to stick to the uterine wall. Rapid division of the trophoblast cells lying against the embryonal knob, takes place. It results in the formation of two distinct layers- syncytiotrophoblast and cytotrophoblast. The outer layer, **syncytiotrophoblast** is syncytium i.e. a layer of protoplasm with many nuclei. It gives out processes which extensively invade the endometrium. The lytic enzymes secreted by the trophoblasts, rupture the endometrial cells thereby making a burrow, into which the embryo begins to get implanted. By the end of the 10th day the whole embryo is deeply embedded into the endometrium, completing the process of implantation.

The inner layer of cells is called **cytotrophoblast** (cells with defined membrane) since, the cells retain their cell boundaries.

Gastrulation :

It is the process of formation of 'gastrula' from the blastocyst. In the gastrula stage, there is slowing of the rate of cleavage or divisions but there are two important events that take place actively :

a. Differentiation of blastomeres : This process results in the formation of three germinal layers i.e. ectoderm, mesoderm and endoderm from the cells of the embryoblast.

b. Morphogenetic movements : These are different types of movements to reach their definite place in the embryo.

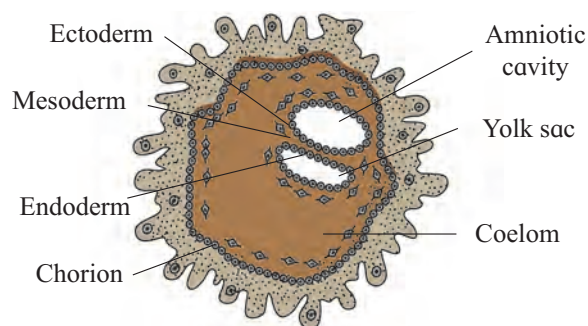


Fig. 2.15 : V. S. of late Gastrula

Gastrulation begins in the embryoblast cells on about 8th day after fertilization. Cell on the free end of inner cell mass called hypoblasts (primitive endoderm) become flatend, start dividing and grows downward towards the blastocoel, cavity of blastocyst. This layer called endoderm is first to differentiate. It grows within the blastocoel and forms a sac called **Yolk sac**. The remaining cell of the inner cell mass, in contact with cells of Rauber are called epiblasts (primary ectoderm). Both layers form a flat, bilaminar embryonal disc.

After formation of endoderm the second layer to be differentiated is the ectoderm. Cells of epiblast divide and redivide and move in such a way that they enclose the amiotic cavity.

Table 2.16 : Fate of germinal layers:

Ectoderm	Mesoderm	Endoderm
Ectoderm gives rise to epidermis of skin, hair, nails, sweat glands, salivary glands, mammary glands, lacrimal glands, sebaceous glands, cornea, lens, retina, conjunctiva, nasal epithelium, enamel of teeth, internal and external ear, foregut, hindgut, adrenal medulla, anterior and posterior pituitary, pineal gland, entire nervous system.	Mesoderm forms all types of muscles (except iris muscles and ciliary muscles of eye which originate from ectoderm), connective tissues, dermis of skin, adrenal cortex, heart, blood, blood vessels, lymphatic vessels, middle ear, dentine of teeth, urinary and reproductive ducts, gonads, kidneys, sclera and choroid of eye.	Endoderm develops into epithelium of mid-gut, glands of stomach and intestine, tongue, tonsils, lungs, trachea, bronchi, larynx, urinary bladder, vagina, liver, pancreas, thyroid gland, parathyroid gland, thymus gland, Eustachian tube, epithelium of urethra, lining of middle ear.

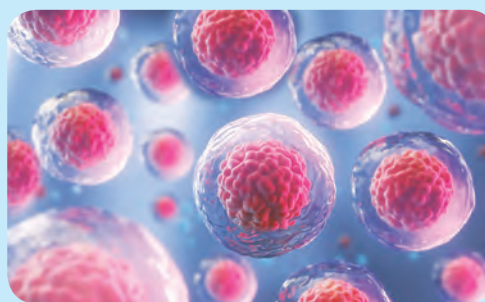
The floor of this cavity has the embryonal disc. The pyramidal cells of the disc towards the amniotic cavity form the embryonal ectoderm. The roof of amniotic cavity is lined by amniogenic cells. Later, these cells divide and redivide to form the amnion. Amnion is an extra embryonic membrane that surrounds and protects the embryo. As a result of all these changes, the bilaminar embryonic disc is positioned in between amniotic cavity and Yolk sac.

Actual gastrulation occurs about 15 days after fertilization, in which the bilaminar embryonic disc is transformed into trilaminar embryonic disc. This transformation occurs by division, rearrangement and migration of cells of epiblast. It begins with formation of primitive streak and a shallow groove on the surface is called primitive groove. This streak progresses from posterior to anterior end of embryo. From site of a primitive streak, a third layer of cells called mesoderm extends between ectoderm and endoderm. Anterior end of primitive groove communicates with yolk sac by an aperture called blastopore (future anus). The embryonal disc now has differentiated into three layers-ectoderm, mesoderm and endoderm. The further process after gastrulation is called histogenesis followed by **organogenesis**.



Do you know ?

Stem cells: These are undifferentiated somatic cells of a multicellular organism. They are capable of giving rise to many more cells of the same type or they can also differentiate into other type of cells. Bone marrow cells, blood stem cells cord cells or umbilical cord cells are examples of stem cells. They can be used in the treatment of Parkinson's disease, Alzheimer's disease, Diabetes, Leukemia, Arthritis, etc.



2.7 Pregnancy :

It is the condition of carrying one or more embryos in the uterus. It is also called **gestation**. It refers to the period between fertilization of the egg, upto parturition. The average period of pregnancy in human lasts for 266 days from fertilization or **280 days** (266+14) counted from LMC- Last Menstruation Cycle. This pregnancy period of approximately nine months is divided into three trimesters of three months each.

First Trimester :

(Period from fertilization to 12th week)

It is the time of most radical changes in mother and embryo. The embryo receives nutrients in the first 2-4 weeks directly from the endometrium. It is the main period of organogenesis or the development of body organs. By the end of eight weeks, the major structures found in the adult are formed in the embryo in a rudimentary form. The embryo is now called foetus and is about 3cm long. The arms, hands, fingers, feet, toes are formed and the foetus can open and close its mouth and fists. CNS is fully formed, working of excretory and circulatory systems begins. Movements of foetus begin but mother can not feel it. Heart beat can be heard from 6th week. At the end of first trimester foetus is about 7-10 cm long.

Meanwhile, the mother's body also undergoes rapid changes. Progesterone level becomes high and menstrual cycle is suspended till the end of pregnancy. High levels of progesterone initiate changes in her reproductive system. The maternal part of placenta grows, the uterus becomes larger. In this period, the mother experiences '**morning sickness**' (nausea, vomiting, mood swings, etc).

Second Trimester :

(Period from 13th to 26th week)

It is the period of rapid growth of foetus. The uterus grows enough for the pregnancy to become obvious. The foetus is very active and grows to about 30 cms. Development of brain begins. Hormone levels stabilize, human chorionic gonadotropin (hCG) declines, the corpus luteum deteriorates (regresses) and the placenta completely takes over the production of progesterone which maintains the pregnancy.

Ultrasound (sonography) at 18-20 weeks shows baby's growth and position. From this estimated due date of delivery can be established. Baby's movements can be easily

felt by the mother. Head has hair, eyebrows and eyelashes appear, pinnae are distinct. The baby reaches half the size of a newborn.

Third (final) Trimester :

(Period from 27th week till the parturition)

The foetus grows to about 3-4 kg in weight and 50 cms in length. Eyes are open. There is gain in body weight. As the foetus grows, the uterus expands around it, the mother's abdominal organs become compressed and displaced, leading to frequent urination, digestive blockages and strain in the back muscles. At the end of third trimester the foetus becomes fully developed and ready for parturition.

2.8 Placenta:

It is a flattened, discoidal organ in the uterus of a pregnant woman. The placenta is a temporary structural and functional connection between foetal and maternal circulation. It is attached to the wall of the uterus and to the baby's umbilical cord. The placenta facilitates the supply of oxygen, nutrients, hormones, antibodies and also the removal of carbon dioxide and excretory wastes produced by the foetus.

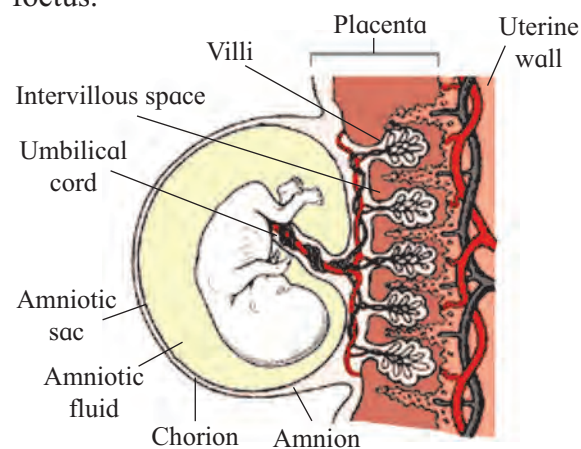


Fig. 2.17 : Placenta

Placenta is the only organ, which is formed of tissues from two different individuals- the mother and the foetus. Part of the placenta contributed by the foetus is called the foetal placenta and it is formed by the **chorionic villi**.

The other part which is rich in blood supply is shared by a part of uterine wall of mother and is termed **maternal placenta**. So human placenta is called **haemochorial**.

The umbilical cord is formed of three blood vessels. Of these three blood vessels, two are small arteries which carry blood towards the placenta and one is a large vein which returns blood to the foetus.

The placenta also acts as an endocrine tissue and produces hormones like hCG, progesterone, estrogen while relaxin is secreted by the ovary in the later phase of pregnancy. Level of hCG increases upto the end of first trimester and then it declines. By the end of first trimester progesterone is produced by placenta. These hormones are required for foetal growth and maintenance of pregnancy.



Know The Institute :

Cord blood bank, Kolkata

India's first Government-run cord blood bank at Kolkata was established in 2001 and is accredited



by AABB (American Association of Blood Bank). The cord blood bank functions according to the central and state government policies, rules and guidelines.

Cord blood (umbilical cord blood) is the blood that remains in the umbilical cord and placenta, post delivery. Cord blood banking is the process of collecting the cord blood, extraction and cryogenically preserving for its stem cells and other cells of the immune system for future potential medical use. Cord blood is rich in stem cells that can transform into all sorts of blood cells. They can be used to treat diseases that harm the blood and immune system e.g. leukemia, certain cancers, sickle cell anemia and some metabolic disorders.



Always Remember

hCG, HPL (Human placental Lactogen), relaxin are produced in women only during pregnancy.

2.9 Parturition:

Humans are viviparous, as they give birth to their young ones. Parturition is the process of giving birth to a baby. The physical activities involved in parturition like uterine and abdominal contractions, dilation of cervix and passage of baby are collectively called labour. Labour is accompanied by localised sensation of discomfort or agony called **labour pains**.

Parturition is controlled by a complex neuroendocrine mechanism. Signals arise from the fully formed foetus and placenta cause mild uterine contractions. It is accompanied by rise in estrogen- progesterone ratio, increase in oxytocin receptors in uterine muscles, .

They cause vigorous contractions of myometrium of uterus at the end of pregnancy. The fully developed foetus gives signals for the uterine contractions by secreting Adrenocorticotrophic Hormone (ACTH) from pituitary and corticosteroids from adrenal gland. This triggers release of oxytocin from mother's pituitary gland, which acts on uterine muscles of mother and causes vigorous uterine contractions. This leads to expulsion of the baby from the uterus. It involves the following three steps :

1. Dilation stage : Uterine contractions begin from top, forcing the baby towards the cervix. Contractions are accompanied by pain caused by compression of blood vessels. Oxytocin induced uterine contractions become stronger and stronger due to stimulatory reflex. As the baby is pushed down in the uterus, its head comes to lie against cervix. Cervix gets dilated. The vagina also shows similar dilation. This stage of labour can normally last upto few hours. It ends in rupturing of amniotic membrane of foetus.

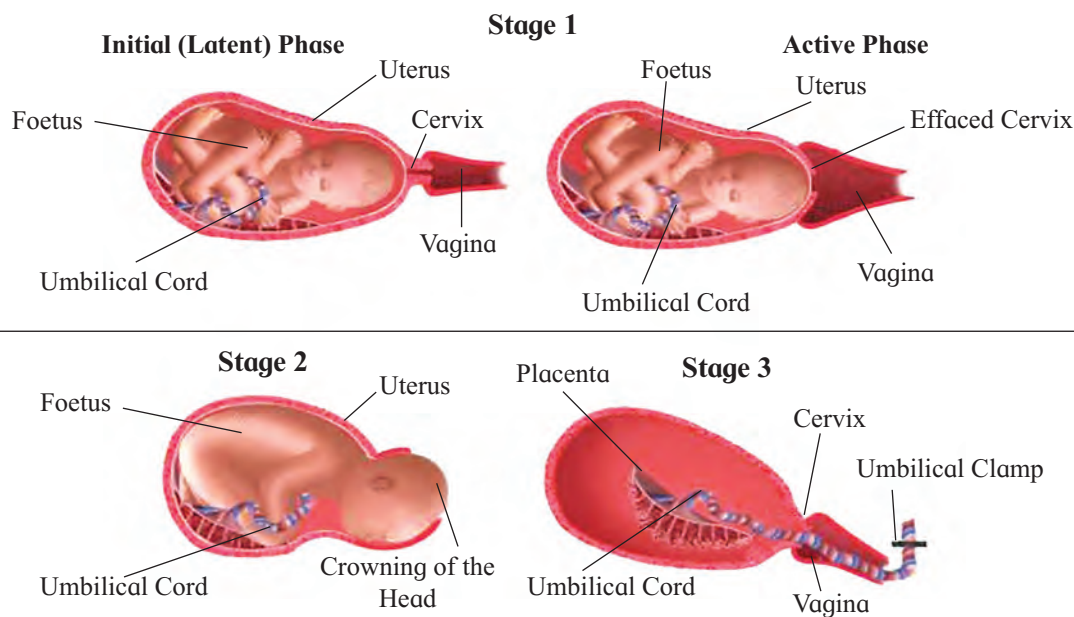


Fig. 2.18 : Parturition

2. Expulsion stage : The uterine and abdominal contractions become stronger. In normal delivery, the foetus passes out through cervix and vagina with head in forward direction. It takes 20 to 60 min. The umbilical cord is tied and cut off close to the baby's navel.

3. After birth : After the delivery of the baby the placenta separates from the uterus and is expelled out as "after birth", due to severe contractions of the uterus. This process happens within 10 to 45 minutes of delivery.

2.10 Lactation:

The mammary glands of the female start producing milk at the end of pregnancy by the process of lactation. Prolactin is the hormone which is responsible for production of milk. Lactation helps the mother in feeding the new born baby. The fluid secreted by the mammary glands soon after child birth is called **colostrum**. **Colostrum:** It is the sticky and yellow fluid secreted by the mammary glands soon after child birth. It contains proteins, lactose and mother's antibodies e.g. IgA. The fat content in colostrum is low. The antibodies present in it helps in developing resistance for the new born baby at a time when its own immune response is not fully developed.



Use your brain power

Why is it said that consumption of mother's milk is safest for the new born?

2.11 Reproductive Health:

According to World Health Organisation (WHO), reproductive health means total wellbeing in all aspects of reproduction- its emotional, behavioural and social aspects along with the physical ones. Therefore, a society with people having physically and functionally normal reproductive organs and normal emotional and behavioural interactions amongst them in all sex-related aspects might be called reproductively healthy society.

Of all the social goals of India, an important one is to attain total reproductive health. India was amongst the first few countries in the world to initiate action plans and programmes at a national level to improve reproductive health. All these improved programmes cover wider areas related to reproduction. These programs are currently in operation under the Reproductive and Child Health Care (RCH) programmes.

Goals of RCH Programmes:

1. To create awareness among people about various aspects related to reproduction.
2. To provide the facilities to people to understand and build up reproductive health.
3. To provide support for building up a reproductively healthy society.
4. To bring about a change mainly in three critical health indicators i.e. reducing total infertility rate, infant mortality rate and maternal mortality rate.

The goals of RCH can be achieved by the following ways:

1. By introduction of sex education in schools. Schools should be encouraged to provide correct information to the young so as to discourage children from believing in myths and clear the misconceptions about sex related aspects. Proper information about safe and hygienic sexual practices, sexually transmitted diseases (STD, AIDS), problems related to adolescence and proper information about reproductive organs.
2. With the help of audio- visual and the print media, government and non- government organisations should take various steps to create awareness about various aspects related to reproduction.
3. By educating the younger generation about birth control measures, pre-natal care of pregnant woman and post-natal care of the mother and child, importance of breast feeding.
4. By developing awareness about problems arising due to uncontrolled population growth, social evils like sex abuse and sex related crimes and take up necessary steps to prevent them.
5. By creating awareness about statutory ban on amniocentesis for sex determination.
6. By creating awareness about child immunization programmes.

7. By educating couples to reduce mortality rate of new borns and maternal mortality rate.

The population in India which was approximately 350 millions at the time of independence, reached close to a billion mark by 2000 and crossed 1.2 billion in May 2011. Now in 2020 population of India has crossed 1.35 billions. The government is taking serious measures to check this population growth. The most important step to overcome this problem, is to motivate society to have smaller families by using various birth control methods.

2.12 Birth control :

The birth control measures which deliberately prevent fertilization are referred to as contraceptives. The contraceptive methods help to prevent unwanted pregnancies. An ideal contraceptive should be easily available, user friendly, effective and with no or least side effects.

Contraceptive methods are of two main types i.e. temporary and permanent.

a. Temporary methods:

These are of following types :

1. Natural method/ Safe period / Rhythm method :

In the natural method, the principle of avoiding chances of fertilization is used. A week before and a week after menstrual bleeding is considered the safe period for sexual intercourse. This idea is based on the fact that ovulation occurs on the 14th day of menstrual cycle. Its drawback lies in having a high rate of failure.

2. Coitus Interruptus or withdrawal : In this method, the male partner withdraws his penis from the vagina just before ejaculation, so as to avoid insemination. This method also has some drawbacks, as the pre-ejaculation fluid may contain sperms and this can cause fertilization.

3. Lactational amenorrhea (absence of menstruation) : This method is based on the fact that ovulation does not occur during the period

of intense lactation following parturition. Therefore, as long as the mother breastfeeds the child fully, chances of conception are almost negligible. However, this method also has high chances of failure.

4. Chemical means (spermicides): In this method, chemicals like foam, tablets, jellies, and creams are used by the female partner. Before sexual intercourse, if these chemicals are introduced into the vagina, they adhere to the mucous membrane, immobilize and kill the sperms. It may cause allergic reaction. This method also has chances of failure.

5. Mechanical means / Barrier methods: In this method, with the help of barriers the ovum and sperm are prevented from physically meeting. These mechanical barriers are of three types.

i) Condom: It is a thin rubber sheath that is used to cover the penis of the male during copulation. It prevents the entry of ejaculated semen into the female reproductive tract. This can prevent conception. It is a simple and effective method and has no side effects. “Nirodh” is the most widely used contraceptive by males. It is easily available and is given free by the government. It should be properly discarded after every use. Condom is also a safeguard against STDs and AIDS.

ii) Diaphragm, cervical caps and vaults: These devices used by the female are made up of rubber. They prevent conception by blocking the entry of sperms through the cervix. The device is inserted into the female reproductive tract to cover the cervix during copulation.

iii) Intra-uterine devices (IUDs) : These clinical devices are plastic or metal objects. A doctor or trained nurse places the IUDs into the uterus. These devices include Lippes loop, copper releasing IUDs (Cu-T, Cu7, multiload 375) and hormone releasing IUDs (LNG-20, progestasert).



Diaphragm



Lippes loop

Fig. 2.19 : Mechanical means

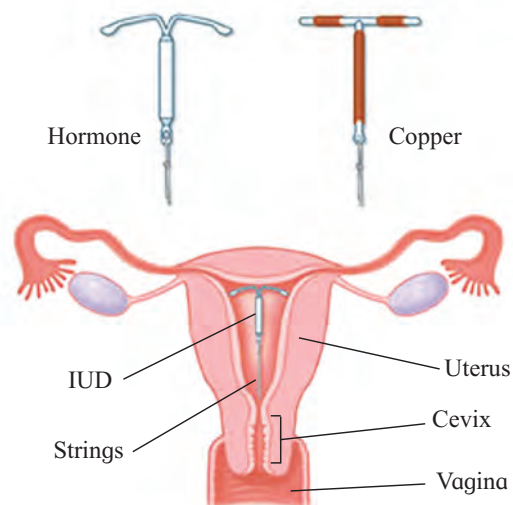


Fig. 2.20 : Copper - T

Lippes loop is a plastic double “s” loop. It attracts the macrophages stimulating them to accumulate in the uterine cavity. Macrophages increase phagocytosis of sperms within the uterus and acts as a contraceptive. Copper releasing IUDs suppress sperm motility and the fertilizing capacity of sperms.

The hormone releasing IUDs make the uterus unsuitable for implantation and cervix hostile to the sperms. It delays pregnancy for longer period. The spontaneous expulsion, occasional haemorrhage and chances of infection are the drawbacks of IUDs.

6. Physiological (Oral) Devices : Physiological devices are used in the form of tablets and hence are popularly called **pills**. It is an oral contraceptive, used by the female. The pill contains progesterone and estrogen. They inhibit ovulation, hence no eggs are released from the ovary of the female using this pill and thus conception cannot occur. They also alter the quality of cervical mucus to prevent the entry of sperms.

The pills have side effects such as nausea, weight gain, tenderness of breast and slight blood loss between menstrual periods. The pill “**Saheli**” is an oral contraceptive for females which is nonsteroidal. Saheli is to be taken once in a week. These pills are sponsored by the Government.



Always Remember

Saheli is now a part of the National Family Programme as an oral contraceptive pill in India.



7. Other contraceptives : The birth control **implant** is a contraceptive used by the female. It is a tiny, thin rod about the size of a matchstick. It is implanted under the skin of the upper arm.

They contain progesterone and estrogen. Their mode of action is similar to that of pills. They prevent pregnancy for 3-4 years.

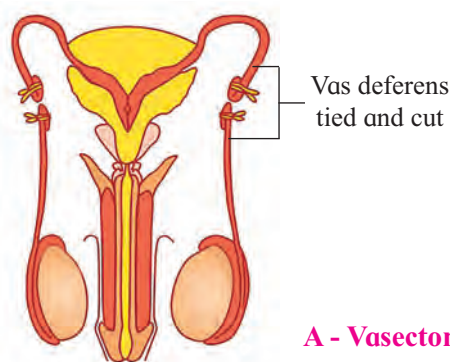


Fig. 2.21 : Implanon/ Nexplanon

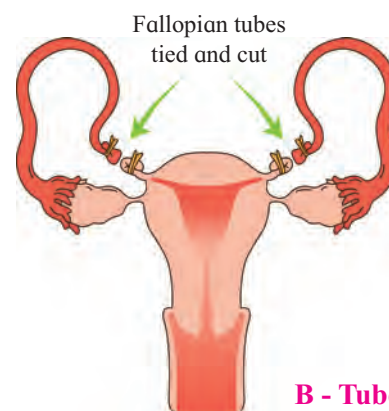
b. Permanent Method:

The permanent birth control method in men is called vasectomy and in women it is called tubectomy.

These are surgical methods, also called sterilization. In vasectomy a small part of the vas deferens is tied and cut where as in tubectomy, a small part of the fallopian tube is tied and cut. This blocks, gamete transport and prevent pregnancy.



A - Vasectomy



B - Tubectomy

Fig. 2.22 : Permanent method

Medical Termination of Pregnancy (MTP) :

An intentional or voluntary termination of pregnancy before full term is called Medical termination of Pregnancy (MTP) or induced abortion. MTP is essential in cases of unwanted pregnancies or in defective development of foetus. It is safe during the first trimester of pregnancy. The defective development of foetus is examined by amniocentesis.

Amniocentesis is a process in which amniotic fluid containing foetal cells is collected using a hollow needle inserted into the uterus under ultrasound guidance. The chromosomes are studied to see the abnormalities in the developing foetus. But the dangerous trend is the misuse of amniocentesis to determine the sex of the unborn child. Frequently, if the foetus is found to be female, it is aborted which is totally illegal. So the Government of India has legalised MTP Act in 1971, with strict conditions to avoid its misuse.

Amniocentesis : Used to extract foetal cells for genetic analysis.

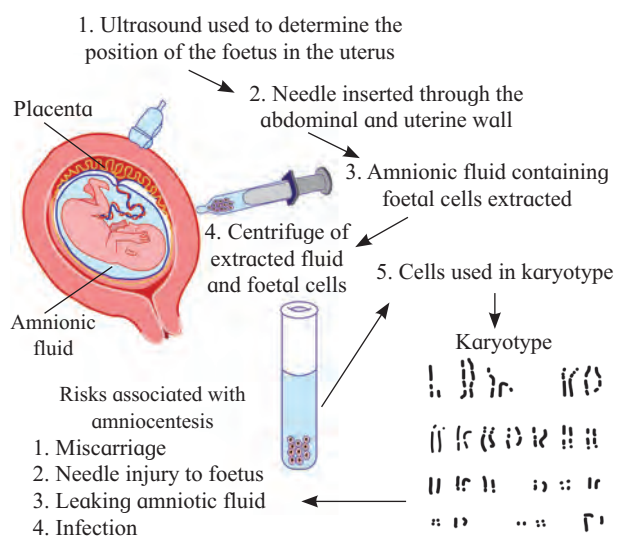


Fig. 2.23 : Amniocentesis process



Always Remember

Act of MTP : The Medical Termination of Pregnancy (MTP) Act 1971 provides the legal framework for MTP Act 2017.



Activity :

In a sonography clinic, we observe a board saying ‘Sex selection and detection is NOT done in this centre and is punishable under PC-PNDT Act;

Find out what is PC-PNDT Act. Why do you think such a mandate is essential?

Medical Termination of Pregnancy (Amendment) Act 2017 under section 3 of the MTP Act 1971 was enacted by Government of India. The intention of MTP Act is to reduce the incidence of illegal abortion and consequent maternal mortality. As per the provisions of the MTP Act, only the consent of woman whose pregnancy is being terminated is required. According to MTP Act pregnancy may be terminated : 1. Within first 12 weeks 2. More than 12 weeks but lesser than 20 weeks. The registered medical practitioner’s opinion is mandatory stating the continuation of the pregnancy would involve a risk to the life of the pregnant woman or grave abnormal physical or mental health or is substantial risk to the child.



Internet my friend

What are the effects of alcohol drinking and smoking on foetus in pregnant women?

2.13 Sexually Transmitted Diseases (STDs) :

Diseases or infections which are transmitted through sexual intercourse are collectively called Sexually Transmitted Diseases (STDs) or Venereal Diseases (VDs) or Reproductive Tract Infections (RTI). The major venereal diseases are syphilis and gonorrhoea.



Internet my friend

Collect information about other sexually transmitted diseases.

Table 2.24 : Sexually Transmitted Diseases (STDs)

Name of Disease	Syphilis	Gonorrhoea
Causative agent	<i>Treponema pallidum</i> (Bacteria)	<i>Neisseria gonorrhoeae</i> (Bacteria)
Incubation period	3-4 weeks	Male – 2 to 14 days Female – 7 to 21 days
Infection site	Mucous membrane in genital, rectal and oral region.	Mucous membrane of urino-genital tract, rectum, throat and eye.
Symptoms	Primary lesion called chancre at the site of infection. Chancre is formed on external genitalia, skin rashes and mild fever, inflamed joints, loss of hair. Paralysis, Degenerative changes occur in the heart and brain.	In male, partial blockage of urethra and reproductive ducts, pus from penis, pain and burning sensation during urination, arthritis, etc. In female, pelvic inflammation of urinary tract, sterility, arthritis, the children born to affected mother suffer from gonococcal ophthalmia and gonococcal vulvovaginitis of girls before puberty.
Preventive measures	Education about sex practices, sex hygiene, avoiding sex with unknown partner or multipartners, using condom during coitus.	Sex hygiene, using condom during coitus, avoiding sex with unknown partner or multipartners.
Treatment	Antibiotic-Penicillin	Antibiotic-Cefixime

2.14 Infertility :

Infertility is defined as the inability to conceive naturally after (one year of) regular unprotected intercourse. The causes of infertility could be physical, congenital, diseases, immunological or even psychological. The common physical causes in females are polycystic ovary syndrome (PCOS), hormonal imbalance, endometriosis while in male, it is less sperm count and small size of penis.

Prior to 1978, infertile couple had two options, adopt or be childless. Today infertile couples have many options to have a child such as fertility drugs, test tube babies, artificial insemination, IUI, surrogate motherhood, etc. The couple could be assisted to have child / children through certain special techniques commonly known as Assisted Reproductive Technologies (ART).

IVF (In Vitro Fertilization) :

It is a process of fertilization where an egg is combined with sperm outside the body in a test tube or glass plate to form a zygote under simulated conditions in the laboratory. The zygote or early embryos (with up to 8 blastomeres) could be then transferred into the fallopian tube for further development.

ZIFT (Zygote Intrafallopian Transfer) :

ZIFT is an infertility treatment used when there is a blockage in the fallopian tubes which prevents the fertilization of egg by the sperm.

In this method, egg is removed from woman's ovary. Fertilization of the egg with sperms is brought about outside the body under sterile conditions to form zygote by the process called *in vitro* fertilization (IVF). The zygote is then transferred to fallopian tube for further development.



Try This

IVF centres : Make a list of IVF centres in Maharashtra.

GIFT (Gamete Intrafallopian Transfer) :

Transfer of an ovum collected from a donor into the fallopian tube of another female who can provide suitable environment for its fertilization and development.

This technique called gamete intrafallopian transfer (GIFT) has been developed for the cases in which only the entrance to the oviducts or the upper segment of the oviducts is blocked. In this procedure ova and sperms are directly injected into regions of the oviduct, where fertilization produces a blastocyst, which enters the uterus via the normal route. GIFT has a success rate of about 30 percent.

ICSI (Intra Cytoplasmic Sperm Injection) :

ICSI is an *in vitro* fertilization procedure in which a single sperm cell is injected directly into cytoplasm of an ovum in the laboratory. Here the sperm has to naturally penetrate the egg.

Artificial Insemination (AI) :

In some infertility cases, the male partner is unable to inseminate the female due to a very low sperm count. This problem can be solved by artificial insemination. In this technique, the sperms are collected from the male and artificially introduced into the cervix of female, for the purpose of achieving a pregnancy through *in vivo* fertilization (inside the body).

IUI (Intra Uterine Insemination) :

In this technique the process is somewhat like that of artificial insemination, the only difference is that the sperms are introduced into the uterine cavity instead of cervix.

Sperm bank / Semen bank:

A sperm bank or semen bank is a place which collects, stores and provides human sperms / semen. The semen is provided by healthy males called **sperm donors**. The sperms are stored in sperm bank by cryopreservation method (at low temperature).



Can you recall?

Surrogate mother :

Some women have problem in implantation of embryo in uterus. Such woman can take help of the modern remedial technique called surrogacy. In this, embryo is implanted in surrogate mother, who is not the biological mother.

Adoption :

Adoption is a legal process by which a couple or a single parent gets legal rights, privileges and responsibilities that are associated to a biological child for the upbringing of the adopted child.

An adoptive parent should be medically fit and financially able to take care of the adopted child. A person wishing to adopt a child must be at least 21 years old but there is no legal upper age limit for adoption.



Always Remember

Tobacco, marijuana and other drugs smoking may cause infertility in both men and women. Nicotine blocks the production of sperm and decreases the size of testicles. Alcoholism by men interferes with the synthesis of testosterone and has an impact on sperm count. Use of cocaine or marijuana may temporarily reduce the number and quality of sperm.



Can you tell?

- Jayesh, a young married man of 26 yrs is suffering from T. B. for the last 2 years. He and his wife are desirous of a child but unable to have one, what could be the possible reason? Explain.
- Neeta is 45 years old and the doctor has advised her not to go for such a late pregnancy. She however wants to be the biological mother of a child, without herself getting pregnant. is this possible and how?



Always Remember

1. Cells of trophoblast do not form any part of the embryo proper.
2. They form ectoderm of the chorion (extra embryonic membrane).
3. They play important role in formation of placenta.



Activity :

1. Prepare concept map on information of male reproductive system.

2. Prepare concept map on information of female reproductive system.



Activity :

3. Prepare concept map on information of menstrual cycle.

Blank area for drawing a concept map on the menstrual cycle.

4. Prepare concept maps on information of gametogenesis.

Spermatogenesis

Oogenesis

Blank area for drawing concept maps on spermatogenesis and oogenesis.

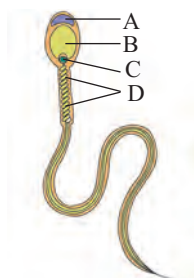
5. Prepare concept map on information of fertilization.

Blank area for drawing a concept map on fertilization.

Exercise

Q. 1 Multiple choice questions.

1. The number of nuclei present in a zygote is
a. two b. one c. four d. eight
2. Which of these is the male reproductive organ in human?
a. sperm b. seminal fluid
c. testes d. ovary
3. Attachment of embryo to the wall of the uterus is known as.....
a. fertilization b. gestation
c. cleavage d. implantation
4. Rupturing of follicles and discharge of ova is known as
a. capacitation b. gestation
c. ovulation d. copulation
5. In human female, the fertilized egg gets implanted in uterus
a. After about 7 days of fertilization
b. After about 30 days of fertilization
c. After about two months of fertilization
d. After about 3 weeks of fertilization
6. Test tube baby technique is called.....
a. In vivo fertilization
b. In situ fertilization
c. In vitro fertilization
d. Artificial insemination
7. The given figure shows a human sperm. Various parts of it are labelled as A, B, C, and D. Which labelled part represents acrosome ?



- a. B b. C c. D d. A

8. Presence of beard in boys is a
a. primary sex organ
b. secondary sexual character
c. secondary sex organ
d. primary sexual character

Q. 2 Answer in one sentence.

1. What is the difference between a foetus and an embryo?
2. Outline the path of sperm upto the urethra.
3. Which glands contribute fluids to the semen?
4. Name the endocrine glands involved in maintaining the sex characteristics of males.
5. Where does fertilization and implantation occur?
6. Enlist the external genital organs in female.
7. Give two differences between blastula and gastrula.
8. What is the difference between embryo and zygote?

Q. 3 Fill in the blanks:

1. The primary sex organ in human male is
2. The..... is also called the womb.
3. Sperm fertilizes ovum in the of fallopian tube.
4. The disc like structure which helps in the transfer of substances to and from the foetus's body is called.....
5. Gonorrhoea is caused by bacteria.
6. The hormone produced by the testis is

Q. 4 Short answer questions.

1. Write a note on budding in *Hydra*.
2. Explain the different methods of reproduction occurring in sponges.
3. Write a note on IVF
4. Comment on any two mechanical contraceptive methods.
5. Write a note on tubectomy
6. Give the name of causal organism of **syphilis** and write on its symptoms.
7. What is colostrum?

Q. 5 Answer the following questions.

1. Describe the phases of menstrual cycle and their hormonal control.
2. Explain the steps of parturition.
3. Explain the histological structure of testis.
4. Describe the structure of blastula.
5. Explain the histological structure of ovary in human.
6. Describe the various methods of birth control to avoid pregnancy.
7. What are the goals of RCH programme.
8. Which hormones are involved in parturition?
9. Which as the function of male accessory glands?
10. What is capacitation? Give it's importance.

Q. 6 Long answer questions.

1. Explain the following parts of male reproductive system along with labelled diagram showing these parts- Testis, vasa deferentia, epididymis, seminal vesicle, prostate gland and penis.
2. Describe female reproductive system of human.
3. Describe the process of fertilization.
4. Explain the process by which zygote divides and redivides to form the morula.

Project :

Prepare a chart showing information about other STDs, mentioning causal organisms, symptoms and control measures.



Can you recall?

1. Is there a similarity between the parents and offsprings?
2. What are the roles of chromosomes in living organisms?
3. How are hybrid seeds produced?
4. Which are the chromosomal disorders?

3.1 Chromosomes and Mechanism of inheritance.

- The transmission of genetic information from generation to generation is known as **heredity** or **inheritance**. The mechanism of inheritance was successfully investigated before chromosomes had been observed or genes were known.
- Gregor Mendel, son of the peasant farmer, was born in Moravia in 1822. Gregor Mendel first gave the accurate explanation for the mechanism of inheritance by using hybridization technique.
- Inheritance of the seven traits in garden pea (Refer the diagram below) plant were

studied individually one at a time or in combination of two or three character at a time. He processed the data mathematically and statistically.

- Mendel postulated the principles of heredity which then became the fundamental laws of heredity, as proposed by Correns (1900). He visualized that the traits as such are not inherited physically but by ‘something’ present inside the gametic cell. To this ‘something’, he coined term ‘factors’ that are responsible for expression of a particular trait/ character. He proposed that factors are particulate in nature. The Mendelian factors are now termed as ‘genes’. These factors occur in pairs in the parents and segregate from each other during gamete formation without blending/ mixing.

Reasons for Mendel’s Success :















- His experiments were carefully planned and involved large samples.
- He carefully recorded the number of plants of each type and expressed his results in terms of ratios.



Do you know ?

Seven pairs of contrasting visible characters in pea plant (*Pisum sativum*)

Pea Plant Traits

	Seed shape	Seed color	Pod shape	Pod color	Flower color	Flower location	Plant height
Dominant	Round 	Yellow 	Inflated 	Green 	Purple 	Axial 	Tall 
Recessive	Wrinkled 	Green 	Constricted 	Yellow 	White 	Terminal 	Short (Dwarf) 

- In the pea plant, contrasting characters can be easily recognized.
- The seven different characters in pea plant were controlled by a single factor each. The factors are located on separate chromosomes and these factors are transmitted from generation to generation.
- He introduced the concepts of dominance and recessiveness.

Before learning about Mendel's experiments let us get acquainted with genetic terms and symbols.

3.2 Genetic Terminology :

Character : It is a specific feature of an organism e.g. height of stem (plant).

Trait : An inherited character and its detectable variant e.g. Tall or dwarf.

Factor : It is a unit of heredity, a particle present in the organism which is responsible for the inheritance and expression of a character. (factor is passed from one generation to the next through gametes). Factor determines a genetical (biological) character of an organism.

Gene : It is a particular segment of DNA which is responsible for the inheritance and expression of that character.

Alleles or Allelomorphs : The two or more alternative forms of a given gene (factor) are called alleles of each other. They occupy identical loci (positions) on homologous chromosomes. Allele is a short form of Allelomorph.

Dominant : It is an allele that expresses its trait even in the presence of an alternative allele i.e. in heterozygous condition only. Alternatively, the allele that expresses in F_1 is called dominant. (It is an allele of a pair that masks the expression of other allele in F_1 generation.)

Recessive : This allele is not expressed in the presence of an alternative allele (in heterozygous condition). It expresses only in the presence of another identical allele. It is an allele that does not express in F_1 hybrid.

Phenotype : The external appearance of an individual for any trait is called phenotype for that trait. It is observable and is determined by different combinations of alleles. e.g. In pea, for the height of stem (plant) tall and dwarf are the two phenotypes (Tall is determined by TT or Tt and dwarf by tt).

Genotype : Genetic constitution or genetic make up of an organism with respect to a particular trait. It is representation of the genetic constitution of an individual with respect to a single character or a set of characters. e.g. tall pea plant can have genotype TT or Tt and dwarf has tt.

Homozygous (pure) : An individual possessing identical alleles for a particular trait, is called homozygous or pure for that trait.

Homozygous breeds true to the trait and produces only one type of gametes e.g., tall with TT and dwarf with tt.

Heterozygous : An individual possessing contrasting alleles for a particular trait, is called heterozygous. Heterozygous does not breed true for that trait and produces two types of gametes e.g. F_1 generation hybrids (Tt). Heterozygous individual is also called hybrid.

Pure line : An individual or a group of individuals (population) which is homozygous or true breeding for one or more traits, constitutes pure line i.e. plant which breeds true for a particular character. It is a descendent of a single homozygous parent produced after self fertilization.

Monohybrid : It is heterozygous for one trait and is produced from a cross between two pure parents differing in single pair of contrasting characters e.g. Hybrid tall produced in a cross between pure tall and pure dwarf parents. It is a heterozygote for a single pair of alleles.

F_1 generation : It refers to the first filial generation. It consists of all off-springs produced from a parental cross. Alternatively, it is first generation from a given mating between pure parents having contrasting characters.

F₂ generation : The second generation (progeny) produced by selfing (inbreeding) of F₁ generation offsprings is called second filial generation. e.g. Progeny produced from a cross between two F₁ individuals (e.g. Tt × Tt).

Punnett square/checker board : It is a probability table representing different permutations and combination of fertilization between gametes of the opposite mating types. In short, it is a diagrammatic representation of a particular cross to predict the progeny of a cross.

Homologous Chromosomes : The morphologically, genetically and structurally essentially identical chromosomes present in a diploid cell, are called homologous chromosomes. Such chromosomes synapse during meiosis.

Back cross : It is a cross of F₁ progeny with any of the parents (e.g. F₁ tall × pure tall; F₁ tall × pure dwarf i.e. Tt × TT/tt).

Test cross : It is a cross of F₁ progeny with homozygous recessive parent (e.g. F₁ tall × pure dwarf i.e. Tt × tt). It is used to test the homozygous/ heterozygous nature of hybrid. It is a kind of back cross.

Phenotypic ratio : It is the ratio of the offsprings produced in F₂ and subsequent generation with respect to their physical appearance e.g. 3Tall : 1 dwarf, is F₂ 'Phenotypic ratio' in monohybrid cross.

Genotypic ratio : It is the ratio of the offsprings produced in the F₂ and subsequent generation with respect to their genetic make up e.g. 1 TT : 2Tt : 1 tt, is F₂ genotypic ratio in monohybrid cross.

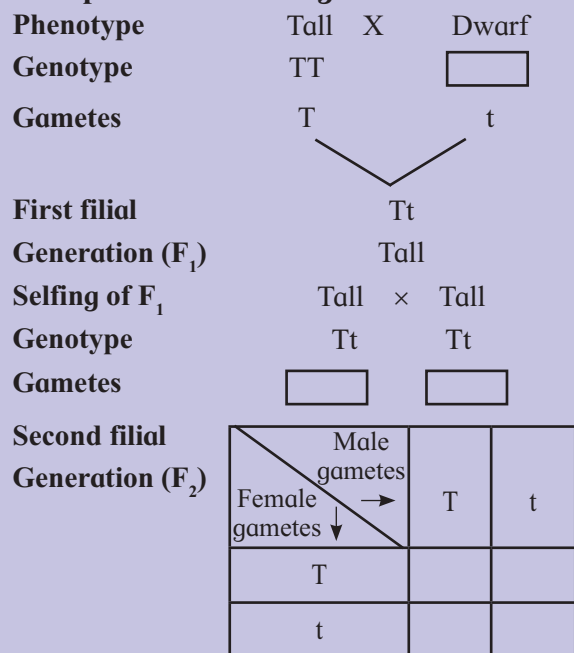
Monohybrid cross :

A cross between parents differing in only one heritable trait is called monohybrid cross. e.g. cross of pure tall and pure dwarf plants. Mendel performed the monohybrid cross between two pea plants with only one pair of contrasting character.



Activity :

Complete the following chart :



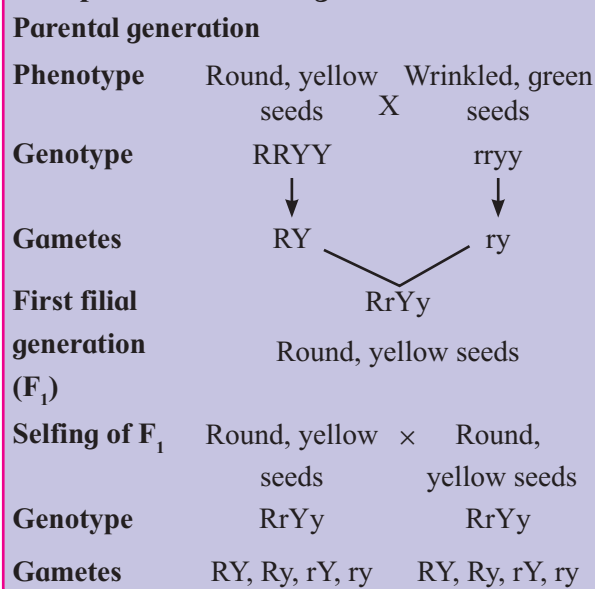
Dihybrid cross :

A cross between parents differing in two heritable traits, is called dihybrid cross e.g. cross of pure tall, round seeded plant with dwarf, wrinkled seeded plant. Mendel also performed the dihybrid cross between pea plants that differed in two pairs of contrasting characters.



Activity :

Complete the following chart :



F₂ Generation :

Male gametes →		RY	Ry	rY	ry
Female gametes ↓					
RY					
Ry					
rY					
ry					



Use your brain power

There are 16 possible individuals in F₂ generation. Try to find out the phenotypes as well as the genotypic and phenotypic ratios.



Can you tell?

Why are farmers and gardeners advised to buy new F₁ hybrid seeds every year?

3.3 Mendel's Laws of Inheritance :

Mendel proposed three basic postulates on the basis of which three laws were formulated. These are described below:

1. Law of Dominance :

In monohybrid and dihybrid crosses, the phenotypic characters are controlled by discrete units, called factors. In a dissimilar pair of factors, one member of the pair dominates (i.e. dominant) over the other (i.e. recessive). The law of dominance is used to explain the expression of only one of the parental characters of a monohybrid cross in F₁ and the expression of both in F₂.

Statement of Law of Dominance : “When two homozygous individuals with one or more sets of contrasting characters are crossed, the alleles (characters) that appear in F₁ are **dominant** and those which do not appear in F₁ are **recessive**”.

2. Law of segregation (Law of purity of gametes) :

This law is based on the fact that the alleles do not show any blending/ mixing and both the alleles (characters) are recovered as such in the F₂ generation, though one of these is not seen at the F₁ stage. During formation of gametes, these two alleles (factors) obviously separate or segregate, otherwise recessive type will not appear in F₂.

The gametes which are formed are always pure for a particular character (trait). A gamete may carry either dominant or recessive factor but not both. That's why it is also called as law of purity of gametes.

Statement of Law of Segregation : The law states that “When hybrid (F₁) forms gametes, the alleles segregate from each other and enter in different gametes”. The gametes formed are pure in that they carry only one allele each (either dominant allele or recessive allele). Hence, this law is also described as “**Law of purity of gametes**”.

3. Law of Independent Assortment :

This law is based on dihybrid cross. It is basic principle of genetics developed by a Mendel. It describes how different genes or alleles present on separate chromosomes independently separate from each other, during formation of gametes. These alleles are then randomly united in fertilization. In dihybrid cross, F₂ phenotypic ratio 9:3:3:1 indicates that the two pairs of characters behave independent of each other. It can be concluded that the two characters under consideration are assorted independently giving rise to different combinations.

Statement of Law of Independent Assortment: The law states that “When hybrid possessing two (or more) pairs of contrasting factors (alleles) forms gametes, the factors in each pair segregate independently of the other pair”.



Try This

Find the ratio of dihybrid test cross by using punnett square.

3.4 Back Cross and Test Cross :

a. Back cross : The F_1 individuals obtained in a cross are usually selfed to get the F_2 progeny. They can also be crossed with one of the two parents from which they were derived (either recessive or dominant). Such a cross is known as **back cross**.

b. Test cross : The cross of F_1 hybrid with the homozygous recessive parent is known as a **test cross**. It is used to test whether an individual is homozygous (pure) or heterozygous (hybrid). Test cross is easy, simple, repeatable and predictable.

Test cross can be used to find out genotype of any plant with dominant expression. But it is not known whether it is homozygous (pure) or heterozygous for that trait. For example, A pea plant having violet (purple) flowers is crossed with a pea plant with white flowers. If all flowers produced are violet, we can conclude that plant is pure or homozygous and if we get violet and white flowers in 1:1 ratio, we can

conclude that plant is heterozygous. Test cross is also used to introduce useful recessive traits in the hybrids of self pollinated plants during rapid crop improvement programs.

Following is the graphic representation of test cross (Fig. 3.1). Recessive parent is crossed to find out unknown genotype.

3.5 Deviations from Mendel's findings :

Few generalizations were arrived at by Mendel, on the basis of his experiments of garden pea plant- such as,

- Single trait \rightarrow Single gene \rightarrow Two alleles.
- Two alleles show interaction in which one is completely dominant.
- Factors (genes) for different traits present on different chromosomes assort independently.

With the passage of time, number of deviations were observed/ identified in the post-Mendelian era, that gave additional information on the patterns of inheritance. These deviations are then described as **Neo-Mendelism**.



Internet my friend

Find out the principle involved in heredity of sheep/ coat colour

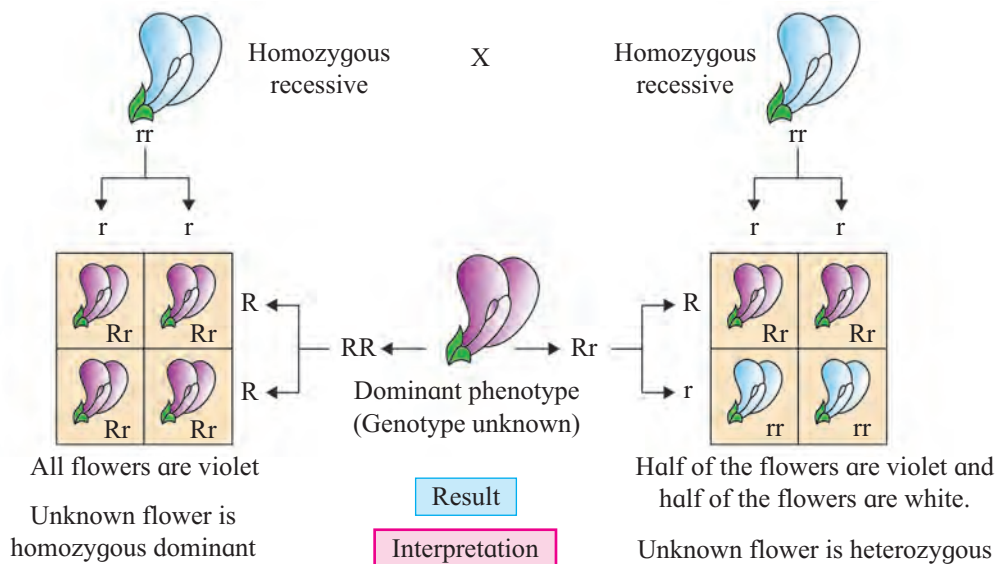


Fig. 3.1 : Graphical representation of test cross



Did you know ?

Deviations from Mendelism are :

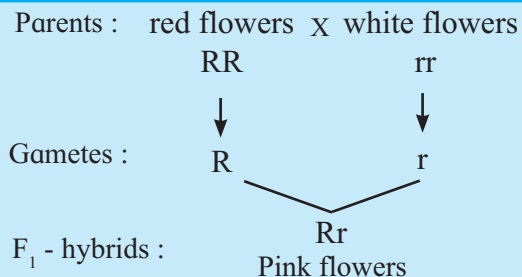
- Single trait \rightarrow Single gene \rightarrow two alleles showing interactions like codominance and incomplete dominance.
- Single trait \rightarrow Single gene \rightarrow more than two alleles (multiple allelism).
- Single trait \rightarrow more than one genes (Polygenic inheritance) showing different epistatic interactions or additive effect.
- Single gene influencing many traits (pleiotropy).

It was observed that the phenotypic expression of a gene can be modified or influenced by another gene. These gene interactions are of two types.

- Intragenic interactions** occur between the alleles of same gene e.g. incomplete dominance and co-dominance. It also occurs between the multiple allele series of a gene.
- Intergenic (non-allelic) interactions** occur between the alleles of different-genes present on the same or different-chromosomes. e.g. pleiotropy, polygenes, epistasis, supplementary and complementary genes, etc. Some of these interactions are discussed below :

a. Incomplete dominance:

In the incomplete dominance, both the alleles (genes) of an allelomorph pair express themselves partially. One allele (gene) cannot suppress the expression of the other allele (gene) completely. In such case, there is an intermediate expression in the F_1 hybrid. A well-known example is the flower colour of *Mirabilis jalapa*. If a red-flowered (RR) plant is crossed with a white-flowered (rr) plant, then F_1 offsprings have pink (Rr) flowers.



F_2 Generation : Selfing of F_1

$\begin{matrix} \text{♂} \\ \text{♀} \end{matrix}$	R	r
R	RR red	Rr pink
r	Rr pink	rr white

Result :

Genotypic ratio - 1RR : 2Rr : 1rr

Phenotypic ratio - 1Red : 2 Pink : 1 White

b. Co-dominance :

In co-dominance, both the alleles (genes) of an allelomorph pair express themselves equally in F_1 hybrids. Such alleles which are able to express themselves equally independently in hybrids, are called co-dominant alleles. Thus in co-dominance both alleles are expressed.

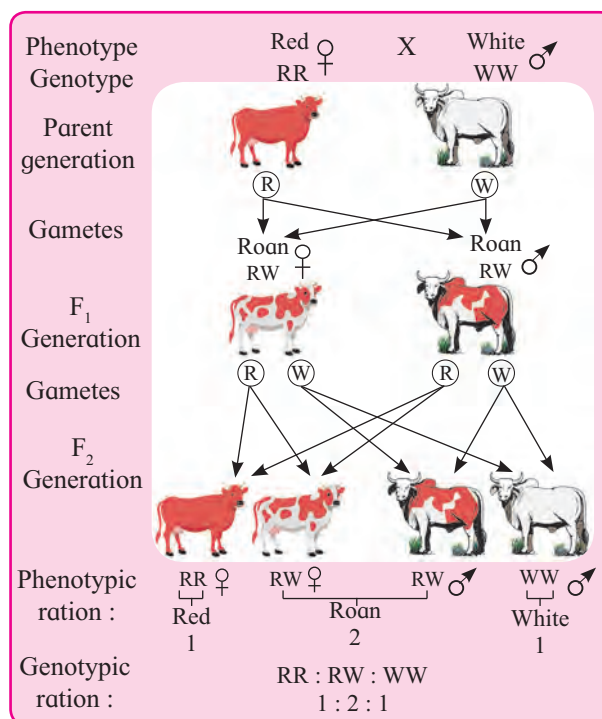


Fig. 3.2 : Representation of co-dominance in cattle

Classic example of co-dominance is coat colour in cattle. There are two types one with red coat (with red colour hair) and other with white coat (with white hair). When red cattles (RR) are crossed with white cattles (WW), F₁ hybrids (RW) are roan.





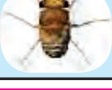
Roans have the mixture of red and white colour hair. Thus both the traits are expressed equally. In F₂ generation red (RR), roans (RW) and white (WW) are produced in the ratio 1:2:1. Thus in Co-dominance, the genotypic and phenotypic ratios are identical.

c. Multiple alleles :

More than two alternative forms (alleles) of a gene in a population occupying the same locus on a chromosome or its homologue, are known as **multiple alleles**. Multiple alleles arise by mutations of the wild type of gene. A gene can mutate several times producing a series of alternative expression. Different alleles in a series show dominant-recessive relation or may show co-dominance or incomplete dominance among themselves. Wild type is dominant over all other mutant alleles.

In *Drosophila*, a large number of multiple alleles are known. e.g. The size of wings from normal wings to vestigial (no) wings, i.e., just stumps, is due to one allele (vg) in homozygous condition. The normal wing is wild type while vestigial wing is recessive type.

Table 3.3 : Few phenotypes and genotypes in *Drosophila*

Phenotype	Genotype
Normal wings 	vg ⁺
Nicked wings 	vg ⁿⁱ
Notched wings 	vg ^{no}
Strap wings 	vg st
Vestigial wings 	vg

Curiosity Box

1. What is qualitative and quantitative inheritance?
2. Find out the traits of quantitative inheritance in humans.

Another good example of multiple alleles is A, B, O blood grouping in human beings.

d. Pleiotropy :

When a single gene controls two (or more) different non-related traits, it is called pleiotropic gene and the phenomenon is called **pleiotropy** or **pleiotropism**. The ratio is 1:2 instead of 3:1 because of the death of recessive homozygote. The disease, sickle-cell anaemia, is caused by a gene Hb^s. Normal or healthy gene Hb^A is dominant. The carriers (heterozygotes Hb^A/Hb^s) show signs of mild anaemia as their RBCs become sickle-shaped i.e. half-moon-shaped only under abnormally low O₂ concentration.

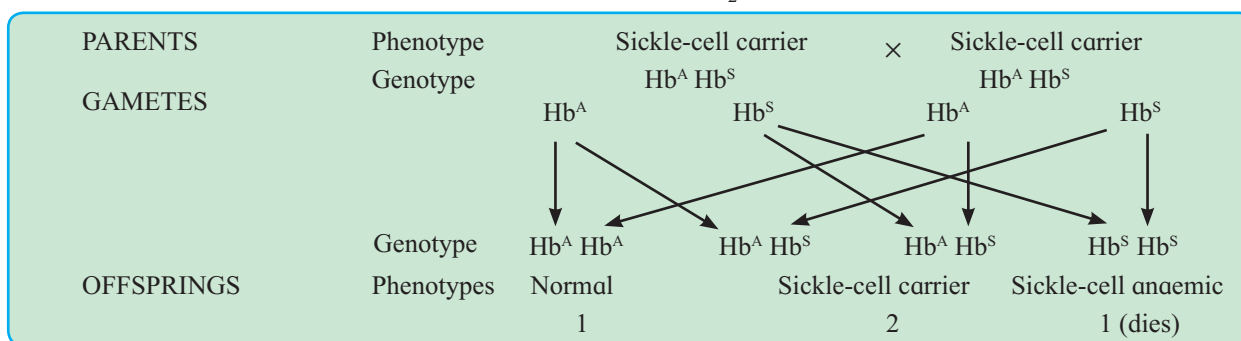


Fig. 3.4 : Representation of Pleiotropy

The homozygotes with recessive gene Hb^s however, die of total anaemia. Thus, the gene for sickle-cell anaemia is lethal in homozygous condition and produces sickle cell trait in heterozygous carrier. Two different expressions are produced by a single gene.

A marriage between two carriers will produce normal, carriers and sickle-cell anaemic children in 1:2:1 ratio. Sickle cell anaemics die leaving carriers and normals in the ratio 2:1. The heterozygotes or carriers can be identified by microscopic examination of blood.



Internet my friend

1. Find out the inheritance pattern of blood groups in human beings.
2. Find out more about pleiotropy-(Sickle-cell anaemia) and Polygenic inheritance - (human skin colour)

3.6 Chromosomal Theory of Inheritance :

Gregor Johann Mendel published his work on inheritance of traits in 1866 but for some reasons, it remained unnoticed or unrecognised till 1900, as communication was not easy in those days. His work was not widely recognized. His approach of using mathematics and statistics to explain biological phenomenon was totally new and unacceptable to the then biologists. As continuous variations were observed in nature, Mendel's concept of factors (genes) as stable and discrete unit which controlled the expression of characters, and that a pair of alleles did not "blend" with each other, was not accepted by his contemporaries. He also did not know the physical location of the 'factors' (genes) in the gametic cell.

In 1900, three scientists Hugo de Vries, Correns and von Tschermak, independently rediscovered Mendel's work on the inheritance of traits. Due to advancements in microcopy, scientists were able to observe cell division and the structure of chromosomes under microscope.

Walter Sutton along with Theodor Boveri (1903) studied the parallel behaviour of Mendel's factors (genes) and behaviour of chromosomes, at the time of meiosis.

Based on these observations, **chromosomal theory of inheritance** was put forth by Sutton and Boveri. This theory identifies chromosomes as the carriers of genetic material.

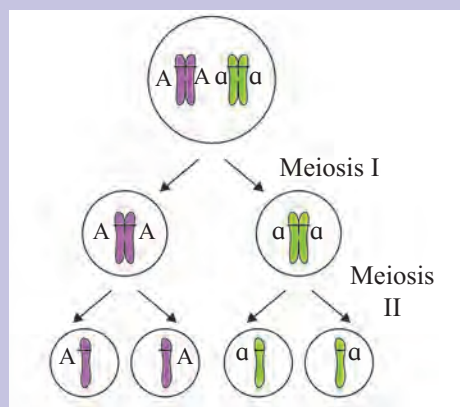
This theory states that the chromosomes are present in pairs in somatic cells. During gamete formation homologous chromosomes pair, segregate and assort independently during meiosis. Thus, each gamete contains only one chromosome from a pair.

Nucleus of gametes contains chromosomes, which carry all hereditary traits. Male and female gametes (sperms and eggs) carry all the hereditary traits. They are the link between parents and offsprings. The fusion of haploid male gamete and haploid female gamete, restores the diploid number of chromosomes of the species.



Activity :

Observe the following diagram and answer the questions given below -



Questions :

1. What is homologous chromosome?
2. In which phase of meiosis-I, homologous chromosomes segregate?
3. Where are genes located?
4. Do genes and chromosomes have similar behaviour? Justify.



Always Remember

1. Genes and chromosomes occur always in pairs in diploid organism.
2. Alleles located on chromosome segregate along with chromosome during gamete formation.



Can you recall?

1. What is chromosome?
2. How many chromosomes are present in human somatic and reproductive cell?

3.7 Chromosomes :

Chromosomes are filamentous bodies present in the eukaryotic nucleus. The term chromosomes (Gr., Chromo = colour, soma = body) was coined by W. Waldeyer (1888). The size of chromosome varies from species to species. Each metaphase chromosome varies from 0.1 to 33 μm in length and 0.2 to 2 μm in thickness. Chromosomes are visible during cell division. They are capable of self replication and play vital role in heredity, mutation, variation, and evolutionary development of eukaryotic species. Chemically eukaryotic chromosomes are made up of DNA, histone and non-histone proteins.

Function :

Chromosomes mainly act as carriers of heredity.

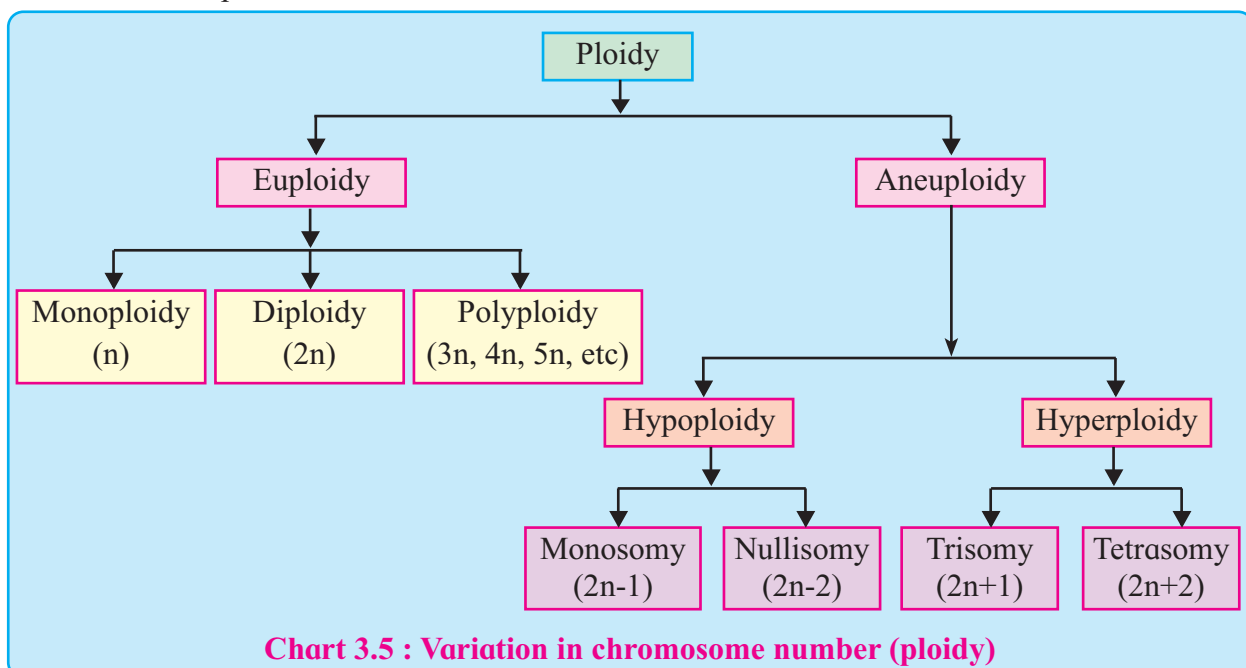
Number of chromosomes :

The number of chromosomes is specific and constant for a particular species, therefore it is of great importance in the study of phylogeny and taxonomy of the species.

The term **Ploidy** speaks for the degree of repetition of the primary basic number of chromosomes (i.e. 'x') in a cell. When the chromosome number in a cell is the exact multiple of the primary basic number, then it is called **euploidy**. Euploids include monoploid/haploid (with one set of chromosomes where $x=n$), diploids ($2n$ -two sets of chromosomes), triploids ($3n$ -three sets of chromosomes), tetraploid ($4n$ -four sets of chromosomes) and so on. When the chromosome number is not the exact multiple of the haploid set, it is described as **Aneuploidy**. Aneuploidy is either addition or deletion of one or more chromosome (s) to the total number of chromosomes in a cell (see the chart 3.5).

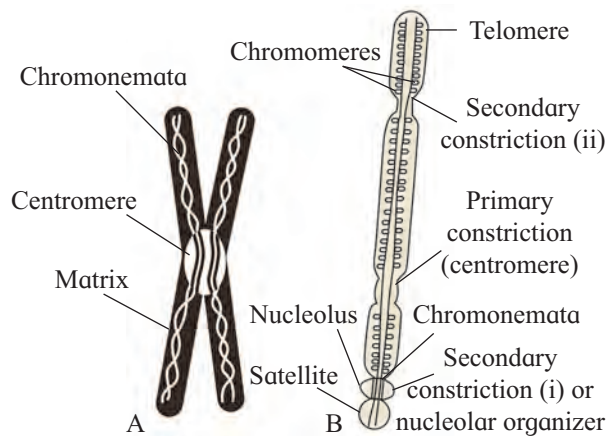
Structure of chromosome :

Chromosomes are best visible under microscope, when the cell is at metaphase stage. It is because at this stage chromosomes are highly condensed. Typical chromosome



consists of a two chromatids joined together at centromere or primary constriction. Primary constriction consists of a disc shape plate called **kinetochore**. It is at the kinetochore, spindle fibres get attached during cell division. Besides primary constriction, some few chromosomes possess additional one or two constrictions called **secondary constriction**. At secondary constriction I, nucleolus becomes organized during interphase. A satellite body (SAT body) is attached at secondary constriction II, in very few chromosomes. Each chromatid in turn contains a long, unbranched, slender, highly coiled DNA thread, called Chromonema, extending through the length of chromatid.

Chromatid consists a double stranded DNA molecule which extends from one end of chromosomes to other.



A : Parts of chromosomes B : Showing secondary constrictions and details

Fig. 3.6 : Structure of Chromosome



Activity :

Study the types of chromosome according to position of centromere. Observe and complete the following table.

Types of Chromosome	Name of Chromosome	Position of Centromere
	Metacentric	-
	-	-
	Acrocentric	-
	-	At one end

Depending upon the position of centromere there are four types (shapes) of chromosomes viz. Acrocentric (j shaped), Telocentric (i shaped), Submetacentric (L shaped) and Metacentric (V shaped). The ends of chromosome (i.e. chromatids) are known as **telomeres**.

Sex Chromosomes :

The chromosomes which are responsible for the determination of sex are known as **sex chromosomes** (Allosomes). Human being and other mammals have X and Y Chromosomes as sex chromosomes.

X chromosome is straight, rod like and longer than Y chromosome. X chromosome is metacentric, while Y chromosome is acrocentric. X chromosome has large amount of **euchromatin** (extended region) and small amount of **heterochromatin** (highly condensed region). Euchromatin has DNA in the extended state, hence it is metabolically more active than heterochromatin. Y chromosome has small amount of euchromatin and large amount of heterochromatin, hence it is genetically less active or inert. Both X and Y chromosome show homologous and non-homologous regions. Homologous regions show similar genes while non-homologous regions show dissimilar genes.

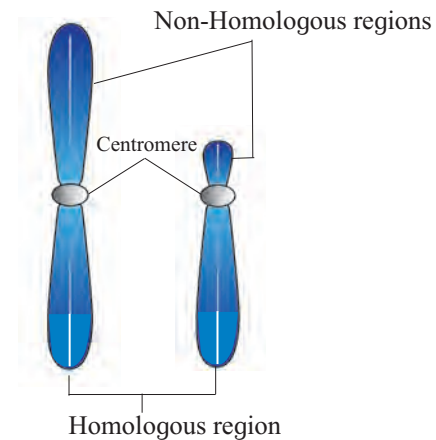


Fig. 3.7 : Structure of X and Y chromosomes (in humans)

Crossing over occurs only between homologous regions of X and Y chromosomes. Non-homologous region of X chromosome is longer and contains more genes than that of

non-homologous region of Y chromosome. X-linked genes are present on non-homologous region of X-chromosome while Y linked genes are present on non-homologous region of Y-chromosome.



Can you tell?

1. What are allosomes?
2. Compare X and Y chromosomes.
3. In which region of chromosomes does crossing over take place?

3.8 Linkage and Crossing Over :

Linkage :

It is a known fact that several genes are closely located on the chromosome. As chromosomes are carriers of heredity, these genes have tendency to be inherited together. Such genes are called **linked genes**. This tendency of two or more genes present on the same chromosomes that are inherited together is known as **linkage**. Linkage was discovered in plants by Bateson and Punnett and in animals by T. H. Morgan. Linkage is of two kinds - complete and incomplete linkage:

I. Complete linkage : The linked genes which are closely located on the chromosome do not separate (no crossing over) and inherit together. They are called completely linked (strongly linked) genes and the phenomenon of their inheritance is called complete linkage. Thus the parental traits are inherited in offsprings. e.g. X chromosome of *Drosophila* males- show complete linkage.

II. Incomplete linkage : The linked genes which are distantly located on the same chromosome and have chances of separation by crossing over, are known as incompletely linked (weakly linked) genes. The phenomenon of their inheritance, is called incomplete linkage. Thus, new traits occur in offsprings. e.g. In *Zea mays* - colour and shape of grain show incomplete linkage.

Linkage Groups :

All the linked genes in a particular chromosome, constitute a linkage group.

The number of linkage groups of a particular species corresponds to its haploid number of chromosomes. e.g. *Drosophila melanogaster* has 4 linkage groups that correspond to the 4 pairs of chromosomes. Garden pea has 7 linkage groups and 7 pairs of chromosomes.

Sex-linkage :

The transmission (inheritance) of X - linked and Y-linked genes from parents to offspring, is called **sex-linked inheritance**. Sex-linked inheritance is of three types viz. X-linked, Y-linked and XY-linked. Sex linkage is of two kinds :

a. Complete sex linkage : It is exhibited by genes located on non-homologous regions of X and Y chromosomes. They inherit together because crossing over does not occur in this region.

Examples of X-linked traits are haemophilia, red-green colour blindness, myopia (near sightedness), Ichthyosis and for Y-linked are hypertrichosis, H-Y antigen gene, etc.

b. Incomplete sex linkage : It is exhibited by genes located on homologous regions of X and Y chromosomes. They do not inherit together because crossing over occurs in this region. Examples of X-Y linked traits are total colour blindness, nephritis, retinitis pigmentosa, etc.

Crossing Over :

Crossing over is a process that produces new combinations (recombinations) of genes by interchanging and exchanging of corresponding segments between non-sister chromatids of homologous chromosomes. It occurs during pachytene of prophase I of meiosis. The term crossing over was coined by Morgan. The mechanism of crossing over consists of four sequential steps such as synapsis, tetrad formation, crossing over and terminalization. This you have already studied in the chapter on cell division in class XI. The phenomenon of crossing over is by and large universal (except in ♂ *Drosophila*). It is necessary for the natural selection, because it increases the chances of variation.



Use your brain power

How many linkage groups are in human being and maize?



Know the Scientist :



T. H. Morgan
(1866-1945)

Thomas Hunt Morgan was an American biologist. He used the fruit fly (*Drosophila melanogaster*) in genetic research and established the chromosomal theory of heredity. He also discovered the principle of linkage, sex linkage and crossing over.

Morgan's work played a key role in the field of genetics. He was awarded the Nobel Prize in 1933, for Physiology and Medicine.

• Morgan's Experiments showing linkage and crossing over :

Morgan used *Drosophila melanogaster* (fruit fly) for his experiments because, *Drosophila* can easily be cultured in laboratory. It's life span is short, about two weeks. Moreover, it has high rate of reproduction.

Morgan carried out several dihybrid cross experiments in fruit fly to study genes that are sex-linked. The crosses were similar to dihybrid crosses, as carried out by Mendel in Pea. For example, Morgan and his group crossed yellow-bodied, white eyed female to the wild type with brown-bodied, red eyed males and intercrossed their F₁ progeny.

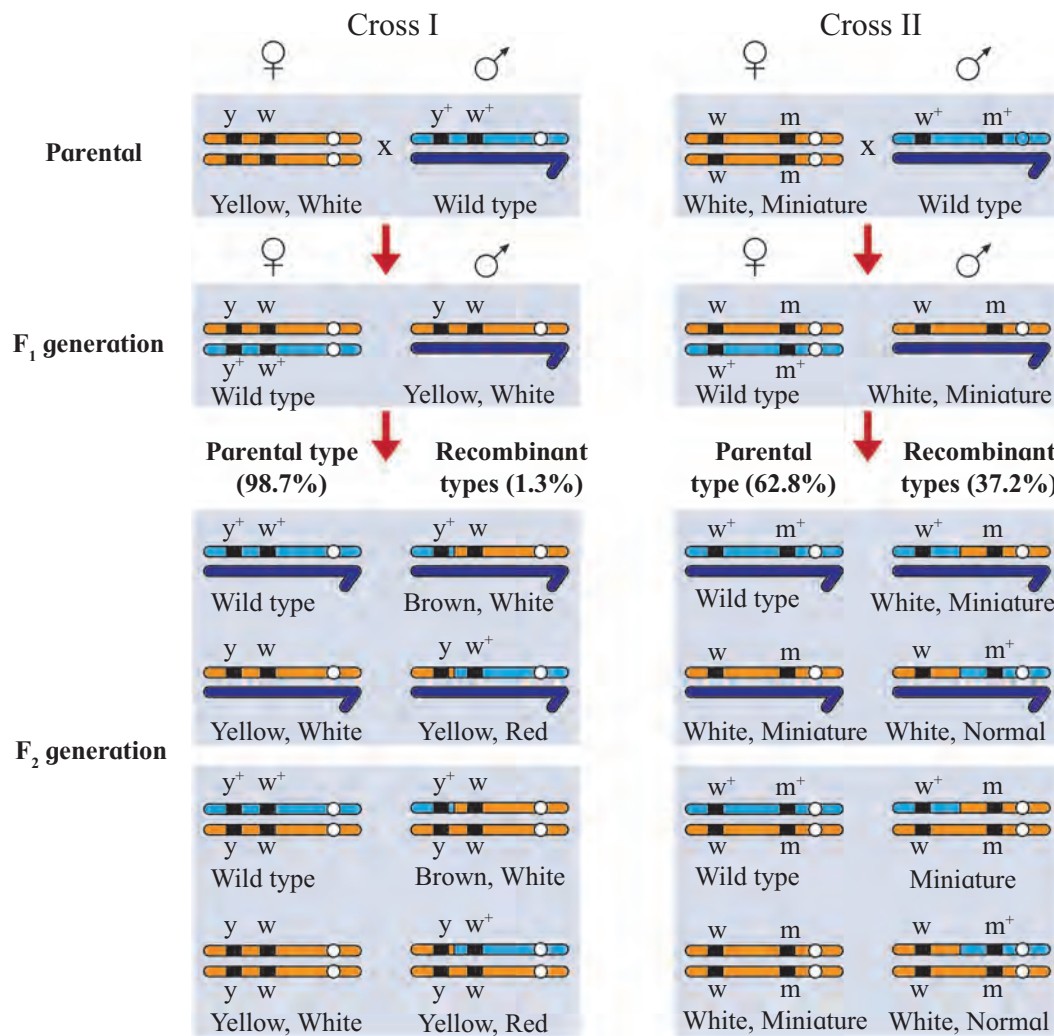


Fig : 3.8 : Linkage and crossing over

He observed that the two genes did not segregate independently of each other and F_2 ratio deviated very significantly from 9:3:3:1 ratio.

Morgan and his group knew that the genes were located on X chromosome and stated that when two genes in a dihybrid cross are situated on the same chromosome, then the proportion of parental combination is much higher than non-parental type. This occurs due to physical association or linkage of the two genes. He also found that, when genes are grouped on the same chromosome, some genes are strongly linked. They show very few recombinations (1.3 %). When genes are loosely linked i.e. present far away from each other on chromosome, they show more (higher) recombinations (37.2 %).

For example, the genes for yellow body and white eye were strongly linked and showed only 1.3 percent recombination (in cross-I). White bodied and miniature wings showed 37.2 percent recombination (in cross-II). Cross I shows crossing over between genes y and w . Cross II shows crossing over between genes white (w) and miniature wing (m). Here dominant wild type alleles are represented with (+) sign.



Always Remember

Parental combinations occur more due to linkage and new combinations less due to crossing over.

3.9 Autosomal Inheritance :

Human somatic ($2n$) cell contains 23 pairs of chromosomes. They can be divided functionally as autosomes and sex chromosomes. A single pair of chromosomes is involved in sex determination and remaining 22 pairs are called autosomes. Autosomes control a variety of traits other than sex. These traits are called autosome linked traits. Transmission of body characters other than the sex linked traits from parents to their offsprings through autosomes, is called **autosomal inheritance**.

Some characters are influenced by dominant genes while some other are by recessive genes, present on autosomes. For example,

- Autosomal dominant traits like Widow's peak and Huntington's disease, etc.
- Autosomal recessive traits like Phenylketonuria (PKU), Cystic fibrosis and Sickle cell anaemia.

a. Widow's peak :

A prominent "V" shaped hairline on forehead is described as widow's peak. It is determined by autosomal dominant gene. Widow's peak occurs in homozygous dominant (WW) and also heterozygous (Ww) individuals.

Individuals with homozygous recessive (ww) genotype have a straight hair line (no widows peak). Both males and females have equal chance of inheritance.

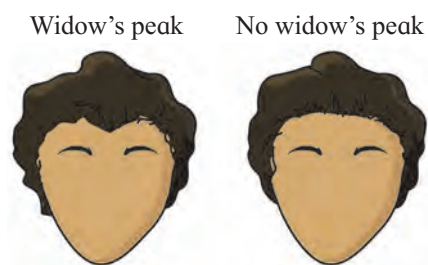


Fig. 3.9 : Widow's peak and straight hair line

b. Phenylketonuria (PKU):-

It is an inborn metabolic disorder caused due to recessive autosomal genes. When recessive genes are present in homozygous condition, phenylalanine hydroxylase enzyme is not produced. This enzyme is essential for conversion of amino acid phenylalanine into tyrosine. Due to absence of this enzyme, phenylalanine is not converted into tyrosine. Hence, phenylalanine and its derivatives are accumulated in blood and cerebrospinal fluid (CSF). It affects development of brain and causes mental retardation. Excess phenylalanine is excreted in urine, hence this disease is called phenylketonuria.

Autosomal recessive traits appear in both sexes with equal frequency. These traits tend to skip generations.

3.10 Sex Linked Inheritance :

Genes located on non-homologous region of sex chromosomes, are called sex-linked genes. The traits that are determined by sex linked genes, are called sex-linked traits.

The inheritance of sex linked genes from parents to their offsprings, is called sex linked inheritance. There are two types of sex-linked genes as X-linked genes and Y-linked genes.

a. X-linked (sex linked) genes :

The X linked genes are located on non homologous region of X chromosome and these gene do not have corresponding alleles on Y chromosome.

Female has two X chromosomes. In female two recessive sex linked genes are required for expression of sex linked traits. If one X chromosome carries a recessive gene for sex-linked trait (defect) its effect is suppressed by the dominant gene present on other X chromosome. The females with one recessive gene are carriers. The carrier female is physically normal as she does not suffer from the disease (disorder).

Male has only one X-chromosome. If X chromosome carries X-linked recessive gene for sex linked trait, then it is expressed phenotypically, because there is no dominant gene on Y chromosome to suppress its effect. Therefore, sex-linked / X-linked traits appear more frequently in males than in the females. Examples of X-linked traits include haemophilia, colour blindness, night blindness, myopia, muscular dystrophy, etc.

b. Y-linked (Holandric) genes :

Genes located on non-homologous region of Y chromosome, are called Y linked genes. The Y-linked genes are inherited directly from male to male. In man, the Y-linked genes such

as hypertrichosis is responsible for excessive development of hair on pinna of ear. This character is transmitted directly from father to son.



Internet my friend

Collect information of Ishihara's Test for colour blindness.

Colour blindness :

Colour blindness is X-linked recessive disorder where person is unable to distinguish between red and green colours as both the colours appear grey. It is caused due to recessive X-linked genes (X^c) which prevents formation of colour sensitive cells, the cones, in the retina of eye.

The homozygous recessive females ($X^c X^c$) and hemizygous recessive male ($X^c Y$) are unable to distinguish between red and green colours. The frequency of colour blind women is much less than colour blind men. Dominant X linked gene (X^C) is necessary for formation of colour sensitive cells in the retina of eye. Thus, genotypes of male and female individuals can be represented as follows-

Sex	Normal	Colourblind	Carrier
Male	$X^C Y$	$X^c Y$	—
Female	$X^C X^C$	$X^c X^c$	$X^C X^c$

The inheritance of colourblindness can be studied in the following two types of marriages:-

1. Marriage between colour blind male with normal female, will produce normal visioned male and female offspring in F_1 . The sons have normal vision but daughter will be carrier for the disease.

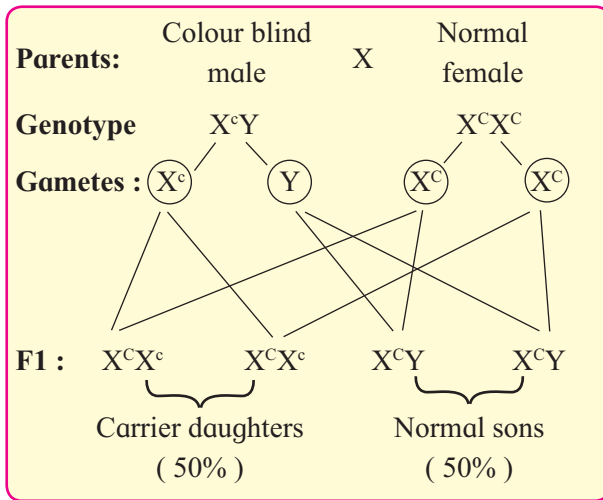


Fig. 3.10 : Sex linked inheritance (colour blindness)

2. Marriage between carrier female (daughter) and normal male will produce female offsprings with normal vision but half of them will be carriers for the disease. Half of male offsprings will be normal while remaining half will be colour blind.

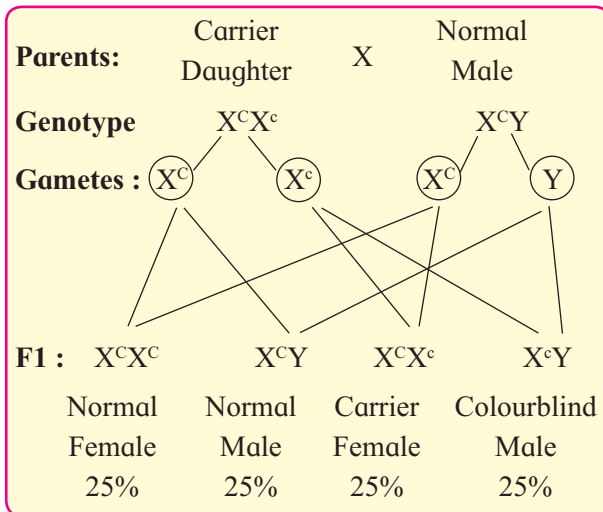


Fig. 3.11 : Sex linked inheritance (colour blindness)

From above example, it is clear that the X linked recessive gene for colour blindness is inherited from colourblind father to his grandson through his daughter. This type of inheritance is called as criss-cross inheritance.

Haemophilia (Bleeder's disease) :

Haemophilia is X-linked recessive disorder in which blood fails to clot or coagulates very slowly. The genes for normal clotting

are dominant over the recessive genes for haemophilia. The person having recessive gene for haemophilia is deficient in clotting factors (VIII or IX) in blood. Even minor injuries cause continuous bleeding, hence haemophilia is also called as bleeder's disease.

The recessive gene for haemophilia is located on non homologous region of X chromosome. As there is no corresponding allele on Y chromosome to suppress its expression, so men suffer from this disease. Women suffers only when both X chromosomes have recessive genes (alleles).

The genotype of male and female individuals can be represented as follow-

Sex	Normal	Haemophilic	Carrier
Male	X^HY	X^hY	—
Female	X^HX^H	X^hX^h	X^HX^h

Like colour blindness, haemophilia also shows criss-cross inheritance. The inheritance of haemophilia can be studied with the help of following examples -

1. Marriage between the Haemophilic male and normal female.

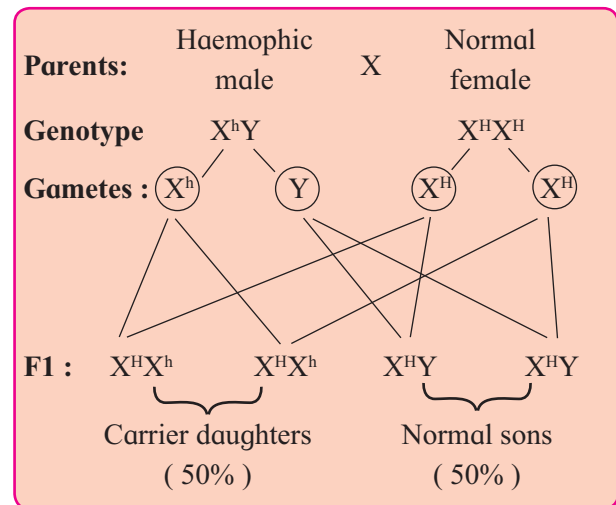


Fig. 3.12 : Sex linked inheritance (Haemophilia)

2. Marriage between carrier female (daughter) and normal male.

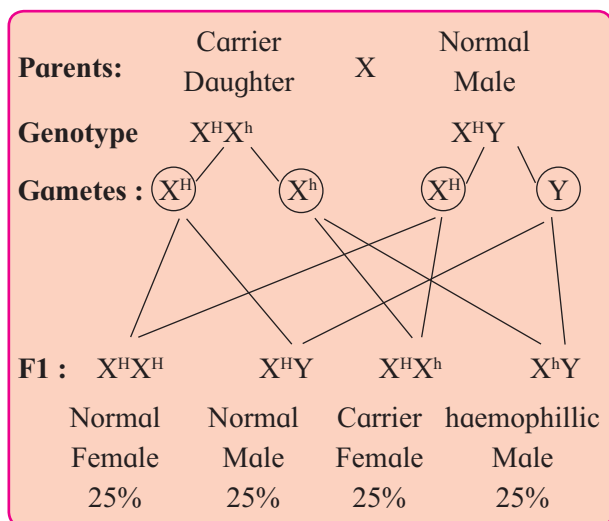


Fig. 3.13 : Sex linked inheritance (Haemophilia)



Do you know ?

Haemophilia is also referred as “The royal disease”, because it affected the royal families of England, Germany, Russia and Spain in the 19th and 20th centuries. Queen Victoria of England, who ruled from 1837-1901, was believed to have been the carrier of haemophilia. She passed the trait on to her three of nine children.



Find out

Aarya shows normal blood clotting but her mother is haemophilic. Ramesh shows normal blood clotting but his father is haemophilic. If Ramesh and Aarya were to marry, then find out the possible phenotypes of their offsprings.

3.11 Sex Determination :

The mechanism by which sex is established is termed as **sex determination**. The term sex refers to sexual phenotype. In some species, both male and female reproductive organs are present in same organism. It is described as **bisexual** or **hermaphrodite** or **monoecious**.

On the other hand, some species in which the organism has either male or female reproductive organs, is said to be **dioecious** or **unisexual**. Humans are dioecious.

German biologist, **Henking** in 1891, while studying spermatogenesis of the squash bug (*Anasa tristis*), noted that 50% of sperms receive the unpaired chromosomes while other 50% sperm do not receive it. Henking gave a name to this structure as the x-body but he could not explain its role in sex determination. Further investigations by other scientists led to conclusion that the “**x-body**” of Henking was infact a chromosome and gave the name ‘X-Chromosome’.

a. Sex Determination in human beings :

The chromosomal mechanism of sex determination in human beings is XX-XY type. In human beings, the nucleus of each somatic cell contains 46 chromosomes or 23 pairs of chromosomes. Out of these, 22 pairs are **autosomes** and one pair of **sex chromosomes**.

Human female has a pair of XX, homomorphic sex chromosomes while male has XY, heteromorphic sex chromosomes.

Thus genotype of :

Female = 44 Autosomes + XX

Male = 44 Autosomes + XY

During gamete formation in male, the diploid germ cells in testis undergo spermatogenesis to produce two types of haploid sperms, 50% sperms contain 22 autosomes and X chromosome while, 50% sperms contain 22 autosomes and Y chromosome.

In Female, the diploid germ cells in ovaries undergo oogenesis to produce only one type of egg. All eggs contain 22 autosomes and X chromosome. Thus human male is heterogametic and female is homogametic.

If sperm containing X chromosome fertilizes egg (ovum), then diploid zygote is formed, that grows into a female child. If

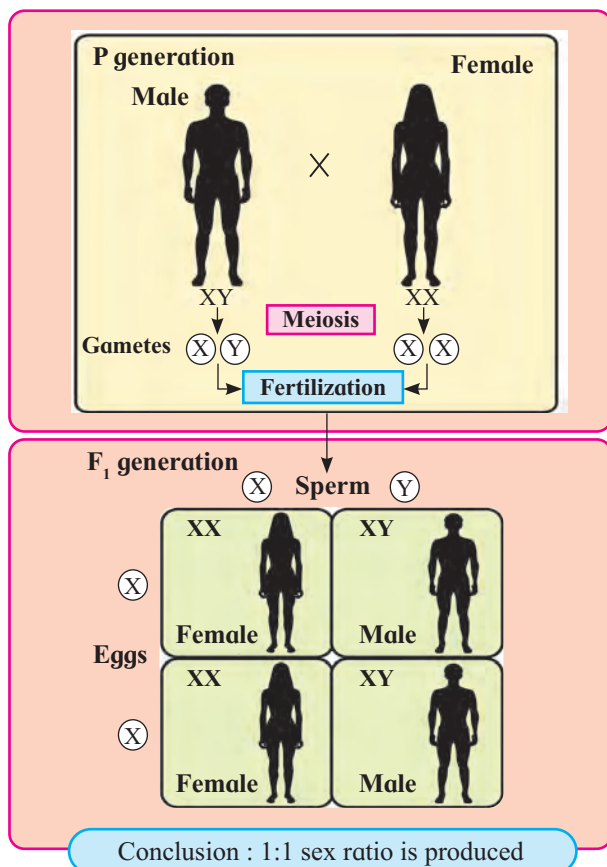


Fig. 3.14 : Sex determination in human beings

sperm containing Y chromosome fertilizes the egg, then diploid zygote is formed that grows into a male child.

This indicates that the sex of a child depends on the type of sperm fertilizing the egg and **hence the father is responsible for determination of sex of child and not the mother**. Due to lack of knowledge, women are often blamed for giving birth to female child.

b. Sex Determination in birds :

In birds, the chromosomal mechanism of sex determination is ZW-ZZ type. In this type females are heterogametic and produce two types of eggs; 50% eggs carry Z- chromosome, while 50% eggs carry W- chromosome.

Males are homogametic and produce one type of sperms. Each sperm carries a Z- chromosome. Thus sex of individual depends on the kind of egg (ova) fertilized by the sperm.

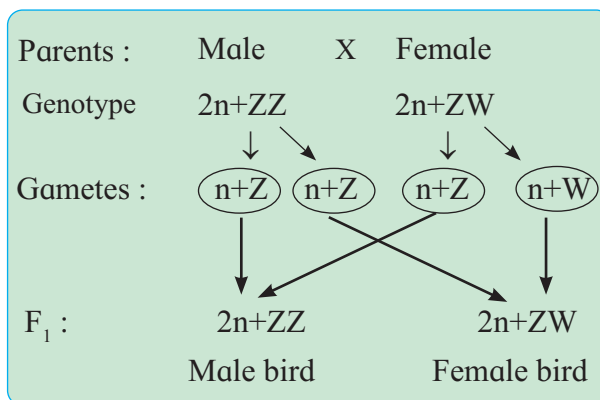


Fig. 3.15 : Sex determination in birds

Something Interesting

In *Bonellia viridis*, the environmental factors determine the sex of individual. The sex of worm *Bonellia viridis* depends on which location the *Bonellia* larva gets settled.

The marine female *Bonellia* worm has about 10 cm long body. She has a proboscis that can extend over a meter in length. If a *Bonellia* larva settles on the seafloor, it becomes a female.

However when, a larva lands on a female's proboscis and enters the female's mouth, it migrates into her uterus and differentiates into a male. Male lives as parasite in uterus of female fertilizing her eggs.



c. Sex Determination in honey bees :

In honey bees, chromosomal mechanism of sex determination is **haplo-diploid type**. In this type, sex of individual is determined by the number of set of chromosomes received. Females are diploid ($2n=32$) and males are haploid ($n=16$). The female produces haploid eggs ($n=16$) by meiosis and male produces haploid sperms ($n=16$) by mitosis. If the egg is fertilized by sperm, the zygote develops into

a diploid female ($2n=32$) (queen and worker) and unfertilised egg develops into haploid male ($n=16$) (Drone) by way of parthenogenesis.

The diploid female gets differentiated into either worker or queen depending on the food they consume during their development. Diploid larvae which get royal jelly as food develops into queen (fertile female) and other develops into workers (sterile females).

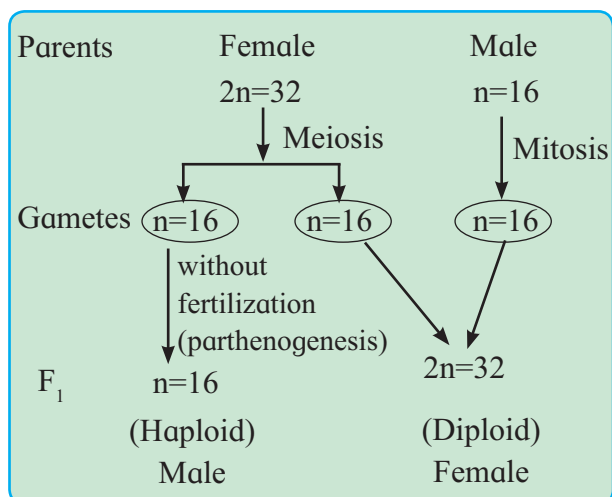


Fig. 3.16 : Sex determination in honey bee

3.12 Genetic Disorders :

Genetic Disorders are broadly grouped into two categories as, **Mendelian disorders** and **chromosomal disorders**, Mendelian disorders are mainly caused due to alteration or mutation in the gene. e.g. thalassemia, sickle-cell anaemia, colourblindness, haemophilia, phenylketonuria, etc. On the other hand, chromosomal disorders are caused due to absence or excess of one or more chromosomes or their abnormal arrangement. For eg, Down's syndrome, Turner's syndrome, Klinefelter's syndrome etc.

Thalassemia :

Thalassemia is an autosomal, inherited recessive disease. Haemoglobin molecule is made of four polypeptide chains- 2 alpha (α) and 2 beta (β) chains. The synthesis of alpha chains are controlled by two closely linked genes (HBA1 and HBA2) on chromosome 16 while the synthesis of beta chain is controlled by a single gene (HBB) on chromosome 11.

Depending upon which chain of haemoglobin is affected, thalassemia is classified as alpha-thalassemia and beta-thalassemia. It is caused due to deletion or mutation of gene which codes for alpha (α) and beta (β) globin chains that result in abnormal synthesis of haemoglobin. In Thalassemia, person shows symptoms like anaemia, pale yellow skin, change in size and shape of RBCs, slow growth and development, dark urine, etc. Massive blood transfusion is needed to these patients. Thalassemia differs from sickle-cell anaemia. The former is a qualitative problem of synthesising few globin molecule, while the latter is a qualitative problem of synthesising an incorrectly functional globin.

Down's Syndrome (21st trisomy) :

Down's syndrome is named after the physician John Langdon Down who first described this autosomal chromosomal disorder in 1866.



Fig. 3.17 : Down's Syndrome

This Syndrome is caused due to an extra copy of chromosome number 21st. It shows presence of three copies of 21st chromosome instead of homologous pair. These individuals will have 47 chromosomes instead of the normal number 46. 21st Trisomy occurs due to non-disjunction or failure of separation of chromosomes (**autosomes**) during gamete formation. The incidence of non-disjunction is distinctly higher in mothers who are over 45 years old.

These patients have mild or moderate mental retardation and skeletal development is poor. Distinct facial features like small head, ears and mouth, face is typically flat and rounded with flat nose, open mouth and protruding tongue, eyes slant up and out with internal epicanthal folds, flat hands and stubby fingers and palm is broad with single palmer crease.

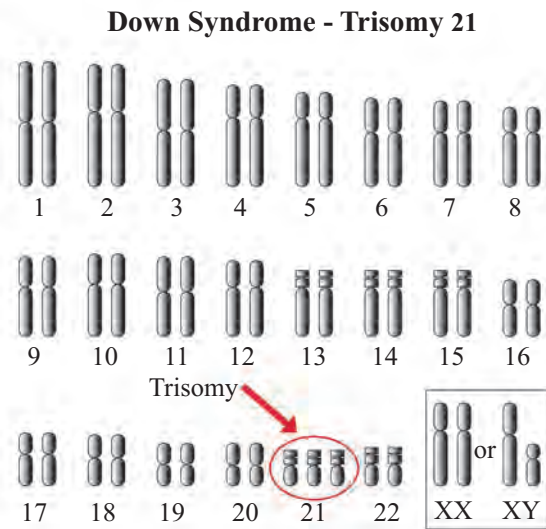


Fig. 3.18 : Karyotype of Down's syndrome

Turner's Syndrome :

(X monosomy / XO females)

It is sex chromosomal disorder caused due to non-disjunction of chromosome during formation of egg. Individual born with Turner's syndrome has 44 autosomes with XO. They are phenotypically female. They have a short stature (height) and webbed neck, lower posterior hair line, broad shield-shaped chest, poorly developed ovaries and breast, and low intelligence.

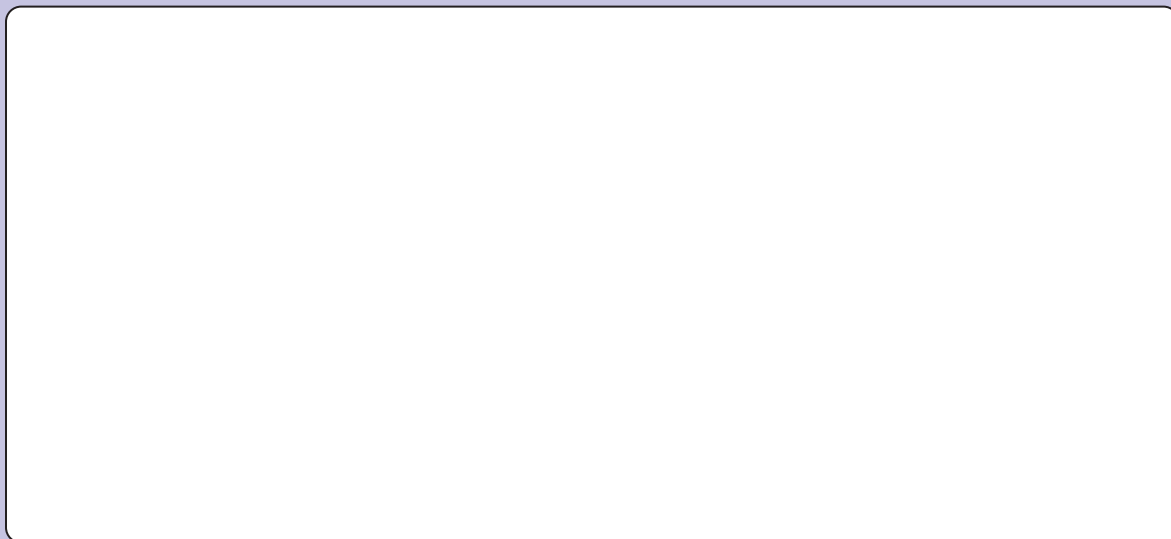
Klinefelter's syndrome (XXY males) :

It is a chromosomal disorder caused due to extra X chromosome in males. Thus genotype of individuals is 44 + XXY. They are described as feminized males. Extra chromosome is a result of non-disjunction of X-chromosome during meiosis in the formation of ovum. Such individual is male and has over all masculine development. Voice pitch is harsh and has under developed testis. They are tall with long arms, showing feminine development (development of breast i.e. Gynaecomastia) and no spermatogenesis, therefore, individuals are sterile.



Activity :

Study the complementary and supplementary interaction (digenic interactions) - both in plants and animals



Exercise

Q. 1 Multiple choice questions.

- F_2 Phenotypic ratio of incomplete dominance in *Mirabilis jalapa*.
a. 2 : 1 : 1 b. 1 : 2 : 1
c. 3 : 1 d. 2 : 2
- In dihybrid cross, F_2 generation offsprings show four different phenotypes while the genotypes are
a. six b. nine
c. eight d. sixteen
- A cross between an individual with unknown genotype for a trait with recessive plant for that trait is
a. back cross b. reciprocal cross
c. monohybrid cross d. test cross
- When F_2 phenotypic and genotypic ratios are the same, then it is an example of
a. incomplete dominance
b. complete dominance
c. Multiple alleles
d. cytoplasmic inheritance
- If the centromere is situated near the end of the chromosome, the chromosome is called
a. Metacentric b. Acrocentric
c. Sub-Metacentric d. Telocentric
- Chromosomal theory of inheritance was proposed by
a. Sutton and Boveri
b. Watson and Crick
c. Miller and Urey
d. Oparin and Halden
- If the genes are located in a chromosome as p-q-r-s-t, which of the following gene pairs will have least probability of being inherited together?
a. p and q b. r and s
c. s and t d. p and s

- Find the mis match pair :-
a. Down's syndrome = 44 + XY
b. Turner's syndrome = 44 + XO
c. Klinefelter syndrome = 44 + XXY
d. Normal female = 44 + XXX
- A colourblind man marries a woman, who is homozygous for normal colour vision, the probability of their son being colourblind is –
a. 0% b. 25%
c. 50% d. 100%

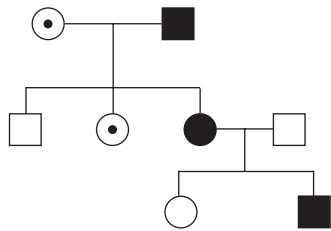
Q. 2 Very Short Answer Questions.

- Explain the statements :
a. Test cross is a back cross but back cross is not necessarily a test cross.
b. Law of dominance is not universal.
- Define the following terms :
a. Dihybrid cross b. Homozygous
c. Heterozygous d. Test cross
- What is allosome?
- What is crossing over?
- Give one example of autosomal recessive disorder.
- What are X-linked genes?
- What are holandric traits?
- Give an example of chromosomal disorder caused due to non-disjunction of autosomes.
- Give one example of complete sex linkage?

Q. 3 Short Answer Questions.

- Enlist seven traits of pea plant selected/ studied by Mendel.
- Why law of segregation is also called the law of purity of gametes?
- Write a note on pleiotropy.
- What are the reasons of Mendel success?

5. "In humans, father is responsible for determination of sex of child and not the mother". Justify.
6. What is linkage? How many linkage groups do occur in human being?
7. Write note on -PKU
8. Compare - X-chromosome and Y-chromosome.
9. Explain the chromosomal theory of inheritance.
10. Observe the given pedigree chart and answer the following questions.



- a. Identify whether the trait is sex linked or autosomal.
- b. Give an example of a trait in human beings which shows such a pattern of inheritance.

Q. 4 Match the column-I with column-II and re-write the matching pairs.

Column-I	Column-II
1. 21 trisomy	a. Turner's syndrome
2. X-monosomy	b. Klinefelter's syndrome
3. Holandric traits	c. Down's syndrome
4. Feminized male	d. Hypertrichosis

Q. 5 Long answer type questions.

1. What is dihybrid cross? Explain it with the suitable example and checker board method.
2. Explain with the suitable example an independent assortment.
3. Define test cross and explain its significance.
4. What is parthenogenesis? Explain the haplo-diploid method of sex determination in Honey bee.
5. In the answer for inheritance of X-linked genes, Madhav had shown carrier male. His answer was marked incorrect. Madhav was wondering why his marks were cut. Explain the reason.
6. With the help of neat labelled diagram, describe the structure of chromosome.
7. What is criss-cross inheritance? Explain with suitable example.
8. Describe the different types of chromosomes.

Project :

Study the genetic traits like Rolling of tongue or Widow's peak in your class and write your own observations.



Can you recall?

1. What is nucleic acid?
2. What are the types of nucleic acid?
3. What are the functions of nucleic acid?
4. What is the difference between DNA in prokaryotes and eukaryotes?

4.1 The Discovery of DNA:

Modern understanding of DNA has evolved from the discovery of nucleic acid to the development of the double-helix model. In 1869, Friedrich Miescher began working with white blood cells which are the major component of pus from infections. He collected a lot of pus from bandages at the local hospital. He used a salt solution to wash the pus off the bandages. When he added a weak alkaline solution to the cells, the cells lysed and nuclei precipitated out of the solution. From the cell nuclei, he isolated a unique chemical substance which he called **nuclein**. Chemically, nuclein has high phosphorus content. Moreover it showed acidic properties. Hence it was named as nucleic acid.

By the early 1900s, we knew that Miescher's nuclein was a mix (mixture) of proteins and nucleic acids. There are two kinds of nucleic acids. DNA (deoxyribonucleic acid) and RNA (ribonucleic acid).

4.2 The Genetic Material is DNA:

By the early 1900s, geneticist knew that genes control the inheritance of traits, that genes are located on chromosomes and that chemically chromosomes are composed mainly of DNA and proteins. Initially, most geneticists thought that proteins are large, complex and varied molecules and store information needed to govern cell metabolism. Hence it was assumed that proteins caused the variations observed within species.

On the other hand the DNA thought to be small, simple molecule whose composition varied little among species. Over the time, these ideas about DNA were shown to be wrong. In fact DNA molecules are large and vary tremendously within and among species. Variations in the DNA molecules are different than the variation in shape, electrical charge and function shown by proteins so it is not surprising that most researchers initially favored proteins as the genetic material.

Over a period of roughly 25 years (1928-1952), geneticists became convinced that DNA and not protein, was the genetic material. Three important contributions that helped cause this shift of opinion.

Griffith's experiments :

In 1928, a British medical officer Frederick Griffith performed an experiment on bacterium *Streptococcus pneumoniae* that causes pneumonia in humans and other mammals. Griffith used two strains or two genetic varieties of *Streptococcus* to find a cure for pneumonia, which was a common cause of death at that time. The two strains used were :

- i. Virulent, smooth, pathogenic and encapsulated S type.
- ii. Non-virulent, rough, non-pathogenic and non-capsulated R type.

Griffith conducted four experiments on these bacteria. First, when he injected bacteria of strain R to mice, the mice survived because it did not develop pneumonia. Second, when he injected bacteria of strain S to mice, the mice developed pneumonia and died. In the third experiment, he injected heat-killed strain S bacteria to mice, once again the mice survived. In fourth experiment, he mixed heat-killed S bacteria with live bacteria of strain R and injected to mice. The mice died and Griffith

recovered large numbers of live strain S bacteria from the blood of the dead mice.

In these four experiments, something had caused harmless strain R bacterium to change into deadly S strain bacterium. Griffith showed that the change was genetic. He suggested that genetic material from heat-killed strain S bacterium had somehow changed the living strain R bacterium into strain S bacterium.

Griffith concluded that the R-strain bacterium must have taken up, to what he called a "transforming principle" from the heat-killed S bacterium, which allowed R strain to get transformed into smooth-coated bacterium and become virulent.

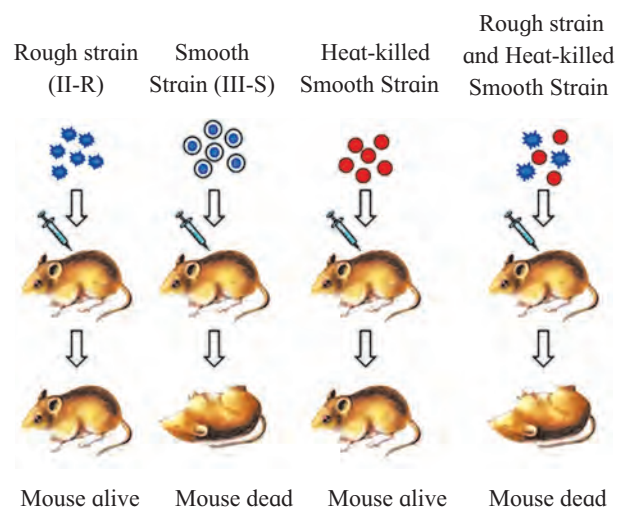


Fig. 4.1 : Griffith's experiment

Avery, McCarty and MacLeod's experiment:

In 1944, after some 10 years of research and experimentation, U. S. microbiologists Oswald T. Avery, Colin M. MacLeod and Maclyn McCarty (all at Rockefeller University in New York) first evidenced to prove the DNA is a genetic material (transforming principle), through the experiments. They purified DNA, RNA, Proteins (enzymes) and other materials from cell free extract of S cells/ strain and mixed with heat killed S strain and R cells separately to confirm which one could transform living R cells into S cells. Only DNA was able to transform harmless strain R into deadly strain S.

They also discovered that protein-digesting enzymes (proteases), RNA-digesting enzymes (RNases) did not affect transformation, so the transforming substance was neither a protein nor RNA. DNA digested with DNase did inhibit the transformation, suggesting that DNA caused the transformation. These experiments proved that the transforming principle is DNA but all biologists were not convinced.

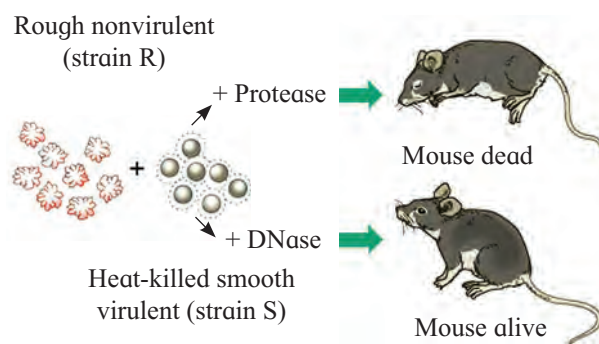


Fig. 4.2 : DNA transforms bacteria

Finally, Alfred Hershey and Martha Chase (1952) proved that DNA is the genetic material and not proteins, by using bacteriophages.

Hershey - Chase Experiment:

Hershey and Chase worked with viruses that infect bacteria i.e. bacteriophages, which are composed of DNA and protein. They used radioactive phosphorous ^{32}P in the medium for some viruses and radioactive sulphur ^{35}S for some others.

They grew some viruses on a medium that contained radioactive phosphorus and some others on medium that contained radioactive sulphur. Viruses grown in the presence of radioactive phosphorus contained radioactive DNA (labelled DNA), but not radioactive proteins because DNA contains phosphorus (labelled DNA) but proteins do not. Similarly, viruses grown on radioactive sulphur contained radioactive protein but not radioactive DNA because DNA does not contain sulphur.

Radioactive phages were allowed to infect *E.coli* bacteria grown on the medium containing normal 'P' and 'S'. Then, as

the infection proceeded, the viral coats were removed with the help of centrifuge. Bacteria which were infected by viruses with radioactive DNA, were radioactive, indicating that DNA was the material that passed from the viruses to the bacteria. Bacteria which were infected by viruses having radioactive sulphur (protein) were not radioactive. This indicates that proteins from the viruses, did not enter the bacteria. DNA is, therefore, the genetic material that is passed from virus to bacteria (fig. 4.3).

In other words, sometime after infection, radioactivity for 'P' and 'S' was tested. Only radioactive 'P' was found inside the bacterial cell, indicating that DNA is the genetic material.

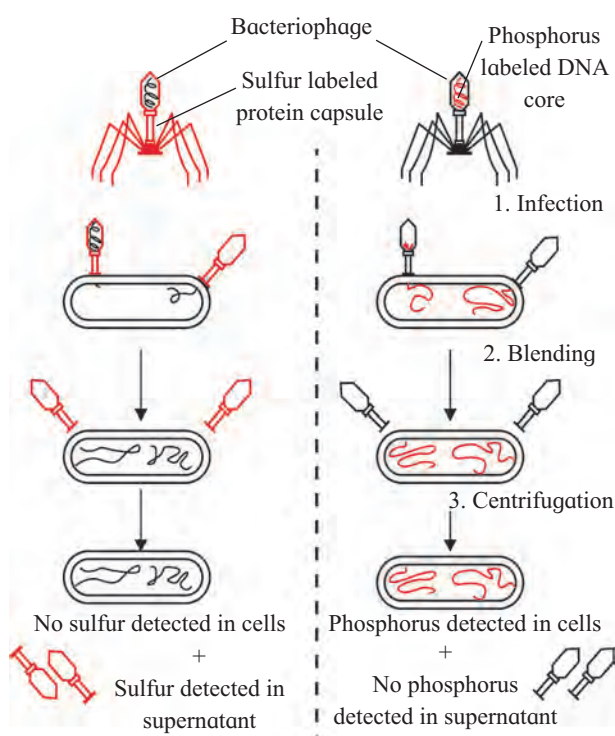


Fig. 4.3 : Hershey - Chase Experiment

Do you know ?

1. Multiple forms of DNA and their differences.
2. Significance of different forms of DNA.



Can you recall?

1. What are the chemical components of DNA?
2. What is a chromosome?

4.3 DNA packaging :

Length of DNA double helix molecule, in a typical mammalian cell is approximately 2.2 meters. (This can be worked out simply by multiplying the total number of base pairs with distance between the consecutive base pairs). Approximate size of a typical nucleus is 10^{-6} m. How this long DNA molecule can be then accommodated in such a small nucleus? It, therefore, must be condensed, coiled and super coiled to fit inside such small nucleus.

Packaging in Prokaryotes:

In prokaryotes like *E. coli*, cell size is almost $2-3\mu$ long. They do not have well organized nucleus. It is without nuclear membrane and nucleolus. The nucleoid is small, circular, highly folded, naked ring of DNA which is 1100μ long in perimeter, containing about 4.6 million base pairs.

The 1100μ long (approximately 1.1 mm, if cut and stretched out) nucleoid is to be fitted or packaged into a cell which is hardly $2-3\mu$ long. Hence the negatively charged DNA becomes circular, reducing the size to 350μ in diameter. This is further reduced to 30μ in diameter because of folding/ looping. 40-50 domains (loops) are formed. Formation of loops is assisted by RNA connectors. Each domain is further coiled and supercoiled, thereby reducing the size down to 2μ in diameter. This coiling (packaging) is assisted by positively charged HU (Histone like DNA binding proteins) proteins and enzymes like DNA gyrase and DNA topoisomerase I, for maintaining super coiled state.

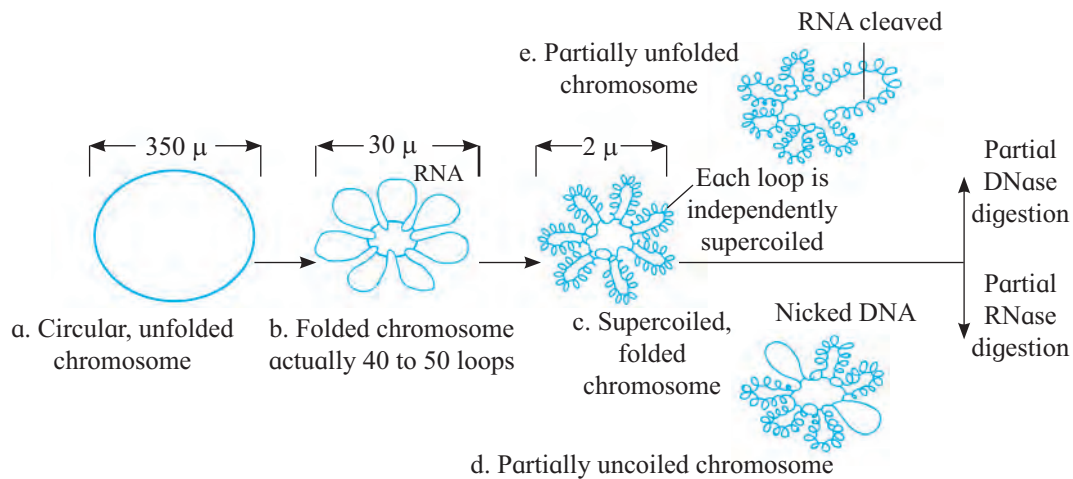


Fig. 4.4 : DNA Packaging in Prokaryotes

Packaging in Eukaryotes:

Eukaryotes show well organized nucleus with nuclear membrane, nucleolus and thread-like material in the form of chromosomes. In the chromosomes, DNA is associated with histone and non-histone proteins as was reported by R. Kornberg in 1974. The organization of DNA is much more complex in eukaryotes. Depending upon the abundance of amino acid residues with charged side chains, a protein acquires its charge. Histones are the proteins that are rich in lysine and arginine residues. Both these amino acid residues are basic amino acids and carry positive charges with them. So, histones are a set of positively charged, basic proteins (histones + protamine). These histones organize themselves to make a unit of 8 molecules known as **histone octamer**.

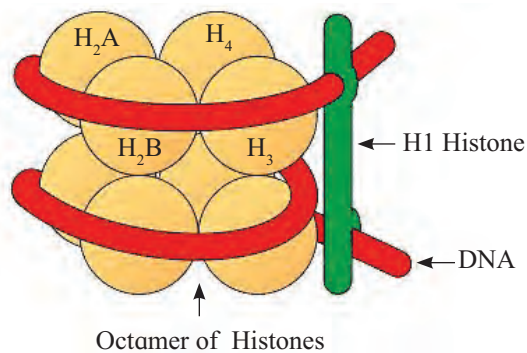


Fig. 4.5 : Nucleosome

The negatively charged helical DNA is wrapped around the positively charged histone octamer, forming a structure known as **nucleosome**. The nucleosome core is an octamer made up of two molecules of each of four types of histone proteins viz. H₂A, H₂B, H₃ and H₄. H₁ protein binds the DNA thread where it enters (arrives) and leaves the nucleosome.

One nucleosome approximately contains 200 base pair long DNA helix of which about 146 bp long segment is wound around each octamer and the remaining bp contribute as linker DNA (Fig. 4.5). Nucleosomes are the repeating units of chromatin, which are thread-like, stained (coloured) bodies present in nucleus. These look like '**beads-on-string**', when observed under an electron microscope (Fig. 4.6). DNA helix of 200 bps wraps around the histone octamer by 1¾ turns. Six such nucleosomes get coiled repeatedly and then form solenoid that looks like coiled telephone wire (Fig. 4.7). The chromatin is 10 nm thick fibre packed to form a **solenoid** structure of 30 nm diameter (300Å⁰) and further supercoiling of solenoid tends to form a looped structure that further coils and condense at metaphase stage to form the **chromosomes**. The packaging of chromatin at higher levels, need additional set of proteins that are called Non-Histone Chromosomal proteins (NHC).

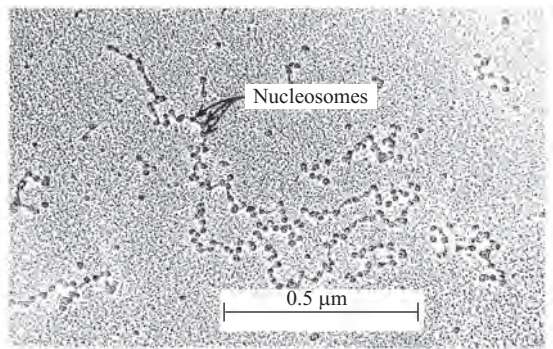
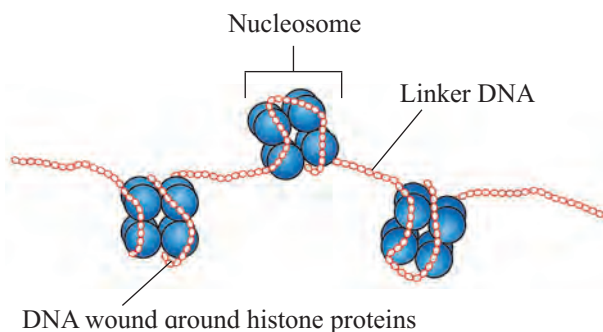
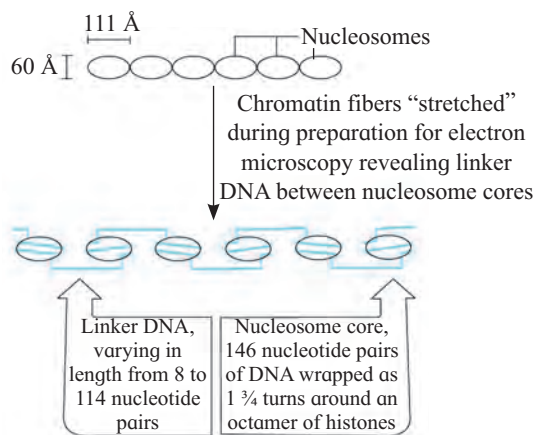


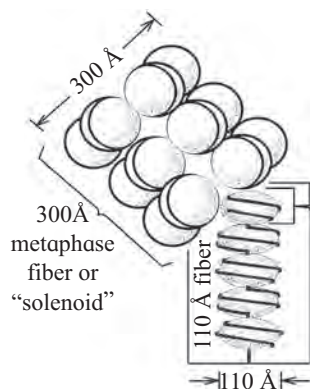
Fig. 4.6 : Chromatin showing beads-on-string



A : Beads-on-string magnified



B : Chain of nucleosomes forming 10 to 11 nm thick fibre



C : Solenoid forming 30 nm thick fibre

Fig. 4.7 : DNA packaging

Non-Histone Chromosomal Proteins (NHC) : These are additional sets of proteins that contribute to the packaging of chromatin at a higher level.

Heterochromatin and Euchromatin :

- 1. Heterochromatin:** In eukaryotic cells, some segments of chromonema/ chromosome during interphase and early prophase remain in a condensed state. These regions constitute heterochromatin. This term was proposed by Heitz. These regions are localized near centromere, telomeres and are also intercalated. It is genetically mostly inactive. It stains strongly and appears dark. Heterochromatin is 2 to 3 times more rich in DNA than in the euchromatin.
- 2. Euchromatin:** The regions of chromonema which are in non-condensed state, constitute euchromatin. Euchromatic regions stain light. Euchromatin is genetically very much active and fast replicating. Euchromatin is transcriptionally active, while heterochromatin is transcriptionally almost inactive.



Can you recall?

1. What is the backbone of the DNA structure?
2. Name the nitrogen bases of DNA.
3. What are Nucleoside and Nucleotide?
4. Is the double helix right or left handed?



Find out

What is the key difference between DNA in prokaryotic and eukaryotic cells?

4.4 DNA Replication :

The DNA molecule regulates and controls all the activities of the cell. Because of its unique structure, it is able to control the synthesis of other molecules of the cell. At the same time when the cell reproduces, the DNA also should duplicate itself to distribute

equally to the daughter cells. As a carrier of genetic information, DNA has to perform two important functions :

a. Heterocatalytic function : When DNA directs the synthesis of chemical molecules other than itself, then such functions of DNA are called heterocatalytic functions. Eg. Synthesis of RNA (transcription), synthesis of protein (Translation), etc.

b. Autocatalytic function : When DNA directs the synthesis of DNA itself, then such function of DNA is called autocatalytic function. Eg. Replication.

The process by which DNA duplicates itself is called replication. Through replication, it forms two copies that are identical to each other and also to the parent DNA.

In eukaryotic organisms, replication of DNA takes place only once in the cell cycle. It occurs in the S- phase of interphase in the cell cycle.

DNA replicates through Semiconservative mode of replication.

The model for Semiconservative replication was proposed by Watson and Crick, on the basis of antiparallel and complementary nature of DNA strands. The process of semiconservative replication is as below:

1. Activation of Nucleotides:

The four types of nucleotides of DNA i.e. dAMP, dGMP, dCMP and dTMP are present in the nucleoplasm. They are activated by ATP in presence of an enzyme **phosphorylase**. This results in the formation of deoxyribonucleotide triphosphates i.e. dATP, dGTP, dCTP and dTTP. The process is known as Phosphorylation.

2. Point of Origin or Initiation point:

It begins at specific point 'O' -origin and terminates at point 'T'. Origin is flanked by 'T' sites. The unit of DNA in which replication occurs, is called replicon. In prokaryotes, there is noly one replicon however in eukaryotes, there are several replicons in tandem.

At the point 'O', enzyme endonuclease nicks one of the strands of DNA, temporarily. The nick occurs in the sugar-phosphate back bone or the phosphodiester bond.

3. Unwinding of DNA molecule:

Now enzyme DNA helicase operates by breaking weak hydrogen bonds in the vicinity of 'O. The strands of DNA separate and unwind. This unwinding is bidirectional and continues as 'Y' shaped replication fork. Each separated strand acts as template.

The two separated strands are prevented from recoiling (rejoining) by SSBP (Single strand binding proteins). SSB proteins remain attached to both the separated strands so as to facilitate synthesis of new polynucleotide strands.

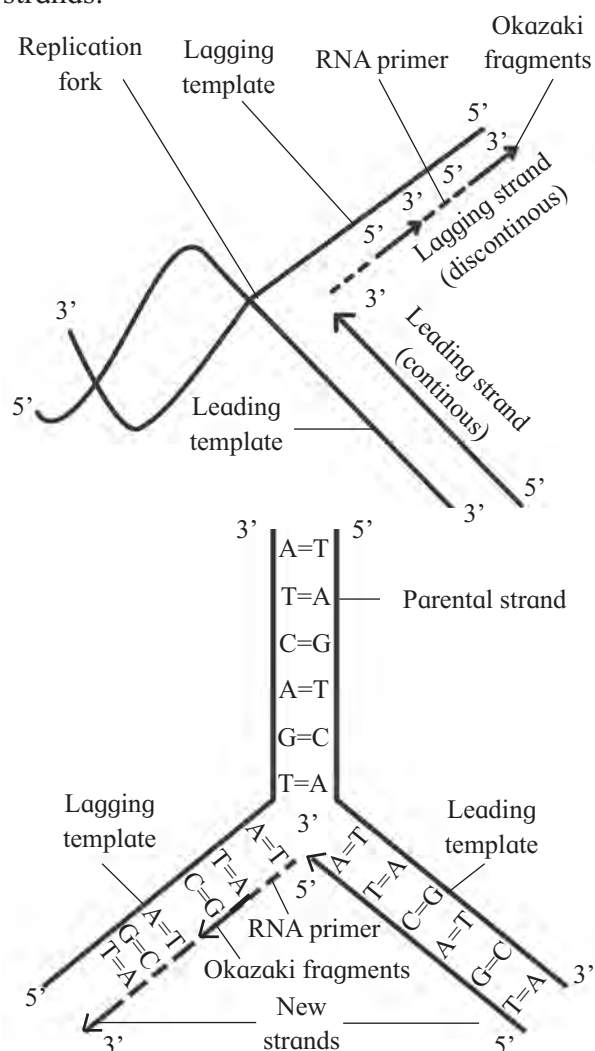


Fig. 4.8 : Semiconservative Replication of DNA

4. Replicating fork:

The point formed due to unwinding and separation of two strands appear like a Y-shaped fork, called replicating/ replication fork. The unwinding of strands imposes strain which is relieved by super-helix relaxing enzyme.

5. Synthesis of new strands:

Each separated strand acts as mould or template for the synthesis of new complementary strand. It begins with the help of a small RNA molecule, called **RNA primer**. RNA primer get associated with the 3' end of template strand and attracts complementary nucleotides from surrounding nucleoplasm. These nucleotides molecules bind to the complementary nucleotides on the template strand by forming hydrogen bonds (i.e. A=T or T=A; G ≡ C or C ≡ G). The newly bound nucleotides get interconnected by phosphodiester bonds, forming a polynucleotide strand. The synthesis of new complementary strand is catalyzed by enzyme DNA polymerase. The new complementary strand is always formed in 5'-3' direction.

6. Leading and Lagging strand:

The template strand with free 3' end is called leading template and with free 5' end is called lagging template. The process of replication always starts at C-3 end of template strand and proceeds towards C-5 end. As both the strands of the parental DNA are antiparallel, new strands are always formed in 5' → 3' direction.

One of the newly synthesized strand develops continuously towards replicating fork is called leading strand. Another new strand develop discontinuously away from the replicating fork is called lagging strand.

Maturation of Okazaki fragments :

DNA synthesis on lagging template takes place in the form of small fragments, called Okazaki fragments (named after scientist

Okazaki). Okazaki fragments are joined by enzyme DNA ligase.

RNA primers are removed by DNA polymerase and replaced by DNA sequence with the help of DNA polymerase-I in prokaryotes and DNA polymerase- α in eukaryotes.

Finally, DNA gyrase (topoisomerase) enzyme forms double helix to form daughter DNA molecules.

7. Formation of daughter DNA molecules:

At the end of the replication, two daughter DNA molecules are formed. In each daughter DNA, one strand is parental and the other one is totally newly synthesized. Thus, 50% is contributed by mother DNA. Hence, it is described as **semiconservative replication**.

Experimental confirmation :

Semiconservative Replication : In newly formed DNA molecule, one strand is old (i.e. conserved) and other strand is newly synthesized. Thus, it is called semiconservative mode of replication.

It is experimentally proved by Matthew Meselson and Franklin Stahl (1958) by using **equilibrium - density - gradient - centrifugation** technique.

1. Meselson and Stahl in 1958 performed an experiment to verify semiconservative nature (mode) of replication.
2. *E. coli* cells growing in ^{14}N were then transferred to ^{15}N medium (heavy isotopic nitrogen) and allowed to replicate for several generations. At equilibrium point density gradient band was obtained, by using 6M CsCl_2 . The position of this band is recorded.
3. The heavy DNA (^{15}N) molecule can be distinguished from normal DNA (^{14}N) by centrifugation in a 6M Cesium chloride (CsCl_2) density gradient. The density

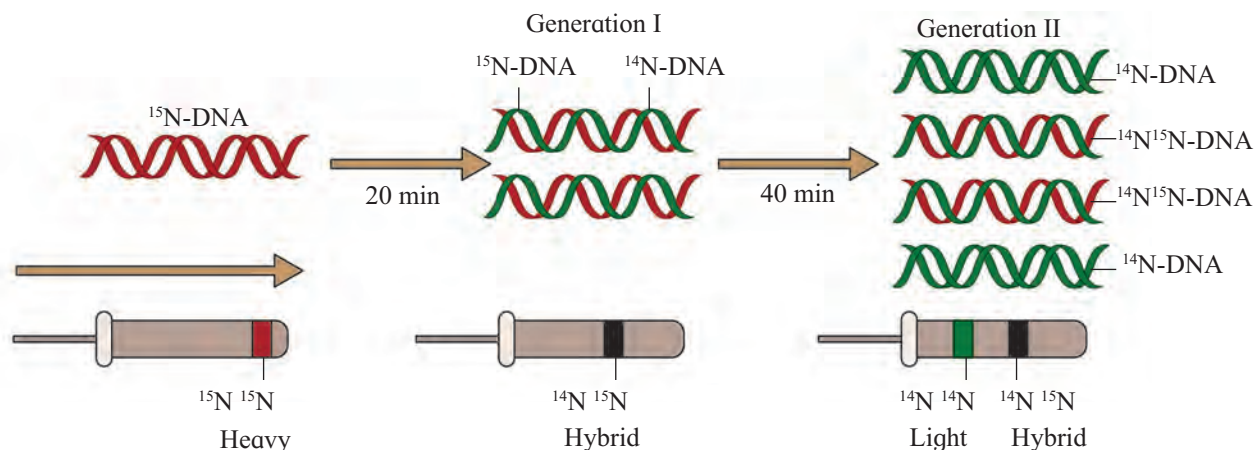


Fig. 4.9 : Interpretation of results of Meselson's experiment on the separation of DNA by equilibrium density gradient centrifugation

gradient value of 6M CsCl₂ and ¹⁵N DNA is almost same. Therefore, at the equilibrium point ¹⁵N DNA will form a band. In this both the strands of DNA are labelled with ¹⁵N.

- Such *E. coli* cells were they transferred to another medium containing ¹⁴N i.e. normal (light) nitrogen. After first generation, the density gradient band for ¹⁴N -¹⁵N (hybrid) was obtained and its position was recorded. After second generation, two density gradient bands were obtained - one at ¹⁴N -¹⁵N position and other at ¹⁴N position.
- The position of bands after two generations clearly proved that DNA replication is Semiconservative.

4.5 Protein synthesis :

Proteins are very important biomolecules. They serve as structural components, enzymes and hormones. The cell needs to synthesize new protein molecules. The process of protein synthesis includes **transcription** and **translation**. The process of copying of genetic information from one (template) strand of DNA into a single stranded RNA transcript, is termed as **transcription**. During this process, synthesis of complementary strand of RNA takes place (Except that the Adenine nitrogen base pairs with the Uracil base instead of Thymine).

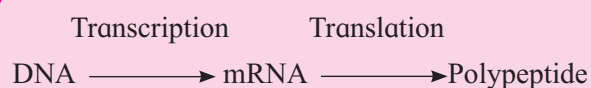
Central Dogma :

Double stranded DNA molecule gives rise to mRNA which acts as a messenger to programme the synthesis of a polypeptide chain (protein). This type of unidirectional flow of information from DNA to RNA to protein/ proteins is referred as **central dogma** of molecular biology. It was postulated by F.H.C. Crick in 1958.

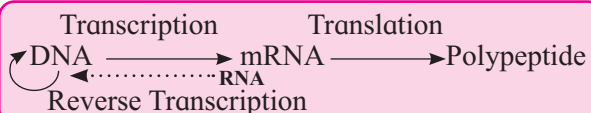


Use your brain power

- List as many different enzyme activities required during DNA synthesis as you can.
- This type of replication is called semi-conservative replication. Considering the meaning of these words, why DNA replication is called semiconservative replication?



The present concept of central dogma in retroviruses or riboviruses is given by Temin (1970) and Baltimore (1970):



Accordingly enzyme RNA dependent DNA polymerase, synthesizes DNA from RNA.



Can you recall?

1. What is transcription?
2. How many nucleotides are present in a codon?
3. Name the molecule which carries anticodon?
4. What is mutation?

A. Transcription:

During transcription, information of only one strand of DNA is copied into RNA. This strand of DNA acts as template. Enzyme RNA polymerase catalyses the formation of RNA transcript.

DNA is located in the nucleoid of Prokaryotes and in nucleus of Eukaryotes. DNA transcription takes place in nucleus in eukaryotes whereas translation occurs in cytoplasm. DNA transfers information to m-RNA which then moves to ribosomes. Transcription occurs in the nucleus during G1 and G2 phases of cell cycle. DNA has promoter and terminator sites. Transcription starts at promoter site and stops at terminator site. Actually the process of transcription, in both Prokaryotes and Eukaryotes, involves three stages viz. Initiation, Elongation and Termination.

Transcription Unit:

Each transcribed segment of DNA is called transcription unit. It consists of i. Promotor, ii. The structural gene, iii. A terminator. Two strands of DNA in the structural gene show following features :

- The promoter is located towards 5' end of structural gene i.e. upstream. It is a DNA sequence that provides binding site for enzyme RNA polymerase. RNA polymerase binds to specific Promotor. In prokaryotes, the enzyme recognizes the promoter by its sigma factor sub unit.
- Structural genes - two strands of DNA have opposite polarity. DNA dependent RNA

polymerase catalyses polymerisation in 5'→3' direction. So the DNA strand having 3'→5' polarity acts as template strand. The other strand of DNA having 5'→3' polarity is complementary to template strand. The sequence of bases in this strand, is same as in RNA (where Thymine is replaced by Uracil). It is the actual coding strand. The information on this strand of DNA is copied on mRNA. This is called sense strand.

- The terminator is located at 3' end of coding strand i.e. downstream. It defines the end of the transcription process.

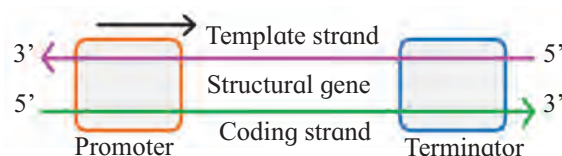


Fig. 4.10 : Transcription unit

After binding to promoter, RNA polymerase moves along the DNA and causes local unwinding of DNA duplex into two chains in the region of the gene. Exposed ATCG bases project into nucleoplasm. Only one strand functions as template (antisense strand) and the other strand is complementary which is actually a coding strand (sense strand). The ribonucleoside tri phosphates join to bases of DNA template chain. As transcription proceeds, the hybrid DNA-RNA molecule dissociates and makes mRNA molecule free.

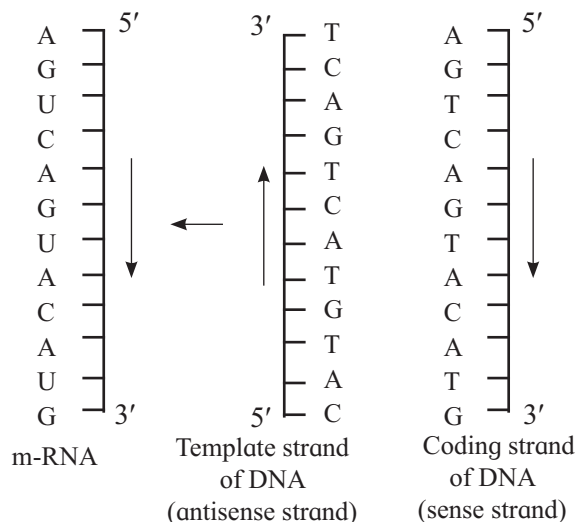


Fig. 4.11 : Formation of Template and Coding strand of DNA

When RNA polymerase reaches the terminator signal on the DNA, it leaves DNA and fully formed mRNA (primary transcript) is released.

As the mRNA grows, the transcribed region of DNA molecule becomes spirally coiled and attains (regains) double helical form.

In bacteria, m-RNA does not require any processing because it has no introns. Prokaryotes possess only one type of RNA polymerase. In eukaryotes, there are three types of RNA polymerases. RNA polymerase-I transcribes r-RNA. RNA polymerase-II transcribes m-RNA (primary transcript) and heterogeneous nuclear RNA (or hnRNA). RNA polymerase-III is responsible for transcription of t-RNA and small nuclear RNA (snRNA).



Do you know ?

1. Many viruses contain RNA as genetic material and replicate by synthesizing first the DNA and then form RNA. This process is called reverse transcription. Such viruses are known as Retroviruses.
2. Human immunodeficiency virus (HIV) is responsible for causing AIDS.
3. In some cases like *E.coli*, a chain terminating protein, the 'rho' factor stops the synthesis of mRNA.
4. The process of transcription as well as translation involves 3 stages - initiation, elongation and termination.



Always Remember

Types of RNA and process of transcription:

In bacteria, there are three types of RNAs: m-RNA provides the encoded message, t-RNA brings specific amino acid, to the site of translation. r-RNA plays a role in providing binding site to mRNA.

There is a single DNA-dependent RNA polymerase that catalyzes transcription of all 3 types of RNA in bacteria. RNA polymerase binds to the promoter and initiates transcription (initiation).

Transcription unit and the gene:

The DNA sequence coding for m-RNA, t-RNA or r-RNA is defined as a gene. A cistron is a segment of DNA coding for a polypeptide. A single structural gene in a transcription unit is said to be **monocistronic** whereas a long segment of DNA having a set of various structural genes in one transcription unit is referred to as **polycistronic**. Structural genes in eukaryotes have interrupted non-coding sequences (**introns**). The coding or expressible sequences are defined as **exons**. Only exons appear in processed mRNA in eukaryotes.

Processing of hnRNA:

In eukaryotes, forms of RNA transcribed from DNA are called **primary transcripts**. Such transcripts undergo changes called processing or maturation before becoming functional. Primary transcript is non-functional and contains both exons and introns. During processing, only introns are removed by the process called **splicing**.

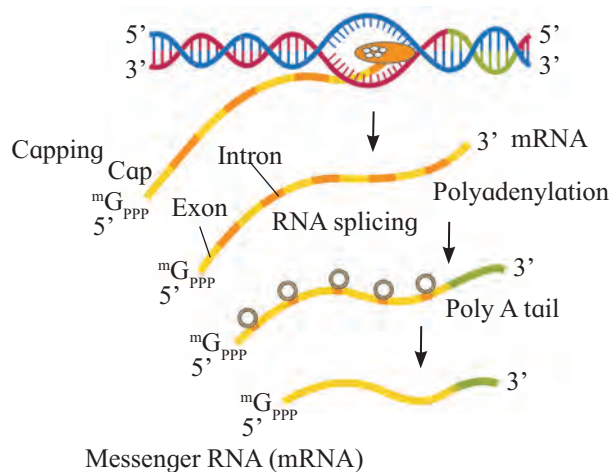


Fig. 4.12 : Transcription and Processing of hnRNA to mRNA in Eukaryotes

Exons are joined in a definite sequence (order) by DNA ligase enzyme. Heterogeneous nuclear RNA, undergoes the process of capping and tailing. In **capping**, methylated guanosine tri phosphate is added to 5' end of hnRNA. In **tailing**, polyadenylation take place at 3' end. It is the fully processed hnRNA, now called m-RNA. For translation m-RNA is transported out of the nucleus through nuclear pore.

Genetic Code:

It is already known that DNA is a master molecule of a cell that initiates, guides, regulates and controls the process of protein synthesis. To perform this complicated function, it must carry the requisite information for the synthesis of proteins. Obviously this information has to be verily located in the DNA itself. The site for storing this information lies in the sequence of nucleotides (i.e. nitrogen bases), as evidenced by Yanofski and Sarabhai (1964).

About, 20 different types of amino acids are involved in the process of synthesis of proteins. DNA molecule has 4 types of nitrogen bases to identify these 20 different types of amino acids. Question arises then, how is it possible that 20 types of amino acids are encoded by 4 types of nitrogen bases?

According to F.H.C. Crick, this information is stored in the form of coded language (cryptogram) called **genetic code**, that contains code words (**codons**) each one specifying (representing) specific amino acid. Genetic code, therefore, is a collection of base sequences that correspond to each amino acid.

A single nitrogen base in a codon (singlet codon) will encode for only four different types of amino acids. A combination of two nitrogen bases (doublet codon) will specify only 16 different types of amino acids. A combination of three nitrogen bases (triplet codon) will specify 64 different types of amino acids. Hence G. Gamov (1954) suggested that in a codon,

there must be combination of three consecutive nitrogen bases that will be sufficient to specify 20 different types of amino acids.

Thus, there would be 64 different codons (code words) in the dictionary of genetic code and that each code word has to be a triplet codon. Every three consecutive nucleotides in DNA will constitute a triplet codon. Genetic code is a triplet code, was evidenced first by Crick (1961) using "frame- shift mutation". However, M. Nirenberg and Matthaei were able to synthesize artificial m-RNA which contained only one type nitrogenous base i.e. Uracil (Homopolymer). This synthetic poly-U sequence was transferred to protein synthesizing enzymes. A small polypeptide molecule was produced/ formed by the linking of phenylalanine molecules. This explains that UUU codes for phenyl alanine. Later different homopolymer codons were deciphered. Codons formed by two or more bases were also tried.

Dr. Har Gobind Khorana : He devised a technique for artificially synthesizing m- RNA with repeated sequences of known nucleotides. By using synthetic DNA, Dr. Khorana prepared chains of polyribonucleotides with known repeated sequences of two or three nucleotides. eg. CUC UCU CUC UCU.

This resulted in formation of polypeptide chain having two different amino acids placed alternately (Leucine and Serine). Similarly polynucleotide chain with three- nitrogen base repeats gave polypeptide chain with only one amino acids. Eg. CUA CUA CUA CUA (leucine). Later, Severo Ochoa established that the enzyme (polynucleotide phosphorylase) was also helpful in polymerising RNA with defined sequences in a template- independent manner (i.e. enzymatic synthesis of RNA).

Finally Nirenberg, Matthaei and Ochoa deciphered all the 64 codons in the dictionary of genetic code.

		Second Letter				
		U	C	A	G	
First Letter	U	UUU } Phe UUC } UUA } Leu UUG }	UCU } UCC } Ser UCA } UCG }	UAU } Tyr UAC } UAA Stop UAG Stop	UGU } Cys UGC } UGA Stop UGG Trp	U C A G
	C	CUU } CUC } Leu CUA } CUG }	CCU } CCC } Pro CCA } CCG }	CAU } His CAC } CAA } Gln CAG }	CGU } CGC } Arg CGA } CGG }	U C A G
	A	AUU } AUC } Ile AUA } AUG Met	ACU } ACC } Thr ACA } ACG }	AAU } Asn AAC } AAA } Lys AAG }	AGU } Ser AGC } AGA } Arg AGG }	U C A G
	G	GUU } GUC } Val GUA } GUG }	GCU } GCC } Ala GCA } GCG }	GAU } Asp GAC } GAA } Glu GAG }	GGU } GGC } Gly GGA } GGG }	U C A G

Fig. 4.13 : Dictionary of genetic code

During replication and transcription, a nucleic acid is copied to form another nucleic acid. These two processes are based on complementarity principle. During translation, genetic information is transferred from a polymer of nucleotides to a polymer of amino acids. Here, complementarity principle does not exist.

It is evident that change in nucleic acid (genetic material) results in the change in amino acids of proteins. This clearly explains that genetic code directs the sequence of amino acids during synthesis of proteins.



Find out

What is the amino acid sequence encoded by base sequence UCA UUU UCC GGG AGU of an m- RNA segment?

Characteristic of Genetic code:

Genetic code of DNA has certain fundamental characteristics –

- i. **Genetic code is a triplet code:** Sequence of three consecutive bases constitute codon, which specifies one particular amino acid. Base sequence in a codon is always in 5' → 3' direction. In every living organism genetic code is a triplet code.
- ii. **Genetic code has distinct polarity :** Genetic code shows definite polarity i.e. direction. It, therefore, is always read in 5' → 3' direction and not in 3'→5' direction. Otherwise message will change e.g. 5' AUG 3'.
- iii. **Genetic code is non-overlapping :** Code is non overlapping i.e. each single base is a part of only one codon. Adjacent codons do not overlap. If non-overlapping, then with 6 consecutive bases only two amino acid molecules will be in the chain. Had it been overlapping type, with 6 bases, there would be 4 amino acid molecules in a chain. Experimental evidence is in favour of non-overlapping nature.

- iv. **Genetic code is commaless** : There is no gap or punctuation mark between successive/ consecutive codons.
- v. **Genetic code has degeneracy** : Usually single amino acid is encoded by single codon. However, some amino acids are encoded by more than one codons. e.g. Cysteine has two codons, while isoleucine has three codons. This is called degeneracy of the code. Degeneracy of the code is explained by Wobble hypothesis. Here, the first two bases in different codons are identical but the third one, varies.
- vi. **Genetic code is universal** : By and large in all living organisms the specific codon specifies same amino acid. e.g. codon AUG always specifies amino acid methionine in all organisms from bacteria up to humans.
- vii. **Genetic code is non-ambiguous** : Specific amino acid is encoded by a particular codon. Alternatively, two different amino acids will never be encoded by the same codon.
- viii. **Initiation codon and termination codon**: AUG is always an initiation codon in any and every mRNA. AUG codes for amino acid methionine. Out of 64 codons, three codons viz. UAA, UAG and UGA are termination codons which terminate/ stop the process of elongation of polypeptide chain, as they do not code for any amino acid.
- ix. **Codon and anticodon** : Codon is a part of DNA e.g. AUG is codon. It is always represented as 5' AUG 3'. Anticodon is a part of tRNA. It is always represented as 3'UAC 5'.

- It is possible to predict sequence of codon on mRNA by studying the sequence of amino acids in a polypeptide chain.

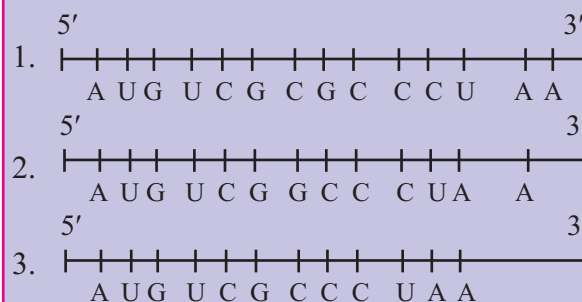


Activity :



m-RNA

Consider given m-RNA strand which has undergone mutation and lost nucleotides A, C, and G sequentially. Resultant mRNA is represented by 1, 2 and 3. With the help of checker board of amino acids, explain the changes in amino acid sequence that will occur due to such mutation.



Mutations and Genetic Code:

Mutation is a phenomenon in which sudden change in the DNA sequence takes place. It results in the change of genotype expressed in terms of phenotype (i.e. character). Along with recombination, mutation is raw material for evolution as it also results in variations. During mutation, possibility of loss (deletion) or gain (insertion/ duplication) of a segment of DNA results in alteration in the chromosome. Mutation can also occur due to change in a single base pair of DNA. This is known as point mutation. Eg. Sickle cell anaemia (Refer to earlier chapter). Deletion or insertion of base pairs of DNA causes frame – shift mutations or deletion mutation. Insertion or deletion of one or two bases changes the reading frame from the point of insertion or deletion. Insertion or deletion of three or multiples of three bases (insert or delete) results in insertion or deletion of amino acids and reading frame remains unaltered from that point onwards.

t-RNA- the adapter molecule:

Scientists considered that there has to be a mechanism in which t-RNA will read the

codon and also simultaneously binds with the amino acid as amino acid does not have any special capacity to read the codon. So t-RNA is considered as an adapter molecule. This role of tRNA was understood much later.

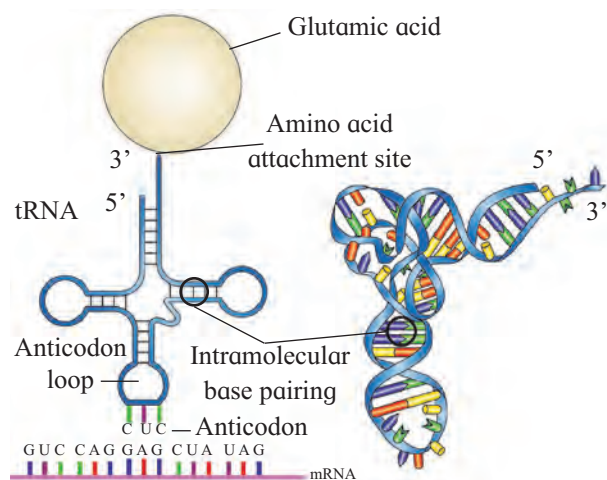


Fig. 4.14: t-RNA - the adapter molecule

Cloverleaf structure (2 dimensional) of t-RNA possess an anticodon loop that has bases complementary to the codon. It is called **anticodon**. It shows amino acid acceptor end (3' end) having unpaired CCA bases (i.e. amino acid binding site) to which amino acid binds. For every amino acid, there is specific t- RNA. Initiator t-RNA is specific for methionine. There are no t-RNA's for stop codons. In the actual structure, the t-RNA molecule looks like inverted L (3 dimensional structure).



Can you tell?

Why t-RNA is called as adapter molecule?



Can you recall?

1. Name different types of RNAs.
2. Name the site of protein synthesis.
3. Which molecule carries information of protein synthesis from gene?
4. Which molecule carries amino acid from cytoplasm to ribosome?

B. Translation - protein synthesis :

Translation is the mechanism in which codons of mRNA are translated and specific amino acids in a sequence form a polypeptide on ribosomes. All types of proteins are synthesised by the cell, within itself (i.e. intracellularly).

Process of translation requires amino acids, mRNA, tRNA, ribosomes, ATP, Mg⁺⁺ ions, enzymes, elongation, translocation and release factors.

- i. Amino acids form raw material for protein synthesis. About 20 different types of amino acids are known to form proteins. These are available in the cytoplasm.
- ii. DNA controls synthesis of proteins having amino acids in specific sequence. This control is possible through transcription of m-RNA. Genetic code is specific for particular amino acid.
- iii. RNAs serve as intermediate molecules between DNA and protein.
- iv. Ribosomes serve as site for protein synthesis. Each ribosome consists of large and small subunits. These subunits occur separately in cytoplasm. Only during protein synthesis, these two subunits get associated together due to Mg⁺⁺ ions.

A ribosome has one binding site for m-RNA and 3 binding sites for t-RNA. They are P site (peptidy t-RNA site), A site (aminoacyl – t-RNA site) and E site (exit site). Only first t- RNA- amino acid complex, directly enters P site of ribosome.

In Eukaryotes, a groove is present between two subunits of ribosomes. It protects the Polypeptide chain from the action of cellular enzymes and also protects mRNA from the action of nucleases.

Mechanism of translation (i.e. synthesis of polypeptide chain) :

It involves three steps :

- i. Initiation, ii. Elongation, iii. Termination

1. Initiation of Polypeptide chain :

- Activation of amino acids is essential before translation initiates for which ATP is essential. Small subunit of ribosome binds (attaches) to the m-RNA at 5' end. Initiator codon, AUG is present on m-RNA which initiates the process of protein synthesis (translation). Initiator t- RNA binds with initiation codon (AUG) by its anticodon (UAC) through hydrogen bonds. It carries activated amino acid methionine (in Eukaryotes) or formyl methionine (in prokaryotes).
- Now the large subunit of ribosome joins with the smaller subunit, that requires Mg^{++} ions.
- Initiator charged t-RNA (with activated amino acid methionine) occupies the P-site of ribosome and A- site is vacant.

2. Elongation of polypeptide chain :

During this process, activated amino acids are added one by one to first amino acid (methionine). Amino acid is activated by utilising energy from ATP molecule. This amino acid binds with amino acid binding site of t-RNA- This results in formation of t-RNA- amino acid complex.

Addition of Amino acid occurs in 3 Step cycle -

- Condon recognition- Amino acyl t- RNA molecule enters the ribosome at A-site. Anticodon binds with the codon by hydrogen bonds.
- Amino acid on the first initiator t-RNA at P-site and amino acid on t-RNA at A-site join by peptide bond. Here enzyme Ribozyme acts as a catalyst. At this time first tRNA at 'P' site is kicked off.
- Translocation- The t- RNA at A-site carrying a dipeptide at A-site moves to the P-site. This process is called **translocation**. In translocation, both the subunits of ribosome move along in relation to tRNA and mRNA. Hence,

tRNA carrying dipeptide now gets positioned at 'P' site of ribosome, making 'A' site vacant. At this site, then next charged tRNA molecule carrying amino acid will be received. During this process, first uncharged tRNA is discharged from E-site.

This process is repeated as amino acids are added to Polypeptide. It takes less than 0.1 second for formation of peptide bond.

Third charged t-RNA with its amino acid, arrives at A-site of ribosome. Anticodon and codon bind by hydrogen bond. Polypeptide bond is formed. Second t-RNA is discharged from P-site to E-site and leaves the ribosome. So the events like arrival of t-RNA- amino acid complex, formation of peptide bond, ribosomal translocation and removal of previous tRNA, are repeated. As ribosome move over the m-RNA, all the codons on mRNA are exposed one by one for translation.

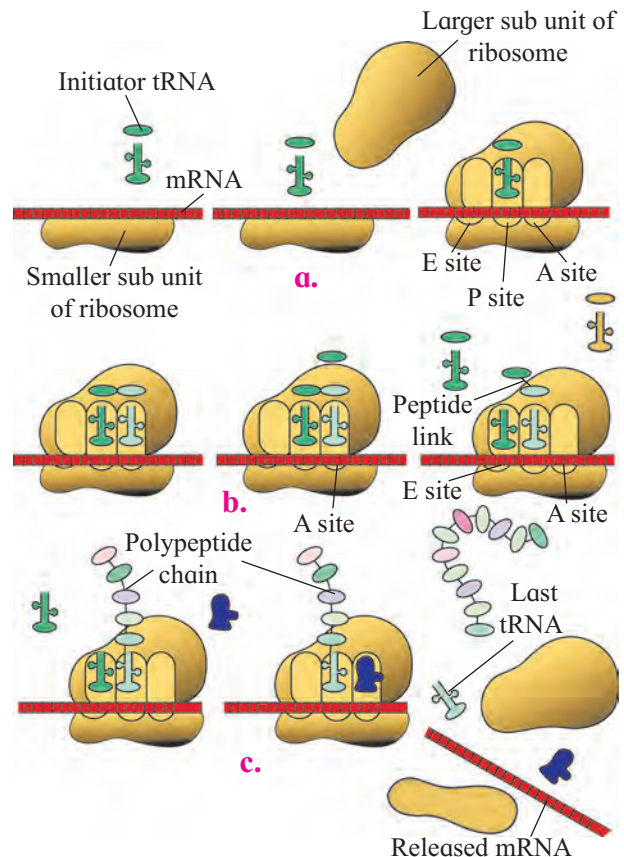


Fig. 4.15 : Translation Protein synthesis
a. Initiation, b. Elongation, c. Termination

3. Termination and release of polypeptide:

At the end of m-RNA, there is a stop codon (UAA/ UAG/ UGA). It is exposed at the A-site. It is not read and joined by anticodon of any t-RNA. The release factor binds to the stop codon, thereby terminating the translation process. The Polypeptide is now released in the cytoplasm.

Two subunits of Ribosome dissociate and last tRNA is set free in the cytoplasm.

m-RNA also has some additional sequences that are not translated and are referred as **untranslated regions** (UTR). The UTRs are present at both 5'-end (before start codon) and at 3'-end (after stop codon). They are required for efficient translation process.

Finally mRNA is also released in the cytoplasm. It gets denatured by nucleases immediately. Hence mRNA is short-lived.



Can you tell?

1. Enlist different steps of protein synthesis.
2. Name the initiator codon of protein synthesis.
3. Explain in brief the process of initiation during protein synthesis.
4. Name three binding sites of ribosome.
5. What is translation?

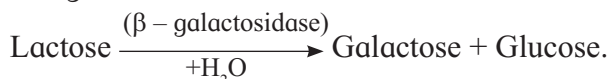
4.6 Regulation of gene expression:

It is the multistep process by which a gene is regulated and its product is synthesized. Thus, gene expression results in the formation of a Polypeptide. Gene expression process is regulated at different levels.

In eukaryotes, the regulation can be at different levels like-

1. Transcriptional level (formation of primary transcript)
2. Processing level (regulation of splicing)
3. Transport of m-RNA from nucleus to the cytoplasm.
4. Translational level.

Genes of a cell are expressed to perform different functions. For eg. An enzyme beta galactosidase is synthesised by *E.coli*. It is used for hydrolysis of lactose into galactose and glucose.



If *E.coli* bacteria do not have lactose in the surrounding medium as a source of energy, then enzyme β -galactosidase is not synthesised. So, it is the metabolic or physiological or environmental conditions that regulate expression of genes. The development and differentiation of embryo into an adult organism, is also a result of the coordinated regulation or expression, of several sets of genes.

Now one has to understand and know the mechanism by which the organisms regulate gene expression in response to changes in the environment. If so, whether single mechanism exists for regulation of the expression of different genes/ sets of genes or different genes are regulated by different mechanisms.

Certain bacteria like *E.coli* adapt to their chemical environment by synthesizing certain enzymes depending upon the substrate present. Such adaptive enzyme is called **inducible enzymes**. A set of genes will be switched on when there is necessity to metabolise a new substrate. This phenomenon is called **induction** and small molecule responsible for this, is known as **inducer**. It is positive control.



Do you know ?

Repressible regulation of gene is seen when the end product of a biosynthetic pathway like amino acid, is provided in the medium. At this time, internal biosynthesis of amino acid stops. It is negative control so the metabolite (amino acid) turns off a set of genes involved in producing that metabolite. This is called feedback repression.

4.7 Operon concept :

It is a transcriptional control mechanism of gene regulation. Francois Jacob and Jacques Monod (1961) explained that metabolic pathways are regulated as a unit.

For example in *E.coli*, when lactose sugar is provided to the culture medium, cell induces production of three enzymes necessary for digestion of lactose. The enzymes are :

- i. **β -galactosidase** : Digests lactose into galactose and glucose.
- ii. **β -galactoside permease** : Permits lactose molecules to enter into the cell.
- iii. **Transacetylase (β -Galactoside acetyltransferase)** : Transfers an acetyl group from acetyl Co-A to galactoside.

Synthesis of these three enzymes, is controlled by a long segment of DNA known as **Operon**. It consists of an operator site O and three structural genes z, y and a .The action of structural genes is regulated by operator site with the help of a **repressor protein**. Repressor protein is produced by the action of gene i (inhibitor) known as **regulator gene**.

The gene expression depends on whether operator is switched on or switched off.

If the operator is switched on, the three genes z, y and a are transcribed by RNA Polymerase into a single m-RNA. Each structural gene is generally known as **cistron** and the transcribed long m-RNA covering various cistrons is known as **Polycistronic**.

Switching on or **switching off** of the operator is achieved (accomplished) by a protein called **repressor**. When this protein is attached to the operator and blocks it, the switch is turned off and structural genes are not expressed.

Lac operon :

Lactose or lac operon of *E.coli* is inducible operon. The operon is switched on when a chemical inducer- lactose is present in the medium.

Lac operon consists of following components :

1. Regulator gene (repressor gene)
 2. Promoter gene
 3. Operator gene
 4. Structural genes
 5. Inducer - It is not a component of operon.
1. **Regulator gene** : This gene controls the operator gene in cooperation with an inducer present in the cytoplasm. Regulator gene precedes the promoter gene. It may not be present immediately adjacent to operator gene. Regulator gene produces a protein called **repressor protein**. Repressor binds with operator gene and represses (stops) its action. It is called **regulator protein**.
 2. **Promoter gene** : This gene precedes the operator gene. It is present adjacent to operator gene. The promoter gene marks the site at which the RNA Polymerase enzyme binds. When the operator gene is turned on, the enzyme moves over the operator gene and transcription of structural genes starts. Promoter gene base sequence determines which strand of DNA acts a template.
 3. **Operator gene** : It precedes the structural genes. This controls the functioning of structural genes. It lies adjacent to the Structural genes. When operator gene is turned on by an inducer, the Structural genes produce m-RNA. Operator gene is turned off by a product of repressor gene.
 4. **Structural gene** : When lactose is added to the *E.coli* culture, the structural genes catalyse (produce) m-RNA which in turn produces polypeptides, on the ribosomes.
The polypeptides formed, act as enzymes to catalyse lactose in the cell. There are 3 structural genes in the sequence lac-z(1), lac-y(2) and lac-a(3). Enzymes produced are β -galactosidase, β -galactoside permease and transacetylase respectively.

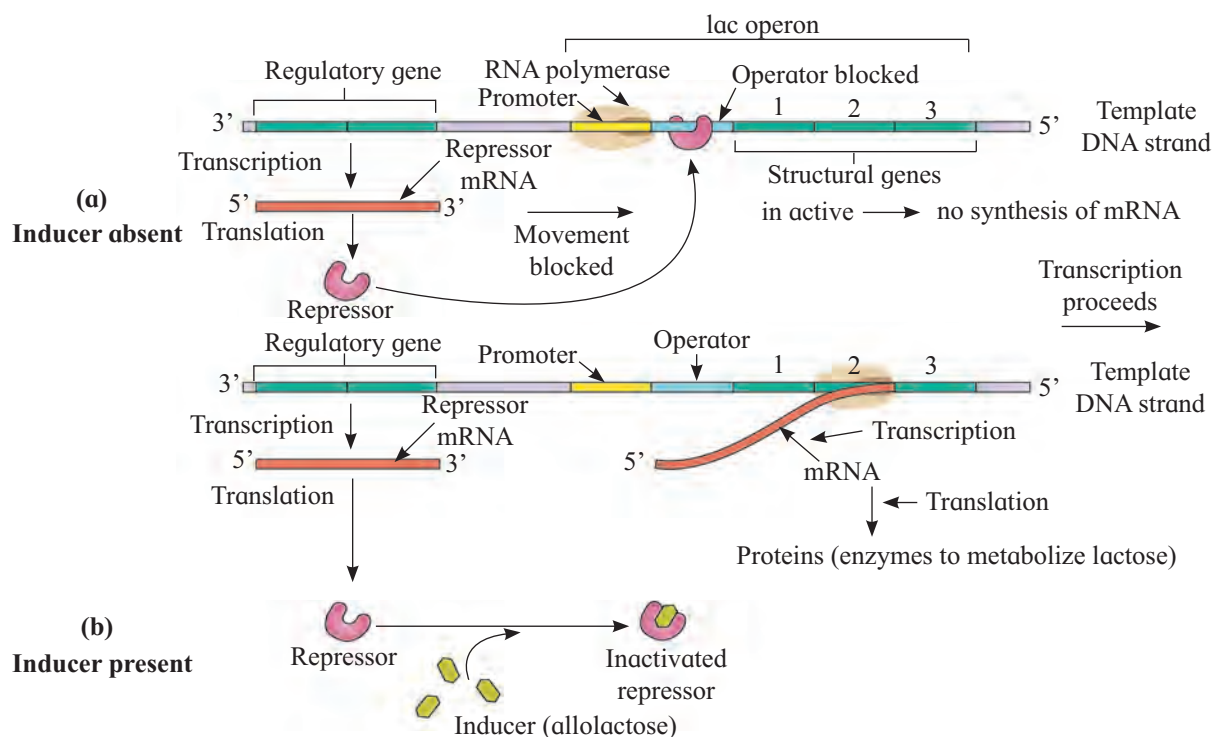


Fig. 4.16 : Working of Lac Operon

5. Inducer : It is a chemical in the cytoplasm (allolactose) which inactivates the repressor. When lac operon is switched on, then inducer joins with repressor protein preventing the binding of repressor to the operator gene. So the Operator gene is free and now enzyme RNA polymerase can move from promoter to structural genes via operator gene.

Role of lactose :

A few molecules of lactose enter into the cell by an enzyme permease. A small amount of this enzyme is present even when operon is switched off. A few molecules of lactose, act as inducer and bind to repressor. This repressor – inducer complex fails to join with the operator gene, which is then turned on. Structural genes produce all enzymes. Thus, lactose acts as an inducer of its own break down. When the inducer level falls, the operator is blocked again by repressor. So structural genes are repressed/ inactivated again. This is negative feedback.



Use your brain power

If operator gene is deleted due to mutation, how will *E.coli* metabolise lactose?



Can you tell?

1. What is the role of a repressor gene?
2. Name the different structural genes in sequence of lac operon.
3. Which molecule does act as inducer molecule in lac operon?
4. In which condition, lac operon is switched off?



Internet my friend

Find out information about Trp-operon, Ara-operon, His-operon, Val-operon.

4.8 Genomics:

The term **Genome** (introduced by H. Winkler in 1920) is the total genetic constitution of an organism. Alternatively, it is a complete copy of genetic information (DNA) or one complete set of chromosomes (monoploid or haploid) of an organism.

The term **Genomics** (term coined by T.H. Roderick in 1986) is the study of genomes through analysis, sequencing and mapping of genes along with the study of their functions.

The sequencing of yeast, *Drosophila* and mouse genome was done in order to facilitate comparative studies between humans and other organisms commonly used for genetic studies, in laboratory. Several additional genomes are now either actively being sequenced or strongly considered for sequencing. These include several microbes, bee, tomato and other crops.

Genomics study may be classified into two types:

- a. **Structural genomics:** It involves mapping, sequencing and analysis of genome.
- b. **Functional genomics:** It deals with the study of functions of all gene sequences and their expression in organisms.

Application of genomics:

Structural and functional genomics are used for different purposes in the improvement of crop plants, human health and livestock. The knowledge and understanding acquired from genomics research can be applied in a number of different sectors, including medicine, biotechnology and social sciences. It helps in the treatment of genetic disorders through gene therapy.

- Genomics is used in agriculture to develop transgenic crops having more desirable characters.
- Genetic markers developed in genomics, have applications in forensic analysis.
- Genomics can lead to introduce new gene in microbes to produce enzymes, therapeutic proteins and even biofuels.

4.9 Human Genome Project :

The human genome project was initiated in 1990 under the International administration of the Human Genome Organization (HUGO). The human genome project is a multinational research project to determine the genomic structure of humans. This project was coordinated by the US department of Energy and National institute of health. Additional contributors included universities across the

United States. International partners included are United Kingdom, France, Germany, Japan, India and China. This Project formally began in 1990 and was completed in 2003.

The main aims of project are –

- I. Mapping the entire human genome at the level of nucleotide sequences.
- II. To store the information collected from the project in databases.
- III. To develop tools and techniques for analysis of the data.
- IV. Transfer of the related technologies to the private sectors, such as industries.
- V. Taking care of the legal, ethical and social issues which may arise from project.

HGP (Human Genome Project) was closely associated with rapid development of a new area in biology, called **Bioinformatics**. The work of human genome project has allowed researchers to begin to understand the blueprint in building and constructing the human genome. As researchers learn more about the functions of genes and proteins, this knowledge will have a major impact in the fields like Medicine, Biotechnology and the Life sciences. Therefore HGP is very important.

Human Genome Project was to provide a complete and accurate sequence of the 3 billion DNA base pairs that make up the human genome and to find out the estimated number of human genes. Now about 33000 genes have been estimated to be present in humans.

The project was also aimed to sequence the genomes of several other organisms such as bacteria e.g. *E.coli*, *Caenorhabditis elegans* (a free living non-pathogenic nematode), *Saccharomyces cerevisiae* (yeast), *Drosophila* (fruit fly), plants (rice and *Arabidopsis*), *Mus musculus* (mouse), etc. Complete genome sequences of these model organisms will be useful for comparative studies that will allow researchers to study gene functions in these organisms.

Table 4.17 : Comparative genome sizes of humans and other model organisms.

Organism	Chromosome number	Estimated gene number	Estimated size (base pairs)
Human (<i>Homo sapiens</i>)	46	33,000	3 billion
Mouse (<i>Mus musculus</i>)	40	25,000	2.9 billion
Fruit fly (<i>Drosophila melanogaster</i>)	8	13,000	165 million
Plant (<i>Arabidopsis thaliana</i>)	10	25,000	157 million
Roundworm (<i>Caenorhabditis elegans</i>)	12	19,000	97 million
Yeast (<i>Saccharomyces cerevisiae</i>)	32	6000	12 million
Bacteria (<i>Escherichia coli</i>)	1*	4400	4.6 million

The secret of our complexity may lie not in the number of our genes but how we use them. It will lead to the understanding of gene structure and function in other species. Since we possess many of the genes same as these of flies, round worms and mice, such studies will lead to a greater understanding of human evolution.



Use your brain power

1. What have we learnt from the Human Genome Project?
2. Why is HGP important?



Can you tell?

Do different organisms have the same DNA?

4.10 DNA Fingerprinting:

Genes present on chromosomes are responsible for determining characters of the organism as well as for inheritance of characters. Due to recombination of paternal and maternal genes, we differ from our parents. Differences also arise due to infrequent mutations that occur during gamete formation (cell division). Due to all these factors, every individual has its unique genetic make-up, which may be called its **Fingerprint**. The technique developed to identify a person with the help of DNA restriction analysis, is known as **DNA profiling** or **DNA fingerprinting**. The technique of finger printing was first given by British geneticist, Dr. Alec Jeffreys in 1984.

DNA fingerprinting technique is based on identification of nucleotide sequence present in this wonder molecule. About 99.9% of nucleotide sequence in all persons, is same. Only some short sequences of nucleotides differ from person to person. In the population, every person shows unusual sequences of 20-100 base pairs, which are repeated several times. They are termed as Variable Number of Tandem Repeats (VNTRs).

The length of the regions having VNTRs is different in each individual and hence is the key factor in DNA profiling. Steps involved in DNA finger printing are as follows:

1. **Isolation of DNA:** The DNA must be recovered from the cells or tissues of the body (host). Only small amount of tissue like blood, hair roots, skin, etc. is required.
2. **Restriction digestion:** The isolated DNA is treated with restriction enzymes. The restriction enzymes cut the DNA into small fragments having variable lengths. This phenomenon is called **Restriction Fragment Length Polymorphism (RFLP)**.
3. **Gel electrophoresis:** The DNA samples are loaded on agarose gel for electrophoresis under an electric influence. The DNA fragments, which are negatively charged move to the positive pole. The movement of these fragments depends on length of the fragments. This results in formation of bands. dsDNA splits into ssDNA by alkali treatment.

- 4. Southern blotting:** It is a technique used for detecting a specific DNA sequence, developed by E.Southern. The separated DNA fragments are transferred to a nylon membrane or a nitrocellulose filter paper by placing it over the gel and soaking them with filter paper overnight.
- 5. Selection of DNA probe:** A known sequence of single- stranded DNA is prepared. It is called DNA Probe. DNA Probe is obtained from organisms or prepared by cDNA preparation method. The DNA probe is labelled with radioactive isotopes.
- 6. Hybridization:** Probe DNA is added to the nitrocellulose filter paper containing host DNA. The single-stranded DNA probe pairs with the complementary base sequence of the host DNA strand. As a

result DNA-DNA hybrids are formed on the nitrocellulose filter paper. Remaining single stranded DNA probe fragments are washed off.

- 7. Photography:** The nitrocellulose filter paper is photographed on an X-ray film by autoradiography. The film is analysed to determine the presence of hybrid DNA.

Application of DNA fingerprinting

- In forensic science, DNA finger printing is used to solve problems of rape and some complicated murder cases.
- DNA finger printing is used to find out the biological father or mother or both, of the child, in case of disputed parentage.
- DNA finger printing is used in pedigree analysis in cats, dogs, horses and humans.

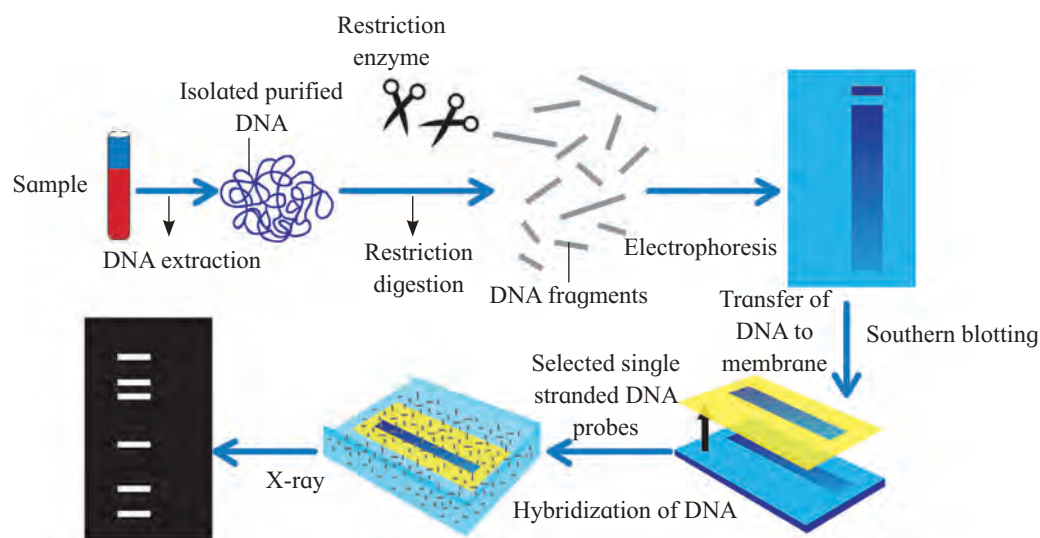


Fig. 4.18 : DNA Fingerprinting



Know the scientists



Dr. Lalji Singh (1947 - 2017)

Father of DNA Fingerprinting in India. He was instrumental in making DNA fingerprinting mainstream in India, for research and its forensic applications. He obtained DNA probe from Y chromosome of female banded krait snake (in this snake female has XY and male has YY chromosome). The unique segment obtained from this chromosome is, banded krait minor (BKM - DNA). It was used to developed probe for the Indigenous DNA fingerprinting technique.

Contributions of Dr. Lalji Singh: i. He installed several dedicated laboratories on aspects of genetics such as population biology, structural biology and transgenic research. ii. His work in the field of DNA fingerprinting technology, contributed for, wildlife conservation, forensics, evolution and phylogeny. iii. Established Centre for DNA Fingerprinting and Diagnostics (CDFD) in late 1990s- making it nodal centre for DNA fingerprinting and diagnostics for all species and several diseases. iv. Founded Laboratory for Conservation of Endangered Species (LaCONES).



Activity :

Prepare physical model of DNA molecule (Watson-Crick model)

Requirements :

Labelled Diagram :

Functions :

Exercise

Q. 1 Multiple Choice Questions

- Griffith worked on
 - Bacteriophage
 - Drosophila*
 - Frog eggs
 - Streptococci*
- The molecular knives of DNA are
 - Ligases
 - Polymerases
 - Endonucleases
 - Transcriptase
- Translation occurs in the
 - Nucleus
 - Cytoplasm
 - Nucleolus
 - Lysosomes
- The enzyme required for transcription is
 - DNA polymerase
 - RNA polymerase
 - Restriction enzyme
 - RNAase
- Transcription is the transfer of genetic information from
 - DNA to RNA
 - tRNA to mRNA
 - DNA to mRNA
 - mRNA to tRNA
- Which of the following is **NOT** part of protein synthesis?
 - Replication
 - Translation
 - Transcription
 - All of these
- In the RNA molecule, which nitrogen base is found in place of thymine?
 - Guanine
 - Cytosine
 - Thymine
 - Uracil
- How many codons are needed to specify three amino acid?
 - 3
 - 6
 - 9
 - 12
- Which out of the following is **NOT** an example of inducible operon?
 - Lactose operon
 - Histidine operon
 - Arabinose operon
 - Tryptophan operon

- Place the following event of translation in the correct sequence
 - Binding of met-tRNA to the start codon.
 - Covalent bonding between two amino acids.
 - Binding of second tRNA.
 - Joining of small and large ribosome subunits.

A. iii, iv, i, ii B. i, iv, iii, ii
C. iv, iii, ii, i D. ii, iii, iv, i

Q. 2 Very Short Answer Questions:

- What is the function of an RNA primer during protein synthesis?
- Why the genetic code is considered as commaless?
- What is genome?
- Which enzyme does remove supercoils from replicating DNA?
- Why are Okazaki fragments formed on lagging strand only?
- When does DNA replication take place?
- Define term- codon and codogen.
- What is degeneracy of genetic code?
- Which are the nucleosomal 'core' histones?

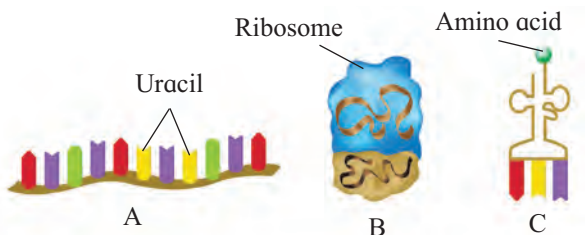
Q. 3 Short Answer Questions:

- Write short note on DNA packaging in eukaryotic cell.
- Enlist the characteristics of genetic code.
- Write a note on applications of DNA finger printing.
- Explain the role of lactose in 'Lac Operon'.

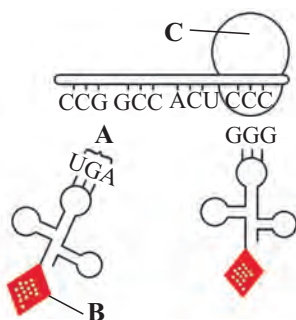
Q. 4 Short Answer Questions:

- Write a note on Human genome project (HGP).

- Describe the structure of 'Operon'.
- In the figure below A, B and C are three types of _____.



- Identify the labeled structures on the following diagram of translation.



Part A is the _____.
 Part B is the _____.
 Part C is the _____.

- Match the entries in column I with those of column II and choose the correct answer.

Column I	Column II
A. Alkali treatment	i. Separation of DNA fragments on gel slab
B. Southern blotting	ii. Split DNA fragments into single strands
C. Electrophoresis	iii. DNA transferred to nitrocellulose sheet
D. PCR	iv. X-ray photography
E. Autoradiography	v. Produce fragments of different sizes
F. DNA treated with REN	vi. DNA amplification

Q. 5 Long Answer Questions:

- Explain the process of DNA replication.
- Describe the process of transcription in protein synthesis.
- Describe the process of translation in protein synthesis.
- Describe the 'Lac-operon'.
- Justify the statements. If the answer is false, change the underlined word(s) to make the statement true.
 - The DNA molecule is double stranded and the RNA molecule is single stranded.
 - The process of translation occurs at the ribosome.
 - The job of mRNA is to pick up amino acids and transport them to the ribosomes.
 - Transcription must occur before translation may occur.
- Guess (i) the possible locations of DNA on the collected evidence from a crime scene and (ii) the possible sources of DNA.

Evidence	Possible location of DNA on the evidence	Sources of DNA
e.g. Eyeglasses	e.g. Ear pieces	e.g. Sweat, Skin
Bottle, Can, Glass	Sides, mouthpiece	-----
-----	Handle	S w e a t , skin, blood
Used cigarette	Cigarette butt	-----
Bite mark	-----	saliva
-----	Surface area	Hair, semen, sweat, urine

Project : Collect information about B and Z forms of DNA. Sketch the diagrams and write the differences between these two forms.



Can you recall?

1. What is evolution?
2. Where was first life formed?
3. Why are many species of plants and animals getting (becoming) extinct?
4. The Lamarck's theory of inheritance of acquired characters.
5. What is Speciation?

5.1 Origin of life : (Protobiogenesis)

The living matter shows attributes or characters like responsiveness, growth, metabolism, energy transformations and reproduction.

As far as origin of life is considered, it has remained an enigma for intellectuals at all times. Despite of advancements in various fields like biochemistry, astrobiology, geography, molecular biology, etc. scientists are unable to ascertain the truth. Various theories and hypotheses have been proposed to find the probable answer to this question.

a. Theory of special creation :

It is the oldest theory and is based on religious belief without any scientific proof. It states that all living organisms are created by a super-natural power.

b. Cosmozoic theory/Theory of Panspermia:

This theory advocates that life did not originate on the planet Earth. It may have descended to the earth from other planets in the form of spores or micro-organisms, called cosmozoa/panspermia. Recently, NASA has reported fossils of bacteria-like organisms on a piece of Martian rock recovered from Antarctica. This theory could not explain the origin of life on the other planets hence not accepted.

c. Theory of spontaneous generation (Abiogenesis) :

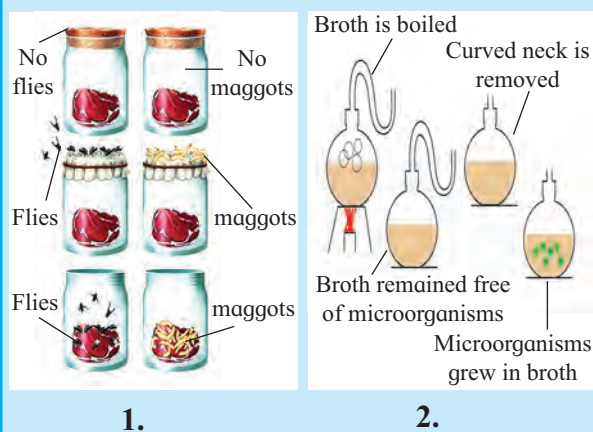
According to this theory, life originated from



Observe and Discuss

1. Redi's Experiment

2. Louis Pasteur's Experiment



non-living (inanimate) material spontaneously. This theory was disproved by Louis Pasteur.

d. Theory of biogenesis :

According to this theory, living organisms are always produced from pre-existing living forms, by process called reproduction.

Theory of biogenesis however could not explain origin of first life on earth but could explain only the continuity of life.

5.2 Chemical Evolution of Life (Self assembly theory of origin of life) :

According to this theory, life originated on earth by combinations of several chemicals through constant chemical reactions over a long period of time. This theory is also called self assembly theory of origin of life or biochemical origin of life.

This theory was first formulated by Haeckel but later developed by the Russian scientist Alexander I. Oparin (1924) and British biologist J. B. S. Haldane (1929). The process of chemical evolution can be divided into following steps :

a. Origin of Earth and Primitive atmosphere:

The origin of universe was explained by the Big-Bang theory of Georges Lemaitre (1931). According to this theory the Universe originated about 20 billion years ago by a single huge titanic explosion. As the universe expanded, the temperature decreased and various galaxies of solid objects were formed. Milky Way is one such galaxy of which our solar system is one small part. Earth is one of the planets of solar system and originated about 4.6 billion year ago. When formed, it was a rotating cloud of hot gases and cosmic dust called Nebula. The condensation and cooling resulted in stratification with heavier elements like nickel and iron passing to the core and lighter ones like helium, hydrogen, nitrogen, oxygen, carbon, etc. remaining on the surface. They formed the atmosphere of the earth. The primitive atmosphere of the earth was quite different from the present one and it was of a reducing type, devoid of free oxygen.

b. Formation of ammonia, water and methane:

Primitive atmosphere was very hot. As it slowly cooled, the lighter elements started to react with each other. The early atmosphere was rich in hydrogen, carbon, nitrogen and sulphur of which hydrogen being more active, it reacted with other elements to form chemicals on earth like CH_4 , NH_3 , H_2O and H_2S .

c. Formation of simple organic molecules :

As temperature of the earth decreased, steam condensed into water that resulted in heavy rain fall and the earth gradually cooled. Rain water got accumulated on the land to form rivers, streams, lakes, seas and oceans. The atmosphere then did not contain ozone layer and thus ultra-violet radiations reached the surface of earth directly. Under the influence of available energy sources such as ultra-violet

rays, radiations, lightening and volcanic activities, the early molecules of hydrocarbons, ammonia, methane and water underwent reactions like condensation, polymerisation, oxidation and reduction. These reactions resulted in formation of simple molecules like monosaccharides, amino acids, purines, pyrimidines, fatty acids, glycerol, etc. All these simple organic molecules accumulated at the bottom of water bodies. Haldane described it as the “**hot dilute soup**” or “**primitive broth**”. It did not show any degradation due to absence of free oxygen and enzymes (non-enzymatic reducing atmosphere).

d. Formation of complex organic molecules:

The primitive broth was neutral and free from oxygen. Polymerisation took place and simple organic molecules aggregated to form new complex organic molecules like polysaccharides, fats, proteins, nucleosides and nucleotides. Polymerisation of amino acids formed **protoproteins** which later formed proteins. Formation of protein molecules is considered as landmark in the origin of life. Proteins (enzymes) accelerated the rate of other chemical reactions.

e. Formation of Nucleic acids :

Nucleotides may have been formed by the reaction between phosphoric acid, sugar and nitrogenous bases (purines and pyrimidines). Number of nucleotides join together to form nucleic acids (RNA, DNA). Nucleic acids acquired self-replicating ability which is a fundamental property of living form.

f. Formation of Protobionts or Procells :

Nucleic acids along with inorganic and organic molecules formed the first form of life called **protobionts**. Protobionts are the prebiotic chemical aggregates having some properties of living system.

Protobionts are formed due to coacervation i.e. aggregation of organic molecules. Oparin (1924) called them **coacervates** and Sidney Fox called **protenoids** or **microspheres**.

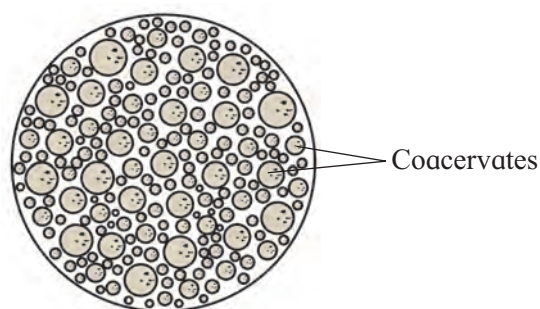


Fig. 5.1 : Coacervates

Coacervates are colloidal aggregations of hydrophobic proteins and lipids (lipid bubbles). Coacervates grew in size by taking up material from surrounding aqueous medium. As they grew, they became thermodynamically unstable and split into smaller units, comparable to daughter cells of budding organisms. Microspheres are protenoids formed from colloidal hydrophilic complexes surrounded by water molecules. These bodies may have outer double-membrane, like primitive cell. Diffusion and osmosis may have occurred across the membrane. They were more stable than coacervates. Coacervates and microspheres were non-living colloidal aggregations of lipids and proteinoids respectively. They had some basic properties of living cells, such as growth and division. These colloidal aggregations turned into first primitive living system called **eobionts** or protocell.

g. Formation of first cell :

When RNA or DNA system developed within protocells, they look like bacteria or viruses. They regulated various metabolic activities. First cell was anaerobic, heterotrophic and obtained energy by chemoheterotrophic processes.

Urey and Miller's Experiment :

Stanley L. Miller and his teacher Harold C.

Urey provided the first experimental evidence in support of chemical evolution theory of Oparin.

They designed a glass-apparatus called spark-discharge apparatus.

The apparatus (Fig. 5.2) was first sterilized and evacuated. Methane, ammonia and hydrogen gases were pumped in the proportion of 1:2:2 into the glass chamber. A tube carrying water vapour was also connected to the chamber. Lightning effect was mimicked by electric discharge carbon arc spark in the chamber. Process of evaporation and precipitation was also simulated by the use of heating mantle and condenser respectively.

The mixture of CH_4 , NH_3 , H_2 was exposed continuously to electric discharge for several days causing the gases to interact, after which these were condensed. The liquid collected in the U-tube turned brown. Chemical analysis of this liquid reported the presence of simple organic compounds. (urea, amino acids, lactic acid, etc). This experiment strongly supports that the simple molecules present in the earth's early atmosphere combined to form the organic building blocks of life.

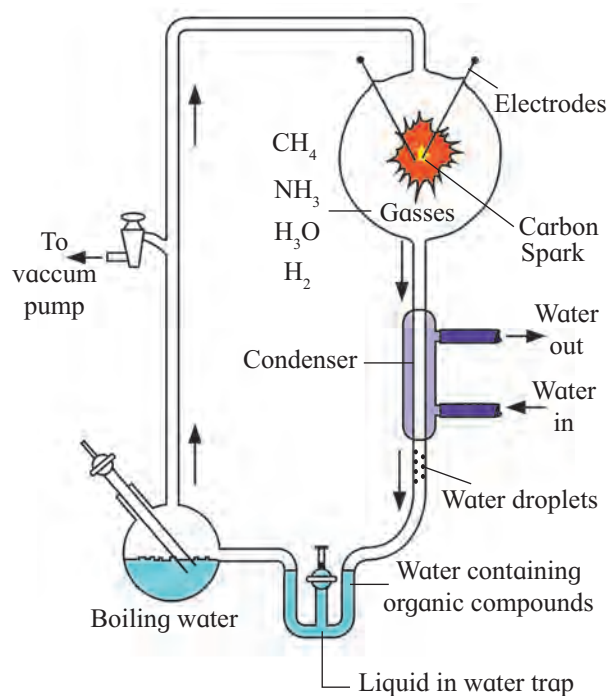


Fig. 5.2 : Urey and Miller's Experiment

RNA World Hypothesis :

Oparin-Haldane theory and Miller-Urey experiment gives us an understanding that pathway of origin of life on earth goes from non-living to living. 'Like begets like' necessitates presence of stable genetic material and cellular machinery to carry out routine activities essential for survival.

We are also aware that certain proteins which we call enzymes, catalyse the chemical reactions in the cell. It was in 1980 that Sidney Altman and Thomas Cech independently found out that RNAs can also act as biocatalysts. These catalytic RNAs are called as Ribozymes. For this discovery, Altman and Cech earned Nobel Prize in chemistry in 1989.

This discovery provided important support for RNA World hypothesis. The hypothesis suggests that early life (first) must have been based exclusively on nucleic acids, most probably RNA. It was first proposed by Carl Woese, Francis Crick and Leslie Orgel in 1960, long before discovery of ribozymes.

Fact is that RNA is found abundantly in all living cells, it is structurally related to DNA and chains of RNA can evolve or undergo mutations, replicate and catalyse reactions, all support this hypothesis.

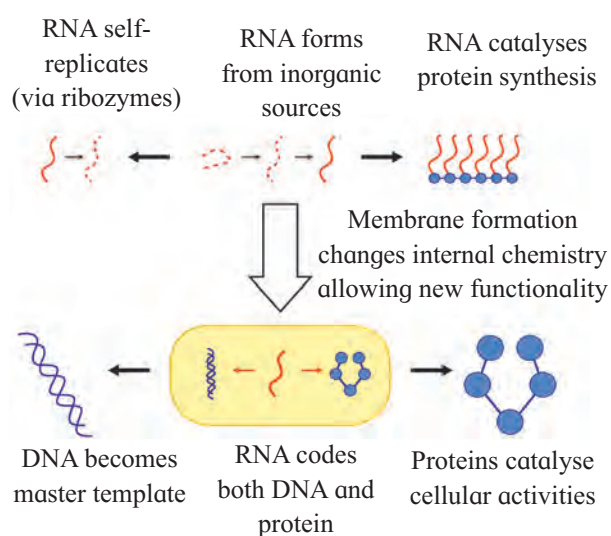


Fig. 5.3 : RNA World Hypothesis

Besides, biomolecules like Acetyl-Co-A have a nucleotide in their molecular structure. Major evidence is existence of ribosome (the protein assembly unit) in the cell. In ribosomes, translation process is catalysed by RNA. (Refer chapter- Molecular Basis of Inheritance).

These molecules might have undergone repeated replication and mutation forming varieties of RNA molecules with varying sizes and catalytic properties. Eventually they might have developed their own protein coats and machinery to survive the assembly of primitive cell. In due course, a double stranded stable structure, the DNA, might have been formed and thus continued the ongoing journey which resulted in rich biodiversity on earth.

5.3 Organic Evolution :

Evolution (Latin word, e = from; volvere = to roll) means the act of unrolling or unfolding of nature. It brings about orderly changes from one form to another. These changes result in descendants becoming different from the ancestors.

Organic evolution can be defined as slow, gradual, continuous and irreversible changes through which the present day complex forms of the life developed (or evolved) from their simple pre-existing forms.

According to **Charles Darwin**, evolution is 'descent with modification'. You have already studied Lamarck's **Theory of inheritance of acquired characters** in 10th std. According to this theory, the traits are acquired due to internal force, changes in environment, new needs and the use and disuse of organs. After several generations, it gives rise to new species.

This theory was disproved by a German biologist August Weismann, who cut the tails of many rats for several generations but could not find any change in size of tail even after

21 generations. He concluded that variations produced in somatic cells (somatoplasm) are not inherited while variations produced in germ cells (germplasm) are inherited to next generation and he proposed the **Theory of continuity of Germplasm**.

5.4 Darwinism :

Before Darwinism, several theories were proposed to explain the process of organic evolution.

Lamarck (1809) published 'Theory of origin of acquired characters' which then was ruled out.

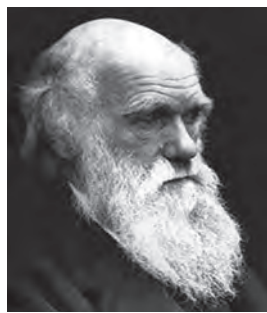


Fig. 5.4 : Charles Darwin (1809 - 1882)

Darwinism (Theory of origin of species by Natural Selection).

Charles Darwin as a naturalist travelled extensively the world over from 27th December 1831 to October 1836 and returned to England. Based on his collections of living and fossil material he published a book, "on the origin of species by means of Natural Selection", in 1859. Before publishing his book, he was very much influenced by C. Lyell (Geologist) who pointed out that natural forces which existed in the past are the same as those existing at present. Darwin also observed the changes in the earth's crust along the coast of South America, caused due to natural forces. He observed variety of strange plants and animals in the Galapagos islands and other nearby islands in the Archipelago in terms of variations between the tortoises and finches. Similar observations were also made by Wallace. He was also influenced by R. Malthus, an Economist, who worked on human population mentioning that

the reproductive potential of humans is greater than others. This resulted in multiplication of human population without corresponding increase in the food supply. This increase lead to competition and struggle for existence of human species.

He was also influenced by plant/animal breeders who by using artificial selection, altered the characteristic of cultivated plant and domestic animals.

Darwinism is based on five main postulates:

- 1. Overproduction (Prodigality of nature)**
- It is the natural tendency to produce more number of progeny in geometric ratio, for perpetuation of the species. He observed prodigality potential of many species of plants and animals e.g. Salmon fish produces about 28 lakh eggs in a single season. In a span of 750 years, single pair of elephants would produce 19,000,000 elephants. However the size of given species in a given area remains relatively constant because of fluctuations that occur seasonally.
- 2. Struggle for existence** - Tendency of over production leads to the struggle for existence between the members of population. The struggle is for limited supply of food or to overcome adverse environmental conditions, for a space and mate, and to escape from enemies etc.
- 3. Organic variations** - The variations speak for all kinds of differences that occur in morphology, physiology, nutrition, habit, behavioural patterns etc. Darwin recognized these variations as raw material for evolution. Variations were observed among members of the same species and even in different species.
- 4. Natural selection** - Organic variations can serve as evidence for some organisms that are better adapted to survive under existing environmental conditions than the others. In the struggle for existence, organisms with favourable variations are

Objections to Mutation Theory :

- i. The large and discontinuous variations observed by Hugo de Vries were actually due to chromosomal aberrations whereas gene mutations usually bring about minor changes.
- ii. Rate of mutation is very low as compared to the requirement of evolution.
- iii. Chromosomal aberrations have little significance in evolution as they are quite unstable.



Always Remember

1. According to Darwin variations are small and directional whereas mutations are large, sudden, random and directionless.
2. Darwin believed that the gradual inheritable variations over a long period of time, lead to speciation (formation of new species) while de Vries believed that mutations are the cause of speciation.
3. A single step large mutation is called saltation.

5.6 Modern Synthetic Theory of Evolution :

- It is the result of true synthesis of all biological disciplines. Studies pertaining to genetical, ecological, anatomical, geographical, palaeontological etc. were pursued to explain mechanism of evolution. Also due importance was given to both mutations and natural selection.
 - R. Fischer, J. B. S. Haldane, T. Dobzhansky, J. Huxley, E. Mayr, Simpson, Stebbins, Fisher, Sewall Wright, Mendel, T. H. Morgan etc. are the main contributors of modern theory of evolution.
 - Stebbins in his book discussed five key factors viz. gene mutations, mutations in the chromosome structure and number, genetic recombinations, natural selection and reproductive isolation. These contributed in the evolution of new species.
- All individuals of the same species constitute a population. The populations occur in small groups of 'interbreeding populations'. Such small interbreeding group of a population is referred as '**Mendelian population**'.
 - The total genetic information encoded in sum total of genes in a Mendelian population is called **gene pool**. Simply, gene pool means the total number of genes of all individuals in a population. The gametes produced by individual furnish a pool of genes of next generation from which the genes will be selected.
 - The migration of population effectively alter the gene pool. The gene pool also changes due to replacement of one generation by another in the Mendelian population. Thus any change in the gene pool affects population.
 - Genes are arranged linearly on the chromosome and occupy a definite position (locus). According to Mendel, every gene that influences a trait has two alleles. The proportion of an allele in the gene pool, to the total number of alleles at a given locus, is called **gene frequency**.
 - Modern synthetic theory comprises five main factors that are broadly divided into three main concepts - a. **genetic variations** caused due to various aspects of mutation, recombination and migration. b. **natural selection** and c. **isolation**. These help in explaining the evolution of species.
- a. **Genetic variations** : The change in gene and gene frequencies, is known as genetic variation. Genetic variations are caused by following factors :
 - i. **Gene Mutation** : A sudden permanent heritable change is called mutation. Mutation can occur in the gene, in the chromosome morphology and in chromosome number. Mutation that occurs

within the single gene, is called **point mutation** or **gene mutation**. This leads to the change in the phenotype of the organism, causing variations.

ii. **Genetic recombination** : In sexually reproducing organisms, during gamete formation, exchange of genetic material occurs between non-sister chromatids of homologous chromosomes. This is called **crossing over**. It produces new genetic combinations which result in variation. Fertilization between opposite mating gametes leads to various recombinations resulting into the phenotypic variations causing change in the frequencies of alleles.

iii. **Gene flow** : Gene flow is movement of genes into or out of a population. Gene movement may be in the form of migration of organism, or gametes (dispersal of pollens) or segments of DNA (transformation). Gene flow also alters gene frequency causing evolutionary changes.

iv. **Genetic drift** : Any random fluctuation (alteration) in allele frequency, occurring in the natural population by pure chance, is called genetic drift. For example, when the size of a population is severely reduced due to natural disasters like earthquakes, floods, fires, etc. thus cause elimination of particular alleles from a population. Smaller populations have greater chances for genetic drift. It will result in the change in the gene frequency. Genetic drift is also an important factor for evolutionary change.

v. **Chromosomal aberrations** : The changes in chromosome number (gains and losses) and the changes in the morphological structure of chromosome/s (deletion, inversion, etc.) constitute the chromosomal aberrations. The changes in the genes arrangement (order or sequence) results in the variations. Chromosomal aberrations occur due to -

a. **Deletion** : Loss of genes from chromosome.

b. **Duplication** : Genes are repeated or doubled in number on chromosome.

c. **Inversion** : A particular segment of chromosome is broken and gets reattached to the same chromosome in an inverted position due to 180° twist. There is no loss or gain of gene complement of the chromosome.

d. **Translocation** : Transfer (transposition) of a part of chromosome or a set of genes to a non-homologous chromosome is called translocation. It is effected naturally by the transposons present in the cell.

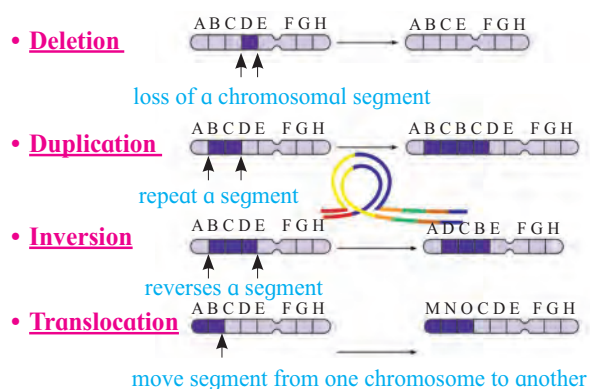


Fig. 5.5 : Chromosomal aberrations

b. **Natural selection** :

According to Darwin, natural selection is the main driving force behind the evolution. This holds that genetic variations rise within the population. The 'fittest' will be at the selective advantage and will be more likely to produce more offsprings than the rest, The 'fit' continues to enjoy greater survival and reproductivity. New species will eventually evolve.

Alternatively, natural selection is the process by which better adapted organisms grow and produce more number of offsprings in the population.

It brings about evolutionary changes by favouring differential reproduction of genes that bring about changes in gene frequency from one generation to next generation.

Selection against harmful mutations leads to a mutation balance, in which allele frequency of harmful recessives remain constant generation after generation.



Fig. 5.6 : Natural selection (*Biston betularia* and *Biston carbonaria*)

Natural selection encourages those genes or traits that assure highest degree of adaptive efficiency between population and its environment. **Industrial melanism** is one of the best example for natural selection. In Great Britain, before industrialization (1845) grey white winged moths (*Biston betularia*) were more in number than black-winged moth (*Biston carbonaria*).

These moths are nocturnal and during day time they rest on tree trunk. White-winged moth can camouflaged (hide in the background) well with the lichen-covered trees that helped them to escape from the predatory birds. on other hand, the black-winged moth resting on lichen covered tree trunks were easy victims for the predatory birds and their number was reduced. During industrial revolution, large number of industries came up in Great Britain. The industries released black sooty smoke that covered and killed the lichens growing on tree and turn the tree black due to pollution.

This change become an advantage to the black-winged moth that camouflaged well with the black tree trunks and their number increased while the white-winged moth become victims to predatory birds due to which their number reduced. Thus natural selection has resulted in the establishment of a phenotypic traits in changing the environmental conditions.

c. Isolation :

Isolation is the separation of the population of a particular species into smaller units which prevents interbreeding between them. Some barrier which prevents gene flow or exchange of genes between isolated populations, is called isolating mechanism.

Number of isolating mechanisms operate in nature and may promote divergence and speciation may occur. The isolating mechanisms are of two types namely, geographical isolation and reproductive isolation.

I. Geographical Isolation :

It is also called as physical isolation. It occurs when an original population is divided into two or more groups by geographical barriers such as river, ocean, mountain, glacier etc. These barriers prevent interbreeding between isolated groups.

The separated groups are exposed to different kinds of environmental factors and they acquire new traits by mutations. The separated populations develop distinct gene pool and they do not interbreed. Thus, new species have been formed by geographical isolation. E.g. Darwin's Finches.

II. Reproductive Isolation :

Reproductive isolations occurs due to change in genetic material, gene pool and structure of genital organs. It prevents interbreeding between population.

Types of Isolating Mechanisms :

A. Pre-mating or pre-zygotic isolating mechanism : This mechanism prevent fertilization and zygote formation.

i. Habitat isolation or (Ecological isolation) : Members of a population living in the same geographic region but occupy separate habitats so that potential mates do not meet.

- ii. **Seasonal or temporal isolation** : Members of a population living in the same geographic region but are sexually mature at different years or different times of the year.
 - iii. **Ethological isolation** : Due to specific mating behaviour the members of population do not mate.
 - iv. **Mechanical Isolation** : Members of two population have difference in the structure of reproductive organs.
- B. Post-mating or Post-zygotic barriers :**
- i. **Gamete mortality** - Gametes have a limited life span. Due to one or the other reasons, if union of the two gametes does not occur in the given time, it results in the gamete mortality.
 - ii. **Zygote mortality** - Here, egg is fertilized but zygote dies due to one or the other reasons.
 - iii. **Hybrid sterility** - Hybrids develop to maturity but become sterile due to failure of proper gametogenesis (meiosis). e.g. Mule is an intergeneric hybrid which is sterile.



Can you tell?

1. What is variation?
2. What is mutation?
3. What is gene frequency?
4. Why variations occur in population?

5.7 Mechanism of organic evolution :

One has to give the importance to the population while considering the mechanism of evolution. It is the population that evolves and not its individual members. Individual's role is to pass its genetic variation to its offspring.

The following are the basic processes which bring about evolution viz. Mutations, gene recombination, gene flow (migration), genetic drift, natural selection, isolation and speciation.

Mutations - These are permanent heritable changes in the genetic material of an organism.

Mutations are already described earlier in this chapter. Gene mutations will alter the genetic make up and the gene pool.

Gene recombination - These are variation produce due to coming together of alleles during sexual reproduction. Gene recombinations occur due to random union of gametes, anaphasic separation of chromosomes and crossing over.

Gene flow - It is the transfer of gene during interbreeding of populations that are genetically different. As explained earlier in this chapter gene flow is due to emigration and immigration. Its brings about changes in the allele frequency.

Genetic drift - Any alteration in allele frequency of a natural population by chance, is called **genetic drift**. Concept of genetic drift was first given by Sewall Wright, hence, called **Sewall Wright effect**. For example, elimination of a particular allele from a population due to events like accidental death prior to mating of the organism. Genetic drifts are random or directionless. The effect of genetic drift is more significant in small population than in large population. Due to genetic drift, some alleles of a population are lost or reduced by chance and some others may be increased. Sometimes, a few individuals become isolated from the large population and they produce a small new population in new geographical area. The allele frequency of the new population become suddenly different. The original members that drifted and established the new population become 'founders' and the effect is called **founder effect**.

The bottle neck effect is a type of genetic drift is seen when much of a population is killed due to a natural disaster (sunami, floods, tornado, disease epidemic, etc.) and only a few individuals are left to begin a new population.

Natural selection - It is a process by which better adapted individuals with useful variations are selected by nature and leave greater or more number of progenies (Differential reproduction).

Type of Natural selection :

a. Stabilizing selection : (Balancing selection)

1. Here more individuals of a population acquire a mean character value.
2. It tends to favour the intermediate forms and eliminate both the phenotypic extreme. For e.g. More number of infants with intermediate weight survive better as compared to those who are over-weight or under-weight.
3. It reduces variations.
4. It does not lead to evolutionary change but tends to maintain phenotypic stability within the population, therefore, it is described as stabilizing selection.
5. Genetically stabilizing selection represents a situation where a population is adapted to its environment.

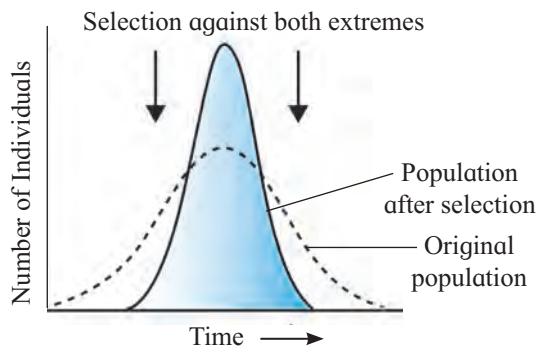


Fig. 5.7 : Stabilising selection

b. Directional selection :

1. In this type, more individuals acquire a value other than the mean character value.
2. Natural selection acts to eliminate one of the extremes of the phenotypic range and favour the other. e.g. systematic elimination of homozygous recessives.
3. Directional selection operates for many generations, it results in an evolutionary trend within a population and shifting of the peak in one direction.
4. Example - Industrial melanism, DDT resistant mosquito etc.

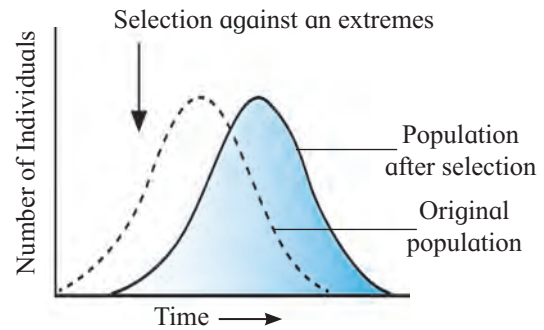


Fig. 5.8 : Directional selection

c. Disruptive Natural selection :

1. Here more number of individuals acquire peripheral character value at both ends of the distribution curve.
2. Nature selects extreme phenotypes and eliminates intermediate. Hence two peaks are formed in distribution of traits.
3. This kind of selection is rare.
4. It ensures the effect on the entire gene pool of a population, considering all mating types or systems.
5. Example - It was observed in the different beak size of African seed cracker finches. The birds have different size of beak and they feed on seeds. The available seeds were of two kinds small and large sized seeds. Large beak sized birds feed on large seeds while small beak sized birds feed on small seeds and their number was increased. Intermediate beak sized birds are unable to feed on either type of seeds so their population was decreased gradually and then eliminated by natural selection.

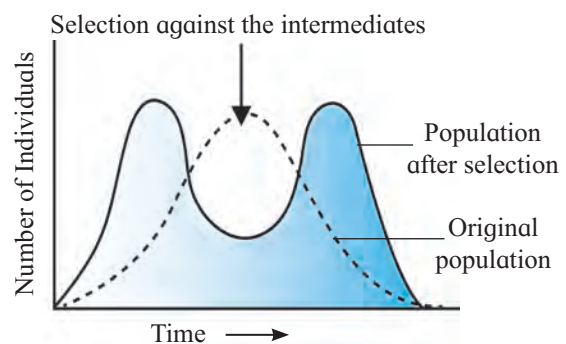


Fig. 5.9 : Disruptive selection

Isolation - It is separation of a single interbreeding population into subunits. Isolation restricts gene flow between discrete (non-continuous) population due to different barriers like geographical barriers. (This part is already explained in detail earlier in this chapter)

Speciation - The sub units of single interbreeding population due to the geographical barriers like river, mountains, desert, sea etc. become isolated in such a way that their interbreeding is prevented. This will finally lead to origin of new species (i.e. speciation). (Discussed in detail ahead in this chapter).

5.8 Hardy-Weinberg's principle :

It is also known as Hardy-Weinberg's equilibrium law. The law states that 'at equilibrium point both the gene (allele) frequency and genotypic frequency remain constant from generation to generation'. It occurs only in the diploid, sexually reproducing, large, free interbreeding population in which mating is random and no selection or other factors are present for changing the allele frequency. e.g. A single locus has two alleles (A and a). The frequencies of these allele are p and q respectively. The allele frequency for any locus is always one. i.e. $p + q = 1$. The genotypic frequencies of both the alleles are represented by $(p + q)^2 = 1$. The binomial expansion of this is $P^2 + 2pq + q^2 = 1$ i.e. $AA = p^2$, $aa = q^2$ and for $2Aa = 2pq$.

Hence $p^2 + 2pq + q^2 = 1$. This is a binomial expansion of $(p + q)^2$.

This can be explained by punnett square as follows :

		Hybrid A a	
		A (p)	a (q)
Hybrid Aa	A (p)	AA (p^2)	Aa (pq)
	a (q)	Aa (pq)	aa (q^2)

Like allele frequency, the genotypic frequencies together are also equal to 1.

$$\text{i.e. } AA + 2Aa + aa = 1$$

There are few factors such as gene migration (gene flow), genetic drift, mutation, genetic recombinations, natural selection, non-random mating, etc. which affect or change the Hardy-Weinberg equilibrium,

If these factors do not occur in the population, then population is genetically stable or non evolving population.



Can you tell?

In which conditions the gene frequency of a population will remain constant?

5.9 Adaptive Radiation :

The process of evolution which results in transformation of original species to many different varieties, is called, adaptive radiation.

Darwin's Finches is one of the best example of adaptive radiation. During his visit to Galapagos Islands, Charles Darwin also noticed a variety of small birds. These birds are called Darwin's finches.



Fig. 5.10 : Darwin's finches

Darwin concluded that the American main land species of bird was the original one from which they migrated to the different islands of Galpagos. They adapted to the different environmental conditions of these islands. From original seed eating features many other forms with altered beaks evolved into insectivorous features.

Another example of adaptive radiation is Australian Marsupials. In Australia, there are many marsupial mammals who evolved from common ancestor.



Can you tell?

1. What is carbon dating and how does it work ?
2. What is fossil ?
3. What are Homologous organs ?
4. What is Embryology ?

5.10 Evidences of organic evolution :

The theory of organic evolution states that the present day complex organisms have originated from earlier simpler forms of life.

The process of evolution is supported by evidences provided by various branches of biology such as : Palaeontology, comparative anatomy, embryology and molecular biology.

A. Palaeontology : The study of ancient life with help of fossils is called palaeontology. Fossils are the dead remains of plants and animals that lived in past in various geological layers.

- The study of fossils provides the most convincing and direct evidence of evolution.
- Fossils are formed in sedimentary rocks, amber (yellowish fossil resin), ice, peat bogs etc.
- During fossilization, the primitive forms of organisms occupy the older, lower layers and the advanced forms occupy the upper, more recent layers of the earth.

Types of fossils :

1. **Actual remains :** These are most common type of fossils. The plants, animals and human bodies got embedded in permafrost of arctic or alpine snow remain preserved in the actual state, e.g. Woolly Mammoth in Siberia. Amber or hardened resin contains preserved bodies of many insects and arthropods.

2. **Moulds :** These are the hardened encasements formed in the outer parts of organic remains which later decayed leaving cavities. Body parts of plants or animals later decay but the impression still remains and becomes permanent. For example Foot – prints are formed in this manner.

3. **Cast :** They are hardened pieces of mineral matter deposited in the cavities of moulds.

4. **Compressions :** Internal structure is absent but a thin carbon film indicates the outline of external features.

Significance of Palaeontology :

1. It is useful in reconstruction of phylogeny.
2. It helps in studying various forms and structures of extinct animals.
3. It provides record of missing link between two groups of organisms.
4. It helps in the study of habits of extinct organisms.
5. Palaeontology provides the following types of evidences.

Connecting link (missing link) :

These are fossil forms transitional or intermediate between two groups of organisms. They show some characters akin to both the groups. Thus it indicates the evolutionary line *Seymouria* (between amphibians and reptiles) and *Archaeopteryx* (between reptiles and birds).

Archaeopteryx lithographica :

It is a fossilized crow size toothed bird found from Jurassic rocks in Germany. It is known as a missing link between reptiles and birds because it shows characters of both.

Reptilian characters :

1. Presence of long tail, claws and scales on the body.
2. Single headed ribs.
3. Abdominal ribs are present which look like ribs of crocodile.
4. Jaws with homodont teeth.

- Sternum without keel.
- Bones are solid (nonpneumatic).
- Hind limbs had four clawed digits.

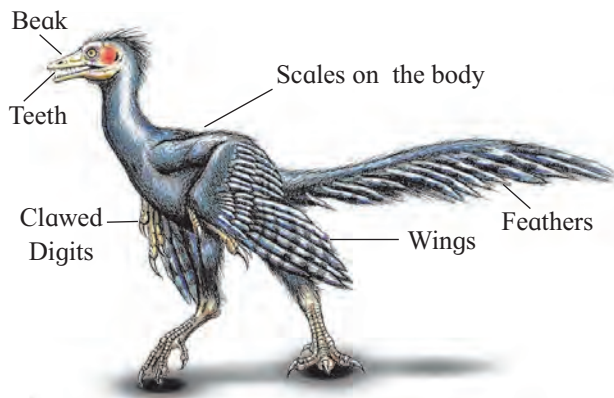


Fig. 5.11 : Archaeopteryx

Avian characters :

- Feathery exoskeleton.
- Forelimbs are modified into wings.
- Jaws are modified into beak.
- Skull bone is completely fused.
- Large rounded cranium.
- Cranium with large orbits and a single condyle.
- Limb bones are bird like.
- Hind limbs with four toes first toe is opposible.

Thus from the above study it is very clear that birds evolved from reptiles. Huxley justified this by calling **birds as glorified reptiles**.



Activity :

Complete the following chart.

Animals	Connecting link between
1. Balanoglossus
2.	annelida and arthropoda
3. Lung fishes
4.	reptiles and mammals
5. Seymouria
6. Ichthyostega	fishes and amphibia

? Do you know ?

First fossil of *Archaeopteryx* was found in Jurassic rocks of Bavaria. It was discovered in 1861 and preserved in British museum by Andreas Wanger.

The second specimen found in 1877, known as *Archaeornis*, is kept in the Berlin museum.

B. Morphology : Morphology deals with study of external structures while, anatomy deals with study of internal structures. From comparative study of morphology and anatomy we can understand the evolutionary aspects in the form of homologous, analogous and vestigial organs.

a. Homologous organs : Homologous organs are those organs, which are structurally similar but perform different functions.

For example :

- Forelimbs of vertebrates such as lizzard, bird, bat, horse, whale and man,

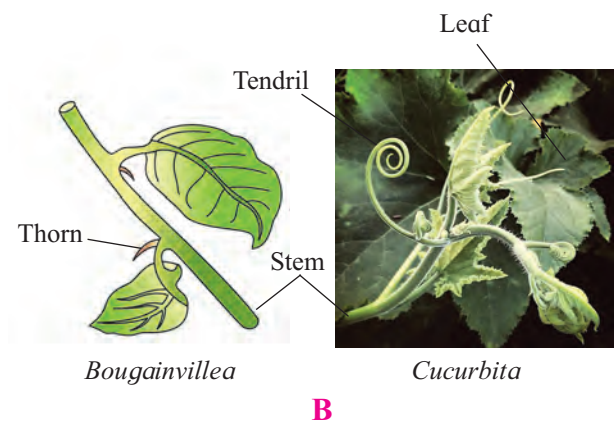
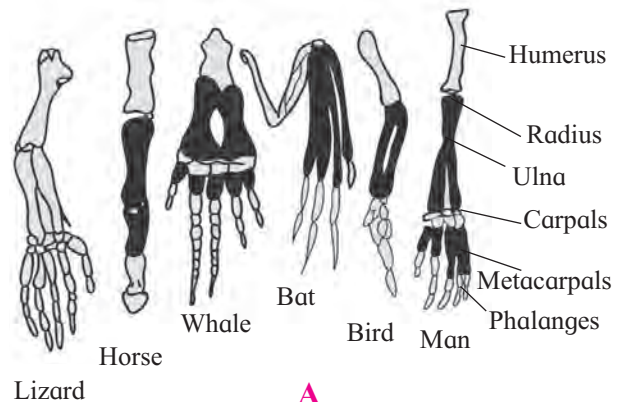


Fig. 5.12 : Homologous organs

all of them have humerus, radius-ulna, carpals, metacarpals and phalanges in their forelimbs. Forelimbs of these vertebrates are structurally similar but perform different functions.

2. Vertebrate heart and brain.
3. In plants, thorns of *Bougainvillea* and tendrils of *Cucurbita* represent homology. Thorns of Bael and tendrils of passion flower are homologous.

The structural similarities between the homologous organs indicates that they have a common ancestry.

Differences in homologous organs are examples of divergent evolution or adaptive radiation.

b. Analogous organs : Analogous organs are those which are structurally dissimilar but functionally similar. These organs have external superficial similarity due to similar functions but they are different anatomically.

For e.g. wings of butterfly (insects) and of birds look superficially alike but they are no anatomically similar structures though they perform similar functions.

Other examples of analogous organs.

1. Eye of the octopus (mollusca) and of mammals. They differ in their retinal position, structure of lens and origin of different eye parts.
2. The flippers of penguins (birds) and dolphins (mammals).
3. Sweet potato (root modification) and potato (stem modification) store food in form of starch.
4. Terminal leaflets of pea and apical bud in *vitis* are modified into tendrils.

Analogous organs leads to convergent evolution i.e. different organisms shows same superficial structural similarities due to similar functions or habitat. These organs do not help to trace the common ancestry. Thus analogous

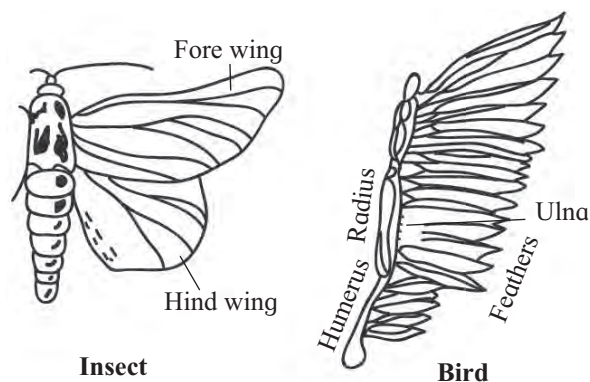


Fig. 5.13 : Analogous organs

organs do not have significant role in evolution.

c. Vestigial organs : (Rudimentary organs)

Vestigial organs are imperfectly developed and non-functional, degenerate structures which were functional in some related and other animals or in ancestors. The vestigial organs are no longer required by the organism but indicate the relationship with those organisms where these organs are fully developed.

Examples : Human beings show some vestigial organs like -

1. Presence of vestigial nictitating membranes.
2. Presence of wisdom teeth (third molars).
3. Coccyx (tail bone) : It is greatly reduced in man since the tail is of no use due to erect posture.

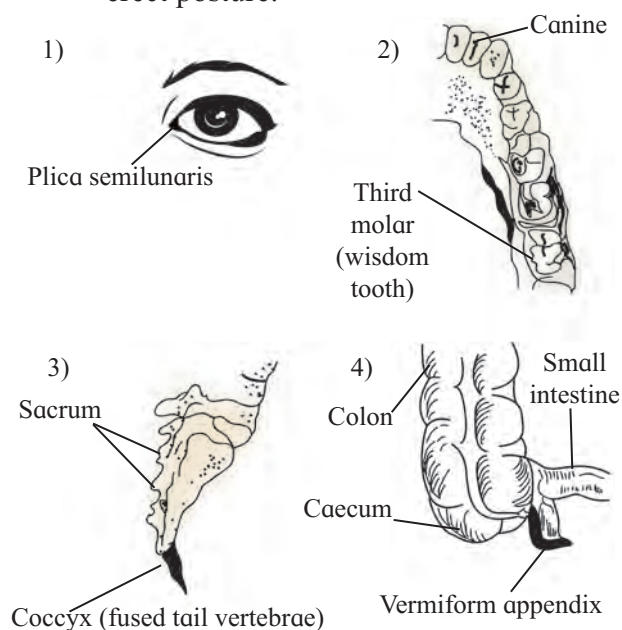


Fig. 5.14 : Vestigial organs

4. Vermiform appendix and the caecum. It is functional in herbivorous mammals for digestion of cellulose. In man due to eating of cooked food it has lost its function.

Presence of these vestigial organs provide evidence that man has (evolved) descended from simple primates.



Can you recall?

Give examples of embryological evidences studied in lower classes.

d. Molecular Evidences :

1. Cell is the basic structural and functional unit of life in all organisms.
2. Similarities in proteins and genetic material performing a similar function among diverse organisms gives evidence of a common ancestry.
3. Basic metabolic activities also occur in a similar manner in all organisms.
4. ATP is the energy source in all living organisms.

5.11 Speciation :

The process of formation of a new species from the pre-existing species is called speciation.

Species is a group of similar organisms that can interbreed and produce a fertile offspring in nature. New species are formed by the following modes.

a. Intraspecific Speciation :

- i. **Allopatric speciation** : It is the formation of a new species due to separation of a segment of population from its original population by distance or by a geographical barrier cutting across the species range. e.g. creeping glaciers, development of mountains. Migration of individuals also causes allopatric speciation. The mode of evolution here is called **adaptive radiation**

e.g. 14 different species of finches in Galapagos islands and several marsupial species in the Australian continent.

- ii. **Sympatric speciation**: It is the formation of species within single population without geographical isolation. These are formed due to reproductive isolation. e.g. *Cichlid* fishes in Lake Victoria. Mutations are helpful in sympatric speciation.

b. Interspecific Speciation :

Hybridisation :

Two different species on crossing may give rise to hybrid that on doubling of chromosomes form a new species. e.g. . *Triticum aestivum* (hexaploid) and *Gossypium herbaceum* (tetraploid).



Can you recall?

1. What are fossils? why should we study fossils?
2. How do we find age of fossils ?
3. Where do we find fossils ?

5.12 Geological time scale :

The planet earth with its present biodiversity was not so when it was born. Study of fossils tells us that life forms were not the same like today's, millions of years ago (MYA). Geological time scale is used to understand the sequence of events that took place on the earth in different ages over a period of time. It is divided into six major 'Eras'. Eras ended with major environmental changes on earth resulting into extinction of some species and emergence of new species. The eras are further divided into periods and epochs based on minor but landmark events in each era. Table 5.15 shows the geological time scale at a glance.

The first life appeared on the earth some 2000 million years ago. It took billions of years for this process to take place, from protenoids to first cells the transition is still a mystery. Once formed, the living forms diversified into various groups. Life began in the sea water and

plants were the first living beings to adapt to terrestrial life. Fishes evolved and diversified. The lobefin group of fishes too got diversified. Some developed stout and strong fins and could go to land and come back to water.

The coelacanth was considered a **living fossils**. It was thought that lobefins are extinct,

but the variety of the lobefin fish, called coelacanth was caught in 1938 in South Africa.

Reptiles evolved from amphibians. They are the first true land vertebrates. They do not have to go to water for reproduction. (Hint : think of amphibian and reptilian eggs). But about 200 million years ago (mya) some reptiles

Table 5.15 : Geological time scale

Era	Period	Time MYA	Epoch	Plant life	Animal life
Cenozoic	Quaternary 0.01-2.0	0.1-0.6	Recent (Holocene)	Angiosperms, Dicots, Monocots. spread of agriculture.	Age of mammals : Development of modern man, birds, fishes and insects. Development of human culture.
		0.6 - 2.0	Pleistocene	Increase in herbs	Extinction of great mammals. Appearance of primitive man.
	Tertiary 2-6.5	2-13	Pliocene	Hard woody plants conifers, bryophytes, monocots, grasslands dominated	Emergence or origin of man. Evolution of ruminants - horse, camel, elephant.
		13-26	Miocene	Abundance of deciduous trees, origin of grasses	Formation of first man like apes. Adaptive radiation or spread of mammals.
		26-38	Oligocene	Rise of monocots and flowering plants	Extinction of Archiac mammals, Appearance of apes and monkeys. Turtles and crocodiles attained development.
		38-54	Eocene	Development of angiosperms	Diversification of placental mammals and modern birds.
Mesozoic	Cretaceous	54-65	Palaeocene	advancement of flowering plants	Arrival of early or first primates, rise of placental mammals.
		65-135		Decline of ferns - sphenopsids (horsetails) and Bennetiales. Ginkos, Gnetales. 1st appearance of flowering plants.	Extinction of Dinosaurs and toothed birds. Appearance of placental mammals and first modern birds
		135-165		Origin of angiosperms Dominance of herbaceous lycopods, ferns, conifers, cycads.	Age of reptiles. Dinosaurs dominant, Appearance of toothed birds (Archaeopteryx) Rise of marsupials.
	Triassic	165-225		Dominance of Bennetiales gymnosperms, extinction, of seed ferns	Appearance and rise of dinosaurs. Extinction of primitive amphibians. Diversification of reptiles. Rise of oviparous mammals. Therapsids, diversification of flies.

Palaeozoic	Permian	225-280		Origin of conifers Decline of arborescent lycopods Abundance of ferns, cycads and Advanced conifers	Rise of modern insects, Disappearance of trilobites, appearance of mammal like reptiles (Pelycosaur). Decline of amphibians.
	Carboniferous	280-345		Development of diverse, arborescent lycopods mosses, seed ferns and primitive conifers. age of ferns and coal forests, different fungal groups	Abundance of amphibians (age of amphibian). Appearance of reptiles and winged insects.
	Devonian	345-400		Appearance of first pro - gymnosperms. Formation of forests, wood decaying fungi, chytrids and origin of bryophytes.	Diversification of fishes. Evolution of amphibians. Appearance of ammonites
	Silurian	400-440		Appearance of lycopods and ferns. Domiance of algae, ascomycetean fungi	Appearance of first terrestrial animals, wingless insects and jawed fish
	Ordovician	440-500		Appearance of first seedless vascular land plants, abundant algae	Abundance of diversified invertebrates. Appearance of first vertebrates jawless fishes, Appearance of corals, giant cephalopods like <i>Nautilus</i> .
	Cambrian	500-590		Rhynia like plants. All types of marine algae	Abundance or age of trilobites, Diversification of invertebrate phyla.
Proterozoic		600-1600		Tracheophyte ancestors, chlorophyte ancestors bacterial single-celled protista, blue green algae	Primitive flat worms, annelids, sponges, coelenterates, primitive metazoans, scanty fossils of prokaryotes
Archaean		1600-3800		No fossil records	Origin of life Simple unicellular forms like viruses, bacteria and algae
Azoic		3800-4600		No life	Absence of living being chemical evolution Formation of earth,

* (MYA = Million Years Ago)

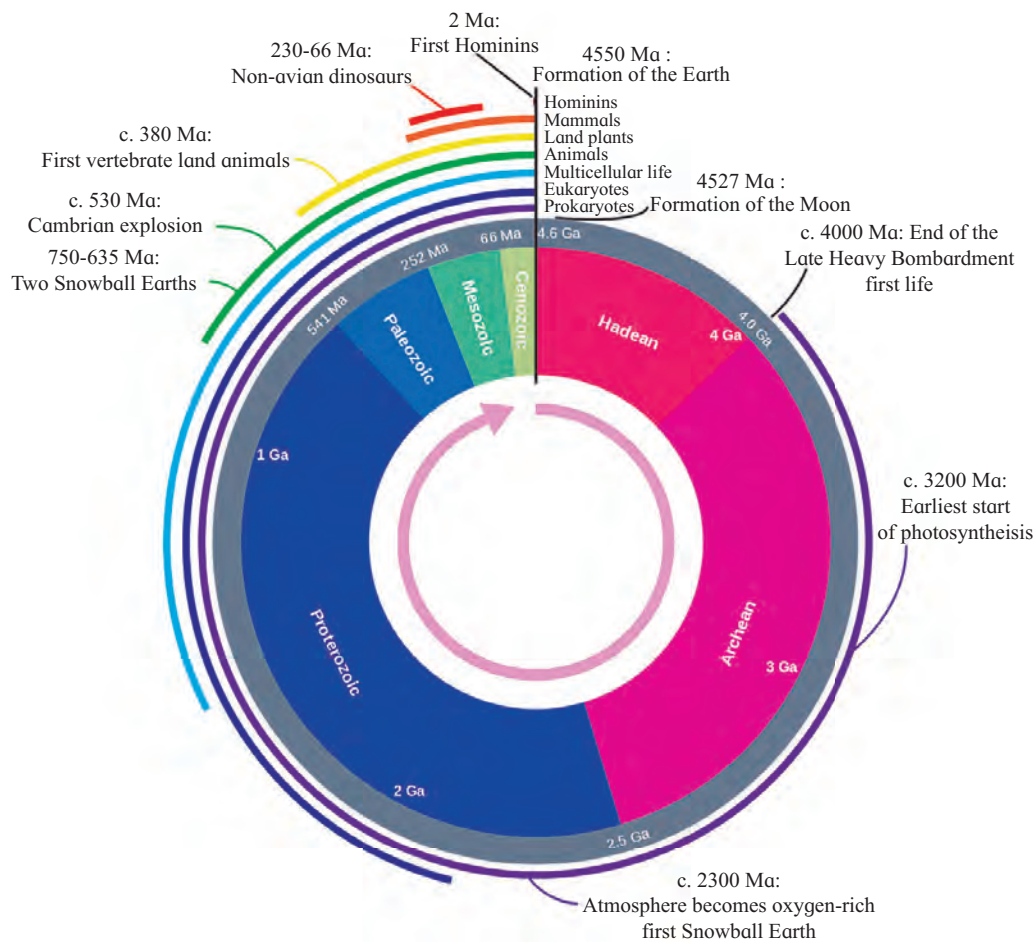


Fig. 5.16 : Geological time scale pie diagram

moved back to aquatic mode of life and gained fish like form as seen in *Ichthyosaurs*. The giant reptiles like Dinosaurs once dominated the earth but are now **extinct**. When was this? Around 65 million years ago! why it must have happened? Can we give affirmative reason for this extinction? At around the same time giant ferns were present on earth. However, they also became extinct and got converted to fossil fuels. How this must have happened?

Decline of giant reptiles marked the beginning of dominance of mammals. These viviparous organisms were more intelligent. They could avoid danger. Early mammals were small shrew like organisms, but this group **diversified**. Whales, dolphins, seals and sea cows live in water, bats are the flying mammals, kangaroo rats are fossorial, lemurs are arboreal. Major physical disturbances led

to phenomenon like the **continental drift** i.e. continents moved from their original place. As a consequence when south America joined north America, ancestral forms of horse, hippos, rabbits, etc. native to south america were dominated by north American animals. At the same time, marsupials diversified into different habitats in Australia. These survived due to lack of competition.

From the fossil records we can trace complete evolutionary history of horse, elephant, dog, etc. Human beings are the most evolved animals on the earth.



Internet my friend

You may gather information out of curiosity about geological events occurred in the past.

5.13 Human Evolution :



Use your brain power

Since your earlier school days you have been solving mysteries/puzzles labelled as **use your brain power**. Did you ever wonder why human brain has such a capacity? Why and how we evolved along these lines? What is the extent of similarity between humans, chimpanzees and monkeys ?

It has been traced that the human evolution appeared to have evolved from a tree dwelling shrew like animal. This process began in Palaeocene epoch. During this period, dwindling forests forced arboreal mammals to adapt to life on land. This descent must have been the driving force. In the following chart, it can be seen that we are most closely related to gibbons, chimpanzees and gorillas.



Do you know ?

There is a difference of only 2.5 % between DNA of chimpanzee and man while between monkey and man it is 10 %.

The major evolutionary trends in transition from ape to man are considered further. Special characteristics have been acquired by man in the course of evolution. Major changes that took place in evolution of man include increase in size and complexity of brain and enhanced intelligence, increase in cranial capacity, bipedal locomotion, opposable thumb, erect posture, shortening of forelimbs and lengthening of hind limbs, development of chin, broadening of pelvic girdle, development of lumbar curvature, social and cultural development (articulated speech, art, development of tools, etc).

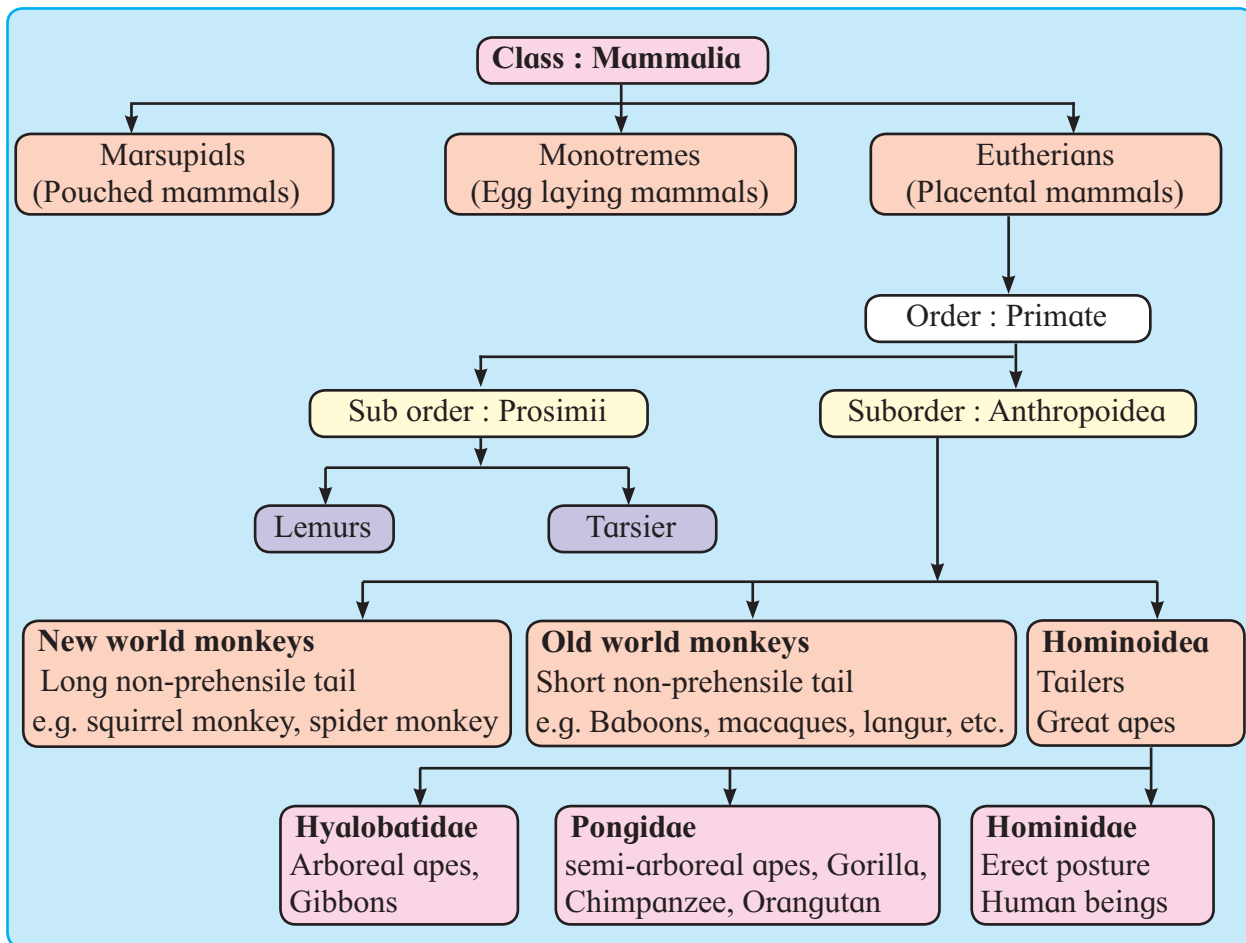


Chart 5.17 : Classification of mammals

Table 5.18 : Human evolution

Heads	Dryopithecus	Ramapithecus	Australopithecus	Homo habilis	Homo erectus	Neanderthal man	Homo sapiens
Appearance	Ape like	Man like	Connecting link between Ape and man	Handy man like	Ape man	Advanced prehistoric man	Modern man
Site of fossil record	Lake victoria of Africa, Haritalyngar of Himachal Pradesh	Shivalik Hills in India and even in Kenya	Toung in South Africa, Ethiopia, Tanzania	Olduvai Gorge Tanzania in Africa	Java and peking	Neanderthal valley, Germany	Africa
Period	Miocene epoch 20 – 25 mya	Miocene and early pliocene epoch 14 -12 mya	late pliocene or early pleistocene epoch about 4 – 1.8 mya	Late pliocene or early pleistocene 2.5 to 1.4 mya	middle of pleistocene epoch 1.5 mya ago	Late pleistocene epoch 100000 to 40000 yrs ago	
Skeltal features			Not taller than 4 feet, jaws larger, prognathus face, chin absent, lumbar curvature present.	Lower jaw, lighty built, dentition more like modern man, smaller molars	5 feet in height prognathus face, massive jaws, Huge teeth, chin absent, Bony eye brow ridges present	Heavy built short prominent brow ridges low forehead, deep jaws, chin absent, outwardly curved thigh bones	
Posture	semi erect	erect	upright	erect	erect	erect	erect
Cranial capacity			450 to 600 cc	650 to 800 cc	900 cc	1400 cc	1450 cc
Special features if any	close similarity to chimpanzee		man with ape brain	probably did not eat meat, made tools from stones, nicknamed handy man	probably ate meat, omnivorous might have used fire	used hide, burried their dead, constructed flint tools	developed distinct races. Developed cave art about 18000 yrs ago

Cranial capacity of human begins increased over a period of time and large size of frontal lobe helped in development of high forehead.

Increase in intelligence necessitated physical development so that body and brain could be used effectively and productively. Freedom of forelimbs from locomotory function and opposable thumb led to better utilization of hands for holding objects effectively and development of motor skills etc.

Bipedal locomotion, upright posture coupled with stereoscopic vision helped man to move around safely on land.

Evolutionary history of man was traced with the help of fossil remains found over a period of time.



Use your brain power

Even though the cranium of elephant is larger than that of humans, why humans are considered more intelligent than elephants ?

Some of our ancestors and their evolutionary history is shown in the table.

The above table clearly shows the gradual increase in cranial capacity, shape of skull and dentition of the ancestral humans till date.

Our journey of evolution still continues.....

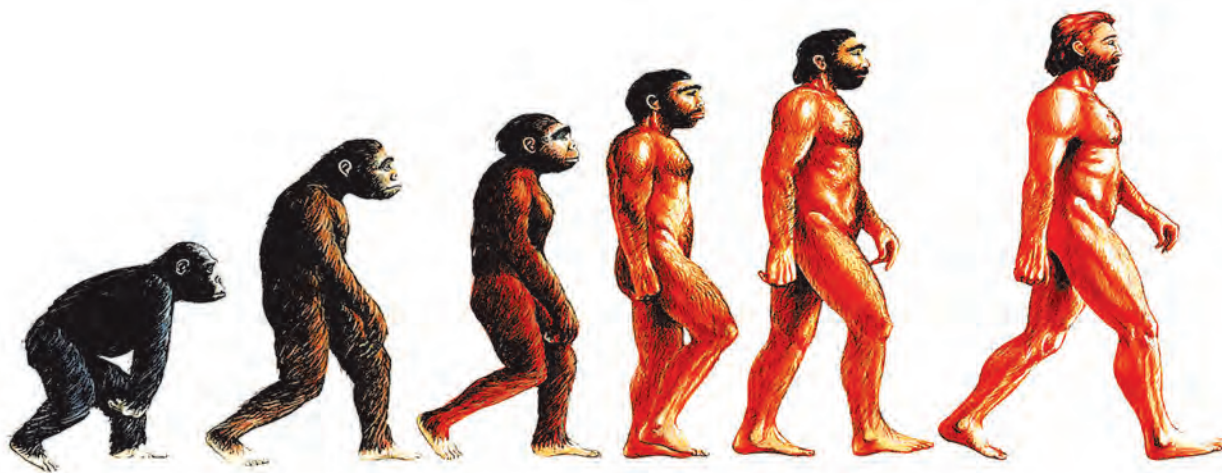


Fig. 5.19 : Human Evolution



Internet my friend

1. Recently a fossil park has been established in Gadchiroli district of Maharashtra state. Find more information about Wadadham fossil park.
2. Find out information about caves in India. One such place is in Madhya Pradesh. It is at Bhimbetka rock shelter in Raisen district. Here we can see cave paintings by prehistoric humans.








Think about it


Human being is said to be most evolved, intelligent living being. Yet we are not self sufficient. Think of various aspects for which we depend on other living beings for our survival.




Activity :

Collect the information about the organisms depicted in the following diagrams and write on the same.

<p>A. </p> <p>B. </p> <p>C. </p> <p>D. </p> <p>E. </p>	
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Exercise

Q. 1 Multiple choice questions.

- Who proposed that the first form of life could have come from pre-existing non-living organic molecules?
 - Alfred Wallace
 - Oparin and Haldane
 - Charles Darwin
 - Louis Pasteur
- The sequence of origin of life may be-
 - Organic materials- inorganic materials – Eobiont- colloidal aggregates- cell.
 - Inorganic materials – organic materials – colloidal aggregates – Eobiont- cell.
 - Organic materials- inorganic materials – colloidal aggregates - cell
 - Inorganic materials- organic materials – Eobiont- colloidal aggregates - cell
- In Hardy - Weinberg equation , the frequency of homozygous recessive individual is represented by :-
 - P^2
 - pq
 - q^2
 - $2pq$
- Select the analogous organs-
 - Forelimbs of whale and bat
 - Flippers of dolphins and penguin
 - Thorn and tendrils of bougainvillea and cucurbita.
 - Vertebrates hearts or brains.
- Archaeopteryx* is known as missing link because it is a fossil and share characters of both-
 - Fishes and amphibians
 - Annelida and arthropoda.
 - Birds and reptiles
 - Chordates and nonchordates.
- Identify the wrong statement regarding evolution.
 - Darwin's variations are small and directional.
 - Mutations are random and non-directional.
 - Adaptive radiations leads to divergent evolution.
 - Mutations are non- random and directional.
- Gene frequency in a population remain constant due to –
 - Mutation
 - Migration
 - Random mating
 - Non- random mating
- Which of the following characteristic is not shown by the ape?
 - Prognathous face
 - tail is present
 - Chin is absent
 - forelimbs are longer than hind limbs
- can be considered as connecting link between between ape and man.
 - Australopithecus
 - Homo habilis
 - Homo erectus
 - Neanderthal man.
- The cranial capacity of Neanderthal man was
 - 600 cc
 - 940 cc
 - 1400 cc
 - 1600 cc

Q. 2 Very short answer question.

- Define the following terms-
 - Gene pool
 - gene frequency
 - Organic evolution
 - Population
 - Speciation

2. What is adaptive radiation?
3. If the variation occur in population by chance alone and not through natural selection and it brings change in frequencies of an allele. What is it called?
4. State the Hardy – Weinberg eqnilibrium.
5. What are homologous organs?
6. What is vestigial organ?
7. What is the scientific name of modern man?
8. What is coacervate?
9. Which period is known as “age of Reptilia”?
10. Name the ancestor of human which is described as man with ape brain.

Q. 3 Short answer question.

1. Write a note on Genetic drift.
2. Enlist the different factors that are responsible for changing gene frequency.
3. Draw a graph to show that natural selection leads to disruptive change.
4. Give the significance of fossils.
5. Write the objections to Mutation theory of Hugo de vries.
6. What is disrruptive selection? Give example.

Q. 4 Match the following.

Columu- I	Column – II
1. August Weismann	a. Mutation theory
2. Hugo de Vries	b. Germplasm theory
3. Charles Darwin	c. Theory of inheritance of acquired characters
4. Lamarck	d. Theory of natural selection

Q. 5 Long answer questions.

1. Would you consider wings of butterfly and bat as homologous or analogous and why?
2. What is adaptive radiation? Explain with suitable example.
3. By taking industrial melanism as one example, explain the concept of natural selection.
4. Describe the Urey and Miller’s experiment.
5. What is Isolation? Describe the different types of reproductive Isolatons.
6. What are Genetic variations? Explain the different factors responsible for genetic variations.

Q. 6 Complete the chart.

Era	Dominating group of animal
1. Cenozoic
2.	Reptiles
3. Palaeozoic
4.	Invertebrates

Project :

Prepare a diagrammatic chart of the chemical evolution of life.

**Can you recall?**

1. Which are the various parts of plant body?
2. What are the functions of various parts of plant body?
3. Which plant tissues are involved in transport of water and minerals?

**Use your brain power**

You know that we need a water pump to lift water at top of the building. but, how does plants lift the water from soil upto canopy without any pump?

Plant obtains variety of substances like water, minerals, nutrients, food and gases like O_2 and CO_2 , from its surroundings. Productivity in plants is mainly affected by the non-availability of water.

Water is considered as '**elixir of life**'. Water constitutes almost 90 to 95% of most plant cells and tissues. Water helps the cells to maintain turgidity and shape. It shows following properties due to which it has great biological importance.

6.1 Properties of water:

It is in the liquid form at room temperature and is the best solvent for most of the solutes. It is inert inorganic compound with neutral pH when in pure form. Due to this, water is best transporting medium for dissolved minerals and food molecules. It is best aqueous medium for all biochemical reactions occurring in the cells. It is an essential raw material for photosynthesis. Water has **high specific heat**, **high heat of vaporization** and **high heat of fusion**. Due to this, it acts as **thermal buffer**. These various properties are due to hydrogen bonds between the water molecules.

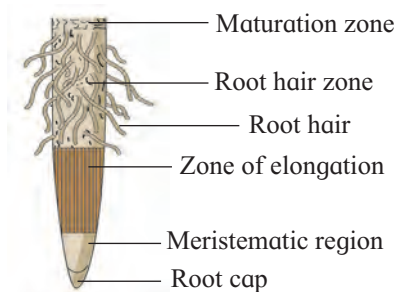
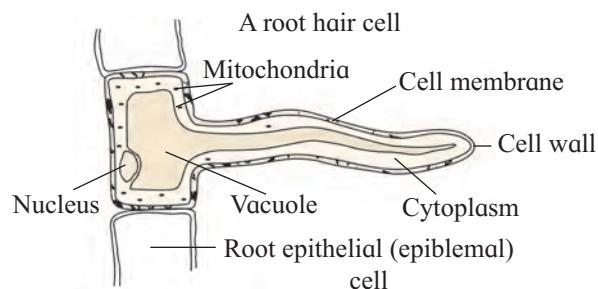
Curiosity Box :

1. What is hydrogen bond?
2. What are the meanings of specific heat, heat of vaporization and heat of fusion?
3. What are adhesive and cohesive forces?

Water molecules have good adhesive and cohesive forces of attraction. Due to high surface tension and high adhesive and cohesive force, it can easily rise in the capillaries. It is therefore, a significant molecule that connects physical world with biological processes.

6.2 Water absorbing organ:**Root :**

Root is the main organ of water and mineral absorption. In terrestrial plants, plants absorb water in the form of liquid from the soil however, epiphytic plants like orchids absorb water vapours from air with the help of epiphytic roots having special tissue called **velamen**. Typical root is divisible into four different regions. In the zone of absorption, epidermal cells (**epiblema cells**) form unicellular hair like extensions called **root hairs**.

**Fig. 6.1 a. : Root tip showing root hair zone****Fig. 6.1 b. : Structure of root hair**

Structure of root hair :

Root hair is cytoplasmic extension (prolongation) of epiblema cell. Each root hair may be approximately 1 to 10mm long and tube like structure. It is colourless, unbranched, short-lived (**ephemeral**) and very delicate. It has a large central vacuole surrounded by thin film of cytoplasm, plasma membrane and thin cell wall, which is two layered. Outer layer is composed of pectin and inner layer is made up of cellulose. Cell wall is freely permeable but plasma membrane is selectively permeable.

6.3 Water available to roots for absorption:

Plants absorb water from the **rhizosphere** (the microenvironment surrounding the root). Water present in the soil occurs as gravitational (free) water, hygroscopic water, combined water and capillary water. Water percolates deep, due to the gravity, in the soil, is called '**gravitational water**'. This is not available to plants for absorption. Fine soil particles imbibe/ adsorb water and hold it. This is called '**hygroscopic water**'. Roots cannot absorb it. Water present in the form of hydrated oxides of silicon, aluminum, etc., is called '**combined water**'. It is also not available to plants for absorption. Some amount of water is held in pores present between the neighbouring soil particles, due to capillarity. This is called **capillary water** that is available for absorption.

6.4 Absorption of water by roots from soil:

Root hair absorbs water by employing three physical processes that occur sequentially- viz. **imbibition, diffusion and osmosis**.



Activity :

Try this at your home.

- Take 10 ml of pure water in a suitable glass vessel and put 2 - 3 raisins in it. Observe the changes in raisins since the time you put them in water till they become fully swollen i.e. turgid. Why did raisins become turgid?
- Take 10 ml of pure water and add 5 gms of either sugar or salt to it. Let it dissolve and then put the same turgid raisins in it and observe the changes in raisins. What changes did occur in raisins and why? Discuss your observations with your teachers.

a. Imbibition:

Imbibition is swelling up of hydrophillic colloids due to adsorption of water. Substance that adsorbs water / liquid, is called as **imbibant** and water/ liquid, that gets imbibed is called as **imbibate**. The root hair cell wall is made up of pectic compounds and cellulose which are hydrophillic colloids. During Imbibition, water molecules get tightly adsorbed without the formation of solution. Imbibition continues till the equilibrium is reached. In other words, water moves along the concentration gradient.

Imbibition is significant in soaking of seeds, swelling up of dried raisins, kneading of flour etc.



Use your brain power

Why do the wooden doors become very hard to close and open in rainy season?

b. Diffusion: Diffusion means to disperse. Diffusion can be defined as the movement of ions/ atoms/ molecules of a substance from the region of their higher concentration to the region of their lower concentration. The movement is due to the kinetic energy of the molecules. Diffusion continues till an equilibrium is reached. Thus, water passes into the cell by diffusion through a freely permeable cell wall. Water is now at the interface of cell wall and plasma membrane.

Diffusion results in the diffusion pressure (D. P.) which is directly proportional to the number of diffusing particles. Diffusion pressure of pure solvent (pure water) is always

more than the diffusion pressure of solvent in a solution. The difference in the diffusion pressures of pure solvent and the solvent in a solution is called **Diffusion Pressure Deficit (DPD)** or **Suction Pressure (SP)**. The term was coined by B.S. Meyer (1938). Now a days, term **water potential** is used for DPD. In colloquial language, the term DPD is actually the thirst of a cell with which it absorbs water from the surroundings. Water around cell wall has more diffusion pressure than cell sap. Due to this, water moves in the cell by diffusion. Diffusion is significant in plants in the absorption of water, minerals, conduction of water against the gravity, exchange of gases and transport and distribution of food.

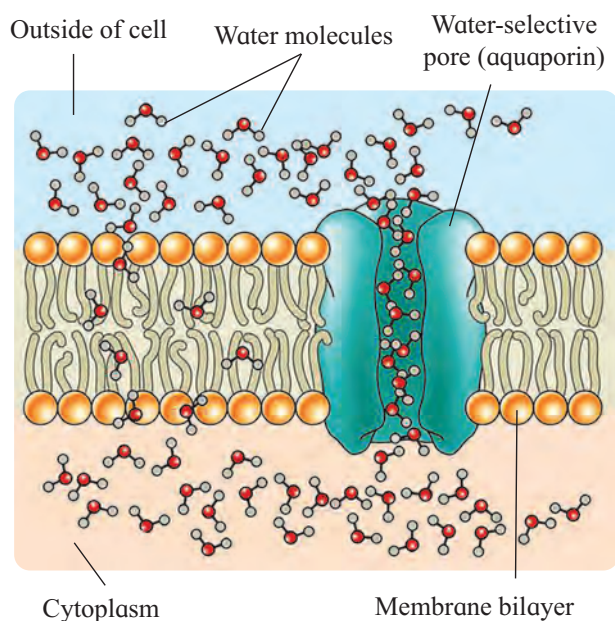


Fig. 6.2 : Diffusion of water into plant cell across the plasma membrane



Use your brain power

1. When you burn an incense stick in one corner of room, its fragrance spreads all over the room in a short time. How does it happen?
2. How does the water come out through the surface of porous earthen pot?

c. Osmosis : It is a process by which water enters into the cytoplasm of the root hair cell. Osmosis is a special type of diffusion of solvent through a semipermeable membrane. The cytoplasm of root hair cell contains minerals, sugars, etc. In other words, solution inside the cell is more concentrated (stronger) than outside the cell (weaker). Therefore, solvent from weaker solution enters into cytoplasm (i.e. to stronger solution) of cell through a semipermeable plasma membrane. This migration of solvent is called **Osmosis**.

Thus, water at the interface of cell wall and plasma membrane, enters into the cytoplasm of the root hair cell due to osmosis.

With respect to the concentration and osmotic migration, three types of solutions are recognized viz,

- i. Hypotonic** (weak solution or strong solvent) having low osmotic concentration.
- ii. Hypertonic** (strong solution or weak solvent) having high osmotic concentration.
- iii. Isotonic** having such a concentration of solution where there is neither gain nor loss of water in an osmotic system. In other words, concentration outside and inside the cell is same.

Osmosis is of two types viz, Exosmosis and Endosmosis.

Exosmosis : It is the diffusion of solvent from the cell outside. It causes flaccidity of cell.

Endosmosis : It is the diffusion of the solvent into the cell. It causes turgidity of cell i.e. cytoplasm becomes turgid. Turgidity increases the **turgor pressure (T. P.)** of the cell. T. P. is the pressure exerted by turgid cell sap on to the cell membrane and cell wall. In a fully turgid cell, DPD is zero. Cell wall being thick and rigid, exerts a counter pressure on the cell sap. This is called **Wall pressure (W. P.)**. In a fully turgid cell, T. P. = W. P. but operating in opposite direction.

Osmotic pressure (O. P.) : The pressure exerted due to osmosis is osmotic pressure.

Osmotic pressure is a pressure of the solution, which is required in opposite direction, so as to stop the entry of solvent molecules into the cell. More simply, osmotic pressure of a solution is equivalent to the pressure which must be exerted upon it to prevent flow of solvent across a semipermeable membrane.

Therefore, $D.P.D. = O.P. - T.P.$
 $(\because T.P. = W.P.)$
 $\therefore D.P.D. = O.P. - W.P.$

In a flaccid cell, T. P. is zero $\therefore DPD = OP$

In a turgid cell, DPD is zero $\therefore TP = OP$



Do you know ?

Importance of T. P. : It keeps cells and organelles stretched; provides support to the non-woody tissues; essential for cell enlargement during growth; maintains shape of cell and facilitates opening and closing of stoma.

Importance of Osmosis : It is responsible for absorption of water into root; maintains turgidity of cell; facilitates cell to cell movement of water; offers resistance to drought, frost, etc; also helps in the drooping of leaflets and leaves in vicinity of “touch me not” plant.

Facilitated diffusion : The passive absorption of solutes when mediated by a carrier, is called Facilitated diffusion. Particles that are lipid soluble can easily diffuse through lipoproteinous cell membrane. The diffusion of hydrophilic solutes has to be facilitated because their diffusion across the membrane is difficult. Membrane proteins provide such sites for facilitated diffusion. These proteins are aquaporins and ion- channels. These proteins help move substances across membranes without the expenditure of energy. Concentration gradient must be present for the molecules to be diffused through facilitated diffusion.

6.5 Water Potential (ψ):

According to the principle of thermodynamics, every component of a system is having a definite amount of free energy which is used to do work. Osmotic movement of water is on the basis of free energy. Free energy per molecule in a chemical system, is called its **chemical potential**.

Chemical potential of water is called **water potential**. It is represented by Greek letter *psi* (ψ). Water potential of protoplasm is equal but opposite in sign to DPD. It has negative value. The unit of measurement is in bars/ pascals/ atmospheres.

Water potential of pure water is always zero. Addition of any solute in it, decreases its *psi* (ψ) value. Therefore, it has negative value.

D. P. D. is now termed as water potential.
 O. P. is now termed as osmotic potential.
 T. P. is now termed as pressure potential. It has always positive value.

Water always flows from less negative potential to more negative water potential (i.e. from high water potential area to low water potential area). Difference between water potential of the adjacent cells decides movement of water through plasmodesmata across the cells.

Factors affecting water absorption:

- Presence of capillary water is essential.
- Rate of water absorption is maximum at soil temperature between 20^o to 30^oC.
- High concentration of solutes in soil water reduces the rate of absorption of water.
- Poorly aerated soil shows poor absorption rate.
- Increased transpiration accelerates the rate of absorption of water in the irrigated soil.

6.6 Plasmolysis:

Exo-osmosis in a living cell when placed in hypertonic solution, is called **plasmolysis**. During plasmolysis, protoplast of cell shrinks and recedes from cell wall. Thus, cell becomes

flaccid. In a plasmolysed cell, a gap is developed between cell wall and the protoplast. This gap is filled up by outer solution.

In a plasmolysed cell, T. P. is always zero. When such cell is placed in hypotonic solution, endo-osmosis occur, making cell turgid. This is called **deplasmolysis**. In a fully turgid cell T. P. = O. P. hence, DPD is always zero.

6.7 Path of water across the root (i.e. from epiblema upto xylem in the stelar region) :

Water is absorbed by root hair cell through imbibition → diffusion → osmosis, sequentially. Consequently the cell becomes turgid. Its turgor pressure increases, but its DPD value decreases. However, the immediately adjacent cortical cell inner to it, has more DPD value, because its O. P. is more. Therefore, cortical cell will suck water from the turgid root hair cell. It then becomes turgid. The flaccid root hair cell now absorbs water from soil.

Water from the turgid cortical cell is sucked by inner cortical cell and the process goes on. Thus, a gradient of suction pressure (DPD) is developed from cells of epiblema to the cortex of the root. Consequently water moves rapidly across the root through loosely arranged living cells of cortex, followed by passage cells of endodermis (in monocot roots) and finally into the cell of pericycle. Protoxylem is in the close proximity with pericycle.

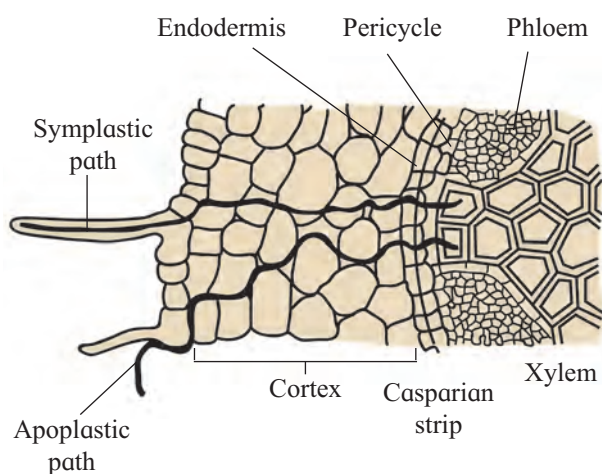
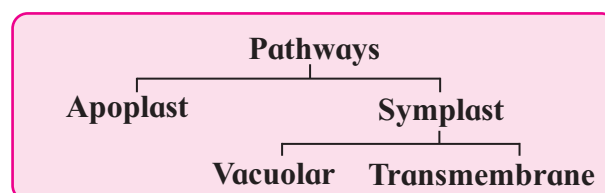


Fig. 6.3 : Pathways for water uptake by the root

Absorption of water being a continuous process, a sort of hydrostatic pressure is developed in living cells of root. This is called **root pressure**. It is due to root pressure, water from pericycle is not only forced into the xylem, but also conducted upwards against the gravity.

Pathway of water across the root essentially occurs in two ways viz, **apoplast** and **symplast**.



When some amount of water passes across the root through the cell wall and the intercellular spaces of cortical cells of root, it is then called **apoplast pathway**. This pathway occurs up to endodermis.

The apoplastic (non-living) pathway provides a route toward the vascular stele through free spaces and cell walls of the epidermis and cortex. An additional apoplastic route that allows direct access to the xylem and phloem is along the margins of secondary roots. Secondary roots develop from the pericycle, a cell layer just inside the endodermis. The endodermis is characterized by the Casparian strip, a suberized layer that forces all to move in the symplast in order to enter the vascular system. Since secondary roots grow through the endodermis, a direct pathway to the xylem and phloem is available that bypasses the Casparian strip and allows to enter the vascular system without moving into the symplast (living tissue).

When water passes across from one living cell to other living cell through plasmodesmata, then it is called **symplast pathway**. It is also called transmembrane pathway.

6.8 Mechanism of absorption of water :

Mainly, there are two ways/ modes of absorption of water viz, passive absorption and active absorption.

a. Passive absorption :

It is the main way of absorbing water **through the roots and not by the roots** from soil into the plant. The driving force is transpiration pull and it thus proceeds through DPD gradient. There is no expenditure of energy (ATP) as water moves in accordance to the concentration gradient. Hence, it is passive absorption. About 98% of the total water absorbed in plants, occur passively. Passive absorption occurs during day time when transpiration is in progress. It stops at night when transpiration stops.

Rapid transpiration creates a tension in the xylem vessel due to negative water potential. This tension is transmitted to xylem in the roots. Consequently water is pulled upwards passively.

During passive absorption, no ATP is utilized. Obviously, the rate of respiration is not affected. In plants, water is mainly absorbed passively.

b. Active absorption :

Here, water is absorbed due to activity of roots. Root cells play active role in the absorption of water. The driving force is the root pressure developed, in the living cells of root. Active absorption occurs usually at night when transpiration stops due to closure of stomata. As water absorption is against the DPD gradient, there is expenditure of ATP (energy) generated through the respiratory activity of cells.

Active absorption may be of two kinds viz, osmotic and non-osmotic :

1. Osmotic absorption : Atkins and Priestly (1922) proposed that water is absorbed from soil into xylem of the root according to the osmotic gradient. To create osmotic conditions, there is an expenditure of energy. But such absorption does not directly require an expenditure of energy.

A gradient of DPD develops from cell of epiblema to pericycle due to activity of living cells of root. As the process is continuous,

a hydrostatic pressure, called root pressure, is developed in root cells. This root pressure forces water from pericycle to xylem and then upwards to the stem.

2. Non-osmotic absorption : Kramer and Thimann (1959) proposed this theory. Sometimes, water is absorbed from soil against the concentration gradient. Such absorption requires an expenditure of energy released during respiration, directly. Poor supply of oxygen retards water absorption. Moreover low temperature retards water uptake because of decrease in the rate of respiration. Use of metabolic inhibitors also retards the rate of respiration and thus the water uptake.

6.9 Translocation of water:

The transport of water with dissolved minerals from root to other aerial parts like stem and leaves, against the gravity, is called translocation or **ascent of sap**.

Translocation of water occurs through the lumen of conducting elements of xylem-tracheids and vessels, in all vascular plants. Ringing experiment has proved that xylem is the path of ascent of sap.

Several mechanisms/ theories have been put forth to explain the mechanism of translocation of water. The theories include-vital force theory, relay pump theory, physical force theory, root pressure theory, etc. We shall consider following three theories :

a. Root Pressure Theory (Vital Theory) :

According to this theory, the activity of living cells of root is responsible for translocation of water. J. Priestly proposed this theory. When a stem of potted plant is cut few inches above the soil by a sharp knife, xylem sap is seen flowing out/ oozing out through the cut end. This exudation at the cut end of stem is a good proof for the existence of root pressure. As water absorption by roots is constant and continuous process, a hydrostatic pressure is developed in the living cells of cortex of root.

This is termed as **root pressure** by S. Hales. It is due to root pressure water along with dissolved minerals is not only forced into xylem but it is also conducted upwards against the gravity.

Root pressure seems to be largely an osmotic phenomenon and its development is an active process. The value of root pressure is +1 to +2 bars which is enough to pump water to a height of 10 to 20 meters. The factors like oxygen, moisture, temperature of soil, salt contents, etc. influence the root pressure.

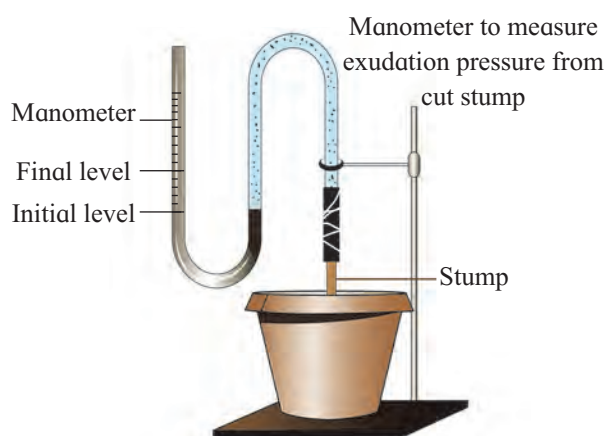


Fig. 6.4 : Experiment to demonstrate root pressure

Objections/ limitations of root pressure theory:

Although, ascent of sap takes place due to root pressure, there are certain objections raised, such as -

- i. It is not applicable to plants taller than 20 meters.
- ii. Ascent of sap can also occur even in the absence of root system.
- iii. Root pressure value is almost nearly zero in taller gymnosperm trees.
- iv. In actively transpiring plants, no root pressure is developed.
- v. Xylem sap under normal condition is under tension i.e. it shows negative hydrostatic pressure or high osmotic pressure.

To sum up therefore, root pressure is not the sole mechanism explaining the ascent of sap in all plants of varying heights..

b. Capillarity theory (physical force theory):

According to this theory, physical forces and dead cells are responsible for ascent of sap.

This theory was put forth by Bohem in (1863). Wick dipped in an oil lamp, shows capillarity due to which oil is raised upwards. The conduction of water in a straw dipped in water, is raised to a certain height because of capillarity. The height to which water is raised depends on the diameter of the straw.

Capillarity is because of surface tension, and forces of cohesion (attraction between like molecules) and adhesion (attraction between unlike molecules). Xylem vessel/ tracheid with its lumen is comparable with straw. Water column exist because of combined cohesive and adhesive forces of water and xylem wall, due to capillarity. It is because of capillarity water is raised or conducted upwards against the gravity, to few centimeters only.

Objections/ Limitations of capillarity theory:

Few important objections are :

- i. Capillary tube (xylem) must be continuously and completely hollow from one end to the other end but tracheids in the xylem show closed end-walls.
- ii. The lower end of capillary tube i.e. xylem must be in direct contact with soil water. However, there exists a barrier of root cortex between xylem and soil water.
- iii. Narrower the capillary tube, greater is the height to which water column is raised. Thus, taller trees should show xylem vessels with very narrow bore (diameter). However, in nature the tall trees show xylem vessels having wider bore.

Hence, to sum up capillarity can not be the sole mechanism to explain ascent of sap in all the plants of varying heights.

c. Cohesion- tension theory (Transpiration pull theory) :

This is presently widely accepted theory explaining ascent of sap in plants. It was put

forth by Dixon and Jolly (1894). This theory is based on two principles.

Cohesion and adhesion, and transpiration pull :

A strong force of attraction between water molecules, is called **cohesive force**. While a strong force of attraction between water molecules and lignified wall of lumen of xylem vessel, is called **adhesive force**.

Due to combined cohesive and adhesive forces a continuous water column is developed (formed) in the xylem right from root upto the tip of the topmost leaf in the plant.

Transpiration pull : The transpiration pull developed in the leaf vessel is transmitted down to root and thus accounts for the ascent of sap.

Excess water is lost in the form of vapour, mainly through the stomata found on leaf. This water loss increases D. P. D. of mesophyll cells. These cells withdraw water ultimately from xylem in the leaf. In other words, due to continuous transpiration, a gradient of suction pressure (i.e. D. P. D.) is developed right from guard cells up to the xylem in the leaf. This will create a tension (called negative pull or transpiration pull) in the xylem. Consequently, water column is pulled out of xylem. Thus, water is pulled upwards passively against the gravity leading to the ascent of sap.

Objections/ Limitations of transpiration pull theory:

- i. For transpiration pull to operate, water column should be unbroken and continuous. However, due to temperature fluctuations during day and night, gas bubbles may enter in water column breaking the continuity.
- ii. This mechanism assumes that tracheids are more efficient than the vessels, as their end walls support water column.

However, vessels are more evolved than tracheids and are more efficient.

iii. If plant leaves are smeared with vaseline in order to stop transpiration, even then ascent of sap occurs.

iv. Ascent of sap also occurs in deciduous plants that have shed all of their leaves.

These observations point to the fact that besides physical forces, activity of living cells seems to be necessary for lifting the water column up.

6.10 Transport of mineral ions :

Soil serves as main source for minerals. Minerals constitute most commonly occurring solid, crystalline inorganic materials obtained from earth's crust. Minerals play an important role in the day to day life of plant. Minerals are absorbed by plants in the ionic (dissolved) form, mainly through roots and then transported.



Do you know ?

- Minerals that play important role in the day to day life, are called essential elements. About 36 to 40 elements are incorporated in the plant's life.
- Some minerals like C, H, O, P, N, S, Mg required in large quantity, are called **macro elements**. While minerals like Cu, Co, Mn, B, Zn required in small quantity, are called **micro elements**.

The analysis of plant ash demonstrates that minerals are absorbed by plants from soil and surroundings. Absorption of minerals is independent of that of water.

Absorbed mineral ions are pulled in upward direction along with xylem sap because of transpiration pull. This could be understood when the ascending sap is analysed. Mineral ions are needed in the areas of the plant viz. apical, lateral, young leaves, developing flowers, fruits, seeds and storage organs. Hence, from the source (root), these are pulled and transported ascendingly through the sap and gets unloaded by fine veins through the

process of diffusion in the vicinity of cells. Cells uptake them actively.

Soil would not be the only source for mineral uptake. Mineral ions can be remobilized within the parts of the plant. Older parts (like leaves in deciduous plants) export their ions to younger leaves before the fall. Most readily mobilized ions are like phosphorus, sulphur, nitrogen and potassium but the ions from structural components like calcium is not remobilized.



Internet my friend

1. The general roles of minerals in the life of plants.
2. The role of essential minerals in the day to day life of plants.

Analysis of xylem exudate also shows that some nitrogen travels as inorganic ions whereas much of it is carried in the organic form like amino acids and related compounds. Small amount of inorganic molecules of phosphorus and sulphur are also carried. It was a belief earlier that xylem transports inorganic and phloem transports organic molecules. However, it is not correct because some exchange of materials also occurs between xylem and phloem.



Do you know ?

- Different modes of passive absorption and active absorption of minerals in plants.
- Carrier concept of active absorption.

6.11 Transport of food :

All the plant parts require continuous supply of food (photosynthate) for nutrition and development. In higher plants, there is a great differentiation and division of labour. Chloroplasts are confined to green cells of leaves where food is synthesized. The non-green parts like root and stem must receive food from leaves. The part where food is synthesized is called **source** and while part where it is utilized, is called **sink**. Food has to travel from source to

sink. This movement of food from one part to the other part, is called translocation of food.

Path of translocation : Food is to be translocated to longer distances in higher plants. Hence plants must have adequate channels for the transport of food. Sieve tubes and vessels are structurally ideally suited for longitudinal (vertical) translocation. The ringing experiment, structure and distribution of phloem, chemical analysis of phloem sap and use of isotope ^{14}C , clearly point out that the phloem tissue is primarily responsible for flow of food in longitudinal downward direction. The horizontal (lateral) translocation occurs from phloem to pith or phloem to cortex via medullary rays in the stem.

Food is always translocated in the form of **sucrose** (soluble form) and always along the concentration gradient from source to sink. The transport of food occurs in **vertical** and **lateral** direction.

Vertical translocation : In vertical (longitudinal) transport, food is translocated in downward direction from leaves (source) to stem and root (sink). It also occurs in upward direction during germination of seed, bulbils, corm, etc. Upward translocation also occurs from leaves to growing point of stem, to developing flowers and fruits situated near the ends of the branches of stem.

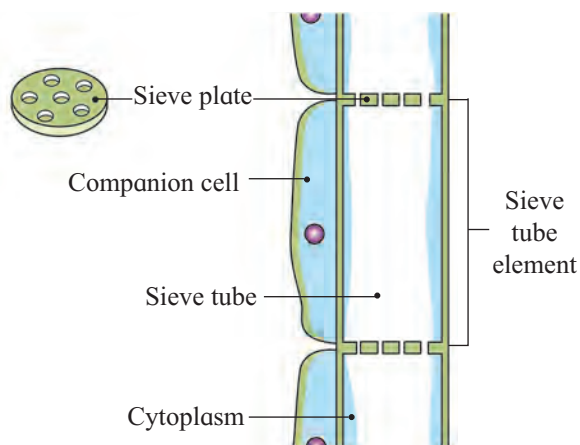


Fig. 6.5 : L. S. of sieve tube

Lateral translocation : It occurs horizontally/ laterally across the root and stem. When food is translocated from phloem to pith, it is called **radial** translocation and from phloem to cortex, it is called **tangential** translocation.

The transport of food through phloem is **bidirectional**. Phloem sap contains mainly water and food in the form of sucrose. But sugars, amino acids and hormones are also transported through phloem.

Mechanism of sugar transport through phloem :

Several mechanisms/ theories like diffusion, activated diffusion, protoplasmic streaming, electro-osmosis, pressure-flow, etc. are put forth. The most convincing theory is Munch's pressure flow theory or mass flow hypothesis.

Ernst Munch proposed that photosynthetic cell synthesizes glucose. Hence, its osmotic concentration increases. Due to endo-osmosis water from surrounding cells and xylem, is absorbed. The cell becomes turgid. Due to increase in turgor pressure, sugar from photosynthetic cell is forced ultimately into the sieve tube of the vein. This is called **loading of Vein**.

At the sink end, root cell utilizes sugar and also polymerizes excess sugar into the starch. Its osmotic concentration is lowered. Exo-osmosis occurs. Water in the root cell is lost to surrounding cells, thereby decreasing the turgidity of cell. Turgor pressure is lowered. Hence, a turgor pressure gradient is developed from sieve tube in the leaf to the root cell. Consequently, food is translocated along the concentration gradient, passively. This is **Vein unloading**. At the sink end sugar is used and excess water exudes into the xylem.

Main objection to this theory is that this mechanism does not explain bidirectional transport of food. More over, according to Munch, pressure flow is purely a physical process.

6.12 Transpiration :

Plants absorb water constantly and continuously. Hardly 5% of the total water absorbed by roots that is utilised for cell expansion and plant growth. Remaining 95% water becomes surplus which is then lost into the atmosphere, through its aerial parts. Hardly 1% of surplus water is lost in the form of liquid and 99% of surplus water, is lost in the form of vapour. The loss of water in the form of liquid is called **guttation**. It occurs through special structures called water stomata or **hydathodes**. The loss of water in the form of vapour is called **transpiration** that occurs through leaves, stem, flowers and fruits. Most of the transpiration occurs through the leaves (called foliar transpiration). The actual water loss during transpiration occurs through three main sites - **cuticle, stomata and lenticels**. Accordingly, three types of transpiration are recognized viz. cuticular, stomatal and lenticular.

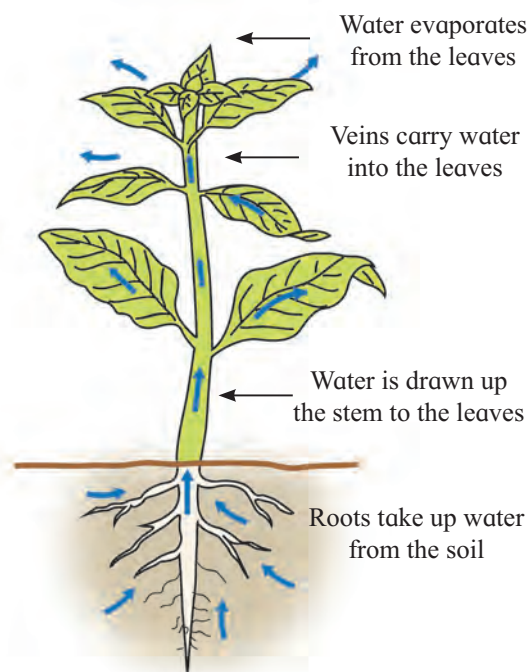


Fig. 6.6 : Transpiration

i. Cuticular transpiration:

Cuticle is a layer of waxy substance- cutin, present on outer surface of epidermal cells of leaves and stem. Cuticular transpiration occurs by simple diffusion and contributes 8-10% of the total transpiration. Cuticular transpiration

occurs throughout the day and its rate is inversely proportional to thickness of cuticle.

ii. Lenticular transpiration:

Lenticels are small raised structures composed of loosely arranged complementary cells. Each lenticel is a porous tissue consisting of cells with large intercellular spaces in the periderm of the secondarily thickened organs and the bark of woody stems and roots of dicotyledonous flowering plants. Lenticels are present in bark of old stem and pericarp of woody fruits but are absent in leaves. Lenticular transpiration contributes only about 0.1-1.0% of total transpiration. Rate of

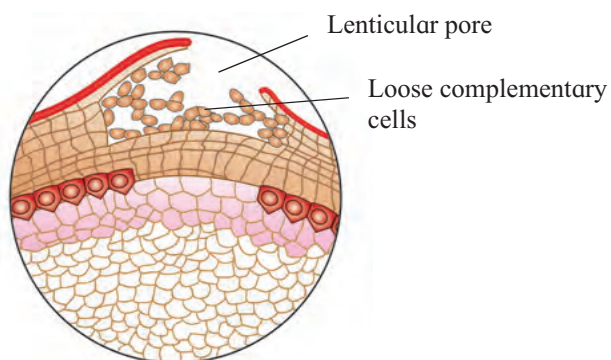


Fig. 6.7 : Structure of lenticel

lenticular transpiration is very slow. It also occurs throughout the day.

iii. Stomatal transpiration:

Stomata are minute apertures formed of two guard cells and accessory cells. They are located in the epidermis of young stem and leaves. Leaves generally show more number of stomata on the lower surface. Depending upon distribution of stomata on leaves, leaves are categorized into three types namely **epistomatic**- on upper epidermis (Hydrophytes- e.g. Lotus), **hypostomatic**- on lower epidermis (Xerophytes- e.g. *Nerium*) and **amphistomatic**- on both surfaces (Mesophytes- e.g. Grass). Stomatal transpiration occurs only during daytime. (Exception: Desert plants).

90 to 93% of total transpiration occurs through stomata and that too during day time only.

Do you know ?

- The number of stomata per unit area of leaf, is called **stomatal frequency**.
- The correlation between the number of stomata and number of epidermal cells per unit area, is called **stomatal index (I)**.

6.13 Structure of stomatal apparatus :

Typical stomatal apparatus consists of two **guard cells**, **stoma** and **accessory cells**.

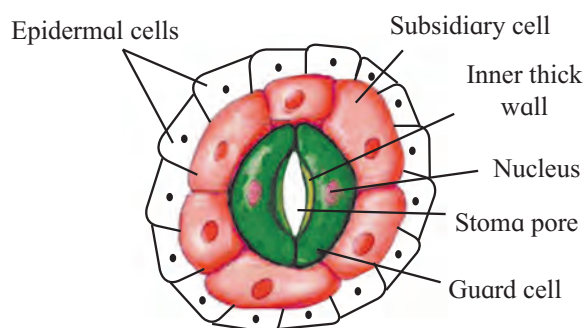


Fig. 6.8 (a) : Structure of guard cell

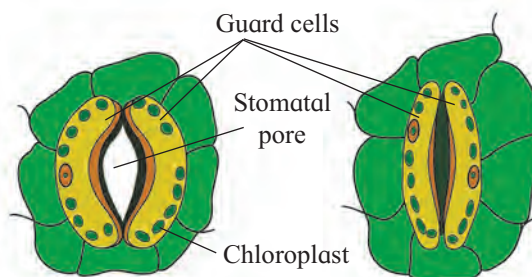
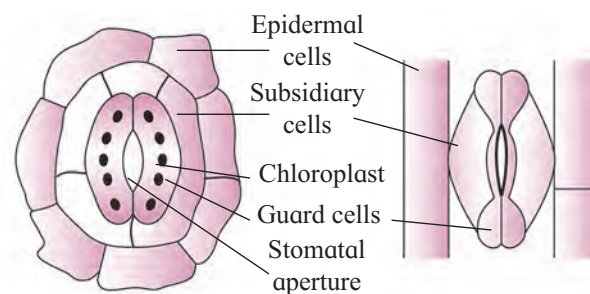


Fig. 6.8 (b) : Open and closed stoma

Stomata are minute, elliptical pores bounded by two kidney/ dumbbell shaped **guard cells**. Guard cell is a type of epidermal tissue which may be called as modified, epidermal parenchyma cell. They are kidney-shaped in dicotyledons and dumbbell-shaped in grasses (monocotyledons/monocots).



A. Kidney shaped B. Dumbbell shaped

Fig. 6.9 : Types of guard cells

In *Cyperus*, both kidney- and dumbbell-shaped guard cells are present.

Guard cells are living, nucleated cells with unevenly thick walls. Inner wall (wall facing stoma) of guard cells is thick and inelastic, and its lateral (outer) wall is thin and elastic. Guard cells contain few chloroplasts which are capable of poor photosynthesis. Guard cells have ability to change their size and form due to which stoma opens (widens) or closes (narrows).

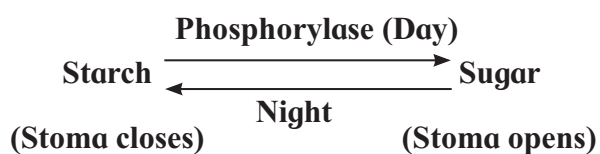
Stoma is an elliptical pore formed due to specific arrangement of guard cells. It is through the stoma, excess water is lost in the form of vapour.

Accessory cells : These are specialized epidermal cells surrounding the guard cells. Their number is variable and are the reservoirs of K^+ ions. These are also called **subsidiary cells**.

Opening and Closing of Stoma :

Opening and closing of stoma is controlled by turgor of guard cells. During day time, guard cells become turgid due to endosmosis. Thus turgor pressure is exerted on the thin walls of guard cells. Being elastic and thin, lateral walls are stretched out. Due to kidney or dumb-bell like shape, inner thick walls are pulled apart to open (widen) the stoma. During night time, guard cells become flaccid due to exosmosis. Flaccidity closes the stoma almost completely. Endosmosis and exosmosis occur due to diurnal changes in osmotic potential of guard cells. Different theories are proposed to explain diurnal changes in osmotic potential.

According to **starch-sugar inter-conversion theory** (Steward 1964), during day time, enzyme phosphorylase converts starch to sugar, thus increasing osmotic potential of guard cells closing entry of water thereby guard cells are stretched and stoma widens. The reverse reaction occurs at night bringing about the closure of stoma.



According to theory of proton transport (Levitt-1974), stomatal movement occurs due to transport of protons H^+ and K^+ ions. During daytime, starch is converted into malic acid. Malic acid dissociates to form Malate and protons. Protons are transported to subsidiary cells and K^+ ions are imported from them. Potassium malate is formed that increases osmolarity and causes endosmosis. Uptake of K^+ ions is always accompanied with Cl^- ions.

At night, uptake of K^+ and Cl^- ions is prevented by abscissic acid, changing the permeability of guard cells. Due to this guard cells become hypotonic and thereby become flaccid.

Significance of Transpiration :

Advantages:

- i. It removes excess of water.
- ii. It helps in the passive absorption of water and minerals from soil.
- iii. It helps in the ascent of sap.
- iv. As stomata are open, gaseous exchange required for photosynthesis and respiration, is facilitated.
- v. It maintains turgor (turgidity) of the cells.
- vi. Transpiration helps in reducing the temperature of leaf and in imparting cooling effect.

Disadvantages:

Excessive transpiration leads to wilting and injury in the plant. It may also lead to the death of the plant.

Transpiration is 'A necessary evil' :

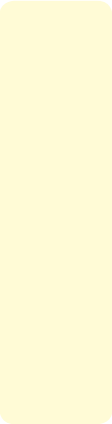
For stomatal transpiration to occur, stoma must remain open, during day time. When stomata are open then only the gaseous exchange needed for respiration and photosynthesis, will take place. If stomatal transpiration stops, it will directly affect productivity of plant through the loss of photosynthetic and respiratory activity. Hence for productivity, stomata must remain open. Consequently transpiration can not be avoided. **Hence, Curtis (1926) regarded transpiration as 'a necessary evil'.**



Activity :

Prepare stomatal frequency chart for any six angiospermic plants in your area.

Plant Name	Details



Exercise

Q. 1 Multiple Choice Questions

- In soil, water available for absorption by root is
 - gravitational water
 - capillary water
 - hygroscopic water
 - combined water
- The most widely accepted theory for ascent of sap is
 - capillarity theory
 - root pressure theory
 - diffusion
 - transpiration pull theory
- Water movement between the cells is due to
 - T. P.
 - W. P.
 - D.P.D.
 - incipient plasmolysis
- In guard cells, when sugar is converted into starch, the stomatal pore
 - closes almost completely
 - opens partially
 - opens fully
 - remains unchanged
- Surface tension is due to
 - diffusion
 - osmosis
 - gravitational force
 - cohesion
- Which of the following type of solution has lower level of solutes than the solution?
 - Isotonic
 - Hypotonic
 - Hypertonic
 - Anisotonic
- During rainy season wooden doors warp and become difficult to open or to close because of
 - plasmolysis
 - imbibition
 - osmosis
 - diffusion
- Water absorption takes place through
 - lateral roots
 - root cap
 - root hair
 - primary root
- Due to low atmospheric pressure the rate of transpiration will
 - increase
 - decrease rapidly
 - decrease slowly
 - remain unaffected
- Osmosis is a property of
 - solute
 - solvent
 - solution
 - membrane

Q. 2 Very short answer questions.

- What is osmotic pressure?
- Name the condition in which protoplast of the plant cell shrinks.
- What happens when a pressure greater than the atmospheric pressure is applied to pure water or a solution?
- Which type of solution will bring about deplasmolysis?
- Which type of plants have negative root pressure?
- In which conditions transpiration pull will be affected?
- Mention the shape of guard cells in *Cyperus*.
- Why do diurnal changes occur in osmotic potential of guard cells?
- What is symplast pathway?

Q. 3 Answer the following questions.

- Describe mechanism for absorption of water.
- Discuss theories of water translocation.
- What is transpiration? Describe mechanism of opening and closing of stomata.

4. What is transpiration? Explain role of transpiration.
5. What is significance of transpiration? Explain root pressure theory and its limitations.
6. Explain capillarity theory of water translocation.
7. Why is transpiration called 'a necessary evil'?
8. Explain movement of water in the root.
9. Define and or explain the terms: Osmosis, diffusion, plasmolysis, imbibition, guttation, transpiration, ascent of sap, active absorption, DPD, turgor pressure, water potential, wall pressure, root pressure.
10. Distinguish between a) Osmotic pressure and turgor pressure b) Diffusion and osmosis.

Q. 4 Long answer questions.

1. Describe structure of root hair.
2. Write on journey of water from soil to xylem in roots.
3. Explain cohesion theory for translocation of water.
4. Write on the mechanism of opening and closing of stoma.
5. Explain the active absorption of minerals.

Project :

1. Prepare powerpoint presentation for different types of transpiration.



Can you recall?

1. Do you think that the growth is property of living beings only?
2. Is there any difference between plant growth and animal growth?

7.1 Plant growth :

Growth is one of the characteristic features of living organisms. Growth as a phenomenon has two aspects viz. **quantitative** and **qualitative**. Quantitative aspect speaks for an increase in the length, breadth, size, volume, body mass or dry weight and number of cells. Growth as a quantitative change is a final end product of successful metabolism.

Qualitative aspect talks about the change in the nature of growth where **development** is an ordered change or progress while **differentiation** leads to higher and more complex state. Growth thus can be defined as permanent, irreversible increase in the bulk of an organism, accompanied by the change of form.

In multicellular (vascular) plants, growth is indeterminate and occurs throughout the life indefinitely. It is restricted to some specific regions called **meristems** which are the regions where new cells are constantly and continuously produced. Meristems are of three types based on location viz. Apical, Intercalary and Lateral.



Do you know ?

Root and stem show indefinite or indeterminate growth. However, in organs like leaves, flowers and fruits, growth is determinate. i.e. they grow up to certain genetically destined size. In unicellular plants, growth is uniform and determinate.



Do you know ?

Apical meristem : In vascular plants, growth is restricted to the apices of root and shoot. It is responsible for growth in length/ height and the differentiation or cell types. It contributes to the primary growth.

Intercalary meristem : It is located at the node or at the base of internode of stem. It is primarily responsible for increasing length of internodes and also for formation of leaf primordia and lateral buds.

Lateral meristem : It is located laterally along the axis of dicotyledons and gymnosperms. It is located as strip in the vascular bundles of stem of dicots. It is called vascular cambium. It is responsible for increase in the girth of the stem due to addition of secondary vascular tissues.

7.2 Phases of growth :

The cells in the meristem divide, enlarge and get differentiated. Corresponding to these three stages, there are three phases of growth :

A. Phase of cell division/ formation : Cells of meristem are thin walled, non-vacuolated having prominent nucleus and granular cytoplasm. Meristematic (particularly cambial) cell undergoes mitotic division to form two new cells. One cell remains meristematic and the other cell undergoes enlargement and differentiation. In this phase, rate of growth occurs at a slower pace (Lag phase).

B. Phase of cell enlargement/ elongation :

The newly formed cell becomes vacuolated, osmotically active and turgid due to absorption of water. The turgidity results in the enlargement of cell - both lengthwise and breadthwise. In this phase new wall materials and other materials are synthesized to cope up with the enlargement. The growth rate in this phase occurs at an accelerated pace (exponential or Log phase).

C. Phase of Cell maturation/ differentiation :

The enlarged cell now becomes specialized to perform specific function and attains maturity - both morphological and physiological. In this phase, rate of growth slows down and comes to a steady state (stationary phase).

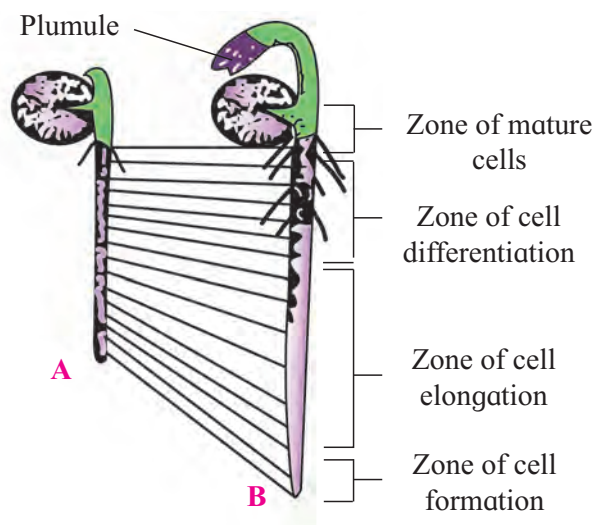


Fig. 7.1 Phases of growth in root - Position of radicle at the beginning (A) and at the end (B)

7.3 Conditions for Growth :

The different environmental and physiological conditions necessary for the growth include - Water, supply of nutrients,

temperature, oxygen, Carbon/ Nitrogen ratio, gravitational force, light and growth hormones. The chief conditions are explained below :

Water is the essential component of protoplasm and maintains turgidity of the cell. It acts as aqueous medium for biochemical reactions. Microelements and Macroelements are **nutrients** required for the proper growth of the plant. Optimum **Temperature** ranges between 25-35°C. **Oxygen** is essential for respiration and the release of energy. **Light** is very much essential for germination of seed and photosynthesis. **Gravitational force** decides the direction of growth of the shoot and root.

7.4 Growth Rate and types of growth :

Growth rate :

It is the increased growth per unit time. It is also called **efficiency index**. Rate of growth can be measured by an increase in the size and area of different plant organs like leaf, flower and fruits.

The ratio of change in the cell number (dn) over the time interval (dt) is called **Absolute growth rate** (AGR). Alternatively, it is the measurement and comparison of total growth per unit time.

Growth in plants can be measured in terms of...

1. Increase in the number of cells produced - e.g. single maize root apical mesistem can give rise to more than 17,500 new cells/Hour.
2. Increase in surface area of the leaf - e.g. growth of dorsiventral leaf.
3. Increase in length - e.g. growth of pollen tube.
4. Increase in volume of a fruit - e.g. In watermelon flower, the ovary after fertilization increases in its size/ volume by upto 3,50,000 times.
5. Increase in girth of shoot.
6. Increase in dry weight of organ.

Various methods for the measurement of linear growth of stem and radicle are as follows:

1. **Direct method** : It can easily be measured with the help of ordinary measuring scale. It is a simple method.
2. **Horizontal microscope** : It is used to measure growth in fields.
3. **Auxanometer** : This equipment is used for precise measurement of linear growth of shoot. There are two types of an auxanometers viz, a) Arc auxanometer b) Pfeffer's auxanometer
4. **Crescograph** : It records primary growth very accurately. It magnifies growth upto 10,000 times giving information of growth per second. It is developed by sir J. C. Bose.

$$AGR = \frac{dn}{dt}$$

The AGR, when divided by total number of cells present in the medium, gives **Relative growth ratio (RGR)**. Alternatively, RGR refers to the growth of a particular system per unit time, expressed on a common basis or it is the ratio of growth in the given time/ initial growth.

$$RGR = \frac{AGR}{n}$$

AGR and RGR are useful in describing the dynamics of cell growth in culture.

Types of growth :

There are two types of growth viz, arithmetic growth and geometric growth.

a. Arithmetic growth : Here, rate of the growth is constant and an increase in the growth occurs in arithmetic progression. i.e. 2, 4, 6, 8 cms etc. In this type of growth, the rate of growth is constant.

After mitosis one of the daughter cell continues to divide and the other cell takes part in the differentiation and maturation. e.g. elongation of root at a constant rate, best explains arithmetic growth. Linear curve is obtained when growth rate is plotted against the time. Arithmetic growth is expressed mathematically by an equation,

It is expressed as,

$$Lt = Lo + rt$$

Where

Lt = Length at time 't'

Lo = Length at time 'Zero'

r = Growth rate

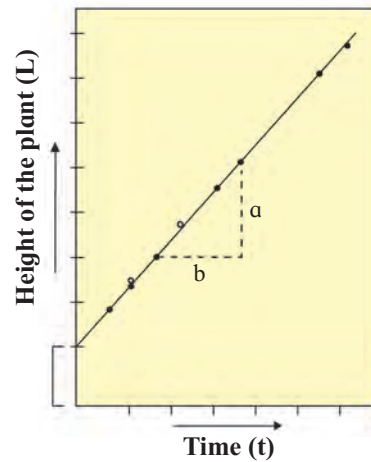
t = Time of growth

When graph of length (L) is plotted against the time (t), a linear curve is obtained as indicated in the diagram.

b. Geometric growth :

Cell divides mitotically into two. Here, both the daughter cells continue to divide and redivide repeatedly. Such growth is called geometric growth. Here, growth rate is slow

initially but later on there is a rapid growth at exponential rate. Geometric growth can be expressed mathematically by an equation.



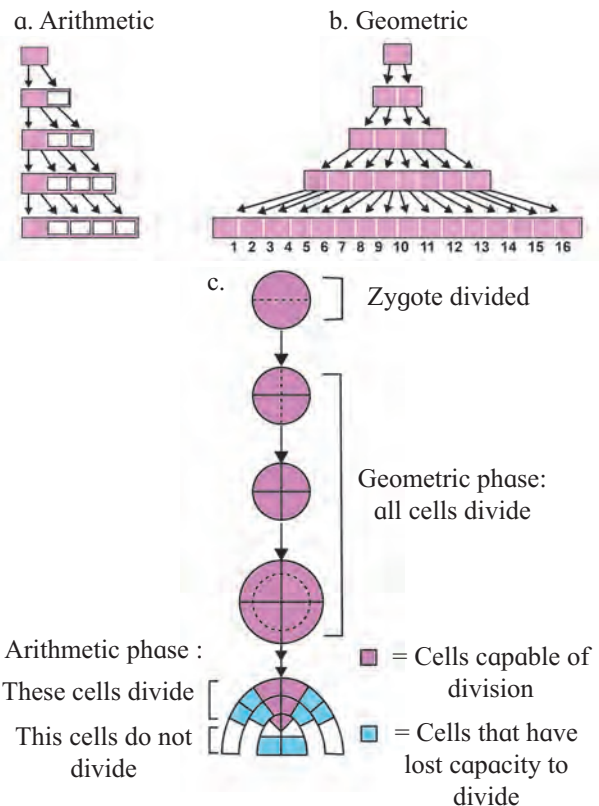
Graph 7.2 : Constant linear growth

$$W_1 = W_0 e^{rt}$$

W₁ = Final size , **W₀** = initial size

r = growth rate, **t** = time of growth

e = base of natural logarithm



Digrammatic representation of : **a.** Arithmetic, **b.** Geometric growth and **c.** Stages during embryo development showing geometric and arithmetic phases

Fig. 7.3 : Growth rate

We can also observe quantitative comparison between the growth of living system in two ways.

Measurement and comparisons of total growth per unit time is called the **Absolute growth rate(AGR)** whereas the growth of the given system per unit time expressed on a common basis per unit initial parameter is called the **Relative growth rate(RGR)**.

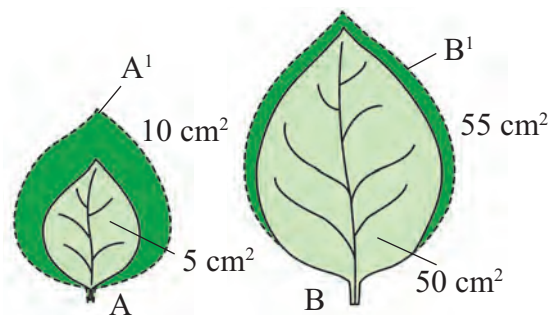


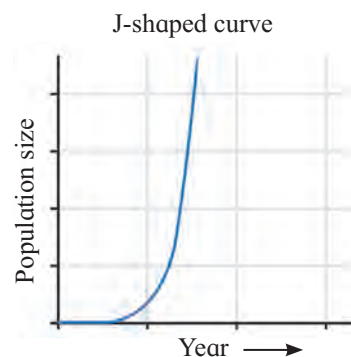
Fig. 7.4 : Growth rate comparison

In the above example, two leaves 'A' and 'B' are of different sizes but show same absolute increase in area in a given time. Both leaves grow and increase their area by 5cm² to produce 'A' and 'B' leaves. 'A' leaf of 5cm² in size grows 5cm²/ day then its RGR would be 100%. If the leaf is 50cm² in size and the growth rate/day is 5cm² then its RGR would become 10%.

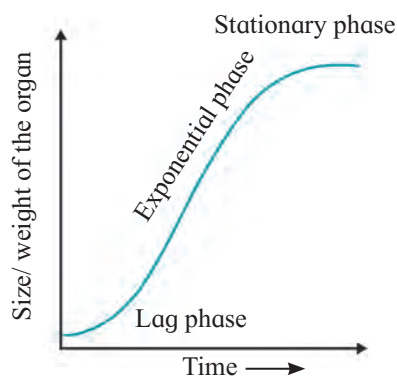
7.5 Growth curve :

It is a graphic representation of the total growth against time. There are three types of curves viz, Linear, Exponential and Sigmoid. Arithmetic growth curve is linear while Geometric growth curve is exponential.

Corresponding to three distinct phases of growth, growth rates differ. In **Lag phase**, growth rate is slow. In **Exponential (Log) phase**, growth rate is faster and reaches its maximum. In **Stationary phase**, growth rate gradually slows down. When a graph of rate of growth against time is plotted for three phases of growth, a 's' shaped i.e. sigmoid curve is obtained.



Graph 7.5 : Exponential Growth curve



Graph 7.6 : Sigmoid Growth curve

The total time (period) required for all phases to occur, is called **Grand Period of Growth**.

7.6 Differentiation, De-Differentiation, Re-Differentiation :

a. Differentiation :

It is maturation of cells derived from apical meristem of root and shoot. Permanent change in structure and function of cells leading to maturation, is called **differentiation**. During cell differentiation, cell undergoes few to major anatomical and physiological changes e.g. Parenchyma in hydrophytes develops large schizogenous interspaces for mechanical support, buoyancy and aeration. The maturation is at the cost of capacity to divide and redivide.

b. Dedifferentiation :

The living differentiated cell which has lost the capacity to divide, may regain the same as per the need and divide. Thus, permanent (mature) cell undergoes dedifferentiation and becomes meristematic e.g. **interfascicular**

cambium and **cork cambium** are formed from parenchyma cells between vascular bundles and inner most layer of cortex, respectively.

c. Redifferentiation :

The cells produced by dedifferentiation once again lose the capacity to divide and mature to perform specific function. This is called **redifferentiation** e.g. secondary xylem and secondary phloem are formed from dedifferentiated cambium present in the vascular bundle.

7.7 Development :

It refers to the ordered or progressive changes in shape, form and degree of complexity. It includes all the changes occurring in sequence from the germination of seed upto the senescence or death during life cycle of plants. Thus development includes growth, morphogenesis, maturation and senescence.



Fig. 7.7 Flow chart of development

7.8 Plasticity :

It is the capacity of being moulded, formed or modeled. It is the ability of plants to form different kinds of structures (i.e. to change) in response to different environmental (external) or internal stimuli, in various phases of life.

In many plants, juvenile stage and mature

stage show different forms of leaves in the same plant e.g. heterophylly in cotton, coriander, larkspur (*Delphinium*). The environmental heterophylly is shown by *Ranunculus flabellasis* (butter cup). The intrinsic plasticity is found in coriander and cotton. Heterophylly is exhibited in the same plant in different growth phases or under different environmental conditions.

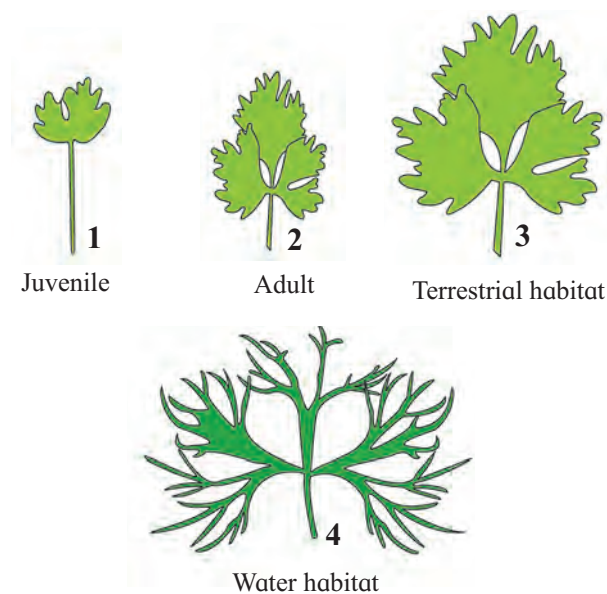


Fig. 7.8 Heterophylly



Can you tell?

1. Is there any relation between phases of growth and regions of growth curve?
2. Which plant organ does show both arithmetic and geometric growth?
3. Differences between arithmetic and geometric growth.

7.9 Growth Hormones :

The term 'hormone' was coined first by Starling (1906) in animal physiology. The internal factors that influence growth are called **growth hormones** or **growth regulators** as they inhibit, promote or modify the growth. **Growth promoters are auxins, gibberellins (GA) and cytokinins (CK). Growth inhibitors in plants are ethylene and abscisic acid (ABA).** All phytohormones are growth regulators.



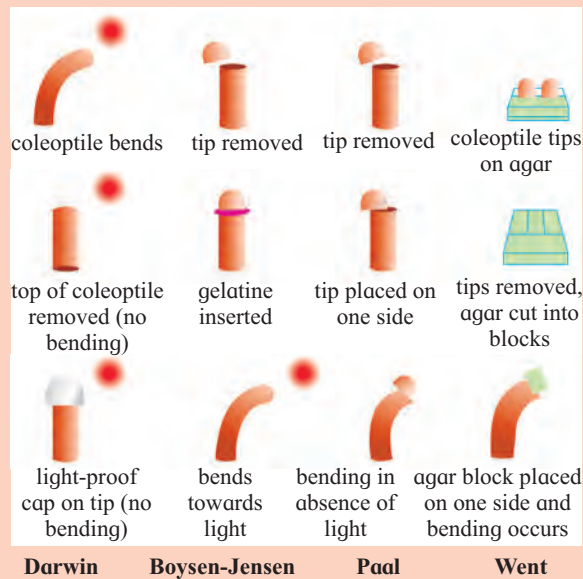
Know the Scientist :

The auxin is the first hormone to be discovered in plants. Discovery of auxins dates back to 19th century when **Charles Darwin** (1886) was studying tropism in plants. He exposed canary grass coleoptile to unilateral light. He concluded that a growth stimulus is developed in the coleoptile tip and transmitted downwards to the growth zone. This has caused bending of the tip towards light.

The Danish plant physiologist Boysen - Jensen (1910) cut off the coleoptile and inserted thin plate of gelatin between the tip and the cut stump. He observed that coleoptile tip still bends towards unilateral light.

Paal (1919) cut off the tip of coleoptile and replaced it asymmetrically on the cut coleoptile stump. He observed that the coleoptile tip bent away from the side bearing tip even in dark.

F.W. Went (1928) successfully isolated natural auxin from *Avena* coleoptile tips. He cut off the tip and placed them on small agar blocks. Then after certain period of time placed the agar blocks asymmetrically on cut coleoptile stump that caused bending. He demonstrated the presence of substance which could diffuse into agar blocks. Went named this substance as **auxin**.



According to **Thimann and Pincus** (1948) "Plant hormones are organic substances produced naturally in higher plants affecting growth or other physiological functions at a site remote from its place of production and active in very minute (optimum) amount". Hormones are transported through phloem parenchyma (Phillips 1971).

a. Auxins (Auxien = to grow):

F. W. Went in 1931, used this term first. Auxin was isolated from urine of a person suffering from Pellagra (Kogl and H. Smit 1931). In plants, it is synthesized in growing tips or meristematic regions of plants from where it is transported to other plant parts. The most common and important natural auxin is Indole-3-acetic acid (IAA). Tryptophan is the primary precursor of IAA in plants. It is the first hormone to be discovered in plants and is primarily responsible for cell elongation. It shows polar transport - Basipetal transport in stem. Now synthetic auxins like IBA (Indole butyric acid), NAA (Naphthalene acetic acid), 2, 4-D (Dichloro Phenoxy acetic acid), etc. are used.

Physiological effects and applications of auxin:

The primary effect is cell enlargement. In most of the higher plants, growing apical bud inhibits the growth of lateral buds. This is called as **apical dominance**. Auxin stimulates growth of stem and root. Auxin induces multiplication of cells, hence used in tissue culture experiments to produce callus. It stimulates formation of lateral and adventitious roots. These are marketed as synthetic herbicides. e.g. 2, 4-D (2,4 dichlorophenoxy acetic acid). It kills dicot weeds without affecting monocot crop plants.

The seedless fruits like orange, lemon, grapes, banana etc. are produced by application of auxin (i.e. induced parthenocarpy). Auxins promote cell division in cambium and also cause early differentiation of xylem and phloem. It

promotes root elongation in low concentration and shooting at higher concentration. It also hastens early rooting in propagation by 'cutting'.

Foliar spray of NAA and 2,4-D induces flowering in litchi and pineapple. Likewise, it prevents premature fruit drop in apples, pear and oranges, and also prevents formation of abscission layer. Auxins play a role in elongation of cell. It is known to increase rate of respiration. Auxins break dormancy in seed and promote quick germination.



Do you know ?

Agent orange : Mixture of two phenoxy herbicides in ester form viz. 2, 4-D and 2,4, 5-T (dioxin), is known as agent orange used in Vietnam war for defoliation of forests.

b. Gibberellins :

It is another growth promoting hormone and is abundant in root tip and developing seeds. It shows non-polar transport through vascular tissue.

Gibberellins were first isolated from the fungus *Gibberella fujikuroi* by a Japanese scientist **Kurosawa** (1926). He observed that when rice plant was infected by fungus *Gibberella fujikuroi*, it shows extensive stem elongation called 'bakane disease'. The crystalline form of Gibberellins were isolated by Yabuta and Sumiki (1938) from the fungus culture. They named it as gibberellin. It is synthesized in young leaves, seeds, roots and stem tips. These are synthesized from mevalonic acid. More than 150 chemical types are known so far. GA₃ (= Gibberellic acid) is most common and biologically active form. Chemically it contains a gibbeane ring- a cyclic diterpene with four isoprene units.

Physiological effects and application of Gibberellins :

Dormancy of bud can be broken by gibberellin treatment. It can promote seed germination in cereals like barley and wheat

by synthesizing hydrolysing enzyme amylase to produce sugar. The most striking effect of it, is the elongation of stem where internodes increase in length and converting genotypic dwarf to phenotypic tall plant. It also promotes bolting i.e. elongation of internodes just prior to flowering in plants those with rosette habit e.g. beet, cabbage. It causes parthenocarpy in tomato, apple and pear, and flowering in long day plants. It is used to increase the fruit size and bunch length of grapes. When gibberellins are applied on genetically dwarf plants like maize, the stem rapidly elongates and acquires the height of normal tall varieties of maize. Application of gibberellins overcomes the requirement of vernalization. Usually, it inhibits growth of root, delays senescence and prevents abscission. It also breaks dormancy of seed and hastens germination. Application of gibberellin causes production of male flowers on female plant.

c. Cytokinin :

It is another growth hormone that promotes cytokinins during cell division. Letham coined the term cytokinin. The first cytokinin was discovered by Skoog and Miller (1954) during investigation of nutritional requirements of callus tissue culture of *Nicotiana tabacum* (Tobacco). They observed that the callus proliferated when the nutrient medium was supplemented with coconut milk and degraded sample of DNA (obtained from herring sperm). They named it as **kinetin**. Chemically kinins are 6-furfuryl amino purine. First natural cytokinin was obtained from unripe maize grains by Letham et al. It is known as Zeatin. 6-benzyl adenine is a synthetic cytokinin hormone. Seven different types of cytokinins are recorded from plants. Natural cytokinins are also reported from plants like Banana flowers, apple and tomato fruits, coconut milk, etc.

Physiological effects and applications of cytokinin :

Besides cell division, it also promotes cell enlargement. High cytokinin promotes shooting.

A low ratio of cytokinin to auxin induces root development but a high ratio causes buds and shoot to develop. Cytokinin and auxin ratio and their interactions controls morphogenic differentiation. It promotes the growth of lateral buds and controls apical dominance by cell division. It delays the senescence or ageing and abscission processes in plant organs. This was reported by Richmond and Lang (1957). Formation of interfascicular cambium and expansion of cells are other functions. It also breaks dormancy and promotes the germination of seeds. Cytokinin reverses apical dominance effect. It induces RNA synthesis and formation of interfascicular cambium.

d. Ethylene :

It is the only gaseous growth regulator. Denny (1924) reported ethylene is effective in fruit ripening. Gane (1934) established that plants naturally synthesize ethylene. Crocker (1930) proposed that ethylene is the plant hormone responsible for fruit ripening. It is a simple gaseous hydrocarbon with essential role in the fruit ripening. The most widely used compound as a source of ethylene is ethephon. It is synthesized in roots, shoot apical meristem, ripening fruits etc.



Do you know ?

Ethret / Ethephon is a 2-chloroethyl Phosphoric acid, which releases ethylene after dissolving in water.

Physiological effects and application of ethylene :

It promotes ripening of fruits like bananas, apples and mangoes. It stimulates initiation of lateral roots in plants and breaks the dormancy of bud and seed.

It accelerates the abscission activity in leaves, flowers and fruits by forming of abscission layer. Ethylene inhibits the growth of lateral buds and causes apical dominance and retards flowering. It is associated with

the enhancement of process of senescence of plants organs. It inhibits flowering in most of the plants except pineapple. It causes **epinasty** (drooping) of leaves and flowers. It increases activity of chlorophyllase enzyme causing degreening effect in banana and *Citrus* fruits.



Internet my friend

Degreening: It is the process of decomposition of green pigment in fruits usually by applying ethylene. This method is called trickle degreening. Collect more information about degreening.

e. Abscissic Acid :

It is a natural growth inhibiting hormone. Carns and Addicott (1961-65) observed that the shedding of cotton balls was due to a chemical substance **abscisin** I and II. Wareing (1963) isolated a substance from buds of **Acer** that can induce bud dormancy and named it **dormin**. These two identical chemical substances were given the common name abscissic acid. It is synthesized in leaves, fruits, roots, seeds etc. Chemically, it is a 15-carbon sesquiterpenoid and is synthesized from mevalonic acid.

Physiological effects and application of ABA:

It promotes abscission of leaves and induces dormancy in many plants. It controls the dormancy in buds and seeds by inhibiting growth processes. It accelerates the senescence of leaves, flowers and fruits. It inhibits and delays cell division and cell elongation and suppresses cambium activity by inhibiting mitosis in vascular cambium. ABA could cause efflux of k^+ ions from the guard cells and result in closure of stomata. So, it is known as an **antitranspirant**. It acts as a stress hormone by inducing the plant to bear the adverse environmental conditions. It inhibits flowering in long day plants but stimulates flowering in short day plants.



Do you know ?

Bioassays for :

Auxins - *Avena* curvature test.
 Gibberellins - Alpha amylase bioassay.
 Cytokinins - Chlorophyll retention test.
 Ethylene - Trippele pea test.
 ABA - Inhibition of alpha amylase test.



Internet my friend

Collect information of synthetic plant hormones.



Can you tell?

1. Why is auxin called a growth regulator?
2. Effect of Gibberellin application on apple.
3. How can we overcome apical dominance?
4. Which is standard bioassay method for auxins?
5. ABA is called as stress hormone. Why?

7.10 Photoperiodism :

Higher plants reproduce sexually by producing special structures called **flowers**. Plants exhibit transition from vegetative growth to reproductive growth during which flowers are produced. Like vegetative growth, reproductive growth is also influenced by several environmental and nutritional factors. Among the environmental factors - light and temperature exert profound influence on flowering. The influence of duration of light is known as **Photoperiodism** and that of temperature, is **Vernalization**.

Light as an environmental factor influences germination of seed, vegetative growth, photosynthesis, etc. Light as a factor has three aspects viz, **Quality**, **Intensity** and **Duration of light**. It is the duration of light that has profound effect on flowering in higher plants.

The term **photoperiodism** was used by Garner and Allard (1920). They were studying

the flowering behaviour in plants - Soyabean and Maryland mammoth variety of tobacco. They found that soyabean plant flowers during late summer and tobacco variety during winter, irrespective of their germination and growing season. They studied effects of different temperatures, nutrition, soil moisture, etc. in respect of flowering. None of these were found to regulate flowering. However, experimentally they found that the exposure to specific duration of light (i.e. photoperiod) had profound influence on flowering. They examined the effect of day length on flowering by using artificial illumination. They concluded that the relative length of the day was most crucial in the growth and development of flowers to which they coined the term **photoperiodism**. Based on the photoperiodic response, plants were classified in three categories viz, **Short Day Plants (SDP)**, **Long Day Plants (LDP)** and **Day Neutral Plants (DNP)**.



Do you know ?

Organs for reception of photoperiodic stimulus - Leaf is the chief organ for receiving the photoperiodic stimulus as demonstrated by Knoff (1934). Defoliated plants will not flower even if the plants are exposed to proper duration of light.

Photoperiodic stimulus - It is a chemical stimulus transported through phloem and is called **florigen** which is hormonal in nature.

Photochemical receptors in the leaves are the biloproteins (pigments) located in the cell membrane. These are called **Phytochromes**.

Blue wavelengths of light influences flowering.

a. Short Day Plants (SDP) :

These plants usually flower during winter and late summer when day length is shorter than the **critical photoperiod** (critical photoperiod is that length of photoperiod above or below which flowering occurs). These are called

long night plants because they require long uninterrupted dark period/ night for flowering. If dark period is interrupted even by a flash of light, SDP will not flower. Some of the short day plants are Dahlia, Aster, Tobacco, *Chrysanthemum*, Soybean (*Glycine max*), Cocklebur (*Xanthium*), etc.

b. Long Day Plants (LDP) :

Plants that flower usually during summer are called long day plants. They require longer duration of light than the critical photoperiod, for flowering. They are called short night plants as they require short dark period. When long dark period is interrupted by a brief flash of light, LD plants can flower e.g. pea, radish, sugar beat, cabbage, spinach, wheat, poppy, etc.

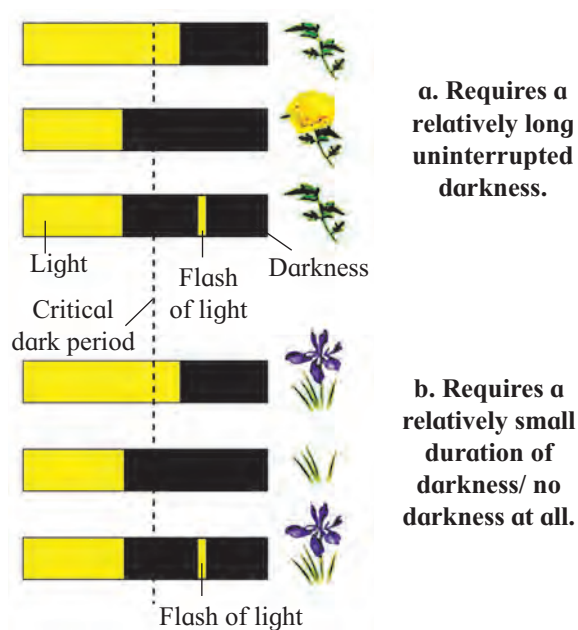


Fig. 7.9 : Photoperiodism

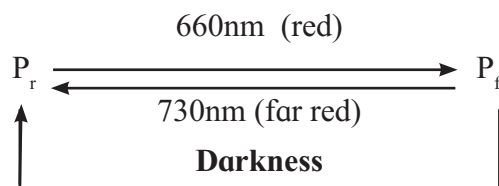
c. Day Neutral Plants (DNP) :

These plants flower throughout the year round, independent of duration of light (photoperiod). They do not require specific photoperiod to flower. Therefore, they are called Day neutral plants e.g. Cucumber, tomato, cotton, sunflower, maize, balsam, etc.

Phytochrome :

Hendricks and Borthwick (1952) observed that flowering in SD plants is inhibited, if dark period is interrupted even by a flash of red light of 660 nm. If it is immediately followed by far red light (730 nm), then SD plants will flower. This observation led them to conclude that some pigment system in plant receives the photoperiodic stimulus. These pigment proteins are called **phytochromes**.

The leaves possess light-receiving proteinaceous pigment called **phytochrome** that induces flowering. It exists in two interconvertible forms viz, red (P_r) and far red (P_{fr}). When P_{fr} absorbs far red light, it is converted into P_r and vice versa. These are located in the cell membrane of green cells.



During day time, P_{fr} accumulates in the plants. It inhibits flowering in SDP but initiates flowering in LDP. During dark period P_{fr} changes into P_r , which stimulates/ promotes flowering in SDP and inhibits flowering in LDP.



Always Remember

Control of morphogenesis by light and phytochrome, is called photomorphogenesis.



Internet my friend

Melchers proposed that chilling treatment may induce formation of hypothetical floral hormone, vernalin.

7.11 Vernalization (Yarovization) :

Temperature as environmental factor influences several physiological processes including reproduction. Temperature as a factor has three cardinal points viz, minimum,

optimum and maximum temperature. It is a low temperature (chilling) treatment that induces early flowering in plants as was evidenced by Klippart (1918). Chouard (1960) defines vernalization as acceleration of the ability to flower by chilling treatment. The term **vernalization** was coined by T.D Lysenko (1928) for the effect of low temperature on flowering in plants.

It is an influence of temperature on development and flowering. Many plants such as cereals, crucifers require a period of cold treatment for flowering. It is the method of inducing early flowering in the plants by pretreatment to their seeds/ seedlings at low temperature (1-6°C for one to one and half months' duration). The site of vernalization is believed to be shoot apical meristem. Generally, vernalization is effective at seed stage in annual plants. Vernalization stimulus is also a chemical stimulus named as **vernalin**. This can be transferred through grafting (Melcher 1939).



Know the Scientist :

The low temperature induces early flowering which was first noticed by **Klippart** (1957) while working on two varieties of wheat- 'winter wheat' and 'spring wheat'. The winter wheat sown in winter, produces flowers in the following summer. The spring wheat sown in the spring and produces flowers during summer. However, if winter variety is sown in spring does not produce flowers during summer but does so in the next year's summer. He observed that if germinated seeds of 'winter wheat' are treated with low temperature, it can behave as spring wheat. He concluded that winter wheat requires low - temperature exposure before the onset of flowering.

The reversion of vernalization by high temperature treatment is called **devernalization**.

Advantages of vernalization :

- Crop plants by shortening juvenile phase attains early flowering.
- Crops can be cultivated in regions where they do not grow naturally.

7.12 Mineral nutrition :

Plant absorbs water, gases, minerals and nutrients, from surroundings. Green plants for the synthesis of their organic food need inorganic substances (elements) which are obtained from soil in the form of minerals. Minerals constitute most commonly occurring solid, inorganic materials obtained from the earth's crust.

Chemical analysis of plant ash clearly indicates that plant absorbs mineral elements from surroundings (soil, air and water) for its use. About 36 to 40 different elements of periodic table are used as minerals by the plants. These are absorbed in ionic (dissolved) form as PO_4 , CO_3 , SO_4 , etc., usually through roots (regions of elongation and growth).

Sources of minerals :

Plants derive necessary elements from the atmosphere, soil and water. Carbon enters the plant as atmospheric carbon-dioxide.

Source of hydrogen is water and oxygen comes from air and water. Carbon, Hydrogen and oxygen are not minerals in origin.

Source of nitrogen is the soil. Plant derives nitrogen from both mineral and non-mineral origin.

Classification of minerals :

Earlier, on the basis of their requirement minerals were classified as **essential** and **non-essential**. Essential minerals are those that are indispensable without which plants can not complete their life cycle e.g. C, H, O, N, P, etc. These elements play structural and physiological roles. Their absence can produce/ cause major **deficiency symptoms**. The non-essential elements are not indispensable and they do not produce/ cause any deficiency symptoms. This classification is obsolete now.

Based on the quantity requirement, minerals are classified as minor or **microelements** and major or **macroelements**. Microelements are required in traces because they function in the catalytic role e.g. Zn, Cu, Al, Si, Mn, B, Mo, Cl etc. as co-factors. Macroelements are required in large quantity. They mainly play the nutritive and structural roles e.g. C, H, O, P, Mg, N, K, S, Ca and Fe. C, H, O are non mineral major elements. This classification is not accepted now.



Know the Scientist :

Criteria for Essentiality of elements proposed by Arnon and Stout (1939) : The elements must be absolutely necessary for supporting normal growth and reproduction. These elements must be having specific functions and should not be replaceable by another element. The element must be directly involved in the nutrition of the plant. If an element satisfies all above three criteria, it is considered to be an **essential** element.



Do you know ?

Liebig showed the Essentiality of mineral nutrients and put forth the 'Law of minimum'. According to this law, the yield of crop plant is determined by the amount of essential element that is present (available) in the minimum quantity.

Now a days minerals are classified on the basis of their biochemical functions.

Table : 7.10 Roles of Mineral Elements in Plants

Element	Region of plant in which required	Functions	Deficiency symptom
Nitrogen NO_2^- or NO_3^- or NH_4^+	Everywhere particularly in meristematic tissues	Constituent of proteins, nucleic acids, vitamins, hormones, coenzymes, ATP, chlorophyll.	Stunted growth, chlorosis.
Phosphorus H_2PO_4^- or HPO_4^{2-}	Younger tissues, obtains from older, metabolically less active cells	Constituent of cell membrane, certain proteins, all nucleic acids and nucleotides required for all phosphorylation reactions.	Poor growth, leaves dull green.

Symptoms of Mineral deficiency in plants :

Any visible deviation from the normal structure and function of the plant, is called **symptom** or hunger sign. The concentration of the essential elements below which plant growth is retarded, is termed as **critical concentration**.

The element is said to be deficient when present below the critical concentration. Certain morphological changes are indicative of the deficiency of particular element. Deficiency symptoms also depend on the mobility of the elements in the plants. The deficiency symptoms appear first in young tissues when elements are relatively immobile e.g. sulphur, calcium.

When the elements are actively mobilized within the plants and exported to young developing tissues, the deficiency symptoms are visible first in the older tissues (senescent leaves). e.g. nitrogen, magnesium, potassium.

Some important deficiency symptoms seen in plants are:

- **Stunting** : The growth is retarded. The stem appears condensed and short.
- **Chlorosis** : It is the loss or non-development of chlorophyll resulting in the yellowing of leaves
- **Necrosis** : It is the localized death of tissue of leaves.
- **Mottling** : Appearance of green and non-green patches on the leaves.
- **Abscission** : Premature fall of flowers, fruits and leaves.

Potassium K^+	Meristematic tissues, buds, leaves, root tips	Helps in determining anion- cation balance in cells involved in protein synthesis, involved in formation of cell membrane and in opening and closing of stomata; increases hardness; activates enzymes and helps in maintenance of turgidity of cells.	Yellow edges to leaves, premature death.
Calcium Ca^{2+}	Meristematic and differentiating tissues, accumulates in older leaves	Involved in selective permeability of cell membranes, activates certain enzymes required for development of stem and root apex and as calcium pectate in the middle lamella of the cell wall.	stunted growth.
Magnesium Mg^{2+}	Leaves, withdrawn from ageing leaves and exported to developing seeds	Activates enzymes in phosphate metabolism, constituent of chlorophyll, maintains ribosome structure.	Chlorosis
Sulphur SO_4^{2-}	Stem and root tips; young leaves remobilised during senescence	Constituent of certain proteins, vitamins (thiamine, biotin CoA) and Ferredoxin.	Chlorosis
Iron Fe^{3+}	Everywhere carries along leaf veins.	Constituents of ferredoxin and cytochrome, activates catalase required for synthesis of chlorophyll.	Chlorosis
Manganese (trace) Mn^{2+}	Leaves and seeds	Activates certain enzymes (carboxylases)	Chlorosis, grey spots on leaves.
Molybdenum (Trace) MoO_2^{2+}	Everywhere, MoO_3^{3+} particularly in roots	Activates certain enzymes in the nitrogen metabolism.	Slight retardation of growth.
Boron (trace) BO_3^{3-} or $B_4O_7^{2-}$	Leaves and seeds	Required for uptake and utilisation of Ca^{2+} , pollen germination and cell differentiation, carbohydrate translocation.	Brown heart disease.
Copper (trace) Cu^{2+}	Everywhere	Activates certain enzymes.	Die-back of shoots.
Zinc (trace) Zn^{2+}	Everywhere	Activates various enzymes especially carboxylases, part of carbonic anhydrase and various dehydrogenases needed for auxin synthesis	Malformed leaves.
Chlorine Cl^-	Everywhere	With Na^+ and K^+ helps to determine solute concentration and anion-cation balance in cells, essential for oxygen evolution in photosynthesis.	Poor growth of the plant



Do you know ?

- **Constituents of protoplasm and cell walls** : C,H,O,N,P,S are important and permanent constituents of the protoplasm and the cell wall.
- **Osmotic potential** : The osmotic pressure of cell is the result of mineral salts and organic compounds dissolved in the cell sap.
- **Catalytic functions** : Many elements like Fe, Cu, Zn, Mg, Mn, Cl, Mn, Mo, B act as catalysts in various enzyme reaction of plant metabolism.
- **Permeability of cytoplasmic membrane:** Permeability of membrane is influenced by the presence of the various 'cations' and 'anions' of the mineral elements in the external medium like Ca^{+} , K^{+} .
- **Components of energy related chemical compounds** : Phosphorus in ATP and Mg in chlorophyll.



Do you know ?

The atmospheric CO_2 is virtually the only source of carbon, which is the basic constituent of all the organic compounds.



Internet my friend

- Collect information about hydroponics and find its use in kitchen garden.
- Collect information on Aeroponics.

Toxicity of Micronutrients :

Mineral ion concentration which reduces the dry weight of tissues by 10%, is considered as toxic. Toxicity of one element may inhibit the uptake of the other e.g. Mn competes with Fe, Mg for uptake, but inhibits Ca translocation to shoot apex. Thus, Mn toxicity develops deficiency symptoms of Ca, Mg and Fe. The phenomenon where toxic effect of one mineral

is counter balanced by the other ion, is called antagonism.

Minerals salt absorption :

Most minerals in the soil are charged particles hence, they can not pass across cell membrane. Hence most of the minerals are absorbed actively with the expenditure energy. Minerals can also be absorbed passively without expenditure of energy. Mineral ion absorption is independent of water absorption.

Mineral ion absorption can occur in two ways :

a. Passive Absorption : Movement of mineral ions into the root occurs by diffusion. Molecules or ions diffuse from a region of their higher concentration to a region of their lower concentration. The movement of mineral ions into root cells as a result of diffusion is without expenditure of energy is called passive absorption. Passive absorption can take place by direct ion-exchange, in direct ion-exchange mass flow and Donnan equilibrium.

b. Donnan equilibrium : It is based on the assumption that certain negatively charged ions (i.e.anions), after their entry into the cell, become fixed on the inner side of the cell membrane and can not diffuse outside through the cell membrane. Therefore, additional mobile cations are required to balance these fixed anions. Obviously concentration of cations become more due to accumulation. This kind of passive absorption of anions/ cations from cell exterior against their own concentration gradient in order to neutralize the effect of cations/ anions, is called **Donnan equilibrium**.

Active Absorption: Uptake of mineral ions against concentration gradient, is called active absorption, such movement requires an expenditure of energy by the absorbing cell. This energy is derived from respiration and is supplied through ATP. When the roots are deprived of oxygen, they show a sudden drop in active absorption of minerals. The mineral ions accumulated in the root hair pass into the cortex and finally reach the xylem.



Internet my friend

Enlist names of symbiotic and free living nitrogen fixing bacteria.

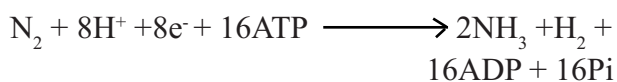


Always Remember

Today nearly 80% of nitrogen found in human tissues originate from the Haber-Bosch process.

Biological Nitrogen fixation : It is carried out by prokaryotes called as 'Nitrogen fixers' or Diazotrophs'. It accounts nearly 70% of natural nitrogen fixation. Nitrogen fixers are either **symbiotic** or **free living**. The cyanobacteria fix significant amount of nitrogen in specialized cells called **heterocysts**.

Nitrogen fixation is high energy requiring process (endothermic reaction) and nitrogen fixers use 16 molecules of ATP to fix each molecule of nitrogen to form ammonia.

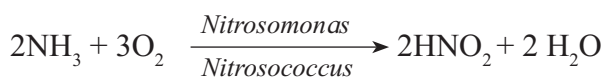


Ammonia is then converted into amino acids.

Nitrification :

Nitrification is exothermic reaction. Most of the soil bacteria participate in converting ammonia into nitrate, the form of nitrogen which can be used by plants and animals. This involves two steps performed by two different types of bacteria.

First a soil bacteria convert ammonia into nitrogen-di-oxide (nitrite) eg. *Nitrosomonas*, *Nitrosococcus*, etc.



Then another type of soil bacterium called *Nitrobacter* adds a third oxygen atoms to create nitrate.



These bacteria are chemoautotrophs. By metabolizing nitrogen along with oxygen, they

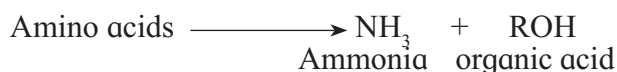
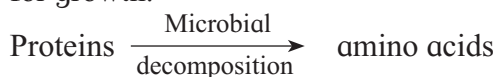
obtain energy to power their own life processes.

Symbiotic N₂ fixation :

The best known nitrogen fixing symbiotic bacterium is *Rhizobium*. This soil living/dwelling bacterium forms root nodules in plants belonging to family Fabaceae e.g. beans, gram, groundnut etc.

Ammonification :

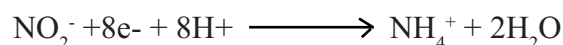
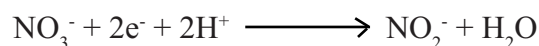
After the death of plants and animals, various fungi, actinomycetes and some ammonifying bacteria decompose the tissues and convert organic nitrogen into amino acid and then to ammonia and back into the ecosystem. Ammonia (NH₄⁺) is now available for uptake by plants and other micro-organisms for growth.



Nitrogen assimilation :

In soil, nitrogen is present as nitrates, nitrites and ammonia (NH₄⁺). It is absorbed by the green plants and converted to nitrogenous organic compounds like amino acids, DNA, etc. This is known as nitrogen assimilation. From plants, nitrogen as biomolecules like amino acids, enters food chain and moves to animals and then to decomposers through the death of animals.

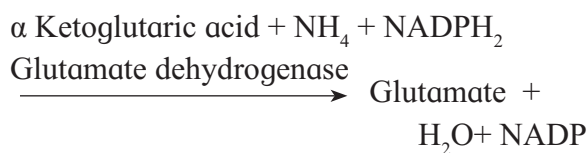
Nitrates are first converted to ammonia but it is highly toxic and immediately used for conversion into amino acids, which are then transported to other parts of the plants for synthesis of proteins.



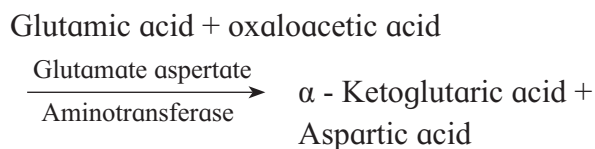
Amino Acid synthesis :

Amino acids are building blocks of proteins. The amino acids are synthesized through :

Reductive amination : Ammonia reacts with alpha ketoglutaric acid to form glutamic acid (glutamate).

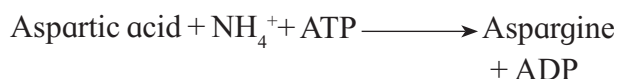
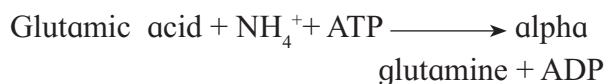


Transamination : Amino group of one amino acid (-CHNH₂) is transferred to keto position (-CO) of other carboxylic acid.



Amides :

Ammonia may be absorbed by amino acid to produce amides. The process is called **amidation**. The amides are the amino acids having two amino groups. Extra amino group is attached to acidic group (-COOH) in presence of ATP.



Amides like glutamine and asparagine are formed from glutamic acid and aspartic acid respectively by addition of another amino group to each. Amides are transported to other parts of plants via xylem vessels.

Denitrification :

It is the process in which anaerobic bacteria can convert soil nitrates back into nitrogen gas. Denitrifying bacteria removes fixed nitrogen i.e. nitrates from the ecosystem and return it to the atmosphere in inert form.

Denitrifying bacteria includes *Bacillus spp.*, *Paracoccus spp.* and *Pseudomonas denitrificans*. They transform nitrates to nitrous and nitric oxides and ultimately to gaseous nitrogen.



Sedimentation : Nitrates of the soil are washed away to the sea or leached deep into the earth along with percolating water.



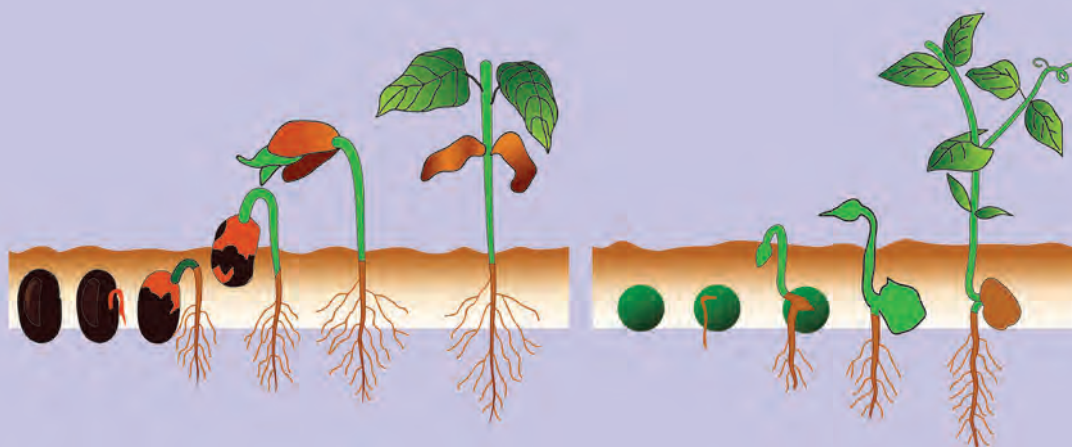
Do you know ?

1. Soil nitrogen is replenished through the nitrogenous waste of animals, (as ammonia, urea and uric acid) ammonification and nitrification.
2. Plastids contain nitrite reductase which reduces it to ammonia.



Activity :

Label the following diagrams and identify the types of germination.



Exercise

Q. 1 Multiple choice questions.

- Which of the hormones can replace vernalization?
 - Auxin
 - Cytokinin
 - Gibberellins
 - Ethylene
- The principle pathway of water translocation in angiosperms is
 - Sieve cells
 - Sieve tube elements
 - Xylem
 - Xylem and phloem
- Absciscic acid controls
 - cell division
 - leaf fall and dormancy
 - shoot elongation
 - cell elongation and wall formation
- Which is employed for artificial ripening of banana fruits?
 - Auxin
 - Ethylene
 - Cytokinin
 - Gibberellin
- Which of the following is required for stimulation of flowering in the plants?
 - Adequate oxygen
 - Definite photoperiod
 - Adequate water
 - Water and minerals
- For short day plants, the critical period is
 - light
 - dark/ night
 - uv rays
 - both a and c
- Which of the following is day neutral plant?
 - Tomato
 - Cotton
 - Sunflower
 - Soybean
- Essential macroelements are
 - manufactured during photosynthesis
 - produced by enzymes
 - absorbed from soil
 - produced by growth hormones
- Function of Zinc is
 - closing of stomata
 - biosynthesis of 3-IAA
 - synthesis of chlorophyll
 - oxidation of carbohydrates
- Necrosis means
 - yellow spots on the leaves
 - death of tissue
 - darkening of green colour in leaves
 - wilting of leaves
- Conversion of nitrates to nitrogen is called
 - ammonification
 - nitrification
 - nitrogen fixation
 - denitrification
- How many molecules of ATP are required to fix one molecule of nitrogen?
 - 12
 - 20
 - 6
 - 16

Q. 2 Very Short Answer Questions :

- Enlist the phases of growth in plants?
- Give the full form of IAA?
- What does it mean by 'open growth'?
- Which is the plant stress hormone?
- What is denitrification?
- Name the bacteria responsible for conversion of nitrite to nitrate.
- What is role of gibberellin in rosette plants?
- Define vernalization.
- Define photoperiodism.
- What is grand period of growth?

Q. 3 Short Answer Questions :

1. Write a short note on :
a. Differentiation b. Redifferentiation
2. Differentiate between Arithmetic and Geometric growth.
3. Enlist the role and deficiency symptoms of :
a. Nitrogen b. Phosphorus c. Potassium
4. What is short day plant? Give any two examples.
5. What is vernalization? Give its significance.

Q. 4 Long Answer Questions :

1. Explain sigmoid growth curve with the help of diagram.
2. Describe the types of plants on the basis of photoperiod required, with the help of suitable examples.
3. Explain biological nitrogen fixation with example.
4. Write on macronutrients and micronutrients required for plant growth.
5. How are the minerals absorbed by the plants?

Project :

1. Grow seed in the deep tray on soil medium and study different stages of germination. Prepare powerpoint presentation with the help of pictures/photographs of the same.
2. Prepare chart differentiating the epigeal and hypogeal germination.
3. Collect the information about the **viviparous germination** in plants growing along seashore.
4. Identify and label the deficiency symptoms in the given diagram.

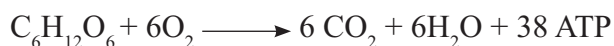


**Can you recall?**

1. Definition of respiration.
2. The types of cellular respiration.
3. Why should we respire all the time?

All living organisms require energy to carry out various life processes. The energy that is stored in the body in the form of complex organic compounds (potential energy) is however not usable by the organisms unless it is converted into usable form. This conversion is achieved through the process of respiration.

Respiration : It is a biochemical process of oxidation of organic compounds in an orderly manner for the liberation of chemical energy in the form of ATP.



For this, the process of gaseous exchange takes place between the organism and the environment. The site of gaseous exchange is called the respiratory surface.

**Use your brain power**

What would happen if respiration take place in one single step?

8.1 Organs of Respiratory Exchange :

Respiratory exchange is a simple physical process. For efficient gaseous exchange, the respiratory surface should have the following features :

- a. It should have a large surface area.
- b. It should be thin, highly vascular and permeable to allow exchange of gases.
- c. It should be moist.

Gaseous exchange in plants :

The shape and structure of plants facilitate gaseous exchange by diffusion. A terrestrial flowering plant has many air spaces between

the cells of stem, leaf and root. These air spaces are continuous. Exchange of gases takes place between the air and the tissue. Oxygen diffuses into the air space through stomata (the pores on leaves and young stems), carbon dioxide and water vapour diffuse out. In the aerated soil, the oxygen dissolves in the film of moisture or water around the root tissue and enters it by diffusion.

Woody flowering plants (trees and shrubs) have an external impervious bark. Here, gaseous exchange occurs through small pores in the stem surface, called lenticels.

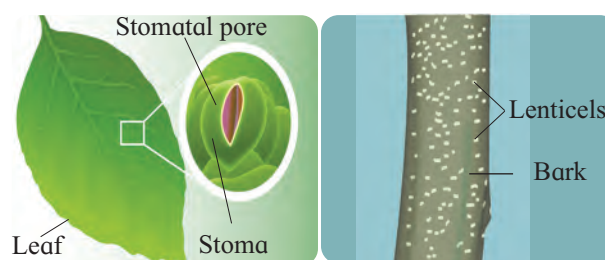


Fig. 8.1 : Organs of gaseous exchange in plants

Curiosity Box :

1. What are the main features of respiratory surface?
2. Which are the parts of plant that help in the process of gaseous exchange?

Respiration in Animals :

As compared to plants, animals show wide variety of respiratory surfaces or organs. The respiratory surfaces differ in various animals. In animals, depending upon the complexity of organization and the surrounding medium, certain parts of the body have become specialized into different types of respiratory organs. In the higher animals, these respiratory organs are also associated with a transport system.

Organism	Habitat	Respiratory surface/ organ
Protists, Sponges and Coelenterates	Aquatic	Plasma membrane
Flatworms like Planaria, Annelids (earthworm, nereis, leech), amphibians (frog)	Aquatic or semiaquatic	Plasma membrane, general body surface (moist skin)
Insects	Terrestrial	Tracheal tubes and spiracles
Arachnids like spiders and scorpions	Terrestrial	Book lungs
<i>Limulus</i> (Arthropod)	Aquatic	Book gills
Amphibian tadpoles of frog, salamanders and newts	Aquatic	External gills
Fish	Aquatic	Internal gills
Reptiles, Birds and Mammals	Terrestrial	Lungs
Turtles	Underwater	cloaca

Table 8.2 : Respiratory surface/ organ in organisms



Use your brain power

Why large animals can not carry out respiration without the help of circulatory system?

8.2 Human Respiratory system:

The respiratory system brings about inspiration, expiration and exchange of gases in the lungs. These are then transported by blood from the lungs to the different tissues and parts of the body. The respiratory system can be divided into an upper respiratory system having external nares, nasal cavities, internal nares, nasopharynx, nose, throat and associated structures. The lower respiratory system refers to the larynx, trachea, bronchi, bronchioles and lungs.

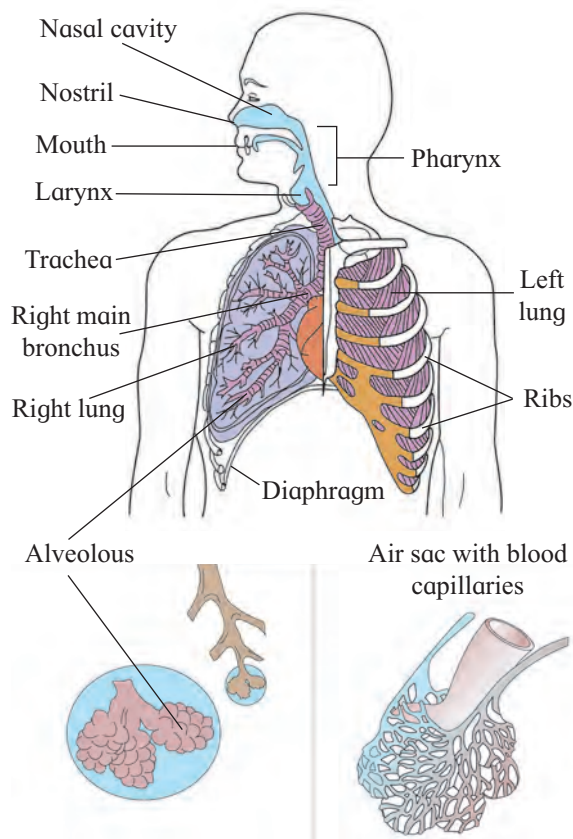


Fig. 8.3 : Human Respiratory system

Nose :

The nose has a pair of slit like openings called external nares or nostrils for entry of air into the nasal cavity. The nasal cavity is divisible into right and left nasal chambers by a **mesethmoid cartilage**. Each nasal chamber is further divided into three regions.

- i. **Vestibule :** It is the proximal part about the nostrils. Its skin has hair for filtering the air and trapping the dust and suspended particles in the inhaled air.
- ii. **Respiratory part (conditioner) :** The middle thin walled highly vascular part for warming and moistening the inhaled air.
- iii. **Olfactory or sensory chamber :** The uppermost part is lined by olfactory epithelium for detection of smell.

Pharynx :

It is divisible into three parts. The nasopharynx is the uppermost part from the nasal chamber it leads into oropharynx (common passage for food and air). This

continues below as the **laryngopharynx**. Between the nasopharynx and oropharynx is the palate bone. The pharynx has a set of lymphoid organs called **tonsils**.

Larynx :

It is called voice box. It is the part of the respiratory tract which contains vocal cords for producing sound. The larynx extends from the laryngopharynx and the hyoid bone to the trachea. It is a hollow, tubular structure. Its wall is made up of cartilage plates held by membranes and muscles. Internally, it is lined by a pair of folds of elastic vocal cords (true vocal cords). Voice is produced by passage of air between the vocal cords and modulations created by tongue, teeth, lips and nasal cavity. The larynx opens into the laryngopharynx through a slit like opening called glottis. This opening of the trachea or wind pipe is guarded by a leaf like flap called **epiglottis**. It prevents the entry of food into trachea.

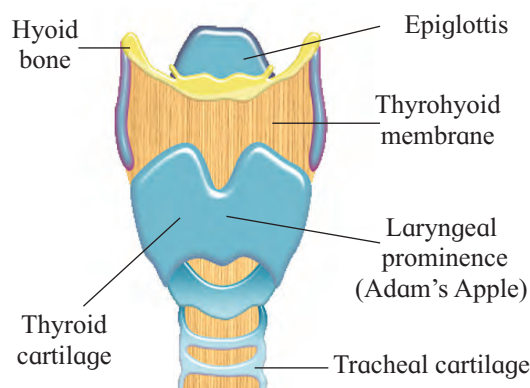


Fig. 8.4 : Larynx

Trachea (wind pipe) :

It is a long tube 10 to 12 cm in length. It runs through the neck in front of the oesophagus and extends into and upto the middle of thoracic cavity. It is supported by 'C' shaped 16 to 20 rings of cartilage which prevent the collapse of trachea. It is lined internally with ciliated, pseudostratified epithelium and mucous glands that trap the unwanted particles preventing their entry into the lungs.



Find out

- Kavya underwent a surgical procedure called Rhinoplasty. What could have been the reason for such a surgery? On which part of the body is it carried out?
- What is the role of tonsils in our body? How many pairs of tonsils do we have?



Do you know ?

- Shreyas choked while eating dinner. How can you help him? What is the immediate help that can be given to him?
- What is a role of epiglottis?

Bronchi :

The trachea divides into right and left primary bronchi as it reaches the middle of the thoracic cavity. The bronchi are supported internally by 'C' shaped incomplete rings of cartilage. The primary bronchi divide to form secondary and tertiary bronchi which lead into terminal bronchioles ending into alveoli.

Lungs :

These are the main respiratory organs of humans. One pair of spongy and elastic lungs are present in the thoracic cavity. Each lung is enclosed and protected by a double pleural membrane, outer parietal and inner visceral membrane. Between the two pleura is a pleural cavity filled with a lubricating fluid called pleural fluid. It is secreted by the membranes. The right lung is larger and divided into 3 lobes, while the left lung is smaller and divided into 2 lobes. Each lobe of the lung has the terminal bronchioles ending in a bunch of air sacs, each with 10 to 12 alveoli.

Alveoli :

These are thin walled lobulated structures, like a bunch of grapes. Each alveolus is surrounded by a network of capillaries of pulmonary arteries and veins. These have highly elastic wall made up of a single layer

of squamous epithelium resting on a basement membrane of connective tissue. There are about 700 million alveoli in the lungs and they provide the surface area for exchange of gases.



Internet my friend

1. What is the importance of pleural fluid?
2. Find the total surface covered by the alveoli.

Diaphragm : It is a muscular septum that separates the thoracic and abdominal cavity. It is dome shaped and on contraction it becomes flattened.

8.3 Mechanism of respiration :

Respiration is a biological process involving exchange of gases between the atmosphere and the lungs and it results in the formation of ATP. It includes the following processes:

- A. Breathing
- B. External respiration
- C. Internal respiration
- D. Cellular respiration



Try This

Count the number of breaths you take in the following situations (a). After a good night's sleep (b). During a vigorous activity (running, climbing stairs etc) (c). After the vigorous activity. Do you find any difference in the count?

A. Breathing :

It is a physical process by which gaseous exchange takes place between the atmosphere and the lungs. It involves inspiration and expiration (see fig. 8.5). Both these steps involved parts of the thoracic cage, the ribs, sternum and the intercostal muscles and muscles of the diaphragm.

Inspiration : During inspiration, the atmospheric air is taken in to the lungs. It occurs due to the pressure gradient formed between the lungs and the atmosphere. It is an active process in which the diaphragm becomes flat and goes downward, the external intercostal muscles contract so the ribs and sternum move upward and outward. This leads to an increase in the thoracic volume and a decrease in pressure of thorax and the lungs. To equalize the low pressure inside the lungs, air from the atmosphere rushes into lungs. This is inspiration.

Expiration : During expiration, the thorax contracts causing air to be exhaled. The diaphragm relaxes and is pushed upwards. It becomes dome shaped. The intercostal muscles also relax pulling the rib cage inward and downward. This causes a decrease in thoracic volume and leads to increase in pressure in the thorax and the lungs as compared to the atmospheric pressure. So air from the lungs rushes out. This is expiration.

One inspiration and one expiration is one breath.

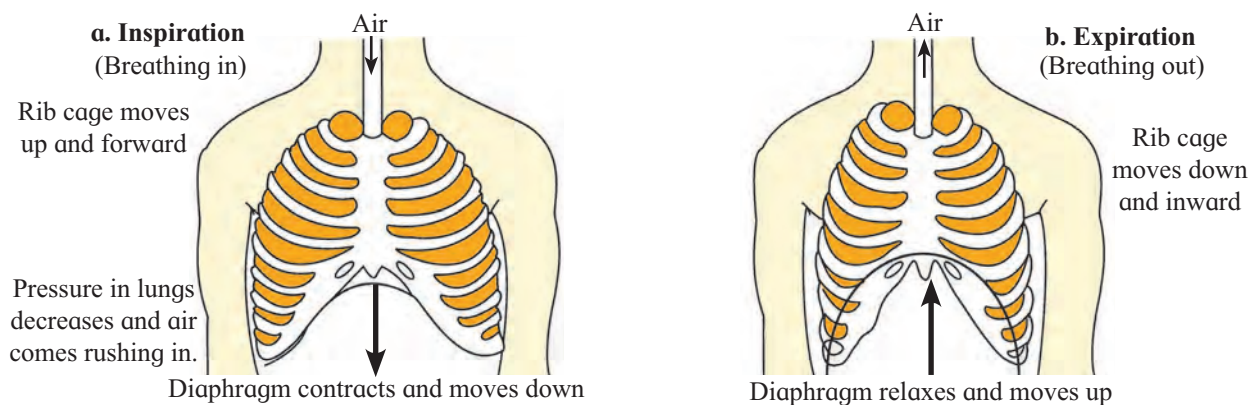


Fig. 8.5 : Breathing

B. External respiration/ Exchange of gases at the alveolar level :

An alveolus consists of a layer of simple squamous epithelium resting on a basement membrane. It is intimately associated with a dense network of capillaries. The capillary wall is also made up of simple squamous epithelium resting on a thin basement membrane. Both the layers have similar structure and are thin walled. Together they make up the respiratory membrane through which gaseous exchange occurs i.e. between the alveolar air and the blood.

Diffusion of gases will take place from an area of higher partial pressure (pp) to an area of lower partial pressure until the partial pressure in the two regions reaches equilibrium.

The partial pressure of carbon-dioxide ($p\text{CO}_2$) of blood entering the pulmonary capillaries is 45 mmHg while partial pressure of carbon-dioxide in alveolar air is 40 mmHg. Due to this difference, carbon dioxide diffuses from the capillaries into the alveolus.

Similarly, partial pressure of oxygen ($p\text{O}_2$) of blood in pulmonary capillaries is 40 mmHg while in alveolar air it is 104 mmHg. Due to this difference oxygen diffuses from alveoli to the capillaries.

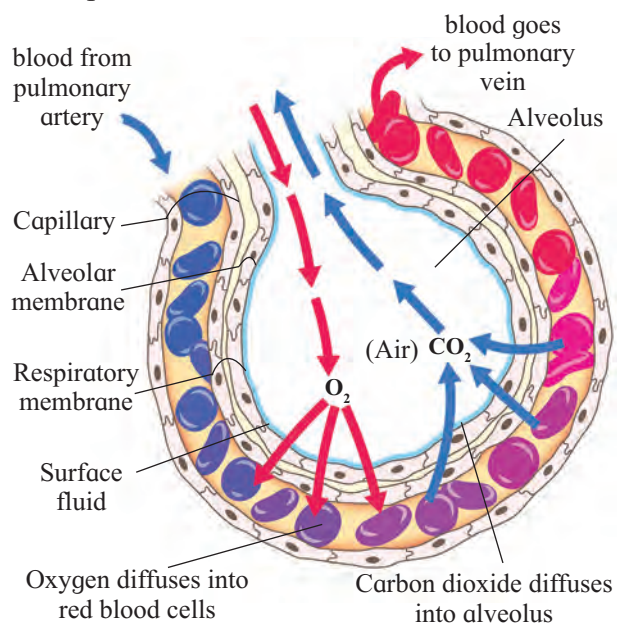


Fig. 8.6 : Exchange of gases between alveolus and capillary

Pulmonary volumes and capacities (Normal values)

Lung Volumes :

Tidal volume (T.V.) : It is the volume of air inspired or expired during normal breathing. It is 500 ml.

Inspiratory reserve volume (IRV) : The maximum volume of air, or the extra volume of air, that is inspired during forced breathing in addition to T.V. Its value is 2000 to 3000ml.

Expiratory reserve volume (ERV) : The maximum volume of air that is expired during forced breathing after normal expiration. Its value is 1000 to 1100ml.

Dead space (DS) : The volume of air that is present in the respiratory tract (from nose to the terminal bronchioles), but not involved in gaseous exchange. It is 150 ml.

Residual volume (RV) : The volume of air that remains in the lungs and the dead space even after maximum expiration. It is 1100 to 1200ml.

Lung capacities :

Total Lung capacity : The maximum amount of air that the lungs can hold after a maximum forceful inspiration (5200 to 5800ml).

Vital capacity (VC) : The maximum amount of air that can be breathed out after a maximum inspiration. It is the sum total of TV, IRV and ERV and is 4100 to 4600ml.



Use your brain power

Why gas exchange in the alveolar region is very rapid?

C. Internal respiration :

The two main components of blood involved in transport of the respiratory gases- CO_2 and O_2 , are the RBCs and the plasma.

i. Transport of oxygen :

Of the total oxygen transported only 3% is transported in a dissolved state by the plasma. The remaining 97% is bound to the haemoglobin (Hb) present in the RBCs.

Haemoglobin acts as the respiratory carrier. It has a high affinity for O_2 and combines with it to form oxyhaemoglobin. Theoretically, one molecule of Hb has 4 Fe^{++} , each of which can pick up a molecule of oxygen (O_2).



Oxyhaemoglobin is transported from lungs to the tissues where it readily dissociates to release O_2 .



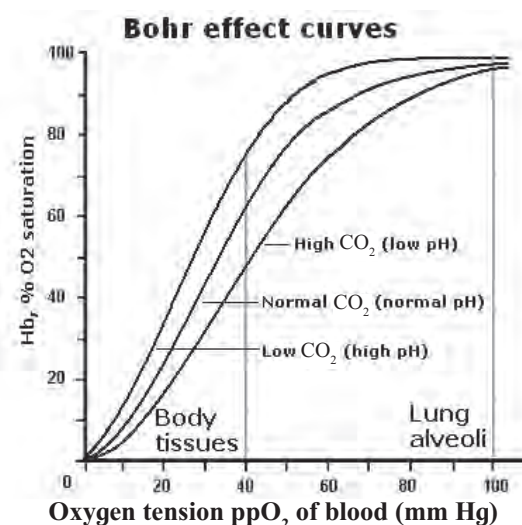
However, the degree of saturation of Hb with O_2 depends upon the O_2 tension i.e. ppO_2 .

- 100% saturation is rare.
- Maximum saturation of 95 to 97% is at ppO_2 in alveoli (100 mmHg).
- Degree of saturation decreases with the drop in ppO_2 . This begins the dissociation of HbO_2 .
- At 30 mmHg of ppO_2 , only 50% saturation can be maintained.
- The relationship between HbO_2 saturation and oxygen tension (ppO_2) is called **oxygen dissociation curve**. This oxygen - haemoglobin dissociation curve is a sigmoid curve and it shifts towards the right due to - increase in H^+ concentration, increase in $ppCO_2$ and rise in temperature and rise in DPG (2, 3 diphosphoglycerate), formed in the RBCs during glycolysis. It lowers the affinity of haemoglobin for oxygen.

Bohr effect : It is the shift of oxyhaemoglobin dissociation curve due to change in partial pressure of CO_2 in blood.

Haldane effect : Oxyhaemoglobin functions as an acid. It decreases pH of blood. Due to increase in the number of H^+ ions, HCO_3^- changes into H_2O and CO_2 .

In the alveoli where ppO_2 is high and $ppCO_2$ is low, oxygen binds with haemoglobin, but in the tissues, where ppO_2 is lower and $ppCO_2$ is high, haemoglobin does not hold as much O_2 . It releases O_2 for diffusion into the tissue cells.



Graph 8.7 : Oxyhaemoglobin dissociation curve (pp = partial pressure)

Carbon monoxide poisoning :

Affinity of haemoglobin for carbon monoxide is about 250 times more, than for oxygen. In the presence of carbon monoxide, haemoglobin readily combines to form a stable compound carboxyhaemoglobin. The haemoglobin is blocked by carbon monoxide, thus preventing oxygen from binding with it as a result haemoglobin. Thus, less haemoglobin is available for oxygen transport depriving the cells of oxygen. This is carbon monoxide poisoning.



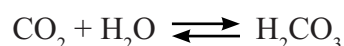
Use your brain power

While working with the car engine in a closed garage, John suddenly felt dizzy and fainted. What is the possible reason?

ii. Transport of CO_2 :

Carbon dioxide is readily soluble in water and is transported by RBCs and plasma in three different forms.

a. By plasma in solution form (7%) : Only 7% of CO_2 is transported in a dissolved form as carbonic acid (which can breakdown into CO_2 and H_2O).



b. By bicarbonate ions (70%) : Nearly 70% of carbon dioxide released by the tissue cells diffuses into the plasma and then into the RBCs.

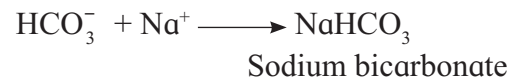
- In the RBCs, CO₂ combines with water in the presence of a Zn containing enzyme, **carbonic anhydrase** to form carbonic acid.
- Carbonic anhydrase enzyme is found in the RBCs and not in the plasma.
- The rate of formation of carbonic acid inside the RBC is very high as compared to its formation in the plasma.
- Carbonic acid being unstable almost immediately dissociates into HCO₃⁻ and H⁺ in the presence of the enzyme carbonic anhydrase (CA) leading to large accumulation of HCO₃⁻ inside the RBCs.



- It thus moves out of the RBCs. This would bring about imbalance of the charge inside

the RBCs. To maintain the ionic balance between the RBCs and the plasma, Cl⁻ diffuses into the RBCs. This movement of chloride ions is known as **chloride shift** or **Hamburger's phenomenon**.

- HCO₃⁻ that comes into the plasma joins to Na⁺ / K⁺ forming NaHCO₃ / KHCO₃ (to maintain pH of blood).



- H⁺ is taken up by protein (haemoglobin).

$$\text{Hb} + \text{H}^+ \longrightarrow \text{HHb}$$

(Reduced Hb)

These H⁺ ions might be expected to lower blood pH, but they are buffered by haemoglobin by the formation of deoxyhaemoglobin (reduced haemoglobin).

- At the **level of the lungs** in response to the low partial pressure of carbon dioxide (ppCO₂) of the alveolar air, hydrogen ion

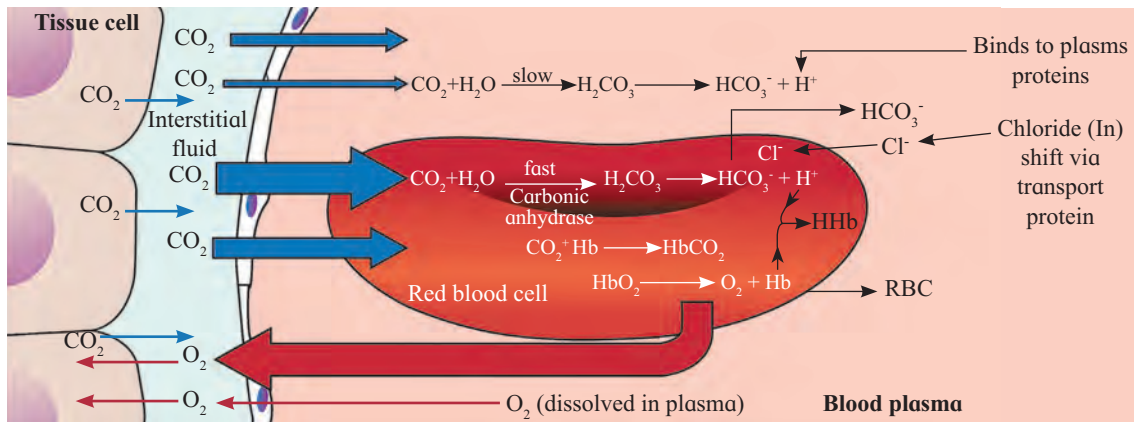


Fig. 8.8 : Oxygen release and carbon dioxide pickup at the tissue

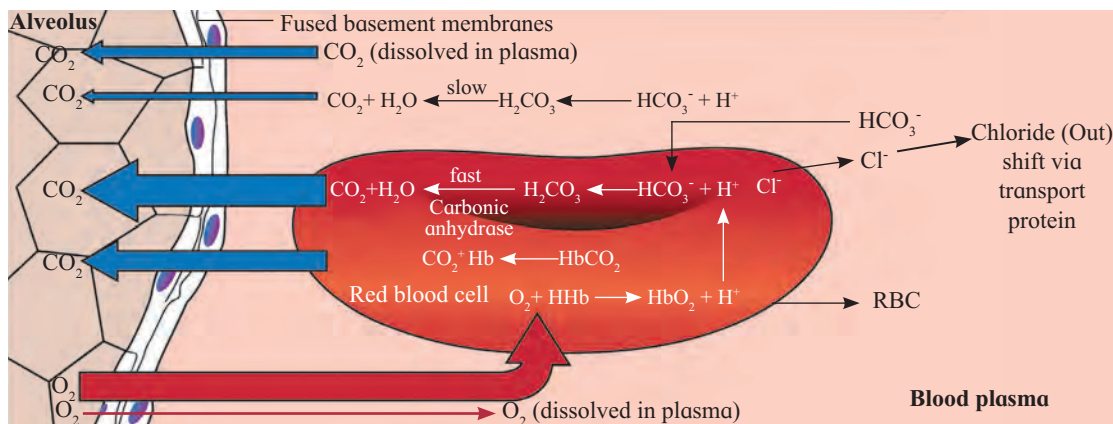


Fig. 8.9 : Oxygen pickup and carbon dioxide release in the lungs

and bicarbonate ions recombine to form carbonic acid and under the influence of carbonic anhydrase yields carbon dioxide and water.



c. By red blood cells (23%) : Carbon dioxide binds with the amino group of the haemoglobin and form a loosely bound compound carbaminohaemoglobin. This molecule readily decomposes in region where the partial pressure of carbon dioxide (ppCO_2) is low (alveolar region), releasing the carbon dioxide.



Use your brain power

1. What is the role of haemoglobin in the transport of oxygen in the blood?
2. Write a note on chloride shift.

D. Cellular Respiration :

It is the last step taking place inside the cell where food is oxidized and ATP is generated. It can be shown by two steps:

a. Oxidation : Breaking down of complex organic molecules into simple inorganic molecules with release of heat energy.



b. Phosphorylation : It involves trapping the heat energy in the form of high energy bond of ATP molecule. ATP is used to carry out vital life processes and so is called as energy currency of the cell.



8.4 Regulation of Breathing :

Respiration is under dual control : nervous and chemical. Human adults breathe about 12 times/minute while a new born about 44 times/minute. Normal breathing is an involuntary process. Steady rate of respiration is controlled by neurons located in the **pons** and **medulla** and are known as the **respiratory centres**. It regulates the rate and depth of breathing. It is divided into three groups : dorsal group of

neurons in the medulla (inspiratory center), ventro lateral group of neurons in medulla (inspiratory and expiratory center) and pneumotaxic center located in pons (primarily limits inspiration, slow wave sleep and rapid eye movement sleep). Apneustic center in the medulla is antagonistic to the pneumotaxic center. It controls non rapid eye movement, sleep and wakefulness.

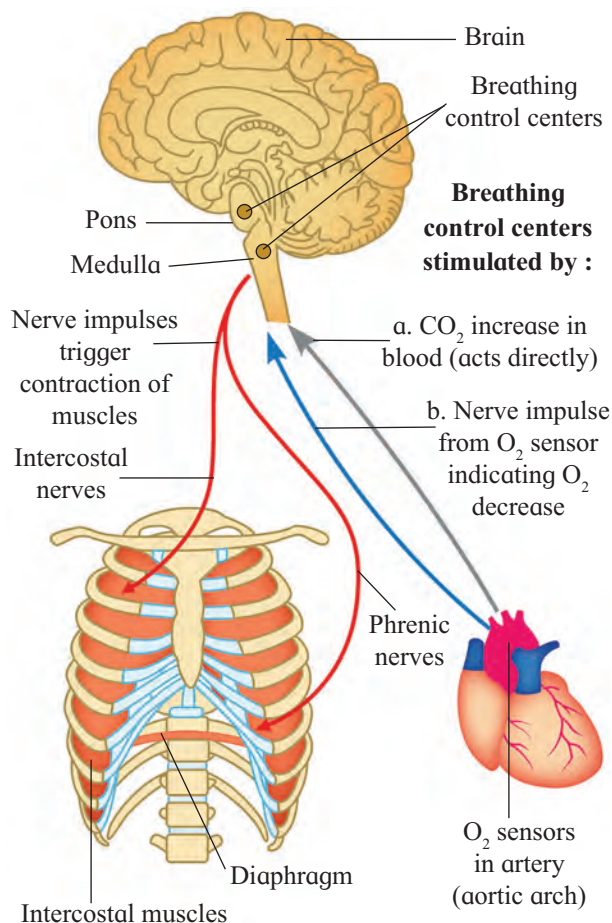


Fig. 8.10 : Regulation of Breathing

During inspiration when the lungs expand to a critical point, the stretch receptors are stimulated and impulses are sent along the vagus nerves to the expiratory centre. It then sends out inhibitory impulses to the inspiratory center.

The inspiratory muscles relax and expiration follows. As air leaves the lungs during expiration, the lungs are deflated and the stretch receptors are no longer stimulated. Thus, the inspiratory centre is no longer inhibited and a new respiration begins. These

events are called the *Hering-Breuer reflex*. The *Hering-Breuer reflex* controls the depth and rhythm of respiration. It also prevents the lungs from inflating to the point of bursting.

The respiratory centre has connections with the cerebral cortex which means we can voluntarily change our pattern of breathing. Voluntary control is protective because it enables us to prevent water or irritating gases from entering the lungs. But the ability to stop breathing is also limited by the build up of carbon dioxide in the blood.

8.5 Modified Respiratory Movements :

Some respiratory movements are different from the normal movements and help express emotion or clear the air passage. Of these

movements some may be reflexes, but others can be initiated voluntarily e.g. coughing and yawning.



Internet my friend

1. Find out information about the various modified respiratory movements and write it in a tabular form.
2. What is the significance of such movements?

8.6 Common disorders of respiratory system:

The given table shows a list of some common respiratory disorders, their symptoms, cause and treatment.

Table 8.11 : Common disorders of respiratory system

Disorder	Symptoms	Cause	Treatment
Emphysema	Breakdown of alveoli, shortness of breath	Smoking, air pollution	Quit smoking, avoid polluted air, administer oxygen to relieve symptoms
Chronic bronchitis	Coughing, shortness of breath	Smoking, air pollution	Quit smoking, avoid polluted air, if possible move to warmer, drier climate
Acute bronchitis	Inflammation of bronchi, shortness of breath, yellow mucous coughed up.	Viruses and bacteria	If bacterial, take antibiotics, cough medicine, use vaporizer
Sinusitis	Inflammation of the sinuses, mucous discharge	Viruses and bacteria	If bacterial, take antibiotics and decongestants, use vaporizer
Laryngitis	Inflammation of larynx, vocal cords, sore throat, hoarseness of voice, mucous build up and cough	Viruses and bacteria	If bacterial, take antibiotics, cough medicines, voice rest, avoid irritants like smoke
Pneumonia	Inflammation of lungs ranging from mild to severe, cough and fever, shortness of breath, chills, sweating, chest pain, blood in mucous	Bacteria, viruses	Consult physician immediately, antibiotics, cough medicines, stay warm
Asthma	Constriction of bronchioles, mucus build up in bronchioles, periodic wheezing, difficulty in breathing.	Allergy to pollen, some foods, food additives, pet hair, etc.	Use of inhalants to open passage ways, avoid irritants
Occupational Respiratory Disorders-silicosis, asbestosis	Inflammation fibrosis, lung damage.	Long term exposure to dust particles, silica and asbestos, particles during occupation	Protective mask and gear during work.



Think about it

Why is it difficult to hold one's breath beyond a limit?

Artificial ventilation :

It is also called artificial respiration. It is the method of inducing breathing in a person when natural respiration has ceased or is faltering. If used properly and quickly, it can prevent death due to drowning, choking, suffocation, electric shock, etc. The process involves two main steps : establishing and maintaining an open air passage from the upper respiratory tract to the lungs and force inspiration and expiration as in mouth to mouth respiration or by mechanical means like ventilator.

Ventilator :

A ventilator is a machine that supports breathing and is used during surgery, treatment for serious lung diseases or other conditions when normal breathing fails. It is mainly used in hospitals as part of life support system. Ventilators do the following,

1. Get oxygen into the lungs.
2. Remove carbon dioxide from the lungs.
3. Help the patient breathe.

8.7 Transportation in living organisms :

All living organisms, whether unicellular or multicellular show an important property of exchange of material with their surrounding as well as between various parts of the their cell or body.



Can you recall?

Which type of circulation is present in cockroach? How is it different from that of humans?

Organisms take up oxygen and nutrients from the surrounding, these are circulated within the body for various metabolic activities.

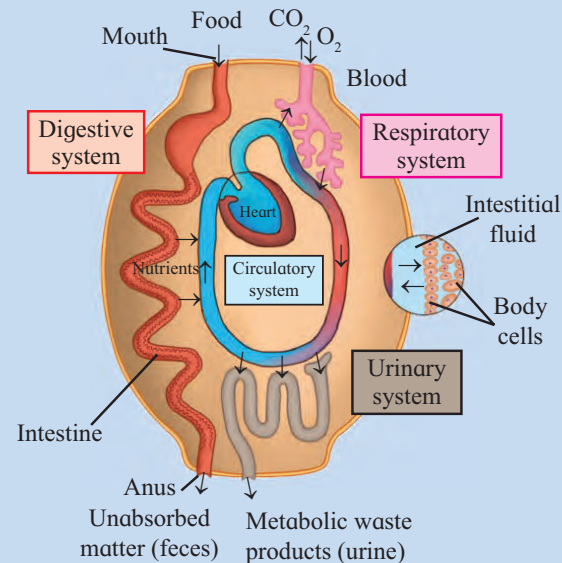
The wastes generated within are given out into the surrounding.

Transportation in organisms and animals occurs by **diffusion** and by **active transport** between the cells. This mechanism is suitable where the surface area of body is large and the distance between parts of the body in the organism is extremely small. **Cyclosis** is the streaming movement of the cytoplasm shown by almost all living organisms e.g. *Paramecium*, *Amoeba*, root hair cells of many plants and WBCs in animals. It is for transportation within the cell or **intracellular transport**. In sponges and coelenterates the surrounding water is circulated through the body cavities. In flat worms there is parenchymal circulation. In round worms there are no blood vessels and the body fluid is moved around the viscera by contraction of body wall and muscles. This is **extracellular transport**.



Observe and Discuss

Observe the diagram and discuss the process with your friends.



Internet my friend

1. Which type of circulation is present in amphibians and reptiles?
2. Enlist organisms without a proper transport system.



Use your brain power

What is the co-relationship between activeness of organism and complexity of transport system?

8.8 Circulation in animals :

In higher animals the circulation is carried out by special fluids - blood and lymph.

Blood vascular system :

Higher animals from Annelida to chordata have a special circulating fluid, the blood which is pumped to the tissues by the heart through the blood vessels.

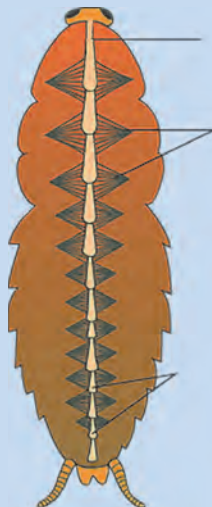
Types of blood vascular system :

1. Open circulation : In animals having an open circulation, blood is circulated through the body cavities (haemocoels). The visceral organs lie in the blood filled body cavity. Exchange of material takes place directly between blood and cells or tissues of the body. The blood flows with low pressure and usually does not contain any respiratory pigment like haemoglobin, so it does not transport respiratory gases. e.g. Arthropods (cockroach, studied in 11th std.) and Molluscs.



Observe and Discuss

Observe the diagram and give appropriate labels.



2. Closed circulation : In all the vertebrates, higher molluscs and annelids, blood is circulated all over the body through a network of blood vessels. In this type of circulation, blood flows within the blood vessels and does not come in direct contact with cells and body tissues. Exchange of material between blood and body tissues is through an intermediate fluid called lymph. Blood flows with high pressure and contains respiratory pigments like haemoglobin for transportation of respiratory gases.

The closed circulation can be divided into two main types : single and double circulation.

Single circulation : In single circulation, the blood passes through heart only once during each cycle as in fishes. Deoxygenated blood is pumped from heart towards gills, where it undergoes oxygenation. This oxygenated blood moves towards various body parts, gets deoxygenated and returns back to heart for next cycle.

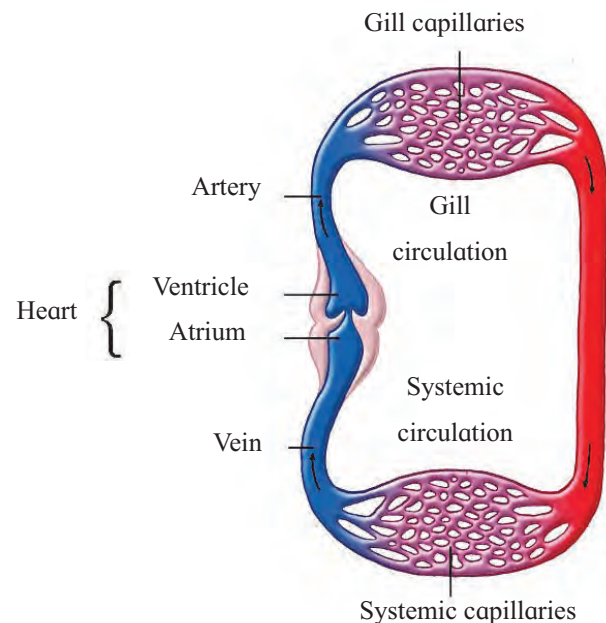


Fig. 8.12 : Single circulation

Since, the heart of fish carries only deoxygenated blood, it also called ‘**venous heart**’.

Double circulation : In double circulation, blood passes through heart twice during each cycle; it occurs in birds and mammals. In these animals, heart pumps deoxygenated blood to lungs for oxygenation and it returns to heart as oxygenated blood. This is ‘**pulmonary circulation**’. The oxygenated blood is pumped from the heart towards various body parts (except lungs) and returns back to the heart as deoxygenated blood. This is ‘**systemic circulation**’. Human heart shows double circulation.

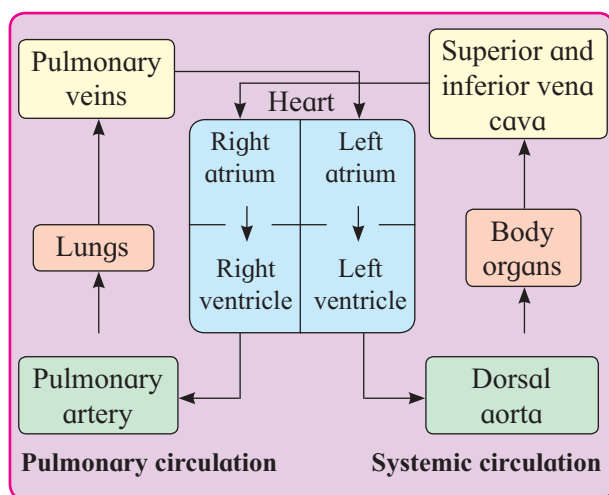


Fig. 8.13 : Double circulation (Diagrammatic)



Internet my friend

1. Find out the difference between coelom and haemocoel?
2. Name respiratory pigments present in the blood of different animals.

8.9 Circulatory System in Human :

The human circulatory system is composed of blood vascular and lymphatic system.

Blood vascular system :

In human beings it consists of blood, heart and blood vessels. It is responsible for various functions like transport, homeostasis and protection.

Blood Composition and Coagulation : Study of blood is called haematology. An average adult has about 4 to 6 liters of blood. It is a red coloured fluid connective tissue derived from embryonic mesoderm. It is slightly alkaline (pH 7.4), salty and viscous fluid. It is heavier than water. It has two main components- the fluid plasma (55%) and the formed elements i.e. blood cells (44%). These can be separated by centrifugation.

Plasma : It constitutes 55% of the blood. It is a straw-coloured, slightly alkaline, viscous fluid and consists of following:

Table 8.14 : Composition of plasma

Contents	Percentage
1. Water	90 %
2. Proteins (albumen, globulin, properdin, prothrombin, fibrinogen)	7 to 8 %
3. Inorganic salts (Na, K, Mg, Ca, Fe, Mn and Cl ⁻ , HCO ₃ ⁻ and PO ₄ ³⁻)	1 %
4. Others :	1 to 2 %
a. Food (glucose, amino acids, fatty acids, triglycerides)	
b. Wastes (urea, uric acid and creatinine)	
c. Regulators (hormones, enzymes, vitamins)	
d. Anticoagulants (heparin)	
e. Cholesterol and antibodies	
f. Dissolved gases (O ₂ , CO ₂ , N ₂)	



Internet my friend

1. Find out the percentage and functions of different blood proteins.
2. What is the clinical importance of haematokrit value?
3. Differentiate between plasma and serum.

Formed elements :

The blood cells that are produced in the body are collectively called formed elements. Human blood contains three types of formed elements as red blood corpuscles (RBCs), white blood corpuscles (WBCs) and platelets.

8.10 Red blood corpuscles / Erythrocytes :

Erythrocytes are the most abundant cells in the human body. They are circular, biconcave and enucleated (in camel and llama they are nucleated).

The red colour of RBCs is due to an oxygen carrying pigment, the haemoglobin, in their cytoplasm. In males, their average number is about 5.1-5.8 million/mm³ (per μ L) and in females about 4.3-5.2 million/mm³. This is called total RBC count. The average life span of RBCs is 120 days. The process of formation of RBCs is called **erythropoiesis**.

RBCs are produced from haemocytoblasts / reticulocytes. The erythropoietic organ of the foetus is the liver and spleen and in the adult, it is mainly the red bone marrow. Vitamin B₁₂, folic acid, iron and heme protein are required for production of RBCs. The old and worn out RBCs are destroyed in the liver and spleen (graveyard of RBCs). Condition with increase in the number of RBCs is called **polycythemia** and with decrease in number of RBCs is called as **erythrocytopenia**. The hormone erythropoietin produced by the kidney cells stimulates the bone marrow for production of RBCs.

Mature erythrocyte is devoid of nucleus, mitochondria or other membrane bound cell organelles. Its cytoplasm (stroma) is rich in haemoglobin and O₂ carrying proteinaceous pigment that gives red colour to the RBCs and blood. It also contains an enzyme, carbonic anhydrase.

Erythrocytes are responsible for the transport of respiratory gases O₂ and CO₂, maintaining pH and viscosity of blood. They also contribute in the process of blood clotting. The **hematocrit** is ratio of the volume of RBCs to total blood volume of blood. It is different for men and women.



Do you know ?

Each erythrocyte approximately contains 270 million molecules of haemoglobin. Normal content of haemoglobin in blood of men is about 14 – 17 gm% and in women it is about 13 – 15 gm%. Condition with less number of RBCs or less amount of haemoglobin or both is called as anaemia.

Each molecule of haemoglobin is a protein-iron complex. It consists of four polypeptide (globin) chains 2 alpha and 2 beta chains. An iron – porphyrin (haem) group is attached to each chain and all four chains are bound together. Each haem group can carry one O₂ molecule and thus one haemoglobin molecule can carry four O₂ forming oxyhaemoglobin. CO₂ interacts with amino acid residues of globin chains and forms carbaminohaemoglobin. After haemolysis, haemoglobin is broken down. Its globin part is broken to recycle the amino acids. Iron of heme group is stored as ferritin in the liver and porphyrin group of heme is converted into green pigment biliverdin and then into red-orange coloured bilirubin. These pigments (mainly bilirubin) are added to bile and finally removed out of body along with faeces.



Internet my friend

1. Compare between myoglobin and haemoglobin.
2. In the erythrocytes, nucleus and mitochondria are absent. Explain?

8.11 White blood corpuscles / Leucocytes:

Leucocytes are colourless, nucleated and amoeboid cells larger than RBCs. Due to their amoeboid movement they can move out of the capillary walls by a process called **diapedesis**. A normal adult has on an average, 5000-11000 WBCs per mm³ of blood. Decrease in number of

WBCs (<4000) is called **leucopenia** (common in HIV, AIDS and TB patients or those exposed to radiations, shock, etc). Temporary increase in number of WBCs is called as **leucocytosis**. It is due to infection. It also occurs during pregnancy and in newborn babies. Uncontrolled increase in number of WBCs is a type of blood cancer called **leukemia**. WBCs are mainly concerned with defense mechanism i.e. protection.

Types of WBCs :

These are colourless, irregular nucleated cells and show **polymorphism** (exist in variable forms). They can be classified into two main types such as granulocytes and agranulocytes.

A. Granulocytes :

These are WBCs with a **granular cytoplasm**, also called **Polymorpho nuclear leucocyte (PMN)** cells. They have lobulated nuclei in different shapes. Granulocytes are formed from myeloid stem cells and once formed, do not divide. Granulocytes constitute about **72%** of total WBCs. Granules are actually secretory vesicles which contain various secretions, enzymes, etc. Depending upon staining property of the granules, these granulocytes are classified into three types as neutrophils, basophils and acidophills.



Think about it

What are the granules in granulocytes?

a. Neutrophils : Granules are very fine, large in number, evenly distributed and stained with neutral stains (dyes). Neutrophils are about **70%** of total WBCs. These cells are spherical and nucleus is several lobed (2-7). These are able to perform amoeboid movements and phagocytosis. They are responsible for destroying pathogens by the process of **phagocytosis**. 'Pus' is mixture of dead neutrophils, damaged tissues and dead microbes.



Think about it

Why and when do the leucocytes perform diapedesis?

b. Basophils / Cyanophils : These cells have very few granules of large size, and stain with basic stains like methylene blue. Basophils are non-phagocytic, small, spherical cells and are about 0.5-1% of total WBCs.. Nucleus is twisted. They are present in infected and allergic conditions only. Basophils secrete heparin, histamine and serotonin.

c. Eosinophils / Acidophills : Acidophills contain lysosomal granules that are stained to red colour with acidic stains like eosin. Eosinophils are about 1 – 3 % of total WBCs. Nucleus is bilobed. They destroy antigen-antibody complex by phagocytosis. Their number increases in allergic condition and they show antihistaminic property. They are also responsible for detoxification as they produce antitoxins.

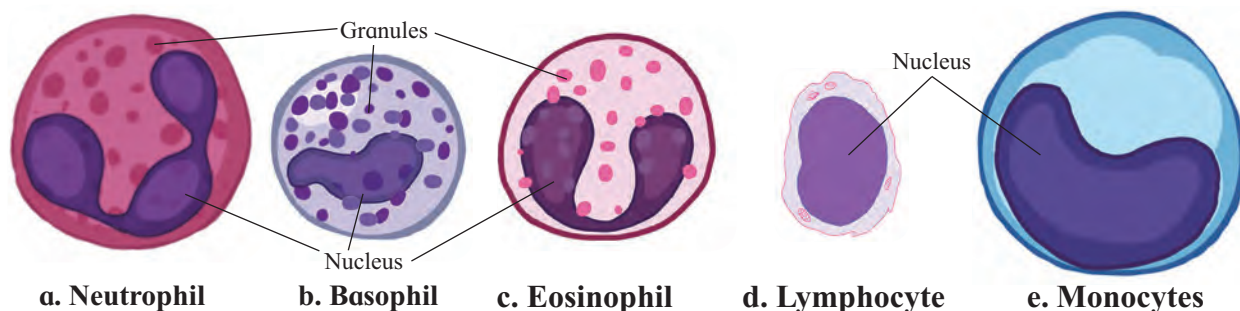


Fig. 8.15 : Granulocytes and Agranulocytes

B. Agranulocytes :

Agranulocytes are about 28% of total WBCs. Cytoplasm of these leucocytes is without granules. They are formed from lymphoid stem cells and can divide by mitosis. Nuclei of agranulocytes are large in size but are not lobulated like the granulocytes. There are two types of agranulocytes - Lymphocytes and Monocytes.

a. Lymphocytes : Lymphocytes are the smallest of all WBCs and have a large spherical nucleus. They constitute about 25-30% of total WBCs. Depending upon function, two types of lymphocytes are present as B-lymphocytes and T-lymphocytes. B-lymphocytes mature in bone marrow and are responsible for antibody production/humoral immunity. It is a highly specific antigen, antibody immunity T-lymphocytes mature in thymus and are responsible for cell-mediated immunity. Helper T-cells, killer T-cell, memory T-cells and suppressor T-cells are four main subtypes of T-lymphocytes.

b. Monocytes : Monocytes are the largest of all the WBCs. Its nucleus is large and bean or kidney shaped. They form 3-5% of WBCs. Monocytes are actively motile and give rise to macrophages. They are mainly phagocytic and destroy the bacteria and dead or damaged tissue by phagocytosis.



Internet my friend

1. What are the reasons for changes in number of neutrophils and the importance of complete blood count (CBC)?
2. Why and when are heparin, histamine and serotonin secreted? Are these biomolecules secreted by any other cell/organ in our body?

8.12 Thrombocytes / Platelets:

Thrombocytes are cellular fragments formed from the large cells called megakaryocytes. These are produced in bone marrow. They are very small, oval shaped cell fragments without nucleus. Normal count of thrombocytes in human blood is about 2.5 – 4.5 lakh / mm³ of blood. If number of thrombocytes decreases than normal, condition is called as **thrombocytopenia**. This condition causes internal bleeding (haemorrhage). Platelets secrete platelet factors which are essential in blood clotting. They also seal the ruptured blood vessels by formation of **platelet plug/thrombus**. They secrete **serotonin** a local vasoconstrictor.

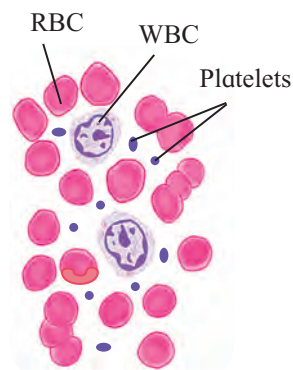
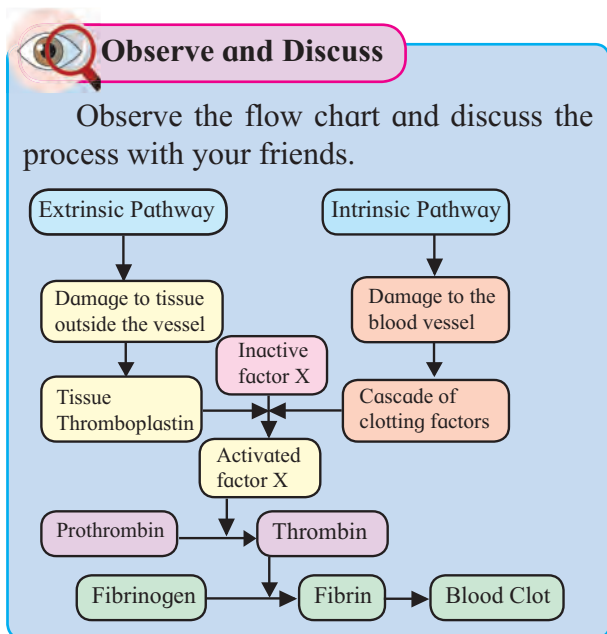


Fig. 8.16 : Blood smear

Blood Clotting/ Coagulation of blood :

Clotting or coagulation is the process of converting the liquid blood into a solid form. This process may be initiated by contact of blood with any foreign surface (intrinsic process) or with damaged tissue (extrinsic process). Intrinsic and extrinsic processes involve interaction of various substances called **clotting factors** by a step wise or cascade mechanism. There are in all twelve clotting factors numbered as I to XIII (factor VI is not in active use). Interaction of these factors in a cascade manner leads to formation of the enzyme thrombin. Thromboplastin, helps in the formation of enzyme prothrombinase. This enzyme inactivates heparin and it also converts inactive prothrombin into its active thrombin.

Thrombin converts soluble blood protein-fibrinogen into insoluble fibrin. Fibrin forms a mesh in which platelets and other blood cells are trapped to form the clot. Blood clotting occurs as shown in the following flowchart.



Curiosity

1. What is blood clotting? How and when does it occur?
2. What is immunity? Name its types.
3. Why does the platelet count decrease in dengue patient?
4. Why does our immune system fail against pathogens like *Trypanosoma* and *Plasmodium*?
5. What is the relation between immunity and organ transplantation?
6. How do monocytes perform amoeboid movement and phagocytosis?
7. How do monocytes modify into macrophages?

8.13 Heart :

Heart is the main pumping organ of the circulatory system. It is reddish brown in colour, hollow, muscular organ, roughly the size of one's fist. Its average weight is about 300gm in males and 250gm in females. It is conical in shape and lies in **mediastenum**- i.e. the space between two lungs. It is broader at upper end

(base) and conical at lower end (apex). Conical end is slightly tilted to left side and rests above the diaphragm.

Heart is enclosed in a membranous sac called **pericardium**. Pericardium is formed of two main layers - outer **fibrous** and inner **serous pericardium**.

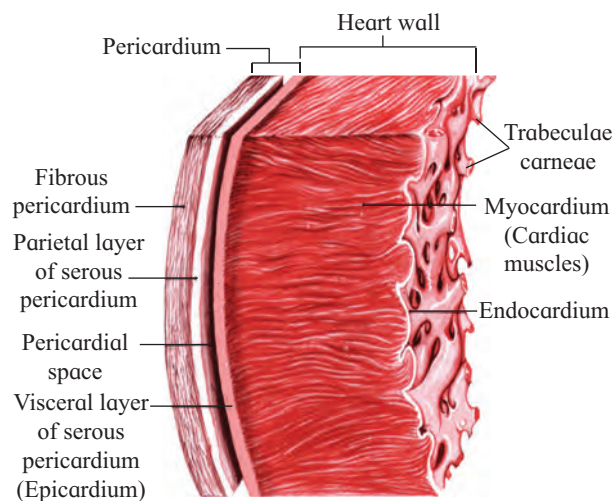


Fig. 8.17 : Heart wall and Pericardium

Serous pericardium is soft, moist and elastic. It is formed of squamous epithelium and is further divisible into two layers as **parietal** and **visceral layer**. Parietal and visceral layers of serous pericardium are separated by a pericardial space. This space is filled with **pericardial fluid** (about 50ml) which acts as a shock absorber and protects the heart from mechanical injuries. It also keeps the heart moist and acts as lubricant.

Heart wall :

The heart is mesodermal in origin. Its wall is formed of three layers, outer **epicardium**, middle **myocardium** and inner **endocardium**. Epicardium is thin and formed of a single layer of flat squamous epithelium resting on basement membrane. Myocardium is the middle thick layer formed of cardiac muscles. Endocardium is a single thin layer formed of squamous epithelium. The epicardium and endocardium are protective in function whereas myocardium is responsible for contraction and relaxation of heart.

External structure of heart :

The human heart is four chambered. The two superior chambers are called **atria** (auricles) and inferior two are called **ventricles**. Externally, the atria are separated from ventricles by a transverse groove called **coronary sulcus** or **atrioventricular groove**. The two ventricles are externally separated from each other by two grooves, the anterior and posterior **inter-ventricular sulci**. Coronary arteries and coronary veins run through these sulci. Pulmonary trunk arising from right ventricle and aorta from left ventricle are present on anterior surface of heart. The

pulmonary trunk bifurcates into right and left pulmonary arteries. Aorta (systemic aorta) is divisible into three regions as ascending aorta, systemic arch /aortic arch and descending aorta. The **Ligamentum arteriosum** joins pulmonary trunk and aortic arch. It is the remnant of an embryonic duct called **ductus arteriosus**. The aortic arch gives out three arteries viz. brachiocephalic (innominate) artery, left common carotid and left subclavian. The right atrium receives **superior and inferior vena cava** along its dorsal surface. **Pulmonary veins** open into left atrium along the dorsal surface of heart.

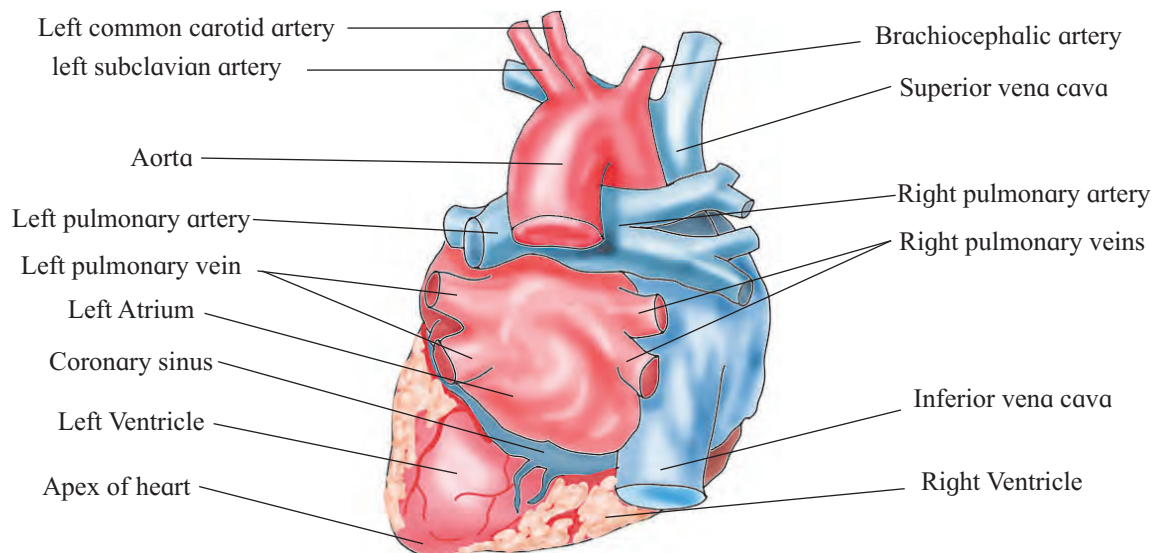


Fig. 8.18 : Posterior (dorsal) view : External structure of human heart

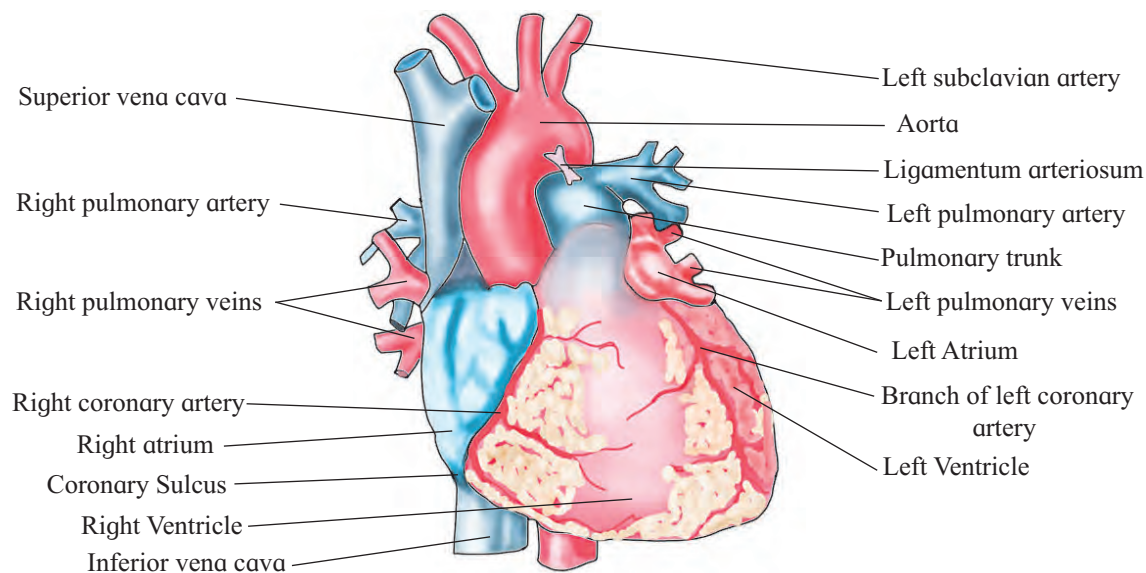


Fig. 8.19 : Anterior (ventral) view : External structure of human heart

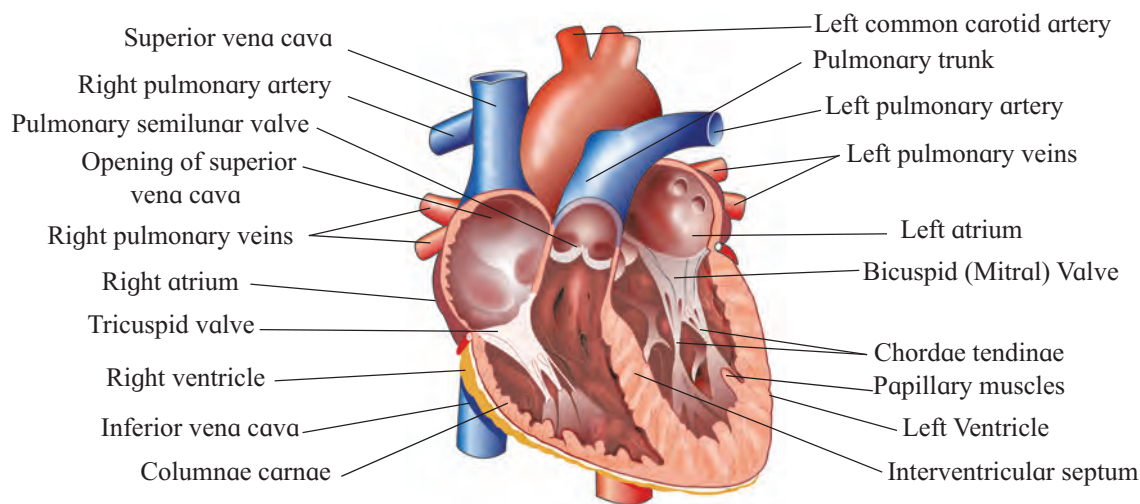


Fig. 8.20 : Internal structure of human heart

Internal structure of heart:

Atria : These are the thin-walled receiving chambers of heart. They are separated from each other by inter-auricular septum. Inter-auricular septum has an oval depression called **fossa ovalis**. It is a remnant of the embryonic aperture called **foramen ovalis**.

Superior vena cava (precaval), inferior vena cava (postcaval) and coronary sinus open into the right atrium. Opening of the postcaval is guarded by a **Eustachian valve** while the **Thebesian valve** guards the opening of coronary sinus into right atrium. Four pulmonary veins open into the left atrium. These openings are without valves.

Both the atria open into the ventricles of their respective sides by atrioventricular apertures. These openings are guarded by cuspid valves. The **tricuspid valve** is present in the right AV aperture and **bicuspid valve** (mitral valve) is present in the left AV aperture. All these heart valves help in maintaining a unidirectional flow of blood. They also avoid back flow of blood.

Ventricles : These are inferior, thick-walled pumping chambers of the heart. The right and left ventricles are separated by an inter-ventricular septum. Wall of the left ventricle is more muscular and about 3-times thicker than the right ventricle. Inner surface of the ventricles

shows several ridges called **columnae carnae** or **trabeculae carnae** which divide the lumen of ventricle into small pockets or fissures. The lumen of ventricles also shows inelastic fibers called **chordae tendinae**. These attach the bicuspid and tricuspid valves to the ventricular wall (papillary muscles) and regulate their opening and closing.

The right ventricle opens into the **pulmonary aorta** and left ventricle opens into the **aorta**. These openings are guarded by three semilunar valves each. These valves prevent the backward flow of blood into the ventricles.

Pumping action of heart :

The heart acts as the main pumping organ of the circulatory system. The pumping action is brought about by a rhythmic contraction and relaxation of the cardiac muscles or heart muscles. Contraction of heart muscles is **systole** and relaxation of heart muscles is **diastole**.

A single systole followed by diastole makes one **heart beat**. The heart beats 70 to 72 times per minute. This is called **heart rate**. During each heart beat ventricles pump about 70 ml of blood this is called **stroke volume**. It means heart pumps about $72 \text{ (heart rate)} \times 70 \text{ ml (stroke volume)} = 5040 \text{ ml} \approx 5 \text{ liters}$ of blood per minute this is called **cardiac output**.

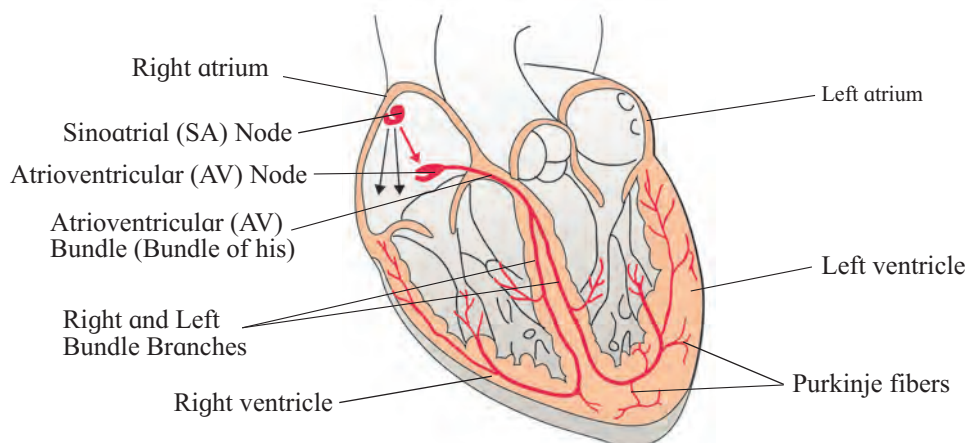


Fig. 8.21 : Conducting system of human heart

Conducting tissue of heart:

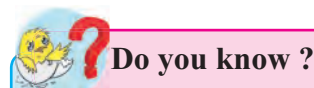
The human heart is **myogenic** i.e. the heart is capable of generating a cardiac contraction independent of nervous input. It also shows auto rhythmicity i.e. it can generate its own rhythm by specialized muscles. A specialized cardiac musculature called the nodal tissue is distributed in the heart. A part of this nodal tissue is present in the upper right corner of the right atrium. It is called **SA Node** or **Sinoatrial node**. It lies at the base of opening of superior vena cava. Another mass of nodal tissue, the modified muscular fibers also called autorhythmic fibers (conducting tissue) control the beating rate of heart. Conducting (nodal) tissue consists of SA node, AV node, bundle of His and Purkinje fibers.

Conducting system of the heart :

SA node (sinu-atrial node) is present in the right atrium. It acts as pacemaker of heart because it has the power of generating a new wave of contraction and making the pace of contraction. SA node passes the contraction to the left ventricle and also to the AV node. AV node (atrio-ventricular node) is present in the right atrial wall near the base of interatrial septum. It acts as pace setter of heart.

Bundle of His/ Tawara branches start from AV node and pass through interventricular septum. Bundle of His forms two branches, the right and left bundles, one for each ventricle.

These branches form network in ventricular walls and these are called Purkinje fibers. Bundle of His and Purkinje fibers spread impulses in ventricles. As a result both the ventricles contract simultaneously.



Do you know ?

Sometimes, valves are defective. Due to this, abnormal or adventitious sound is heard, called 'heart murmuring' or 'hissing.'

8.14 Working mechanism of human heart : Cardiac Cycle :

Human heart alternately contracts and relaxes. Contraction is called **systole** and relaxation is called **diastole**. Atria and ventricles contract alternately. Consecutive systole and diastole constitutes a single heartbeat or **cardiac cycle**. It is completed in **0.8 sec**. On an average, 72 beats are completed in one minute in an adult, at rest.

a. Atrial systole (AS):

Right atrium receives deoxygenated blood and left atrium receives oxygenated blood. When both the atria are completely filled with blood, pressure is exerted on the wall. In response to this pressure, SA node gets excited and generates cardiac impulse. Due to this, cardiac muscles in the atrial wall contract causing atrial systole. During atrial systole, blood is pumped into ventricles. Blood

is prevented from going back to the veins and coronary sinus by Eustachian and Thebesian valve respectively. After completing systole the atria go into diastole.

In normal conditions, atrial systole (AS) is for 0.1 sec. and atrial diastole (AD) is for 0.7 sec.

b. Ventricular systole (VS):

The impulse which started from SA node now reaches the AV node and it gets excited. AV node sends impulses to bundle of His and from bundle of His to Purkinje fibers. Purkinje fibers spread impulses all over the wall of ventricles. Due to this, ventricular wall contracts causing ventricular systole. During ventricular systole, right ventricle pumps deoxygenated blood into pulmonary trunk and left ventricle pumps oxygenated blood into aorta. During ventricular systole the cuspid valves close both the atrioventricular apertures preventing blood flow into atria (lubb sound is heard).

In normal conditions, ventricular systole lasts for 0.3 sec. and ventricular diastole (VD) lasts for 0.5 sec. During ventricular diastole, semilunar valves are closed, preventing backflow of blood from pulmonary trunk and systemic aorta into ventricles (dub sound is heard).

For about 0.4 second, both atria and ventricles are in diastole. When all the chambers of heart are in diastole, this condition is called **joint diastole** or **complete diastole**. Thus, duration of one cardiac cycle is 0.8 sec.

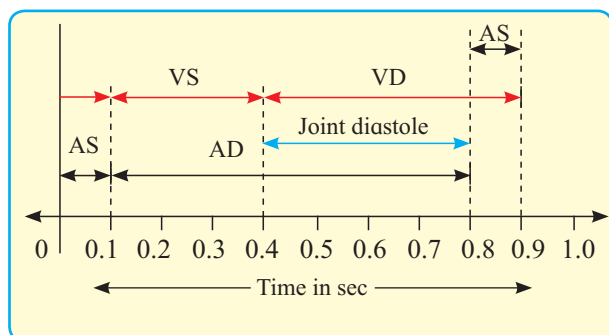


Fig. 8.22 : Diagrammatic representation of cardiac cycle

Right side of heart contains deoxygenated and left contains oxygenated blood. Total volume of blood pumped during one ventricular systole is called stroke volume (SV) and it is approximately 70 ml.

Cardiac output (CO):

It is the volume of blood pumped out per min. For a normal adult human being it is calculated as follows :

$$\begin{aligned} \text{(CO)} &= \text{SV} \times \text{HR} \\ &= 70 \times 72 = 5040 \text{ ml/min} \end{aligned}$$

Regulation of cardiac activity :

Though human heart is **myogenic**, it is also under dual control, the nervous as well as hormonal. The nervous control includes the part of autonomic nervous system. Sympathetic system (with hormone epinephrine as neurotransmitter) increase the rate of heartbeat during emergency. Parasympathetic system (with acetylcholine as neurotransmitter) reduces rate of heartbeat.

Nervous control includes the part of the autonomous nervous system- its cardiovascular center lies in the medulla oblongata. It controls rate of heart beat in response to inputs from various receptors like proprio-receptors (which monitor the position of limbs and muscles), chemoreceptors (monitoring chemical changes in blood) and baroreceptors (monitoring the stretching of main arteries and veins).

Chemical control of the heart rate includes the conditions like hypoxia, acidosis, alkalosis causing decreased cardiac activity, hormones like epinephrine and norepinephrine enhance the cardiac activity. Besides, concentration of cations like K^+ , Ca^{++} and Na^+ have major effect on cardiac activity. Cardiac activity decreases with the elevated blood level of K^+ and Na^+

8.15 Blood vessels :

There are three main types of blood vessels in the human circulatory system viz, arteries, veins and capillaries.

Arteries :

These blood vessels carry blood from heart to various parts/organs of the body, there they branch into arterioles and further into fine capillaries. They normally carry oxygenated blood to all parts of the body (except the pulmonary artery which carries deoxygenated blood). They are usually situated deep in the body except a few like the radial, brachial, femoral, etc. which are superficially located. In a **T. S. of artery**, its wall shows three layers.

1. Tunica externa or tunica adventitia
2. Tunica media
3. Tunica interna or intima

The outermost tunica externa is a thick, tough layer of collagen fibers. The tunica media is made up of smooth muscles and elastic fibres. This thick muscular and elastic layer makes the arterial wall pulsatile. The innermost tunica interna is a single layer of flat compact endothelial cells surrounding the lumen. The angular margin around the lumen shows **tesselations**. Arterial lumen is devoid of valves and blood flows through it rapidly and with high pressure.

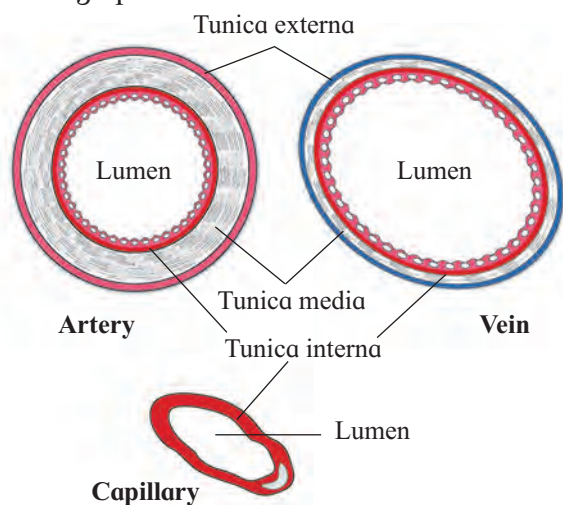


Fig. 8.23 : T. S. of Artery, Vein and Capillary

Veins :

Veins are thin walled, mostly superficial vessels which carry blood from the organs towards the heart. The capillaries around the various organs join to form the veins. Except for the pulmonary veins or other veins of the body carry deoxygenated blood towards the heart.

Portal vein : A portal vein e.g. hepatic portal vein, differs from the other normal veins in that its starts as capillaries from one organ and capillarises in some intermediate organ e.g. liver, before taking the blood towards the heart. Histologically, the veins also show the three layers like in the arteries. The tunica externa, tunica media and tunica interna. However, the tunica media is comparatively thinner and their lumen is wide and narrow. Internal valves at regular intervals can be seen. Blood flows with flow pressure and the valves prevent backflow of blood.

Capillary :

These are a network of minute blood vessels. They are thin walled having a single layer of flat squamous epithelium resting on a single basement membrane. They are mainly involved in exchange of materials. Wall of capillaries is formed of single layer of squamous epithelium and it is stretchable. Blood flows through the capillaries under high pressure. Wall of capillaries bear small endothelial pores or fenestrae through which blood cells (WBCs) can escape by the process called as diapedesis.

Pulse : It is a series of pressure waves that travel through the arteries due to ventricular systole. It is the strongest in arteries closer to the heart and gradually becomes weak in arteries away from heart. It can be felt easily in the superficial arteries like radial artery in the wrist and carotid artery in the neck. The pulse can be felt at particular points on the body. All locations where the pulse can be felt are shown in the figure 8.24.

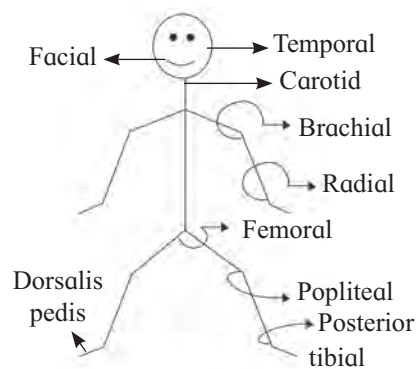


Fig. 8.24 : Pulse points

Pulse rate is equal to heart rate. Pulse rate higher than normal (above 100 beats/min) is called tachycardia and slower pulse rate (below 60 beats/min) than normal is called bradycardia.



Use your brain power

Why do the veins have valves?

8.16 Blood pressure (B. P.):

The pressure exerted by blood on the wall of the blood vessels is called blood pressure. It is measured by the sphygmomanometer. It is usually measured from the arteries.

Arterial Blood Pressure:

Pressure exerted by blood on the wall of artery is arterial blood pressure. Pressure on arterial wall during ventricular contraction (systole) is systolic pressure (SP). For a normal healthy adult the average value is 120 mmHg.

Pressure on arterial wall during relaxation of ventricles is diastolic pressure (DP). For a normal healthy adult it is 80 mmHg.

$$B P = SP / DP \\ = 120/80 \text{ mmHg}$$

Blood pressure is normally written as 120/80 mmHg. Difference between systolic and diastolic pressure is called **pulse pressure**. Normally, it is 40 mmHg.

Deviations from normal blood pressure value indicate malfunctioning of heart. It may be due to high or low blood volume, arterial inelasticity or hardening of arteries

(arteriosclerosis), deposition of fats like cholesterol in the arteries (atherosclerosis), renal diseases and emotion induced hormonal changes, obesity, etc. Blood pressure lower than normal i.e. below 90/60 mmHg is called hypotension and blood pressure higher than normal i.e. above 140/90 mmHg is hypertension.

Various factors that affect the blood pressure are cardiac output, peripheral resistance, blood volume, length and diameter of blood vessels, viscosity of blood, age, gender, venous return, sleep, emotions, exercise, anxiety, etc.



Think about it

Why are obese persons prone to hypertension?

Normal cardiac output is 5 lit/min. Increase in cardiac output increases systolic pressure. Peripheral resistance depends upon the diameter of blood vessels. Decrease in diameter of arterioles and capillaries under the effect of vasoconstrictors like vasopressin or ADH cause increase in peripheral resistance and thereby increase in blood pressure. Blood loss in accidents decreases blood volume and thus the blood pressure. Blood pressure is directly proportional to Viscosity of blood.

Blood pressure increases with age due to increase in inelasticity of blood vessels. Amount of blood brought to the heart via the veins per unit time is called the venous return and it is directly proportional to blood pressure. Blood pressure is also directly proportional to the total length of the blood vessel. Blood pressure can also be affected by vaso constriction or vaso dilation. Female (before menopause) has slightly lower BP than males of her age. However, the risk of high B. P. increases in the females after menopause sets in.



Internet my friend

Surf the internet for video-clips of angiography, angioplasty and by-pass surgery. Gather more information about these medical procedures.



Think about it

1. Why the heart-recipient has to rely upon life-time supply of immunosuppressants?
2. Why the transplanted heart beats at higher rate than normal?

Measurement of blood pressure:

Blood pressure is measured with the help of an instrument called **sphygmomanometer**. This instrument consists of inflatable rubber bag cuff covered by a cotton cloth. It is connected with the help of tubes to a mercury manometer on one side and a rubber bulb on the other side. During measurement, the person is asked to lie in a sleeping position. The instrument is placed at the level of heart and the cuff is tightly wrapped around upper arm. The cuff is inflated till the brachial artery is blocked due to external pressure. Then pressure in the cuff is slowly lowered till the first pulsatile sound is heard.

At this moment, pressure indicated in manometer is systolic pressure. Sounds heard during measurement of blood pressure are

called as *Korotkoff* sounds. Pressure in the cuff is further lowered till any pulsatile sound cannot be heard due to smooth blood flow. At this moment, pressure indicated in manometer is diastolic pressure. An optimal blood pressure (normal) level reads **120/80 mmHg**.

Hypertension :

Persistently raised blood pressure higher than the normal is called hypertension. 140/90 mmHg is called as threshold of hypertension and the 180/120 mmHg and higher readings are dangerous to the health. It may damage the heart, brain and kidneys.

Under the condition of hypertension, heart uses more energy for pumping which causes angina pectoris- the chest pains due to lowered blood supply to cardiac muscles and may lead to myocardial infarction. There are more chances of brain hemorrhage due to hypertension as arteries in brain are less protected by surrounding tissues as compared to other organs. In kidney, hypertension may cause kidney failure.

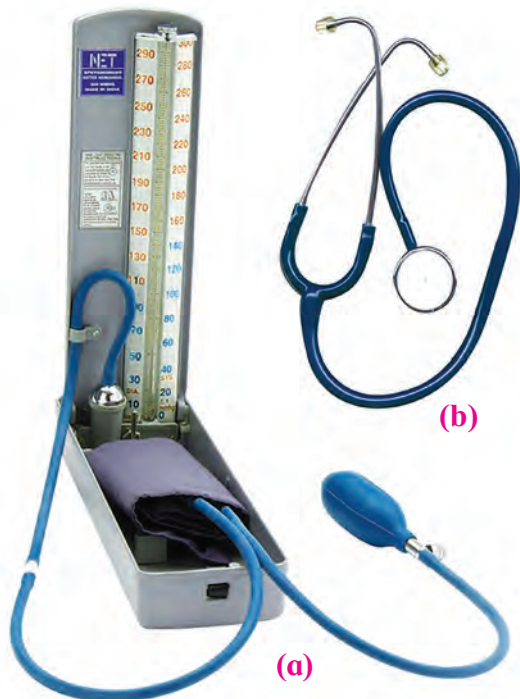
Coronary Artery Disease (CAD) :- It is also known as atherosclerosis. In this disease, calcium, fat cholesterol and fibrous tissues get deposited in blood vessels supplying blood to the heart muscles, making the lumen narrow.

Angina Pectoris :

It is the pain in the chest resulting from a reduction in the blood supply to the cardiac muscles because of atherosclerosis or arteriosclerosis. It is characterized by severe pain and heaviness in the chest. The pain may spread to the neck, lower jaw, left arm and left shoulder. The pain usually results from exertion, when there is more demand of oxygen by the heart, but the supply does not meet the requirement.

Angiography :

X-ray imaging of the cardiac blood vessels to locate the position of blockages



**Fig. 8.25 : (a) Sphygmomanometer
(b) Stethoscope**

is called angiography. Depending upon the degree of blockage, remedial procedures like angioplasty or by-pass surgery are performed. In angioplasty, a stent is inserted at the site of blockage to restore the blood supply while in by-pass surgery, the atherosclerotic region is by-passed with part of vein or artery taken from any other suitable part of the body, like hands or legs.

Heart Transplant :

Replacement of severely damaged heart by normal heart from brain-dead or recently dead donor is called heart transplant. Heart transplant is necessary in case of patients with end-stage heart failure and severe coronary arterial disease.

Silent Heart Attack :

Silent heart attack, also known as silent myocardial infarction is a type of heart attack that lacks the general symptoms of classic heart attack like extreme chest pain, hypertension, shortness of breath, sweating and dizziness. Symptoms of silent heart attack are so mild that a person often confuses it for regular discomfort and thereby ignores it. It has been studied that men are more affected by silent heart attack than women.



Know the Scientist :

Akash Manoj, a teenager from Chennai invented the non-invasive technique to predict the possibility of a silent heart attack. Interestingly, he invented this technique when he was in class-X.

For his innovation, he had been invited to the Rashtrapati Bhavan as a guest of the President of India under the Innovation Scholars In-Residence Programme.

His innovative kit analyses the level of FABP3 (Fatty Acid Binding Protein-3) with the help of UV light. It is the smallest protein in the blood.

Find out more information about.....

8.17 Electrocardiogram:

Graphical recording of electrical variations detected at the surface of body during their propagation through the wall of heart is **electrocardiogram (ECG)**. This recording may be in the form of printout or onscreen display. The instrument used for this recording is the **ECG machine** or **electrocardiograph**. This instrument detects and amplifies the signals.

Various electrodes are used for recording of signals. Four electrodes are positioned on limbs; two on arms and two on legs. These are limb electrodes. Six electrodes are positioned on chest. These are chest electrodes.

In a normal record, three different waves are recognized as P-wave, QRS complex and T-wave. **P-wave** is a small upward deflection from baseline of graph. It represents the atrial depolarization. The **QRS complex** starts as a slight downward deflection from baseline, continues as sharp and large upright wave and ends as a downward wave. QRS complex represents the ventricular depolarization. **T-wave** is small, wide and upwardly elevated wave. It represents the ventricular repolarization.

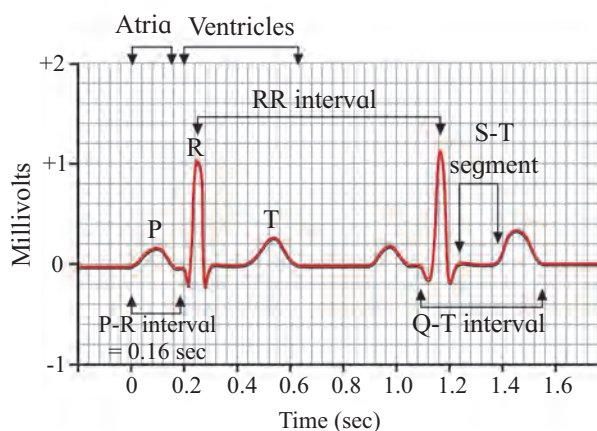


Fig. 8.26 : Normal ECG

ECG helps to diagnose the abnormality in conducting pathway, enlargement of heart chambers, damages to cardiac muscles, reduced blood supply to cardiac muscles and

causes of chest pain. A physician can find out the defect in the heart by examining the wave pattern and the time interval between them.

8.18 Lymphatic System :

Lymphatic system consists of lymph, lymphatic vessels, some organs and tissues. The word 'lymph' means 'clear water' and it is a fluid connective tissue with almost similar composition to the blood except RBCs, platelets and some proteins. Fluid from intercellular spaces of the body tissue enters into the lymphatic vessels, from here it is discharged into the blood vessels (veins) through the thoracic duct and the right lymphatic duct.

Curiosity

1. What is depolarization and repolarization?
2. What is the correlation between depolarization and repolarization as well as contraction and relaxation of the heart?
3. How are the signals detected and amplified by electrocardiograph?
4. Who discovered ECG?

Internet my friend

Location of lymph nodes in human body.

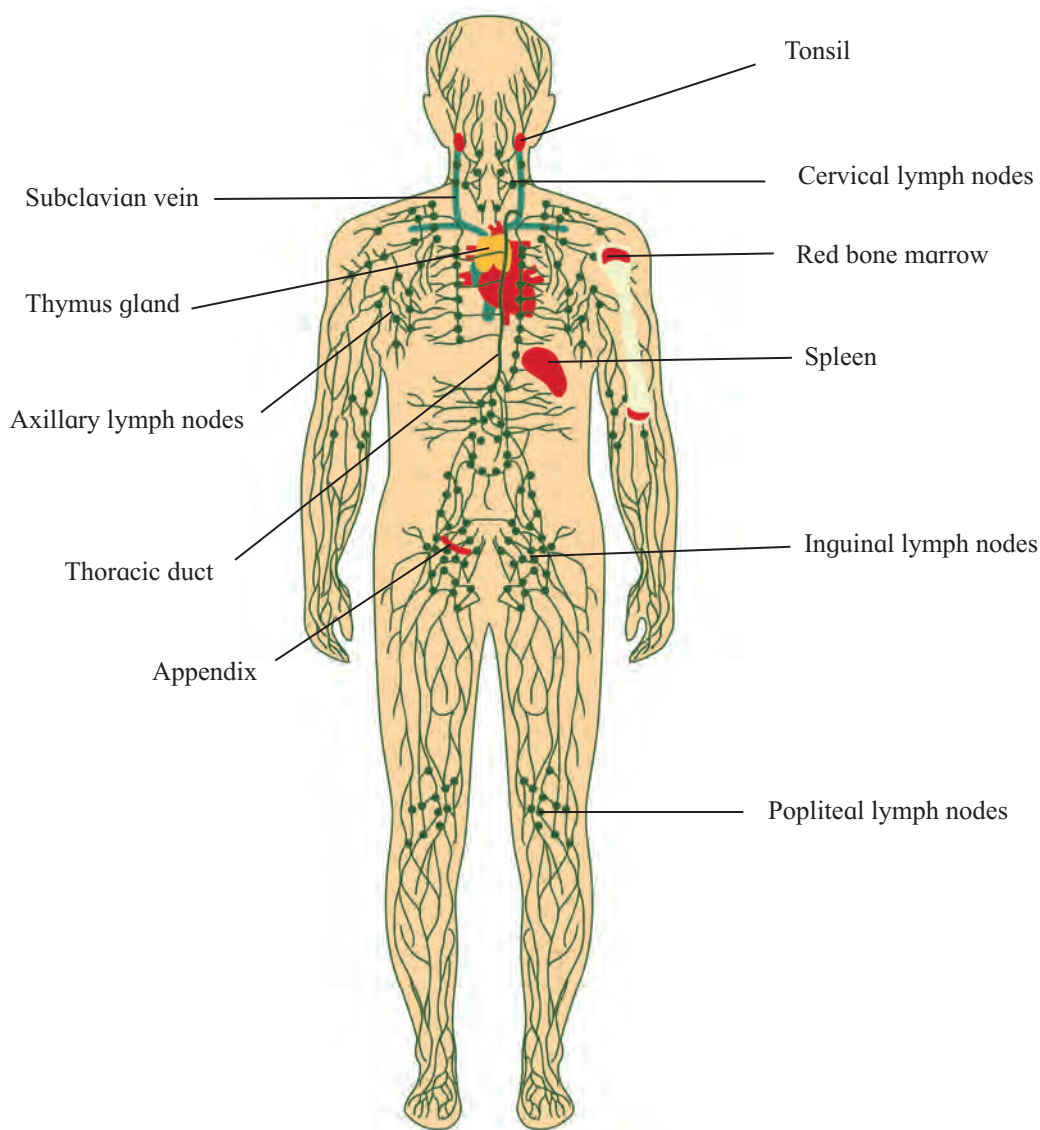


Fig. 8.27 : Lymphatic System



Do you know ?

Tonsils are small lymphatic nodules in pharyngeal region. Normally there are five tonsils strategically positioned to fight against inhaled and ingested foreign substances. Inflammation of tonsils is called as tonsillitis. It is caused due to viral or bacterial infection. Symptoms include sore throat, fever, swollen lymph nodes, nasal congestion, difficulty in swallowing, headache, etc. Viral tonsillitis cures naturally but bacterial tonsillitis needs antibiotic treatment. Tonsillectomy is performed in some patients who do not respond to the treatment.

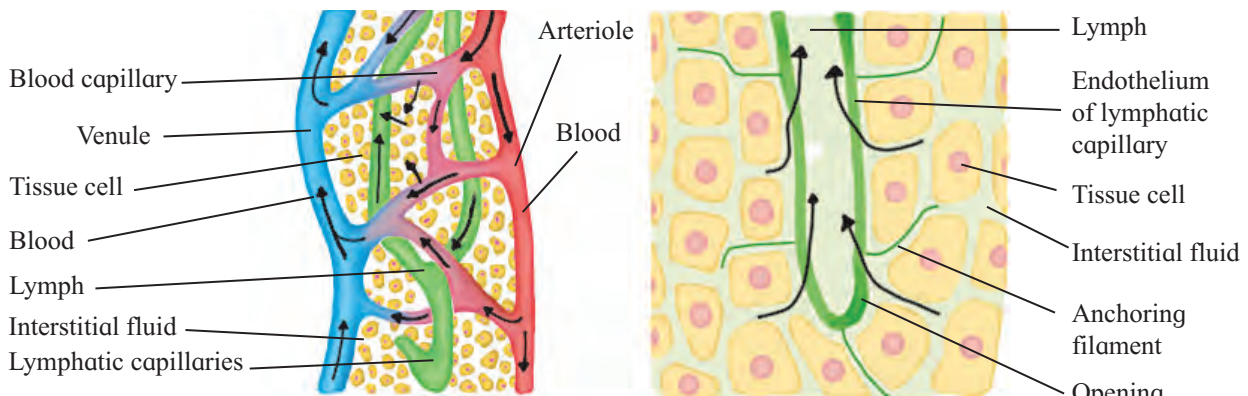
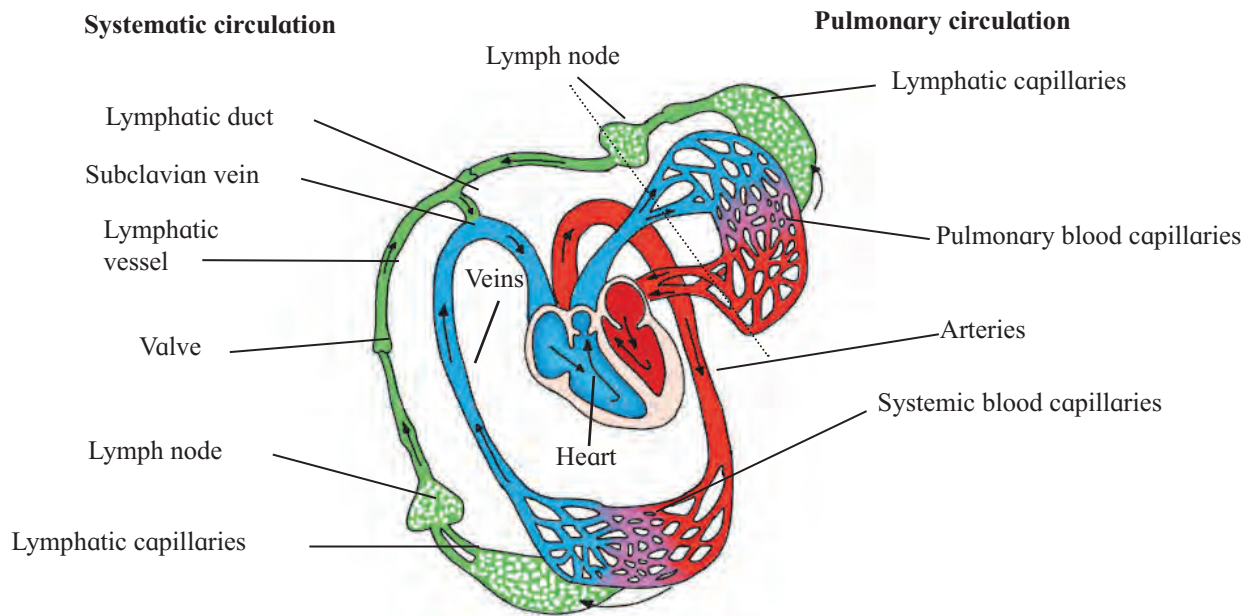


Fig. 8.28 : Circulation and Lymphatic System



Activity :

1. Prepare chart for different types of heart muscles and their functions.

2. Differentiate between the functioning of heart in frog and human.

Human Heart	Frog Heart

Exercise

Q.1 Choose the correct alternatives from those given below and complete the statements.

1. The muscular structure that separates the thoracic and abdominal cavity is _____.
a. pleura b. diaphragm
c. trachea d. epithelium
2. What is the minimum number of plasma membrane that oxygen has to diffuse across to pass from air in the alveolus to haemoglobin inside a R.B.C.?
a. Two b. Three c. Four d. Five
3. _____ is a sound producing organ,
a. Larynx b. Pharynx
c. Tonsils d. Trachea
4. The maximum volume of gas that is inhaled during breathing in addition to T.V is _____.
a. residual volume b. I.R.V.
c. G.R.V. d. vital capacity
5. _____ muscles contract when the external intercostal muscles contract
a. Internal abdominal
b. Jaw
c. Muscles in bronchial walls
d. Diaphragm
6. Movement of cytoplasm in unicellular organisms is called _____.
a. diffusion b. cyclosis
c. circulation. d. thrombosis.
7. Which of the following animals do not have closed circulation?
a. Earthworm. b. Rabbit
c. Butterfly d. Shark
8. Diapedesis is performed by _____.
a. erythrocytes b. thrombocytes
c. adipocytes d. leucocytes
9. Pacemaker of heart is _____.
a. SA node b. AV node
c. His bundle d. Purkinje fibers
10. Which of the following is without nucleus?
a. Red blood corpuscle b. Neutrophill
c. Basophill d. Lymphocyte
11. Cockroach shows which kind of circulatory system?
a. Open b. Closed
c. Lymphatic d. Double
12. Diapedesis can be seen in _____ cell.
a. RBC b. WBC
c. Platelet d. neuron
13. Opening of inferior vena cava is guarded by _____.
a. bicuspid valve
b. tricuspid valve
c. Eustachian valve
d. Thebesian valve
14. _____ wave in ECG represent atrial depolarization.
a. P b. QRS complex
c. Q d. T
15. The fluid seen in the intercellular spaces in Human is _____.
a. blood b. lymph
c. interstitial fluid d. water

Q. 2 Match the Respiratory surface to the organism in which it is found.

Respiratory surface	Organism
Plasma membrane	Insect
Lungs	Salamander
External gills	Bird
Internal Gills	Amoeba
Trachea	Fish

Q. 3 Very short answer questions.

1. Why does trachea have 'C' shaped rings of cartilage?
2. Why is respiration in insect called direct respiration?
3. Why is gas exchange very rapid at alveolar level.
4. Name the organ which prevents the following the entry of food into the trachea while eating.

Q 4. Short answer questions.

1. Why is it advantageous to breathe through the nose than through the mouth?
2. Identify the incorrect statement and correct it,
 - a. A respiratory surface area should have a large surface area.
 - b. A respiratory surface area should be kept dry.
 - c. A respiratory surface area should be thin, may be 1mm or less.
3. Given below are the characteristics of some modified respiratory movement. Identify them.
 - a. Spasmodic contraction of muscles of expiration and forceful expulsion of air through nose and mouth.
 - b. An inspiration followed by many short convulsive expiration accompanied by facial expression.
4. Write a note on blood plasma.
5. Explain blood clotting in short.
6. Describe pericardium.
7. Describe valves of human heart.
8. What is role of papillary muscles and chordae tendinae in human heart?
9. Explain in brief the factors affecting blood pressure.

Q. 5 Give scientific reason.

1. Closed circulation is more efficient than open circulation.
2. Human heart is called as myogenic and autorhythmic.
3. Person who has undergone heart transplant needs lifetime supply of immunosuppressants.
4. Arteries are thicker than veins.
5. Left ventricle is thick than all other chambers of heart.

Q. 6 Distinguish between :

1. Open and closed circulation.
2. Artery and vein.
3. Blood and lymph.
4. Blood capillary and lymph capillary.

5. Intrinsic and extrinsic process of clotting.

Q. 7 Long answer questions.

1. Smita was working in a garage with the doors closed and automobiles engine running. After some time she felt breathless and fainted. What would be the reason? How can she be treated?
2. Shreyas went to a garden on a wintry morning. When he came back, he found it difficult to breath and started wheezing. What could be the possible condition and how can he be treated?
3. Why can you feel a pulse when you keep a finger on the wrist or neck but not when you keep them on a vein?
4. A man's pulse rate is 68 and cardiac output is 5500 cm³. Find the stroke volume.
5. Which blood vessel of the heart will have the maximum content of Oxygen and why?
6. If the duration of the atrial systole is 0.1 sec and that of complete diastole is 0.4 sec, then how does one cardiac cycle complete in 0.8 sec?
7. How is blood kept moving in the large veins of the legs?
8. Describe histological structure of artery, vein and capillary.
9. What is blood pressure? How is it measured? Explain factors affecting blood pressure.
10. Describe human blood and give its functions.

Project :

- Visit pathological laboratory to study various blood tests like Hb detection, CBC, blood groups.
- Visit hospital to study how to take ECG, stress test, measurement of BP, etc.
- Evaluation of ECG on broad basis.
- Use of stethoscope.
- Differential count of WBCs.



Can you recall?

1. What is the need for control and co-ordination in multicellular animals ?
2. How do plants carry out control and co-ordination ?

Unicellular organisms have a simple organization of their life processes. However a multicellular organization of the body organs and organ systems required both, a control over their life processes as well as a coordination between the various systems in order to maintain homeostasis of the organism. Plants and animals both show a control and coordination mechanism. In plants, this is done by sending chemical signals and bringing about various types of movements (e.g. phototropic, chemotactic, thigmotactic, etc). Animals, specially the higher vertebrates show a gradual increase in the complexity of their control and coordination by giving both electrical and chemical signals. In this chapter, now you will study about development of nervous system in different animal groups and details of the system in humans.

The porifera (sponges) are the most primitive of the animal phyla.



Can you recall?

1. Do sponges have tissues and organs ?
2. Can sponges coordinate their various functions ?

Even though there are different types of cells in sponges for carrying out different functions, a proper nervous system is lacking. However to bring about efficient working of the body these multicellular animals show division of labour among the cells. This leads to specialization of cells for the various activities like digestion, respiration, excretion and others. Later in the higher animal phyla, the different

cells form tissues, organs and systems which must work in coordination with each other for smooth internal functioning of the body. Also the organism must be able to respond and coordinate with respect to various stimuli or changes in the external environment.

In the lower animals like *Hydra* and *Planaria* the nervous system achieves this function, while in higher and more complex animals, it is done by two coordinating systems - the **nervous (electrical) system** and **endocrine (chemical) control system**. These two systems will be studied separately though they work in coordination with each other.

NERVOUS COORDINATION

9.1 Nervous System in *Hydra* :

Hydra, a cnidarian shows the diffused nervous system, which is the most primitive nervous system. The cnidarians are thus the first animal group showing true simple nervous system. It consists of the sensory cells and the nerve cells or neurons along with their fibres. The nerve cells are scattered or distributed throughout the body and inter connected to each other by synapses between their fibres to form the nerve net. There are two nerve nets both in the mesoglea, one connected towards the epidermis and second towards the gastrodermis. There are sensory cells scattered in the

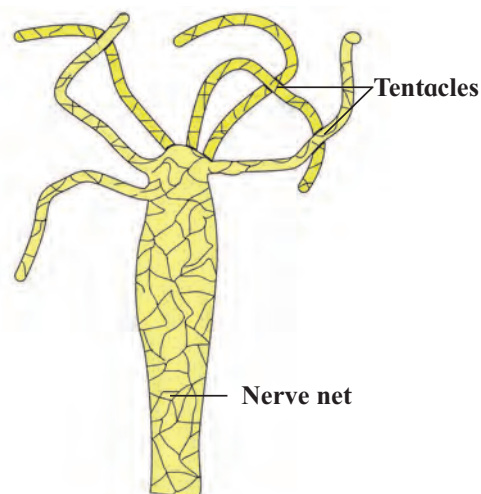


Fig. 9.1 : Nerve net in *Hydra*

body wall and tentacles, but sense organs are lacking. Neurons have fibres but there are no sensory and motor nerves. The nerve impulse shows no polarity or direction. Thus in *Hydra*, activation of sensory cells can happen at any point, and from this point impulse can be carried through out the body in any direction, thus bringing movements of the body or tentacles eg. catching of prey during feeding.

The diffused types of nervous system is the first important landmark in the nervous system. It is seen in the ctenophora as well as in the enteric system or gut wall of higher animals including man.

9.2 Nervous System in *Planaria* (flatworm) :

Planaria is a flatworm belonging to the phylum platyhelminthes. It is the most primitive animal with a central nervous system (CNS) located on the ventral side of body. It consists of a mass of cerebral or cephalic ganglion appearing like an inverted U shaped brain. These lie in the anterior or head region and from each ganglion arise nine branches towards the outer side. Ventrally from below the ganglia arise a pair of Ventral Nerve Cords (VNC) or long nerve cords. These are inter connected to each other by transfer nerve or commissure in a ladder like manner. The peripheral nerve plexus arising laterally from VNC. The PNS

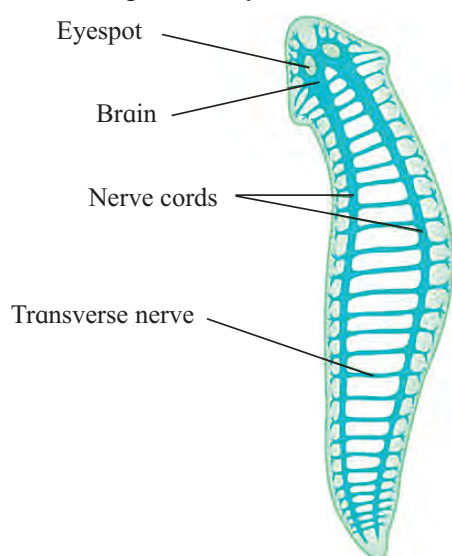


Fig 9.2 Ladder type nervous system in Flatworms e.g. *Planaria*

include sensory cells arranged in lateral cords in the body. A pair of photosensory structure, the eyes are located on dorsal side of the brain. Also there are single sensory cells scattered in the body.

In the above examples of *Hydra*, *Planaria* and the earlier studied examples of cockroach and humans, we have seen the gradual evolution or changes of the neural system. There is a high level of specialization in the formation of neurons as electrically signalling cells and also in the entire system, gradually from a diffuse neural system to a centralized nervous system. The expansion into a properly organized system involving the brain, its gradual expansion in size and functions. This has led to centralization of various sense organs assisting in coordinating the internal environment with that of the external environment. Also there is evolution of a complex networking system which efficiently transmits signals between one part or organ of the body and another.



Activity :

Note the changes taking place in the internal environment of the body, when a person goes from resting state into a state of physical activity.

9.3 Neural tissue

The neural tissue consists primarily of two types of cells viz. the neurons (nerve cells) and the neuroglia or glial cells. The word 'nerve' is used for a bundle of axons outside the CNS while inside the CNS for the same, the word 'tract' is used. The nerves may be sensory or motor or mix type i.e. having both the types - sensory and motor fibres.

All these along with nervous organs make up the nervous system of the higher animal and bring about coordination and control of various activities of the body. This is done through the receptors which bring in sensory inputs towards the central nervous system. Processing is carried out in the CNS and then through the

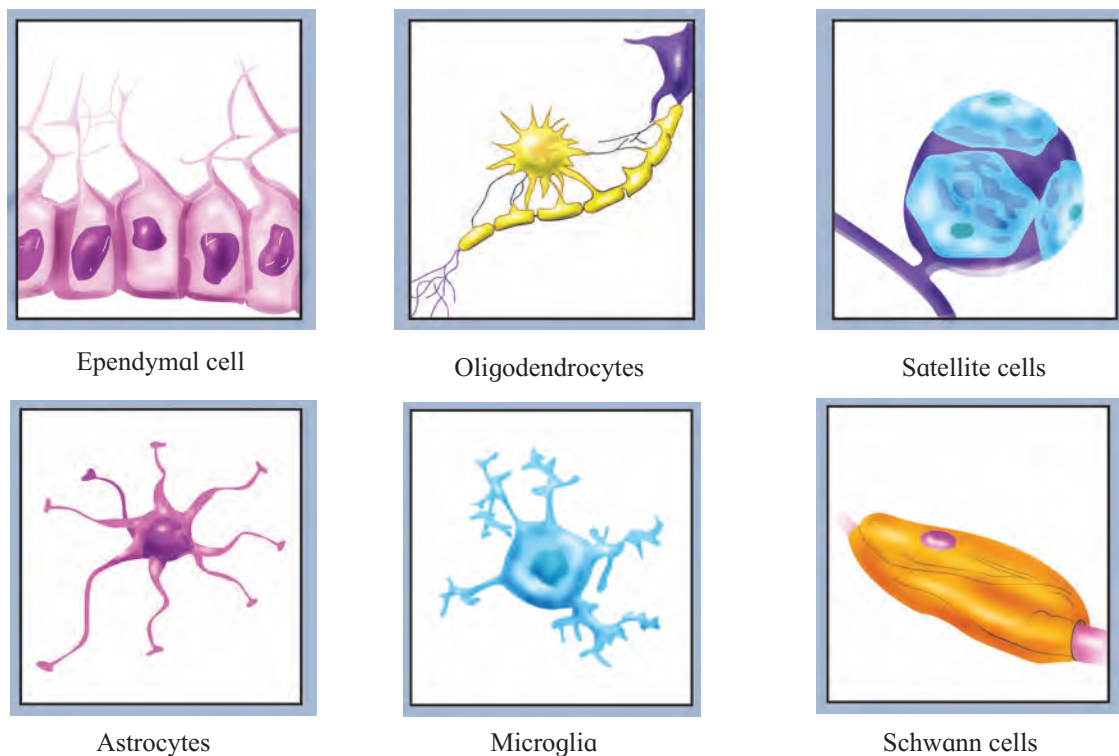


Fig. 9.3 : Types of Neuroglial cells

motor commands, the response is sent out. The nerves arising from the cytons of the CNS, travel throughout the body transmitting the nerves impulses to or from the CNS.

Curiosity Box:

Find out about the nervous system in Earthworm.



Internet my friend

1. How many neurons are present in the human body and specifically in the brain?
2. Normally what percentage of cranial capacity is used by an average human ?
3. What is the ratio between neurons and neuroglia ?

Neurons/Nerve cells : These are the structural and functional units of the nervous system. (You may recall the structure of a neuron studied in earlier classes). Each multipolar neuron has three parts - cyton or cell body, dendron and axon. The **cyton** has a distinct central nucleus with a nucleolus

and neuroplasm. A clear film of cytoplasm surrounds the nucleus around which there are neurofibrils, Nissl's granules and other cell organelles. Nissl's granules are riboprotein components. They play an important role in the synthesis of the enzyme required for formation of the neurotransmitter. Neurofibrils play an important role in transmission of nerve impulse. **Dendrons** are many small conical processes arising from the cyton. These are highly branched into fine dendrites. Nissl's granules and neurofibrils both can be seen at the base of the dendrons. They transmit message towards the cyton. **Axon** is a single long, usually unbranched process arising from the cyton at the axon hillock. It consists of a bundle of neurofibrils. Nissl's granules are absent. Terminally, the axon gives out branches called telodendrons. The axons carry the messages away from the cytons. The axons may give out lateral branches called collaterals. The terminal branches attach to a muscle, gland, skin or telodendrites of another neuron. The interconnection between two neurons or neuron with motor organ is called **synapse**. It is

Table 9.4 : Types of Neuroglial cells and their functions

Location	Cell type	Functions
CNS (Central Nervous System)	Oligodendrocytes	These cells have few branches and mainly form the myelin sheath around the central axons, which form the white matter of CNS. Myelin an insulating sheath is made up of protein and fatty substances. It allows quick transmission of electrical impulses.
	Microglia or brain macrophages	Small sized cells with few branches. These are derived from monocytes and act as macrophages. They go to the site of injury, dead neurons and cell debris in the CNS. They mediate immune response in the CNS.
	Astrocytes	Star shaped cells and the most abundant glial cells of CNS. They have varied roles in the brain, secretion and absorption of neural transmitter and maintenance of blood-brain barrier BBB. Regulate the transmission of electrical impulses with the brain.
	Ependymal cells	Form single layer of squamous or columnar, often ciliated epithelial cells lining the ventricles or brain cavities and central canal of spinal cord. Mainly responsible for production and probably also for circulation of CSF in brain ventricles and central canal.
PNS (Peripheral Nervous System)	Schwann Cells	These are the most abundant glial cells of PNS. They produce myelin sheath around medullated nerves of PNS.
	Satellite cells	They support the functions of neurons.

usually axo-dendronic or may be axo-axonic, axo-somatic or dendro-dendronic.

The cytons are generally found inside the brain, spinal cord (CNS) and in the ganglia. Small groups of cell bodies inside the white matter of brain are called **basal nuclei**. A bundle of axons called nerve may be covered only by **neurilemma** in the **non-medullated** nerves while in the medullated nerves it is covered both by medullary sheath and on the outside by neurilemma. Conduction of impulse by the medullated nerves is 50 times faster than in the non-medullated nerves. The connective tissue covering around the nerve fascicule is called **endoneurium**. Few nerve fasciculi with

endonurium are surrounded by connective tissue, called **perineurium** and a still large bundle of nerves is covered on the outer side by **epineurium**. Blood is supplied to all the nerves to provide oxygen and nutrients.

Neuroglial cells :

The neuroglial cells are far greater in number than the neurons. Most of the supporting cells of the nervous system are derived from the same embryonic tissue layer (ectoderm) that produces neurons. The term neuroglia refers to the supporting cells of the Central Nervous System (CNS) and Peripheral Nervous System (PNS).

9.4 Synapse :

It is a junction between two nerve cells with a minute gap (synaptic cleft) in between them which allows transmission of impulse by a neurotransmitter bridge.

1. Properties of nerve fibres :

a. Excitability/Irritability - The nerve fibres, on account of presence of a polarised membrane, have the ability to perceive stimulus and enter into a state of activity.

b. Conductivity - It is ability to transmit the excitation.

c. Stimulus - It is any detectable, physical, chemical, electrical change in the external or internal environment which brings about excitation in a nerve/muscle/organ/organism. In order to be effective, the stimulus must have a minimum intensity called **threshold stimulus**. **Subliminal** (weak) stimulus will have no effect while **supraliminal** (strong) stimulus will produce the same degree of impulse as the threshold stimulus.

d. Summation effect - A single subliminal stimulus will have no effect but if many such weak stimuli are given in quick succession,

they may produce an impulse due to addition or summation of stimuli.

e. All or none law - The nerve will either conduct the impulse along its entire length or will not at all conduct the impulse, as in case of subliminal or weak stimulus.

f. Refractory period - It is the time interval (about millisecond) during which a nerve fails to respond to a second stimulus however strong it is.

g. Synaptic delay - The impulse requires about 0.3 to 0.5 milliseconds to cross a synapse. This time is required for release of neurotransmitter from the axon terminal and excitation in the dendron of the next neuron.

h. Synaptic fatigue - The transmission of nerve impulse across the synapse halts temporarily due to exhaustion of its neurotransmitter.

i. Velocity - The rate of transmission of impulse is higher in long and thick nerves. It is higher in homeotherms than in poikilotherms. The velocity of transmission is higher in voluntary fibres (100 - 120 m/second in man) as opposed to autonomic or involuntary nerves (10-20 m/second). Similarly it is faster in medullated

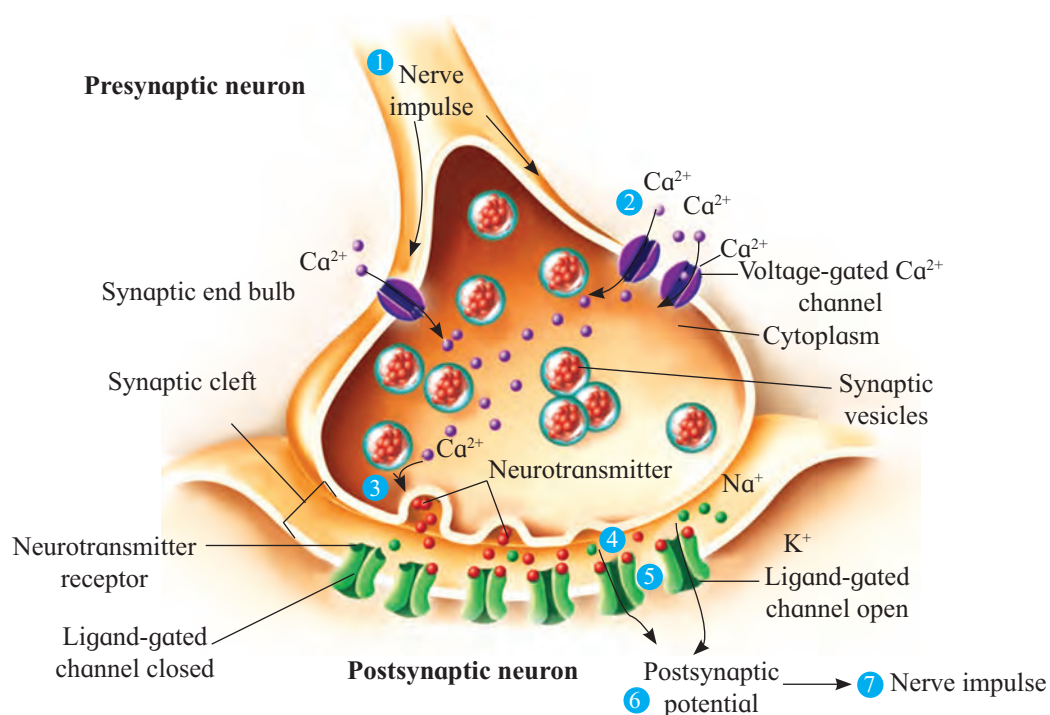


Fig. 9.5 : Synaptic transmission

nerve, as the impulse has to jump from one node of Ranvier to the next. At the synapse where the neurons communicate with one another. The neuron carrying an impulse to the synapse is the pre-synaptic neuron. The neuron receiving input at the synapse is the post synaptic neuron or generator region (gland or muscle). A synaptic cleft or a small intercellular space lies in between two cells having a width about 20-30 nm between them. .

The process by which the impulse from the pre-synaptic neuron is conducted to the post-synaptic neuron or cell is called **synaptic transmission**. It is a one way process carried out by neurotransmission.

2. Types of synapses

a. Electrical synapse : In this type of synapse gap between the neighbouring neurons is very narrow. The synapse between such closed neurons is mechanical. The electrical conductive link is formed between the pre - and post synaptic neurons. At the gap junction, the two cells are within almost 3.8 nm distance of each other. Transmission across the gap is faster but depends on the connection located at the gap junctions between the two neurons. Electrical synapses are found in those places of the body requiring fastest response as in the defence reflexes. They are bidirectional allowing transmission in either direction or may be unidirectional.

b. Chemical synapse : These are specialized junctions through which cells of the neural system send chemical signal to the other neurons and to non-neuronal cells, such as gland and muscle. Synaptic gap is larger than that in electrical synapse and it is 20-40 nm.

A chemical synapse between a motor neuron and a muscle cell is called a neuromuscular junction. There are three components of a typical chemical synapse. 1. The presynaptic terminal (mostly axonic terminal), 2. The synaptic membrane of the post synaptic cell (usually on the dendrite of the next neuron/

gland cell/ muscle) and 3. The post synaptic neuron.

The impulse travels along the axon of the pre-synaptic neuron to the axon terminal. Most presynaptic neurons or axons have several synaptic knobs at their ends or terminals. These knobs have arrays of membranous sacs, called synaptic vesicles, that contain neurotransmitter molecules.

When an impulse reaches a synaptic knob, voltage sensitive Ca^{++} channels open and calcium (Ca^{++}) diffuses inward from the extracellular fluid.

The increased calcium concentration inside the cells, initiates a series of events that fuse the synaptic vesicles with the cell membrane of presynaptic neuron, where they release their neurotransmitters by **exocytosis**. Once the neurotransmitters bind to the receptors of the post-synaptic cell, the action is either excitatory (turning a process on) or inhibitory (turning a process off). This is dependent on the nature of the neurotransmitter involved.

Once the impulse has been transferred across the synapse, the enzyme like **cholinesterase** destroys the neurotransmitter and the synapse is ready to receive a new impulse.

9.5 Transmission of nerve impulse :

The neurons are cells with some special features. The cells are excitable. The nerve impulse is a wave of bioelectrical or electrochemical disturbances passing along a neuron. The transmission of the nerve impulse along the long nerve fibre/axon tube is a result of electrical charges across the neuronal membrane during conduction of an excitation. Each neuron has a charged cellular membrane with a voltage which is different on the outer and inner side of the membrane. The plasma membrane separates the outer and inner solutions of different chemical compounds but having approximately the same total number of ions. The external tissue fluid has both Na^+

and K^+ but there is predominance of Na^+ and Cl^- , while K^+ is predominant within the fibre or in the intracellular fluid. This condition of a resting nerve is also called **polarised state** and it is established by maintaining an excess of Na^+ on the outside. On the inside there is an excess of K^+ along with large negatively charged protein molecules and nucleic acid. Some amount of Na^+ and K^+ always leaks across the membrane. The Na^+/K^+ pump in the membrane **actively** restores the ions to their appropriate side. Against the concentration and electrochemical gradient, Na^+ is being forced out and K^+ is being forced inside the membrane. This process is called **sodium pump** or **Na-K exchange pump**. This active process requires ATP energy. The difference in distribution of Na^+ and K^+ on the two sides of the membrane produces a potential difference of -50 to -100 millivolts (average is -70 millivolts).

This potential difference seen in a resting nerve is thus called resting potential. (-70 millivolts) and it is mainly due to differential permeability of the resting membrane which is much more permeable to K^+ than to Na^+ . This results in slightly more K^+ diffusing out than Na^+ moving inside and causing slight difference in polarity. Also ions like negatively charged proteins and nucleic acids inside the cell make the overall charge negative on the inside and positive charge on the outside.

The nerve membrane not only has leakage channels but also has many gated channels for Na^+/K^+ . These are also called voltage gated channels. These channels enable the neuron to change its membrane potential to active potential in response to a stimulus. The Na^+/K^+ gated channels are separate so transport of both these ions is separately done. However during resting potential, both these gates are closed and the membrane resting potential is maintained.

Generation of nerve impulse :

1. Depolarization : The origin and maintenance of resting potential depends on the original perfect state. Any change or disturbance to the membrane will cause Na^+ to enter into the membrane and lower the potential difference (lesser than -70 millivolts). This makes the membrane more permeable to Na^+ , so there will be rapid influx of Na^+ . This property is peculiar to a nerve membrane.

The voltage gated Na^+/K^+ channels are special in 2 ways : They can change the potential difference of the membrane as per the stimulus received and also the gates operate separately and are self closing.

During resting potential, both gates are closed and resting potential is maintained. However during depolarization the Na^+ gates open but not the K^+ gates. This causes Na^+ to rush into the axon and bring about a depolarisation (opposite of polarity). Extra cellular fluid (ECF) becomes electronegative with respect to the inner membrane which becomes electropositive. The value of action potential is $+30$ millivolts to $+60$ millivolts. This triggers depolarisation in the next part while it itself starts going to repolarisation.

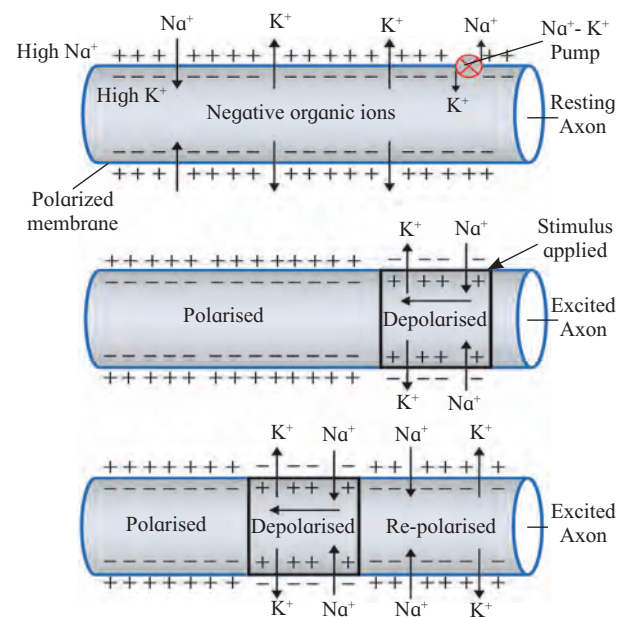


Fig. 9.6 : Polarisation and Depolarisation along a nerve

2. Re-polarization : Change in the polarity from depolarized, back to the original state is done by the process of repolarization. It occurs after a short interval called **refractory period**. The large number of Na^+ on the inside causes a drop in the permeability of membrane to Na^+ and at the same time making it more permeable to K^+ ions by opening the K^+ voltage gates and slowly closing the Na^+ gates. This action is a localized activity. K^+ ions pass out very rapidly as compared to slow entry of Na^+ .

In this period, Na^+ gates are closed, K^+ gates are open and $\text{Na}^+ - \text{K}^+$ pumps become operational. This process of producing a wave of stimulation \rightarrow causing depolarization \rightarrow repolarization is repeated continuously up to the end of axon terminal. It is a self-propagating process.

In myelinated nerves, the insulating fatty myelin sheath prevents flow of ions between the axoplasm and ECF. The transport pump and gated channels can operate only in the region of nodes of Ranvier, where myelin sheath is absent.

The action potential cannot travel as a wave of membrane depolarization; it has to jump from node to node. This process called **saltatory conduction**, is at the rate of 120 m/second. It is faster than the continuous conduction in non-medulated fibre (50 :1).



Do you know ?

$\text{Na}^+ - \text{K}^+$ pump operates actively and by use of carrier (for every 3 Na^+ pumped out 2 K^+ are pumped in). It is electrogenic.



Always Remember

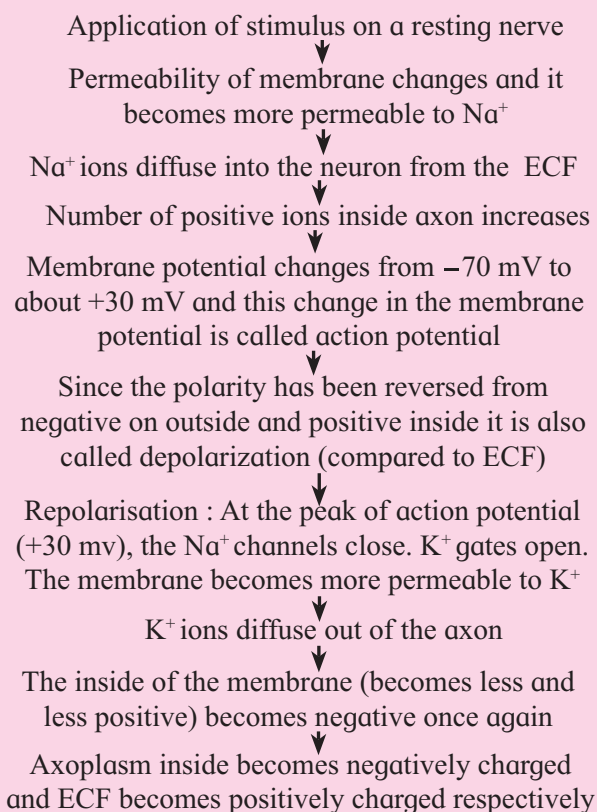
The resting potential of axon is -70 mV. $\text{Na}^+ - \text{K}^+$ Pump pumps out 3 Na^+ ions for every 2 K^+ ions they pump into the cell.



Can you tell?

1. Explain how is impulse transmitted through a synapse?
2. Describe the conduction of a nerve impulse in the neuron.

Chart 9.7 : Steps in generation and conduction of nerve impulse



9.6 Human Nervous System :

The nervous system in humans is well developed and complex. It is broadly classified into three parts, viz, CNS, PNS and ANS.

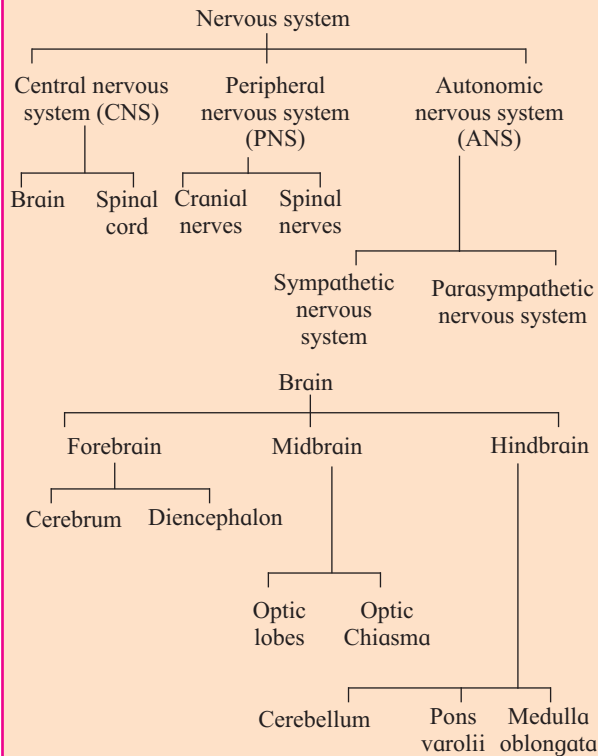
1. Central nervous system (CNS) : The brain and spinal cord are the parts of CNS which lie along the mid dorsal axis. Brain is enclosed within the brain box/**cranium** of the skull, whereas the spinal cord occupies the vertebral canal of the **vertebral column**. Inner to these bony coverings, are three protective membranes called **meninges**. That protect the brain and spinal cord.

a. Dura mater : It is the outermost tough, non vascular, thick and fibrous meninx and is attached to the inner side of the cranium. It is separated from the underlying arachnoid mater by the subdural space, filled with a serous fluid.

b. Arachnoid mater : It is the middle, thin and non vascular layer of connective tissue having web like appearance. It is separated from the



Can you recall?



pia mater below by a narrow subarachnoid space filled with cerebro spinal fluid - CSF.

c. Pia mater : It is the innermost delicate, highly vascular membrane lies in close contact with the CNS.

CSF (cerebrospinal fluid) : It is lymph like extra cellular fluid secreted by choroid plexuses of pia mater present inner to subarachnoid space in the ventricles of the brain within the central canal of spinal cord. The CSF is secreted by the pia mater, the choroid plexuses and the ependymal cells lining the ventricles of the brain and central canal of spinal cord. The three openings in the roof of **medulla oblongata** help in draining out the CSF from brain to the outside. The meninges and CSF act as a shock absorber and protect the brain and spinal cord from mechanical injuries. It also maintains constant pressure inside cranium. It helps in exchange of nutrients and wastes between blood and brain tissue. It helps in the

supply of oxygen to the brain. It protects the brain from desiccation.

CSF is slightly alkaline fluid with a specific gravity of 1.005. A total of 100 -120 cc of CSF is present in and around the CNS.

Curiosity Box:

CSF is released by the pia mater. Inside the ventricles/cavities of the brain, there is release of CSF by the choroid plexuses and the ependymal cells.

Does this CSF remain enclosed inside the ventricles ?

What can be the outcome of such a situation ?



Do you know ?

Blood brain barrier (BBB) -

Keeps a check on passage of ions and large molecules from the blood to the brain tissue. Endothelial cells lining the blood capillaries help in this process along with the astrocytes.



Always Remember

CSF is continuously generated by the ependymal cells lining the ventricles and central canal and simultaneously drained out of the brain into the blood stream. CSF maintains a constant pressure inside the cranium. The nervous tissue is without lymphatic vessels.



Think about it

During extraction of a tooth, the dentist gives an injection of Anaesthesia to the patient before extraction. Is the action potential generated ? How does the local anaesthesia work? What is the effect of pain killer on the nervous system.



Can you recall?

Label the various parts of the diagram and recall their functions.



A. The Human brain :

The study of all aspects of the brain is called encephalology. The brain can be divided into three main parts - forebrain, midbrain and hindbrain.

a. Forebrain : Forebrain consists of olfactory lobes, cerebrum and diencephalon.

i. Olfactory lobes :

These are highly reduced in human brain and covered by cerebrum from all sides except ventral. Each lobe consists of a olfactory peduncle and olfactory bulb.

ii. Cerebrum :

It is a largest part of the brain, making up about 85 % of total brain. It is divided into right and left cerebral hemisphere by means of a deep median, long fissure. The two hemispheres internally connected to each other by a thick

band of nerve fibres called **corpus callosum**.

The outer surface of cerebrum is called **cerebral cortex** while the deep inner part is **cerebral medulla**. The cerebral cortex has outer thin region composed of grey matter and inner medulla composed of white matter.

The surface of each cerebral hemisphere is greatly folded by many convolutions or **gyri** and grooves called **sulci**. These greatly increase total surface area for accomodation of the vast number of nerve cells.

Each cerebral hemisphere is further divided into four main lobes by three deep sulci. These are -

Centre sulcus which demarcates frontal lobe from the parietal lobe.

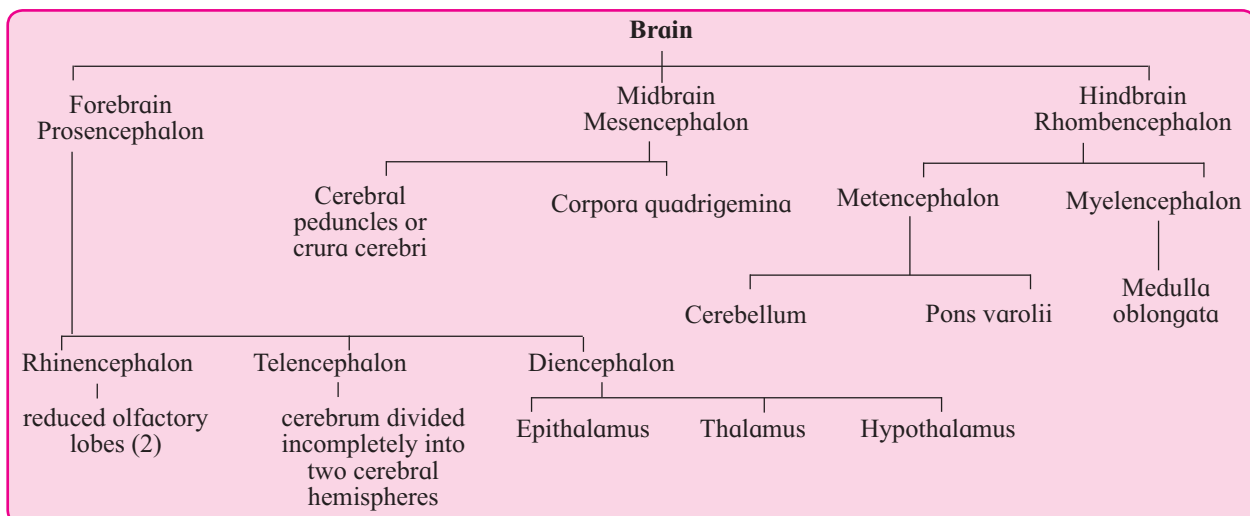
Parieto-occipital sulcus separates the parietal from occipital lobe.

The lateral or **sylvian sulcus** demarcates the temporal lobe from the frontal and parietal lobes

Since these three sulci are not complete, the lobes are not clearly demarcated from each other. A fifth median lobe called **insula** or **insular cortex** is folded deep within the lateral sulcus.

The grey matter of cerebral cortex mainly consists of cell bodies of billions of neurons along with non-medulated fibres and dendrons. The white matter mainly has axons of myelinated nerves.

Chart 9.8 Parts of Brain





Do you know ?

Corpus callosum is typically seen in mammalian brain.

Corpus callosum is the largest commissure of the brain.

It has an anterior and posterior fold called **genu** and **splenium** respectively.

Why an injury to the right cerebrum affects the functioning of the left side of body?

Functional areas of cerebrum :

1. Frontal lobes : They have motor area which controls voluntary motor activities or movements of muscles. The premotor area is higher centre for involuntary movements and autonomic nervous system. **Association area** is for coordination between sensation and movements. **Broca's area/motor speech area.** It translates the thoughts into speech. Expression of emotions, intelligence, will-power, memory, personality areas are located in the frontal lobe.

2. Parietal lobes : They are mainly for somesthetic sensation of pain, pressure, temperature, taste (gustatoreceptor).

3. Temporal lobes : It contains centres for smell (olfactory), hearing (auditory), speech and emotions.

4. Occipital lobes : They have visual area mainly for sense of vision.

Area of contact between temporal, parietal and occipital lobes is centre for Wernicke's area or intelligence centre. It helps in the understanding of spoken and written words.

The cerebrum, thus shows all three types of areas sensory, motor and association area.

Basal nuclei or basal ganglia are grey masses present within the white matter or lying on the lateral sides of thalamus. The basal ganglia or nuclei of cerebrum receive neurotransmitters from various parts. They help the cortex in the execution of activities at the subconscious level e.g. writing slow or rapid typing. **Corpus striatum** at the floor of cerebrum is the largest basal nucleus.

Curiosity Box:

- Find out how different functional areas of the brain can be mapped ?
- What is EEG ? What information can be obtained from the EEG ?
- Which are silent areas of the brain?

iii. Diencephalon :

It is the part of the forebrain that contains the epithalamus, thalamus and hypothalamus. It lies below the corpus callosum and above the midbrain. It encloses a single cavity termed **third ventricle/Diocoel** which communicates with the two lateral ventricles of cerebrum through a narrow opening called **foramen of Monro**.

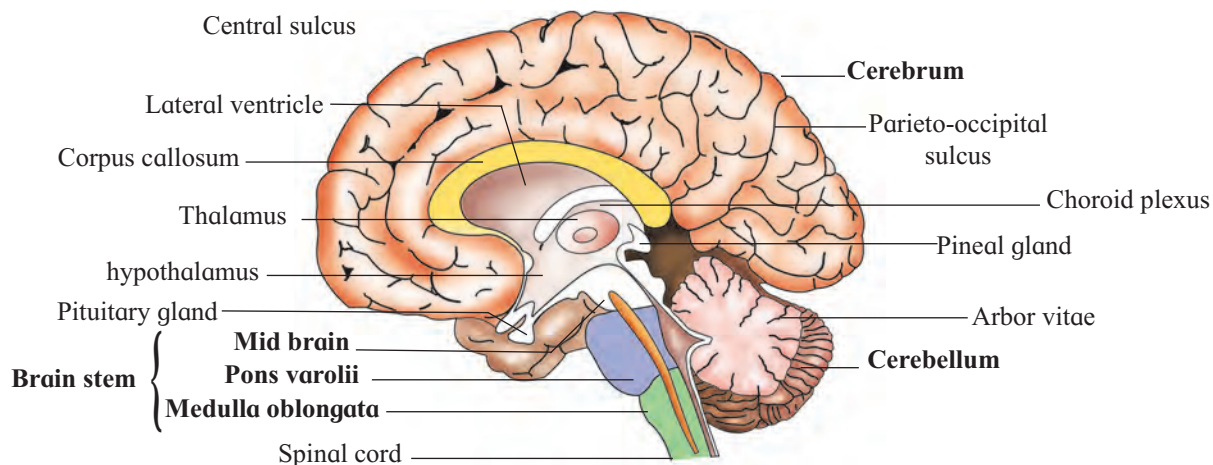


Fig. 9.9 Sagittal section of brain

The **epithalamus** is the thin non nervous roof of the diencephalon. Anteriorly it is fused with the pia mater to form the anterior choroid plexus and from its dorsal wall it is connected to pineal gland through a pineal stalk. (Pineal gland is discussed in detail in chemical coordination). Earlier this gland was thought to be vestigial, but later it has been found to produce the hormone melatonin (sleep inducing hormone; also related to reproductive behaviour). The lateral thick walls of diencephalon form the thalami. They mainly contain grey matter. The habenular commissure connects two thalami. Different parts of the brain are interconnected by the RAS (Reticular Activating System) through the thalami. It is called relay centre as it transmits all sensory impulses except those of olfactory (smell) to the cerebrum (gatekeeper of cerebrum connecting the anterior lateral ventricle to the **iter** posteriorly). The narrow cavity of diencephalon is called **IIIrd ventricle or diocoel**. It connects anteriorly to the two lateral ventricles by a single opening called Foramen of Monroe and posteriorly to the **IVth ventricle or metacoel** through a narrow **duct of Sylvius or iter**.

Hypothalamus : It forms a floor of the **diencephalon**. It is richly supplied with blood vessels (Hypothalamo-hypophyseal portal vein) helps in feedback mechanism for hormonal control). It maintains homeostasis, internal equilibrium of the body and

involuntary behaviour control. Like in the cerebrum, the hypothalamus also contains hypothalamic nuclei in its white matter (refer fig. 9.33 pituitary gland) with neuro-secretory cells involved in the production of hormones oxytocin and vasopressin.

The hypothalamus is a link between the nervous and the endocrine system. It has higher centres for endocrine system. It regulates heart rate, respiration, blood pressure (B.P.), body temperature, water and electrolyte balance. It has centres for hunger, thirst, sleep, fatigue, satiety centre, secretion of glands of stomach and intestine. It also produces neurohormones that stimulate the pituitary gland. A complex neuronal circuit called the **limbic system** is formed by the hypothalamus amygdala, parts of epithalamus and thalamus, hippocampus and other areas. It appears to be responsible for emotional reactions, motivational drives and memory. The floor of the hypothalamus continues as a downward projection called hypophyseal stalk or infundibulum which connects it to the hypophysis (pituitary gland) both physically and functionally by secretion of neurotransmitters (details in chemical coordination). The inferior surface of hypothalamus also bears the optic chiasma (crossing of the two optic nerves) and a pair of mammillary bodies (unique to mammalian brain and responsible for recollective memory).

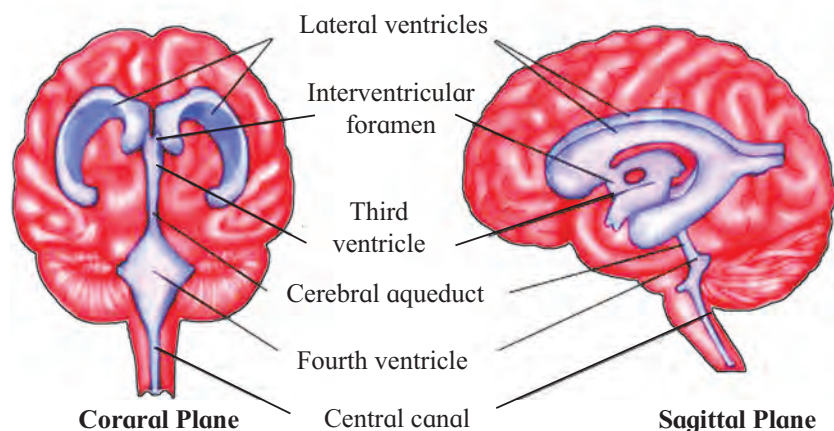


Fig. 9.10 : Ventricles of brain

b. Mid brain :

It is located between diencephalon and the pons varolli. It contains the cerebral aqueduct or iter that connects the third and fourth ventricles. The **corpora quadrigemina** are four rounded elevations on the dorsal surface of the mid brain. The two superior colliculi are involved in visual reflexes and the two inferior colliculi are relay centres for auditory reflexes that operate when it is necessary to move the head to hear sounds more distinctly.

The mid brain also contains on its inferior surface two thick fibrous tracks called cerebral peduncles or **crura cerebri**. These tracts of ascending and descending nerve fibres from RAS and connect the cerebrum mid brain. Near the centre of the mid brain is a mass of grey matter scattered within the white matter. It is called the **red nucleus**. It plays an important role in controlling posture and muscle tone, modifying some motor activities and motor co-ordination.



Always Remember

Wernicke's area : It is the sensory speech area responsible for understanding and formulating written and spoken language.

Broca's area : It is the motor speech area and translates thoughts into speech and controls movement of tongue, lips and vocal cords.

c. Hind brain :

The posterior region of the brain is called **hind brain**. It consists of pons varolli, cerebellum and medulla oblongata. The **pons varolli** appears as a rounded bulge on the underside of the brain stem (brain stem consist of mid brain, pons and medulla and continues upto spinal cord), and contains a cross band of nerve fibres connecting cerebrum, cerebellar lobes, medulla oblongata and spinal cord. It also contains several nuclei. The **cerebellum** is the second largest part of the brain and consists of two lateral hemispheres and a central vermis. It is composed of white matter with a thin layer

of grey matter, the cortex. The white matter intermixes with the grey matter and shows a tree-like pattern called **arbor vitae**.



Can you tell?

Explain - "Cerebellum is well developed in humans".

The surface of cerebellum shows convolutions (gyri and sulci). A number of nuclei lie deep within each lateral or cerebellar hemisphere. Over 30 million neurons lie in the cortex. Three pairs of myelinated nerve bundles called cerebellar peduncles connect cerebellum to the other parts of CNS. It is an important centre which maintains equilibrium of body, posture, balancing orientation, moderation of voluntary movements, maintainance of muscle tone. It is a regulatory centre for neuromuscular activities and controls the rapid activities like walking, running, speaking etc. All activities of cerebellum are involuntary (though may involve learning in early stages). The **medulla oblongata** is the posterior conical part of the brain and continues as the spinal cord. It has inner grey matter and outer white matter. It controls involuntary vital functions like heart beat, respiration, vasomotor activities and peristalsis. It also controls non-vital reflex activities like coughing, sneezing, swallowing, vomiting, yawning etc. The cavity of medulla is called IVth ventricle or metacoel. It's roof has the posterior choroid plexes for secretion of CSF. The posterior choroid plexes also shows 3 openings - a pair of lateral foramen of Luschka and a median foramen of Magendie.



Internet my friend

- Find out the role of the foramina mentioned above.
- What is ataxia ?



Can you tell?

1. The functions of fore brain
2. Injury to the medulla oblongata causes sudden death - Explain.
3. Distinguish between cerebrum and cerebellum.
4. About the mid brain.

B. Spinal Cord :

Spinal cord is the part of central nervous system and forms the lower extension of the medulla oblongata of the brain. Like the brain, it is covered and protected by bony covering and membranes. It lies within the neural canal of the vertebral column and is surrounded by three meninges - dura mater, arachnoid mater and pia mater. The Cerebro Spinal Fluid (CSF) secreted by pia mater, forms a fluid cushion around the spinal cord and within it inside the central canal.

Externally, the spinal cord appears as long cylindrical rod, 42 to 45 cm long and 2.0 to 2.5 cm broad. The spinal cord is broadest at its anterior end gradually tapers into **conus medullaris** (L1 to L2) and continues as a thread like **filum terminale** end posteriorly. Spinal cord shows two swellings along its length called cervical and lumbar swelling.

31 pairs of spinal nerves arise from lateral sides of the spinal cord. These nerves are concentrated in the region of cervical and lumbar swelling and around the conus medullaris. The bunch of nerves in the hind part of the spinal cord, along with the filum terminale, appear like a horse's tail, so called **cauda equina**.

T. S. of spinal cord

The spinal cord is dorsoventrally flattened due to the presence of deep, narrow posterior fissure and shallow, broad anterior fissure. A central canal can be seen in the centre. The fissures divide the spinal cord incompletely into a right and left side. The grey matter is somewhat H-shaped or butterfly shaped and

is on the inner side, while the white matter is on the outer side. The fissures divide the grey matter into six horns, namely dorsal, lateral and ventral horns, while the white matter is divisible into 6 columns or funiculi, namely dorsal, lateral and ventral funiculi. The dorsal and ventral horns extend out of the spinal cord as dorsal root and ventral root of spinal cord respectively. Of these, the dorsal root is connected to the dorsal root ganglion. (It lies just outside and lateral to the spinal cord). It has an aggregation/collection of unipolar sensory neurons.

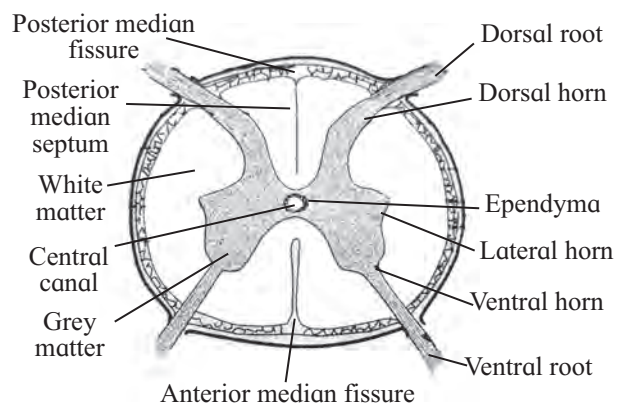


Fig. 9.11 : T. S. of spinal cord

The association or inter-neurons lie inside the grey matter. They receive signal from the sensory nerve, integrate it and direct the response towards motor neurons lying towards the ventral horn. The lateral horns have neurons of autonomic nervous system (ANS). The nerves arising from these neurons, emerge out from the ventral root of spinal nerve.

The white matter consists mainly of bundles of myelinated nerve fibre called ascending and descending tracts. The ascending tracts conduct sensory impulses from spinal cord to the brain and these lie in the dorsal column/funiculi. The descending tracts conduct motor impulses from brain to the lateral and ventral funiculi of spinal cord.

Functions : The spinal cord is the main centre for the most reflex actions. It provides pathway for conduction of sensory and motor impulses to and from the brain. It provides nervous connection to many parts of the body.

2. Peripheral Nervous System (PNS) : The peripheral nervous system connects the central nervous system to the different parts of the body having receptors and effectors. Depending

on the connection to the CNS, the peripheral nerves are classified into two main types -
Cranial nerves - connected to the brain.
Spinal nerves - connected to the spinal cord.

Table 9.12 Cranial nerves - nature and functions

No.	Name	Type	Origin	Organs Innervated	Functions
I.	Olfactory	Sensory	Olfactory bulb	Epithelium of Nose	Smell
II.	Optic	Sensory	Side of diencephalon	Retina of Eye	Vision
III.	Oculomotor	Motor	Floor of mid brain	Eye muscles (4 of 6 eye muscles)	Movement of eye ball
IV.	Pathetic	Motor	Floor of mid brain	Eye muscles (1 of 6 eye muscles, forehead scalp)	Rotation and movement of eye ball
V.	Trigeminal (Dentist's nerve)	Mixed	Ventral side of pons	-	Sensation of skin touch, taste, jaw movement.
	a. Ophthalmic	Sensory	-	Nasal cavity, Upper eyelids, forehead, scalp, conjunctiva, lacrimal gland, scalp	
	b. Maxillary	Sensory	-	Mucosa of nose, palate, upper teeth, upper lip, lower eye lid parts of pharynx	
	c. Mandibular (largest)	Mixed	-	Lower teeth, skin over mandible cheek, side of head in front ear, muscles of mastication	
VI.	Abducens	Motor	Pons	Muscles of eye ball, lateral rectus muscle	Movement of eye
VII.	Facial (bearing geniculate ganglion)	Mixed	Pons	facial, scalp and neck muscles, lacrimal, sublingual, submandibula, nasal and palatine glands	Facial expression, movement of neck, secretion of tears, taste, salivary secretion.
VIII.	Auditory (vestibulo-cochlear)	Sensory	Pons	Internal Ear	Hearing and equilibrium
	i. Vestibular		-		
	ii. Cochlear	-			
IX.	Glossopharyngeal	Mixed	Side of medulla oblongata	Pharynx, tongue, salivary glands	Taste, salivation and swallowing

X.	Vagus (Pneumogastric)	Mixed	Side of medulla oblongata	Larynx, trachea, pharynx, alimentary canal, heart, lungs, pancreas, blood vessels,	Visceral sensations and visceral movements like breathing cardiac, slowing, gastric and pancreatic secretion, gastrointestinal movements
XI.	Spinal accessory	Motor	Side of medulla oblongata	Neck and shoulder muscles, reflexes of thoracic and abdominal viscera larynx, pharynx	Movements of larynx, pharynx, neck and shoulder
XII.	Hypoglossal	Motor	Side of medulla oblongata	Tongue muscles	Movement of tongue

Cranial Nerves : These nerves develop from the brain, in all amniotes (reptiles, birds and mammals). There are 12 pairs of cranial nerves. Roman numbering I to XII is used to denote them. These nerves originate from or terminate into the brain.

According to their function, these are classified as **sensory** (I, II, VIII), **motor** (III, IV, VI, XI, XII) and **mixed** (V, VII, IX, X) nerves.

The details of cranial nerves present in the human body is presented in Table 9.12.

Spinal Nerves : Thirty one pairs of spinal nerves originate from the spinal cord. They are mixed nerves and they provide two way communication between the spinal cord and part of the upper and lower limbs, neck and trunk.

Table 9.13 : Number and types of spinal Nerves

Group	No. of pairs	Region of origin from vertebral column
Cervical	8 (C1-C8)	Neck
Thoracic	12 (T1-T12)	Thorax
Lumbar	5 (L1-L5)	Abdomen
Sacral	5 (S1-S5)	Pelvis
Coccygeal	1 (Co1)	Coccyx



Do you know ?

1. Of the 12 pairs of cranial nerves, only the X (vagus) passes into the body and innervates internal organs.
2. Vagus has the maximum number of branches and longest distribution.
3. V/trigeminal/Dentist's nerve is the largest cranial nerve
4. VI/abducens is the smallest cranial nerve.

Formation of a typical spinal nerve : All spinal nerves are of the mixed type i.e. they have some nerve fibre as sensory and some motor. Each spinal nerve is formed inside the neural canal of vertebral column by two roots - the posterior or dorsal sensory root and anterior or ventral root. Anterior root receives the sensory nerve from the dorsal root ganglion (cell bodies of sensory neurons are located in the ganglion), while the anterior/ventral root gives out the motor nerve. The dorsal sensory and the ventral motor nerves together form the mixed spinal nerve. It emerges out from both sides of the spinal cord through the intervertebral foramen. As soon as it emerges out of vertebral column, it shows three branches viz.

- **Ramus dorsalis** : from skin and to muscles of dorsal side
- **Ramus ventralis** : the largest of the three supplies the organs and muscles on lateral and anterior side
- **Ramus communicans** : the smallest of the three and given out from 1st thoracic upto 3rd lumbar (L3) spinal nerve. It joins the sympathetic ganglia.

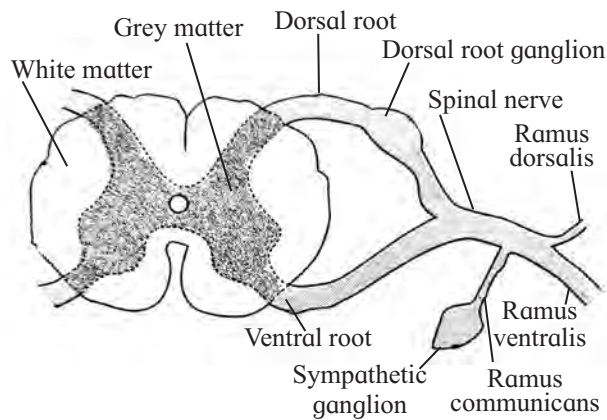


Fig. 9.14 Formation of Spinal Nerve

Reflex Action : It is a sudden, spontaneous automatic, involuntary response to stimulus. The response to stimulus is said to be involuntary as it is carried out without any conscious effort by the brain. The path along which the action is carried out is called **reflex arc**.

[Note : In your earlier classes you have studied about the reflex actions.]

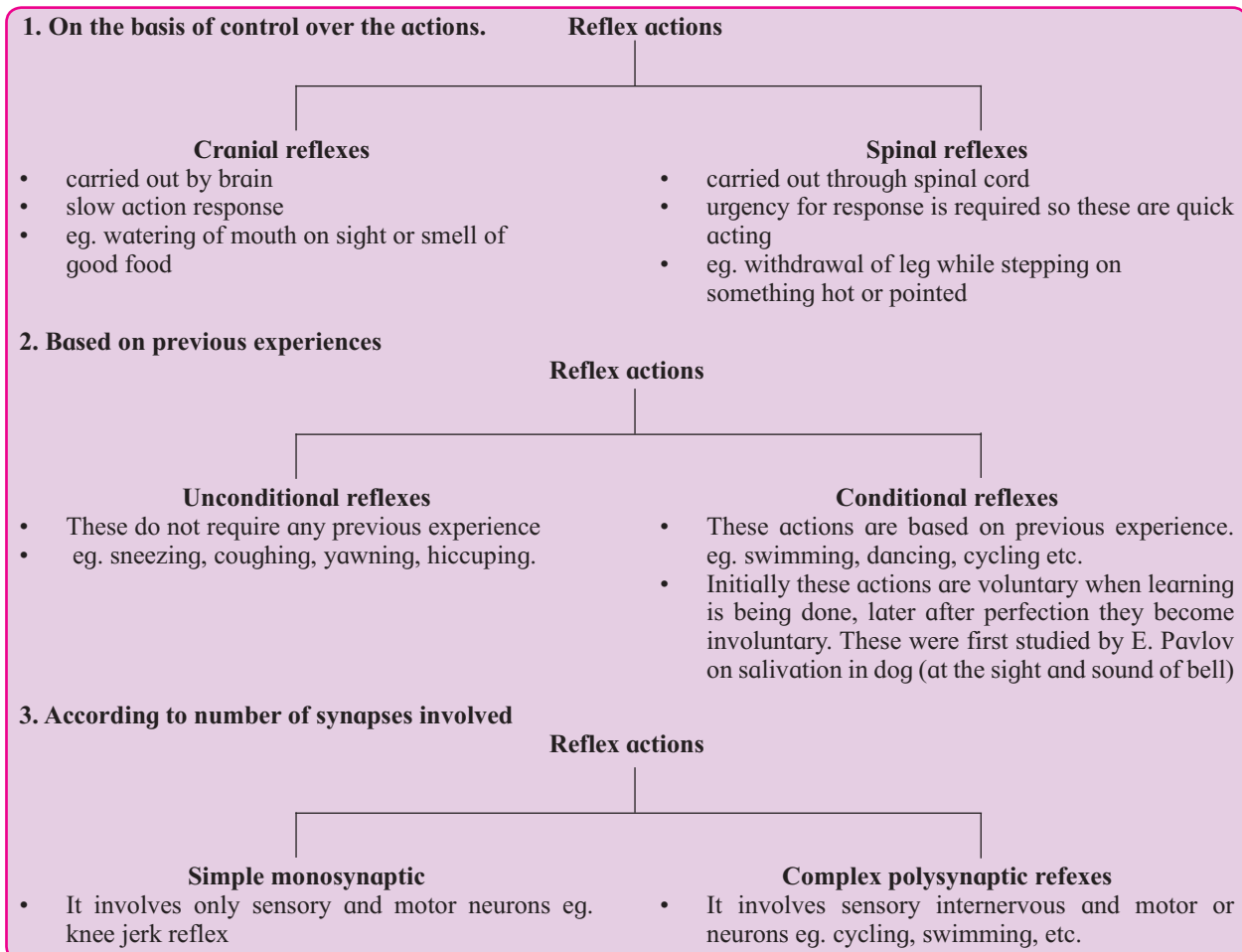
Human nervous system is divided into CNS, PNS, ANS. PNS consist of network of nerves arising from or going to the CNS. Accordingly peripheral nerves are classified as

a. Afferent nerves

b. Efferent nerves

Afferent nerve fibres transmit sensory impulse from tissue or organ to the CNS and the efferent nerve fibres transmit regulatory or motor impulses from the CNS to the various peripheral tissues and organ.

Chart 9.15 Types of Reflex actions



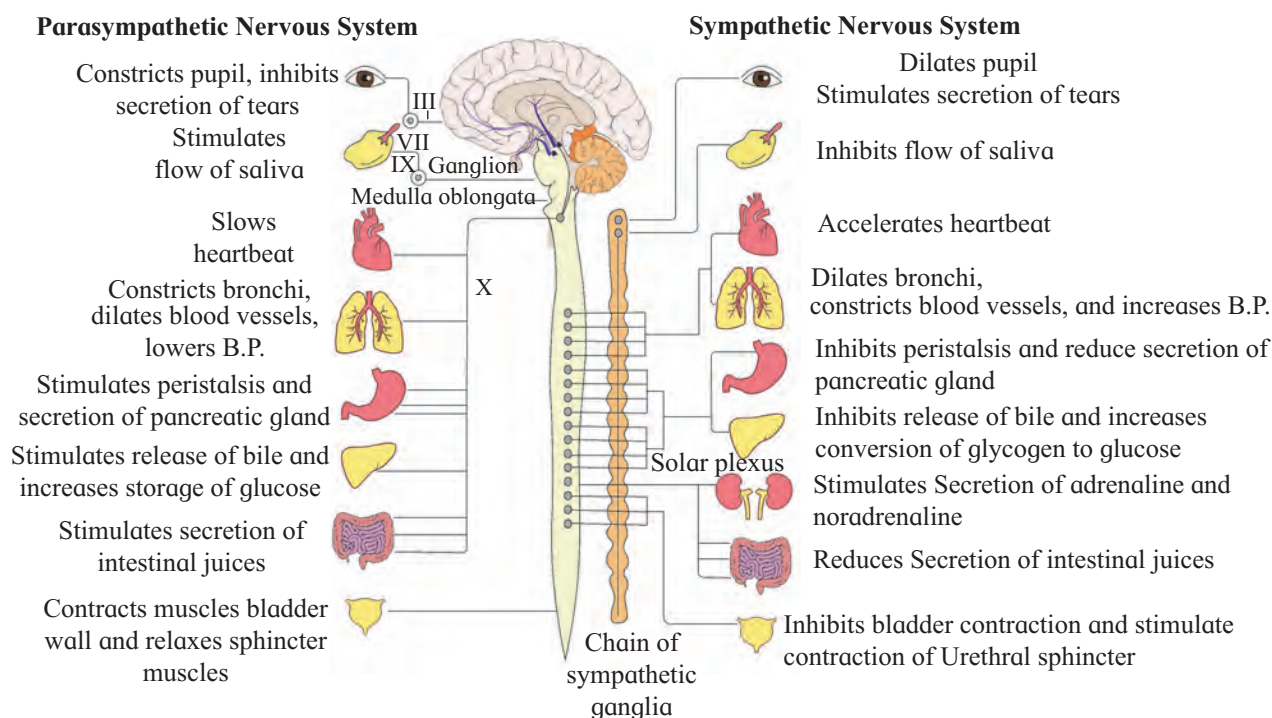


Fig. 9.16 Autonomic Nervous System

However according to recent studies, the extent of PNS has been broadened to incorporate the ANS.

According to this view, the PNS is divided into

- i. somatic nervous system and
- ii. autonomic nervous system

The somatic nervous system relays impulses from CNS to the skeletal or voluntary muscles of the body.

3. Autonomic Nervous System (ANS) :

Autonomic nervous system transmits impulses from CNS to the involuntary organs and smooth muscles of the body.

ANS consists of a special set of peripheral nerves that regulate the activities of involuntary organs like cardiac muscles, smooth muscles, glands etc. In this, impulses are conducted from the Central Nervous system by an axon that synapses with an autonomous ganglion. It is preganglionic neuron. The second neuron in this ganglionic pathway has an axon that extends from the autonomic ganglion to an effector organ and is known as postganglionic neuron.

Autonomic nervous system consists of sympathetic and parasympathetic nervous system.



Use your brain power

Mr. Sharma suffered from a stroke and the right side of his body was paralysed. However his response was normal for knee jerk reflex with either leg. Explain how and why ?

a. Sympathetic Nervous System (SNS):

It is also called thoraco-lumbar outflow. It originates in the thoracic and lumbar region of spinal cord (T1 to L3) and consists of 22 pairs of sympathetic ganglia which lie on a pair of sympathetic cords on lateral sides of the spinal cord.

The pre-ganglionic nerve fibres are short and post ganglionic nerve fibres are long. Adrenaline and Noradrenaline is produced at the terminal ends of postganglionic nerve fibres at the effector organ, hence it is also called **Adrenergic fibres**. Sympathetic nervous system controls body activities during emergencies (fight or flight response). It has excitatory and stimulating effect on most organs of the body except in the digestive and the excretory organ.

Table 9.17 : Comparison between sympathetic and Parasympathetic Nervous System

Organ/Region	Sympathetic effect	Parasympathetic effect
Heart beat	Increases	Decreases
Blood vessels	Constricts	Dilates
Arterial B.P.	Increases	Decreases
Pupil of Eye	Dilates	Constricts
Gastrointestinal movements (stomach and intestine)	Retards peristalsis	Accelerates peristalsis
Urinary bladder	Relaxes the bladder	Contracts the bladder

b. Parasympathetic Nervous System :

It is also called cranio-sacral outflow. It consists of the branches from the cranial (III, VII, IX, X) nerves, sacral (II, III) and spinal (IV) nerves. It consists of ganglia which are very close or within the wall of the effector organs. The pre-ganglionic nerves are long and post-ganglionic nerves are short. Acetylcholine is produced at the terminal end of post-ganglionic nerve at the effector organ, hence these are also called **cholinergic fibres**.

Parasympathetic nervous system is antagonistic to sympathetic nervous system. It brings back to normal, all activities which are stimulated by the sympathetic system. Hence it is also called housekeeping system. It has an inhibitory effect on most organs. However, the activities like those associated with digestion, peristalsis and micturition, which are inhibited by sympathetic system are thus accelerated by the parasympathetic system.

The answers to all these questions are the presence of a sensory system in our body. It consists of simple to complex structures called sensory receptors.

9.7 Sensory Receptors :

Sensory receptors are some specialized structures in the body to receive the various stimuli from the external or internal environment. The nature of the receptor is defined by the

type of stimulus that it can respond to e.g. a photoreceptor responds to light. What is the receptor? It implies that when a specific type of stimulus is received by the sensory neuron (receptor), it generates an action potential in it and this action potential is conducted in the form of an impulse. These impulses are conducted to the different functional areas of the brain for processing and interpretation.

Curiosity Box:

1. Ever wondered as to how we are able to understand the smell of the first showers of rain, or the sudden changes in the climate ?
2. How are we able to hear the chirping of the birds and recognize the sound of the bird ?
3. How can we see and enjoy the beautiful colours of the nature after the sunrise ?



Internet my friend

Find out the fifth category of taste called Umami apart from the four recognized ones - salty, sour, sweet, bitter.

Classification of receptors :

Receptors are classified on the basis of their location, function and their sensitivity to specific stimuli. Their classification is given in the following chart.

Chart 9.18. Types of receptors

No.	Name/Type of receptor	Location	Function
I. Exteroceptors : Receive external stimuli			
	a. Phonoreceptors	Internal Ear - organ of corti	Sound reception
	b. Statoreceptors	Internal Ear- semicircular canals	Receptors for maintaining balance and equilibrium
	c. Photoreceptors	Retina of Eye	Receives sensory stimuli for vision
	d. Thermoreceptors	Skin	Receives sensory stimuli for heat (caloriceptors) and cold (trigidocetptors)
	e. Mechanoreceptors	Skin	Sensitive to mechanical stimuli like touch, pain, pressure, deep pressure, etc.
	f. Chemoreceptors <ul style="list-style-type: none">• Gustatoreceptors• Olfactory receptors	Taste buds of tongue Olfactory Epithelium of Nose	Sensitive to taste of sweet, salt, sour, bitter and umami. Sensitive to about 10,000 different smells
II. Interoceptors : Receive stimuli coming from within the body			
	a. Enteroceptors	from internal body organs	Sensitive to stimuli coming from internal organs like hunger, thirst, pain, osmotic change
	b. Proprioceptors	Joints, muscles and tendons	Detect changes in the movements of joints, tendons and muscles; pain, tension and sensitive to vibrations
	c. Baroreceptors (*These are also considered as mechanoreceptors, receiving signals from internal organ)	Present in walls of atria, venae cavae, aortic arch, carotid sinus	Sense changes in B.P. so as to restore homeostasis through vasodilation or vasoconstriction

Eye :

The eyes are a pair of sensory organs of vision. These are located in the orbit of skull with a cushion of fat around them. Each eye is spherical/rounded and called eyeball. The eyes are protected within bony sockets and by eyebrows, upper and lower eyelids, eyelashes and the lacrimal (tear) glands. Movement of the eyeball within the orbit is controlled by 6 sets of muscles.

Wall of the eyeball is made up of 3 layers (1) sclera, (2) choroid and (3) retina

1. Sclera/sclerotic : It is the outermost layer made of dense fibroelastic connective tissue with collagen fibres. It provides attachment to the eyeball muscles. The anterior thick, transparent part of sclera is cornea. It is slightly bulged out for focussing light on the retina. The sclera is provided with blood vessels, however the cornea is devoid of them. Cornea is nourished by aqueous humour and also by lacrimal secretion. The exposed part of sclera and the entire cornea are covered by a transparent membranous covering called **conjunctiva**. It provides protection and lubrication to the cornea.

2. Choroid /Uvea : It is the middle, vascular and pigmented layer. It is not a complete layer and can be divided into 3 regions - a. Choroid proper b. Ciliary body c. Iris

a. The Choroid proper : It lines the sclera. Due to its pigmented nature it prevents internal reflection. The blood vessels of choroid provide nutrition and oxygen to the retina.

b. Ciliary body : It is a thick, muscular, ring like structure at the junction of choroid and iris. Its epithelium secretes aqueous humor. Attached to the ciliary body are suspensory ligaments which hold the lens. The ligaments and muscles of the ciliary body help in the adjustment of the size of lens.

c. Iris : At the junction of the sclera and cornea, the vascular part of choroid sharply bends into the cavity of eyeball, forming a thin and coloured partition called iris. It is perforated in the middle by an aperture called pupil. Smooth muscles of the iris help in regulating the size of pupil depending on the intensity of light entering the eyeball. The pigment in the iris determines the colour of the eye.

Lens : It is a transparent, elastic and biconvex structure. It is suspended in the eyeball by the suspensory ligaments. The lens and suspensory ligaments divide the cavity of the eyeball into a small anterior aqueous chamber, filled with a clear watery fluid aqueous humor and a posterior large vitreous chamber, filled with a jelly like vitreous humor. It maintains shape of the eyeball and maintain pressure for keeping the lens in position.

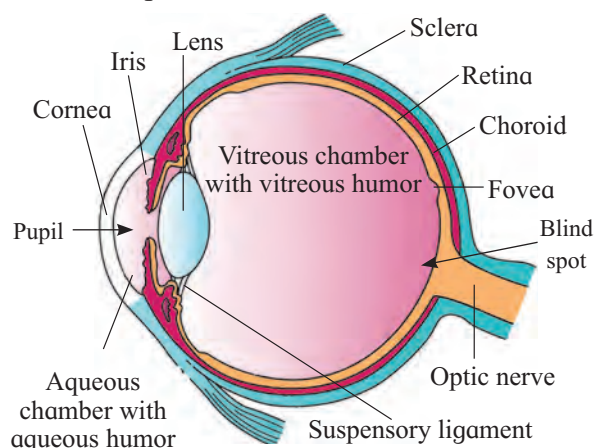


Fig. 9.19 : Eye

3. Retina : It is the innermost, delicate, non vascular light sensitive layer. It has 2 regions (a) single layer of pigmented non sensory part lining the iris and ciliary body (b) sensory part lining the choroid. It has an outer pigmented part and an inner nervous part. The inner nervous part is transparent and made of 3 layers (1) outer photosensitive layer made of rod and cone cells. (2) middle layer of bipolar nerve cells (3) inner layer of ganglion cells. The nerve fibres from the basal end of the ganglion cells collectively form the optic nerve.



Do you know ?

Why do eyes of cats and dogs glow in the night ? The glowing of eyes in some animals is due to presence of a reflecting layer behind the retina, called tapetum lucidum.

The **blind spot** is an area diagonally opposite to the lens. It is the area of retina from where the optic nerve and blood vessels leave the eyeball. There are no rod and cone cells in this region. An area, lateral to and above the blind spot is called **yellow area** or **macula lutea**. At its centre is a depression called **fovea centralis**. It has maximum density of cone cells and is the place of formation of sharpest vision.

The rod and cone cells lie deep in the retina, so that light has to pass through the ganglion and bipolar cells before reaching them.

Photo receptor cells : These are of two types (a) Rod cells (b) cone cells.

They contain light sensitive proteins termed as photopigments. The cones are responsible for daylight (photopic) vision and colour vision. While the rods function in dim light (Scotopic) vision. The purple red protein called rhodopsin is present in the rods which is vitamin A derivative. The cones are of three types having their own characteristic photopigments that respond to red, green and blue lights. Various combinations of these cones and their photopigments produce sensation of different colours. The sensation of white light is produced due to the simultaneous equal stimulation of these three types of cones.

The Optic nerve consists of the fibres arising from the base of ganglion cells. It leaves the eye ball from the posterior side and carries visual impulses from the retina to the brain.

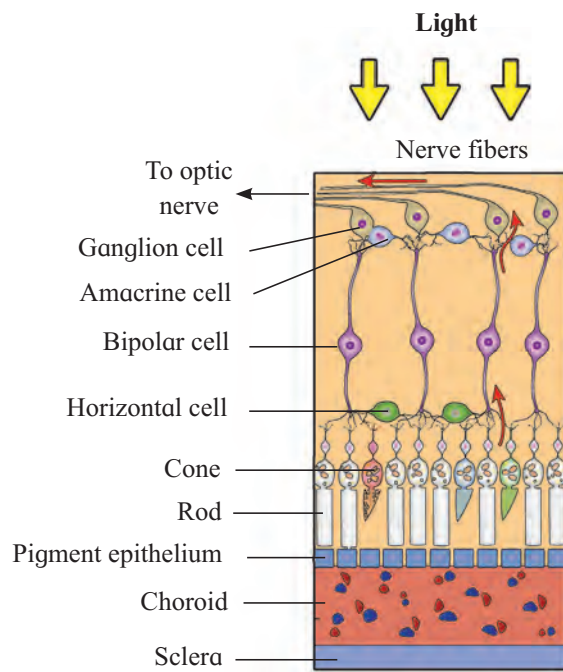


Fig. 9.20 : Structure of retina

Generation of image :

The light rays from the object pass through the conjunctiva, cornea through the pupil upon the lens and is focused on the retina to form an image. In the visual area of cerebrum, the nerve impulses are analysed and the image formed is recognized.



Always Remember

Accommodation : The lens makes fine adjustments to bring a sharp focus on retina. The ability of the lens by which the light ray from far and near objects is focused on the retina is called accommodation power of the lens.



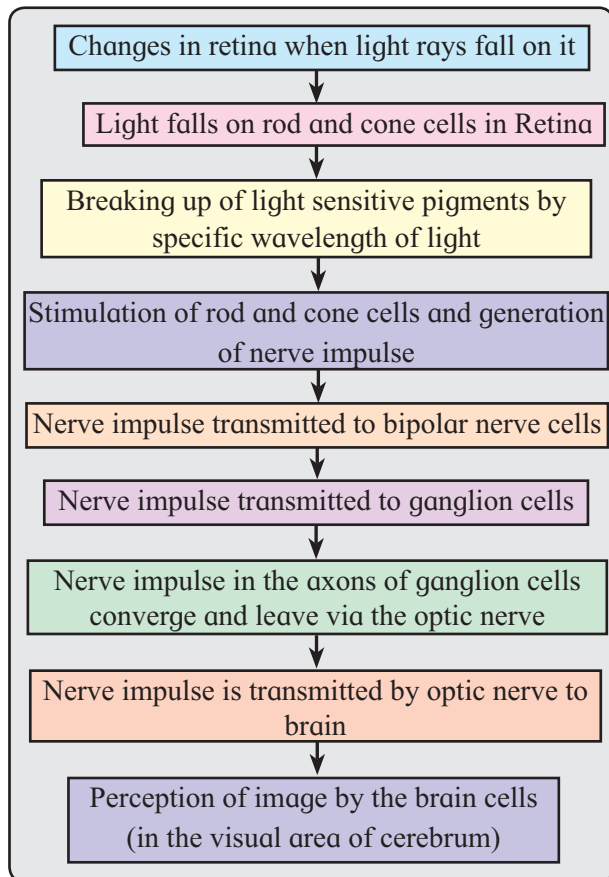
Internet my friend

- Find out information about those who can donate eyes ?
- Is there any age limit for donating eyes ?
- Who cannot donate eyes ?
- Facts about eye donation.



Activity :

Collect information about causes and corrections for myopia, hypermetropia, astigmatism, presbyopia, cataract, night blindness.



Ear :

The human ear is called stato-acoustic organ and it has two functions - hearing and body equilibrium. Anatomically the ear is made up of three parts : the external ear, middle ear and inner ear.

The **external ear** consists of ear pinna, auditory canal and tympanic membrane. In humans, the ear **pinna** is an immovable part, supported by elastic cartilage structure. It leads into an auditory canal. The pinna collects and sends the sound waves into the auditory canal. The **auditory canal** ends at the ear drum. It transfers the sound waves to the ear drum. There are very fine hair and wax secreting sebaceous glands in the skin of pinna and

auditory canal. The **tympanic membrane** is a delicate, membranous structure which transmits the sound waves to the middle ear. It is formed of connective tissue covered with skin on the outside and mucous membrane on the inside.

The **middle ear** consists of a chain of three ear ossicles called **Malleus** (hammer), **Incus** (anvil) and **Stapes** (stirrup-the smallest bone). On receiving the vibrations from the tympanic membrane, the ear ossicles amplify the vibrations and transfer these to the **cochlea**. A short **eustachian tube** connects the middle ear to the pharynx. It equalises air pressure on both sides of the ear drum.

The **internal ear** consists of the labyrinth. The labyrinth consists of bony labyrinth and membranous labyrinth. These are filled with perilymph and endolymph respectively. The membranous labyrinth consists of three parts: vestibule, semicircular ducts and cochlea.

The **cochlea** is a coiled duct and contains three fluid filled chambers separated by **Reissner's membrane** and **basilar membrane**. The upper chamber towards **vestibule** is called **scala vestibuli** and the bottom chamber **scala tympani** are filled with perilymph. The middle chamber is the **scala media**. It is filled with endolymph while scala vestibuli and scala tympani are filled with perilymph. The **organ of Corti** is a pea sized structure located on basilar membrane (floor of scala media).

The organ of Corti has a sensory epithelium over the basilar membrane. The sensory epithelium is in contact with a gelatinous **tectorial membrane**. The sensory cells have sensory hair on their free end so also called hair cells. In between the rows of hair cells are present supporting cells.

Hair cells have long stiff microvilli called **stereocilia** on their apical surfaces. Above these stereocilia, is a jelly like membrane

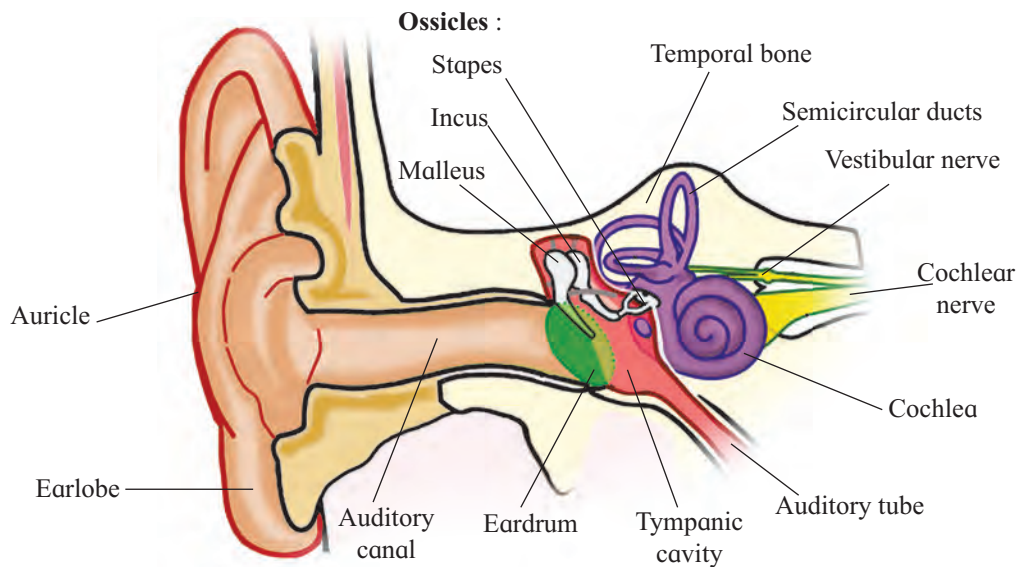


Fig. 9.21 : Structure of human ear

called **tectorial membrane**. This organ acts as a transducer, converting sound vibrations into nerve impulses.

Internal Ear and the mechanism of balance :

Besides the cochlea, the inner ear also has the vestibular apparatus which is composed of three **semi-circular ducts or canals** and the utriculo saccular region with the otolith organ. All three semi-circular canals lie in different planes at right angle to each other. These canals are filled with endolymph. The base of each of the canal has an ampulla in which there is a sensory spot called **crista**. The cristae help in maintaining equilibrium. The vestibule has two sensory spots viz. **macula of saccule** and of **utricle**. The macula consist of hair cells and supporting cells. Tips of the hair and cilium project into a thick gelatinous sheath called **otolithic membrane**. Within this membrane are secreted minute particles, the **otoliths or otoconia**. These are made of CaCO_3 and protein. The macula and crista are the receptors sensitive to the position of the head with respect to gravity. The three semi-circular canals are arranged such that the movement in any plane can be detected by these cells and thus the balance and posture of the body is maintained. Receptors for dynamic balance lie in the cristae of ampullae while for static/linear

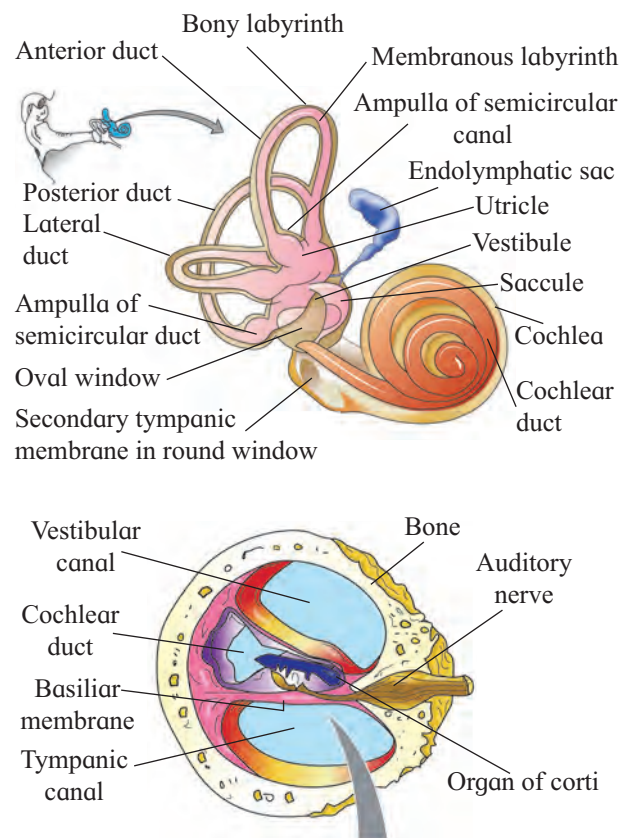


Fig. 9.22 : Human Internal Ear



Activity :

The auditory centre of the brain analyses the impulses received and the sound is perceived.

Draw flow chart of mechanism of hearing.

balance these are in the maculae of utriculus and sacculus.

Mechanism of Hearing :

Pinna of the ear receives the sound waves and directs them to eardrum. Eardrum vibrates and these vibrations are amplified and transmitted through the ear ossicles to the endolymph inside cochlea. This generates wave in the endolymph. These waves induce ripples in the basilar membrane. These movements in the basilar membrane cause the hair cells to press against tectorial membrane. This generates nerve impulse in the afferent neurons. Impulse is sent to the brain via the auditory nerve. Auditory cortex of the brain decodes the sound.



Use your brain power

1. What is the function of tympanic membrane?
2. Enlist the various receptors found at various location in the body.

9.8 Disorders of nervous system :

Psychological disorders :

Commonly called mental disorders, are a wide range of conditions that affect the mood, thinking or behaviour. These affect multiple areas of life and create distress for the person suffering from it. Some of the major categories of psychological disorders are - ID (Intellectual disability, (formerly known as mental retardation), ASD (Autism spectrum disorder), BD (bipolar disorder), depression, anxiety disorder, ADHD (Attention Deficit Hyperactivity Disorder) and stress related disorders.

Parkinson's disease :

Degeneration of dopamine producing neurons in the CNS causes Parkinson's disease.

Symptoms develop gradually over the years. Symptoms are tremors, stiffness, difficulty in walking, balance and co-ordination.

Alzheimer's disease :

It is the most common form of dementia. It's incidence increases with the age, showing the loss of cognitive functioning i.e. thinking, remembering, reasoning and behavioral abilities to such an extent that it interferes with the persons daily life and activities. It occurs due to loss of cholinergic and other neurons in the CNS and the accumulation of amyloid proteins. There is no cure for Alzheimer's, but treatment slows down the progression of the disease and may improve the quality of life.

CHEMICAL COORDINATION

The cells and organisms communicate with each other through chemical signals. Also they are broadly of four types as follows :

Autocrines : Cells release secretion to stimulate itself.

Paracrines : Cells release secretion to stimulate neighbouring cells.

Endocrines : Cells release secretion to stimulate distant cells.

Pheromones : Organs release secretions to stimulate other organism.

Higher animals have complex body organization. Due to this, in addition to the nervous coordination, there is need of chemical coordination. Chemical coordination is carried out by secretions of ductless glands. This chemical coordination system is also called the endocrine system.

9.9 Endocrine system :

The endocrine system controls body activities by means of chemical messengers called hormones. Hormones are released directly into the blood. The hormone is carried all over the body via blood. However the message is relayed only to the target organs which are stimulated to carry out specific process which include activities like growth and development.

Chemical nature of hormones

- I. Amines :** These are simple amines. Catecholamines secreted by adrenal medulla, epinephrine and non-epinephrine and melatonin from pineal gland. Some are modified from the amino acids. e.g., Thyroxine.
- II. Peptide hormones :** These hormones consist of long or short chains of amino acids. e.g. Hormones of hypothalamus oxytocin, ADH, GnRH.
- III. Protein hormone :** Insulin, glucagon TSH, FSH, LTH, GH, relaxin.
- IV. Fatty acid derivatives :-** Prostaglandin
- V. Steroid hormones :** These hormones are lipid soluble and derived from cholesterol and other steroids. e.g. estrogen testosterone, aldosterone. Action of these hormones is concerned with long lasting responses.
- VI. Gas :** NO (Nitric Oxide)

Properties of Hormones :

They act as chemical messengers and are effective in very low concentration. Hormones can function as regulators that inhibit or stimulate or modify specific processes. Some hormones interact with receptors present on plasma membrane of target cells where as some enter the nucleus to interact with genes. Hypersecretion or Hyposecretion of hormones leads to various disorders.

These are metabolised after their function. Thus cannot be reused. Hormone secretion is regulated by positive or negative feedback mechanism.

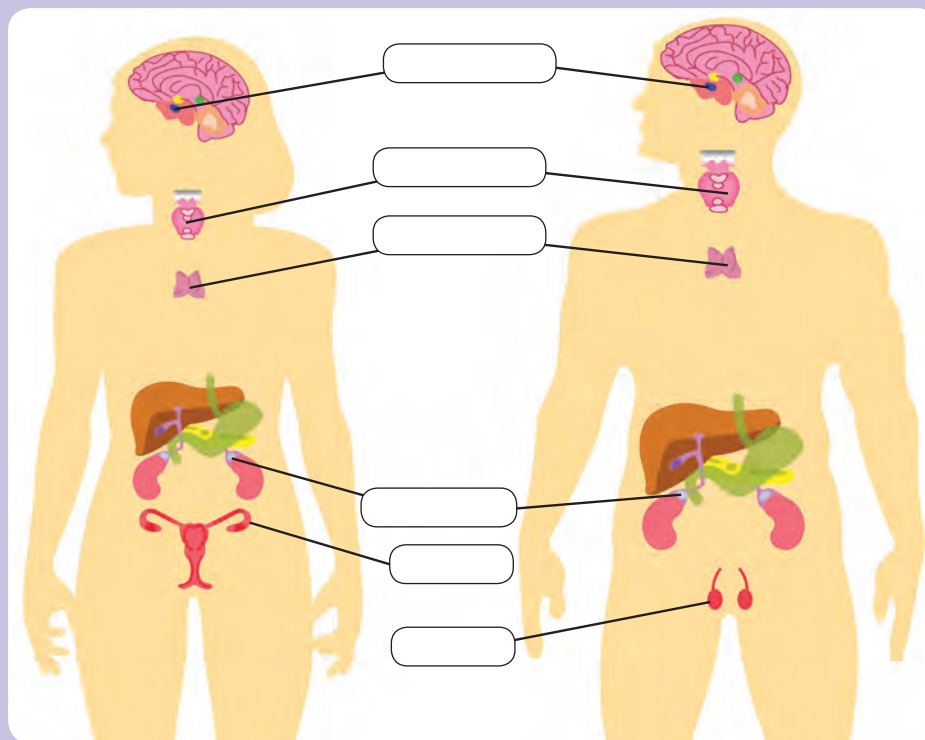
Mechanism of hormone action :

Hormones are released in a very small quantity. They produce their effect on the target organs / cells by binding to hormone receptors. The hormone receptors may be on the cell membrane or may be intracellular.



Activity :

Identify the glands and state their functions.



A hormone receptor complex is formed and this leads to biochemical changes in the target tissue.

A. Mode of hormone action through membrane receptors :

Hormones like catecholamines, peptide and polypeptide hormones are not lipid soluble. Therefore they cannot enter their target cells through plasma membrane.

These non steroid water soluble hormones interact with surface receptor, which initiate metabolic activity. Molecules of amino acid derivatives, peptide hormones bind to specific receptor molecules located on the plasma membrane. The hormone receptor complex causes the release of an enzyme adenylate cyclase from the receptor site. This enzyme forms cyclic AMP from ATP of the cell. cAMP activates enzymatic actions. The hormone acts as the first messenger and cAMP is the second messenger. Other kind of second messengers are Ca^{++} , cGMP and IP_3 (Inositol triphosphate).

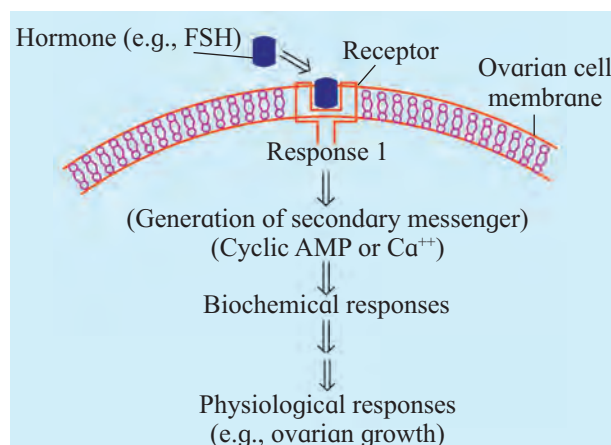


Fig. 9.23 : Mechanism of hormone action through membrane receptor



Internet my friend

What is negative and positive feedback of hormone action ?

B. Mode of action through intracellular receptors :

Steroid and thyroid hormones are lipid soluble and easily pass through plasma membrane of target cell into the cytoplasm. In the cytoplasm, they bind to specific intracellular receptor proteins forming a hormone-receptor complex that enters the nucleus. In the nucleus, the hormone receptor complex binds to a specific regulatory site of DNA. The activated genes transcribe mRNA which directs protein synthesis and enzymes in the cytoplasm. Action of lipid soluble hormones is slower but long lasting.

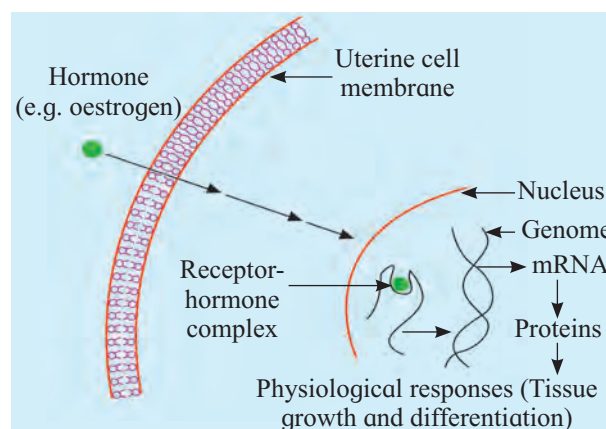


Fig. 9.24 : Mechanism of hormone action through intracellular receptor

9.10 Major endocrine glands :

A. Hypothalamus :

It is ectodermal in origin. It is located at the floor of diencephalon. Major function of hypothalamus is to maintain homeostasis. It controls the secretory activity of pituitary gland by the releaser (RH) and inhibitor (IH) hormones. All hormones of hypothalamus are peptide hormones. They are secreted by the neurosecretory cells so they are called neurohormones. The hormones secreted by hypothalamus are : ADH, Oxytocin.

1. **Adrenocorticotropin Releasing Hormone** : It stimulates the release of ACTH by the anterior pituitary gland.
2. **Thyrotropin Releasing Hormone** : It stimulates the release of TSH by anterior pituitary gland.
3. **Gonadotropin Releasing Hormone (GnRH)** : It stimulates pituitary to secrete gonadotropins.
4. **Prolactin Inhibiting Hormone (Prolactostatin)** : It inhibits prolactin release by anterior pituitary gland.
5. **Somatostatin** : It inhibits the release of growth hormone.
6. **Somatotropin** stimulates release of growth hormone.
7. **Gastrin Releasing Peptide (GRP) and**
8. **Gastric Inhibitory Polypeptide (GIP)**

B. Pituitary gland or hypophysis gland :

The pituitary gland is the smallest gland. It is a pea sized reddish-grey coloured gland. It controls almost all other endocrine glands, hence earlier it was called the master endocrine gland. It is located just below the hypothalamus and is attached to it by a stalk called **infundibulum** or hypophyseal stalk. Pituitary gland remains lodged in a bony depression called **sella turcica** of the sphenoid bone. Pituitary gland consists of two lobes called **anterior lobe (Adenohypophysis)** and **posterior lobe (Neurohypophysis)**. Both the lobes develop from different parts of embryo. Hence it has dual origin.

Adenohypophysis is an outgrowth from the roof of buccal cavity. This outgrowth is called Rathke's pouch. It grows upward towards the brain. The neurohypophysis grows as a downward extension of hypothalamus. The two outgrowths together form the pituitary gland. The connection of Rathke's pouch with pituitary gland is lost in embryo. Intermediate lobe (Pars intermedia) is a small reduced part

lying in the cleft between the anterior and posterior lobe.

Neurohypophysis is connected directly to the hypothalamus by axon fibres. Adenohypophysis and intermediate lobes are connected to the hypothalamus through **hypophyseal portal system**.

Hypophyseal portal system :

Various hormones secreted by hypothalamus reach the pituitary gland through the hypophyseal portal system. The portal vein collects blood from various parts of hypothalamus and opens into anterior lobe of pituitary. From pituitary, the vein finally carries the blood into the superior vena cava.

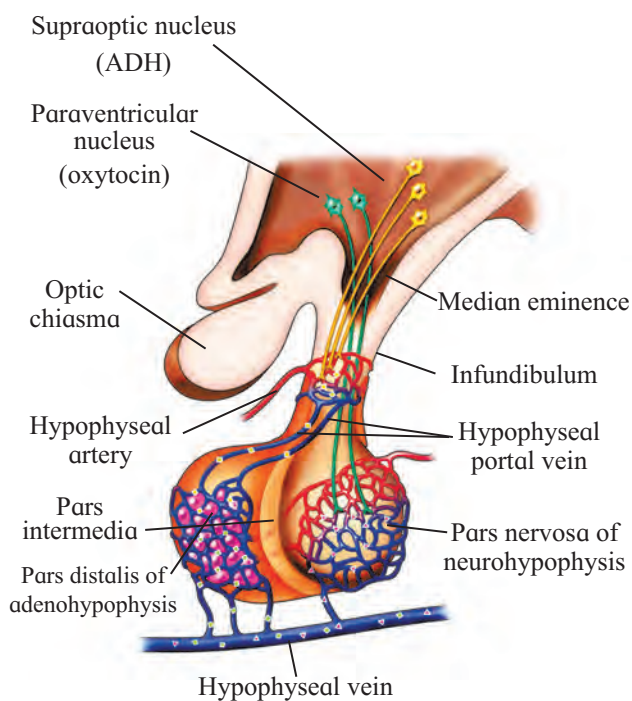


Fig. 9.25 : Pituitary gland

Adenohypophysis: It is the larger lobe of pituitary gland. It is a highly cellular and vascular part of pituitary gland. It contains various types of epitheloid secretory cells, acidophils, basophils and chromatophores. It is differentiated into three parts – pars distalis, pars intermedia and pars tuberalis.

The hormones of adenohipophysis are as follows :

1. Somatotropin /Somatotropic Hormone / STH / Growth hormone / GH :

This hormone stimulates growth and development of all tissues by accelerating protein synthesis and cell division. Highest secretion of GH is seen till puberty and then its secretion becomes low. However, it is continuously secreted through out life for repair and replacement of body tissue or cells.

Improper secretion of growth hormone produces various disorders. Hyposecretion of growth hormone since childhood results in stunted physical growth and the condition is called **pituitary dwarfism**.

Hypersecretion of growth hormone in childhood causes **Gigantism** a condition of overgrowth. The individual attains abnormal height. When the pituitary gland produces excess growth hormone in middle aged adults, it results in disproportionate growth causing disfigurement and enlargement of bones of nose, lower jaw, hands, fingers and feet. The condition is called **Acromegaly**.

2. Thyrotropin / Thyroid stimulating Hormone / TSH : Its primary action is to stimulate the thyroid gland for secretion of the hormone thyroxine.

3. Adreno corticotropic hormone / ACTH / Adrenocorticotropin : It stimulates adrenal cortex to produce and secrete its hormones. It maintains functioning of adrenal cortex.

4. Prolactin / Luteotropin / Mammotropin: Prolactin is unique among pituitary hormones as it is under predominant inhibitory control from hypothalamus. Prolactin activates growth of breasts during pregnancy (mammotropin) and stimulates the milk production and secretion of milk (lactogenic) by mammary gland after child birth.



Internet my friend

Collect the information about Simmond's disease and Gorilla rib.

5. Gonadotropin :

a. Follicle stimulating hormone/ FSH :

It stimulates growth of ovarian follicles in the females, while in males, it is concerned with the development of seminiferous tubules.

b. Leutinizing Hormone/ LH :

In female, the leutinizing hormone helps in growth, ripening and discharge of ovum (ovulation) from graafian follicle. FSH and LH are responsible for stimulation of ovaries to produce oestrogen. LH also induces the ruptured follicle to develop into corpus luteum for production of progesterone.

c. Interstitial cell stimulating hormone / ICSH :

In males, ICSH stimulates the testes to produce the androgen called testosterone. Testosterone is responsible for development of secondary sexual characters.

Neurohypophysis: It is differentiated into three parts, 1. Pars nervosa/ neural lobe 2. Infundibulum 3. Median eminence.

The pars nervosa acts as storage area for the secretions of hypothalamus. It stores and releases oxytocin and vasopressin.

1. Oxytocin : It stimulates contraction of uterus during parturition. It also stimulates the contraction of mammary glands to initiate ejection or release of milk. So is called birth hormone or milk ejecting hormone.

2. Antiduretic Hormone (ADH)/ Vasopressin:

It stimulates the re-absorption of water in distal convoluted tubule and collecting ducts of uriniferous tubules of the kidneys. It decreases loss of water by reducing the urine quantity. It increases blood pressure by causing vaso constriction.

Deficiency of ADH reduces water re-absorption and increases urine output. This condition is called **diabetes insipidus**. No glucose is lost in the urine. Excessive micturition causes excessive thirst. This condition is called polydipsia.

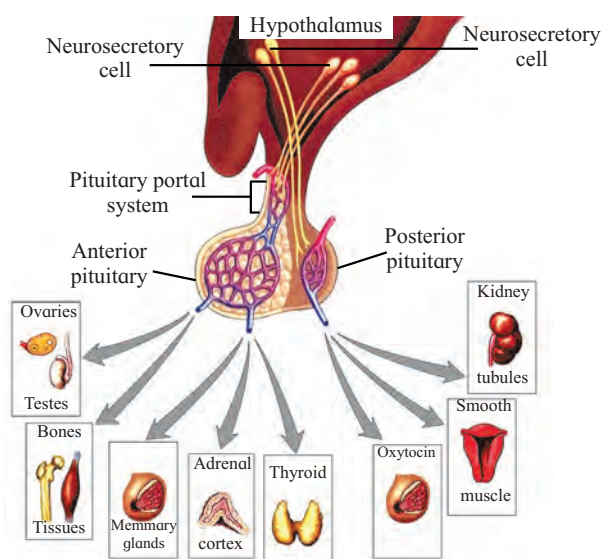


Fig. 9.26: Hypothalamus, Adenohypophysis and Neurohypophysis

Pars intermedia : It is poorly developed in human beings. It secretes Melanocyte Stimulating Hormone (MSH) in some lower vertebrates like fish and amphibians.

MSH stimulates the dispersion of melanin granules in melanocytes and is responsible for skin pigmentation.



Can you tell?

1. State properties of hormones ?
2. Explain the mode of action of steroid hormones ?
3. Describe neurohormonal regulation of pituitary and thyroid gland ?
4. Give names and functions of hormones secreted by adenohypophysis ?

C. Pineal gland : The pineal gland is given off from the roof of diencephalon and is located between the two cerebral hemispheres. The pineal gland is sensitive to the biochemical signals of light. It secretes a hormone called **melatonin** also known as sleep hormone. Melatonin is derived from tryptophan and plays a very important role in the regulation of **Biological Clock** i.e. 24 hour diurnal rhythm of body. It helps in maintaining the normal rhythm of sleep-wake cycle and also influences body temperature, metabolism and reproductive cycles.

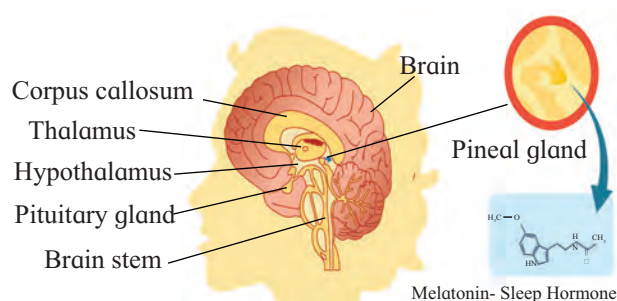


Fig. 9.27 : Pineal gland

D. Thyroid gland :

It is the largest endocrine gland weighing about 25 gms. This bilobed gland is situated in front of the trachea just below the larynx. It is richly supplied with blood vessels. The two lobes of thyroid gland are connected by a non-secretory band called **isthmus**. The thyroid lobes are composed of rounded follicles held together by interfollicular connective tissue called **stroma**. The stroma contains blood capillaries and small groups of **parafollicular cells** or 'C' cells (**clear cells**). Thyroid **follicles** are composed of a single layer cuboidal epithelium resting on a basement membrane and its lumen is filled with a gelatinous hormone rich **colloid**.

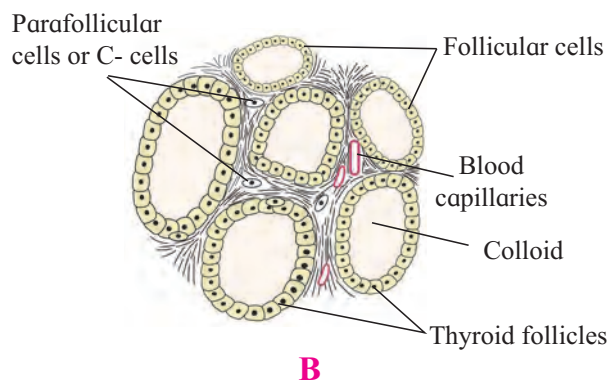
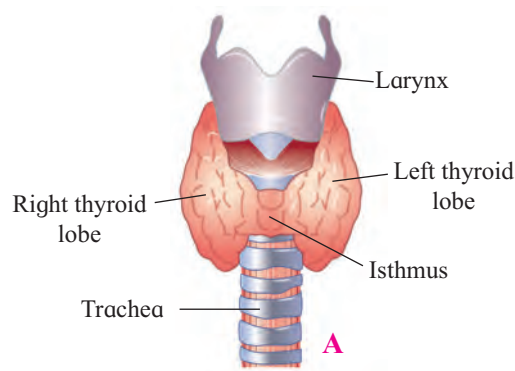


Fig. 9.28 Thyroid gland

Thyroid gland is stimulated to secrete its hormones by thyroid stimulating hormone (TSH). The two hormones secreted by the follicular cells are Thyroxine/tetra iodothyronine/ T_4 (four atoms of iodine) and Triiodothyronine or T_3 (three atoms of iodine). Thyroxine is synthesized by attaching iodine to amino acid tyrosine by enzymatic action. The amino acid tyrosine molecule binds to an atom of iodine to produce Monoiodothyronine (T_1) or to two atoms of iodine to produce Diiodothyronine (T_2). T_1 and T_2 molecules bind end to end to make colloidal mass inside the follicle. They are further metabolised to prepare T_3 and T_4 . Triiodothyronine or T_3 is secreted in small quantity. It is physiologically more active than T_4 . Thyroid gland is the only gland that stores its hormones. T_3 and T_4 hormones are stored before secretion and are regulated by thyrotropin of pituitary gland by negative feedback mechanism.



Think about it

1. Patient suffering from hypothyroidism shows increased level of TSH. Why ?
2. Why do we use iodized salt?
3. In which part thyroid gland stores its hormones?

Thyroxine regulates the basal metabolic rate of body. It also regulates metabolism by stimulating protein synthesis and promotes growth of body tissues. It helps in thermoregulation by increasing heat production. It increases activity of the neurotransmitters-adrenaline and nor-adrenaline. It also supports the process of RBC production and maintenance of water and electrolyte balance. It also regulates reproductive cycles in females.

Calcitonin : It is secreted by the parafollicular or 'C' cells of the thyroid gland. It regulates the concentration of calcium and phosphorus in the blood. It is under feedback control of plasma calcium concentration and is secreted when concentration of calcium rises in the blood. It lowers concentration of calcium and phosphorus in the plasma by decreasing their release from the bones and accelerating the uptake of calcium and phosphorus by the bones.

Disorders related to thyroid gland:

a. Hyperthyroidism :

It is caused by increase in the levels of thyroid hormones. This increases the overall metabolic rate and body temperature leading to sweating, hot flushes, rapid respiration, nervous sensitivity, bulging of eye balls, and affects other physiological activities.



Fig. 9.29 Bulging eyeballs

Grave's disease (Exophthalmic goitre) : Hyperthyroidism in adults is characterised by protruding eyeballs, increased BMR and weight loss. Increased BMR produces a range of effects like increased heart beat, increased BP, higher body temperature, nervousness, irritability, tremor of fingers and bulging eyeballs.

b. Hypothyroidism : It is caused by deficiency of thyroid hormones or removal of thyroid gland (Thyroidectomy).

Cretinism : Hyposecretion in infants leads to cretinism. A **cretin** has reduced BMR and low oxidation. They are short statured because the skeleton fails to grow. They are mentally retarded. They show dry skin, thick tongue, prolonged neonatal jaundice, lethargy and constipation. This can be treated by early administration of thyroid hormones. The cretin shows stunted physical growth delayed puberty and mental retardation.

Myxoedema : It is the deficiency of thyroid hormones in adults. It is characterised by a peculiar thickening and puffiness of skin and subcutaneous tissue particularly of the face and extremities. Patient lacks alertness and intelligence, suffers from slow heart rate, low B.P., low body temperature and retarded sexual development.

Do you know ?

During pregnancy, hypothyroidism causes defective development and maturation of growing baby.

Simple goitre : (Iodine deficiency goitre) Iodine is needed for synthesis of thyroid hormone. If there is deficiency of iodine in the diet, it causes enlargement of thyroid gland leading to simple goitre. This disease is common to people in hilly areas. Addition of iodine to table salt prevents this disease. Size of the thyroid gland

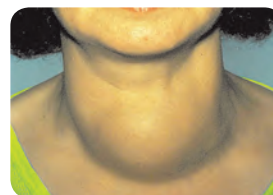


Fig. 9.30 Simple goitre

is increased but total output of thyroxine is decreased.



Find out

Why are African pygmies diminutive. Is it due to lack of GH, thyroxine or absence of thyroxine receptors on their cell surface?

E. Parathyroid gland :

Parathyroid gland is situated on the posterior surface of the lobes of thyroid gland. Parathyroids are four in number and named as superior and inferior parathyroid glands. The cells of parathyroid glands are arranged in a compact mass.

The parathyroids secrete a peptide hormone called **parathormone (PTH)**. It is also called **Collip's hormone**. It regulates calcium and phosphate balance between blood and other tissues. Release of parathormone increases blood calcium level by taking calcium from bones, increasing calcium absorption in the digestive tract and reducing loss of calcium in the urine. Secretion of parathormone is under feedback control of blood calcium level.

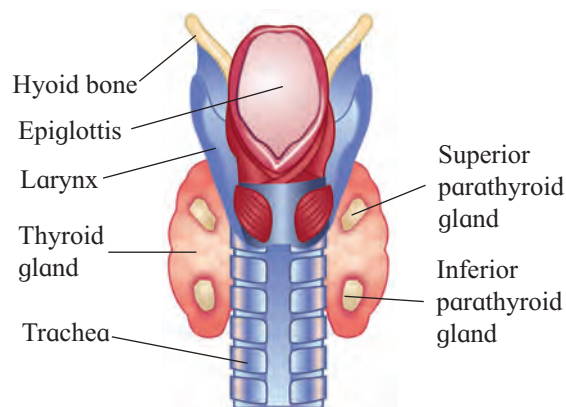


Fig. 9.31 Parathyroid gland

Concentration of calcium and phosphate is maintained by parathormone and calcitonin. These two hormones form an antagonistic pair like insulin and glucagon.

Hyposecretion of parathormone lowers the concentration of calcium in the blood. This increases excitability of nerves and muscles causing muscle twitch and spasm. This is called **parathyroid tetany** or **hypocalcaemic tetany**. Hypersecretion of parathormone is responsible for more absorption of calcium from bones i.e., demineralization of bones resulting in softening, bending and fracture of bone. This is called osteoporosis. It is common in women who have reached menopause.



Can you tell?

1. With the help of a suitable diagram describe the structure of thyroid gland.
2. How does a fall or rise in blood calcium stimulate secretion of parathyroids?

F. Thymus gland :

Thymus gland is located in the upper part of thorax on the dorsal side of the heart. It is a soft, pinkish and bilobed mass of lymphoid tissue. It is a prominent gland at birth but gets gradually atrophied in the adult, so it is called a temporary gland.

It secretes the hormone **thymosin**. It has an important role in the development of immune system by maturation of T-lymphocytes.

It also promotes production of antibodies by providing humoral immunity.

G. Adrenal gland/Suprarenal gland :

Adrenal glands have dual origin from mesoderm and ectoderm. They are located on the upper border of each kidney. Adrenal glands are small, conical yellowish glands and show two distinct regions, outer cortex and inner medulla.

1. Adrenal cortex :

The adrenal cortex is derived from embryonic mesoderm and it secretes many hormones collectively called **corticoids**. It is differentiated into three concentric regions.

a. Zona glomerulosa : It is the outer thin part of adrenal cortex and it secretes **Mineralocorticoids**. They are released for regulating sodium and potassium ion concentration. They regulate salt-water balance, blood volume and blood pressure. **Aldosterone** (salt retaining hormone) is the main mineralocorticoid. It balances Na-K levels.

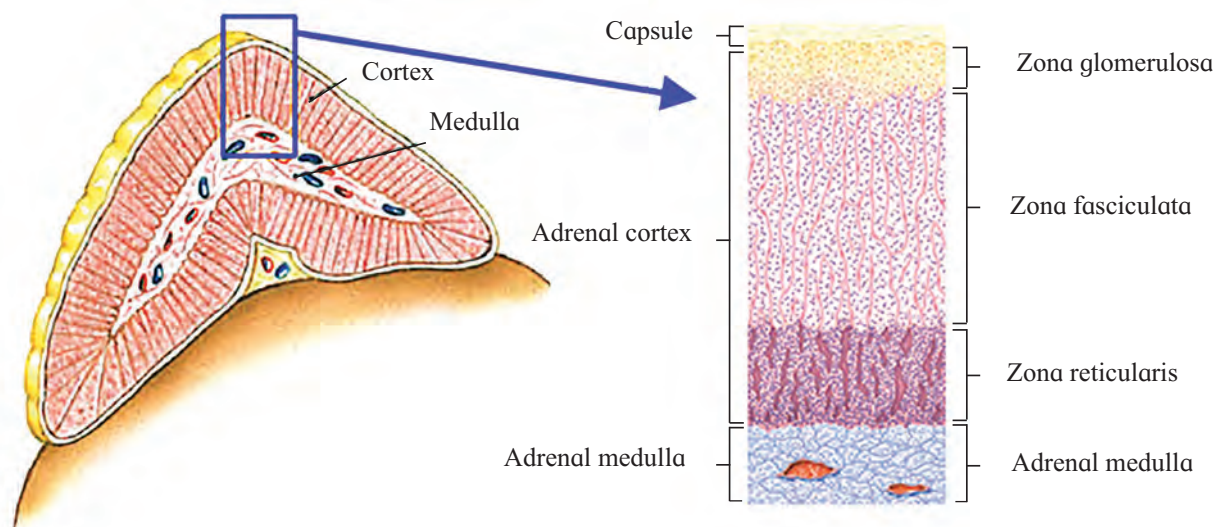


Fig. 9.32 Histology of Adrenal gland

b. zona fasciculata : It is the middle thick part and is responsible for secretion of **Glucocorticoids** like cortisol. It regulates metabolism of carbohydrates, proteins and lipids. Cortisol is an important glucocorticoid. It is responsible for increase in blood glucose level. It is also immuno suppressive. It suppresses synthesis of antibodies. So it is used in treatment of allergy. It prepares animals to face emergencies in nature.

c. Zona reticularis : It is the inner thin responsible for production of sex corticoids (**Gonadocorticoids**). In males, they have a role in development and maintenance of external sex characters. Excess of sex corticoids in a female causes adrenal virilism and hirsutism (excess hair on face) while in males it causes gynaecomastia i.e. enlarged breast. Androgens and estradiols are produced by the adrenal cortex.

Disorders related to Adrenal cortex :

a. Hyposecretion of mineralocorticoids and glucocorticoids are responsible for **Addison's disease**. Characteristic features of this disease are low blood sugar, low Na^+ and high K^+ concentration in plasma, increased loss of Na^+ and water in urine. It leads to weight loss, weakness, nausea, vomiting and diarrhoea.

b. Hyper secretion of glucocorticoids produces **Cushing's disease**. It leads to high blood sugar level, excretion of glucose in urine, rise in Na^+ , rise in blood volume, high blood pressure, obesity and wasting (gradual loss) of limb muscles.

Adrenal medulla : It develops from ectoderm. It secretes two hormones adrenaline (epinephrine) and noradrenaline (norepinephrine). **Adrenaline** also known as emergency hormone, is also called 3F hormone – (fight, flight and fright). **Noradrenaline** regulates the blood pressure under normal condition. It also acts as vasoconstrictor.

H. Pancreas :

It develops from an endoderm. It is both an exocrine (studied in digestive System) and an endocrine gland. Endocrine cells of pancreas form groups of cells called **Islets of Langerhans**. There are four types of hormone producing cells in each islet of Langerhans which secrete hormones.

- I. Alpha (α) cells (20%) secrete **glucagon**. It stimulates liver for glycogenolysis to increase blood glucose level.
- II. Beta (β) cells (70%) secrete **insulin**. It stimulates liver and muscles for glycogenesis. This lowers blood glucose level.
- III. Delta (δ) cell (5%) secrete **somatostatin** which inhibits the secretion of glucagon and insulin. It also decreases the gastric secretions, motility and absorption in digestive tract.
- IV. PP cells or F cells (5%) secrete pancreatic polypeptide (PP). It inhibits the release of pancreatic juice.

Disorders related to pancreas :

Diabetes mellitus (Hyperglycemia)

This is the most common metabolic endocrine disorder of pancreas. It leads to increase in blood glucose level. This is due to under activity of Beta cells resulting in reduced secretion of insulin. In children, such a condition is called **insulin dependent diabetes mellitus/ Type I (IDDM)** The other form of diabetes is **Non insulin dependent diabetes mellitus/ Type II (NIDDM)**. It is caused due to failure of insulin to facilitate the movement of glucose into cells. Reduced sensitivity to insulin is called **insulin resistance**.



Can you tell?

1. Distinguish between glucocorticoids and mineralcorticoids.
2. Name the hormones secreted by the adrenal cortex and state their role.
3. Write a note on Islets of Langerhans.
4. Pancreas is both exocrine as well as endocrine gland. Give reason.

In both disorders, blood glucose level increases. Some of the glucose is excreted in urine. It also causes excessive urination and dehydration of body tissues. Degradation of fats increases formation of ketone bodies (ketosis). Administration of insulin lowers blood glucose level.

I. Gonads :

Gonads are sex organs (testes and ovaries). Besides producing gametes, they also produce various hormones.

i. Ovaries : They produce the following hormones.

1. **Estrogen :** These are secreted by developing follicle. Estradiol is the main oestrogen. It is responsible for secondary sexual characters in female.
2. **Progesterone :** It is secreted by corpus luteum of the ovary after ovulation. This hormone is essential for thickening of uterine endometrium, thus preparing the uterus for implantation of fertilized ovum. It is responsible for development of mammary glands during pregnancy. It inhibits uterine contractions during pregnancy.
3. **Relaxin :** It is secreted by the corpus luteum of the ovary at the end of gestation period. It relaxes the cervix of the pregnant female and ligaments of pelvic girdle for easy birth of young one.
4. **Inhibin :** It is secreted by the corpus luteum. Inhibin inhibits the FSH and GnRH production.

ii. Testes :

Testes secrete male sex hormones called androgens such as testosterone.

Testosterone : It is secreted from interstitial cells or Leydig cells by the influence of luteinising hormone (LH). Rise in testosterone level in blood above normal inhibits LH secretion.

It is also responsible for appearance of secondary sexual characters such as facial and pubic hair, deepening of voice, broadening of shoulders, male aggressiveness, etc. It also helps in maintenance of testes.

J. Diffuse endocrine glands

Placenta :

It is the intimate connection between foetus and uterine wall of the mother for physiological exchange of the material. Placenta is a temporary endocrine gland.

During pregnancy, placenta secretes hormones such as estrogen, progesterone, hCG (Human Chorionic Gonadotropin) and human placental progesterone. These hormones check the contraction of uterine muscles and also maintain the thickness of uterine endometrium thus they help to maintain pregnancy.

Gastro intestinal tract :



Do you know ?

Presence of hCG in urine sample indicates pregnancy.

In the gastro-intestinal mucosa, certain cells are endocrine in function. These cells produce hormones which play vital role in digestive processes and flow of digestive juices.

1. **Gastrin :** It stimulates gastric glands to produce gastric juice.
2. **Secretin :** It is responsible for secretion of pancreatic juice and bile from pancreas and liver respectively.
3. **Cholecystokinin CCK and Pancreozymin PZ :** These hormones stimulate the pancreas

to release its enzymes and also stimulates gall bladder to release bile.

4. Entero-gastrone / Gastric inhibitory peptide (GIP) : It slows gastric contractions and inhibits the secretion of gastric juice.

Kidney : It produces **renin, erythropoietin** and **calcitriol** (calcitriol is the active form of vitamin cholecalciferol (D_3)).

Heart : It produces Atrial natriuretic hormone (ANH) which increases sodium excretion by kidneys and reduces blood pressure.

Hormone therapy/ HT : Hormone therapy is the use of hormones in medical treatment. HT is applied in Pregnancy, Menopause, Osteoporosis, Growth hormone deficiency, Insulin Resistance, Cancer, etc.



Can you tell?

1. Give significance of relaxin and inhibin.
2. Enlist hormones secreted by GI tract and state their role.
3. Mention the role of heart and kidney in hormone secretion.



Do you know ?

Chemicals that operate between members of the same species are social hormones or pheromones. These are commonly also called sex attractants or external hormones.

A pheromone is a volatile substance produced and discharged by an organism, which induces a physiological response in the other organisms of the same species. Pheromones are produced by many species of insects. Some pheromones enhance the chances of mating between the sexes. These are called signalling pheromones used to induce a behavioral response. Social insects such as ants make use of signalling pheromones to locate food sources and warn of danger.

Worker bees are females maintained in a sterile state by the pheromone called anti-queen factor produced by queen. The factor spreads among the workers preventing maturation of the ovaries of workers as long as the queen is present in the bee hive. Increase in colony size results in dilution of pheromones and second queen may develop.



Activity :

1. Categorise given activities into appropriate type of reflex action.

Swimming	
Dancing	
Cycling	
Salivation	
Blinking of eyes	
Sneezing	

2. Prepare concept map of mechanism of hearing.

- Name the type of hormones which bind to the DNA and alter the gene expression.
- What is the cause of abnormal elongation of long bones of arms and legs and of lower jaw.
- Name the hormone secreted by the pineal gland.
- Which endocrine gland plays important, role in improving immunity?

Q. 3 Match the organism with the type of nervous system found in them.

- | | |
|----------------|--------------|
| 1. Neurons | a. Earthworm |
| 2. Ladder type | b. Hydra |
| 3. Ganglion | c. Flatworm |
| 4. Nerve net | d. Human |

Q. 4 Very short answer questions.

- Describe the endocrine role of islets of Langerhans.
- Mention the function of testosterone?
- Give symptoms of the disease caused by hyposecretion of ADH.

Q. 5 Short answer questions

- Rakesh got hurt on his head when he fell down from his motorbike. Which inner membranes must have protected his brain? What other roles do they have to play?
- Give reason - Injury to medulla oblongata may prove fatal.
- Distinguish between the sympathetic and parasympathetic nervous system on the basis of the effect they have on:
 - Heart beat
 - Urinary Bladder
- While holding a tea cup Mr. Kothari's hands rattle. Which disorder he may be suffering from and what is the reason for this?
- List the properties of the nerve fibres.
- How does the tongue detect the sensation of taste?
- State the site of production and function of Secretin, Gastrin and Cholecystokinin.

- An adult patient suffers from low heart rate, low metabolic rate and low body temperature. He also lacks alertness, intelligence and initiative. What can be this disease? What can be its cause and care?
- Where is the pituitary gland located? Enlist the hormones secreted by anterior pituitary.
- Explain how the adrenal medulla and sympathetic nervous system function as a closely integrated system.
- Name the secretion of alpha, beta and delta cells of islets of langerhans. Explain their role.
- What are the two types of goitre? Mention their causes.
- Name the ovarian hormones and give their functions.

Q. 6 Answer the following.

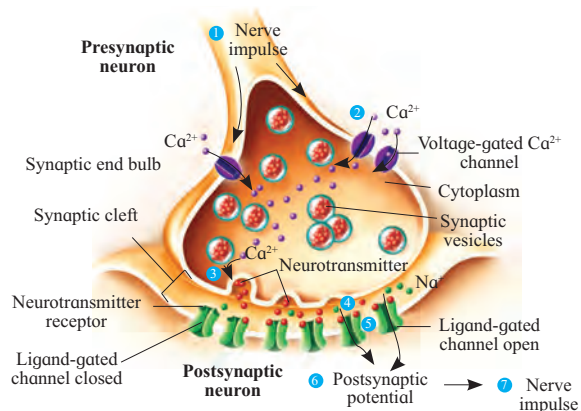
- Complete the table.

Location	Cell Type	Function
PNS	Produce myelin sheath
PNS	Satellite cells
.....	Oligodendrocytes	Form myelin sheath around central axon
CNS	Phagocytose pathogens
CNS	Form the epithelial lining of brain cavities and central canal.

Q. 7 Long answer questions.

- Explain the process of conduction of nerve impulses upto development of action potential.
- Draw the neat labelled diagrams of :
 - Human ear
 - Sectional view of human eye
 - L. S. of human brain
 - Multipolar Neuron

3. Answer the questions after observing the diagram given below.



- What do the synaptic vesicles contain?
 - What process is used to release the neurotransmitter?
 - What should be conditions for the next impulse to be conducted?
 - Will the impulse be carried by post-synaptic membrane even if one pre-synaptic neuron is there?
 - Can you name the channel responsible for their transmission?
- Explain the Reflex Pathway with the help of a neat labelled diagram.
 - Krishna was going to school and on the way he saw a major bus accident. His heart beats increased and his hands and feet became cold. Name the part of the nervous system that had a role to play in this reaction.
 - What will be the effect of thyroid gland atrophy on the human body?
- Write the names of hormones and the glands secreting them for the regulation of following functions.
 - Growth of thyroid and secretion of thyroxine.
 - Helps in relaxing pubic ligaments to facilitate easy birth of young ones.
 - Stimulate intestinal glands to secrete interstitial juice.
 - Controls calcium level in the blood
 - Controls tubular absorption of water in kidneys.
 - Urinary elimination of water.
 - Sodium and potassium ion metabolism.
 - Basal Metabolic rate.
 - Uterine contraction.
 - Heart beat and blood pressure.
 - Secretion of growth hormone.
 - Maturation of Graafian follicle.
 - Explain the role of hypothalamus and pituitary as a coordinated unit in maintaining homeostasis?
 - What is adenohipophysis? Name the hormones secreted by it?
 - Describe in brief, an account of disorders of adrenal gland.
 - Explain action of steroid hormones and proteinous hormones.
 - Describe in brief an account of disorders of the thyroid.

Project : Prepare animated powerpoint presentation to explain mechanisms of hormonal action.



Can you recall?

1. Generally individuals are conscious about their health. So define health.
2. Define infectious and non infectious disease? Give their examples.

Health does not simply mean ‘absence of disease’ or physical fitness. In fact, it is difficult to aptly define health. According to the World Health Organization, health is defined as the state of complete physical, mental and social well-being and not merely the absence of disease or infirmity. In short, health is birth right of humans. Health also reflects the metabolic and functional efficiency of living organisms. Hygiene is a science of rules of health. To achieve good health, it is therefore, necessary that we have hygienic balanced diet, clean drinking water, personal and community hygiene, regular physical exercise, knowledge about diseases and their effect on body, proper disposal of waste and control of vectors.

Everyday we are exposed to various foreign bodies, including infectious agents like bacteria, viruses, etc. Despite constant exposure to variety of pathogens, most of us remain healthy. This is due to fact that the human body has ability to resist almost all type of these foreign bodies. The system which protects us from various infectious agents, is called **immune system**. Resistance is the ability to prevent the damage or disease, through our defense mechanism.

10.1 Immunity:

The term ‘immunity’ has traditionally referred to as the resistance exhibited by the host towards injury caused by pathogens and their products. However, protection against infectious diseases is only one of the many

consequences of immune response, which is entirely concerned with the reaction of the body against any foreign **antigen**.

Immunity is in fact the “freedom” or “exempt”. The concept of immunity is believed to be started by **Edward Jenner** in England. He developed cowpox vaccine for the protection against the attack of small pox (virus). **Immunology** is a branch of science which deals with the study of immune system, immune responses to foreign substances and their role in resisting infection by pathogens.

The most important characteristic of immune system is that it distinguishes self (body’s own cells) and non-self (foreign molecules or invading cells). So, the immune system differentiates between the body cells and the invaders. Any foreign substance invading body and capable of stimulating an immune response, is called an **antigen**. The protective chemicals produced by immune cells in response to antigens are called **antibodies**.

A. Types of immunity:

There are two types of immunity as Innate or Inborn (inherited) immunity and Acquired or Adaptive immunity.

i. Innate immunity or Inborn immunity:

Innate immunity is the resistance to infections that an individual possesses by virtue of his or her genetic make-up. It is the natural (inborn) defense system of the body. It is not affected by prior contact with microorganisms or immunization. It is nonspecific, when it indicates a degree of resistance to infection in general, or specific where resistance to a particular pathogen is concerned. One form of innate immunity comprises the various types of barriers which prevent entry of foreign agents into the body.

a. Epithelial surface :

The intact skin and mucous covering the body, protect it considerably against invasion by microorganism(s).

The healthy skin possesses bactericidal activity due to the presence of high concentrations of salt in drying sweat. Sebaceous secretions and long chain of fatty acids have bactericidal and fungicidal properties. The mucosa of the respiratory tract has several innate mechanisms of defense. The nose prevents entry of microorganisms to a large extent, the inhaled particles being arrested through hair at or near the nasal orifices. Those that pass beyond are held by mucus lining the epithelium and are swept back to pharynx where they tend to be swallowed or coughed out.

The cough reflex is an important defence mechanism of respiratory tract. The mouth is constantly bathed in saliva which has inhibitory effect on microorganisms. The acidity of gastric secretions in the stomach destroys microorganisms. The flushing action of urine eliminates bacteria from the urethra. Spermine and zinc present in semen are antibacterial.

Knowledge Enhancer:

The conjunctiva is freed of foreign particles by the flushing action of lachrymal secretions. Eyes become susceptible to infection when lachrymal secretion is absent. Tears contain the antibacterial substance Lysozyme (1st described by A. Fleming). This is a thermolabile, low molecular weight, basic protein which acts as a muramidase. Lysozyme is active only against some pathogenic Gram +ve bacteria.

b. Antimicrobial substances in blood and tissues:

The complement system contains more than 30 serum proteins, circulating in the blood in an inactive state. The presence of microbial

pathogens activates the 'Complement cascade' to eliminate pathogens. The interferons are a class of cytokines (soluble proteins) released by cells infected with viruses and certain white blood cells to stimulate other cells to protect themselves from viral infection.

c. Cellular factors in innate immunity :

Natural defence against the invasion of blood and tissues by microorganisms and other foreign particles, is mediated to a large extent by phagocytic cells which ingest and destroy them. Phagocytic cells (discovered by Metchnikoff in 1882) are grouped as microphages and macrophages. They remove foreign particles that enter the body. A class of lymphocytes called Natural killer (NK) cells is important in nonspecific defence against viral infections and tumors.

d. Fever :

Increase in the body temperature following the infection is a natural defense mechanism. It helps to accelerate the physiological processes to destroy the invading pathogens. Fever stimulates the production of interferon and helps in recovery from viral infections.

e. Acute phase proteins (APPs):

Infection or injury leads to a sudden increase in concentration of certain plasma proteins, collectively called acute phase proteins. These include C Reactive Protein (CRP), Mannose binding protein, Alpha-1-acid glycoprotein, Serum Amyloid P, etc. APPs are believed to enhance host resistance, prevent tissue injury and promote repair of inflammatory lesions.

ii. Acquired immunity:

The resistance that an individual acquires during life is known as "Acquired immunity". It is also known as Adaptive or Specific immunity". It involves the formation of antibodies in the body, which neutralize the antigens. Acquired or Adaptive immunity has the following unique features.

- a. **Specificity:** It can produce specific antibody or T-lymphocyte against a particular antigen/pathogen.
- b. **Diversity:** It can recognize a vast variety of diverse pathogens or foreign molecules.
- c. **Discrimination between self and non-self:** It differentiates between own body cells (self) and foreign (non-self) molecules.
- d. **Memory:** When the immune system encounters a specific foreign agent for the first time, it generates an immune response and eliminates the invader. This is called first encounter. The immune system retains the memory of the first encounter. As a result, a second encounter with same pathogen brings about quicker and stronger immune response.

Types of Acquired Immunity:

Acquired immunity is of two types Active and Passive.

a. Active immunity:

It is the resistance developed by individuals as a result of an antigenic stimulus. It also known as “Adaptive immunity”. Active immunity may be natural or artificial.

1. **Natural Acquired Active immunity:** Immunity acquired due to infection is called natural active immunity. It is developed after entry of pathogens in the body. It is long-lasting immunity. e.g. person who has recovered from attack of measles develops natural acquired active immunity to measles, for the life time.
2. **Artificial Acquired Active immunity:** It is the resistance induced by vaccines. Vaccine is introduced into the body to stimulate the formation of antibodies by the immune system. e.g. Polio vaccine, BCG vaccine etc. such immunity may be temporary or permanent.

b. Passive immunity:

Passive immunity is acquired when ready-made antibodies are received by the body cells. i.e. Body cells do not take any active part in the production of immunity. Passive immunity can be acquired either naturally or artificially.

1. Natural Acquired Passive immunity:

Before birth maternal antibodies are transferred from mother to foetus through placenta. After birth, antibodies are transferred from mother to infant through colostrum (first milk of mother) and continue throughout the period of breast feeding. The antibodies received by baby from mother remain in the body for a short time. Therefore, natural acquired passive immunity is short lived.

2. Artificially Acquired Passive immunity:

This immunity is developed by injecting previously prepared antibodies using serum from humans or animals. For e.g. Antibodies obtained from hyper immunised horses are injected to humans against rabies pathogens. It is short lived.

B. Cells of Immune System:

There are two main types of cells involved in the working of Immune system. They are (a) Lymphocytes and (b) Antigen Presenting cells.

a. Lymphocytes:

Lymphocytes are the main cells of the immune system. They, like the other blood corpuscles, arise from the stem cells, the haemocytoblasts, present in liver of the foetus and in the bone marrow in adult. Some of them undergo differentiation in the gut – associated bursal lymphoid tissues (Tonsils, Peyer’s patches) and are called **Bursal** or **B-lymphocytes**; others are differentiated in the thymus gland and are termed as **T-lymphocytes**. The mature lymphocytes pass into body fluids (blood and lymph) and circulate in the body.

Many of them stay in the lymph nodes. The B-lymphocytes and T-lymphocytes form humoral or antibody-mediated immune system (AMIS) and cell-mediated immune system (CMIS) respectively. Both the immune systems need antigens to come into action, but they respond in different ways.

Mechanism of response of T-lymphocytes to antigens: On coming in contact with an antigen, a T-lymphocyte forms clones of T-cells which are similar but they perform different functions. The clone has four types of T-lymphocytes:

- i. **Helper T-cells:** Sensitized helper T-cells produce lymphokines for performing several types of functions like proliferation of other T-cells, stimulation of B-lymphocytes, macrophages, etc.
- ii. **Killer T-cells or Cytotoxic T-cells:** They directly attack and destroy invading microbes, infected body cells and cancer cells. Killer T-cells bind to infected cell and secrete perforins. Then perforins form a hole in infected cell. It also releases substances that kill the cell, hence the name cytotoxic T-cell.
- iii. **Suppressor T-cells:** These cells suppress entire immune system against attack on the own body cells.
- iv. **Memory T-cells:** These are previously sensitized cells which retain the sensitization memory for long time in the future.

Mechanism of action of B-lymphocytes to antigens:

B-lymphocytes are sensitized directly by both antigens as well as by helper T-cells. Activated B-lymphocyte multiplies very fast to produce clone of plasma cells and memory B-cells. The plasma cells produce specialized glycoproteins, called antibodies which are circulated through body fluids (humor) like blood and lymph. The antibody molecules may bind to a cell membrane or they remain free.

The free antibodies have three main functions:

- i. **Agglutination** of particulate matter, including bacteria and viruses. The immobilized mass is then engulfed by phagocytes.
- ii. **Opsonisation** or coating of bacteria to facilitate their subsequent phagocytosis by macrophages.
- iii. **Neutralization** of toxins released by bacteria e.g. tetanus toxin.

Each antibody is specific for a particular antigen.

b. Antigen Presenting cells:

Antigen presenting cells engulf invading pathogens and process the antigens. Then the processed antigens are presented on their own surface. These cells are able to deliver a stimulatory signal that is necessary for activation of helper T-cell.



Can you tell?

Which is kind of immunity provided by vaccination?



Can you recall?

1. Why are vaccines considered as antigen containing material?
2. How are vaccines produced?
3. Who was Edward Jenner?

C. Vaccination :

Administration of vaccine (i.e. inactivated pathogen or antigenic protection of particular pathogen) to protect against a particular pathogen, is called vaccination.

The body's immune system helps to protect against pathogens that cause infection. It's an efficient system, most of the time,. It either keeps microorganisms out or tracks them down and gets rid of them. However, some pathogens can overwhelm the immune system. When this happens, it can cause serious illness. The pathogens most likely to cause problems, are the ones the body doesn't recognize.

Vaccination is a way to “teach” the immune system as to how to recognize and eliminate pathogenic organism. That way, body is always prepared if you are ever exposed.

Vaccination is an important form of primary prevention, that can protect people from getting sick. Vaccination has allowed us to control diseases like measles, polio, tetanus and whooping cough that once threatened many lives. It’s important that as many people as possible get vaccinated. Vaccinations don’t just protect individuals, when enough people are vaccinated. It also helps to protect the society.



Do you know ?

How does vaccination work?

Healthy immune system defends against invaders. Immune system consists of several types of cells. These cells defend against harmful pathogens. However, they have to recognize an invader. Vaccination teaches the body to recognize new pathogens causing diseases. It stimulates the body to make antibodies against antigens of pathogens. It also primes immune cells to remember the types of antigens. This allows a faster response to the pathogen in future.

Vaccines work by exposing you to a safe version of pathogen. Vaccines may be in the form of:

- A protein or sugar from the pathogen.
- A dead or inactivated form of a pathogen.
- A toxoid containing toxin produced by a pathogen.
- A weakened (attenuated) pathogen.

When the body responds to the vaccine, it builds an adaptive immune response. This helps to equip the body to fight off an actual infection.



Internet my friend

Find out different types of vaccines available in the market and their significance.

Vaccinations are safe:

Vaccines are considered to be safe. They are rigorously tested and go through many rounds of study, examination, and research before they are used for the general public.

Extensive research and evidence shows that vaccines are safe their side effects are rare and typically mild.



Internet my friend

1. Whether vaccination can be done during or before pregnancy?
2. Will it be helpful to protect the mother and baby both?
3. Which vaccines can be administered before pregnancy? When can it be?
4. How will you increase awareness in the society on this issue?

10.2 Structure of Antibody:

Antibodies are glycoproteins which are highly specific to specific antigens. They are also known as **Immunoglobulins** (Igs), produced in response to antigenic stimulation. Antibodies are produced by plasma cells which in turn are formed by B-lymphocytes. The mature plasma cells produce antibodies at an extremely rapid rate i.e. about 2000 molecules per second.

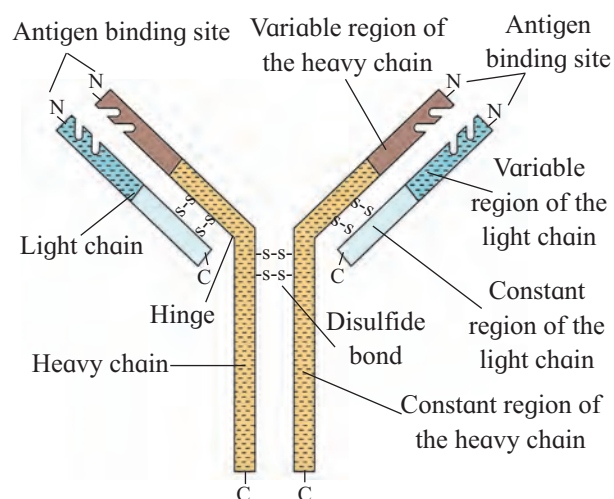


Fig. 10.1 : Structure of antibody

Structure:

Antibody is a 'Y' shaped molecule. Each immunoglobulin molecule is made up of four polypeptide chains. There are two heavy or **H-chains** and two light or **L-chains**. The four polypeptide chains are held together by **disulfide bonds** (-s-s-) to form a 'Y' shaped structure. The region holding together arms and stem of antibody, is termed as hinge. Each chain of the antibody includes two distinct regions, the variable region and the constant region. **Variable regions** constitute the antigen-binding site (**paratope**). This part of antibody recognizes and binds to the specific antigen to form an antigen-antibody complex. Since most antibodies carry two antigen binding sites, they are said to be bivalent.



Find out

Collect information about IgG, IgA, IgM, IgD and IgE antibodies from internet / reference book / teacher and prepare a chart / power point presentation.

Formation of antigen-antibody complex:

Study of antigen-antibody interactions is called **serology**. Each antibody is specific for a particular antigen. Combining sites of antigen, called antigenic determinants (**epitopes**) react

with the corresponding antigen binding sites of antibodies called **paratopes**. The antigen binding sites (paratopes) are located on the variable regions of the antibody. Small variations in the variable regions make each antibody highly specific for a particular antigen. The variable region enables the antibody to recognize the specific antigen and bind to specific antigen in a lock and key manner forming an antigen-antibody complex.

Antigen on Blood Cells:

There are several known antigens on the surface of human red blood cells. These antigens give rise to different blood groups. There are many genetically determined blood groups system like ABO, Rh, Duffy, Kidd, Lewis, P, MNS, Bombay blood group, etc.

ABO Blood Groups:

The A, B and O blood groups were discovered by Karl Landsteiner in 1900. Later on, the blood group AB was discovered by Landsteiner's students Decastello and Sturli in 1902. Landsteiner was awarded the Nobel prize for his discovery of human blood groups. He found two antigens or agglutinogens on the surface of human red blood cells and named them as antigen A and antigen B. He also noticed the corresponding antibodies or agglutinins in the serum called 'a' and 'b'.

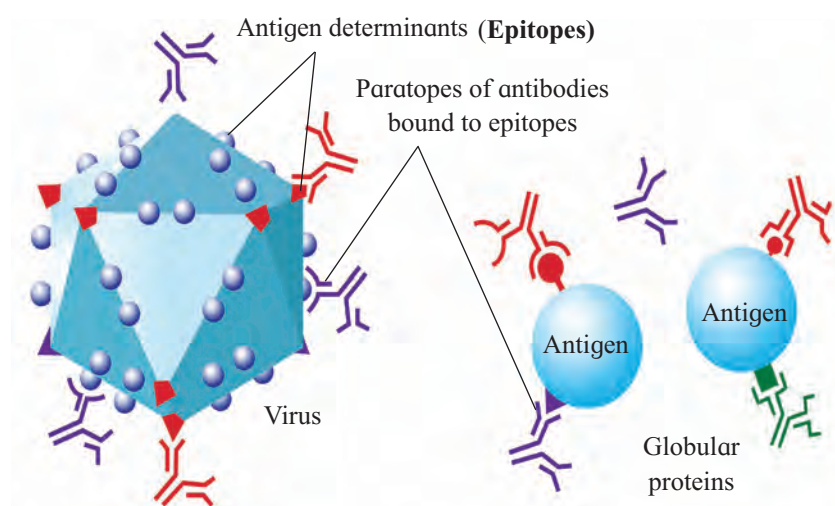


Fig. 10.2 : Formation of antigen- antibody complex

Table 10.3 : ABO Blood groups in man

Blood group	Genotype	Antigen on Surface of RBC	Antibody in Serum	Can donate blood to	Can receive blood from
A	$I^A I^A$ or $I^A I^O$	A	Antibody b	A, AB	A, O
B	$I^B I^B$ or $I^B I^O$	B	Antibody a	B, AB	B, O
AB (universal acceptor)	$I^A I^B$	A and B	Nil	AB	A, B, AB, O
O (universal donor)	$I^O I^O$	Nil	Both Antibody a and Antibody b	A, B, AB, O	O

In ABO system, the blood groups are determined by the presence or absence of antigen A and antigen B, the blood group of person is classified into four groups A, B, AB and O.

Blood group A : Individuals, with blood group ‘A’ have the antigen A on the surface of their red blood cells (RBCs) and antibody ‘b’ in their plasma.

Blood group B : Individuals with blood group ‘B’ have the antigen B on the surface of their RBCs and antibodies ‘a’ in their plasma.

Blood group AB : Individuals with blood group ‘AB’ have both antigens A and B on the surface of their RBCs and no antibodies in their plasma.

Blood group O – Individuals with blood group ‘O’ lack both antigens A and B on the surface of their RBCs and show presence of both ‘a’ and ‘b’ antibodies in their plasma.

Rh factor:

Rh is the most complex of the blood group system. Rh factor is an antigenic protein present on the surface of the red blood cells in the human beings. It was first discovered by Landsteiner and Wiener (1940), on the surface of RBCs of Rhesus monkey, so it is called Rh factor (also called D antigen). Person having Rh factor (D antigen) are called Rh positive (Rh +ve) and those lacking D antigen are called Rh negative (Rh -ve).

Rh (D) antigen induces a strong immunogenic response when introduced into Rh-ve individuals. Rh blood group is an important factor in blood transfusion and is involved in haemolytic diseases of the newborn (HDN), which is called **erythroblastosis foetalis** (destruction of the erythrocytes of the foetus). It occurs when an Rh -ve mother conceives Rh+ve foetus.

The Rh +ve RBCs from the foetus may enter the mother’s circulatory system during child birth, causing her to produce anti-Rh antibodies. As a result, subsequent Rh+ve fetuses will be exposed to the anti-Rh antibodies produced by mother, which result in HDN. In order to prevent HDN, Rh -ve mother is injected with the anti-Rh antibody during all pregnancies carrying Rh +ve foetus.



Can you tell?

1. The blood group of Krutika is O Rh +ve. What would be the possible blood groups of her parents?
2. Mrunmayi is called as universal blood acceptor. What is her blood group?



Use your brain power

Can a person with blood group O Rh+ve donate blood to a patient with blood group O Rh-ve? Why?

10.3 Common Human Diseases:

Disease is defined as condition of disturbed or deranged functioning of one or more organs or organ systems of the body, caused due to infections, defective diet or heredity. All human diseases can be broadly categorized into **congenital** diseases and **acquired** diseases.

Congenital diseases are present from birth; may be caused by genetic abnormality or metabolic disorder. They may be permanent and were practically incurable. However, modern research has helped to cure some inborn diseases through gene therapy, enzyme replacement therapy, etc.

Acquired diseases develop after the birth and can be subdivided into (a) Communicable or infectious diseases or (b) Non- Communicable or Non-infectious diseases.

The diseases which are transmitted from infected person to another healthy person either directly or indirectly, are known as **Communicable** or Infectious diseases. Malaria and other diseases which are to be studied in this chapter, are examples of this type.

The diseases that cannot be transmitted from infected person to another healthy one either directly or indirectly are known as **Non- Communicable** or Non - Infectious diseases. Cancer and deficiency diseases are examples of this type.

Communicable diseases are caused by pathogens like viruses, bacteria, fungi, helminth worms, etc. All the disease causing organisms are called '**Pathogens**'.

Many pathogens use another organism, the '**Vector**' to reach us. These vectors are actually parasites which we regularly come across. **Parasite** is an organism that lives in or on the body of another organism and derives its nutrition from that of host organism. Parasites are of two categories viz. : 1. Ectoparasite e.g. bedbug 2. Endoparasite. e.g. *Plasmodium*.

A. Malaria :

It is a vector borne infectious disease caused by the protist, the *Plasmodium*.

There are four species of *Plasmodium* as *P. vivax*, *P. ovale*, *P. malariae*, *P. falciparum*. Only *P. falciparum* causes serious illness while others are rarely fatal. Based on the species, there are 4 - types of malaria.

Signs and symptoms of malaria :-

Symptoms of malaria begin to appear about 7 to 15 days after the bite of infective mosquito.

- Initial symptom are fever, headache, and chills, may be difficult to recognize as malaria.
- Classical symptoms of malaria is cyclic occurrence of high fever followed by sweating and sudden shivering. Such entire episode lasts for four to six hours and recurs every two days or three days.
- Vomiting and convulsions.
- Arthralgia (joint pain), anaemia due to rupturing of RBCs.
- Haemoglobinuria, hepatomegaly (liver enlargement).
- Retinal damage (eye).
- Cerebral malaria (brain infection).

Table 10.4 : Table showing different species of *Plasmodium*

Species of <i>Plasmodium</i>	Incubation period	Pattern of high fever
<i>P. vivax</i>	14 days	High fever after 48 hr interval
<i>P. malariae</i>	28 days	High fever after 72 hr interval
<i>P. ovale</i>	17 days	High fever after 48 hr interval
<i>P. falciparum</i>	12days	High fever at irregular intervals between 22-48 hrs

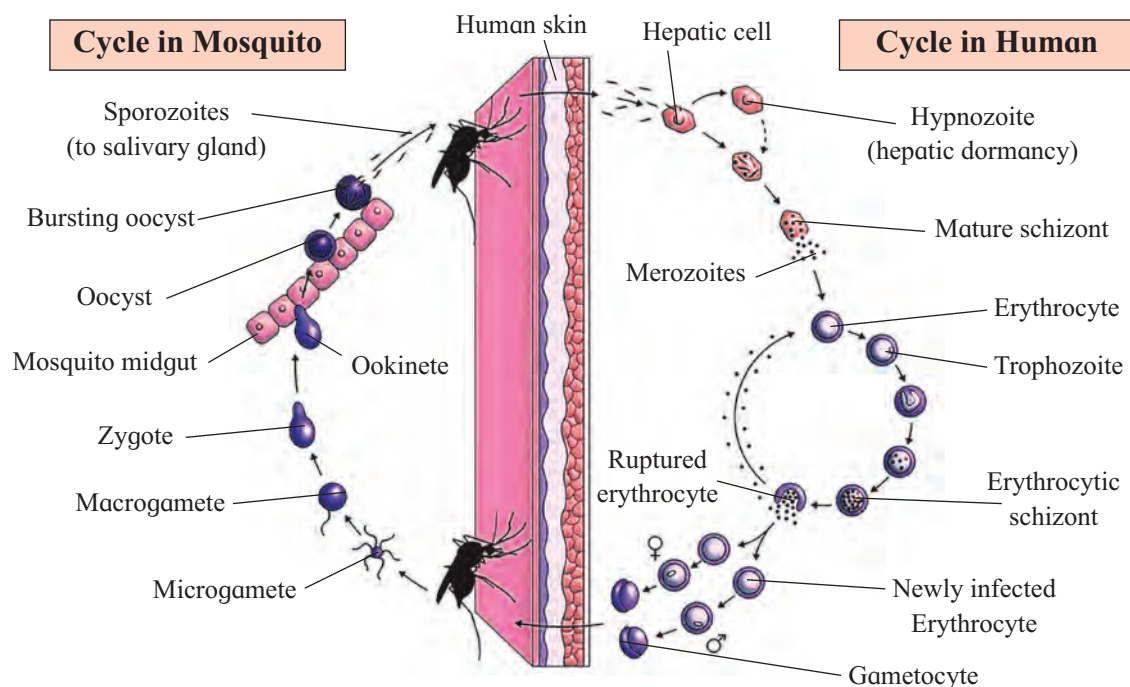


Fig. 10.5 : Stages in the life cycle of *Plasmodium* in mosquito and human

Mode of transmission :

Plasmodium is transmitted from one person to other through an insect vector- female *Anopheles* mosquito.

When infected female *Anopheles* sucks the human blood, it may transfer sporozoites to human circulation. Sporozoites reproduce asexually through fission (schizogony) in the liver cells or erythrocytes. The cells formed are now called merozoites. The cells formed within erythrocytes function as gametocytes (gamogony). Besides, it forms gametocytes within erythrocytes (gamogony).

Gametocytes if taken up by female *Anopheles*, fertilization occurs in its gut. Diploid zygote is formed which transforms into oocyst. Oocyst forms large number of haploid sporozoites through meiosis (sporogony). Sporozoites migrate to salivary glands and are ready to infect new human host.

Diagnosis and Treatment :

Malaria can be diagnosed by microscopic study of blood smear. Besides, other rapid diagnostic tests based on nucleic acid amplification techniques are also used.

Treatment of malaria includes Artemisinin based combination therapies (ACTs). WHO has recommended 5 different ACTs which includes various combinations of artesunate, sulfadoxine, pyrimethamine, etc. In addition, quinine is also used.

Prevention and Control :

1. Prevention of mosquito bite by using mosquito nets and insect repellents.
2. Spraying insecticides and draining stagnant water where mosquito lays eggs.
3. Mosquito larvae can be controlled by using *Gambusia* fresh water fish (biocontrol).



Always Remember

The time interval from the invasion of a pathogen to the development of clinical manifestations, is known as Incubation period.

B. Amoebiasis :

- Amoebiasis is also known as Amoebic dysentery. It's a common infection of human gastro-intestinal tract, which affects 15 % population of India.

- Amoebiasis is caused by a protist parasite- *Entamoeba histolytica*.

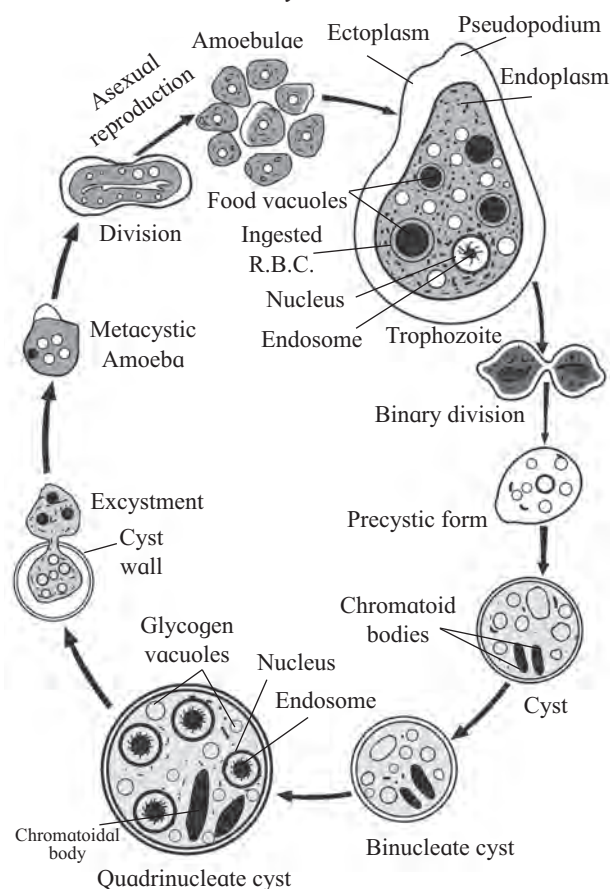


Fig. 10.6 : Life cycle of *Entamoeba histolytica*

Signs and symptoms:

- Diarrhoea, flatulence, stool with mucus and abdominal pains (cramps) are common.
- Passing of blood with stool is common in severe cases.
- Hepatomegaly occurs if parasite enters the liver. Liver develops amoebic liver abscess accompanied with fever and pain in right abdomen.

Mode of transmission :

- Faeco-oral route.
- Eating with dirty hands.
- Contaminated food and water.

Diagnosis and Treatment : Diagnosis of amoebiasis is made through microscopic examination of the stool sample.

Amoebiasis is treated by the use of Metronidazole and Tinidazole which can destroy the *E. histolytica* in the digestive tract as well as other tissues.

Prevention and Control : Wash hands with hot water and soap after using toilets and changing baby's diaper.

Drink boiled water. Otherwise, water must be chlorinated and filtered. Avoid eating unhygienic food. Vegetables must be properly washed and cooked. Proper sanitary facilities including sewage disposal help in prevention.



Internet my friend

Gather information about trophozoite and its occurrence in the life cycle of other parasitic protozoans.

C. Ascariasis :

It is an infectious disease of human intestinal tract, caused by roundworm- *Ascaris lumbricoides*. *Ascaris lumbricoides* is an endoparasitic round worm or nematode.

Signs and symptoms :

- Gastro-intestinal discomfort accompanied with vomiting and fever.
- Presence of live worms in fecal matter.
- Pulmonary disorders occur in some patients. Pneumonitis (inflammation in alveolar wall).
- Loss of appetite and weight loss.
- Eosinophilia (number of eosinophils is increased).

Mode of transmission: Food and drinks contaminated with the eggs of these worm is the main mode of transmission. Eggs hatch inside the intestine of the new host. The larvae pass through various organs and settle as adults in the digestive system.

Diagnosis and Treatment: Diagnosis can be done by microscopic examination of the stool.

Anti-helminthic drugs like Piperazine, Mebendazole, Levamisole, Pyrantel are effective against *Ascaris lumbricoides*.

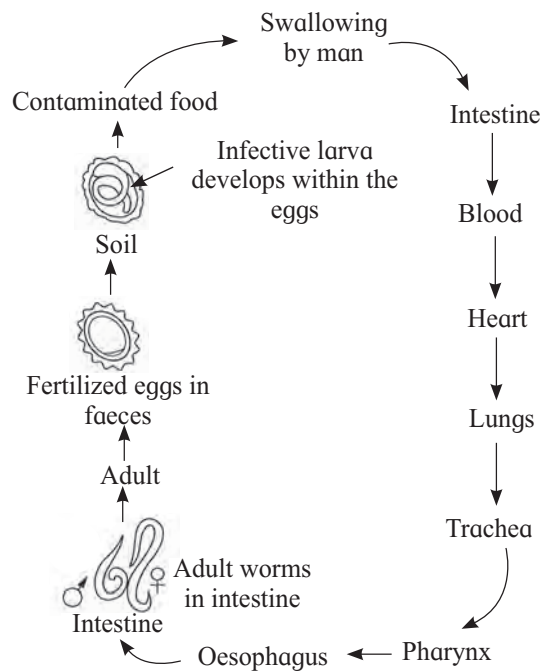


Fig. 10.7 : Life cycle of *Ascaris lumbricoides*

Prevention and Control : Avoiding defaecation in open space, prevents the spread of *Ascaris*. Personal hygienic habits like washing hand with water and soap after using toilet are also important. Washing vegetables thoroughly before cooking and avoiding raw vegetables is important for prevention of ascariasis.

D. Filariasis/ Elephantiasis :

Filariasis is caused by thread like worms-nematodes. These nematode parasites are transported from person to person via mosquito bite. Filariasis can be divided into 3 subtypes as 1) Lymphatic Filariasis, 2) Subcutaneous Filariasis (e.g. *Loa loa*, *Mansonella* spp.) and 3) Serous (abdominal)cavity Filariasis (e.g. *Mansonella* spp.).

Lymphatic Filariasis (Elephantiasis) is caused by the worms - *Wuchereria bancrofti*, *Brugia malayi*, *Brugia timori*.

Signs and symptoms :-

- Edema with thickening of skin and underlying tissue.
- *Wuchereria bancrofti* affects the legs, arms, breasts, scrotum, etc.
- In lymphatic filariasis, worms infect lymphatic system and causes enlargement of lymph vessels and nodes. This is elephantiasis i.e. limbs are swollen like legs of elephant.
- Lymphedema i.e. accumulation of lymph fluid in tissue causing swelling.

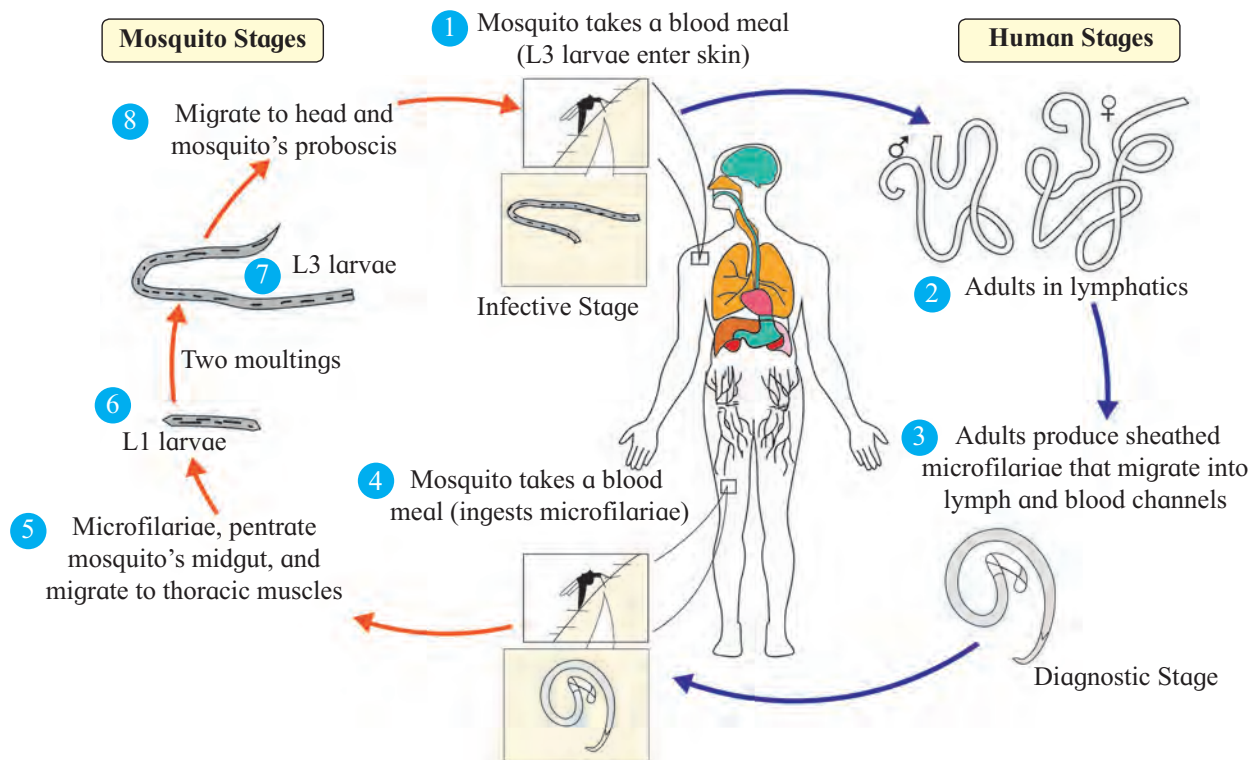


Fig. 10.8 : Mode of transmission - Filariasis (*Wuchereria bancrofti*)

- Hydrocele i.e. testis are enlarged due to accumulation of lymphatic fluid in testis.

Mode of transmission : *Wuchereria bancrofti* is transmitted to human body by female *Culex* mosquito.

The larvae escape mosquito body and arrive on the human skin. They penetrate the skin, undergo two moultings before they become adults and settle in the lymphatic system. Incubation period can be as long as 8-16 months.

Diagnosis and Treatment : Use of diethyl-carbamazine citrate twice a day for three weeks and thereafter for five days every six month is effective against filarial worms.

Prevention and Control : Avoid mosquito bite by using mosquito nets and insect repellents. Eradication of mosquitoes is essential for control of filariasis.

E. Typhoid :

Typhoid is an acute infection of intestine. It is caused by *Salmonella typhi*. It is Gram -ve bacteria found in intestinal lumen of infected person. Pathogenicity is due to “O”- antigen, a lipopolysaccharide (LPS), present on surface coat. Flagella contains “H”- antigen.

Signs and Symptoms :

- Prolonged fever as high as 104°F.
- General nausea, fatigue, headache.
- Abdominal pain, constipation or diarrhoea.
- Rose-coloured rash on skin.
- White coat on tongue, cough.
- Anorexia (loss of appetite).
- If not treated- breathlessness, irregular heartbeats, haemorrhage.

Mode of transmission :

- It is a food and water borne disease.
- Insects like housefly and cockroaches feeding on fecal matter, may transfer the bacteria to food material.
- Poor hygiene habits and poor sanitation conditions are responsible for the spread of typhoid.

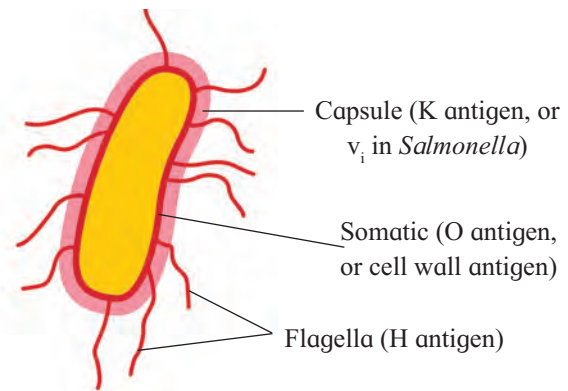


Fig. 10.9 : *Salmonella typhi*

Diagnosis and Treatment :- Widal test is used for diagnosis of typhoid. Treatment of typhoid involves surgical removal of gall bladder in severe cases. Antibiotics like Chloromycetin is helpful treatment.

For prevention of typhoid WHO recommends two vaccines as oral (Ty21a vaccine) and injectable (Typhoid polysaccharide vaccine) sold as- typhim vi and typherix.

F. Pneumonia :

Pneumonia is an inflammatory condition of lungs or alveoli of lungs. It is caused by a variety of pathogens which may be viruses like influenza virus, adenovirus, para influenza and Respiratory Syncytial Virus (RSV) or bacteria like *Streptococcus pneumoniae* or fungal pathogens e.g. *Pneumocystis jirovecii* and *Pneumocystis carinii*. Pneumonia can also be caused by chemical burns or physical injury to lungs.

Signs and Symptoms :-

- Cough produces yellow or greenish sputum or phlegm.
- High fever.
- Shortness of breath (Dyspnea).
- Chest pain during deep breath or coughing.
- Loss of appetite, fatigue, headaches, vomiting, joint pains and muscle aches.

Mode of transmission : Pneumonia mostly spreads by direct person to person contact.

It can also spread via droplets released by infected person or even by using shared clothes and utensils.

Diagnosis and Treatment :- Course of treatment depends upon pathogen leading to the disease. For bacterial pneumonia, antibiotics like Benzyl penicillin, Ampicillin and Chloramphenicol are effective.

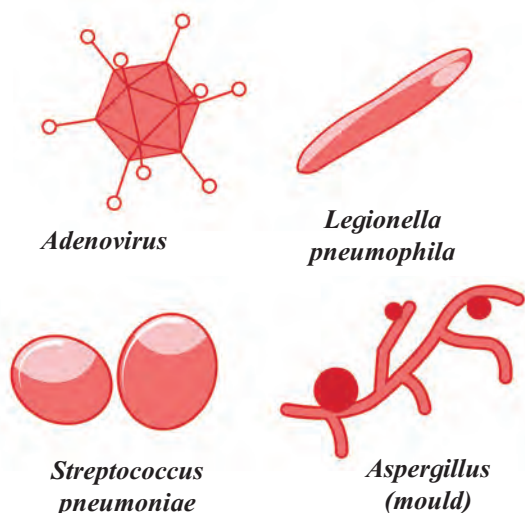


Fig. 10.10 : Pneumonia -Infectious agents

Prevention and Control :-

- Vaccination is important prevention in both children and adults.
- Vaccines against *Haemophilus influenzae* and *Streptococcus pneumoniae* in first year of life, help greatly to reduce the chances of causing Pneumonia.



Internet my friend

Find out other modes of infection by which a pathogen spreads from person to person.

G. Common Cold :

It is a viral infectious disease of upper respiratory region. It is also known as **nasopharyngitis**, acute viral rhinopharyngitis, acute coryza or a cold .

It is caused by a group of viruses known as Rhinoviruses and Coronaviruses.

Signs and Symptoms :

- Cough , sore throat, running nose and fever.
- Nasal congestion, sneezing.
- Conjunctivitis (red eyes)
- Muscle rashes, fatigue, headache, shivering and loss of appetite.

Prevention and Control :

- Staying away from person suffering from common cold.
- Washing hands with soap and water.
- Use of handkerchief to cover the nose and mouth during coughing and sneezing.
- Alcohol based hand sanitizer can also be used.



Can you tell?

Why do we suffer from common cold repetatively in our life, but other viral diseases like Influenza or Small pox only once?

H. Ring Worm (Dermatophytosis) :

It is fungal infection of skin. It is caused by many fungal species belonging to the genera *Trichophyton* and *Microsporum*. These fungi feed on keratin in skin, hair and nails.

Signs and Symptoms :

- Infected skin shows enlarged, red ring caused due to **ringworm**.
- Appearance of dry, scaly lesions on various parts of the body. These red patches cause intense itching.
- Infection to nails is termed as **onychomycosis**, in which nails become thick, discoloured and disfigured.
- Athlete's foot is the fungal infection that usually begins between the toes.

Mode of transmission :- Ringworm spreads by sharing of clothes, comb of infected person, etc. Close contact with infected person is another mode of infection.

Diagnosis and Treatment: Diagnosis is by physical examination and treatment uses drugs like nystatin, fluconazole, itraconazole, etc.

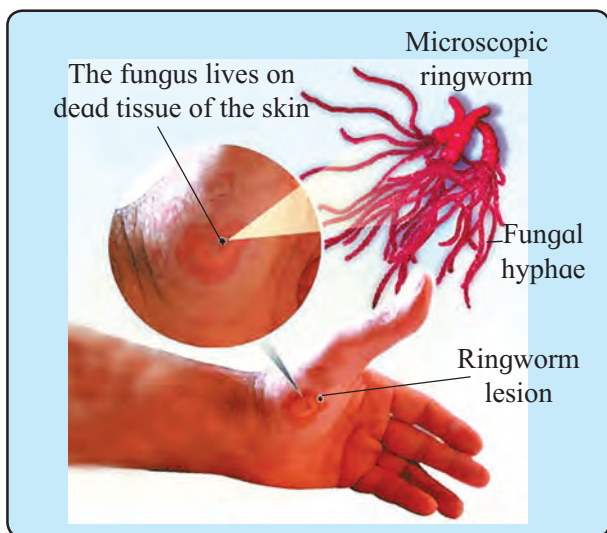


Fig. 10.11 : Ring Worm

Prevention and Control :

Avoid close contact, sharing of clothes, sport equipments. Washing clothes in hot water with fungicidal soap help to destroy fungal pathogen.

I. Dengue :

Dengue fever is a painful, debilitating vector borne disease, caused by any one of four closely related dengue viruses. Dengue virus is transmitted through the bite of female *Aedes* mosquito. The mosquito takes up the dengue virus when it sucks blood of a person suffering from dengue. It can not spread directly from one person to another person.



Can you tell?

1. Which are the diseases that can be avoided by eradication of mosquitoes in your area?
2. What are the various ways in which mosquitoes can be eradicated from any area?
3. What precautions will you take if you are travelling in an area which has lot of mosquitoes?

J. Cancer:

Abnormal, uncontrolled and purposeless division of cells may lead to the formation/development of mass of undifferentiated cells

i.e. tumor. When tumor is malignant, it is described as cancer and has ability to invade other tissues.

Cancer harms the body when cancerous cells divide uncontrollably to form new lumps or masses of tissue called **neoplasm** (except in the case of leukemia). Tumors can grow and interfere with the normal functioning of various organs. They also release secretions which alter body function(s).

It is one of the main killer diseases nowadays. Physicians and researchers who specialize in the study, diagnosis, treatment and prevention of cancer are called **oncologists**.

Tumors may develop anywhere in the body. However, all tumors are not cancerous. There are two types of tumors: **benign** or nonmalignant and cancerous or **malignant**.

1. Benign or Nonmalignant Tumor :

It grows slowly, may attain quite a large size, but it remains restricted to the site of its origin (localized) and does not spread to other part of the body. This does not necessarily mean that the benign tumors are not troublesome. Some benign tumors are harmful and fatal e.g. brain tumor (A brain tumor may cause death because the brain is squeezed against the hard skull). Moreover, the benign tumors may sometimes become malignant. e.g. Adenoma and Fibroid.

2. Malignant tumor or cancer :

The growth rate of this tumor is rapid and mortality rate is comparatively more than benign tumor. Rapid growth of tumors causes overcrowding and disruption of normal cells. The cancerous cells compete with the normal cells for nutrients and finally kill them.

These cells are spread from one organ to other via blood or lymph and form new tumors called secondary tumors. This migratory process is called **metastasis**.

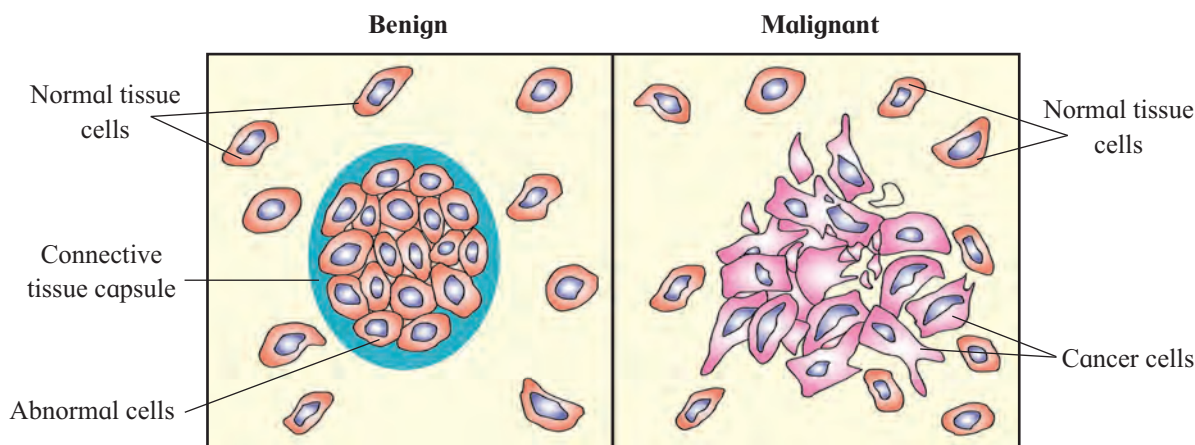


Fig. 10.12 : Benign and Malignant Tumor

a. Types of Cancer:

There are five main types of cancers according to the type of tissue affected. Cancers are named according to the tissue from which they arise.

- i. **Carcinoma:** Cancer that arise from epithelial tissue covering or lining the body organs is known as carcinoma. It include breast cancer, lung cancer, cancer of stomach, skin cancer. etc.
- ii. **Sarcoma:** Cancer that arises from connective tissue is called sarcoma. It include bone tumors (osteosarcoma), muscle tumors (myosarcoma), cancer of cartilage (chondrosarcoma) and cancer of adipose tissue (liposarcoma).
- iii. **Lymphoma:** Cancer that arises from lymphatic tissue, is called lymphoma. It occurs in the lymphatic nodes, spleen and tissues of immune system.
- iv. **Leukemia:** It is a type of blood cancer in which there is excessive formation of white blood cells (WBCs) or leucocytes in the bone marrow. People suffering from leukemia have very high leucocyte count. The blood contains millions of abnormal immature WBCs or leucocytes that are incapable of fighting infections. There are various types of leukemia such as monocytic leukemia, lymphoblastic leukemia etc.

- v. **Adenocarcinoma:** Adenocarcinoma cancer arises in thyroid, pituitary, adrenal and other glandular tissues.



Find out

Collect information about different types of lymphoma and hereditary cancer syndromes.

b. Causes of Cancer:

Although, it is still not very clear as to how the cancer is caused, several factors are now known to be cancer- causing i.e. **carcinogenic**. These factors are as follows.

- i. **Chemicals :** Several chemicals are known to induce cancer. These include nicotine, caffeine, products of combustion of coal and oil. Several polycyclic hydrocarbons, some sex hormone and steroids, if given or secreted in large amounts, may cause cancer. Breast cancer seems to have hormonal relationship. It is more commonly observed in women who avoid breast feeding.
- ii. **Radiation :** The x-rays, gamma-rays cosmic rays, ultra-violet rays etc. are carcinogenic. Incidence of skin cancer is higher in the people working in very sunny areas due to UV radiation in the sunlight.
- iii. **Viruses :** Viruses causing cancer have genes called viral oncogenes (v-onc genes).

These viruses are also called oncogenic viruses. e.g. EBV (Epstein-barr virus), HPV (Human papilloma virus) etc.

- iv. **Oncogenes** : Several genes called cellular oncogenes (c-onc genes) or proto-oncogenes have been identified in normal cells which when activated under certain condition could lead to oncogenic transformation of cells.
- v. **Addiction** : Different types addictions like smoking, chewing of tobacco lead to cancer of mouth, lips and lungs. Alcohol consumption may result in cancer of oesophagus, stomach, intestine and liver. Drugs also cause cancer e.g. Marijuana, anaerobic steroids etc.



Try This

1. Find the Oncocenters nearby your area.
2. Prepare a chart of types of cancer and their preventive measures.
3. Organize a street-play on awareness about cancer and present it in your area.

Table 10.13 : Carcinogens and Organ Affected

Carcinogen	Organ affected
1. Soot	Skin, lungs
2. Coal tar (3,4 benzopyrene)	Skin, lungs
3. Cigarette smoke (N-nitrosodimethlene)	Lungs
4. Cadmium oxide	Prostate gland
5. Aflatoxin (a metabolite of <i>Aspergillus flavus</i> , a mould)	Liver
6. 2-naphthylamine and 4-aminobiphenyl	Urinary bladder
7. Mustard gas	Lungs
8. Nickel and chromium compounds	Lungs
9. Asbestos	Lungs
10. Diethylstilbestrol (DES)	Vagina
11. Vinylchloride (VC)	Liver

c. Treatment of Cancer:

Cancer treatment consists of combination of a number of therapies which are follows:

- i. **Chemotherapy:** Chemotherapy comprises administration of certain anticancer drugs. The anticancer drugs check cell division by inhibiting DNA synthesis or are more toxic to cancerous cell than to normal cells. Chemotherapy may lead to hair loss or anaemia but both get corrected after the treatment is stopped.
- ii. **Radiotherapy:** It is used in addition to chemotherapy. In radiotherapy, the basic principle is to bombard the cancerous tissue or cells with the rays from radioactive materials. For treatment, the cancer tissue or cells are exposed to radiations from radioactive materials such as cobalt, iridium and iodine. The x-rays, gamma rays and charged particles are used to destroy the cancerous tissue or cells but cause minimum damage to the surrounding normal tissue or cells.
- iii. **Surgery:** In surgery, the entire cancerous tissue or cells are removed surgically. It has limited utility. In certain cases such as breast tumor or uterine tumor, the surgery is most effective, but other treatments are also given to kill any cancerous cell that may have been escaped in surgery.
- iv. **Immunotherapy:** Tumor cell have been shown to avoid detection and destruction by immune system. Therefore, the patients are given substances called biological response modifiers such as α -interferon which activates their immune system and helps in destroying the tumor.
- v. **Supportive therapy:** Supportive therapy is used to treat symptoms of cancer and side effects of cancer treatments. Objective of this therapy is to improve the quality of life of cancer patient. This therapy varies depending upon condition of individual patient.

Do you know ?

The Tata Memorial Hospital is situated at Kharghar (Mumbai), Maharashtra in India. It is a special cancer treatment and research centre, closely associated with the Advanced Centre for Treatment, Research and Education in Cancer. It is recognized as one of the leading cancer centres in South Asia. It is an autonomous body funded and controlled by Department of Atomic Energy, Government of India which also oversees the administration of Institute since 1962.



K. AIDS :

AIDS, the **acquired immuno deficiency syndrome**, is a usually fatal illness caused by a retrovirus (ss RNA) known as the **human immuno deficiency virus (HIV)** which weakens the body's immune system, leaving the victim vulnerable to life-threatening opportunistic infections, neurological disorders and unusual malignancies. AIDS can be called a modern pandemic (world wide), affecting both industrialized and developing countries.

AIDS was first noticed in USA in 1981. In India, first confirmed case of AIDS was in April 1986 from Tamil Nadu.

Structure of HIV:

HIV is 100 to 140 nm in diameter. It is spherical. Virus particle shows centrally located two ss RNA molecules along with reverse transcriptase enzymes. It is covered by two layers of proteins. The outer layer is of matrix protein (p17) while in inner layer is capsid protein (p24).

Over the matrix protein layers there is an additional layer of lipids. Impregnated with glycoprotein GP120 and GP 41 (Refer Fig. 10.14). The virus replicates in actively dividing T_4 lymphocytes and can remain in a latent state/stage in the lymphoid cells. The virus has unique ability to destroy human T_4 lymphocytes.

HIV is found in greatest concentration in blood, semen and cerebrospinal fluid (CSF) and to lesser extent in tears, milk, urine, saliva, cervical and vaginal secretions.

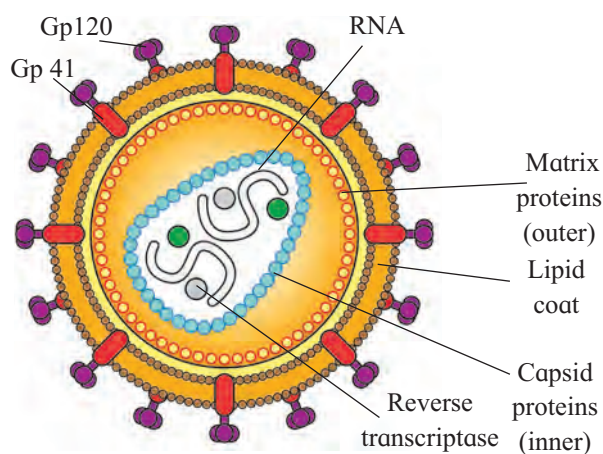


Fig. 10.14 : Structure of HIV

The virus can be transmitted through :

1. **Unsafe sexual contact:** Including oral, vaginal and anal sex.
2. **Blood :** Via blood transfusions or needle sharing.
3. **From mother to child (Transplacental):** A pregnant woman can transmit the virus to her foetus through their shared blood circulation, or a **nursing mother** can transmit it to her baby from her breast milk.

Other methods of spreading of the virus are rare and include accidental needle injury, artificial insemination with infected donated semen and transplantation with infected organs.

AIDS virus has been found in urine, tears, saliva, breast milk and vaginal secretions but it seems not to be transmitted by these fluids unless it gets into injuries.



Always Remember

HIV infection is not spread by:

Causal contact such as hugging, insect bite (mosquitoes), participation in sports, touching items previously touched by a person infected with the virus, hand shake, sharing clothes, swimming pools etc.

Clinical manifestations:

The clinical manifestations (symptoms) of AIDS have been classified into four broad categories.

- i. **Initial infection** with the virus and formation of antibodies, usually 2-8 weeks after initial infection.
- ii. **Asymptomatic carrier state** in which no signs of disease, are seen. Incubation period ranges for 6 months to 10 years.
- iii. **AIDS related complex (ARC)** with one or more of the following clinical signs: recurrent fever for longer than one month, fatigue, unexplained diarrhea, night sweats, shortness of breath, loss of more than 10 per cent body weight, etc.
- iv. **AIDS** is the end stage of HIV infection. It is characterised by life threatening opportunistic infections (like pneumonia, tuberculosis, kaposi sarcoma, etc.).

Preventive measures:

AIDS has no cure, hence prevention is the best choice. The following steps help in preventing this dreadful disease-

- i. People, particularly those in high –risk group, should be educated about HIV transmission.
- ii. Disposable needles and syringes should be used and disposed off properly and immediately.
- iii. Sexual habits should be changed immediately.
- iv. High-risk groups should refrain from donating blood.
- v. Tooth brushes, razors, other articles that can become contaminated with blood should not be shared.
- vi. Before receiving blood, ensure that it has been screened for not containing HIV infections.
- vii. Routine screening must be done for –
 - Blood donors.
 - Organ donors (kidney, liver, lung, cornea).
 - Donors of semen and growth hormone.
 - Patients undergoing hemodialysis and females in high risk group who are pregnant or contemplating pregnancy.



Do you know ?

- Every year, December 1st is observed as World AIDS Day.
- India started a National AIDS control Programme in 1987.
 - “A red ribbon” is worn on World AIDS Day as a universal symbol of awareness and support for people living with HIV.
- In India, 5 AIDS reference centers have been established-
 1. AIIMS – New Delhi.
 2. National Institute of Virology, Pune.
 3. National Aids Research Institute, Bhosari, Pune (NARI)
 4. Center of advanced research on Virology, Vellore.
 5. National institute of communicable diseases, New Delhi.
- National AIDS Control Organization (NACO) was set up in year 1992 by the Ministry of Health and Family Welfare. The aim of this organization is to prevent further transmission of HIV, to decrease morbidity and mortality associated with HIV infection and to minimize the socio-economic impact resulting from HIV infection.



Laboratory diagnosis:

At first a test is used to detect the HIV antibodies, while a second confirmatory test is used to weed out any false positive results. The first test is **ELISA** (Enzyme-Linked Immunosorbent Assay). The confirmatory test, usually a **Western Blot**, is a highly specific test. It is based on detecting specific antibody to viral core protein and envelope glycoprotein.

Treatment of AIDS :

Although AIDS has no cure, certain medicines called as Antiretroviral drugs can help in reducing the viral load and prolong the life of HIV patient. Examples of these drugs used in Antiretroviral therapy (ART) are TDF (tenofovir), EFV(Efavirenz), Lamivudine (3TC), etc.

The advancements made in biological sciences have helped us to deal effectively with many infectious diseases. The use of vaccines and immunization programmes have enabled us to eradicate completely the dreadful diseases like small pox. A large number of other infectious diseases like polio, diphtheria, pneumonia and tetanus have been controlled to a large extent by the use of vaccines. Biotechnology is on the verge of making available newer and safer vaccines. Discovery of antibiotics and various drugs has also enabled us to treat effectively infectious diseases like tuberculosis.

10.4 Adolescence :

It is the period of beginning with the appearance of secondary sexual characters and the termination with cessation of somatic i.e. body growth. It can also be regarded as a transitional stage of physical and mental development of child occurring between puberty and the legal adulthood between 10 to 19 years of age where individual is no longer a child but not yet an adult.

Adolescence in fact is a phase rather than fixed time period in the life. It is a phase of

development on many fronts like sexual and reproductive maturity, mental development, adult identity and transition from socio-economic and emotional dependence to relative independence.

Stages of Adolescence :

Adolescents are defined as individuals of 10 to 19 year age group. The government of India in its National youth policy defines adolescents as 13-19 years. Adolescence is divided in three stages (periods) viz. early stage, middle stage and late stage.

i. Early period (10 to 14 years) :

The changes include beginning of the appearance of secondary sexual characters ; growth reaches to its peak; rapid physical growth; concrete thinking; defining boundaries of dependence/independence; self exploration; developing body image; development of intense friendship; seeking to counter instability and evaluation.

ii. Middle period (15 to 17 years) :

It is characterized by almost complete development of secondary sexual characters; physical growth slows down, approximately 95 % of the adult stature is attained; thinking is more abstract; concrete thinking under stressful conditions; capability for long range thinking; reestablishing of body image; sense of leadership and all powerfulness; preoccupied with romantic fantasy and attraction towards opposite sex; establishing peer group that helps in defining behavioral code; etc.

iii. Late period (18 to 19 years) :

It is characterized by establishment of total physical maturity, established abstract thinking, intellectual and functional identity; peer group recedes in favor of individual relationship, stable relationship and change from childhood to adulthood relationship.

Physical changes of adolescence :

- **Growth spurt** occurs in both boys and girls. In boys muscles develop, skin become oily, broadening of shoulders, cracking of voice, development of underarm and chest hair, pubic hair, facial hair, enlargement of penis and testis. In girls development of breast, widening of hip, development of underarm and pubic hair, enlargement of uterus and ovaries.
- **Sexual development** : Sex organs mature and enlarge, sexual desire, erection of penis in boys, sperm production, ejaculation, ovulation, menstruation and initiation of sexual behaviour.

Emotional and social changes :

These include - establishing own identity, fantasy, day dreaming, attention seeking behavior, emotional instability, full of energy, sexual attraction, rapid mood changes, conflicts with family, behavioural code (influence of peer group), self exploration and evaluation, formation of new relationships, peer pressure, etc.



Always Remember

- Avoid looking at your parents as the enemy. Try to understand that your parents are human beings, with their own needs and feelings.
- Listen to your parents with an open mind, and try to see their point of view.
- Share your feelings with your parents so that they can understand you better.
- Live up to your responsibilities at home and in school so that your parents will be more inclined to grant you the kind of independence you want and need.
- Bolster your criticisms of family, school and government with suggestions for practical improvements.
- Be as courteous and considerate to your own parents as you would be to the parents of your friends.

Mental Health and Adolescence:

Many of the emotional and social changes have implication on the mental health. Most of the mental health issues that people confront as adults begin to appear in adolescence . The mental health implication includes confusion, irritation, moodiness, frustration, nausea, less concentration, hyperactivities, anger, effects on life styles like obesity, addictions, accidents, leading to ill health, etc. The mental illness in fact is in terms of different forms of depression like insomnia and loss of energy.

Thus, mental illness (disorders or unfavourable changes) is associated with psychological or behavioural manifestations. These are broadly classified as either psychoses or neuroses. Psychoses include delusions, hallucinations, disturbance in the thinking process, etc. The psychoses lead to Alzheimer's disease, schizophrenia, depressive psychosis, amnesia (loss of memory), bulimia (extreme over indulgence in food), anxiety (fear or apprehension), anorexia nervosa (emotional aversion to food) and depression (sadness, inactivity, reduced interest to enjoy life), etc. Neuroses include schizophrenia, illusions, hallucination, etc.

Adolescence is thus characterized by number of cognitive, emotional, behavioural, physical and attitudinal changes. These may lead to positive personality development which depend upon the relation with parents or to other conditions, or to conflicts with others. Infact, adolescents view their friends, peer group more important and influential than their parents. This may lead to various kinds of addictions like smoking, taking drugs, etc.

Treatment :

Treatment of such disorders should be preferentially with non-pharmacological approach with due respect to the rights of children. WHO has recommended evidence based guidelines under mental health Gap Action Programme (mhGAP).

10.5 Addiction :

This is a complex condition characterised by compulsive use of substance despite of its harmful consequences. Addiction may involve the use of substances (drugs) such as alcohol, opioids, cocaine, nicotine and others or behaviours such as gambling.

There is scientific evidence that the addictive behaviours share key neurobiological features. They intensely involve brain pathways of reward and reinforcement, affecting motivation, which involve the neurotransmitter dopamine. It is important to know that the neurological changes are reversible after the substance-use or behaviour is discontinued.

Addiction overall result in the impairment of physical, physiological and psychological functions of the body.

Causes of substances abuse during Adolescence

- Insufficient parental supervision and monitoring.
- Lack of communication between child and parents.
- Poorly defined rules.
- Family conflicts.
- Favorable parental attitudes towards alcohol and drug uses.
- Expectations from drugs use.
- Risk taking behavior.

Methods /measures to control drug abuse :-

1. Always remember 'Prevention is better than cure'.
2. Avoid undue pressure – A child should not be forced to perform beyond his /her capacities in studies, sports and other activities.
3. Education and Counselling of child to face problems and stress, to accept

disappointments and failures as a part of life. Channelize the energy of child in sports, studies and other constructive activities.

10.6 Drugs Abuse :

Surveys and statistics show that use of drugs and alcohol has been on the rise especially among the youth. This is really a cause of concern as it could result in many harmful effects. Proper education and guidance would enable youth to safeguard themselves against these dangerous behaviour pattern and follow healthy lifestyle. **The drugs, which are commonly abused, are opioids, cannabinoids and alkaloids of coca.**

a. Opioids :

These drugs binds to opioid receptors present in central nervous system and gastrointestinal tract. **Heroin**, otherwise called **smack** is chemically di-acetyl morphine. It is extracted from latex of poppy plant *Papaver somniferum*. Heroin is depressent and slows down the activity of body.

b. Cannabinoids :

Interact with receptors present in brain. Inhalation and ingestion of these substances affect the cardiovascular system. These are obtained from inflorescences and the parts of *Cannabis sativa*. Marijuana, hashish, charas and ganja are other different forms of drugs obtained from this plant.

c. Cocain :

This is an alkaloid obtained from coca plant- *Erythroxylum coca*. It increases level of neurotransmitter-dopamine. Its excessive dosage causes extreme happiness, irritability, paranoia.

Hallucinogens (mind expanding drugs) :

These are alkaloids causing day-dreaming. Lysergic acid and cannabis are hallucinogenic substances. *Atropa bellodona* and *Datura*

spp. also have hallucinogenic properties. Hallucinations are unreal perceptions of unreal objects due to the disorder of the nervous system.



Do you know ?

Drugs like barbiturates, amphetamine, benzo-diazepines, lysergic acid diethylamide (LSD) are derived from the plant *Cannabis sativa* and are used as medicine. They help the patient in contracting insomnia and depression.

Addiction and Dependence :

Because of the perceived benefits, drugs are used repeatedly. The most important thing, which one fails to realize, is the inherent addictive nature of alcohol and drugs. Addiction is a psychological attachment to certain effects—such as euphoria and a temporary feeling of well-being—associated with drugs and alcohol. These drive people to take them even when these are not needed, or even when their use becomes self-destructive. With repeated use of drugs, the tolerance level of the receptors present in our body increases.

Consequently the receptors respond only to higher doses of drugs or alcohol leading to greater intake and addiction. However, it should be clearly borne in mind that use of these drugs even once, can lead to addiction. In the absence of any guidance or counselling, the person gets addicted and becomes dependent on their use. Dependence is the tendency of the body to manifest a characteristic and unpleasant **withdrawal syndrome** if regular dose of drugs/ alcohol is abruptly discontinued. This is characterized by anxiety, trembling, nausea and sweating, which may be relieved when use is resumed.

Effects of Drug/ Alcohol Abuse :

The immediate adverse effects of drugs and alcohol abuse are manifested in the form of reckless behaviour, vandalism and violence.

Excessive doses of drugs may lead to coma and death due to respiratory failure, heart failure or cerebral hemorrhage. A combination of drugs or their intake along with alcohol generally results in overdose and even deaths.

The most common **warning signs** of drug and alcohol abuse among youth include drop in academic performance, unexplained absence from school/college, lack of interest in personal hygiene, withdrawal, isolation, depression, fatigue, aggressive and rebellious behaviour, deteriorating relationships with family and friends, loss of interest in hobbies, change in sleeping and eating habits, fluctuations in weight, appetite, etc.

There may even be some far-reaching implications of drug/alcohol abuse. If an abuser is unable to get money to buy drugs/ alcohol he/she may turn to crime. At times, a drug/alcohol addict becomes the cause of mental and financial distress to his/her entire family and friends.

Those who take drugs intravenously (direct injection into the vein using a needle and syringe) are likely to acquire serious infections like HIV and hepatitis B. Use of alcohol during adolescence may also have long-term effects like loss of balance, liver cirrhosis, pancreatitis. It could lead to heavy drinking in adulthood.

Chronic use of drugs and alcohol damages the nervous system and liver (**cirrhosis**). Use of drugs and alcohol during pregnancy adversely affects the foetus.

Another misuse of drugs is that certain sports persons use drugs to **enhance performance**. They (mis)use narcotic analgesics, anabolic steroids, diuretics and certain hormones to increase muscle strength and bulk and to promote aggressiveness and overall improvement in their performance. Side-effects of the use of **anabolic steroids** in females include masculinization (features like males), increased aggressiveness, mood

swings, depression, abnormal menstrual cycles, excessive hair growth on the face and body, enlargement of clitoris, deepening of voice.

In males it includes acne, increased aggressiveness, mood swings, depression, and reduction of size of the testicles, decreased sperm production, kidney and liver dysfunction, breast enlargement, premature baldness, enlargement of the prostate gland. These effects may be permanent with prolonged use.

Prevention and Control :

The age-old adage (i.e. proverb) is ‘prevention is better than cure’ holds true for all addictions. It is also true that habits such as smoking, taking drug or alcohol are more likely to be taken up at a young age, more during adolescence. It is best to identify the situations that push an adolescent towards use of drugs or alcohol, and to take remedial measures well in time. **In this regard, the parents and the teachers have a special responsibility.**



Use your brain power

Deaddiction may be difficult but not impossible. Collect information about NGOs, working in the field of deaddiction.



Activity :

1. Identify the name of the plant. Enlist different types of drugs derived/obtained from the same.



2. Prepare chart of different stages of adolescence mentioning the changes in growth, sexual development and emotional and social changes.

Changes	Stages	I		II		III	
		Boy	Girl	Boy	Girl	Boy	Girl
Growth	1.						
	2.						
	3.						
Sexual development	1.						
	2.						
	3.						
Emotional and social	1.						
	2.						
	3.						

Exercise

Q. 1 Multiple Choice Questions

- Which of the following is NOT caused by unsterilized needles?
 - Elephantiasis
 - AIDS
 - Malaria
 - Hepatitis B
- Opium derivative is
 - Codeine
 - Caffeine
 - Heroin
 - Psilocybin
- The stimulant present in tea is
 - tannin
 - cocaine
 - caffeine
 - crack
- Which of the following is caused by smoking?
 - Liver cirrhosis
 - Pulmonary tuberculosis
 - Emphysema
 - Malaria
- An antibody is
 - molecule that binds specifically to an antigen
 - WBC which invades bacteria
 - secretion of mammalian RBC
 - cellular component of blood
- The antiviral proteins released by a virus-infected cell are called
 - histamines
 - interferons
 - pyrogens
 - allergens
- Both B-cells and T-cells are derived from
 - lymph nodes
 - thymus glands
 - liver
 - stem cells in bone marrow
- Which of the following diseases can be contracted by droplet infection?
 - Malaria
 - Chicken pox
 - Pneumonia
 - Rabies

- Confirmatory test used for detecting HIV infection is
 - ELISA
 - Western blot
 - Widal test
 - Eastern blot
- Elephantiasis is caused by
 - W. bancrofti*
 - P. vivax*
 - Bedbug
 - Elephant
- Innate immunity is provided by
 - phagocytes
 - antibody
 - T- Lymphocytes
 - B- Lymphocytes

Q. 2 Very Short Answer Questions

- What is the source of cocaine?
- Name one disease caused by smoking?
- Which cells stimulate B-cells to form antibodies?
- What does the abbreviation AIDS stand for?
- Name the causative agent of typhoid fever?
- What is Rh factor?
- What is schizont?
- Name the addicting component found in tobacco.
- Name the pathogen causing Malaria.
- Name the vector of Filariasis.
- Give the name of the causative agent of ringworm.
- Define health.

Q. 3 Short Answer Questions:

- What are acquired diseases?
- Differentiate between antigen and antibody.
- Name the infective stage of *Plasmodium*. Give any two symptoms of malaria.
- Explain the mode of infection and cause of elephantiasis.
- Why is smoking a bad habit?

6. What do the abbreviations AIIMS and CMIS denote?
7. What is a carcinogen? Name one chemical carcinogen with its target tissue.
8. Distinguish between active immunity and passive immunity.

Q. 4 Short Answer Questions

1. Differentiate between B-cells and T-cells.
2. What are the symptoms of malaria? How does malaria spread?
3. Write a short note on AIDS.
4. Give the symptoms of cancer.
5. Write a note on antigens on blood cells.
6. Write a note on antigens-antibody complex.
7. What are the various public health measures, which you would suggest as safeguard against infectious diseases?
8. How does the transmission of each of the following diseases take place?
 - a. Amoebiasis
 - b. Malaria
 - c. Ascariasis
 - d. Pneumonia
9. What measure would you take to prevent water-borne diseases?
10. Write a short note on typhoid.

Q. 5 Match the following.

Column I	Column II
a. AIDS	i. Antibody production
b. Lysozyme	ii. Activation of B-cells
c. B-cells	iii. Immunoglobulin
d. T-helper cells	iv. Tears
e. Antibody	v. Immuno deficiency

Q. 6 Long Answer Questions

1. Describe the structure of antibody.
2. Write a note on Vaccination.
3. What is cancer? Differentiate between benign tumor and malignant tumor. Name the main five types of cancer.
4. Describe the different type of immunity.
5. Describe the ill-effects of alcoholism on health.
6. In your view, what motivates the youngsters to take to alcohol or drugs and how can this be avoided?
7. Do you think that friends can influence one to take alcohol/drugs? If yes, how may one protect himself/herself from such an influence?

Project :

1. Collect information about the, symptoms and preventive measures treatments for Dengue, Swine flu and Tuberculosis (TB).
2. Browse the information about **COVID - 19** with respect to
 - a. Structure
 - b. Modes of infection
 - c. Preventive measures
 - d. Treatment





Can you recall?

1. What are hybrid plants?
2. Different examples of microbes used in household production.
3. The types or groups of micro-organisms that produce antibiotics.
4. What are biofertilizers?
5. What is the source of silk thread?

11.1 Improvement In Food Production :

Food is one of the basic needs as it gives us energy for everything. It keeps us alive, strong and healthy. It can be defined as any thing solid or liquid, which when swallowed, digested and assimilated in the body, keeping us well. It is organic, energy rich, non-poisonous, edible and nourishing substance.

Green plants synthesize their own food through photosynthesis. But animals including humans can not synthesize their food on their own, hence are dependent on plants directly.

The famine is responsible for dearth of food, besides the rapid and dramatic increase in world population over the time. To meet the increasing demand for food, there is need for improvement of food production, both quantity wise and quality wise (nutritive). **Plant breeding** and **animal breeding** mainly help us to increase the food production.

11.2 Plant breeding :

The improvement or purposeful manipulation in the heredity of crops and the production of new superior varieties of existing crop plants, constitute what is called **plant breeding**. It is, therefore, an applied branch of botany. It is infact an art and the science of changing and improving the heredity of plants. Plant breeding can be carried out by using the applications of principles of genetics, taxonomy, physiology, pathology, agriculture, rDNA technology, etc.



Do you know ?

Primary aim of plant breeding is to obtain a new crop variety superior to the existing type, in all characters. This purposeful manipulation incorporates various objectives which differ from plant to plant depending upon its type and the use. Some objectives are common- like yield, quality, resistance, life cycle span, etc., while some objectives will differ on the basis of the type and use of plant.

Plant breeding is a method of altering the genetic pattern of plants to increase their value and utility for human welfare. The plant breeding is done to increase crop yield, improve quality, increase tolerance to environmental stresses, make the plants resistant to pathogens and increase tolerance to insect pest. Green Revolution was the result of a sequence of scientific breakthroughs and developmental activities that successfully fought hunger by increasing food production. Seeds with superior quality, use of chemicals - pesticides and fertilizers, and multiple cropping system supported by the use of modern farm machinery and proper irrigation system, helped for the development of high-yielding and disease resistant varieties in wheat, rice, maize, etc.

In fact, plant breeding dates back to about 10,000 years ago. The present day crops are the result of domestication and acclimatization.



Do you know ?

Different methods of plant breeding include Introduction, Selection, Hybridization, Mutation breeding, Polyploidy breeding, Molecular plant breeding, Tissue culture, rDNA technology, SCP, etc.



Know the Scientist :

Dr. Norman E. Borlaug:

An American biologist, who has been called “Father of the Green Revolution”, “Agriculture’s greatest spokesperson” and “The Man Who Saved a Billion Lives”, Dr. Borlaug, a 1970 Nobel Laureate, was honoured for his work in the ‘Green Revolution,’ saving millions of lives from famine in India, Mexico, and the Middle East.

Dr. M. S. Swaminathan:

He has been called the “Father of Green Revolution in India” for his role in introducing and further developing high-yielding varieties of wheat in India. He advocated moving India to sustainable development, especially using environmentally sustainable agriculture, sustainable food security and the preservation of biodiversity. He is pioneer in mutation breeding in India. He developed new varieties of wheat like Sonora, NP 165 and Sharbati.

A. Hybridization and its technique :

It is the chief method that offers greater possibilities in the crop improvement than other methods. The use of this method is the only effective means of combining together the desirable characters of two or more varieties. By this method, one can create new genetic combinations of already existing characters and new genetic variations. It also exploits and utilizes hybrid-vigour.



Internet my friend

Try to know more about the hybrid-vigour.

Hybridization can be **intravarietal**, **intervarietal** (between two varieties of the same species), **interspecific** (between two species of the same genus) and **intergeneric** (between two genera of the same family). As parental plants are distantly related, such crosses are also

called **wide/ distant crosses**. Interspecific and intergeneric hybrids are seldom to occur in the nature.

The main steps of the plant breeding program (Hybridization) are as follows :

1. Collection of Variability :

Wild species and relatives of the cultivated species having desired traits, should be collected and preserved. The entire collection having all the diverse alleles (i.e. variations) for all genes in a given crop, is called *germplasm collection*. Variations are useful in the selection. Germplasm conservation can be done in following ways-

- *In situ* conservation : It can be done with the help of forests and Natural Reserves.
- *Ex situ* conservation : It is done through botanical gardens, seed banks, etc.

2. Evaluation and Selection of Parents :

It is an important and essential step. The collected germplasm is evaluated (screened) to identify plants with desirable characters. The two plants selected as parents must be healthy, vigorous and should show desirable but complementary features. The selected parents are selfed for three to four generations to make them pure or homozygous. It is made sure that only pure lines are selected, multiplied and used in the hybridization.

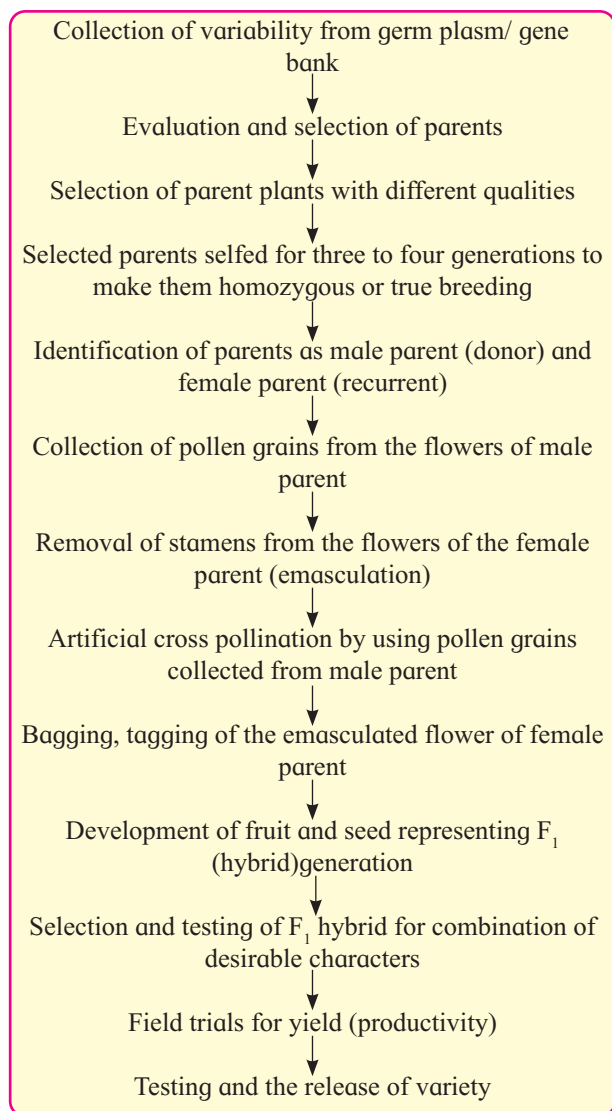
3. Hybridization :

The variety showing maximum desirable features is selected as female (recurrent) parent and the other one as male parent (donor) which lacks good characters found in recurrent parent.

The pollen grains from anthers of male parent are collected and then artificially dusted over stigmas of emasculated flowers of female parent. Pollination is followed by seed and fruit formation in due course. The seed, thus obtained represents the hybrid generation.

The hybrid F_1 progeny is selected and evaluated for the desired combinations of characters.

Chart 11.1 : Steps of Hybridization technique



4. Selection and Testing of Superior Recombinants :

The F_1 hybrid plants showing superiority over both the parents and having high hybrid-vigour, are selected. Such hybrids are then selfed for few generations to make them homozygous for the said desirable characters till there is a state of uniformity, so that the characters will not segregate further.

5. Testing, Release and Commercialization of New Cultivars :

The newly selected lines are evaluated for the productivity and other features like disease resistance, pest resistance, quality, etc. Initially, these plants are grown under controlled conditions of water, fertilizers, etc. and their

performance is recorded. The selected lines are then grown for three generations at least in natural field, in different agroclimatic zones. Finally variety is released as new variety for use by the farmers.

The cultivation of many high yielding, hybrid varieties of rice, wheat, sugarcane, millets, developed through hybridization, fertilizers pesticides and proper irrigation have helped farmer community to attain record agricultural production in India since 1961. This is called **green revolution**.

Indian Hybrid Crops :

1. Wheat and Rice :

In 1960s, wheat and rice production increased tremendously. Norman E. Borlaug developed semi-dwarf varieties of wheat. *Sonalika* and *Kalyan Sona* are two of the hybrid wheat varieties, grown in India. Semi-dwarf rice varieties were taken from IR-8 (International Rice Research Institute) and Taichung native-I (from Taiwan) and introduced in India. *Jaya*, *Padma* and *Ratna* are the better-yielding, semi-dwarf rice varieties that were developed later.

2. Sugarcane :

- *Saccharum barberi* is a native of North India and *S. officinarum* belongs to South India.
- *S. officinarum* has thicker stem and high sugar contents, but it does not grow well in North India.
- These two varieties were crossed to get the desirable qualities of both (high sugar content, thicker stem and the ability to grow in North India). CO -419, 421, 453 are high yielding and having high sugar contents are developed in India at Coimbatore (Tamilnadu).

3. Millets :

- Hybrid maize (Ganga-3), *Jowar* (CO-12), and *Bajra* (Niphad) have been successfully developed in India.
- These varieties are high yielding and resistant to water stress.

Plant Breeding for Disease Resistance :

Some of the diseases caused in plants are-

Pathogen	Plant disease
Fungi	Brown rust of wheat Red rot of sugarcane Late blight of potato
Bacteria	Black rot of crucifers
Viruses	Tobacco mosaic virus

The basic objective of breeding for disease resistance is to develop inherent quality in the plant to prevent the pathogen from causing the disease. Such varieties of plants are called disease resistant plants. The basic technique used is the same as for normal hybridization process.

Some disease resistant plants developed are :

Crop	Variety	Resistant to Disease
Wheat	Himgiri	Leaf and stripe rust, hill bunt
<i>Brassica</i>	Pusa Swarnim	White rust
Cauliflower	Pusa Shubra	Black rot and Curl blight black rot
Chilli	Pusa Sadabahar	Chilli mosaic virus, Tobacco mosaic virus and leaf curl

B. Mutation Breeding :

Mutation is sudden heritable change in the genotype, caused naturally. It can also be induced by application of chemicals.

Natural (physical) mutagens are : High temperature, high concentration of CO₂, X rays, UV rays.

Chemical mutagens are : Nitrous acid, EMS (Ethyl Methyl Sulphonate), Mustard gas, Colchicine, etc.

Seedlings or seeds are irradiated by CO-60, exposed to UV bulbs, X ray machines, etc.

Mutagens cause gene mutations and chromosomal aberrations. The treated seedlings are then screened for resistance to diseases/ pests, high yield, etc. e.g. Jagannath

variety of rice, NP 836 variety of wheat (rust resistant), Indore-2 variety of cotton (resistant to bollworm), Regina-II variety of cabbage (resistant to bacterial rot), etc.

Plant Breeding for Developing a Resistance to Insect Pest:

Insects being herbivores, incur heavy loss in the quantity and quality of crops. Resistance in crops can be developed by following ways :

- Development of morphological characters like hairy leaves in cotton and wheat develop vector resistance from jassids and cereal leaf beetle, respectively.
- Solid stem in wheat leads to resistance to stem borers.
- Biochemical characters provide resistance to insects and pests. For example, the high aspartic acid, and low nitrogen and sugar content in maize, lead to resistance against stem borers.
- The nectar-less cotton having smooth leaves develop resistance against bollworms.

Some pest-resistant varieties are-

Crop	Variety	Insect pest
<i>Brassica</i>	Pusa Gaurav	Aphids
Flat bean	Pusa sem 2 Pusa sem 3	Jassids, aphids and fruit borer
Okra	Pusa Sawani, Pusa A-4	Shoot and fruit borer

11.3 Tissue culture :

It is actually a collection of different techniques. It is infact, emerged as a technique of plant biotechnology. Here, isolated cells, tissues, organs are grown '*in vitro*' on a solid/ liquid nutrient medium, under aseptic and controlled conditions of light, humidity and temperature, for achieving different objectives. The part of plant used in tissue culture is called **explant**.

Plant tissue culture is based on principle of **Totipotency** which is an inherent ability of living plant cell to grow, divide, redivide and give rise to a whole plant. Haberlandt (1902)

for the first time conceived this idea and developed the concept of *in vitro* cell culture (plant morphogenesis).

The plant tissue culture medium contains all essential minerals, sources for carbohydrates, proteins and fats, water, growth hormones, vitamins and agar (for callus culture). The most commonly preferred medium for tissue culture is MS (Murashige and Skoog) medium.

Based on the nature of explant : There are three types viz, cell culture, organ culture and embryo culture.

Based on the type of *in vitro* growth : There are two types viz. Callus culture (solid medium) and Suspension culture (liquid medium).

Maintenance of aseptic conditions : Aseptic condition is essential so as to avoid contamination by other harmful micro-organisms. It is accomplished by **sterilization** of : **glass ware** (use of detergents, hot air oven), **nutrient medium** (by autoclave under constant pressure of 15 lb/sq inch for continuous 20 minutes), **Explant** (by treatment of 20% ethyl alcohol and 0.1% HgCl₂), **Inoculation chamber** (Laminar air flow) - by using UV ray tube for 1 hour before performing actual inoculation of explant on the sterilized nutrient medium.

Other conditions maintained are : Temperature - 18°C to 20°C, pH of nutrient medium 5 to 5.8 and aeration particularly for suspension culture.

In **callus culture**, the solid medium is used. The development and organisation of tissue is lost. Hence, the cells of explant, divide and redivide to form a mass of undifferentiated cells, called **callus**. It is maintained on solid medium. Callus can be induced to form organs like root (rhizogenesis) and shoot (caulogenesis) and thus the plantlet. No shaker (agitator) is needed.

In **suspension culture**, small groups of cells or a single cell are used as explant in the liquid medium. The liquid medium is constantly agitated by using shakers (agitators) so that there is constant mixing of medium and the explant.

Both the callus and suspension cultures die in due course of time. Therefore, subculturing is necessary for continuation of the technique.

Micropropagation (Clonal Propagation) :

Organogenesis via shoots is considered as one of the most widely used commercial method of regeneration of plant.

Micropropagation is also known as **clonal propagation**. It is the only process adopted by Indian plant biotechnologists in

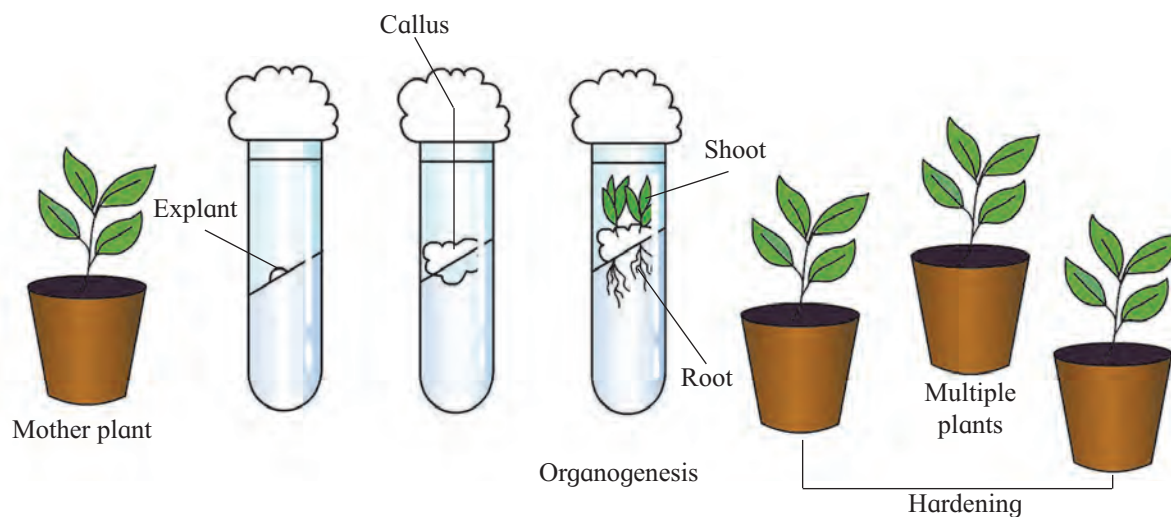


Fig. 11.2 : Steps in plant tissue culture (callus culture)



Do you know ?

Applications of tissue culture :

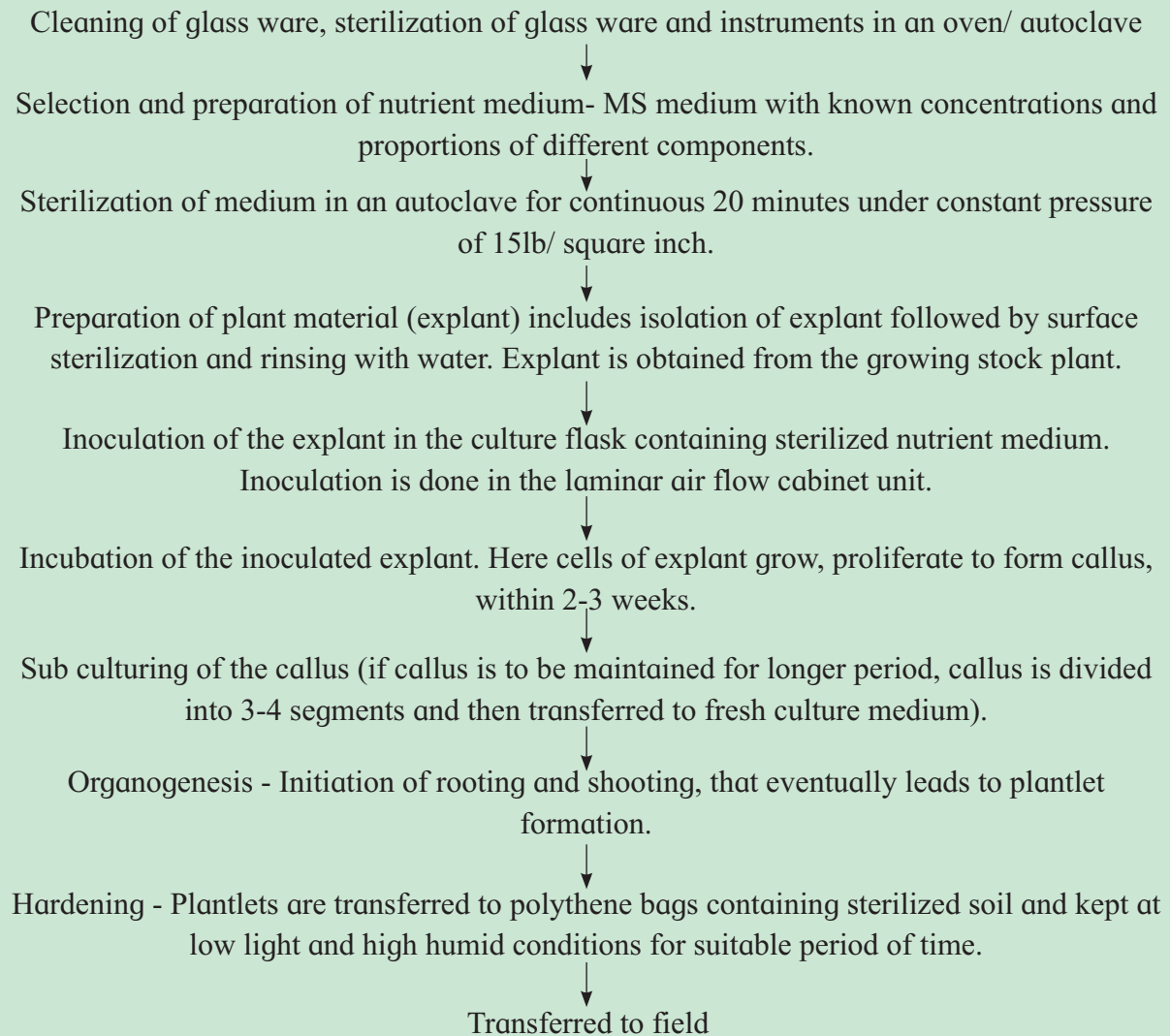
There are various application of tissue culture in forestry, agriculture, horticulture, genetic engineering, physiology, etc. The different applications of tissue culture include - Production of disease free plants and haploid plantlets, micropropagation, production of secondary metabolites, protoplast culture, culture of rare plants, somaclonal variations, production of stress resistant plants, etc.

different industries, mainly for the commercial production of ornamental plants like orchids, *Chrysanthemum*, *Eucalyptus*, etc. and fruit plants like banana, grapes, *Citrus*, etc.

Advantages of micropropagation:

1. It helps in rapid multiplication of plants.
2. A large number of plantlets are obtained within a short period and in a small space.
3. Plants are obtained throughout the year under controlled conditions, independent of seasons.
4. Genetically identical plants (clones) are produced (formed) by this method. Therefore desirable characters (genotype) and desired sex of superior variety are kept constant for many generations.
5. The rare plant and endangered species are multiplied by this method and such plants are saved.

Chart 11.3 : Flow chart for tissue culture technique.



6. With the help of somatic hybrids (cybrids), we are able to obtain new variety in short time span.



Do you know ?

High yielding varieties of Banana viz. 'Shrimati', 'Basarai' and G- 9 are mostly used in Maharashtra.

11.4 Single cell protein (SCP) :

By 2050, the world would need to produce 1,250 million tonnes of meat and dairy products per year, to meet global demand for animal-derived protein at current consumption levels. However, growing demand for protein will not be met sustainably by increasing meat and dairy production because of the low efficiency of converting feed to meat and dairy products.

More over, human population in underdeveloped and even in the developing countries is suffering from protein malnutrition, resulting into variety of nutritional diseases. To fight with this, efforts are undertaken by conventional methods to increase the food yield by different methods of crop improvement, use of biofertilizers, biopesticides, chemical fertilizers and high yielding varieties (green revolution). The efforts in other direction are also undertaken in non-conventional way. One such way is production of SCP- single cell proteins. Importance of SCP was realised during World War I.

Single-cell protein refers to the crude, or a refined edible protein, extracted from pure microbial cultures or from dead or dried cell biomass.

Microorganisms like algae, fungi, yeast, and bacteria have very high protein content in their biomass. These microbes can be grown using inexpensive substrates like agricultural waste viz. wood shavings, sawdust, corn cobs, paraffin, N-alkanes, sugarcane molasses, even human and animal wastes.

The microorganisms utilize the carbon and nitrogen present in these materials and convert them into high-quality proteins that can be used as a supplement, in both human and animal feed. Besides proteins, SCP is also rich in vitamins, vitamin B complex, minerals and fats. The single-cell proteins can be readily used as fodder for achieving fattening of calves, pigs, in breeding fish and even in poultry and cattle farming. The microorganisms used for the production of SCP are as follows :

Fungi : *Aspergillus niger*, *Trichoderma viride*

Yeast : *Saccharomyces cerevisiae*, *Candida utilis*

Algae : *Spirulina spp*, *Chlorella pyrenoidosa*

Bacteria : *Methylophilus methylotrophus*,
Bacillus megasterium.

Advantages of Single-Cell Protein :

- Microorganisms have a high rate of multiplication that means a large quantity of biomass can be produced in a comparatively short duration.
- The microbes can be easily genetically modified to vary the amino acid composition. They have high protein contents- 43% to 85% (W/W basis).
- A broad variety of raw materials, including waste materials, can be used as a substrate for SCP. This also helps in decreasing the number of pollutants.
- SCP serves as a good source of vitamins, amino acids, minerals, crude fibres, etc.

11.5 Biofortification

It is a method in which crops are bred (produced) for having higher levels of vitamins, minerals and fats (i.e. better nutritive value). It can also be achieved by supplementing nutrients from outside, besides breeding. Due to this, problem of malnutrition can be overcome. Following objectives were considered for the breeding program :

- Protein content and quality
- Oil content and quality
- Vitamin content
- Micronutrient content and quality

Biofortification can be achieved through conventional selective- breeding practices and also through r-DNA technology. It focusses on making plant food more nutritive as plants grow or develop.

Some examples of biofortification :

- Fortified Maize having twice the amount of amino acids- lysine and tryptophan.
- Wheat -Atlas 66 has a high protein content and Iron-fortified rice has 5 times more iron than normal.
- Vegetable crops like carrot and spinach have more vitamin A and minerals.
- Vitamin C enriched bitter gourd and tomato have been developed by IARI.

11.6 Animal husbandry :

Animal husbandry is an agricultural practice of breeding and raising livestock. It is not only a skill of farmers but also is as much a science, as it is an art.

Animal husbandry deals with care and breeding of livestock like buffaloes, cows, pigs, horses, cattles, sheeps, camels, goats, etc. which are useful to humans. It also includes poultry farming, fish farming, bee keeping, sericulture, lac culture, etc. Animals like honey bees, silk worms, prawns, crabs, birds, fish, pigs, cattle, sheep and camels have been used by humans for the products like milk, eggs, meat, wool, honey, silk, etc.

During the conventional practices of animal breeding, just taking care is not enough to give maximum yield. Although India and China have 70% of the world's livestock population, surprisingly the productivity serves only 25% of the world farm-produce.

So professional approach is needed to boost the production. It requires management procedures, new technologies to be employed in various farm system to achieve improvement in quality and productivity. Industrial principles of production, processing and marketing are to be employed.

Management of farms and farm animals :

Farm management starts from selection of high yielding breeds, their food requirements, supply of adequate nutritional sources, cleanliness of the environment and maintenance of health. Management of farm animals includes veterinary supervision, vaccination, high yielding cross breed development, production and preservation of products, distribution and marketing.

A. Animal breeding :

Breeding of animals is an important aspect of animal husbandry. Animal breeding aims at increasing the yield of animals and improving the desirable qualities of the products.

Breed:

A group of animals related by descent and similar in most characters like general appearance, features, size, configuration, etc., are said to belong to a breed.

Animal breeding is done for getting improved breeds with desirable qualities of product and also to increase yield of animals, Desirable characters such as increased production of milk, quality of product, quality of meat or maximum yield of eggs per year etc., are necessarily achieved through animal breeding.



Can you tell?

Find out the name of common breeds of cattle and poultry in the farms, found in your area.

Breeding can be of two main types - inbreeding and outbreeding :

a. Inbreeding : It involves breeding of closely related individuals for 4 to 6 generations. Inbreeding increases homozygosity. By inbreeding, pure lines of animals can be obtained. Inbreeding is helpful in the elimination of harmful recessive genes and for the accumulation of superior genes. Inbreeding has the demerit that it usually reduces the fertility and productivity.

b. Outbreeding: It involves breeding of unrelated animals. The animals may be of the same breed but having no common ancestors for 4 to 6 generations.

Outcrossing involves breeding between the animals of different species. It is also known as interspecific hybridization. Outcrossing helps to remove the inbreeding depression.

Crossbreeding involves the breeding of superior male of one breed with superior female of another breed. By cross-breeding, new animal breeds of desirable characters are developed. e.g. Hisardale is a new breed of sheep developed from crossing of Bikaneri ewe and Marino rams in Punjab.

Interspecific hybridization involves breeding of animals of two different but related species. It result in the formation (production) of animals with desirable characters from both the parents. But such breeding is not always successful e.g. Mule is a breed obtained from horse and donkey.

Artificial insemination technique involves controlled breeding experiments. Semen from selected superior males is collected and preserved in frozen state or injected into the genital tract immediately. It is useful to overcome the problem of normal mating and inconvenience of transportation.

Multiple Ovulation Embryo Transfer (MOET) involves the technology which provides the chances of successful production of hybrids. In this method, cow is administered with FSH like hormone, to induce follicular maturation and then the super ovulation is brought about. In each cycle, 6 to 8 eggs mature simultaneously. The cow is either mated with an elite bull or artificially inseminated. The blastocysts at 8 to 32 cell stage are recovered non-surgically and transferred to surrogate mothers. This technology is successfully used in cattles, sheeps, rabbits, buffaloes, etc. High milk yielding breeds of female and high

quality meat yielding bulls have been found to be successful, to increase herd size in a short period.

B. Dairy farm management :

Dairy industry involves production, processing and distribution of milk and milk products. Milk is a valuable food stuff universally consumed by human beings. Milk yield mainly depends on the quality of breeds in the farm. Selection of good breeds having high yielding potential under the climatic conditions of inhabiting area, and disease resistance is the basic requirement. In India, cows and buffaloes are mainly used for dairy farms. Sahiwal, Sindhi, Gir are Indian breeds and Jersey, Brown Swiss, Holstein are exotic breeds, which are used in dairy farming. Buffaloes are restricted to some part of Asia only.

In India, six breeds occur viz, Jaffarabadi, Mehsana, Murrah, Nagpuri, Nili, Surati, which are all good milk producers. Cattles have to be well looked after. Quality and quantity of fodder in proper ratio, should be given. Silage made from legumes and grasses, maize and jowar, makes good feed. Silage is supplemented with oil cakes, minerals, vitamins and salts. Cleanliness and hygiene of the cattles and handlers is of more importance while milking, storage and transport of milk and milk products. In recent years, much of these processes are mechanised, which reduce the chance of direct contact with the product. The shed must be cleaned daily. It should be spacious with adequate facilities for feeding, watering and lighting.



Do you know ?

Regular visit of veterinary doctor to dairy farm is mandatory, why ?

Milk processing, marketing and distribution, play an important role in dairy industry. Variety of milk product like curd, cream, butter, ghee, condensed milk, khoa,

cheese must be prepared from extra milk and on demand. An additional income can be obtained in the cattle farms from cow dung, manure, fuel cakes and gobar gas for cooking and lighting.

C. Poultry farm management :

Poultry includes number of bird species such as chicken, ducks, turkey, and fowls which are domesticated for their eggs and meat.



Can you tell?

1. What is a layer ?
2. What is a broiler ?

Allied professions to poultry include processing of eggs and meat, marketing of poultry products, compounding and sale of poultry feed, poultry equipment, pharmaceuticals, feed additives, etc.

Selection of proper and disease free breed, suitable and safe farm condition, proper feed and water, hygiene and health care, are important requirements for poultry farm management. On the basis of their origin, different types of poultry breeds are : **American breeds** Plymouth Rock, New Hampshire, Rhode Island Red; **Asiatic breeds** are Brahma, Cochin and Langshan; **Mediterranean breeds** are Leg horn, Minorca; **English breeds** include Australorp; **Indian breeds** are Chittagong, Aseel, Brahma, and Kadaknath.

Leghorn is best layer (for eggs) while Playmouth rock, Rhode Island Red, Aseel, Brahma and Kadaknath, are preferred as broilers (for meat).

Management of layers, requires purchase of high yielding chicken, well ventilated farms, proper feed, debeaking, lighting, waterer, sanitation, culling and vaccination. Management of broilers requires selection of breed, housing, temperature, ventilation, lighting, floor space and broiler feed. Different types of poultry diseases are:

i. **Viral diseases** include Ranikhet, Bronchitis, Avian influenza (bird flu), etc. Few years ago, bird flu had seriously influenced poultry farming and caused human infection too.

ii. **Bacterial diseases** mainly include Pullorum, Cholera, Typhoid, TB, CRD (chronic respiratory disease), Enteritis, etc.

iii. **Fungal diseases** are Aspergillosis, Favus and Thrush.

iv. **Parasitic diseases** include infections by lice, round worm, caecal worm, etc.

v. **Protozoan diseases** e.g. Coccidiosis is a protozoan disease.

D. Apiculture or bee keeping :

Apiculture or bee keeping deals with an artificial rearing of honey bees to obtain bee products like honey, wax, pollens, bee venom, propolis (bee glue) and royal jelly as well as pollinating agents for crop plants.



Fig. 11.4 : Bee hive

Bee keeping is an ancient cottage industry. Honey is a food of high nutritive value and also finds uses in the indigenous system of medicines.



Can you tell?

Why are honey bees called as best pollinators?

The four species of honey bees commonly found in India are *Apis dorsata* (rock bee or wild bee), *Apis florea* (little bee), *Apis mellifera* (European bee) and *Apis indica* (Indian bee).

Knowledge enhancer :

For bee keeping, *Apis mellifera* and *Apis indica* are the suitable species, hence they are known as domesticated species.

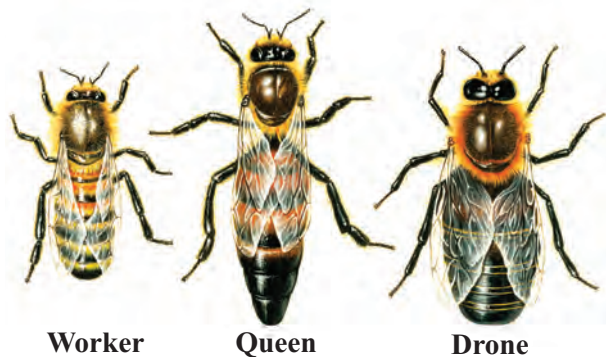


Fig. 11.5 : Honey bee (*Apis mellifera*)

Polymorphism in honey bee :

Bee keeping is practiced in the areas where sufficient wild shrubs, fruit orchards and cultivated crops are present. Bee keeping requires the equipments like bee hive boxes, with comb foundation sheets, bee veil, smoker, bee brush, gloves, gumshoes, uncapping knife, swarm net, queen excluder, overall hive tool, etc.

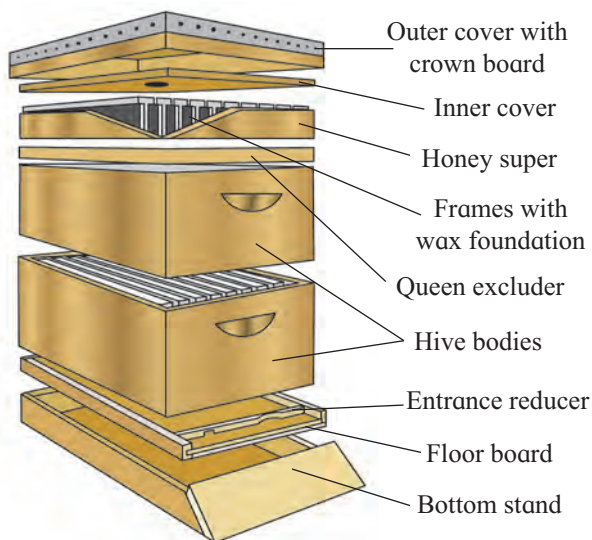


Fig. 11.6 : Artificial bee hive

Artificial bee hive :

For successful bee keeping one must be familiar with the habits of bees, selection of suitable location, catching and hiving of swarms, management of hives during different

seasons, handling and collection of honey, bee wax and other products. Periodic inspection for cleanliness of hive boxes, activity of bees and queen, condition of brood, provision of water, is very much necessary.

Many Indian crop fields need the services of honey bees as the pollinators. Bee keeping in the crop field of sunflower, mustard, safflower, chilly, cabbage, cucumber, legumes, fruits like apple, mango, citrus, etc. help in increasing the productivity of honey as well as crops tremendously.

E. Fishery :

Fishery is a branch of applied biology which deals with the catching, processing, fish farming and marketing of fish, and other useful aquatic animals such as, prawns, lobsters, oysters, mussels and crabs. Three division of fishery are- inland fishery, marine fishery and estuarine fishery.

Inland fishery includes culturing and capturing of fish from fresh water bodies like ponds, lakes, dams and river. Inland aquatic area of our country covers about 40 to 50 lakh acres. The common fresh water fish are *Labeo rohita* (rohu), *Catla* (catla), *Cirrhina mrigala* (mrigala) and other carps.

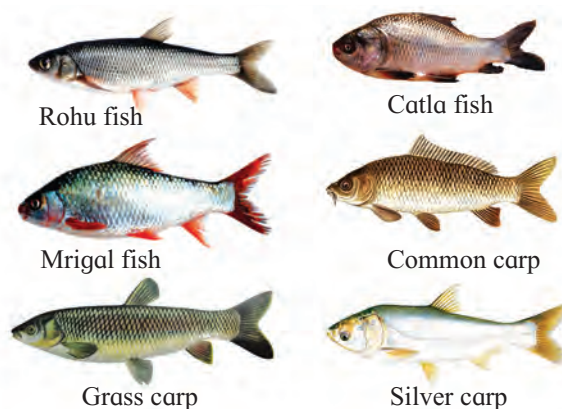


Fig. 11.7 : Fresh water fish forms

Knowledge enhancer :

Fish farming or culturing of edible and commercially important fish is only possible in fresh water bodies.

Marine fishery includes capture of fish from sea water. Indian coastal line is about 7500 km long. The common marine fish are *Harpadon* (Bombay duck), *Sardinella* (sardine), *Rastrelliger* (mackerel) and *Stromateus* (pomphret).



Can you tell?

1. Give the names of estuaries found in Maharashtra and mention their locations.
2. Enlist the names of different fish found at an estuary.

Estuarine fishery includes capture of fish from estuary. Estuary is a place where river meets the sea. e.g. Sunderban area in west bengal.

The common factors for the maintenance of fish farm, includes selection of suitable site, excavation of ponds, requirements of hatchery tank, nursery tank, rearing tank, stocking tank or ponds, water source, manures, supplementary feed, etc. The culture fishery may be monoculture (only one species) or polyculture (many species) type.

After catching of fish, the fish spoilage is prevented by different preservation methods like chilling, freezing, freeze drying, sun drying, smoke drying, salting and canning. In addition to the source as nutritious food, fish yield a number of by-products which are of commercial value. They are fish oil, fish meal, fertilizers, fish guano, fish glue and isinglass. These by-products are widely used in paints, soaps, oils, and medicine. Prawns and Lobsters have market value all over the world. Fishery provides good job opportunities and self employment to many people.

F. Sericulture :

Sericulture is the branch of applied zoology which deals with rearing of silkworm and production of silk. Like other farming, sericulture also involves skill and scientific knowledge for rearing and development. It require less investment and can be started in small space. It is the oldest business and large number of families are associated with the production of silk in India. Disabled, older persons, handicapped people can successfully do this job.

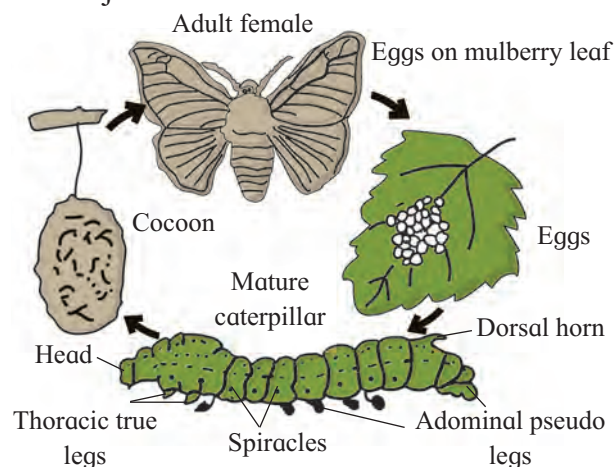


Fig. 11.8 : Life cycle of silk moth (Mulberry silk worm)



Internet my friend

Collect information about life cycle of silk moth.

The best quality silk called mulberry silk, is produced by silkworm *Bombyx mori*, while Tussar silk and Eri silk are of inferior quality. The quality and quantity of silk depends on the quality of mulberry leaves on which the larvae feed.

Rearing, development and looking after the silkworms, involve skill and labour for constant watch. A little negligence can spoil the complete industry. Silkworm larvae may be infected by protozoans, viruses and fungi. Besides these, ants, crows, birds, and other predators are ready to attack these insects, hence the cages of these larvae must be managed to prevent predators attack.

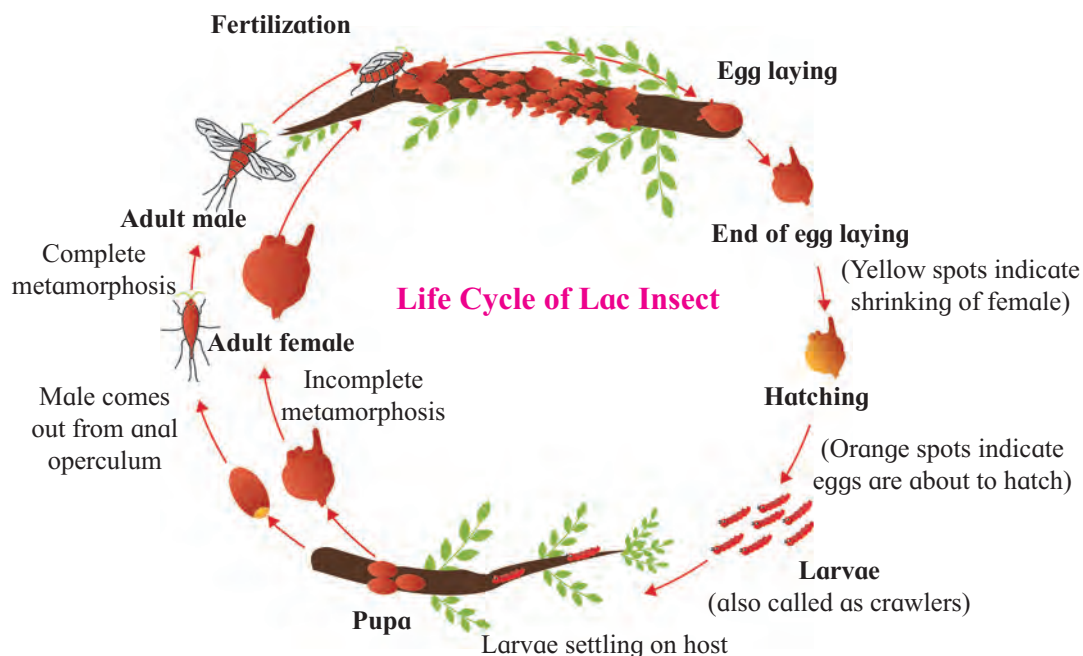


Fig. 11.9 : Life Cycle of Lac Insect



Can you tell?

1. What are the different stages found in life cycle of silkworm?
2. Process of cocoon formation.
3. Which process is involved in silk production from cocoon ?

G. Lac culture :

Lac is produced by an insect *Trachardia lacca*, which is quite small in size and colonial in habit. Resin like substance is produced by Dermal glands of female lac insect. Insect feeds on succulent twigs of certain plants like ber, people, palas, kusum, babool, etc and secretes pink coloured resin, that hardens on coming in contact with air forming lac. It is produced on a large scale all over India.

Lac is a complex substance having large amount of resin together with sugar, water, minerals and alkaline substances.

Natural lac is always contaminated. Shellac is pure form of lac obtained by washing and filtering. Lac insect is a native of India and our share is 85 % of total lac produced in the world. Products of lac play a vital role in the economy of the farmers. Lac is used in bangles, toys, woodwork, inks, mirrors, etc. Production

of lac requires an artificial inoculation of plants which give better and regular supply of good quality and quantity of lac.

11.7 Microbes in human welfare :

Biotechnology is the applications of ‘Scientific and Engineering principles for the processing of materials by biological agents to provide goods and service to humans or for human welfare’.

There are variety of microorganisms like algae, fungi, bacteria, viruses, protozoans, nematodes, etc. Their products having beneficial activities are used for welfare of humans in regard to food, health, industry, agriculture, medicine, biocontrol, etc. These organisms are used variously in food and feed technology, industry, waste utilization, energy, etc.

Microbes in food preparation :

The development of biotechnology occurred in two phases viz, **traditional** (upto 1970) and **modern** (after 1970). Traditional biotechnology is based on the fermentation principle by using fermenting bacteria. These were used in the preparation of variety of indigenous fermented food products.

1. Dosa, Dhokla and Idli:

The dosa, idli and dhokla are fermented products produced due to activity of bacteria. They are fermented preparation of rice and black Gram with air borne *Leuconostoc* and *Streptococcus* species of bacteria. CO₂ produced during fermentation causes puffing up of the dough.



Can you tell?

Name the microbes used in fermentation of dhokla.



Find out

Names of some edible mushrooms and poisonous mushrooms.

2. Microbes as the Source of Food :

Some microbes or their fruiting bodies are directly used as a source of food, as they are rich in vitamins and proteins. The term “SCP” or “single cell protein” denotes, dead and dried cells of microbes like bacteria, algae, molds and yeasts.

Some mushrooms and truffles are directly used as food. They belong to higher fungi. They produce large, fleshy fruiting bodies which are edible. Fruiting bodies are sugar free, fat free but rich in proteins, vitamins, minerals and amino acids. The food in the fruiting body is low calori ed.

3. Dairy Products:

Lactic acid bacteria (LAB) like *Lactobacillus* are added to milk. It ferments lactose sugar of milk into lactic acid. Lactic acid causes coagulation and partial digestion of milk protein casein. Milk is changed into curd, yoghurt and cheese. The starter or inoculum used in preparation of milk products actually contains millions of lactic acid bacteria (LAB).

i. Curd:

Indian curd is prepared by inoculating milk with *Lactobacillus acidophilus*. It also checks growth of disease causing microbes.

ii. Yoghurt (= yogurt):

It is produced by curdling milk with the help of *Streptococcus thermophilus* and *Lactobacillus bulgaricus*.

iii. Butter Milk:

The acidulated liquid left after churning of butter from curd, is called **butter milk**.

iv. Cheese:

It is the partially degraded concentrate of milkfat and protein. The milk is coagulated with lactic acid bacteria and the curd formed is filtered to separate whey. The solid mass is then ripened with growth of mould that develops flavour in it. Different varieties of cheese are known by their characteristic texture, flavor and taste which are developed by different specific microbes. The ‘Roquefort and Camembert cheese’ are ripened by blue-green molds *Penicillium roquefortii* and *P. camembertii* respectively. The large holes in Swiss cheese are developed due to production of a large amount of CO₂ by a bacterium known as *Propionibacterium shermanii*.

11.8 Role of Microbes in Industrial Production:

During fermentation, variety of products like alcoholic beverages, organic acids, vitamins, growth hormones, enzymes, antibiotics, etc. are produced.



Do you know ?

These are actually the secondary metabolites produced during **idio phase** and are not required by micro-organisms for their growth. The type of substrate and the type of micro-organism result into the production of particular secondary metabolites.

Production on an industrial scale requires growing microbes in very large vessels, called **fermenters**. The main function of a fermenter is to provide a controlled environment for growth of a microorganism, or a defined mixture of microorganisms, to obtain the desired product.

a. Production of Alcoholic Beverages :

Beverage is alcoholic or non-alcoholic liquid produced or prepared and used for drinking e.g. tea, coffee, beer, wine, which acts as stimulant. Alcoholic beverages are the products of alcoholic fermentation of specific substrates. Use of microbes in making alcoholic beverages, is known since vedic period- 5000 to 7000Bc.

Microbes especially yeast have been used from time immemorial for the production of beverages like wine, beer, whiskey, brandy or rum. For this purpose, the yeast *Saccharomyces cerevisiae* var. *ellipsoidis* (commonly called Brewer's Yeast) is used for fermenting malted cereals and fruit juices to produce ethanol. Among the beverages, Wine and Beer are produced without distillation whereas whiskey, brandy and rum are distilled beverages. Tubular tower fermenter is used for large scale production of alcohols.

Several traditional drinks and foods are also made through fermentation by yeast. Today, a traditional drink of the coastal region, in South India, is made by fermenting the sugar sap extracted from palm plants and coconut palm. The famous wine of Goa, traditionally known as 'fenny' is made by fermenting fleshy pedicels of cashew fruits.

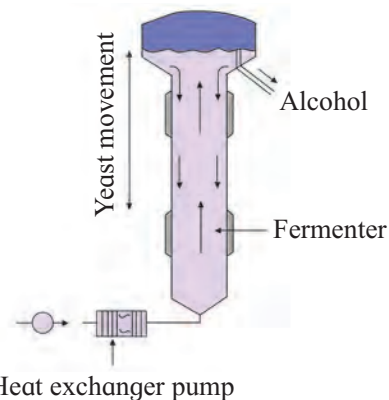


Fig. 11.10 : Tubular tower fermenter

b. Production of organic acids:

Microbes are also used for the commercial and industrial production of certain **organic acids**. These compounds can be produced

directly from **glucose** (e.g. **gluconic acid**) or formed **as end products** from **pyruvate or ethanol**.

Organic acid	Microbes used
i. Citric acid	<i>Aspergillus niger</i>
ii. Gluconic acid	<i>Aspergillus niger</i>
iii. Fumaric acid	<i>Rhizopus arrhizus</i>
iv. Acetic acid (vinegar)	<i>Acetobacter aceti</i>

The organic acids are further used variously e.g. citric acid is used in confectionary, fumaric acid in resins as wetting agents and gluconic acid in medicine for solubility of Ca^{++} .

c. Production of vitamins:

Vitamins are some complex organic nitrogenous compounds required in small amount and are capable of performing many life-sustaining functions inside our body. These compounds cannot be synthesized by humans (except vitamin D), and therefore they have to be supplied in small amounts in the diet.

Microbes are capable of synthesizing the vitamins and hence they can be successfully used for the commercial production of many of the vitamins e.g. thiamine, riboflavin, pyridoxine, folic acid, pantothenic acid, biotin, vitamin B_{12} , ascorbic acid, beta-carotene (pro-vitamin A) and ergosterol (provitamin D).

Vitamins are manufactured by fermentation using different microbial sources as mentioned below:

Name of the vitamin	Microbial source
i. Vitamin B_2	i. <i>Neurospora gossypii</i> ii. <i>Eremothecium ashbyi</i>
ii. Vitamin B_{12}	<i>Pseudomonas denitrificans</i>
iii. Vitamin C	<i>Aspergillus niger</i>



Can you recall?

1. What are antibiotics?
2. Who invented first antibiotic?

d. Production of Antibiotics:

Antibiotics are probably the most important group of compounds synthesized by industrial

microorganisms. Most antibiotics are secondary metabolites. They have therapeutic importance and are used in medical treatment. These are produced in small amounts by certain microbes (like bacteria, fungi and few algae), which inhibit growth of other microbial pathogens. Therefore, they are used in medicine. The antibiotics are antibacterial, antifungal, etc.

Antibiotics have greatly improved our capacity to treat deadly diseases such as plague, whooping cough, diphtheria, leprosy, etc.



Use your brain power

Can antibiotics kill viruses?

Some common antibiotics and their microbial sources are listed below :

Table 11.11 : Antibiotic producing microbes

Antibiotic produced	Microbial sources
Chloromycetin	<i>Streptomyces venezuelae</i>
Erythromycin	<i>Streptomyces erythreus</i>
Penicillin	<i>Penicillium chrysogenum</i>
Streptomycin	<i>Streptomyces griseus</i>
Griseofulvin	<i>Penicillium griseofulvum</i>
Bacitracin	<i>Bacillus licheniformis</i>
Oxytetracycline / Terramycin	<i>Streptomyces aurifaciens</i>



Can you recall?

What are Enzymes? How are they classified?

e. Production of Enzymes:

In living beings, enzymes play a key role in metabolic reactions and therefore, these are essential for the survival of such beings.

Enzymes are proteins known as biocatalysts due to their ability to promote reactions more quickly at body temperature and more efficiently. Many microbes **synthesize and excrete large quantities of enzymes into the surrounding medium**. Using this feature of these tiny organisms, many enzymes are produced commercially. These enzymes are **Amylase, Cellulase, Protease, Lipase, Pectinase, Streptokinase** and many others.

Several industrial sectors, use enzymes from microorganisms for specific applications. In the textile industry, enzymes are able to improve the quality of the fabrics. In the pulp and paper industry, they are involved in biomechanical pulping and bleaching, in the food industry, they are used in the fermentation processes for the production of bread and drinks such as wine and beer, also they participate in the extraction of substances, such as carotenoids and olive oil; lipases are used in detergent industry, because they have superior cleaning properties, increasing the brightness and removing of oil stains; they are also used in cosmetics, animal feed and agricultural industries, among others. Following are the few examples of enzymes used in industrial food processing:

Name of the enzyme Microbial source

Invertase	<i>Saccharomyces cerevisiae</i>
Pectinase	<i>Sclerotinia libertine,</i> <i>Aspergillus niger</i>
Lipase	<i>Candida lipolytica</i>
Cellulase	<i>Trichoderma konigii</i>

f. Gibberellin production:

Gibberellin is a group of growth hormones mainly produced by higher plants and fungi to promote growth by stem elongation. The first gibberellin was isolated by two Japanese scientists -Yabuta and Sumiki (1938) from rice seedlings infected with the fungus *Gibberella fujikouri*.

Extra information: *Streptokinase* enzyme (TPA) is produced by the bacterium *Streptococcus spp.* It has fibrinolytic effect. Hence, it is used as a 'clot buster' for clearing blood clots in the blood vessels of patients, which may cause heart attack.

Statins produced by the yeast *Monascus purpureus* have been produced on commercial scale. It is a blood-cholesterol lowering agent. This agent acts as competitive inhibitor of the enzyme responsible for synthesis of cholesterol.

About 15 different types of gibberellins have been isolated. Gibberellins have many practical applications. They are used to induce parthenocarpy in apple, pear, etc. They are used in breaking the dormancy of seed and also in inducing flowering in Long Day Plants (LDP). They are also used to enlarge the size of grape fruits.

11.9 Microbes in Sewage Treatment:

Sewage is a matter carried off in drainage. It is a municipal waste containing human excreta, house hold waste, dissolved organic matter and even pathogenic microbes (bacteria, viruses, protozoans, nematodes and microfungi). It also includes discharged water from hospital waste, slaughter house waste, animal dung, etc. Discharge from industrial waste (contains toxic dissolved organic and inorganic chemicals), tannery, pharmaceutical waste, etc. also add to sewage.

Extra Information : Sewage is also a potential source of pathogenic bacteria, viruses and protozoa. The causative agents of dysentery, cholera, typhoid, polio and infectious hepatitis may occur in sewage. The bacteria from the soil are also present in the sewage. During the course of sewage decomposition, initially aerobic and facultative anaerobic organisms predominate which are then followed by strict anaerobic especially methanogenic bacteria that produce methane (CH₄) and CO₂.



Do you know ?

The oil eating bacteria can clean up crude spills. Collect more information about these bacteria.

Composition of Sewage:

Sewage consists of approximately 99.5% to 99.9% water and 0.1 to 0.5% inorganic and organic matter in suspended and soluble form. Composition of sewage varies depending upon the type of waste discharged into water from different industries. e.g. textile, chemicals, pharmaceuticals, dairy, canning, brewing, meat packing, tannery, oil refineries and meat industries, etc.

Microorganisms in Sewage:

Various types of micro-organisms are also present in sewage. Bacteria, viruses, fungi, protozoa, nematodes, algae, etc. are found in sewage. However, their number and type, fluctuate depending upon the sewage composition and source of sewage. Raw sewage may contain millions of bacteria per ml. These include coliforms, fecal *Streptococci*, anaerobic spore forming *bacilli* and other types originating in the intestinal tract of humans.

Before waste water is made available for human use, it has to be treated properly, so as to remove organic matter, inorganic salts and pathogens as well. **Sewage treatment process includes four basic steps as follows:**

1. Preliminary Treatment:

The preliminary treatment includes Screening and Grit Chamber.

i. Screening : Sewage and waste water contains plenty of suspended, floating materials, coarse and solid particles along with dissolved substances. The suspended objects are filtered and removed. This is done in screening chambers. The sewage is passed through screens or net in the chambers. Larger suspended or floating objects are held back in the screening

chambers. These have to be removed before the biological treatment.

ii. Grit Chamber : After screening, the filtered sewage is then passed into series of grit chambers. These chambers contain large stones (pebbles) and brick-ballast. Coarse particles settle down by gravity. Thus, passage of filtered sewage removes much of the coarse particulate matter.

2. Primary treatment (physical treatment):

After the preliminary treatment, the sewage water is pumped into the **primary sedimentation tank**. The sedimentation of suspended solid or organic matter occurs in this tank. About 50-70% of the solids settle down. There is reduction of about 30-40% (in number) of *coliform* organisms. The organic matter which is settled down, is called **primary sludge** which is removed by mechanically operated devices. The supernatant (effluent) in the primary sedimentation tank still contains large amount of dissolved organic matter and micro-organisms which can then be removed by the secondary treatment.

3. Secondary treatment (biological treatment):

The primary effluent is passed into large aeration tanks. Here it is constantly agitated

mechanically and air is pumped into it. Aerobic bacteria grow vigorously and form **flocs**. Flocs are the masses of bacteria held together by slime and fungal hyphae to form mesh like masses. These aerobic microbes consume the major part of the organic matter present in the effluent, as they grow. Due to this BOD (Biochemical Oxygen Demand) of the effluent is significantly reduced.

4. Tertiary treatment:

Once the BOD of waste water is reduced, it is passed into a **settling tank**. Here the bacterial flocs are allowed to sediment. The sediment is now called **activated sludge**. Small part of this is passed back in to aeration tank and the major part is pumped in to large tanks called **anaerobic sludge digesters**. In these tanks, anaerobic bacteria grow and digest the bacteria and fungi in the sludge. During this anaerobic digestion, gases such as methane, hydrogen sulphide, CO₂, etc. are produced. Effluents from these plants (digester) after chlorination, are released in natural water bodies like rivers and streams. Chlorination kills pathogenic bacteria. Digested Sludge is then disposed.

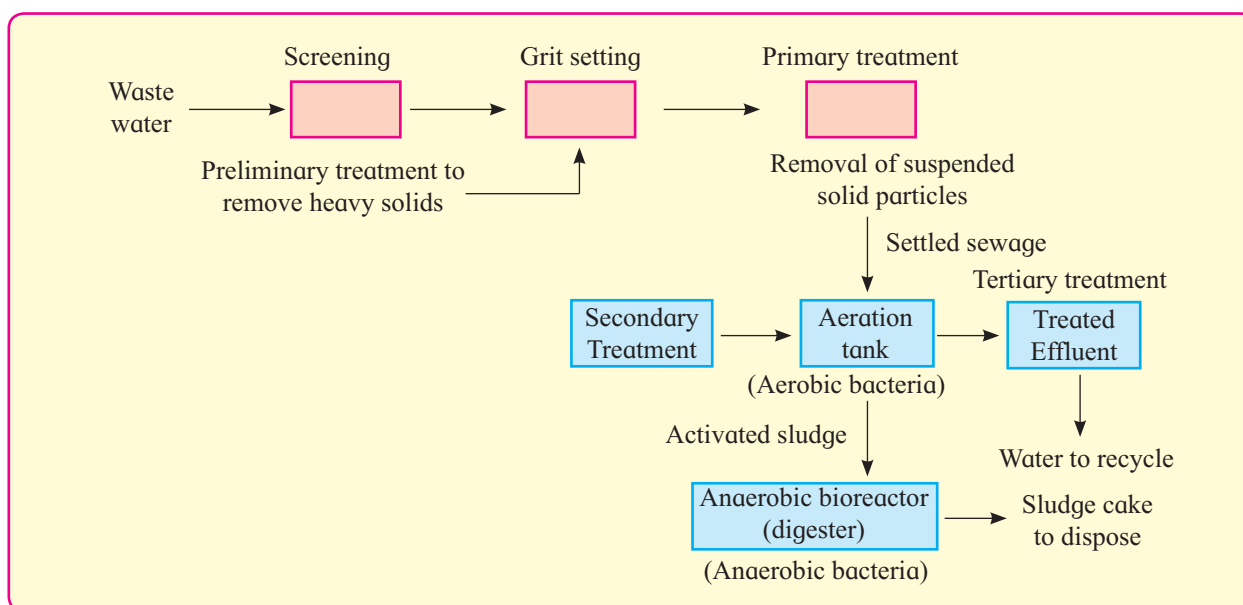


Fig. 11.12 : Diagrammatic representation of various stages in wastewater treatment

11.10 Microbes in Energy Generation :

Many developing countries are encouraging for installation of biogas plants to meet out the requirement of energy. Biogas is used as a domestic as well as industrial fuel. It is a non-conventional and renewable source of energy and is obtained by microbial fermentation. Biogas is a mixture of methane CH_4 (50-60%), CO_2 (30-40%), H_2S (0-3%) and other gases (CO , N_2 , H_2) in traces. Biogas is highly inflammable and is used as a source of energy.

Plant wastes, animal wastes, domestic wastes, agriculture waste, municipal wastes, forestry wastes, etc. are commonly used for biogas production. Cattle dung is most commonly employed substrate for biogas production. It is a rich source of cellulose from plants.

Biogas Production:

Most commonly used models of biogas plants are KVIC and IARI. The digester used for biogas production is called **Biogas Plant**. A typical biogas plant using cattle dung as a raw material, consists of digester and gas holder. Digester is made up of concrete bricks and cement, or steel. There is cylindrical gas holder or gas tank above it to collect gases. Digester has a side opening (charge pit) into which raw material as cow dung is fed. The digester is partly burried in the soil.

Anaerobic digestion involves in three processes :

i. Hydrolysis or solublization : In initial stage raw material (cattle dung) is mixed with water in equal proportion to make slurry which is then fed into the digester. Here anaerobic hydrolytic bacteria (e.g. *Clostridium*, *Pseudomonas*) hydrolyse carbohydrates into simple sugars, proteins into amino acids and lipids into fatty acids.

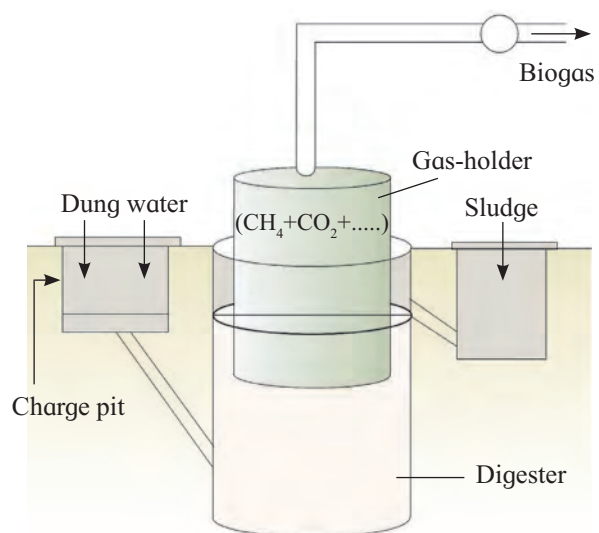


Fig. 11.13 : Biogas plant

ii. Acidogenesis : In this stage, facultative anaerobic, acidogenic bacteria and obligate anaerobic organisms, convert simple organic material into acids like formic acid, acetic acid, H_2 and CO_2 .

iii. Methanogenesis :

This is last stage in which anaerobic Methanogenic bacteria like *Methanobacterium*, *Methanococcus* convert acetate, H_2 and CO_2 into Methane, CO_2 and H_2O and other products.

- $12\text{mol CH}_3\text{COOH} \longrightarrow 12\text{CH}_4 + 12\text{CO}_2$
(acetic acid) methane
- $4\text{mol H.COOH} \longrightarrow \text{CH}_4 + 3\text{CO}_2 + 2\text{H}_2\text{O}$
(formic acid)
- $\text{CO}_2 + 4\text{H}_2 \longrightarrow \text{CH}_4 + 2\text{H}_2\text{O}$

Benefits:

- It is a cheap, safe and renewable source of energy. It can be easily generated, stored and transported.
- It can be used for domestic lighting, cooking, street lighting as well as small scale industries.
- It burns with blue flame and without smoke.
- It helps to improve sanitation of the surrounding.
- It is eco-friendly and does not cause pollution and imbalance of the environment. Sludge which is left over is used as a fertilizer.

11.11 Role of Microbes as Biocontrol Agents:

The term biocontrol refers to the use of biological methods to control diseases and pests. The natural method of eliminating and controlling insects, pests and other disease-causing agents, is by using their natural, biological enemies. This is called **biocontrol** or **biological control**.

The agents which are employed for this are called **biocontrol agents**. Microbes are one among them. These microbes include bacteria, fungi, viruses and protozoans. Microbes as biocontrol agents act in three ways, either they cause the disease to the pest or compete or kill them. Chemicals, insecticides and pesticides are extremely harmful to human beings and also pollute our environment. Hence, the use of biocontrol measures will greatly reduce our dependence on toxic chemicals and pesticides.

Examples of Microbial bio-control :

- i. *Bacillus thuringiensis* (Bt) is used to get rid of butterfly, caterpillars where dried spores of *Bacillus thuringiensis* are mixed with water and sprayed onto vulnerable plants such as *Brassicacae* and fruit trees. These spores are then eaten by the insect larvae. In the gut of the larvae, the toxin (cry protein) is released and the larvae get killed eventually.
- ii. *Trichoderma* species are free-living fungi found in the root ecosystem (rhizosphere). These are effective as biocontrol agents of several soil borne fungal plant pathogens. The fungus produces substances like viridin, gliotoxin, gliovirin, etc. that inhibit the other soil borne pathogens attacking root, rhizomes, etc. causing rot disease.

Four groups of biocontrol agents are known. They are bacteria, fungi, viruses and protozoans.

I. Microbial Pesticides and their host :

The correlation is depicted as per the following table :

Table 11.14 : Microbes and their host

Pathogen (biocontrol agent)	Host (target pest)
Bacteria: <i>Bacillus thuringiensis</i> (Bt), <i>B. papilliae</i> and <i>B. lentimorbus</i>	Caterpillars, cabbage worm, adult beetle, etc.
Fungi: <i>Beauveria bassiana</i> , <i>Entomophthora pallidroseum</i> , and <i>Zoopthora radicans</i>	<i>Aphid crocci</i> , <i>A. unguiculata</i> , mealy bugs, mites, white flies etc
Protozoans: <i>Nosema lacustae</i>	Grasshopper, caterpillars, crickets
Viruses: Nucleopolyhedrovirus (NPV) and Granulovirus (GV)	Caterpillars and Gypsy moth, ants, wasps and beetles.

Bioherbicides :

Weeds are the unwanted plants that grow in agricultural fields, ponds, lakes, etc. Weeds compete with the main crop in the farm-land for water, space, minerals, light, air, etc. and also act as collateral hosts for several pathogens.

Microbes are also used as herbicides. Many dicot herbs that grow as weeds in the field of cereals, can be killed by certain microbes. Some examples are as below :

II. Microbial Herbicides and Their Host :

1. Pathogenic fungi as mycoherbicides:
 - i. *Phytophthora palmivora* - controls milk weed in orchards.
 - ii. *Alternaria crassa* - controls water hyacinth.
 - iii. *Fusarium spp.* - control most of the weeds.
2. Bacterial pathogen as herbicides:
 - i. *Pseudomonas spp.* - attacks several weeds
 - ii. *Xanthomonas spp.* - attacks several weeds
 - iii. *Agrobacterium spp.* - attacks several weeds

3. Insects as herbicides :
 - i. *Tyrea* moth - controls the weed *Senecio jacobea*
 - ii. *Cactoblastis cactorum* - controls cacti weeds.

11.12 Role of Microbes as Biofertilizers:

Fertilizers are nutrients which are necessary for the growth of plants and thus for the productivity of cultivated plants. Use of fertilizers for increasing productivity is one of the aspects of green revolution. Fertilizers are classified as **inorganic** (chemical) and **organic** (biological). Inorganic fertilizers are synthetic where mineral salts of NPK are mixed in definite proportion and then dusted in the field. Non-judicious or excessive use of such fertilizers lead to pollution of soil, air and ground water. Soil becomes acidic.

Organic fertilizers are biological in origin and include Farm Yard Manure (FYM), compost and green manure. Use of these fertilizers increases the fertility of soil.

Now a days for better and sustainable agricultural production, farmers use **biofertilizers** and practise organic farming. Biofertilizers are mostly N_2 fixing, living microorganisms which enrich the nutrient quality of soil. They include bacteria, cyanobacteria and fungi.

Biofertilizers are commercial preparation of ready-to-use **live** bacterial or fungal formulations. Their application to plant, soil or composting pits, helps to enrich the soil fertility due to their biological activity.

Use of Biofertilizers is cost effective and eco-friendly. They play a vital role in maintaining a long term soil fertility and sustainability.

Types of Biofertilizers:

On the basis of nature and function biofertilizers are divided into following groups-

1. **N_2 fixing Biofertilizers:** The nitrogen fixing microorganisms which convert atmospheric nitrogen into nitrogenous compounds like nitrites and nitrates via ammonia. Nitrogen fixing microorganisms, also called **diazotrophs**, are of two types:
 - i. Symbiotic N_2 fixing microorganisms: for eg. *Rhizobium*, *Anabaena*, *Frankia*. These are always associated generally with underground parts i.e. roots of higher plants.
 - ii. Free-living or Non- Symbiotic N_2 fixing microorganisms: e.g. *Azotobacter*, *Nostoc*, *Clostridium*, *Beijerinckia*, *Klebsiella*, etc.
2. **Phosphate solubilizing biofertilizers:** These are the bacterial species which solubilize the insoluble inorganic phosphate compound, such as rock phosphate. For eg. *Pseudomonas striata*, *Bacillus polymyxa*, *Agrobacterium*, *Micrococcus*, *Aspergillus spp.*, etc.
3. **Compost making biofertilizers:** Composting is a natural process that turns organic material into a dark rich substance called as compost or humus. The composting process is dependent on microorganisms to break down organic matter into compost. There are many types of microorganisms found in active compost such as bacteria, fungi, actinobacteria, protozoa and rotifers.
4. **Cyanobacteria as biofertilizers :** Many cyanobacteria are aquatic and terrestrial, free-living or symbiotic, aerobic, photosynthetic, N_2 fixing, heterocystous or non-heterocystous forms. e.g. *Anabaena*, *Nostoc*, *Plectonema*, *Oscillatoria*, etc. *Anabaena*, *Nostoc* and *Tolypothrix* are associated with lichens while *Anabaena* is associated with plants like *Azolla* and *Cycas*.



Do you know ?

Classification of Biofertilizers :

On the basis of **nature** or group of organisms, biofertilizers are classified as **bacterial** fertilizers and **fungal** fertilizers.

Bacterial fertilizers include eubacteria and cyanobacteria. On the basis of **function**, bacterial fertilizers are further grouped as **nitrogen fixing, phosphate solubilizing** and **compost making** biofertilizers. Cyanobacterial biofertilizers, on the basis of function, are nitrogen fixing type.

Fungal biofertilizers include mycorrhizal fungi. On the basis of function, they are classified as **ectomycorrhizae** and **endomycorrhizae**.

3. Fungal biofertilizers:

Mycorrhiza is a fungus. It forms symbiotic association with the underground parts like rhizomes and roots of higher plants occurring in thick humid forests. These were discovered by Frank (1885). There are two types viz, Ectomycorrhizae and Endomycorrhizae.

I. **Ectomycorrhizae:** They have well developed mycelium that forms mantle on the outside of the roots. This increases absorptive surface area of roots and accelerates uptake of water and nutrients (N, P, Ca and K). Due to this the plant vigour, growth and yield, increase. Some hyphae of mycorrhizal fungus, penetrate into the root and forms **hartig-net** in the intercellular spaces of root cortex.

II. **Endomycorrhizae:** They grow in between and within the cortical cells of roots. Fungal hyphae penetrate the cells and form finely branched **arbuscules** intracellularly and form **vesicles** mostly in the intercellular spaces of cortical cells. Hence they are called Vesiculo Arbuscular Mycorrhizae or VAM. Now a days they are described as AM fungi. The plants with VAM grow luxuriantly in

less irrigated lands. The association of VAM with crop plants helps in conversion of less productive field into more productive field.

Benefits of Mycorrhiza :

1. Selective absorption of P, Zn, Cu, Ca, N, Mn, Br and Fe.
2. Enhance water uptake.
3. Induce growth by secreting hormones.
4. Offer protection to host plant from other microbes, by secreting antibiotics.



Do you know ?

Now a days, mycorrhiza are classified into 8 different types viz, Ectomycorrhizae, Endomycorrhizae, Ectendomycorrhizae, Orchidaceous mycorrhizae, Ericoid mycorrhizae, Arbutoid mycorrhizae, Monotropoid mycorrhizae and Ophioglossoid mycorrhizae.



Use your brain power

Why are healthy root nodules pink in colour?

Biofertilizer microorganisms:

1. **Rhizobium:** Rhizobia are rod shaped, motile, aerobic, gram negative, non spore forming, nitrogen-fixing bacteria containing *Nod* genes and *Nif* genes. They form symbiotic association with roots of leguminous plants. They bring about nodule formation on the roots and multiply inside the nodule. They fix atmospheric nitrogen into organic forms, which can be used by plants as nutrients. For eg. *R. leguminosarum* is specific to pea. and *R. phaseoli* to beans.
2. **Azotobacter:** It is an important and well known free living, nitrogen fixing, aerobic, non-photosynthetic, non-nodule forming bacterium which is intimately associated with roots of grasses and certain plants. It is used as a bio-fertilizer for all non-leguminous plants especially rice, cotton, vegetables, etc.

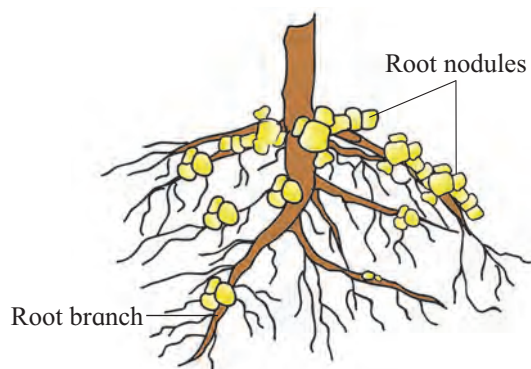


Fig. 11.15 : Root system of Leguminous plant

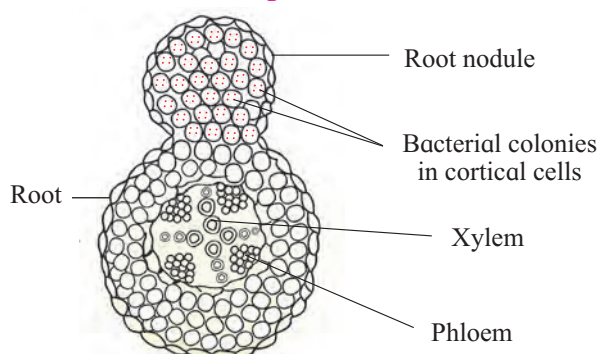


Fig. 11.16 : T. S. of root nodule



Fig. 11.17 : Azolla

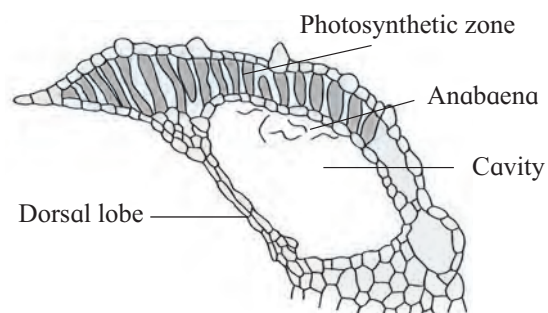


Fig. 11.18 : L. S. of Azolla leaf showing filamentous Anabaena

3. **Azospirillum:** It is free living, aerobic nitrogen fixing bacterium associated with roots of corn, wheat and jowar. It fixes the considerable quantity of nitrogen (20-40kg N/ha) in non – leguminous plants such as cereals, millets, cotton, oilseed, etc.
4. **Anabaena :** It is a genus of multicellular, filamentous cyanobacteria that exists as plankton. It has ability to fix nitrogen and also forms symbiotic relationships with certain plants, such as the coralloid roots of *Cycas* and *Anthoceros* thallus. It has some specialized and colourless cells, called **Heterocysts** which are the sites for nitrogen fixation.
5. **Azolla:** *Azolla* is a free-floating water fern. *Azolla* plant consist of a floating rhizome (stem) with small overlapping bi-lobed leaves and roots. The leaf shows dorsal and ventral lobe.

In the dorsal lobe, *Anabaena* filaments are present in the aerenchyma, which fixes nitrogen. *Azolla* can be used as biofertilizer in the rice field.

Benefits of Biofertilizers :

1. Low cost and can be used by marginal farmers.
2. Free from pollution hazards.
3. Increase soil fertility.
4. BGA as biofertilizers secrete growth promoting substances, organic acids, proteins and vitamins.
5. *Azotobacter* supply nitrogen and antibiotics in the soil.
6. Biofertilizers increase physico-chemical properties of soil- like texture, structure, pH, water holding capacity of soil by providing nutrients and organic matter.

Now in our country many biofertilizers are available in market to reduce the use of chemical fertilizers and thus, the pollution.

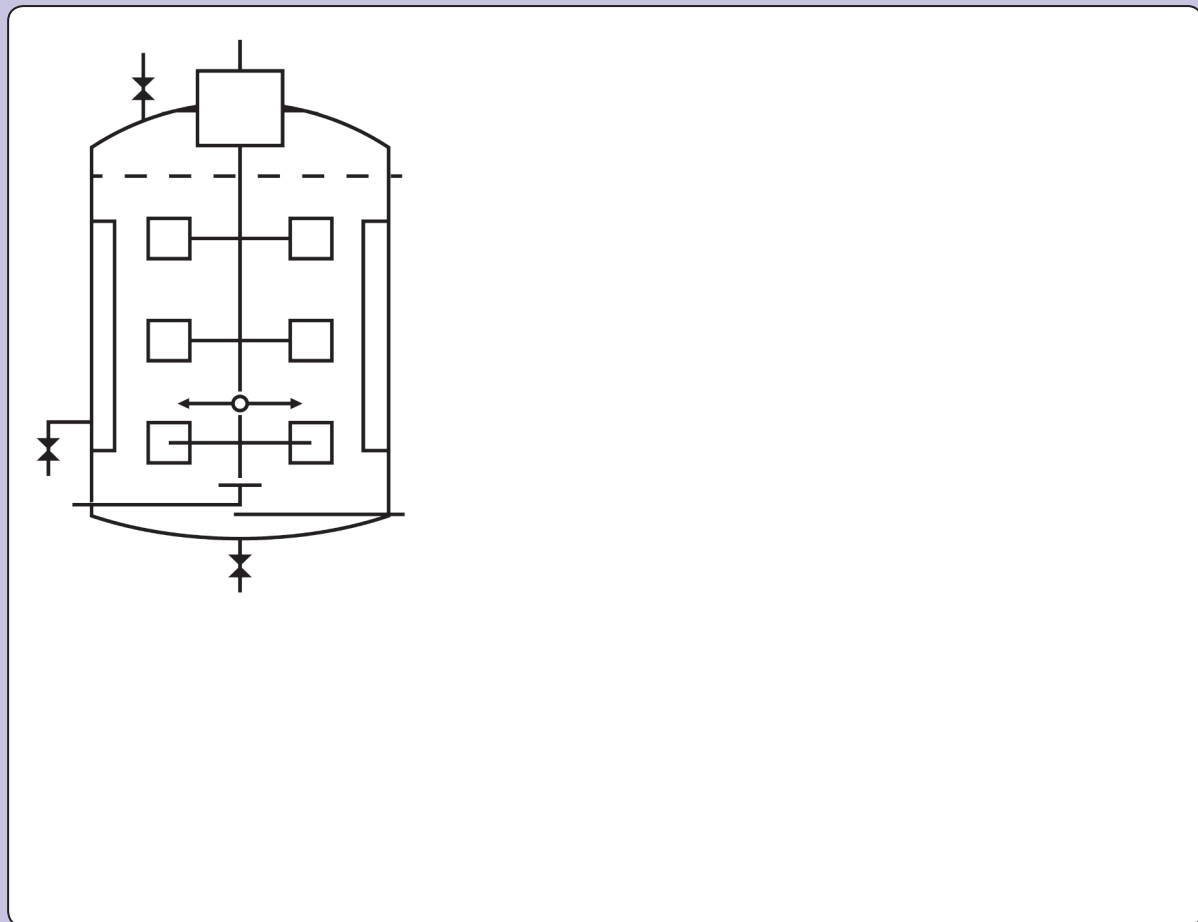


Activity :

1. Visit fish market and enlist different types of fresh water and marine water fish mentioning their local and scientific names along with their salient features.

Type	Local Name	Scientific Name	Detail Information
1.			
2.			
3.			
4.			
5.			
6.			
7.			

2. Label the different parts of the fermenter and mention functions of each part.



Exercise

Q. 1 Multiple Choice Questions.

- Antibiotic Chloromycetin is obtained from
 - Streptomyces erythreus*
 - Penicillium chrysogenum*
 - Streptomyces venezuelae*
 - Streptomyces griseus*
- Removal of large pieces of floating debris, oily substances, etc. during sewage treatment is called
 - primary treatment
 - secondary treatment
 - final treatment
 - amplification
- Which one of the following is free living bacterial biofertilizer?
 - Azotobacter*
 - Rhizobium*
 - Nostoc*
 - Bacillus thuringiensis*
- Most commonly used substrate for industrial production of beer is
 - barley b. wheat
 - corn d. sugarcane molasses
- Ethanol is commercially produced through a particular species of
 - Aspergillus*
 - Saccharomyces*
 - Clostridium*
 - Trichoderma*
- One of the free-living anaerobic nitrogen-fixer is
 - Azotobacter* b. *Beijerinckia*
 - Rhodospirillum* d. *Rhizobium*
- Microorganisms also help in production of food like
 - bread b. alcoholic beverages
 - vegetables d. pulses
- MOET technique is used for
 - production of hybrids
 - inbreeding
 - outbreeding
 - outcrossing
- Mule is the outcome of
 - inbreeding
 - artificial insemination
 - interspecific hybridization
 - outbreeding

Q. 2 Very Short Answer Questions:

- What does make idlies puffy?
- Name any two bacterial biofertilizers.
- What is the microbial source of vitamin B₁₂?
- What is the microbial source of enzyme Invertase?
- Milk start to coagulate when Lactic Acid Bacteria (LAB) is added to warm milk as a starter. Mention any two other benefits of LAB.
- Name the enzyme produced by *Streptococcus* bacterium. Explain importance in medical sciences.
- What is breed?
- Define estuary.
- What is shellac?

Q. 3 Short Answer Questions.

- Many microbes are used at home during preparation of food items. Comment on such useful ones with examples.
- What is biogas? Write in brief about the production process.
- Write a note on biocontrol agents.
- Name any two enzymes and antibiotics with their microbial source.
- Write principles of farm management.
- Give economic importance of fishery.
- Enlist the species of honey bee mentioning their specific uses.

8. What are A, B, C and D in the table given below :

Types of microbe	Name	Commercial product
Fungus	A	Penicillin
Bacterium	<i>Acetobacter aceti</i>	B
C	<i>Aspergillus niger</i>	Citric acid
Yeast	D	Ethanol

Q. 4 Long Answer Questions.

1. Explain the process of sewage water treatment before it can be discharged into natural bodies. Why this treatment is essential?
2. Write a note on lac culture.
3. Describe various methods of fish preservation.
4. Give an account of poultry diseases.
5. Give an account of mutation breeding with examples.
6. Describe briefly various steps of plant breeding methods.

Project :

Collect information about :

- A) Different types of worms/organisms used in sericulture.
- B) How is silk obtained and isolated ?
- C) Types of silk.
- D) Job potential of sericulture.



Can you recall?

1. What is Biotechnology?
2. How do genetically modified organisms are produced?
3. Which are the benefits of Biotechnology?

You are already aware of what biotechnology is. It is the product of interaction between the biological science and technology. It is infact, an applied branch of biology. The term biotechnology was first used by Karl Ereky in 1919 to describe a process for large scale production of pigs.

12.1 Biotechnology

It is defined by different organizations in different ways. It has been brodly defined as ‘the development and utilization of biological forms, products or processes for obtaining maximum benifits to man and other forms of life’. According to OECD (Organization for Economic Cooperation and Development, 1981)- ‘It is the application of scientific and engineering principles to the processing of materials by biological agents to provide goods and service to the human welfare’. It uses scientific priciples of microbiology, genetics, biochemistry, chemical engineering, mathematics, statistics, computers, industrial processes, etc. Biological agent means plants and animal cells, microorganisms, enzymes or their products.

History of origin of biotechnology is as old as human civilization. Development of biotechnology in terms of its growth, occurred in two phases viz, Traditional biotechnology and Modern biotechnology.

Traditional biotechnology (old biotechnology) was primarily based on fermentation technology using microorganisms as in the preparation of curd, ghee, soma,

vinegar, yogurt (yoghurt), cheese making, wine making, etc. It became an art of kitchen in indian houses. It was more an art than science. Till that time people did not know as to how exactly the process occurs and the organisms causing this process. The contributions made by several chemists, biochemists and microbiologist, over the time, could explain the mechanism of process and also the nature of microorganisms causing the process.

During 1970 a new technique of ‘recombinant DNA technology was developed and then established by Stanley Cohen and Herbert Boyer in 1973. This technique has changed the overall outlook, then. The technique permits to change/ modify genetic (heritable) material for getting new specific products. The combination of biology and production technology based on genetic engineering evolved into **modern biotechnology** (new biotechnology).

There are two major features of technology that differentiate modern biotechnology from classical or old biotechnology viz,

- i. Capability of science to change the genetic material for getting new specific products through rDNA technology, polymerase chain reaction (PCR), microarrays, cell culture and fusion, and bioprocessing.
- ii. Ownership of technology and its socio-political impact.

Now the conventional industries, pharmaceutical industries, agro industries, food industries, etc. are also focussing attention to produce biotechnology- based products.

12.2 Principles and Processes of Biotechnology :

Modern biotechnology is based on two core techniques viz. genetic engineering and chemical engineering.

Genetic engineering deals with alteration of genetic material (DNA and RNA) while chemical engineering deals with maintaining sterile environment for manufacturing variety of useful products including vaccines, antibodies, enzymes, organic acids, vitamins, therapeutics, etc.

Genetic engineering is defined as the manipulation of genetic material towards a desired end and in a directed and predetermined way, using “in vitro” process. Manipulation of genes involve repairing of the defective genes or replacing of defective genes by healthy genes or normal genes; artificially synthesizing a totally new gene; transfer of genes into a new location or into a new organism; introducing an altogether new gene; manipulation of genes for improvement of living organisms; combining of genes from two organisms, altering the genotype; gene cloning etc. Therefore, the genetic engineering is alternatively called **recombinant DNA technology** or **gene cloning**.

Extra information :

John E. Smith (1996) gave definition of genetic engineering as ‘the formation of new combination of heritable material by the insertion of nucleic acid molecule produced by whatever means outside the cells, into any virus, bacterial plasmid or other vector system so as to allow their incorporation into a host organism in which they do not occur naturally but in which they are capable of continued propagation’.

Technique of gene cloning and rDNA technology :

In gene cloning, a gene of known function can be transferred from its normal location into a cell (that of course does not contain it) via a suitable vector. The transferred gene is replicated normally and is handed over to the next progeny.

A. Tools and techniques for gene cloning/ rDNA technology :

Before we venture into a procedure of gene cloning, let us know briefly, the basic requirements for the technique.

I. Different instruments (devices) :

Macromolecule such as DNA, RNA, proteins, etc. are synthesized in the living cells which vary in their molecular weight, solubility, presence of charges, absorbance of light, etc. Several techniques are used to isolate and characterize the macromolecules. The size of different types of molecules varies and therefore their molecular weights also vary. The techniques used on the basis of molecular weight, are gel permeation, ion exchange chromatography, spectroscopy, mass spectrometry, electrophoresis, etc. Electrophoresis is the separation of charged molecules, applying an electric field. It is applied for the separation of DNA, RNA and proteins. DNA being negatively charged, migrates to anode. Small fragments of DNA molecules, move faster and thus separate faster. Use of Agarose gel electrophoresis, PAGE, SDS PAGE are the different methods of electrophoresis.

Polymerase chain reaction (PCR) :

Polymerase chain reaction (PCR) is another device used for gene cloning or gene multiplication in vitro. It is the amplification of gene of interest, through PCR.

In 1985, Kary Mullis made an important discovery (contribution) in the form of an extremely powerful technique called polymerase chain reaction (PCR). PCR can generate a billion copies of the desired segment of DNA or RNA, with high accuracy and specificity, in a matter of few hours. The process of PCR is completely automated and involves automatic thermal cycles for denaturation and renaturation of double stranded DNA.

The device required for PCR is called thermal cycler.

PCR is *in vitro* amplification of a desired DNA segment, which requires: DNA containing the desired segment to be amplified, several molecules of four deoxyribonucleoside triphosphates (dNTPs), excess of two primer molecules, heat stable DNA polymerase and appropriate quantities of Mg^{++} ions.



Internet my friend

Collect the information from internet about PCR machine.



Mechanism of PCR:

At the start of PCR, the DNA segment, and excess of two primer molecules, four deoxyribonucleosides triphosphates and the thermostable DNA polymerase are mixed together in 'ependorf tube' and the following operations are performed sequentially (Figure).

Step i : The reaction mixture is heated to a temperature (90–98 °C) to separate two strands of desired DNA. This is called denaturation.

Step ii : The mixture is allowed to cool (40–60 °C) that permits pairing of the primer to the complementary sequences in DNA. This step is called annealing.

Step iii : The temperature (70–75 °C) allows thermostable Taq DNA polymerase to use single-stranded DNA as template and adds nucleotides. This is called primer extension. It takes around two minutes duration.

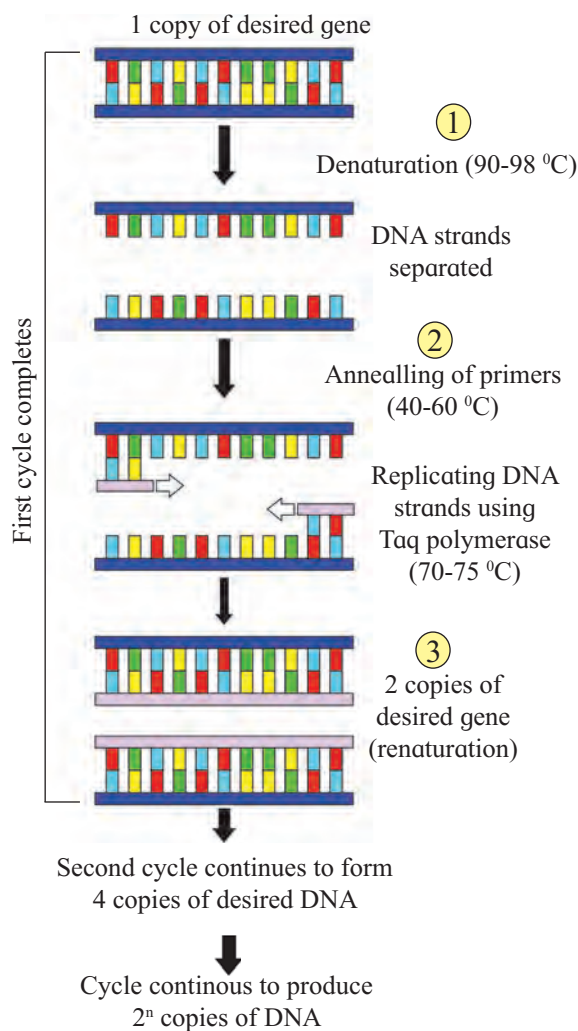


Fig. 12.1 : DNA replication through polymerase chain reaction.

One cycle takes around 3 to 4 minutes. To begin second cycle, DNA is again heated to convert double stranded DNA into single strands.

In an automatic thermal cycler, the above three steps are automatically repeated 20-30 times. Thus, at the end of 'n' cycles 2^n copies of DNA segments, are produced. The machine performs the entire operations automatically and precisely.

Once the desired number of cycles is completed, the amplified DNA segment is purified by gel electrophoresis. After its sequencing, the amplified DNA segment can be inserted into a cloning vector. Desired gene can also be obtained from gene library.

II. Biological tools :

There are three types of biological tools used viz, enzymes, cloning vectors (vehicle DNA) and competent host (cloning organisms) for transformation with recombinant DNA.

A. Enzymes : Different enzymes include Lysozymes, Nucleases such as exonucleases endonucleases, restriction endonucleases, DNA ligases, DNA polymerases, alkaline phosphatases, reverse transcriptases, etc.

i. Restriction enzymes :

Enzymes that cut the phosphodiester bonds of polynucleotide chains are called **nucleases**. These are of two types - exonuclease and endonuclease. Exonucleases cut nucleotides from the ends of DNA strands whereas endonuclease cut DNA from within. During the 1970s, it was found that bacteria contain nucleases that would recognize short nucleotide sequence with duplex DNA and cut.

The phosphodiester back bone at highly specific sites on both strands of duplex, is cut by these enzymes, called **restriction endonucleases (REN)** or simply **restriction enzymes (RE)**. They were given this name because they are used by the bacteria to destroy various viral DNAs that might enter the cell, thereby restricting the potential growth of the virus.

Thus, restriction enzymes serve as defence mechanism. The bacteria protect its own DNA from nucleolytic attack by methylating the bases at susceptible sites, a chemical modification that blocks the action of the enzyme.

The restriction enzymes are thus the molecular scissors that are used to recognize and cut DNA at specific sequences. The sites recognized by them, are called **recognition sequences** or **recognition sites**. Different restriction enzymes found in different organisms recognize different nucleotide sequences and therefore cut DNA at different sites. Table encloses list of some restriction endonucleases and the site at which they cleave DNA.



Activity :

Find out the biological source of following restriction enzymes and discuss their recognition sequences :

Pst I, Sal I, Taq I, Xer III, Mbo II, Hpa I, BgII, Kpn I, Not I.

ii. Recognition sequences :

The sequences recognized by restriction enzymes are 4 to 8 nucleotides long and characterized by a particular type of internal symmetry. Consider the particular sequence recognized by the enzyme EcoRI .

Table 12.2 : Source and recognition sequences (indicated by arrow) of various restriction enzymes:

Restriction Enzyme	Source (Organism and strain)	Recognition sequence	Product	End products produced
Alu I	<i>Arthobacter luteus</i>	$ \begin{array}{c} 5' \text{ ---A- G- } \downarrow \text{ -C -T--- } 3' \\ 3' \text{ ---T- C- } \uparrow \text{ -G-A--- } 5' \end{array} $	$ \begin{array}{cc} \boxed{5' \text{ ---A-G}} & \boxed{\text{C-T---} 3'} \\ \boxed{3' \text{ ---T-C}} & \boxed{\text{G-A---} 5'} \end{array} $	Blunt ends
Bam HI	<i>Bacillus amyloliquefaciens H</i>	$ \begin{array}{c} 5' \text{ ---G- } \downarrow \text{ -G-A-T-C-C--- } 3' \\ 3' \text{ ---C-C-T-A-G- } \uparrow \text{ -G--- } 5' \end{array} $	$ \begin{array}{cc} \boxed{5' \text{ ---G}} & \boxed{\text{G-A-T-C-C---} 3'} \\ \boxed{3' \text{ ---C-C-T-A-G}} & \boxed{\text{G---} 5'} \end{array} $	Sticky ends
EcoRI	<i>Escherichia coli</i> Ry13	$ \begin{array}{c} 5' \text{ ---G- } \downarrow \text{ -A-A-T-T-C--- } 3' \\ 3' \text{ ---C-T-T-A-A- } \uparrow \text{ -G--- } 5' \end{array} $	$ \begin{array}{cc} \boxed{5' \text{ ---G}} & \boxed{\text{A-A-T-T-C---} 3'} \\ \boxed{3' \text{ ---C-T-T-A-A}} & \boxed{\text{G---} 5'} \end{array} $	Sticky ends
Hind II	<i>H. influenzae</i> Rd	$ \begin{array}{c} 5' \text{ --G-T-C- } \downarrow \text{ -G-A-C-- } 3' \\ 3' \text{ --C-A-G- } \uparrow \text{ -C-T-G-- } 5' \end{array} $	$ \begin{array}{cc} \boxed{5' \text{ --G-T-C}} & \boxed{\text{G-A-C--} 3'} \\ \boxed{3' \text{ --C-A-G}} & \boxed{\text{C-T-G--} 5'} \end{array} $	Blunt ends

3'----- C T T A A G-----5'

5'----- G A A T T C-----3'

When one reads the sequence in opposite direction (3' to 5' or 5' to 3') it is identical/same.

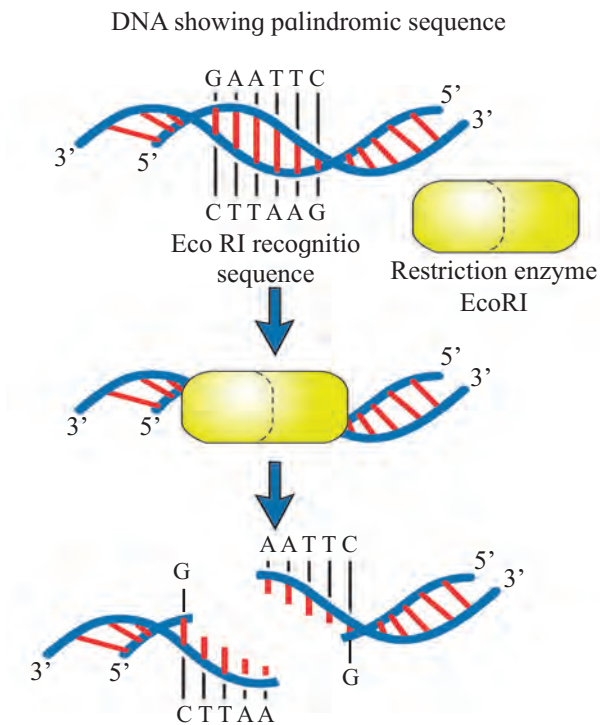


Fig. 12.3 : Mode of action of EcoRI to cleave DNA

A sequence with this type of symmetry is called a **palindrome**. When the enzyme EcoRI attacks this palindrome, it breaks each strand at the same site in the sequence, which is indicated by the arrow between the A and G residues.



Restriction enzymes either cut straight across the DNA in the region of palindrome to give blunt ends or cuts producing short, single stranded projections at each end of DNA to produce, cohesive or sticky ends or staggered ends.

B. Cloning vectors (vehicle DNA) - Vectors are DNA molecules that carry a foreign DNA segment and replicate inside the

host cell. Vectors may be plasmids, bacteriophages (M13, lambda virus), cosmids, phagemids, BAC (bacterial artificial chromosome), YAC (yeast artificial chromosome), transposons, baculoviruses and mammalian artificial chromosomes (MACs). Most commonly used vectors are plasmid vectors (pBR 322, pUC, Ti plasmid) and bacteriophages (lambda phage, M13 phage). Plasmids and bacteriophages are most commonly used as vectors.

Do you know ?

Smith, Nathan and Arber achieved the discovery of restriction enzymes. For this spectacular achievement, they were awarded Nobel Prize for physiology and medicine in 1978. Since then restriction enzymes have been used for genetic manipulation by dissecting, analyzing and re-configuration of the genetic information at the molecular level.

Do you know ?

There are three types of restriction enzyme viz,
 Type I - Which function simultaneously as endonuclease and methylase e.g. EcoK.
 Type II - Which have separate activities for cleaving and methylation; they are more stable and are used in rDNA technology e.g. EcoRI, BgII; these enzymes cut DNA at specific sites within the palindrome.
 Type III - Which cut DNA at specific non-palindromic sequences e.g. HpaI, MboII.
 Thousands of type II REs. are recognized.

A good vector should have the ability of independent replication so that as the vector replicates (through *ori* gene), large number of copies of the DNA insert will be formed. Moreover the vector should be able to easily get introduced into host cells.

A vector should have marker genes for antibiotic resistance; must contain unique cleavage sites for restriction enzymes; it should have at least suitable control elements like promoter, operator, ribosomal binding sites, etc. The plasmids obtained naturally do not possess all the characteristics. Hence, they are constructed by inserting a gene for antibiotic resistance. e.g. pBR 322, pBR 320, pACYC 177 are the constructed plasmids. pBR 322 is mostly used in rDNA technology in plants.

i. Plasmid : The plasmids most commonly used in recombinant DNA technology are those that replicate in *E. coli*. Investigators have engineered these plasmids to optimize their use as vectors in DNA cloning.

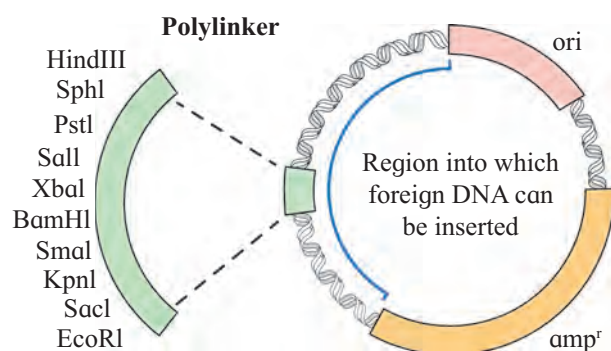


Fig. 12.4 : Plasmid cloning vector showing a replication origin (ori), a drug resistance gene (amp^r) and a region in which foreign DNA can be inserted.

ii. Plasmid vectors for plants : An important vector for carrying new DNA into many types of plants is a plasmid that is found in *Agrobacterium tumefaciens*. This bacterium lives in the soil and causes a plant disease called crown gall, which is characterized by the presence of over-growths, or tumors, in the plant. *A. tumefaciens* contains a plasmid called Ti (for tumor-inducing). The Ti plasmid contains a transposon, called T DNA, which inserts copies of itself into the chromosomes of infected plant cells. The

transposon, with the new DNA, can still be inserted into the host cell's chromosomes. A plant cell containing this DNA, can then be grown in culture or induced to form a new, transgenic plant.

C. Competent hosts (cloning organisms) used are usually the bacteria like *Bacillus haemophilus*, *Helicobacter pylori* and *E. coli*.

Mostly *E. coli* is used for the transformation with recombinant DNA

12.3 Methodology for rDNA technology :

The steps involved in gene cloning are as follows :

a. Isolation of DNA (gene) from the donor organism :

i. The desired gene to be cloned has to be obtained from the source organism (donor). Initially the cells of the donor organism are sheared with the blender and treated with suitable detergent. Genetic material from the donor is removed, isolated and purified by using several techniques. Isolated DNA can be spooled on to a glass rod.

ii. Isolated purified DNA is then cleaved by using restriction enzymes particularly Restriction Endonucleases (RE). These enzymes cleave DNA at specific sites, called **restriction sites** and break the DNA into fragments. There are several types of restriction endonucleases. Cleaved DNA fragments have cohesive, **sticky**, staggered ends or **blunt** ends.

From cleaved DNA fragments, a fragment containing desired gene is isolated and selected for cloning. This is now called **foreign DNA** or **passenger DNA**. A desired gene can also be obtained directly from genomic library or cDNA library.

? Do you know ?

Gene library is a collection of different DNA sequences from an organism where each sequence has been cloned into a vector for ease of purification, storage and analysis. There are two types of gene libraries on the basis of the source of DNA used.

- **Genomic library :** It is a collection of clones that represent the complete genome of an organism. The genomic library of prokaryotes can be constructed by using plasmid vector. It is because prokaryotic genome does not contain repetitive DNA.
- **cDNA library :** It represents the library of eukaryotic organisms only. DNA is produced from isolated mRNA by reverse transcription. The DNA so made is called complementary DNA (cDNA). The library is called cDNA library. Eukaryotic DNA genome contains introns, regulatory genes and repetitive DNA. Hence, the establishment of genomic library in eukaryotes is not meaningful.

b. Insertion of desired foreign gene into a cloning vector (vehicle DNA) :

The foreign DNA or passenger DNA is now inserted into a cloning vector or vehicle DNA. The most commonly used cloning vectors are plasmids of bacteria and the bacteriophage viruses like lambda phage and M13. The most commonly used plasmid is pBR 322.

Plasmids are isolated from the vector organisms i.e. bacterium. By using same RE (which is used in the isolation of the desired gene from the donor), plasmid i.e. vector DNA is cleaved.

Now by using enzyme DNA ligase, foreign DNA is inserted/ integrated into the vector DNA. The combination of vector DNA and foreign DNA is now called **Recombinant DNA** or **Chimeric DNA** and the technology is referred to as rDNA technology.

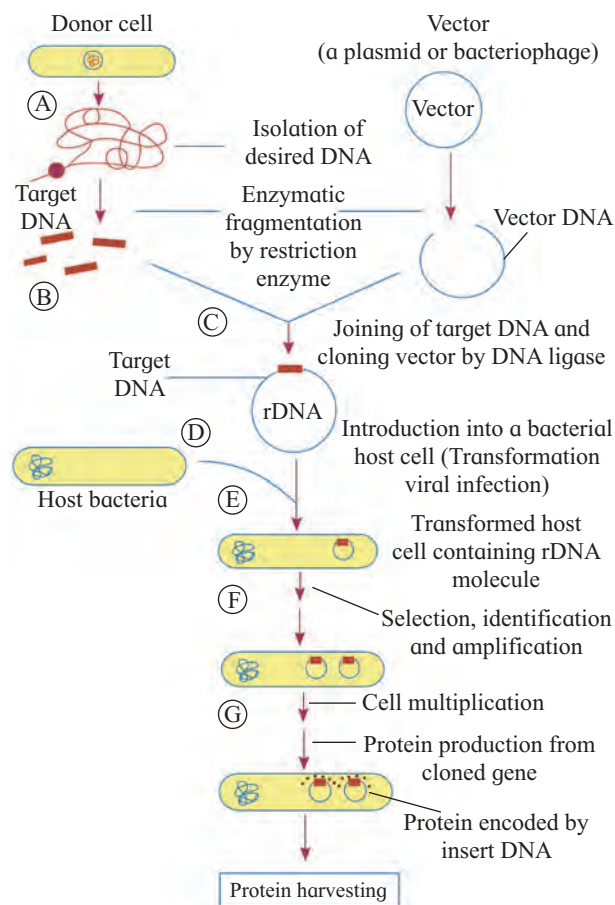


Fig. 12.5 : Outline of the process of recombinant DNA technology

c. Transfer of rDNA into suitable competent host or cloning organism :

Finally the recombinant DNA is now introduced i.e. transferred for expression into a competent host cell of the suitable cloning organism which is usually a bacterium. Host cell takes up naked rDNA by process of ‘transformation’ and incorporates into its own chromosomal DNA which finally expresses the trait controlled by passenger DNA. The transfer of rDNA into a bacterial cell is assisted by divalent Ca^{++} . The cloning organisms used in plant biotechnology are *E.coli* and *Agrobacterium tumifaciens*. The host/ competent cell which has taken up rDNA is now called **transformed cell**.

Foreign DNA can also be transferred directly into the naked cell or protoplast of the competent host cell, without using vector. This

is done by using techniques like electroporation, microinjection, lipofection, shot gun, ultrasonification, biolistic method, etc. But in plant biotechnology the transformation is through Ti plasmids of *A. tumifaciens*.

d. Selection of the transformed host cell :

The transformation process generates a mixed population of transformed (recombinant) and non-transformed (non-recombinant) host cells. For isolation of recombinant cell from non-recombinant cell, marker genes of plasmid vector is employed. For example, pBR322 plasmid vector contains different **marker genes** (Ampicillin resistant gene and Tetracycline resistant gene). When Pst1 RE is used, it knocks out Ampicillin resistant gene from the plasmid, so that the recombinant cell become sensitive to Ampicillin.

e. Multiplication of transformed host cell:

Once transformed, host cells are separated by the screening process. In this step the transformed host cells are introduced into fresh culture media.

At this stage the host cells divide and redivide along with the replication of the recombinant DNA carried by them.

f. Expression of the gene to obtain the desired product:

The next step involves the production of desired products like alcohol, enzymes, antibiotics, etc. Finally the desired product is separated and purified through downstream processing using suitable bioreactor.



Do you know ?

The Centre for Cellular and Molecular Biology (CCMB) located in Hyderabad, is a premier research organization in frontier areas of modern biology including DNA fingerprinting and Molecular approaches in animal breeding. The objectives of the Centre are to conduct high quality basic research and training in frontier areas of modern biology, and promote centralized national facilities for new and modern techniques in the interdisciplinary areas of biology.

12.4 Applications of Biotechnology:

Biotechnology is an umbrella term covering a broad spectrum of scientific applications used in many sectors, such as health and agriculture, industry, environment and genomics.

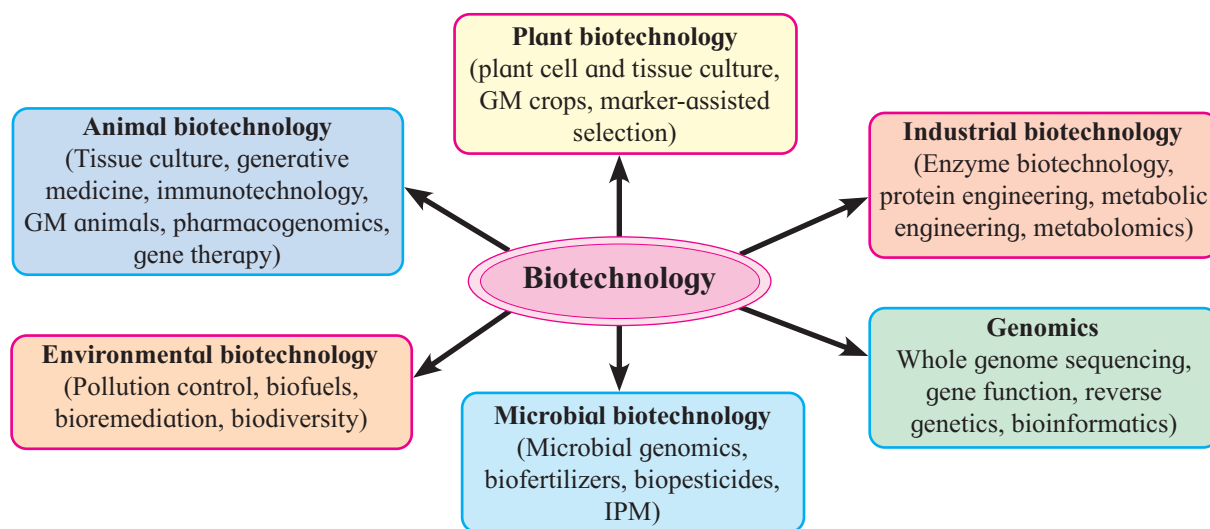


Fig. 12.6 : Applications of Biotechnology

Tabel 12.7 : Human proteins produced by rDNA technology to treat human diseases

Disorder/ Diseases/ Health condition	Recombinant protein(s)
1. Anaemia	Erythropoietin
2. Asthma	Interleukin-1 receptor
3. Atherosclerosis	Platelet derived growth factor
4. Parturition	Relaxin
5. Blood clots	Tissue Plasminogen Activator (TPA) Urokinase
6. Cancer	Interferons, tumour necrosis factor interleukins, macrophage activating factor
7. Diabetes	Insulin
8. Emphysema	α_1 - Antitrypsin
9. Haemophilia A	Factor VIII
10. Haemophilia B	Factor IX
11. Hepatitis B	Hepatitis B vaccine

a. Healthcare Biotechnology :

It refers to a medicinal or diagnostic product or a vaccine. This technology has a tremendous impact on meeting the needs of patients. Biotechnology offers patients a variety of new solutions such as: unique, targeted and personalized therapeutic and diagnostic solutions for organ transplant. Stem cell technology, genetic counselling, forensic medicine, gene probes, genetic fingerprinting and karyotyping are the outcomes of biotechnology.

Human insulin :

Insulin is a peptide hormone produced by β -cells of islets of Langerhans of pancreas. It was discovered by Sir Edward Sharpey Schafer (1916) while studying Islets of Langerhans.

Insulin is essential for the control of blood sugar levels. Diabetes mellitus is a disease in which some people cannot make insulin themselves. Hakura et al (1977), chemically

synthesized DNA sequence of insulin for two chains A and B and separately inserted into two pBR322 plasmid vector. Insulin production by recombinant DNA technology is designed by Gilbert and Villokomaroff in 1978.

The genes for chain A and B are inserted next to B galactoside gene separately in separate plasmid vectors. The recombinant plasmids were then separately transformed into *E. coli* host. The host produced penicillinase + pre-pro insulin. Insulin is later separated by trypsin treatment.

Vaccine production:

A vaccine is a biological preparation that provides active acquired immunity against a certain disease. Usually a vaccine consists of a biological agent that represents the disease-causing microorganism. It is often made from a weakened or killed form of the microorganism, its toxins or one of its surface protein antigens.

Biotechnology has offered modern diagnostic test kits - rickettsial, bacterial and viral vaccines along with radiolabelled biological therapeutics for imaging and analysis.

Vaccines have eliminated small pox, polio and other deadly diseases for the last several decades. Biotechnology has made advancements in vaccination by making recombinant vaccines that have the potential to eradicate non-communicable diseases like cancer. Naked DNA vaccines, viral vector vaccines and plant-derived vaccines are found to be most effective against a number of bacterial and viral disorders.



Internet my friend

Collect the information pertaining to recombinant vaccines and their types - protein vaccines and DNA vaccines.

Oral vaccines: a novel approach :

The latest hot spot in the field of vaccine research is the development of vaccine which can be taken orally. Immunogenic protein of

certain pathogens is found to be active when administered orally. The gene corresponding to such proteins is isolated and a gene construct is produced. This is introduced and expressed in a plant genome, which results in production of such immunogenic proteins in the parts of the plant where it is expressed. These when fed into animals or mainly humans, the person becomes vaccinated against certain pathogen. Such vaccines are also known as **edible vaccines**.

An exciting invention is production of '*melt in the mouth*' vaccines that can be administered by placing them under your tongue that delivers it into the blood stream. The most important example is the production of flu vaccine by *Bacillus subtilis* which melts in the mouth. The tremendous benefit of such vaccines, is the comfort of administration, low cost and ease of storage.

b. Agriculture :

Applications of biotechnology in agriculture involves the use of genetically modified crops through the gene manipulation. The GM crops that are pest resistant, stress tolerant etc. are used in improving productivity.

Tissue Culture is used in Micropropagation i.e. large-scale propagation of plants in very short durations. Tissue culture technique is also the best method for storing germplasm and maintaining a specific genetic type (Clone). This technique is used in those plants, which produce recalcitrant seeds or produce highly variable seeds.

Recalcitrant means the reduction in the seed moisture contents below certain levels and freezing drastically reduces the survival and thus present difficulty in storage. Here, subcellular damage of seeds occur accompanied by consequent loss of viability, when dried.

c. Gene therapy:

A gene is a stretch of DNA required to make a functional product such as part or all of a protein. During gene therapy, DNA that codes for specific genes is delivered to individual cells in the body.

Gene therapy is the treatment of disease (disorder) by replacing, altering or supplementing a gene that is absent or abnormal and whose absence or abnormality is responsible for the disease (disorder).

Most, if not all, disorder have a genetic factor. The genetic factor can be wholly or partially responsible for the disorder. For example, in disorders such as cystic fibrosis, haemophilia, and muscular dystrophy, changes in a gene directly result in the condition.

In other conditions, such as high cholesterol and high blood pressure, genetic and environmental factors interact to cause disease. There are more than 5000 different human genetic diseases known to be caused by single gene defects e.g. sickle cell anaemia, thalassaemia, Tay-sach's disease, cystic fibrosis, Huntington's chorea, haemophilia, alkaptonuria, albinism, etc.

Gene therapy is being used in many ways. For example, to:

- Replace missing or defective genes;
- Deliver genes that speed up the destruction of cancer cells;
- Supply genes that cause cancer cells to revert back to normal cells;
- Deliver bacterial or viral genes as a form of vaccination;
- Deliver DNA to antigen expression and generation of immune response;
- Supply of gene for impairing viral replication;
- Provide genes that promote or impede the growth of new tissue; and
- Deliver genes that stimulate the healing of damaged tissue.



Do you know ?

Delivery of genes into cells :

Genes can be delivered by three ways :

- i. *Ex vivo* delivery where cells are removed from the patients and then gene is introduced using viral or non-viral vectors e.g. **Parkinson's disease**, a neurological disorder.
- ii. *In vivo* delivery where therapeutic genes are directly delivered in the cells at the target sites of the diseased tissue in the patient- like intravenous infusion genes to treat cancer are injected directly into tumor.
- iii. Use of virosomes (Liposome + inactivated HIV), bionic chips are the other methods of gene delivery.

Forms of gene therapy :

a. Germ line gene therapy : In this method healthy genes can be introduced into germ cells like sperms, eggs, early embryos. It allows transmission of the modified genetic information to the next generation. Though it is highly effective in counteracting the genetic disorders, it is not encouraged for application in human beings due to a variety of technical and ethical reasons.

b. Somatic cell gene therapy : In this type the gene is introduced only in somatic cells like bone marrow cells, hepatic cells, fibroblasts endothelium and pulmonary epithelial cells, central nervous system, endocrine cells and smooth muscle cells of blood vessel walls. Modification of somatic cells only affects the person being treated and the modified chromosomes cannot be passed on the future generations. Somatic cell gene therapy is the only feasible option and the clinical trials have already employed for the treatment of acquired disorders such as cancer and rheumatoid arthritis and blood disorders including SCID, Gaucher's

disease, familial hypercholesterolemia, haemophilia, phenylketonuria, cystic fibrosis, sickle cell anaemia, Duchenne muscular dystrophy, emphysema, thalassemia etc.

d. Genetically Modified Organisms (GMOs):

These are living organisms whose genetic material has been artificially manipulated in a laboratory through genetic engineering. This creates combinations of plant, animal, bacteria, and virus genes that do not occur in nature or through traditional crossbreeding methods.

Most GMOs have been engineered to withstand the direct application of herbicide and/or to produce an insecticide. However, new technologies are now being used to artificially develop other traits in plants, such as a resistance to browning in apples, and to create new organisms using synthetic biology. Despite biotech industry promises, there is no evidence that any of the GMOs currently on the market, offer increased yield, drought tolerance, enhanced nutrition, or any other consumer benefit.

I. Transgenic Plants :

The human race is very dependent on agriculture and as world populations continue to expand, there must be continuous reassessment of agricultural practices to optimize their efficiency. Since early times human beings have sought to improve the quality and productivity of agriculturally important plants. This was done by selection and traditional breeding procedures that were painstakingly slow and difficult. Traditional breeding programmes involve sexual crosses, which resulted in the high quality of present day food plants such as wheat, rice, corn, potato, etc. More recently, biotechnological approaches have been applied to these plants to create genetic variations that are beneficial for mankind.

First transgenic plant produced was tobacco. More than 60 transgenic dicot plants and several monocot plant like maize, oat,

rice, wheat are known. Tomato, soybean, potato, sugar beet, grapes, brinjal, cotton are other transgenic plants. Transgenic plants are being looked up as bioreactors for molecular farming i.e. for production of novel drugs like interferons, edible vaccines, antibodies, amino acids, immunotherapeutic drugs, etc.

Advantages of GM food-plants :

The ways in which one thinks that genetically modified plants can help, are listed as follows:

a. Insect pest resistance:

It can help farmers to reduce their use of chemical pesticides, which in turn can reduce the cost of producing food. However, an alternative has been available for more than 30 years which is a biological insecticide from the bacterium, *Bacillus thuringiensis* (Bt). However, the use of *B. thuringiensis* sprays is limited because of low stability of the protein in the field.

Bt cotton is one of the best transgenic plants known for its insect resistance property.

Insect resistant plants contain either a gene from *B. thuringiensis* or the cowpea trypsin inhibitor gene. The gene called cry gene present in *B. thuringiensis* produces a protein that forms crystalline inclusions in bacterial spores. When ingested by a susceptible insect, a combination of high pH and the enzyme proteinase of the insect's midgut, processes them hydrolytically to release the core toxic fragments.

The effect of these fragments is seen within minutes of ingestion, beginning with midgut paralysis and ending with disruption of midgut cells of insect. *Bt* toxin activity has been against many species of insects within the orders of Lepidoptera, Diptera, and Coleoptera.

Golden rice - a transgenic food crop used to reduce vitamin A deficiency disease.

Similarly, the gene of α -amylase inhibitor (α AI-Pv) has been isolated from adzuki bean (*Phaseolus vulgaris*) and transferred to tobacco and this gene works against pests like *Zabrotus subfasciatus* and *Callosobruchus chinensis*.

b. Improved nutritional qualities (biofortification):

Transgenic plants have also been produced to provide functional food and nutraceuticals. For millions of people in developing countries, rice is the main item in their diet. Because rice does not contain many essential nutrients, malnutrition is very common in these countries.

Especially terrible is the night blindness that results from a lack of vitamin A. This vitamin is abundant in milk and in vegetables such as carrots, which most of the poor people of the world do not easily afford. To solve this problem, Swiss researchers created transgenic rice (golden rice) and transgenic mustard (golden mustard) varieties that are high in vitamin A. The golden colour is due to beta carotene, precursor of vitamin A. They hoped that this rice, if grown and eaten in developing countries, would reduce the diseases associated with vitamin A deficiency (VAD).

Table 12.8 : Some transgenic plants produced for functional food and nutraceuticals

Substance	Potential benefit	Crop	Transgene
Provitamin A	Anti-oxidant	Rice	Phytoene synthase, lycopene cyclase
Vitamin E	Anti-oxidant	Canola	γ - tocopherol methyl transferase
Flavonoids	Anti-oxidants	Tomato	Chalone isomerase
Fructants	Low calories	Sugarbeet	1-sucroese: sucrose fructosyl transferase
Iron	Iron fortification	Rice	Ferritin, metallothioein, phytase

Improvement in oil content and oil quality of oil crops like soybean, oil palm, rapeseed and sunflower, have been achieved by transfer of 'Arabidopsis genes'.

Iron deficiency is also a serious nutritional problem, affecting an estimated 30% of the world population. For production of transgenic crops that will produce food rich in iron, an iron storage protein (ferritin) is targeted. Ferritin is found in many animals, plants and bacteria. The genes for ferritin protein isolated from soybean and *Phaseolus* have been transferred to rice.



Do you know ?

- Genetically engineered **herbicide tolerant** plants are developed as in maize, wheat and many other monocot plants.
- Genetically engineered **disease resistant plants** (against bacterial and viral pathogens) are also developed in crop plants like tomato, potato and tobacco.
- Plants deficient in amino acids like methionine, lysine and tryptophan are genetically engineered by introducing genes from other sources so as to make the seeds protein rich. e.g. Leguminous plants (pulses), maize, etc.
- Similarly, genetically engineered plants tolerant to abiotic stresses such as high temperature, water, cold, etc. are also developed.

c. Modification in Post-harvest characteristics:

Diseases and pests, bruising on soft fruits and vegetables, heat and cold storage, over-ripeness, loss of flavours and odours, etc. lead to great deal of losses during storage and transport of crops.

Most of these physiological changes are due to endogenous enzyme activity. Genetic engineering has made it possible to slow down these activities. In the tomato the enzyme polygalacturonase breaks down the cell wall constituent- pectin, leading to softening of fruit during ripening. Thus, the fruits are easily bruised and damaged on shipment. By inhibiting the polygalacturonase by antisense genes, the tomato (genetically modified tomatoes are called "Flavr savr" tomatoes) can remain on the vine until mature long shelf life and be transported in a firm solid state.

d. Plants as factories :

To produce novel biochemicals and vaccines (**Biopharmaceuticals**), plants are potential factories or bioreactors for high value biochemicals like starch, sugar, lipids, proteins, and products like fine chemicals, perfumes and adhesive compounds as well as industrial lubricants, biodegradable plastic and even 'renewable' energy crops to replace fossil fuels.

Biopharmaceuticals are proteins, hormones, antibodies, vaccines or enzymes isolated from transgenic plants. Some of the proteins that are being produced by transgenic crop plants:

- Human growth hormone with the gene inserted into the chloroplast DNA of tobacco plants.
- Humanized antibodies against such infectious agents as HIV, Respiratory syncytial virus (RSV), Herpes simplex virus (HSV), the cause of "cold sores"
- Protein antigens to be used in vaccines for e.g. Patient-specific antilymphoma (a cancer) vaccines. B-cell lymphomas are clones of malignant B cells expressing on their surface a unique antibody molecule.

Currently many novel products have been commercially exploited for their products such as:

- a. A ‘superglue’ produced by tobacco plants with genes encoding for powerful adhesive proteins, enables marine mussels to stick to rocks. It will be especially valuable as a biochemical glue for body repairs during surgery.
- b. Transgenic plants, containing oil-encoding gene from marine algae, produce oil that has nutritional value similar to cod- liver oil.
- c. Plant that will produce the antimalarial drug, Artemisinin.
- d. Product of Genetically engineered opium (poppy) used to produce more powerful painkillers. i.e. analgesics and sedatives.

e. Transgenic plants producing edible vaccines:

Genetically altered plants can provide protection to infectious diseases. Plant products acting as vaccines would be inexpensive to produce and thus, can easily be made available in developing countries. GM Potatoes, tomatoes, bananas, soybeans, alfalfa and cereals are the most common foods proposed for edible vaccine delivery. Fruits of GM bananas can be used as oral edible vaccine against hepatitis B.

II. Transgenic animals:

Many transgenic animals such as mice, rats, rabbits, pig, sheep, cows, fowls, fish have been

produced through rDNA technology. The term transgenic animal refers to an animal in which there has been a deliberate modification of the genome - the material responsible for inherited characteristics - in contrast to spontaneous mutation. Foreign DNA is introduced into the animal, using recombinant DNA technology, and then must be transmitted through the germ line so that every cell, including germ cells, of the animal contain the same modified genetic material.

A representative, but non-inclusive, list of purposes for which transgenic animals have been used, indicates the wide-ranging application of this biotechnology:

- in medical research, transgenic animals are used to identify the functions of specific factors in complex homeostatic systems through over- or under-expression of a modified gene (the inserted transgene);
- in toxicology: as responsive test animals (detection of toxicants);
- in mammalian developmental genetics;
- in molecular biology, the analysis of the regulation of gene expression makes use of the evaluation of a specific genetic change at the level of the whole animal;
- in the pharmaceutical industry, targeted production of pharmaceutical proteins, drug production and product efficacy testing;

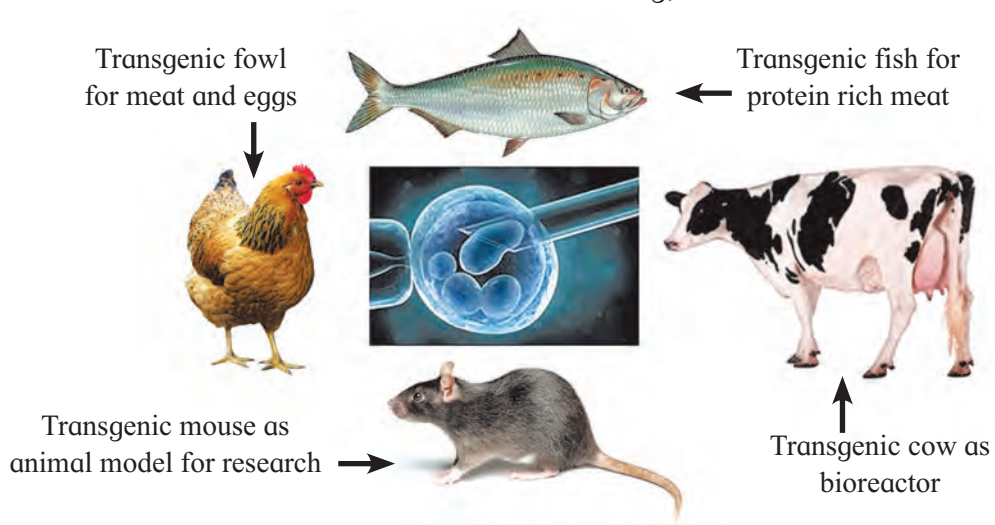


Fig. 12.9 : Transgenic animals

- in biotechnology: as producers of specific proteins;
- genetically engineered hormones to increase milk yield, meat production; genetic engineering of livestock and in aquaculture affecting modification of animal physiology and/or anatomy; cloning procedures to reproduce specific blood lines; and
- developing animals specially created for use in xenografting.

a. Transgenic mice and cancer research :

Through laboratory investigations with transgenic mice that have been modified using a particular oncogene (cancer causing gene) and thus developed a certain type of cancer, questions concerning the relationship between oncogenes and cancer development could be answered. Theoretically, such animals can also be used for research into cancer treatment and prevention of malignancy.

In the laboratory of Philip Leder in Harvard (USA) the transgenic mouse model for the investigation of the breast cancer was developed. The oncogenes *myc* and *ras* were analysed to find out if they lead to breast cancer in mice transformed with these genes.

b. Transgenic farm animals :

With the advent of technology, the intention of researchers is diverted to produce transgenic farm animals from which the mankind can derive greater benefits. Many of the farm animals are improved for their meat production ability while some of them are improved for milk yields and quality, and disease-free status. At the beginning of the century, a dairy cow provided 2,000 to 3,000 liters of milk a year.

Today, Holstein cow provides 6,000 liters on average and up to 8,000 – 10,000 for the best ones. A century ago, a hen laid about 70 eggs a year whereas today the best races lay up to 250 eggs per year. This could be possible because of the advent of biotechnology.

The main objectives for improved animal breeding programmes coupled with this new technology of gene transfer are given below :

- Efficiency of meat production
- Improved quality of meat
- Milk quality and quantity
- Egg production
- Wool quality and quantity
- Disease resistance in animals
- Production of low-cost pharmaceuticals and biologicals

c. Transgenic cattle for food production :

Of the few research reports describing the use of transgenic technologies in cattle only one is directed towards a food production application. Researchers introduced additional copies of bovine beta or kappa casein into dairy cattle and evaluated the effect on milk production and composition. Transgenic offspring had an 8 to 20% increase in beta casein and a two-fold increase in kappa casein.

d. Transgenic cattle for human therapeutic production :

A second application for genetically modified cattle is the production of human therapeutic proteins. Human proteins that have been expressed in milk include human lactoferrin, human alpha lactalbumin, human serum albumin and human bile salt stimulated lipase. The mammary gland in dairy cows is an excellent protein production factory. On the other hand, one transgenic cow would be more than sufficient for production of annual world supply of factor IX (plasma thromboplastin component) that is used in the treatment of haemophilia.

In 1990 Tracy, the transgenic cow was born in Scotland, and could produce a human protein in her milk for human therapeutics.

Antibodies are currently used for many different human clinical applications; including treatment of infectious disease, cancer,

transplanted organ rejection, autoimmune diseases and for use as antitoxins. To make a human antibody product, the genetically modified cows are immunized with a vaccine containing the disease agent.

e. Transgenic Sheep :

Gene transfer technology is applied to sheep to produce transgenic sheep which are able to achieve better growth and meat production as well as to serve as bioreactors. Human growth hormone gene is introduced in sheep for promoting the growth and meat production.

Bacterial genes, *cys E* and *cys M*, are concerned with biosynthesis of cysteine amino acids involved in formation of keratin protein found in wool. Both these genes are identified, cloned and introduced in sheep to increase wool production and to improve the quality of wool.

f. Transgenic pigs :

The objective of gene transfer in pigs is to increase growth and meat production and to act as bioreactors.

Pigs are regarded as the most suitable animals to be bred for heart transplant because a pig's heart is about the same size as a human heart, and pig heart valves have been used in human heart surgery for over a decade. The pig clone is the first step towards providing animal organs and tissues for human transplants (xenotransplantation).

g. Transgenic fish :

The commercially important fish like Atlantic salmon, catfish, goldfish, *Tilapia*, zebra-fish, common carp, rainbow trout, etc. are transfected with growth hormone, chicken crystalline protein and *E.coli* hygromycin resistance gene. Transgenic fish showed increased cold tolerance and improved growth and it is the quantity and quality of fish proteins as well as its preservation, are the factors affecting the economic value of fish.

h. Transgenic chicken :

Also carry and express foreign genes. They could be used to improve the genetic make-up of existing strains with respect to built-in (*in vivo*) resistance to viral and coccidial diseases, better feed efficiency, lower fat and cholesterol levels and high protein containing eggs, and better meat quality.

12.5 Bioethics, bio-piracy and bio-patent :

Bioethics:

Ethics usually deals with the matters related to socially acceptable moral duty, conduct and judgement. In other words, it helps to regulate the behaviour of community by some set of standards. However the concepts differ according to culture and traditions. Moreover concepts change with the time due to shifting of perception of values which are affected by progress in science and technology. Bioethics helps to study moral vision, decision and policies of human behaviour in relation to biological phenomena or events. Ethics deals with 'Life' e.g. *in vitro* fertilization, sperm bank, gene therapy, cloning, gene manipulations, euthanasia, death, maintaining those who are in comatose state, prenatal genetic selection, etc.

The era of biotechnology has brought wide spectrum on new topics like cloning, transgenic, gene therapy, eugenics, rDNA technology, etc. The use of all these has drawn a wide range of reactions in the society. The reactions are based on individual's own perception and moral. Ethical aspects pertaining to the use of biotechnology seems to be more controversial and frightening. These concerns are broadly summarized below :

Use of animals causes great sufferings to them; violation of integrity of species caused due to transgenesis; transfer of human genes into animals and vice versa; indiscriminate use of biotechnology pose risk to the environment, health and biodiversity.

The introduction of Genetically Modified Organisms (GMOs) has led to a wider debate on bioethical concerns affecting social, economic and environmental spheres. These include the effects on non-target organisms, insect resistance crops, gene flow and the loss of diversity as well as the issue on interfering with nature in which the modification process itself is disrupting the natural process of biological entities. Ethics in biotechnology also includes the general subject of what should and should not be done in using recombinant DNA techniques

12.6 Effects of Biotechnology on the Environment :

a. Herbicide Use and Resistance :

Effects on the environment are of a particular concern with regard to GMO crops and food production. One area of development involves adding the ability to produce pesticides and resistance to specific herbicides. These traits are helpful in food production, allowing farmers to use fewer chemicals, and to grow crops in lesser ideal conditions. However, herbicide use could be increased, which will have a larger negative effect on the surrounding environment. Also unintended hybrid strains of weeds and other plants can develop resistance to these herbicides through cross-pollination, thus negating the potential benefit of the herbicide. One such herbicide that has already been added is RoundUp. Crops of RoundUp-ready soybeans have already been implemented into agricultural practices, possibly conferring RoundUp resistance to neighboring plants.

b. Effects on Untargeted Species :

Bt corn, which produces its own pesticide, is also in use today. It has adverse effects on Monarch butterfly populations, which are not the original target of the pesticide. It can also have unintentional effects on neutral or even beneficial species.

12.7 Effects of Biotechnology on Human Health :

a. Allergies :

GMO crops could potentially have negative effects on human health as well. Consumers have developed unexpected allergic reactions. e.g. Researchers used a gene from the Brazil nut to increase the production of Methionine in soybeans. The insertion of this gene inadvertently caused allergic reactions to the soybean in those with known nut allergies (“Biotech Soybeans”).

b. Long-Term Effects :

Because GMO technology has been available for such a short amount of time, there is relatively little research which has been conducted on the long-term effects on health which we cannot anticipate at this point.

c. New Proteins :

Proteins that have never been ingested before by humans are now part of the foods that people consume every day. Their potential effects on the human body are as of yet unknown. e.g. Whey proteins.

d. Food Additives :

GMOs also present us with possibilities of introducing additional nutrients into foods, as well as antibiotics and vaccines. This availability of technology can provide nutrition and disease resistance to the people of those countries that don't have the means to provide these, otherwise.

However, there is possibility of the creation of antibiotic and vaccine-resistant strains of pathogens.

This shows that the vast advances in life sciences and our multicultural and pluralistic modern societies create numerous bioethical problems requiring some stringent regulation. In terms of GMOs, the Indian Government has set up the **Genetic Engineering Approval Committee (GEAC)**. This organization makes decisions regarding the validity of research involving GMOs and addresses the safety of GMOs introduced for public use.

12.8 Biopatent and Biopiracy :

a. Biopatent :

Patent is a special right granted to the inventor by the government. Patent is a personal property of inventor. It can be sold like any other property. A patent consists of three parts - **grant** (agreement with the inventor), **specification** (subject matter of invention) and **claims** (scope of invention to be protected).

Biopatent is a biological patent. Biopatents are awarded for strains of microorganisms, cell lines, genetically modified strains, DNA sequences, biotechnological processes, product processes, product and product applications.

Biopatents are awarded to recognize real innovative contributions made by the inventor to the cause of human welfare. The awards are given to inculcate encouragement and values in developing scientific culture and in emphasizing the role of biology in shaping human society.

Indian patent allows 'process patent' and not the 'product patent'. Biopatent allows the patent holder to exclude others from making, using, selling or importing protected invention for a limited period of time. Duration of biopatents is five years from the date of the grant or seven years from the date of filing the patent application, whichever is less.

First biopatent was patented pertaining to genetically engineered bacterium '*Pseudomonas*' used for clearing oil spills. Patent under the title 'control of plant gene expression' was issued jointly to Delta and Pineland company and U. S. department of agriculture. Patent is based on a gene that produces a protein toxic to plant thus, do not allow seeds to germinate. However, this patent was not granted by Indian government. Such a patent is considered morally unacceptable and fundamentally unequitable. This is because financially powerful corporations would acquire monopoly over biotechnological process. This in turn would pose a threat to global food security.

b. Biopiracy :

Pirates in general terms are those who steal and kill others to enrich themselves. Biopirates are those who do not kill but steal the patent (misuse the patent). Biopiracy is defined as 'theft of various natural products and then selling them by getting patent without giving any benefits or compensation back to the host country'. In short, it is unauthorized misappropriation of any biological resource and indigenous knowledge.

The developed, industrialized and financially rich nations are poor in biodiversity and traditional knowledge whereas developing and underdeveloped nations have ample of biodiversity and traditionally, they know better the use of their bio-resources. Traditional knowledge naturally includes a deep understanding of ecological processes and the ability to sustainably extract useful products from the local habitat. Traditional knowledge is handed over through the generations. This helps them to develop modern, commercial applications that save the makers time, money and effort.

Components of **Traditional Knowledge** that are especially relevant to our global survival include knowledge of:

- Food, crop varieties and agricultural/farming practice
- Sustainable management of natural resources and conservation of biological diversity
- Biologically important medicines

The conservation of species, habitat, and biodiversity are essential to the continued survival of tribal people. By conserving the customs and habitat of tribal people, we concurrently reduce emissions from deforestation and ecosystem degradation. Furthermore, the opportunity for cultural survival is a basic human right. The traditional knowledge is facing a problem of biopiracy.

The act of Piracy is unauthorized publication or reproduction of another person's work or material. When someone indulges in piracy, the accused is using someone's work illegally or without taking any permission. The innovations and discovery of the pharmaceutical and agricultural researches are not new as to qualify as invention as they are based on centuries of knowledge of the traditional societies.

Examples of Biopiracy :

• Patenting of Neem (*Azadirachta indica*) :

The people of India in a variety of ways have used neem, since time immemorial. Indians have shared the knowledge of the properties of the neem with the entire world. Pirating this knowledge, the USDA and an American MNC W.R. Grace in the early 90s sought a patent from the European Patent Office (EPO) on the "method for controlling on plants by the aid of hydrophobic extracted neem oil." The patenting of the fungicidal properties of Neem, was an example of biopiracy.

• Patenting of Basmati :

Basmati is a long-grained, aromatic variety of rice indigenous to the Indian subcontinent.

In 1997 the US Patent and Trademark Office (USPTO) granted a patent to a Texas based American company Rice Tec Inc for "Basmati rice line and grains" having trade

name **Texmati**. The patent application was based on 20 very broad claims on having "invented" the said rice. Due to peoples movement against Rice Tec in March 2001, the USPTO has rejected all the claims.

• Haldi (Turmeric) Biopiracy :

Two American researchers of Indian origin of the University of Mississippi Medical Center, put a claim to the US Patent and Trademark Office, maintaining that they had discovered *haldi's* healing properties. Surprisingly, they were granted a patent in March 1995 for something you had known for years and our ayurvedas for centuries.

It meant they had exclusive rights over any such *haldi* drug and were in a position to make millions of dollars. The Council of Scientific and Industrial Research (CSIR) applied to the US Patent Office for a re-examination and they realized the mistake and cancelled the patent. This was after Indian scientists shouted from rooftops about how we are losing our traditional knowledge to marauding foreign companies who have started poaching on our ancient healing techniques.

It is the need of hour to launch genetic literacy movement in Indian school and colleges for better understanding of opportunities and risks related to biotechnology and also to promote the safe and meaningful use of technologies of modern life sciences.



Activity :

Collect information on the use of Biotechnology in pollution control.

Exercise

Q. 1 Choose the correct option

- The bacterium which causes a plant disease called crown gall is
 - Helicobacter pylori*
 - Agrobacterium tumifaciens*
 - Thermophilus aquaticus*
 - Bacillus thuringiensis*
- The enzyme nuclease hydrolyses of polynucleotide chain of DNA.
 - hydrogen bonds
 - phosphodiester bonds
 - glycosidic bonds
 - peptide bonds
- In vitro amplification of DNA or RNA segment is known as
 - chromatography
 - southern blotting
 - polymerase chain reaction
 - gel electrophoresis
- Which of the following is the correct recognition sequence of restriction enzyme *Hin d* III.
 - 5' ---A-A-G-C-T-T---3'
3' ---T-T-C-G-A-A---5'
 - 5' ---G-A-A-T-T-C---3'
3' ---C-T-T-A-A-G---5'
 - 5' ---C-G-A-T-T-C---3'
3' ---G-C-T-A-A-G---5'
 - 5' ---G-G-C-C---3'
3' ---C-C-G-G---5'
- Recombinant protein is used to dissolve blood clots present in the body.
 - insulin
 - tissue plasminogen activator
 - relaxin
 - erythropoietin

- Recognition sequence of restriction enzymes are generally nucleotide long.
 - 2 to 4
 - 4 to 8
 - 8 to 10
 - 14 to 18

Q. 2 Very short answer type questions.

- Name the vector which is used in production of human insulin through recombinant DNA technology.
- Which cells from Langerhans of pancreas do produce a peptide hormone insulin?
- Give the role of Ca^{++} ions in the transfer of recombinant vector into bacterial host cell.
- Expand the following acronyms which are used in the field of biotechnology.
 - YAC
 - RE
 - dNTP
 - PCR
 - GMO
 - MAC
- Fill in the blanks and complete the chart.

GMO	Purpose
i. Bt cotton
ii.	Delay the softening of tomato during ripening.
iii. Golden rice
iv. Holstein cow

Q. 3 Short answer type questions.

- Explain the properties of a good and ideal cloning vector for rDNA technology.
- In a PCR machine the temperature can be raised upto 100 °C but after that it is not able to lower the temperature below 70 °C automatically. Which step of PCR will be hampered first in this faulty machine? Explain why?
- In the process of rDNA technology, if two separate restriction enzymes are used to

cut vector and donor DNA then which problem will arise in the formation of rDNA or chimeric DNA? Explain.

4. Match and write the pairs.

Recombinant protein	It's use in or for
i. platelet derived growth factor	a. Anemia
ii. α -antitrypsin	b. cystic fobrosis
iii. Relaxin	c. Haemophilia A
iv. Erythropoietin	d. Diabetes
v. Factor VIII	e. Emphysema
vi. DNA ase	f. Parturition
	g. Atherosclerosis

Q. 4 Long answer type questions.

- Define and explain terms.
 - Biopiracy
 - Biopatent
 - Bioethus
- Explain the steps in process of rDNA technology with suitable diagrams.
- Explain the gene therapy. Give two methods of the same.
- How are the transgenic mice used in cancer research?
- Give the steps in PCR or polymerase chain reaction with suitable diagrams.
- What is a vaccine? Give advantages of oral vaccines or edible vaccines.
- Enlist different types of restriction enzymes commonly used in rDNA technology? Write on their role.
- Enlist and write in breif about the different biological tools required in rDNA technology.

Project :

Visit the tissue culture laboratory in your area. Prepare powerpoint presentation on tissue culture methodology and its applications.



Can you recall?

1. What is an ecosystem?
2. What are the abiotic and biotic components of an ecosystem?

Natural world around us shows amazing diversity of forms and complexity of relations. To understand these, we have to study levels of organization in the living world viz. macromolecules, cells, tissues, organs, individual organism, population, communities, ecosystems and biomes.

You have already studied in school, that ecology is a study of the interactions among organisms and between the organisms and their physical (abiotic) environment. Term ecology was first used by Reiter but E. Haeckel gave substance to the term (introduced) ecology.

Ecological grouping of organisms is nothing but ecological hierarchy. There are four sequential levels with increasing complexity of ecological (biological) organizations viz, Organism, Populations, Communities and Biomes. Individual **organism** is the basic unit of ecological hierarchy. Organisms of same kind inhabiting a geographical area constitute **population**. Several populations of different species in a particular area constitute **community** that interact with one another in several ways. **Biome** constitutes a large regional terrestrial unit delimited by a specific climatic zone having major vegetation zone (plant communities) and the associated fauna. There are six major groups of terrestrial biomes. We shall explore first two levels viz, organism and populations.

13.1 Organisms and the environment around:

Ecology at the level of organism is basically the study of animal or plant physiology which helps us to understand how the organisms

are adapted to their environments, not only for their survival but also for propagation (multiplication).

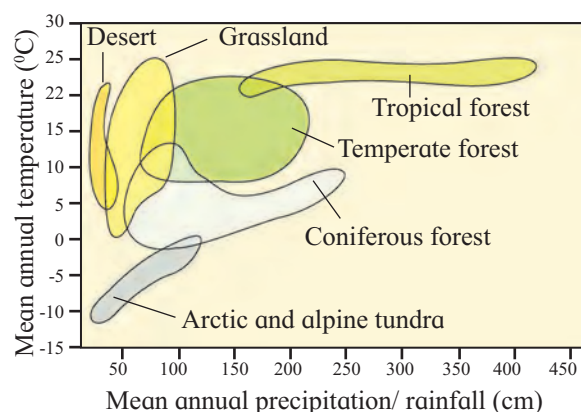


Fig. 13.1 : Distribution of major Biomes with respect to annual temperature and precipitation

You have studied in earlier classes about the rotation of earth around the Sun and the tilt of its axis, cause seasons. These seasons with annual variation in precipitation in the form of rain and snow, gives rise to formation of major biomes of the earth like desert, rain forest, grassland, tundra, etc.

Regional and local variations within each biome lead to the formation of a variety of habitats. Major biomes of earth are shown in Fig. 13.1. On the Earth, life exists even in extreme and harsh habitats like scorching deserts of Rajasthan, perpetually rain-soaked forests of North Eastern states and high mountain tops of Himalayas.



Find out

What are the key differences that make such a great variation in the physical and chemical conditions of different habitats?

Here, we must remember that it is not only the physico-chemical (abiotic) components that make up the habitat of an organism, but the habitat also includes biotic components like

plants, pathogens, parasites, and predators of the organism. We assume that over a period of time, the organism had through natural selection, evolved adaptations to optimize its survival and reproduction in its habitat.



Do you know ?

Ethology - The term was coined by Hilaire (1854) but was popularised by W. M. Wheeler (1902). The term denotes (speaks for) the study of behaviour of animals in relation to their environment.

Ecology - The term was introduced by E. Haeckel (1865) for the relationship of animals and plants with their surroundings.

Bionomics - Lankester (1890) coined this term for the study of relation between organisms to their environment.

Environmental biology (modern ecology) - The term was introduced in 20th century (G. L. Clarke 1964, Odum 1969) giving emphasis on the functional or physiological interrelationships between the organism and their surroundings.

Biosphere - All the ecosystems on earth constitute biosphere.

Habitat and Niche :

Habitat is a place or the set of environmental conditions around the organism to which it must adapt to survive and prosper. The term **Niche** is used to denote the functional role played by an organism in its environment (J. Grinnell 1917). Niche includes various aspects of the life of an organism like diet, shelter, etc.

A habitat defines the physical space of an organism with the other living or non-living factors, while niche describes how that organism is linked with its physical and biological environment. In colloquial language habitat is a postal address while niche is the profession of organism.

Table 13.2 : The differences between Habitat and Niche can be summarized as :

Habitat	Niche
A habitat is an area, where a species lives and interact with the other factors and prosper.	A niche is a concept, of how an organism lives or survives in the environmental conditions.
Habitat consists of numerous niches.	Niches do not contain such components.
Habitat has effect of temperature, rainfall and other abiotic factors on its population.	Niche shows impact of organisms upon the habitat.
Habitat supports numerous species at a time.	Niche supports a single species at a time.
Habitat is a physical place.	Niche is an activity or role performed by organisms.
Habitat is not species specific.	Niche is species specific.

Definition of Habitat :

Place or area where a particular species lives, is its habitat. Factors like the sunlight, average rainfall, annual temperatures, type of soil present and other abiotic (topographic) factors, affect the presence of organisms. These factors help in determining the presence of the particular type of species in the environment.

Pond, river, ocean, etc. are the examples of habitat as many organisms are found in the same place or habitat. These habitats can be arboreal, terrestrial, aerial, aquatic, etc. The immediate surrounding of an organism, sometimes also referred to as **microhabitat**, is an important concept to remember when working with sedentary or weakly motile organisms.

Definition of Niche :

The term niche was first time used by 'J. Grinnell'. The term ecological niche is still not well understood and is sometimes even misused.

Niche is described as a position of a species in the environment like, what they do for their survival? how they fulfill their needs of shelter, food? etc. Niche deals with the flow of energy from one organism to another and hence, it is important to understand, what an organism eats, how it interacts with other organisms, etc. As soon as the niche is left vacant, other organisms fill that position. The niche is specific to each species, which means no two species can share the same niche.

If the species creates its own unique niche in an ecosystem, it would be helpful in reducing competition for resources among species. By taking an example of a bird, it can be understood that how these birds differ in their eating habits, where some birds eat only insects, some only fruits and some can eat both and anything they come across. So here we can conclude that these birds living in the same habitat differ in their niches because of different eating habits.

Three types of niches are found:

a. Spatial or habitat niche : It deals with the physical space occupied by the organisms.

b. Trophic niche : It is on the basis of trophic level of an organism in a food chain.

c. Multidimensional or hypervolume niche: It considers number of environmental factors (both biotic and abiotic), the resulting space will be a hypervolume; not something that can be perceived by the human mind. This space is called the **hypervolume niche**. Alternatively, it is the position of an organism in the environmental gradient.

For every species, there is a fundamental niche and a realized niche. **Fundamental niche** is the niche in the absence of all competitors, this is highly improbable in nature. Hence, **realized niche** is more realistic approach, in the presence of competition for the resources available in the habitat.

We have seen earlier in the chapter that each habitat type is regulated by a number of abiotic or physico- chemical factors.

Key abiotic factors that influence any habitat are ambient temperature, availability of water, light and type of soil.

13.2 Major Abiotic Factors :

Temperature:

It is the most ecologically relevant environmental factor. Average temperature on land varies from subzero levels in polar areas and high altitudes, upwards upto 50°C in tropical deserts in summer. Temperature also varies seasonally. It decreases progressively from the equator towards the poles and from plains to the mountain tops. There are some unique habitats such as hot springs (80 to 100°C) and deep-sea hydrothermal vents where average temperatures usually 400° C. Ambient temperature affects the enzyme kinetics of the cell and thus, the entire metabolism, activity and other physiology of the organism.

Only few organisms can tolerate and thrive in a wide range of temperatures (eurythermal), but, a vast majority of them are restricted to a narrow range of temperatures (stenothermal).



Find out

1. Give names of eurythermal and stenothermal animals and plants?
2. What will be the effect of increasing global temperatures on the different habitats and the organisms found in those habitats?

Water: Availability of water is an important factor affecting the organisms. As we know, life on earth originated in water, its availability is so limited in deserts that only special adaptations are required to survive there. The productivity and distribution of plants are also heavily dependent on water.

Organisms living in water bodies such as oceans, lakes and rivers, have their own water-related problems. For aquatic organisms the chemical composition and pH of water are important.

The dissolved salt concentration (measured as salinity in parts per thousand), is less than 5ppt in fresh waters of streams, lakes and rivers, and 30-35ppt in the seas and oceans. It may go up to 100ppt in some hypersaline lagoons.

Some organisms are tolerant for a wide range of salinities (euryhaline) but others are restricted to a narrow range (stenohaline). Many fresh water animals cannot live for long in sea water and vice versa because of the osmotic problems, they would face.



Find out

Give examples of an animal and plant that can survive in fresh water as well as sea water?

Light: Plants use light for photosynthesis, which is only source of energy for the entire ecosystem. Photosynthesis can occur only in presence of sunlight. Many species of small plants (herbs and shrubs) growing on forest floor are adapted to perform photosynthesis optimally under very low light conditions because they are constantly overshadowed by tall trees.

For animals too, diurnal and seasonal variations in light intensity and duration (photoperiod) are clues for timing their foraging, reproductive and migratory activities. The availability of light on land is closely linked with that of temperature, since the sun is the source for both.



Find out

What is the source of energy for the life in deep ocean trenches where sunlight does not reach?

Soil: The nature and properties of soil are dependent on the climate, the weathering process.

Various characteristics of the soil such as soil composition, grain size, determine the percolation and water holding capacity of the soil. These characteristics along with pH, mineral composition and topography, determine the vegetation of an area. Vegetation in turn dictates the type of animals.

The abiotic factors that determine the type of habitat, also show considerable diurnal and seasonal variations. The plants and animals must adapt to these changes in order to survive and flourish in the habitat. During the course of their evolution, many species have evolved a relatively constant 'internal' environment that permits all biochemical reactions and physiological functions to proceed with optimum rate, and allow the species to flourish. The organisms try to maintain the constancy of its internal environment (homeostasis) despite variations in the external environmental conditions. To survive and flourish in any environment, organisms must adapt to the changes in the environment for which there are following possibilities :

- i. **Regulate:** Some organisms are able to maintain homeostasis by physiological and behavioural changes which ensure constant body temperature, constant osmotic concentration, etc. All birds and mammals are capable of such regulation (thermoregulation and osmoregulation).
- ii. **Conform:** Most of animals and plants cannot maintain a constant internal environment. Their body temperature changes with the ambient temperature. In aquatic animals, the osmotic concentration of the body fluids changes with that of the ambient water osmotic concentration. These animals and plants are simply conformers.

Some species have evolved the ability to regulate, within a limited range of environmental conditions, beyond which they simply conform. If the stressful environment is localized or only for a short period of time, the organism may migrate or suspend its activities.

- iii. **Migrate:** The organism can move away temporarily from the stressful habitat to a more hospitable area and return when stressful period is over. Many animals, particularly birds, during winter undertake long-distance migrations to more hospitable areas.
- iv. **Suspend:** In plants, seeds serve as means to tide over periods of stress; they germinate to form new plants under favourable moisture and temperature conditions. They do so by reducing their metabolic activity and going into a state of 'dormancy'. In animals, the organism, if unable to migrate may go into **hibernation** during winter e.g. polar bear. Some snails and fish go into **aestivation** to avoid summer heat.



Internet my friend

Find out the difference between hibernation and aestivation.



Do you know?

1. Adaptation of plants for aquatic and desert habitats.
2. Adaptations of animals for aquatic and desert habitats.

13.3 Adaptation :

To cope up with extreme variations in their environment, some organisms respond through physiological adjustments, while others do so behaviourally (like migration). These are their adaptations. Therefore, we can say that adaptation is an attribute of the

organism (morphological, physiological, and behavioural) that enables the organism to survive and reproduce in its habitat.

Many desert plants have a thick cuticle on their leaf surfaces and have their stomata in deep pits to minimize loss of water through transpiration. They also have a special photosynthetic pathway (CAM - Crassulacean acid metabolism) that enables their stomata to remain closed during daytime. Some desert plants like *Opuntia*, have their leaves reduced (modified) to spines and the photosynthetic function is taken over by the flattened stems.

Mammals from colder climates generally have shorter snout, ears, tail and limbs to minimize the loss of body heat (Allen's Rule.) In the polar seas, aquatic mammals like seals have a thick layer of fat (blubber) below their skin acting as an insulator to reduce loss of body heat.



Can you tell?

1. What is homeostasis?
2. Why do animals need to maintain homeostasis?
3. What are the adaptations in animals living under crushing pressure at great depths of ocean?

Some organisms show behavioural responses to cope with variations in their environment. Desert lizards manage to keep their body temperature fairly constant by behavioural adaptations. They bask in the sun and absorb heat, when their body temperature drops below the comfort zone, but move into shade, when the ambient temperature starts increasing. Some species burrow into the sand to hide and escape from the heat.

13.4 Population :

In nature, we rarely find isolated, single individuals. They live in groups in a well-defined geographical area, share or compete

for similar resources, potentially interbreed and thus form a population.

Although, it is an individual organism that has to cope with changes in the environment, it is at the population level that natural selection operates to evolve the desired traits. **Population ecology** is, therefore, an important area of ecology because it links ecology to population dynamics, genetics and evolution.

A population has certain attributes of its own, which are different from those of an individual. Basic physical characteristics of population are - its **size** and **density**. Size speaks for the number of individuals in a population while density tells us number of individuals present per unit space, in a given time. Besides size and density, the other characteristics include natality, mortality, immigration, emigration, age pyramids, expanding population, population growth forms and biotic potential. An individual has birth and death, but a population has birth rate and death rate.

Natality is the birth rate of a population. It has the greatest influence on a population's growth. Natality is a crude birth rate or specific birth rate. **Crude birth rate** is used when calculating population size (number of births per 1000 population/year), whereas **specific birth rate** is used relative to a specific criterion such as age. E.g. If in a pond, there were 200 carp fish last year and through reproduction 800 new fish are added, taking the current population to 1000, we calculate the birth rate as $800/200 = 4$ offspring per carp per year.

- **Absolute Natality** : the number of births under ideal conditions (with no competition, abundance of resources such as food and water, etc.).
- **Realized Natality** : the number of births when environmental pressures come into play.

It must be remembered, that absolute natality will be always more than realized natality.

Mortality is the death rate of a population. Mortality rate or death rate, is a measure of the number of deaths (in general, or due to a specific cause) in a particular population, in proportion to the size of that population, per unit of time. Mortality rate is typically expressed in deaths per 1,000 individuals per year. Thus, a mortality rate of 9.5 (out of 1,000) in a population of 1,000 would mean 9.5 deaths per year in that entire population, or 0.95% out of the total.

- **Absolute Mortality** : the number of deaths under ideal conditions (with no competition, abundance of resources such as food and water, etc.).
- **Realized Mortality** : the number of deaths when environmental pressures come into play.

It must be remembered that absolute mortality will be always less than realized mortality.

Sex ratio: The sex ratio of the population affects and is reciprocally affected by birth, death, immigration and emigration rates. It is measured as the ratio of the number of individuals of one sex to that of the other sex. The males and females in a ratio of 1:1 is generally the most common **evolutionary stable strategy (ESS)**.



Can you tell?

What can be the causes of deviation from 1: 1 sex ratio in natural habitat?



Internet my friend

1. Find out the sex ratio of Indian population and the state of Maharashtra.
2. What are the reasons behind deviation of sex ratio among Indian population?

Age distribution and Age pyramid:

A population is composed of individuals of different ages. If the age distribution (per cent individuals of a given age or age group) is plotted for the population, the resulting structure is called an **age pyramid**. For the purpose of simplicity, the entire population is divided into three age groups as **Pre-Reproductive** (age 0-14 years), **Reproductive** (age 15-44 years) and **Post-reproductive** (45-85+ years).



Use your brain power

From the age pyramids given below (Fig.13.3)- what will be your forecast for 15 years from now for the populations of 1. Kenya, 2. Australia, 3. Italy and 4. Hungary.

The size of the population it can support tells us a lot about its status in the habitat. The population size, in natural habitat, could be as low as less than 10 (Siberian cranes in bird sanctuary) or go into millions (*Chlamydomonas* in a pond).

Population size, more technically called population density (designated as N), need not necessarily be measured in numbers only. Although total number is generally the most appropriate measure of population density, it is in some cases either meaningless or difficult to determine.

In an area, if there are millions of termites / ants but only a few animals which feed on them, stating that the population density of these predators is low, will be misleading.

In such cases, the biomass is a more meaningful measure of the population size. Total number is again not an easily adoptable measure, if the population is huge and counting is impossible or very time-consuming. Sometimes, for certain ecological investigations, there is no need to know the absolute population densities; relative densities serve the purpose equally well. For instance, the number of birds / insects caught per trap is good enough measure of their total population density. We are mostly obliged to estimate population sizes indirectly, without actually counting them or seeing them. The tiger census in our national parks and tiger reserves is often based on pug marks and fecal pellets.

Population Growth :

The size of a population for any species is a dynamic parameter. It keeps changing with time, depending on various factors including food, predation pressure and adverse weather. In fact, these changes in population density that give us some idea whether it is flourishing or declining.

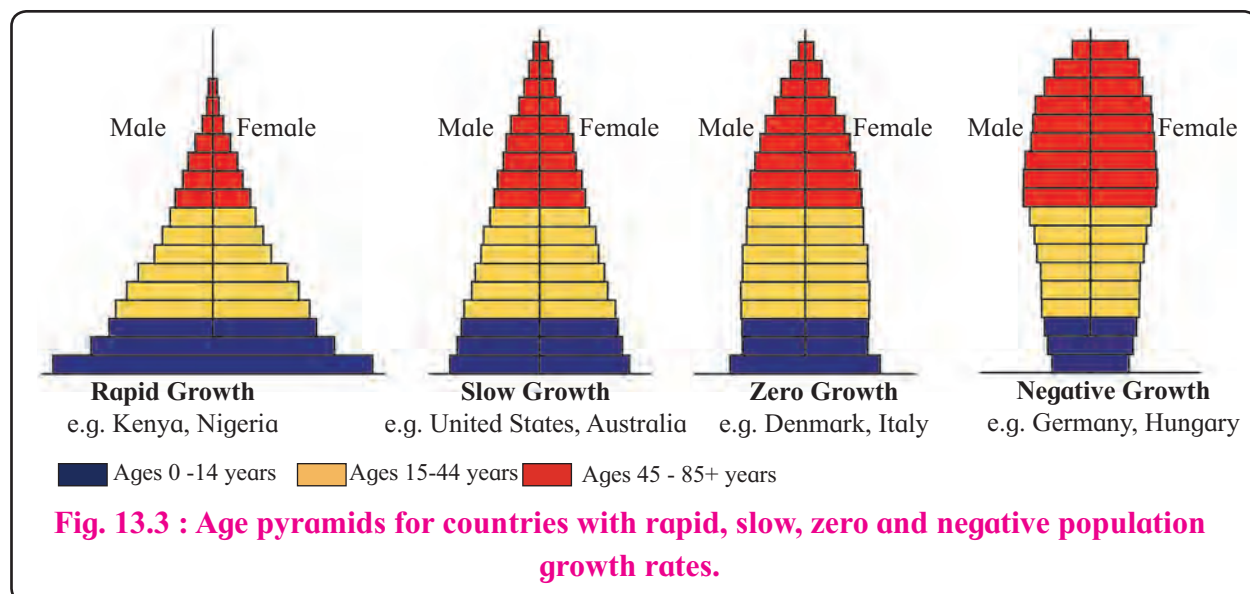


Fig. 13.3 : Age pyramids for countries with rapid, slow, zero and negative population growth rates.

Density of population in a habitat during a given period, fluctuates due to changes in four basic processes. New births (B) and Immigration contribute to an increase in population density. Deaths (D) and Emigration lead to decrease in population density. **Immigration** (I) is the number of individuals of the same species that have come into the habitat from elsewhere during the time period under consideration. **Emigration** (E) is the number of individuals of the population who left the habitat during the time period.

So, if N is the population density at time 't', then its density at time 't + 1' can be calculated as,

$$N_{t+1} = N_t + [(B + I) - (D + E)]$$

Growth Models: Does the growth of a population with time show any specific and predictable pattern? We have been concerned about unbridled human population growth and problems created by it in our country and it is therefore natural for us to be curious if other animal populations in nature behave the same way or show some restraint on growth.

i. Exponential growth:

Resources like food, space are essential for any growth of a population. Ideally, when resources in the habitat are unlimited, each species has the ability to fully realize its innate potential to grow in numbers. Then the population grows in an exponential or geometric proportion.

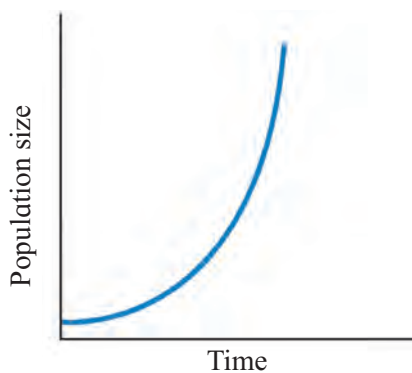


Fig. 13.4 : Exponential growth curve of population

Every species is capable of growing exponentially under unlimited resource conditions, and reach enormous population densities in a short time. Darwin showed how even a slow growing animal like elephant could reach enormous numbers. (provided food and space remain unlimited).

But resources like food and space are not always unlimited. They may be in the beginning; but as the population density increases, so does the competition for those resources, resulting in slowdown in the rate at which the original population was growing. This results in logistic or sigmoid growth curve.

ii. Logistic growth: competition between individuals for limited resources will weed out the 'weaker' ones. Only the 'fittest' individuals will survive and reproduce. In nature, a given habitat has enough resources to support a maximum possible number, beyond which no further growth is possible. Let us call this limit as nature's carrying capacity (K) for that species in that habitat.

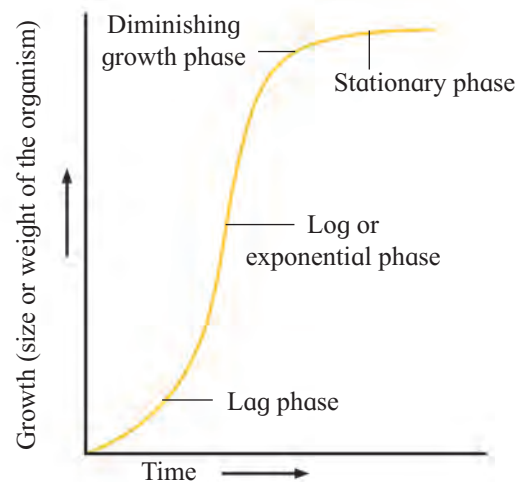


Fig. 13.5 : Logistic growth curve of population

A population growing in a habitat with limited resources show initially a lag phase, followed by phases of acceleration and deceleration and finally an asymptote, when the population density reaches the carrying capacity. A plot of population density (N) in

relation to time (t) results in a sigmoid curve. This type of population growth is called Verhulst-Pearl Logistic Growth.

Since resources for growth of most animal populations, are finite and become limiting sooner or later, the logistic growth model is considered a more realistic one.



Internet my friend

Gather from Census-data the population figures for India for the last 100 years, plot them and check which growth pattern is evident?



Think about it

1. What will happen when carrying capacity of any habitat is exceeded?
2. What could be the reasons behind enormous increase in human population?

Under a particular set of selection pressures, organisms evolve towards the most efficient reproductive strategy. Some produce a large number of small-sized offspring (Oysters, pelagic fishes) while others produce a small number of large-sized offspring (birds, mammals).



Think about it

What can be the reason behind the different reproductive strategies adopted by monocot plants like cereals/ pulses and dicot plants like mango?

13.5 Population Interactions :

There is no natural habitat, which has only one species or animals or plants. For any species, the minimal requirement is another species as food. Even a plant species, which has photosynthetic abilities, cannot survive alone; it needs soil microbes to break down the organic matter in soil and release the inorganic nutrients.

It is obvious that in nature, animals, plants and microbes do not and cannot live in isolation but interact in myriad ways to form a biological community. Interactions may be **intraspecific** i.e. existing between organisms of same population, and **interspecific** -between members of different species. The interspecific interactions occur between minimum two organisms- plants/ animals/ plant and animal. Such interaction may be classified as four types viz, neutralism, negative (harmful), positive (beneficial), and both positive and negative interactions. Even in simplest communities, many interactions exist, not all may be easily seen. Interspecific interactions arise from the interaction of populations of two different species. These interactions could be beneficial, detrimental or neutral (neither harm nor benefit) to one of the species or both.

Table 13.6 : Interspecific Interactions

Sr. no.	Type of interactions with subdivisions	Species	
		A	B
I	Neutralism - no significant effect	O	O
II	Negative interactions		
	a. Competition - direct interference type	-	-
	b. Competition - resource - use type	-	-
III	Positive interactions		
	a. Symbiosis (Mutualism)	+	+
	b. Commensalism	+	O
IV	Both positive and negative interactions		
	a. Parasitism	+	-
	b. Predation	+	-

+ = benefited

- = inhibited

O = not affected

The various types of interactions are classified as per the nature of these interactions to one or both the species. Both the species are benefited in **mutualism** and one or both may be harmed in **competition**.

In **parasitism** and **predation** only one species benefits (parasite and predator, respectively) and the interaction is detrimental to the other species (host and prey, respectively). The interaction where one species is benefitted and the other is neither benefitted nor harmed is called **commensalism**. In **amensalism**, on the other hand one species is harmed whereas the other is unaffected.

Mutualism:

This interaction is obligatory and interdependent. It benefits both the species. Lichens represent an intimate, mutualistic relationship between a fungus and photosynthetic algae or cyanobacteria. The most spectacular and evolutionarily fascinating examples of mutualism are found in plant-animal relationships. Plants need the help of animals for pollinating their flowers and dispersing their seeds. Animals obviously have to be rewarded in the form of pollen and nectar for pollinators and juicy and nutritious fruits for seed dispersers.

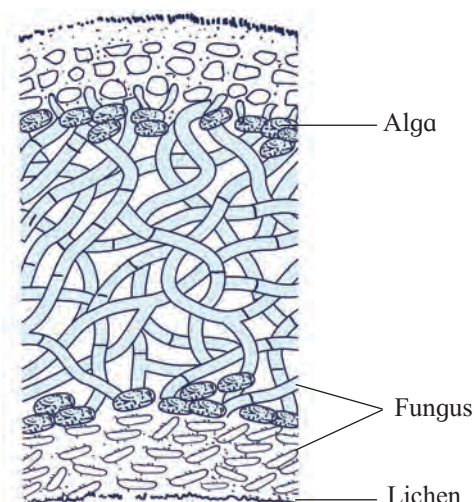


Fig. 13.7 : V. T. S. of Lichen thallus showing Algal and fungal components

But the mutually beneficial system should also be safeguarded against ‘cheaters’, for example, animals that try to steal nectar without aiding in pollination. Plant-animal interactions often involve co-evolution of the mutualists, that is, the evolutions of the flower and its pollinator species are tightly linked with one another.

Competition:

Competition is the type of interaction where both the species are at a loss. Totally unrelated species may compete for the same resource e.g. in shallow creeks on the west coast of Mumbai, visiting flamingos and resident fish compete for their common food, the zooplankton. Secondly, resources need not always be limiting for competition to occur. In competition, the feeding efficiency of one species is reduced due to the interference or inhibitory presence of the other species, even if resources (food and space) are abundant, e.g. Leopards do not hunt in close proximity of lion pride. Therefore, competition is best defined as a process in which the fitness of one species is significantly lower in the presence of another species.

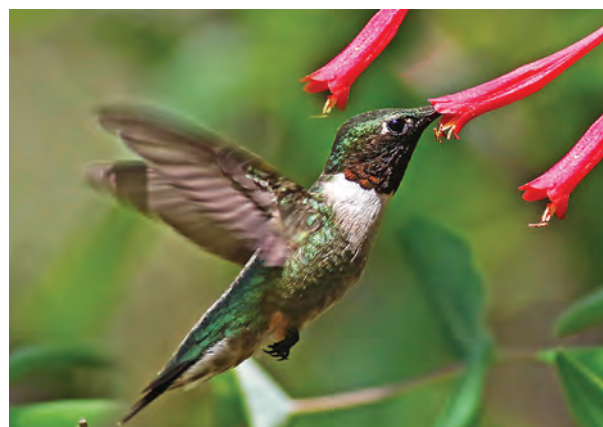


Fig. 13.8 : Co-evolution of humming birds and plants

Gause’s ‘Competitive Exclusion Principle’ states that two closely related species competing for the same resources cannot co-exist indefinitely and the competitively inferior one will be eliminated eventually. This may be

true if resources are limiting, but not otherwise. In interspecific competition with sufficient resources, species facing competition will evolve mechanisms that promote co-existence rather than exclusion. One such mechanism is 'resource partitioning'. If two species compete for the same resource, they could avoid competition by choosing different times for feeding.



Fig. 13.9 : Lion standing over dead leopard- removing competition

A species whose distribution is restricted to a small geographical area because of the presence of a competitively superior species, is found to expand its range when the competing species is removed.

Parasitism:

Parasitism has evolved in so many taxonomic groups from plants to higher vertebrates. Many parasites have evolved to be host-specific (they can parasitize only a single species of host) in such a way that both host and the parasite tend to co-evolve, against each other. In accordance with their life styles, endoparasites evolved special adaptations such as the loss of unnecessary sense organs, presence of adhesive organs or suckers to cling on to the host, loss of digestive system and high reproductive capacity.

The life cycles of parasites are often complex, involving intermediate hosts or vectors to facilitate transfer to the host. The malarial parasite *Plasmodium vivax* needs a vector (mosquito) to spread to other hosts.

Majority of the parasites harm the host. They may reduce the survival, growth and reproduction of the host and may lead to death of the host, thus reducing its population density. They might render the host more vulnerable to predation by making it physically weak.



Use your brain power

1. Should an ideal parasite be able to thrive within the host without harming it?
2. Why didn't natural selection lead to the evolution of such totally harmless parasites?

Parasites that feed on the external surface of the host organism are called **ectoparasites**. The most familiar examples of this group are the lice on humans and ticks on dogs. Many marine fish are infested with ectoparasitic copepods. *Cuscuta*, a parasitic plant that is commonly found growing on hedge plants, has lost its chlorophyll and leaves in the course of evolution. It derives its nutrition from the host plant which it parasitizes.

Brood parasitism in birds is a fascinating example, in which the parasitic bird lays its eggs in the nest of its host bird and lets the host bird incubate them. During the course of evolution, the eggs of the parasitic bird have evolved to resemble the host's egg in size and colour to reduce the chances of the host bird detecting the foreign eggs and ejecting them from the nest. Eggs of the parasitic bird (Asian koel) hatch before that of its host (Common Indian crow).



Fig. 13.10 : Crow feeding Asian Koel hatching in its own nest.

Predation:

When we think of predator and prey, most probably it is the tiger and the deer that readily come to our mind, but a sparrow eating any seed is no less a predator.

Although grazers are animals eating plants, classified as herbivores, they are, not very different from predators. Predators play many important roles. They keep prey populations under control. Without them, prey species would reach very high population densities and cause ecosystem instability e.g. in absence of frogs, locusts increase in density and destroy large tracts of agricultural lands. Biological control, methods adopted in agricultural pest control are based on the ability of the predator to regulate prey population. Predators also help in maintaining species diversity in a community, by reducing the intensity of competition among competing prey species.

When certain exotic species are introduced accidentally or intentionally into a new geographical area, they become invasive and start spreading rapidly due to absence of natural predator, e. g. zebra mussels in the intertidal zone of North America.



Internet my friend

Name the invasive species of plants and animals in India.

If a predator over exploits its prey, then the prey might become extinct and following it, the predator will also become extinct for lack of food. This is the reason why predators in nature are 'prudent'.

Prey species have evolved various defenses to reduce the impact of predation. It is a kind of 'evolutionary arms race'. Better camouflage for concealment, faster speed for escape are for the prey and the predator has to counter these, by its own means. Some species of insects and frogs are cryptically coloured (camouflaged) to avoid being detected easily. The Monarch

butterfly is highly distasteful to its predator (bird) because of a special chemical present in its body. Interestingly, the butterfly acquires this chemical during its caterpillar stage by feeding on a poisonous weed.



Fig. 13.11 : Adult Monarch butterfly displaying warning colour pattern

For plants, herbivores are the predators. Plants therefore have evolved variety of morphological and chemical defenses against herbivores. Thorns (*Acacia*, Cactus) are the most common morphological means of defense. Many plants produce and store chemicals that make the herbivore sick. When chemicals/ produce are eaten, they inhibit feeding or digestion of predator and disrupt reproduction or even kill it. *Calotropis* growing in abandoned fields, produces highly poisonous cardiac glycosides and that is why you never see any cattle or goats browsing on this plant. A wide variety of chemical substances that we extract from plants on a commercial scale, (nicotine, caffeine, quinine, strychnine, opium, etc.,) are secondary metabolites produced by them actually as defences against grazers and browsers.


Commensalism:

This is the interaction in which one species benefits and the other is neither harmed nor benefited. An orchid growing as an epiphyte on

a branch of mango tree, will get benefit while the mango tree derives no benefit. The cattle egret and grazing cattle in close association, is a classic example of commensalism. Cattle egrets always forage close to cattle, as cattle move they flush out insects that might be difficult for the egrets to find and catch.

Another example of commensalism is the interaction between sea anemone that has stinging tentacles and the clown fish that lives among them.

The fish gets protection from predators which stay away from the stinging tentacles. The anemone does not appear to derive any benefit by hosting the clown fish.

 **Do you know ?**

The instrument used to measure the height of forest trees is called hypsometer.

World Environment day - 5th June

World Population day - 11th July

World Earth day - 22nd April

World Ozone day - 16th September



Fig. 13.12 : Commensalism- Cattle egret with buffalo and Clown fish in the tentacles of Sea anemone



Activity :

Prepare chart for various interactions existing in the plant community or animal community around you and paste the photographs of the same.

Exercise

Q. 1 Multiple choice questions.

- Which factor of an ecosystem includes plants, animals and microorganisms?
a. Biotic factor b. Abiotic factor
c. Direct factor d. Indirect factor
- An assemblage of individuals of different species living in the same habitat and having functional interactions is
a. Biotic community
b. Ecological niche
c. Population
d. Ecosystem
- Association between sea anemone and clown fish is that of
a. Mutualism b. Commensalism
c. Parasitism d. Amensalism
- Select the statement which explains best parasitism.
a. One species is benefited.
b. Both the species are benefited.
c. One species is benefited, other is not affected.
d. One species is benefited, other is harmed.
- Growth of bacteria in a newly inoculated agar plate shows
a. exponential growth
b. logistic growth
c. Verhulst-Pearl logistic growth
d. zero growth

Q. 2 Very short answer questions.

- Define the following terms :
a. Commensalism b. Parasitism
c. Camouflage
- Give one example for each :
a. Mutualism
b. Interspecific competition

- Name the type of association:
a. Clown fish and sea anemone
b. Crow feeding the hatchling of Koel
c. Humming birds and host flowering plants
- What is the ecological process behind the biological control method of managing with pest insects?

Q. 3 Short answer questions.

- How is the dormancy of seeds different from hibernation in animals?
- If a marine fish is placed in a fresh water aquarium, will it be able to survive? Give reason.
- Name important defense mechanisms in plants against herbivores.
- An orchid plant is growing on the branch of mango tree. How do you describe this interaction between the orchid and the mango tree?
- Distinguish between the following:
a. Hibernation and Aestivation
b. Ectotherms and Endotherms
c. Parasitism and Mutualism
- Write a short note on :
a. Adaptations of desert animals
b. Adaptations of plants to water scarcity
c. Behavioural adaptations in animals
- Define Population and Community.

Q. 4 Long answer questions.

- With the help of suitable diagram describe the logistic population growth curve.
- Enlist and explain the important characteristics of a population.

Project :

Study the age pyramid of human population in your area.



Can you recall?

1. What is environmental biology?
2. Meaning of the term habitat.
3. Importance of ecology for humans.

An ecosystem is a self regulatory and self sustaining structural and functional unit of nature (biosphere). It contains both biotic and abiotic components. Biotic components interact with each other and also with the surrounding environment. Tansley (1935) coined the term **ecosystem**. Ecosystems vary greatly in size from a small pond to large oceans or small farmland to village. Entire biosphere can be considered as one global ecosystem, made up of many local ecosystems. Since the earth ecosystem is too big and too complex to be studied, it is divided into two basic categories, viz. **terrestrial** and **aquatic**. Forest, grassland and desert are the types of terrestrial ecosystems while lakes, wetlands, rivers, seas, oceans and estuaries are the types of aquatic ecosystems. The ecosystems can also be classified as Natural ecosystems and Artificial ecosystems. **Natural ecosystems** do not require any human inputs, in other words they are self-sustainable. **Artificial ecosystems** e.g. a farm land, a fish tank or even a large pond used for rearing fish, require constant input in terms of energy or materials.

In this chapter, we will study and analyse the structure and functions of the ecosystem, in order to appreciate the input (net productivity), transfer of energy (food chain/web, nutrient cycling) and the output (as degradation and energy loss). We will also look at the relationships, chains and webs that are created because of the energy flows within the system.

14.1 Ecosystem :

Structure and Function :

We have already studied the various biotic and abiotic components of the environment. We know that all these biotic and abiotic components influence each other. Let us now look at these components with an integrated approach and see how the flow of energy takes place in ecosystem. Interaction of biotic and abiotic components, results in a physical structure that is characteristic for each type of ecosystem. Identification and enumeration of plant and animal species of an ecosystem gives its species composition.

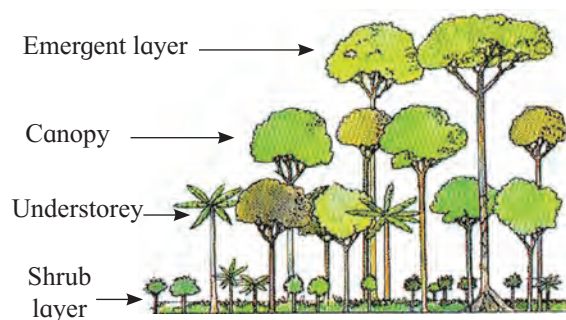


Fig. 14.1 : Stratification of plants in forest

Biotic and abiotic components differ as the locations vary in space and time. The variation due to space results in **spatial pattern**. There are two types of spatial patterns. viz. Stratification and Zonation.

Vertical distribution of different species of plants and animals occupying different levels, is known as **stratification**. For example, trees occupy top vertical strata or layer of a forest, shrubs the second and herbs and grasses occupy the bottom layer. Similarly, stratification is also observed in the open seas as epipelagic, meso-pelagic, bathy-pelagic and benthic zones.



Find out

Stratification of animals in amazon rain forest.

Horizontal distribution of plants and animals on land or in water, is called **zonation**. Zonation is observed in aquatic (wetlands) as well as terrestrial ecosystems, but it is easily seen at the junction of the two. Edges of a large lake or beach show pronounced zonation in the form of Inter-tidal, Littoral, Sub-littoral zones.

Zonation

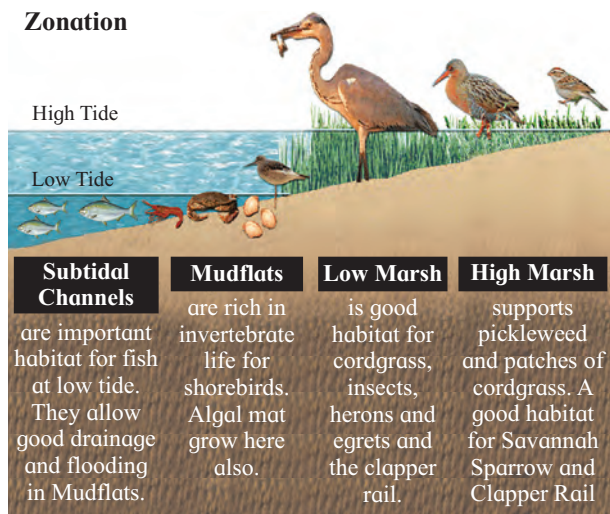


Fig. 14.2 : Zonation in wetland



Think about it

Why is zonation more pronounced at the edges of habitat?

The biotic and abiotic components of an ecosystem are all linked together to function as an 'ecosystem unit' through various processes like, **Productivity, Decomposition, Nutrient cycling and Energy flow**. In fact, these are functional aspects of ecosystem.

Any ecosystem must perform these four processes for its sustenance (**to be self-sustaining**). The ecosystem under study may be as small as a pond or entire biosphere as a whole. The process of **productivity** involves conversion of inorganic chemicals into organic material with the help of the radiant energy of the sun by the autotrophs and consumption of the autotrophs by heterotrophs. The **Decomposition** is the break down of dead organic material and mineralization of the dead matter. The **nutrient cycling** is the storage

and transport of nutrients. (minerals released in decomposition process are used again by autotrophs). The **energy flow** is unidirectional flow of energy from producers to consumers and finally dissipation and loss as heat.

Example- Think of a small pond ecosystem. It is fairly a self-sustainable unit that explains the complex interactions which exist in any aquatic ecosystem. A pond is a shallow water body in which all the above four aforesaid basic processes of an ecosystem are observed. The abiotic component is water with all the dissolved inorganic and organic substances and also the rich soil deposit at the bottom of the pond. The solar input, the cycle of temperature, day-length and other climatic conditions regulate the rate of function of the entire pond. The producers include the phytoplankton, algae and other aquatic plants. The consumers are represented by the zooplankton, aquatic insects and fish. The decomposers are the fungi, bacteria located at the bottom of the pond.

a. Productivity :

A constant input of solar energy is the basic requirement for any ecosystem to function and sustain. **Productivity** refers to the rate of generation of biomass in an ecosystem. It is expressed in units of mass per unit surface (or volume) per unit time, for instance grams per square metre per day ($g/m^2/day$). The mass unit may relate to dry matter or to the mass of carbon generated.

It can be divided into **gross primary productivity**(GPP)and netprimaryproductivity (NPP). **Gross primary productivity** of an ecosystem is the rate of production of organic matter during photosynthesis. Plants themselves use a considerable proportion of this GPP for their respiration. Hence, gross primary productivity minus respiratory losses (R) constitute the net primary productivity (NPP).

$$GPP - R = NPP$$

Net primary productivity is the available biomass for the consumption, to heterotrophs (herbivores, carnivores and decomposers). The annual net primary productivity of the whole biosphere is approximately 170 billion tons (dry weight) of organic matter. Of this, the productivity of the oceans is only 55 billion tons. Rest of course, is from land based ecosystems.



Use your brain power

What could be the reason for the low productivity of ocean?

Primary productivity (GPP) depends on the plant species inhabiting a particular area. It also depends on a variety of environmental factors, availability of nutrients and photosynthetic capacity of plants. Therefore, it varies in different types of ecosystems. **Secondary productivity** is defined as the rate of formation of new organic matter by consumers. Alternatively, it is the rate of assimilation of food energy at the level of consumers. It is the amount of energy available to consumer for transfer to the next trophic level.

b. Decomposition :

Decomposers break down complex organic matter into inorganic substances like carbon dioxide, water and nutrients, and the process is called **decomposition**. Dead remains of plants and animals, including fecal matter, constitute detritus, which is the raw material for decomposition. The important steps in the process of decomposition are fragmentation, leaching, catabolism, humification and mineralization.

Detritivores (e.g. earthworm) break down detritus into smaller fragments or particles. This process is called **fragmentation**.

By the process of **leaching**, water soluble inorganic nutrients go down (percolate) into the soil horizon and get precipitated as unavailable salts. Bacterial and fungal enzymes degrade detritus into simpler inorganic substances. This process is called as **catabolism**. It is important to note that all the above steps in decomposition operate simultaneously on the detritus.

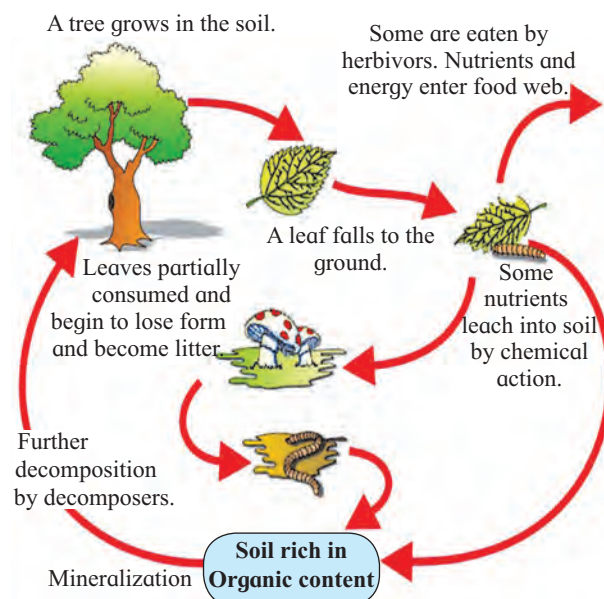


Fig. 14.3 : Decomposition cycle

Humification and mineralization occur during decomposition in the soil. Humification leads to accumulation of partially decomposed, a dark coloured, amorphous, colloidal organic substance called **humus** that is resistant to microbial action and undergoes decomposition at an extremely slow rate. Humus formation changes soil texture and increases water holding capacity of soil.

Being colloidal in nature, humus serves as a reservoir of nutrients. The humus is further degraded by some microbes and release of inorganic nutrients occurs by the process known as **mineralisation**.

Decomposition as a process requires oxygen. Temperature and soil moisture are the most important factors that regulate decomposition indirectly to help soil microbes. Warm and moist environment favours decomposition whereas low temperature and anaerobic conditions inhibit decomposition.



Can you recall?

1. What is a food chain?
2. What are trophic levels in a food chain?

14.2 Energy Flow :

Sun is the only source of energy for all ecosystems on the earth except for the deep-sea ecosystems. Of the total incident solar radiation, less than 50 % of it is **photosynthetically active radiation (PAR)**. Plants and photosynthetic bacteria (autotrophs) fix energy to prepare food from simple inorganic materials. Plants capture only 2-10 % of the PAR and this small amount of energy sustains the entire living world.

Therefore, it is very important to know how the solar energy captured by plants flows through different organisms of an ecosystem. Directly or indirectly, all organisms are dependent for their food on producers. Hence there is **unidirectional flow of energy** from sun to producers and then to consumers. The direction can not be reversed. Energy can be used only once in the ecosystem.

The autotrophs need a constant supply of energy to synthesize the molecules they require. The autotrophs are called **producers**. In a terrestrial ecosystem, major producers are herbaceous and woody plants. Likewise, producers in an aquatic ecosystem are phytoplankton and algae.



Find out

1. Is there any presence of living organisms in the perpetual darkness of deep oceanic trenches?
2. In absence of solar radiation, what is their source of energy?
3. Which organisms do serve as producers in the food chains of deep oceans?

All animals directly or indirectly depend on plants for their food. They are hence

called **consumers** (heterotrophs). If they feed directly on the plants, they are called **primary consumers**, and if the animals eat other animals which eat plants, they are called **secondary consumers**. Likewise, you could have **tertiary consumers** too.

The primary consumers are also known as **herbivores**. Some common herbivores are insects (grasshopper, aphids), birds (parrot) and some mammals (sheep, cattles, goat, donkey) in terrestrial ecosystem and molluscs in aquatic ecosystem. The consumers that feed on these herbivores are **carnivores**, (secondary consumers). Those animals that depend on the primary carnivores for food are called secondary carnivores.

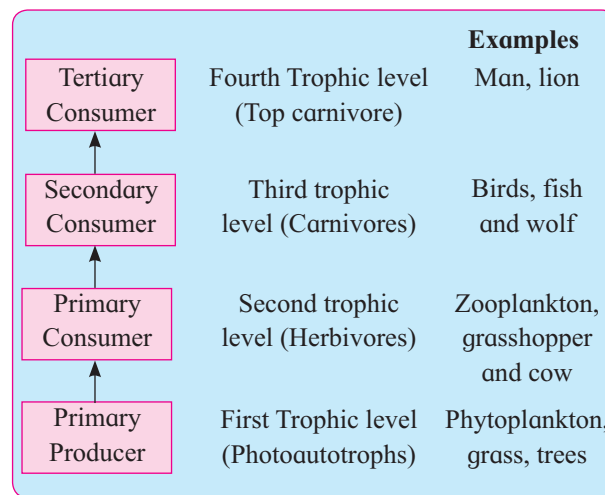
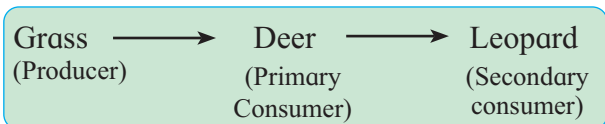


Chart 14.4 : Trophic levels

You have studied several food chains and food webs that exist in nature. Food chains are always straight and usually have four or five trophic levels. There are three types of food chains viz. **grazing**, **detritus** and **parasitic**. Starting from the plants (or producers) food chains and food webs are formed such that an animal feeds on a plant or on another animal and in turn is food for another. The energy trapped by the producer, is either passed on to a consumer or remains trapped till the producer organism dies. Death of organism is the beginning of the detritus food chain/web.

A simple grazing food chain (GFC) is depicted below:



The detritus food chain (DFC) begins with dead organic matter. It is composed of decomposers which are heterotrophic organisms, mainly fungi and bacteria. They meet their energy and nutrient requirements by degrading the detritus. These are known as **saprotrophs (Saprobies)**. Decomposers secrete enzymes that breakdown dead organic materials into simple, inorganic materials, which are absorbed by them. Detritus food chain may be connected with the grazing food chain at some levels. In a natural ecosystem, some animals like cockroaches, crows, bears, man, etc. are **omnivores**. Omnivores eat producers as well as consumers. These natural interconnection of food chains make it a **food web**.

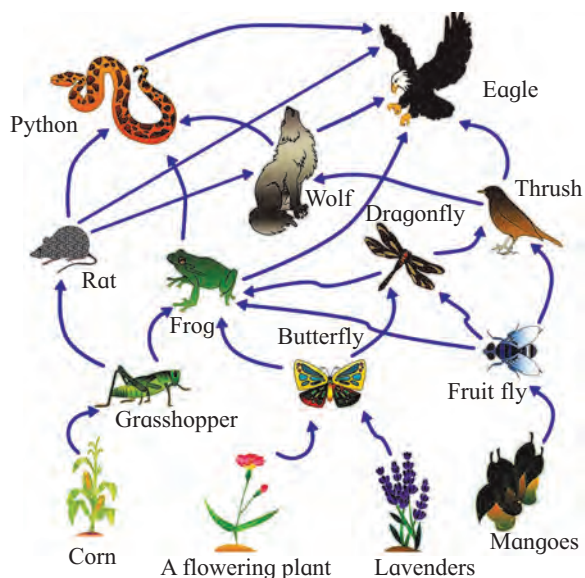


Fig. 14.5 : Food web



Can you tell?

From the given food web diagram, give the trophic levels where the eagle is present.



Use your brain power

1. What could be the connecting points between the GFC and DFC?
2. How will you classify man as carnivore (primary/ secondary) or omnivore? Why?
3. How many trophic levels human beings function in a food chain?

Every organism occupies a place in ecological community according to the source and method of obtaining its food. Organisms occupy a specific place in the food chain that is their **trophic level**. Producers belong to the first trophic level, herbivores (primary consumer) to the second and carnivores (secondary consumer) to the third trophic level.

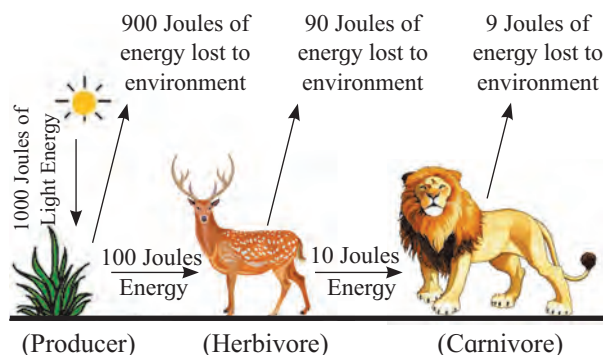


Fig. 14.6 : Energy flow showing increasing trophic level

The amount of energy available decreases at each successive trophic level. The number of trophic levels in any food chain is restricted as the transfer of energy follows ‘**10% Law**’ (R. Lindermann, 1942). The law states that ‘only 10 % of the energy is transferred to each trophic level as net energy, from the previous trophic level’. In nature, it is possible to have different trophic levels - producer, herbivore, primary carnivore, secondary carnivore, tertiary carnivore and ultimate carnivore.



Think about it

What is the maximum number of trophic levels in a food chain?

Beyond secondary carnivores, however the amount of energy available is too less, hence, there is no tertiary carnivore that feeds exclusively on secondary carnivore, even though the secondary carnivore many times will feed on herbivores directly. This is the reason why food chains do not exist in isolation, but are always interconnected to form food web that maintains the stability of an ecosystem.

14.3 Ecological Pyramids :

Ecological pyramid is a graphic representation of the relationship between the organisms of various successive trophic levels with respect to energy, biomass and number.

Pyramid is a structure which has broader base that gradually narrows upwards forming an upright cone like structure. This concept was developed by C. Elton in 1927.

The base of each pyramid represents the producers or the first trophic level while the apex represents tertiary or top level consumer. Any calculations of energy content, biomass, or numbers, has to include all organisms at that trophic level.

The three ecological pyramids which are usually studied are : **Pyramid of biomass**, **Pyramid of numbers** and **Pyramid of energy**.

The relative number of individuals per unit area at different trophic levels, constitutes the **number pyramid**; of biomass/ unit area, is **biomass pyramid** and of amount of accumulated energy per unit area, is **energy pyramid**.

In most well balanced ecosystems, all the pyramids, of number, energy and biomass are upright, i.e. producers are more in number and biomass than the herbivores, and herbivores are more in number and biomass than the carnivores. There are exceptions to this, e.g. oceanic ecosystem show inverted biomass pyramid.

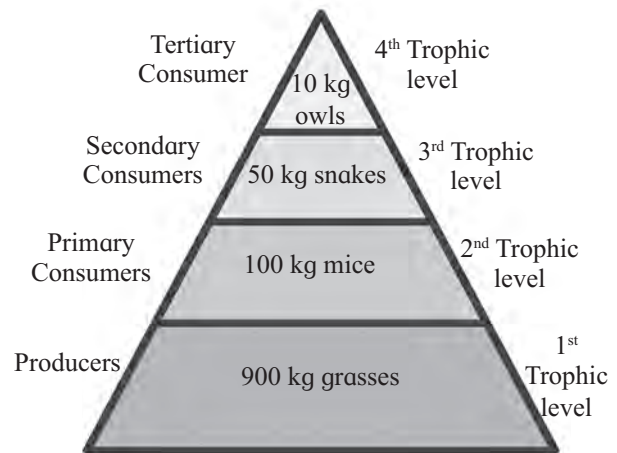


Fig. 14.7 : Pyramid of Biomass



Think about it

How would you explain inverted pyramid of biomass in oceanic ecosystem?

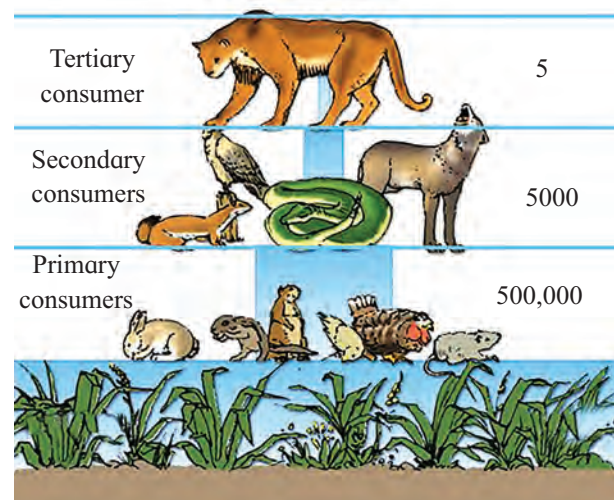


Fig. 14.8 : Pyramid of Numbers

Similarly, pyramid of numbers can also be sometimes inverted. e.g. if we plot the number of insects on a single tree, smaller birds feeding on insects, and parasites on those birds, we get an inverted pyramid.



Use your brain power

What will happen, if in the above example, we substitute larger bird of prey feeding on small insect eating birds?

Pyramid of energy is always upright. It can never be inverted, because when energy flows from a particular trophic level only 10% pass on as net energy to the next trophic level, and large amount of energy is always lost as heat at each step. In smaller food chains, more energy is available than in the longer food chains.

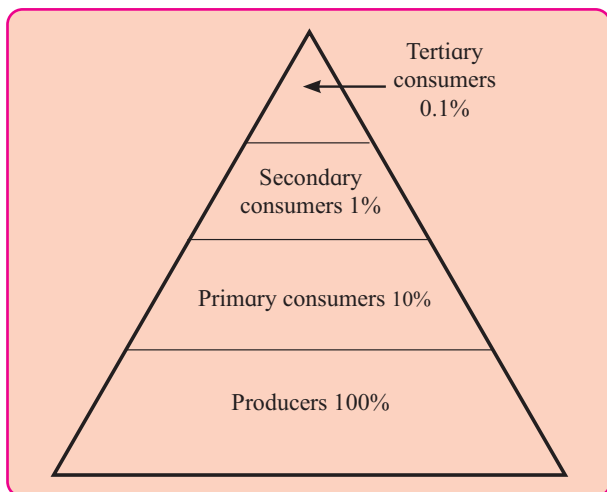


Fig. 14.9 : Pyramid of Energy

There are certain limitations of ecological pyramids. It assumes a simple food chain, something that almost never exists in nature. It does not accommodate a food web. Moreover, saprophytes are not given any place in ecological pyramids even though they play a vital role in the ecosystem.

A given species may occupy more than one trophic levels in the same ecosystem at the same time, e.g. a sparrow is a primary consumer when it eats seeds, fruits, peas, and a secondary consumer when it eats insects and worms.

14.4 Nutrient Cycles :

We have studied in earlier classes that all organisms need a constant supply of nutrients to grow, and reproduce. The amount of nutrients, such as carbon, nitrogen, phosphorus, calcium, etc. present in the soil. Their concentration varies in different kinds of ecosystems on a seasonal basis. These nutrients, which are never lost from the ecosystems, are recycled indefinitely. The movement of nutrient elements through

the various components of an ecosystem, is called **nutrient cycling**. Another name of nutrient cycling is **biogeochemical cycle**. Here, essential elements are cycled from abiotic to biotic world and back.

Types of Nutrient cycles : There are two types of nutrient cycles viz. (a) gaseous and (b) sedimentary. The reservoir for gaseous type of nutrient cycle (e.g., nitrogen, carbon cycle) is the atmosphere and for the sedimentary cycle (e.g. phosphorus cycle) the reservoir is Earth's crust. The function of the reservoir is to meet with the deficit, which occurs due to imbalance in the rate of influx and efflux in any ecosystem.

Carbon Cycle :

All life forms on earth are carbon based because carbon is the main component of all the organic compounds of protoplasm. It constitutes 49% of dry weight of organisms. If we look at the total quantity of global carbon, we find that 71% carbon is found dissolved in oceans. This oceanic reservoir regulates the amount of carbon dioxide in the atmosphere.

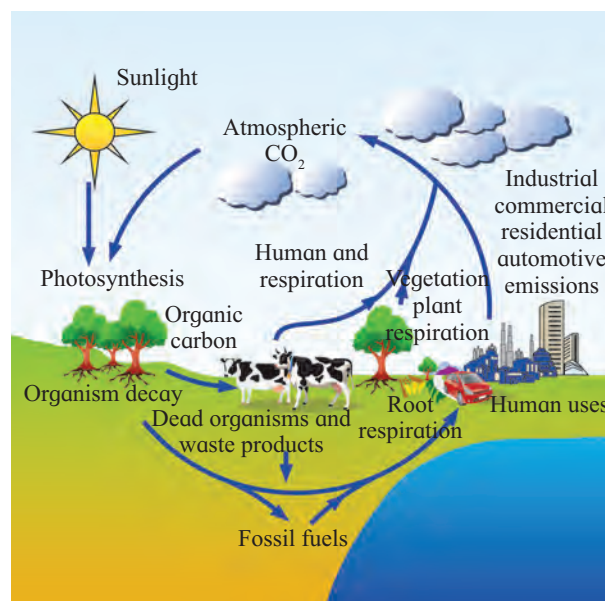


Fig. 14.10 : Carbon cycle

Carbon that is a part of rocks and fossil fuels like oil, coal and natural gas, has been away from the rest of the carbon cycle for a long time. These long-term storage places

are known as “sinks”. When fossil fuels are burned, carbon that had been underground is released back into the air as carbon dioxide.

The element carbon is a part of seawater, the atmosphere, rocks such as limestone and coal, soils, as well as all living things.

- Carbon as CO_2 moves from the atmosphere to plants. Through the process of photosynthesis, carbon dioxide is pulled from the air to produce food.
- Carbon moves from plants to animals, through food chains, i.e. the carbon present in plants moves to the animals.
- Carbon moves from living things to the atmosphere. Each time you exhale, you are releasing carbon dioxide gas (CO_2) into the atmosphere.
- Decomposers also contribute substantially to CO_2 in atmosphere, by their processing of waste materials and dead organic matter of land and oceans.
- When fossil fuels burn to power factories, power plants, motor vehicles, most of the carbon quickly enters the atmosphere as carbon dioxide gas. Each year, 5.5 billion tons of carbon is released through combustion of fossil fuels. Of this massive amount, 3.3 billion tons stays in the atmosphere. Most of the remainder is dissolved in seawater and deposited as calcium or magnesium carbonate compounds which make up shells of marine animals.
- Burning of wood, forest fire and combustion of organic matter, fossil fuel and volcanic activity, are additional sources for releasing CO_2 in the atmosphere.
- Carbon moves from the atmosphere to the oceans. The oceans and other water bodies, absorb some carbon in the form of CO_2 from the atmosphere. The carbon is dissolved into the oceanic water. Some amount of the fixed carbon is lost to sediments and removed from circulation.

- Fossil fuels represent a reservoir of carbon. Carbon cycling occurs through atmosphere, ocean and through living and dead organisms. Human activities have significantly influenced the carbon cycle.
- Rapid deforestation and massive burning of fossil fuel for energy and transport, have significantly increased the rate of release of carbon dioxide into the atmosphere.

Thus, the entire carbon cycle is run by basic processes viz. Photosynthesis, Respiration, Decomposition, Sedimentation and Combustion.

Phosphorus Cycle :

Cyclic movement of phosphorus through hydrosphere, lithosphere and biosphere constitutes phosphorus cycle.

Phosphorus is a major constituent of biological membranes, nucleic acids and cellular energy transfer systems. Many animals also need large quantities of this element to make shells, bones, hooves and teeth.

The natural reservoir of phosphorus is rock, which contains phosphorus in the form of phosphates. When rocks are weathered, minute amounts of these phosphates dissolve in soil solution and are absorbed by the roots of the plants. Herbivores and other animals obtain this element from plants. The waste products and the dead organisms are decomposed by phosphate-solubilizing bacteria releasing phosphorus. Unlike carbon cycle, there is no respiratory release of phosphorus into atmosphere.

There are two major differences between carbon and phosphorus cycle. Atmospheric inputs of phosphorus through rainfall are much smaller (measure) than carbon inputs, and secondly, exchanges of phosphorus between organism and environment are negligible as compare to carbon.

Unlike carbon and nitrogen, Phosphorus is always in short supply and hence acts as

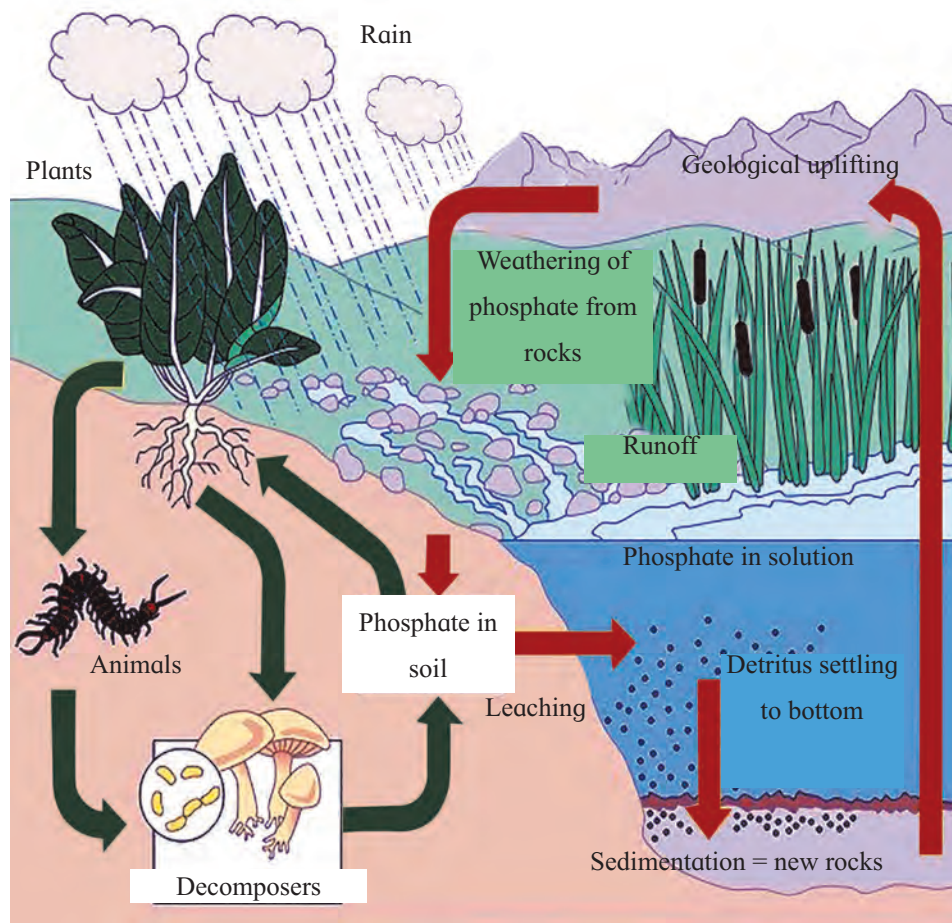


Fig. 14.11 : Phosphorus Cycle

limiting factor for the plant growth. Sudden influx of phosphorus in the form of agricultural runoff or industrial effluents rich in phosphate content, leads to **eutrophication** in water bodies. Eutrophication is due to overgrowth of algae at the instance of high phosphorus dissolved in water. The overgrowth of algae kills or harms the aquatic life.



Internet my friend

What are the long term and short-term effects of eutrophication in water bodies?

14.5 Ecological Succession :

Succession : It is a spatial pattern which occurs over the time. Time is a crucial factor. The ecosystem is initially occupied by the pioneer species. As time passes, species diversity increase giving way to more complex organisms.

Eventually, it leads to **climax community**. Climax community does not evolve further.

The gradual and predictable change in the species composition of a given area is called **ecological succession**. The change is sequential and environmentally regulated.

Process of succession involves sequential steps like Nudation, Invasion, Ecesis, Aggregation, Competition and co-action, Reaction and stabilization.

During succession, some species colonize an area and their populations become more numerous, whereas populations of other species decline and even disappear.

The entire sequence of communities that successively change in a given area, constitute what is called **sere(s)**. Alternatively, it is an entire gradient of organisms from pioneer stage to climax stage. The individual transitional

communities are termed **seral communities**. In the successive seral stages, there is a change in the diversity of species of organisms, increase in the number of species and organisms as well as an increase in the total biomass.

The present day communities in the world have come to be, because of succession that has occurred over millions of years since life started on earth. Succession is hence a process that starts where no living organisms were present before - like on a newly formed volcanic island. This is called **primary succession**.

Other examples of areas where primary succession occurs are newly cooled lava, rocks and newly created pond or reservoir. The establishment of a new biotic community is generally very slow. Before a biotic community of diverse organisms can become established, there must be soil. Depending mostly on the climate, it takes natural processes, several hundred to several thousand years to produce fertile soil on bare rock.

Secondary succession begins in areas where natural biotic communities have been destroyed such as in abandoned farm lands, burned or cut forests, lands that have been flooded, etc. Since some soil or sediment is present, succession is faster than in primary succession. Description of ecological succession usually focuses on changes in vegetation. However, these vegetational changes in turn affect food and shelter for various types of animals. Thus, as succession proceeds, the numbers and types of animals and decomposers also change. At any time during primary or secondary succession, natural or human induced disturbances (fire, deforestation, etc.), can convert a particular seral stage of succession to an earlier previous / preceding stage. Also, such disturbances create new conditions that encourage some species and discourage or eliminate other species.

Succession of Plants :

Based on the nature of the habitat – whether it is water (or very wetland areas) or it is on very dry areas – succession of plants, is called

hydrarch (hydrosere) or xerarch (xerosere), respectively.

Hydrarch succession takes place in wetter areas and the successional series progress from hydric to the mesic conditions. As against this, xerarch succession takes place in dry areas and the series progress from xeric to mesic conditions. Hence, both hydrarch and xerarch successions lead to medium water conditions (mesic) – neither too dry (xeric) nor too wet (hydric).

The species that invade a bare area, are called **pioneer species**. In primary succession on rocks these are usually crustose lichens which are able to secrete acids to dissolve rock, helping in weathering of rocks and soil formation. These pave the way for bryophytes, mosses that are able to take hold in the small amount of soil.

They are, with time, succeeded by herbaceous plants, and after several more stages, ultimately a stable climax forest community is formed. The **climax community** remains stable as long as the environment remains unchanged.

In **primary succession** of aquatic habitat, (steps 1 to 7 in fig. 14.13) the pioneers are the small phytoplankton. They are replaced with time by rooted-submerged plants (e.g. *Hydrilla*), rooted-floating angiosperms (e.g. Lotus) followed by free-floating plants (e.g. *Pistia*), then reed swamp (e.g. *Typha*), marsh-meadow (e.g. *Cyperus*), scrub (e.g. *Alnus*) and finally the trees (e.g. *Quercus*). The climax again would be a forest. With passage of time, the water body is converted into land.

In **secondary succession**, the species that invade depend on the condition of the soil, availability of water, the environment as also the seeds or other propagules present. Since soil is already there, the rate of succession is much faster and hence, climax is also reached more quickly. Figure 14.14 shows the sequence of stages 1 to 8 in a forest ecosystem after fire.

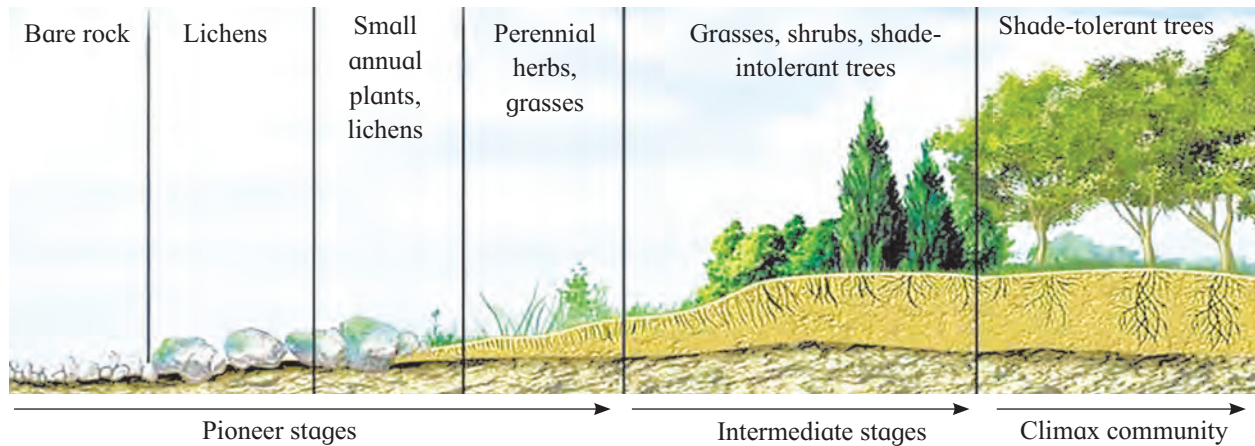


Fig. 14.12 : Xerarch Succession of Plants

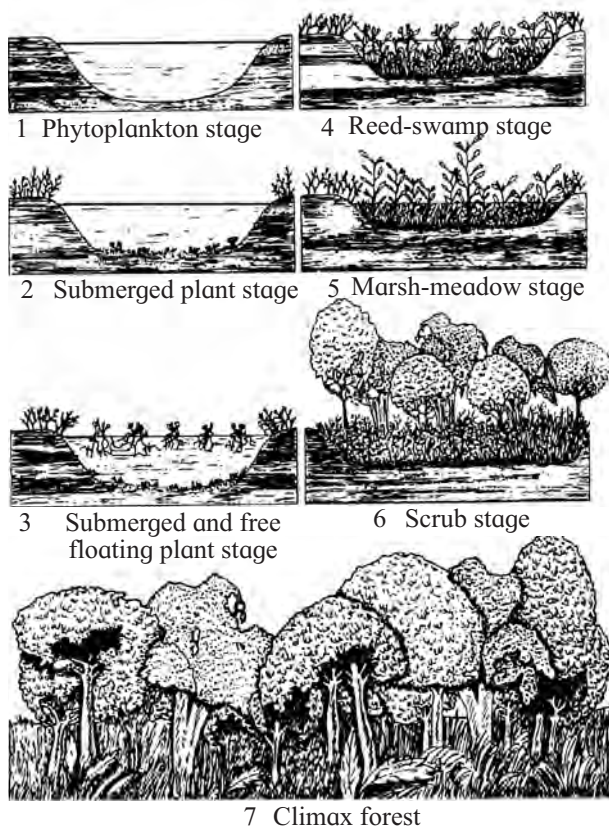


Fig. 14.13 : Hydrarch Succession of Plants

It is important to understand that succession, particularly primary succession, is a very slow process, maybe requiring thousands of years for the climax to be reached.

Another important fact to understand is that all the successions whether taking place in water or on land, proceed to a similar climax community – the mesic.

14.6 Ecosystem Services :

Healthy ecosystems are the base for a wide range of economic, environmental and aesthetic goods and services. The products of ecosystem processes are named as **ecosystem services**, for example, healthy forest ecosystems purify air and water, mitigate droughts and floods.

The Millennium Ecosystem Assessment report 2005 defines *Ecosystem services* as benefits people obtain from ecosystems and identifies four categories of ecosystem services as follows.

- **Supporting services** include services such as nutrient cycling, primary production, soil formation, habitat provision and pollination maintaining balance of ecosystem.
- **Provisioning services** include food (including seafood), raw materials (including timber, skins, fuel wood), genetic resources (including crop improvement genes, and health care), water, medicinal resources (including test and assay organisms) and ornamental resources (furs, feathers, ivory, orchids, butterflies, etc.)
- **Regulating services** include Carbon sequestration, Predation regulates prey populations, Waste decomposition and detoxification, Purification of water and air, and pest control.

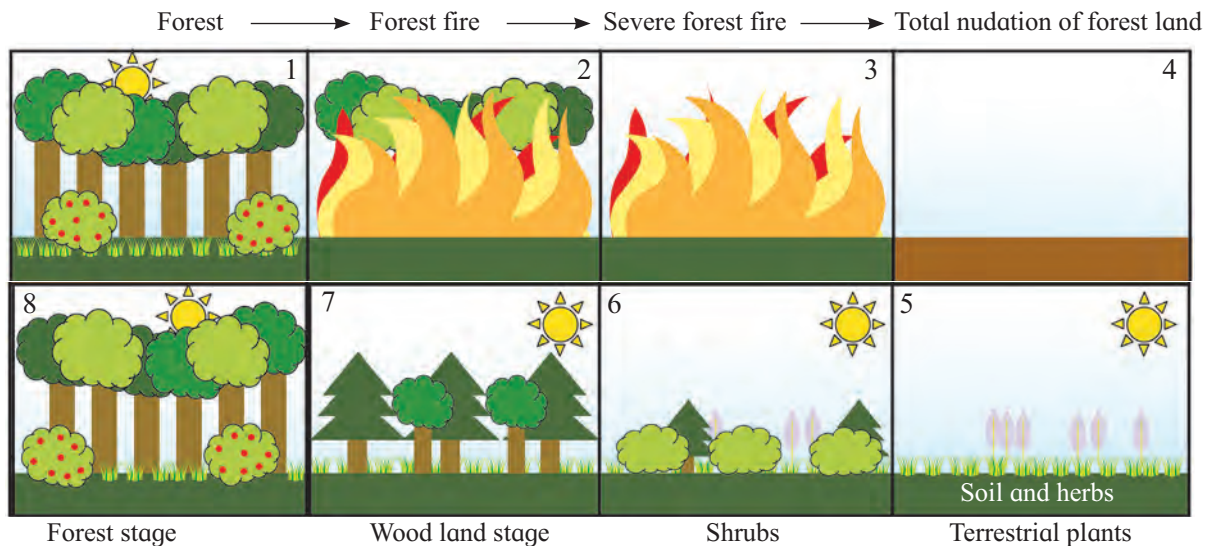


Fig. 14. 14 : Secondary succession in a forest ecosystem after fire

- **Cultural services** include cultural, spiritual and historical, recreational experiences, science and education, and Therapeutics (including animal assisted therapy)

Following are the main ecological services :

Fixation of atmospheric CO₂ and release of O₂ are the most important services provided by an ecosystem. Photosynthetic activity of photoautotrophs sequesters carbon (in CO₂ form) from the atmosphere and releases O₂ as a

byproduct. O₂ not only purifies air but is also used for respiration by all aerobes.

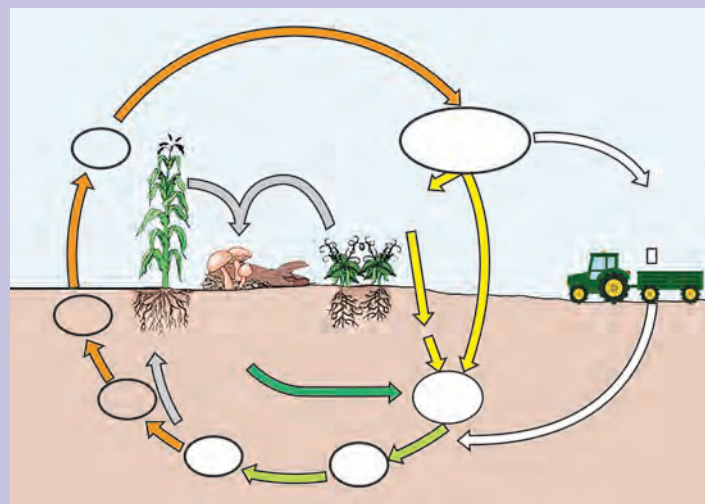
Pollination of plants brought about by wind, water or other biotic agencies, is also an important ecosystem service, without which there would be no crops and no fruits.

Though the value of all such services of biodiversity is difficult to determine, it seems reasonable to think that biodiversity should carry a hefty price tag.



Activity :

1. Fill in the blanks in the given diagram and identity.



2. Collect information and prepare a chart of the sequential steps involved in the ecological succession, explaining each step.

Exercise

Q. 1 Multiple choice questions

1. Which one of the following has the largest population in a food chain?
 - a. Producers
 - b. Primary consumers
 - c. Secondary consumers
 - d. Decomposers
2. The second trophic level in a lake is _____
 - a. Phytoplankton
 - b. Zooplankton
 - c. Benthos
 - d. Fishes
3. Secondary consumers are _____
 - a. Herbivores
 - b. Producers
 - c. Carnivores
 - d. Autotrophs
4. What is the % of photosynthetically active radiation in the incident solar radiation?
 - a. 100%
 - b. 50 %
 - c. 1-5%
 - d. 2-10%
6. Give the term used to express a community in its final stage of succession?
 - a. End community
 - b. Final community
 - c. Climax community
 - d. Dark community
7. After landslide which of the following type of succession occurs?
 - a. Primary
 - b. Secondary
 - c. Tertiary
 - d. Climax
8. Which of the following is most often a limiting factor of the primary productivity in any ecosystem.
 - a. Carbon
 - b. Nitrogen
 - c. Phosphorus
 - d. Sulphur

Q. 2 Very short answer question.

1. Give an example of ecosystem which shows inverted pyramid of numbers.
2. Give an example of ecosystem which shows inverted pyramid of biomass.
3. Which mineral acts as limiting factor for productivity in an aquatic ecosystem.
4. Name the reservoir and sink of carbon in carbon cycle.

Q. 3 Short answer questions.

1. Distinguish between upright and inverted pyramid of biomass
2. Distinguish between Food chain and Food web.

Q. 4. Long answer questions.

1. Define ecological pyramids and describe with examples, pyramids of number and biomass.
2. What is primary productivity? Give brief description of factors that affect primary productivity.
3. Define decomposition and describe the processes and products of decomposition.
4. Write important features of a sedimentary cycle in an ecosystem.
5. Describe carbon cycle and add a note on the impact of human activities on carbon cycle.

Project :

1. Collect the information on the various services offered by dense forest ecosystem.
2. Collect the information of various types of pollinators and the impact of human activity on them.



Can you recall?

1. You have learnt about Chipko movement and efforts of Shri. Jadav Payeng in school. What is the importance of such activities?
2. Why should we have national parks and sanctuaries?
3. We have read about how Indian scientists won the battle for patent of Turmeric and Basmati rice. Why was gaining these patents essential?
4. What is *in situ* and *ex situ* conservation?

Diversity is variety. This variety of life is called biodiversity. **Biodiversity** includes a vast array of species of microorganisms- viruses, algae, fungi, plants and animals occurring on Earth, either in terrestrial or aquatic habitat and the ecological complexes of which they are part.

The diversity is with respect to size (microscopic to macroscopic), shape, colour, form, mode of nutrition, type of habitat, reproduction, motility, duration of life cycle span, etc. This is actually due to the attempt of living beings to accommodate with different environmental conditions (like climatic, edaphic, topographic, geographic, etc.) or situations, solely for their survival and perpetuation. In doing so, living organisms adapt themselves to overcome different situations and thus develop distinct but different features and that has actually led to the diversity in them. The diversity in features become infused in the life cycle. In short, these adaptations in different environments serve as basis for diversity.

Definition of Biodiversity : It is the part of nature which includes the differences in the genes among the individuals of a species; the variety and richness of all plants and animal species at different scales in a space - local regions, country and the world; and the types of ecosystem, both terrestrial and aquatic, within a defined area. In nutshell, biodiversity is the totality of genes, species and ecosystem in a region (as per IUCN, UNEP and WRI).

The term biodiversity was actually coined by Walter Rosen (1982) but the term was popularised by sociologist Edward Wilson to describe combined diversity at all the levels of biological organisation.

Biodiversity that we see today, is the outcome of over 3.5 billion of years of evolutionary history mainly influenced by natural processes and of late by influence of humans.

In this chapter, we shall study the basic concepts of biodiversity such as levels and patterns of biodiversity, expanse, importance, loss and conservation methods and efforts undertaken.

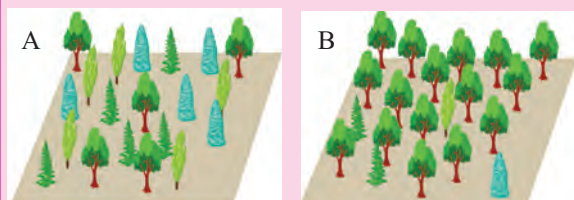
15.1 Levels of Biodiversity:

Diversity of living world can be observed at various levels, ranging from molecular to ecosystem level. Major hierarchical and interrelated levels are genetic diversity, species diversity (community), and ecosystem diversity (ecological).



Can you tell?

What can you say about species diversity A and B?



a. Genetic diversity:

It is the **intraspecific diversity**. It is the diversity in the number and types of genes as well as chromosomes present in different species and also the variation in the genes and their alleles in the same species. It includes variation within a population and diversity between populations that are associated with adaptation to local conditions. Genetic variations (e.g. allelic genes) lead to individual differences within species. Such variations pave way to evolution. They also improve chances of continuation of species in the changing environmental conditions or allow the best adapted to survive. Existence of subspecies, races are examples of genetic diversity. Greater the diversity, better would be sustenance of a species. There are about 1000 varieties of mangoes and 50,000 varieties of rice or wheat in India.

Another case of genetic diversity is a medicinal plant *Rauwolfia serpentina* which secretes active component reserpine, is found in different Himalayan ranges. This plant shows variations in terms of potency and concentration of active chemical, from location to location.

Genetic diversity or variability is essential for a healthy breeding population of a species.

b. Species diversity:

It is the **interspecific diversity**. The number of species of plants and animals that are present in a region, constitutes its species diversity. Some areas or regions are richer in species than in the other regions. **Species diversity** deals with variety of species (**species richness**) as well as number of individuals of different species (**species evenness**) observed in area under study. E.g. amphibian species diversity is more in western ghats than in eastern ghats. Natural undisturbed tropical forests have much greater species richness than monoculture plantation of timber plant, developed by forest plantation. India is one among 15 nations that are rich in species diversity.

c. Ecological (Ecosystem) diversity:

It is related to the different types of ecosystems/ habitats within a given geographical area. There are a large variety of ecosystems on Earth having their own complement of distinctive interlinked species, based on the differences in the habitat. It can be described for a specific geographical region. Generally, there may be one or many different types of ecosystems in a region. Thus, ecosystem diversity is very high in India while it is quite low in Norway. In India, we can find a great variety of ecosystems - deserts, rain forests, deciduous forests, estuaries, wetlands, grasslands, etc. The Western ghats show great ecosystem diversity while regions like Ladakh and Rann of Kutch do not show variance like we observe in Western ghats.

The diversity of life at all the three levels is now rapidly being modified by modern man.

15.2 Patterns of Biodiversity:

There are two patterns viz, Latitudinal and Altitudinal gradient, and species-area relationship.



Think about it

What are latitudes and longitudes? Which of these imaginary lines are more significant with reference to diversification of living beings? Why?

a. Latitudinal and altitudinal gradients :

Biodiversity, barring Arid/ Semiarid and aquatic habitat, show latitudinal and altitudinal gradient.

Latitudinal : Ecological studies have revealed that the distribution of diversity is not uniform around the Globe. Species richness exhibits latitudinal gradient for many plants and animals (if not all). It has been observed that species richness is high at lower latitudes and there is a steady decline towards the poles.

Factors like overall stability of tropical regions for millions of years, lesser climatic changes throughout the year and availability of plenty of sunlight that favoured speciation. Tropical areas have less often experienced drastic disturbances like periodic glaciations observed at poles. Such a stability over millions of years might have favoured speciation. Lesser migrations in tropics might have reduced gene flow between geographically isolated regions and favoured speciation. Scientists also have considered availability of more intense sunlight, warmer temperatures and higher annual rainfall in tropics, as factors responsible for bountifulness of these regions. In more or less constant climatic conditions and abundance of resources, some animals enjoy food preferences. For e.g. fruits being available throughout the year in rain forests, variety of frugivorous organisms is obviously more as compared to the temperate regions.

In short, species richness or diversity for plants and animals decreases as we move away from equator to the poles. It is maximum in tropical rain forests e.g. Amazon rain forest (40,000 plants, 1300 birds, 427 mammals, 3000 species).

Altitudinal : It speaks for the height from mean sea level (MSL) upwards. Species diversity is more at lower altitudes than at the higher altitudes. It is because at higher altitudes, change in the climatic conditions and drastic seasonal variations, lead to the decrease in the species diversity.

b. Species-Area relationships :

Scientists have tried to establish relationship between species diversity and the size of the habitat. It is considered that number of species present is directly proportional to the area. It is understood that larger areas may have more resources that can be distributed amongst the inhabitant species. Does this always hold true?

German naturalist Alexander Von

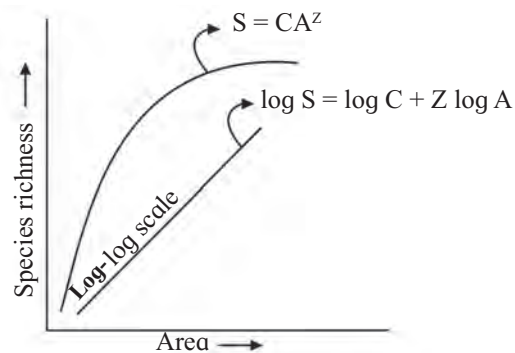


Fig. 15.1 : Graph showing species area relationship

Humboldt observed that species richness does increase with the increase in area but upto certain limit. Observe the graph for species-area relationship. For many species, this relationship is in the form of rectangular hyperbola. If we consider S to be species richness, A as area under study, C as the Y intercept and Z as the slope of the line, this relationship can be described by the equation,

$$\log S = \log C + Z \log A.$$

On logarithmic scale this relationship is a straight line, as observed in the figure above.

For smaller areas, value of Z ranges between 0.1 to 0.2 regardless of species or region under study.

But for the larger areas like the entire continents, slopes are closer to vertical axis i.e. steeper. This observation indicates that in very large areas, number of species found, increase faster than the area explored.



Can you tell?

1. What is biodiversity? Explain genetic diversity with suitable example.
2. Species richness goes on decreasing as we move from equator to pole. Explain.

Importance of species diversity to the ecosystem:

Let us now understand whether we really need all the diversity around us. What if few species are lost permanently?

A community is said to be stable, if average biomass production remains fairly constant over a period of time. It should be strong enough to withstand disturbances and recover quickly. It also must be resistant to invasive species. David Tillman carried out various field experiments and proved that species richness does help the stability of an ecological community. Rich diversity leads to lesser variation in biomass production over a period of time. This is called **Productivity-Stability Hypothesis**.

Paul Ehrlich, an ecologist from Stanford gave an analogy to explain significance of diversity. It is called **Rivet Popper Hypothesis**. He compared Aeroplane to ecosystem and the species as rivets that keep all parts of the aeroplane together. Ofcourse, there are thousands of rivets needed to hold all the parts of the aeroplane together. If each passenger decides to pop even one rivet or in other words, if one species gets extinct, initially not much of the turbulence will be experienced but slowly, as number of popped rivets will increase, there will be a serious threat to the safety of the aeroplane. Also, which rivets are removed will also matter.

Suppose, rivets at key positions such as the ones that bind the wings to the body of the aeroplane, are removed, situation will become serious. Thus, we can say that relationship between diversity and well being of ecosystem is not linear. But it is certain that loss of species may not pose threat to the ecosystem only initially. Loss of key species will certainly cause threat in very short span of time. It will affect food chains, food web, energy flow, natural cycles, etc. In short it will affect the balace of ecosystem.

15.3 Biodiversity Current Scenario

How many species do really exist on earth and how many of them are found in India? It is difficult to come to consensus about the exact

number of species that are present on earth today. Though over 1.5 million species have been documented as per IUCN data (2004) so far, we are yet to study lot more than these. We are also unaware about speciation process that might have continued. Most of the studies that have been carried out are in temperate regions.

Tropical rain forests, the major diversity hubs, are yet to be explored completely. Some exorbitant numbers like existence of 20 to 50 million varieties have been made. But Robert May has given convincing estimate of about 7 million species round the globe. Observe the given pie charts and find out the relative share of various plant and animal groups in the existing knowledge of biodiversity.

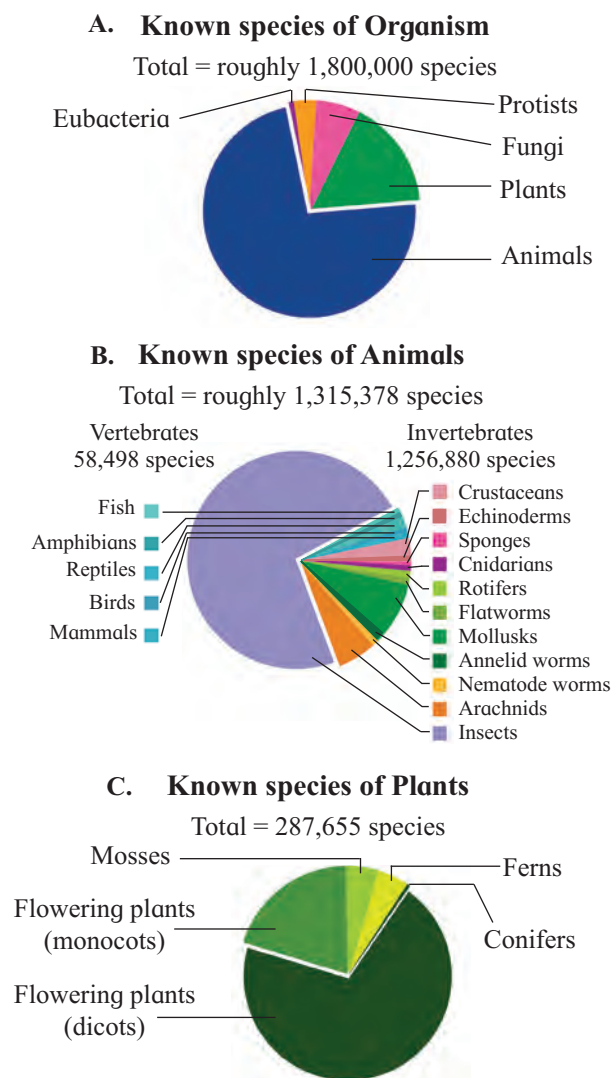


Fig. 15.2 : Graphic representation of known animal and plant groups

In the diagrams, we do not find any data of prokaryotes. Several moneran species are not cultivable under laboratory conditions. Also, conventional taxonomic methods are not suitable for identification of prokaryotic species.

India boasts a handsome share of 8.1% of total biodiversity wealth of the earth. One of the 12 megadiversity countries of the globe, India has 2.4% of total land area of the world. We have identified around 45000 plant species and nearly double the number of animal varieties from our natural wealth. If we consider May's estimate of global biodiversity, we have recorded only 22% of our natural wealth. This situation underlines the need of taxonomists to study the biodiversity. But major concern is the possibility of loss of these varieties before we identify them because of activities like reclamation and deforestation.



Do you know ?

Recently a group of naturalists proved that a lizard from Amboli ghat of Maharashtra was misidentified. They proved it on the basis of DNA profiling as well as number of glands and scales present on the legs of the lizard. Earlier thought to be *Hemidactylus brookii*, the lizard is now renamed as *Hemidactylus varadgirii* in the honour of renowned herpetologist and conservationist, Dr. Varad Giri. They also proved that *brookii* variety is not found in India.



Internet my friend

1. Find out information about Zoological Survey of India and Botanical Survey of India.
2. You may visit Zoological Survey of India, Pune office and find out how information about natural wealth is conserved.

15.4 Loss of Biodiversity:

Loss of biodiversity leads to the overall imbalance in the ecosystem. The chief serious aspect of loss of biodiversity is extinction of species. There are three types of extinction viz, **natural** extinction, **mass** extinction and **man-made** (anthropogenic) extinction.

Damage to biodiversity takes place due to both, natural and manmade reasons. Natural reasons include forest fires, earthquakes, volcanic eruptions etc. Manmade reasons are habitat destruction, hunting, settlement, overexploitation and reclamation (of mangrove lands).

Major mass extinction events occurred between pleistocene and holocene epoch; between Cretaceous and Tertiary period; between Triassic and Jurassic period; between Permian and Triassic period; between Devonian and Carboniferous period; between Ordovician and Silurian period in the geological time scale where plants as well as animal groups underwent mass extinction.

We are aware of six mass extinctions during various stages of history of earth (e.g. ice age). The current loss of biodiversity is considered to be the 7th extinction which is progressing at an alarming rate which is estimated to be 100 to 1000 times faster than prehuman times. Ecologists blame this to the human intervention in natural habitats. They do not forget to warn that if the current trend continues, we might lose about 50% of diversity. Loss of biodiversity in any area can lead to the decline in plant production, lower resilience to environmental disturbance like flood. It may also lead to alteration in environmental processes like disease cycles, plant productivity etc.

Causes of Biodiversity losses:

There are four major causes popularly known as, 'The Evil Quartet'.

i. Habitat loss and fragmentation:

It is the prime cause of destruction. Reduction in vast natural habitats and local degradation by pollution, create crisis situation for the living beings. Loss of local habitat due

to human activities, creates threat to migratory birds as well as those animals that need larger territories. Tropical rain forests are being lost at an alarming rate. Tropical rain forest cover has reduced from 14% to 6% over the years.



Do you know ?

Great Indian Bustard alias *Maldhok* is a critically endangered bird. It is considered as flagship species of grasslands. We lost last *Maldhok* from Maharashtra in 2018. Now this bird is found in Rajasthan and Gujrat. It is an example of local extinction due to habitat loss and hunting.

ii. Over-exploitation:

Basic difference between human beings and other animals is that, humans have the tendency to accumulate beyond their needs. This has resulted in the overexploitation of



a. Dodo bird

b. Stellar sea cow



c. Passenger pigeon

Fig. 15.3 : Examples of extinction due to over-exploitation

resources which in turn causes threats to various organisms. Can you correlate this with dirth of fish, we observe now a days? Dodo bird, stellar sea cow and passenger pigeon are few examples of extinction due to overexploitation.

iii. Alien species invasion:

When a new species gets introduced into any ecosystem accidentally or intentionally, there are chances that it proves harmful for existing species. Sometimes, it can lead to extinction of local species. In such a case, it is called as **invasive species**. E.g. the carrot grass (*Parthenium*), *Lantana* and water hyacinth (*Eichhornia*). Introduction of **predator fish** - Nile perch in Lake Victoria, proved deleterious for 200 **local species** of Cichlid fish.

In India, introduction of African catfish *Clarias gariepinus* for aquaculture purpose has proved harmful to endemic catfish varieties. One of the major reasons of such a harmful effect of alien species is, lack of local predator.

iv. Co-extinctions:

Many a times, organisms are associated with each other in obligatory way. In such cases, extinction of one variety leads to loss of associate variety from the ecosystem. e.g., Extinction of host fish causes extinction of unique parasites. Coevolved plant-pollinator, also will have such a threat.

We are aware of threat to diversity and loss of species from earth. When any species is totally eliminated from earth, it is called **extinct**. e.g. Dinosaurs. When the number of members of a species starts dwindling, it is said to be **endangered**. The International Union for Conservation of Nature and Natural Resources (IUCN) maintains a Red Data Book also known as Red List, where conservation status of plant and animal species is recorded.



Internet my friend

What is ballast water and how can it bring about introduction of alien species into any ecosystem?



Do you know ?

Many a times, we read about leopard attacks on humans or about elephants from Karnataka destroying agricultural lands and orchards in Sindhudurg region of Maharashtra. With increase in human population, man started encroaching forest land. Animals either out of sheer curiosity (in case of young leopards) or for lack of sufficient resources, venture out from their original place. This results in Man-Animal conflict.

Various measures are adapted by forest department to minimise this tussle. e.g., Government not only gives monetary compensation to farmers affected by elephant attacks, but even the forest department conducts community meetings to train locals to face the attack. Also, measures like appointing experts to tame the wild elephants with the help of domesticated elephants, and sensitising people towards wild life are most important part of such activities.



Do you know ?

The IUCN system uses a set of five quantitative criteria to assess the extinction risk of a given species. These criteria are : The rate of population decline; The geographic range; Whether the species already possesses a small population size; Whether the species is very small or lives in a restricted area; and Whether the results of a quantitative analysis indicate a high probability of extinction in the wild.

After a given species has been thoroughly evaluated, it is placed into one of following several categories.

1. **Extinct (EX)**, a designation applied to species in which the last individual has died or is not recorded.
2. **Extinct in the Wild (EW)**, a category containing those species whose members survive only in captivity
3. **Critically Endangered (CR)**, a category containing those species that possess an extremely high risk of extinction with very few surviving members (50).
4. **Endangered (EN)**, a designation applied to species that possess a very high risk of extinction as a result of rapid population decline of 50 to more than 70 percent over the previous 10 years (or three generations).
5. **Vulnerable (VU)**, a category containing those species that possess a very high risk of extinction as a result of rapid population decline of 30 to more than 50 percent over the previous 10 years (or three generations).
6. **Near Threatened (NT)**, a designation applied to species that are close to becoming threatened or may meet the criteria for threatened status in the near future.
7. **Least Concern (LC)**, a category containing species that are pervasive and abundant after careful assessment
8. **Data Deficient (DD)**, a condition applied to species in which the amount of available data related to its risk of extinction, is lacking in some way.
9. **Not Evaluated (NE)**, a category used to include any of the nearly 1.9 million species described by scientists, but not assessed by the IUCN



Can you tell?

1. Explain how loss of species diversity can harm ecosystem?
2. Give various categories of endangered species explained by IUCN.
3. What do you understand by invasive species? How does it affect local population?

15.5 Conservation of Biodiversity :

Conservation of biodiversity means protection, upliftment and scientific management of biodiversity to maintain its optimum level and also to derive sustainable benefits for the present and future strategies.

Why to conserve Diversity?

The reasons for conservation of biodiversity can be classified into three categories:

a. Narrowly utilitarian reasons:

Since time immemorial, humans are reaping material benefits from biodiversity. It may be deriving resources for basic needs such as food, clothes, shelter or industrial products like resins, tannins, perfume base etc. For aesthetic use as in ornaments or artefacts. Medicinal use of plants and animals, is another major factor. It shares 25% of global medicine market. Around 25000 species are put to use by tribals worldwide as traditional medicines. Several are yet to be explored for their potential as medicinal plants.

Now a days bioprospecting of economically important species is carried out. Bioprospecting is systematic search for development of new sources of chemical compounds, genes, micro-organisms, macro-organisms, and other valuable products from nature.

b. Broadly utilitarian reasons:

If we find out the cost of oxygen cylinder and try to calculate the value of oxygen we breathe with such ease; we will understand what nature is giving us for free! Animals play a crucial role in pollination and seed dispersal.

Amazon forest is estimated to produce 20% of total oxygen of earth's atmosphere. We need to consider recreational use of diversity too.

You must have come across the news about devastating fires in Amazon rainforest in August 2019.

These are mainly caused in Brazil and are more manmade than natural. The slash

and burn policy of locals to reclaim forestland has caused a towering 906000 hectares of forest devastation, only in the year 2019. We the humans, need to rethink about our attitude towards nature!

c. Ethical reasons:

We have no right to destroy the diversity simply because we share the earth with them! All living beings have equal right to survive irrespective of their known or prospective economic use.



Internet my friend

Visit www.gotul.org.in to find out information about various efforts for biodiversity conservation in Maharashtra.

How do we conserve biodiversity?

Conservation means sustainable use of natural resources. There are two main types of conservation strategies :

a. *In situ* conservation:

Protection of an organism will automatically take place, if its natural habitat is protected. e.g. Announcing Kanha forest as tiger reserve. This is called ***in situ* conservation**. This is the most appropriate method of conservation. It is nothing but conservation 'at home'. Around **34 Biodiversity hotspots** have been identified by the conservationists. These are the regions with high species richness as well as density. These areas need to be protected strategically by setting legislative measures apart from awareness and conservation.

In situ conservation also includes introduction of varieties traditionally used into farming and horticulture. E.g. In Maharashtra, Pawra tribals in Satpuda have protected varieties of corn with different coloured kernels.

India has four of world's biodiversity hotspots (the areas with high density of biodiversity), **Western ghats, Indo-Burma region, sundaland and Eastern-Himalayas**. It has been estimated that protection of these diversity rich hotspots could reduce extinction rate by almost 30%.

India, at present has 14 biosphere reserves, 90 national parks and 448 wildlife sanctuaries. In Maharashtra, there are 5 national parks and 11 sanctuaries.



Internet my friend

Collect the information about protected areas, biosphere reserves, national parks and wild life sanctuaries.



Can you recall?

What are sacred groves?

Indian culture and traditions are always connected with nature and rituals are laid down to protect biodiversity. In many cultures, stretches of forests were set aside and protected in the name of Almighty, which are called **sacred groves**.

Such sacred groves are found in Khasi and Jaintia hills in Meghalaya, Western ghat regions of Maharashtra and Karnataka, Aravali hills of Rajasthan and Bastar, and Chanda and Sarguja areas of Madhya Pradesh.



Do you know ?

Dr. Akira Miyawaki studied native forests of Japan especially the old shrine groves and developed a technique of growing dense plantations in short time. It is a technique for restoration of natural vegetation on degraded land. In this technique, after soil testing, the landmass is dugout and soil is mixed with local biomass and humus. Plantation is done in layers and saplings are planted close to each other. Due to this, sunlight doesn't reach soil retaining the moisture. Close proximity of plants leads to faster vertical growth than lateral. Also, it promotes natural selection. Native varieties are planted and the forest develops at almost ten times faster than the natural way. It requires a caring period of 3 years after which it grows and develops its own diversity naturally. In India, this technique is adapted at several places including remote districts like Chandrapur as well as metro cities like Mumbai and Bengaluru. Though there is debate about feasibility of the technique, it is certainly helping in retaining and recharging groundwater table, supporting local biodiversity and curbing air pollution by adding to green cover.

Sacred groves serve the only chance of survival for some endangered varieties of animal and plant species. Tribals do not allow to cut even a single branch of tree from sacred grove. But with the increasing lust and greed, are sacred groves safe? We must think about it.

b. *Ex situ* conservation:

Sometimes when a species is critically endangered, special measures have to be under taken to protect it. It might be protected in captivity, as one of the measures of protection. This is called *Ex situ* conservation. In this type of conservation, living beings are protected away from their natural habitats in special settings. Wild life safari parks, zoological parks and botanical gardens serve this purpose. Animals which have decreased in number, are allowed to breed in captivity in order to protect them. e.g. crocodile bank of Chennai.

Seed banks are established to conserve wild varieties of food grains and vegetables. Now a days, modern techniques like tissue culture, *in vitro* fertilization of eggs and cryopreservation (preservation at low temperature -196°C) of gametes, are used to protect endangered species.

By now we have, thus, understood the immense importance of biodiversity and dire need to protect it.



Internet my friend

Find out information about LACONES, Hyderabad.

15.6 Biological diversity Act 2002:

India participated in Earth Summit, Rio de Janeiro and is a party to Convention on Biological Diversity (CBD-1992).

In order to provide framework for the sustainable management and conservation of our country's natural resources, government passed Biological Diversity Act (BD Act) in the year 2002 in compliance with CBD. The law broadly defines biodiversity, as plants, animals and microorganisms and their parts, their genetic materials and by-products. It excludes value added products and human genetic material.

Regulation of access to Indian biological resources as well as scientific cataloguing of traditional knowledge about ethnobiological materials, were the main objectives for proposing this act.

A three-tier system has been established with **National Biodiversity Authority (NBA)** at the national level, the **State Biodiversity Boards (SBBs)** at the state level, and **Biodiversity Management Committees (BMCs)** at the local level for approval of utilization of any biological resource for commercial or research purpose. It is mandatory for foreigners, NRIs as well as Indian citizens and institutions to seek permission from NBA before exploiting local resource. NBA has powers of civil court. Not seeking approval of NBA, can incur jail and fine up to 10 lakh rupees.



Can you tell?

1. Differentiate between *in situ* and *ex situ* conservation.
2. Name any two modern methods of *ex situ* conservation of species.
3. Write a note on BD Act 2002.



Know the Conservationist



Rahibai Popere, seed mother of Maharashtra. Hailing from remote village in Ahmednagar district, Rahibai runs seed bank for 54 crops and 116 varieties. Crops include wild varieties of brinjal, guava, mango, spinach, methi, millets, pulses, hyacinth beans and peas. She also trains farmers and students for seed selection, enhancement of soil fertility, pest management and control. She is among 3 Indians on BBC list of '100 women, 2018'. She was awarded Padma Shri in 2020.



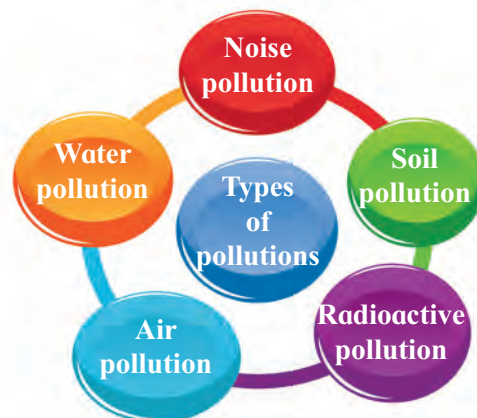
Can you recall?

1. What is pollution? Enlist its types.
2. Define pollutant. How are our daily activities responsible for pollution?

15.7 Environmental issues:

Exponential growth of human population coupled with industrial development, has resulted in the rampant loss of natural resources over last ten decades.

This uncontrolled exploitation of nature disturbed the delicate balance between living and non-living components of biosphere. Utilization and production of synthetic materials and construction activities have pumped several undesired substances in ecosphere. This has resulted in severe pollution.



Any substance - chemical or form of energy that adversely affect health of living organisms including humans, is called **pollutant**. In order to protect and improve the quality of our environment, the Government of India has passed the Environment Protection Act 1986.

a. Air Pollution :

Effect of air pollution :

Respiratory surfaces of living beings are constantly interacting with air. Any unfavourable alteration in air quality, affects the respiratory system. Severity of damage depends on concentration of pollutant, duration of exposure and the organism. Even in case of plants, air pollution results in poor yield of crops and premature death of plants. Nowadays automobiles are omnipresent. They are major cause for atmospheric (air) pollution. Regular maintenance of vehicles and use of lead-free petrol or diesel can reduce pollutant from exhausts.

Types of air pollutants :

Air pollutants are of two types – **particulate** pollutants and **gaseous** pollutants.

Particulate air pollutants are solids and also as fine liquid droplets called **aerosols**. Particles with diameter 10 μm may settle in the soil but particles with 1 μm or less remain suspended in the air. According to Central Pollution Control Board (CPCB), particulate matter of size 2.5 μm or less in diameter (PM2.5) are responsible for causing the greatest harm to humans.

These fine particulates can be inhaled deep into the lungs and are responsible for irritation, inflammation and damage to lungs. In addition to this, it causes breathing and respiratory disorders and premature deaths.

Smoke, smog, pesticides, heavy metals, dust and radioactive elements are the examples of particulate pollutants.



Use your brain power

Does particulate matter help to reduce atmospheric temperature?

Gaseous pollutants include CO_2 , CO , SO_2 , NO , NO_2 etc.

Carbon di-oxide is a greenhouse gas. In the past, levels of CO_2 in the atmosphere remained low. Due to burning of fossil fuels, as well as increasing deforestation the levels of CO_2 are rising at alarming rate. Photosynthesis process balances CO_2 : O_2 ratio of the air to a great extent. CO_2 is also removed from the air by weathering of silicate rocks forming limestone. A jet plane in a single trip across the Atlantic uses 35 tonnes of oxygen and releases 70 tonnes of CO_2 . We are going to discuss the role of CO_2 in global warming later in this chapter.

Carbon monoxide (CO) :

It is a poisonous gas produced by incomplete combustion of fuel such as coal or wood. Vehicular exhausts are the largest source of CO .



Use your brain power

State the effects of CO on human body.

Nitrogen di oxide (NO_2) and nitrogen monoxide (NO):

These are released by automobiles and chemical industries as waste gases. NO_2 when combines with water vapours forms nitric acid. It causes irritation to eyes and lungs. At high concentration, it causes injury to lungs, liver and kidneys.

Control measures:

Various ingenious mechanisms have been developed to control emission of gaseous and particulate pollutants through vehicles and industries. Few examples are explained below:

Electrostatic precipitator :

It is most widely used for removing particulate matter like soot and dust present in industrial exhaust. It can remove almost 99% particulate matter present in exhaust from a thermal power plant.

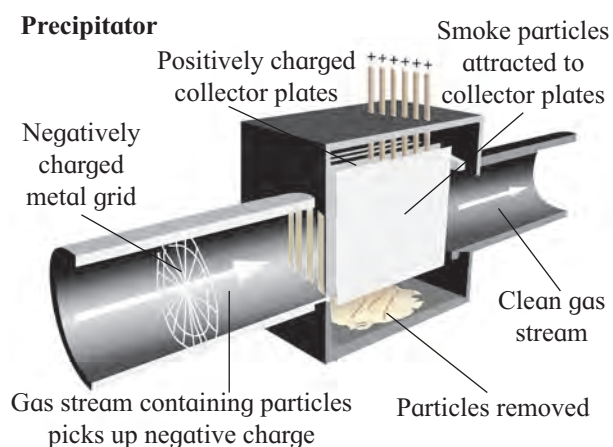


Fig. 15.4 : Electrostatic precipitator

In this method, high voltage is applied and electric discharge takes place which causes ionisation of air in the smokestack. Free electrons in the ionised air get attached to the gaseous/dust particles moving up the stack. These negatively charged particles move towards the positive electrode and settle down there. These particles are removed by vibrations of the electrodes and collected in the reservoir.



Internet my friend

Find out information about carbon foot print. Think about measures to reduce your own carbon foot print.

Exhaust gas Scrubbers are used to clean air for both dust and gases by passing it through dry or wet packing material. It can remove gases like SO_2 . In the scrubber, the exhaust is passed through a spray of water or lime.



Fig. 15.5 : Exhaust gas Scrubber

Catalytic converters:

Motor vehicles equipped with catalytic converter should use unleaded petrol because lead in the petrol, inactivates the catalyst.

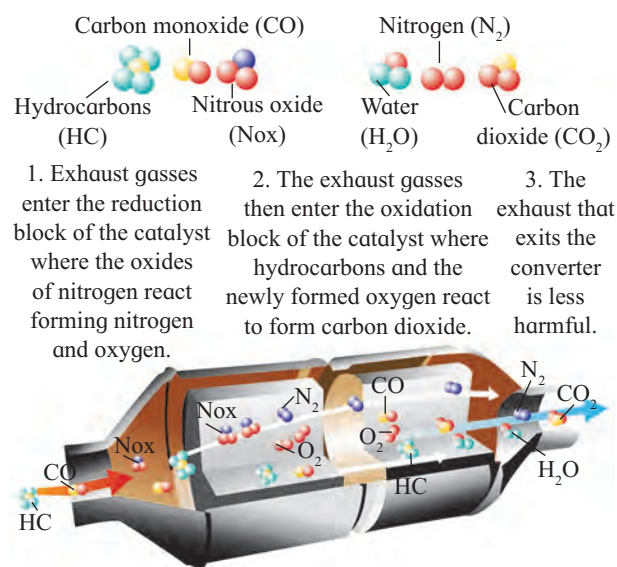


Fig. 15.6 : Catalytic converter

Controlling Vehicular Air Pollution : A case study of Delhi

In the year 1990, Delhi ranked fourth among 41 most polluted cities of the world. In response to PIL (Public Interest Litigation), Supreme court of India sent orders to Delhi government to take appropriate actions.

In response, several measures were taken by Delhi Government. By 2002, all the city buses were converted to run on CNG (**compressed natural gas**). CNG is advantageous over other fuels because it is economic, burns efficiently and is adulteration proof fuel.

According to new fuel policy, the norms are set to reduce sulphur and aromatic content of petrol and diesel. Another provision is upgradation of engines. For this, **Bharat stage emission standards (BS)** are set. These standards are equivalent to Euro norms and have evolved on similar lines as Bharat Stage II (BS II) to BS VI from 2001 to 2017. Let us observe how the norms have changed in the following table 15.7

Table 15.7 :Bharat stage emission standards in cities of India.

Vehicle	Norms	Cities of Implementation
4 wheelers	Bharat Stage II	All metro cities
4 wheelers	Bharat Stage III	Throughout the country since October 2010
4 wheelers	Bharat Stage IV	13 mega cities (Delhi and NCR, Mumbai, Kolkata, Chennai, Bengaluru, Surat, Kanpur, Agra, Lucknow, Solapur) since April 2010.
2 wheelers	Bharat Stage III	Throughout the country since October 2010
3 wheelers	Bharat Stage III	Throughout the country since October 2010



Do you know ?

Pradhan Mantri Ujjwala yojana (Pradhan Mantri clean fuel programme)

It was launched by Prime Minister of India, on 1st May 2016 to distribute 50 million LPG connections to women of BPL families. In many BPL families, chullhas are used where incomplete combustion of wood and coal leads to CO poisoning. Use of LPG helps in reducing such household air pollution.

Have you noticed that BS V is missing in above table? Note that, in 2001, Bharat stage II emission norms were set for CNG and LPG vehicles. It stipulates that emission of sulphur be controlled at 50 ppm in diesel and 150ppm in petrol. Aromatic hydrocarbons should be just 42% in concerned fuel. The aim is to reduce sulphur emission to 50ppm in petrol and diesel along with aromatic hydrocarbons to 35%. Government of India directly adapted BS VI in the year 2018, skipping BS V. These efforts decreased the levels of CO₂ and SO₂ in Delhi.



Do you know ?

Inspite of all the measures, New Delhi and adjoining areas witnessed great smog between 1st to 9th November 2016. Air pollution at this time peaked on both 2.5PM and 10PM levels. This is reported as one of the worst levels of air quality since 1999.

Colder weather and stagnant winds, trap smoke from various sources like firecrackers, burning crop stubbles, lit garbage and road dust. Citizens suffered breathlessness, chest muscle contraction, irritation in eyes, asthma and allergy. Administration took certain measures like closing educational institutions, suspending of construction or demolition work, undertaking vacuum cleaning of roads etc. Even Badarpur thermal power plant was temporarily closed down. Do you think mere setting standards is not enough? We must encourage means like car pooling and use of public transport.



Can you tell?

1. Describe any 2 particulate and gaseous pollutants.
2. Explain various technologies used for removing particulate matter from different sources of air pollution.
3. What are the ill effects of noise pollution on human health?
4. Give any norms for reducing sulphur and aromatic contents of petrol and diesel.

b. Noise pollution :

In India, the Air (Prevention and control of pollution) Act 1981, amendment 1987, includes **noise** as an air pollutant. Noise is an undesired high level of sound. Noise causes psychological and physiological changes in human beings. There is dire need of creating awareness about noise pollution caused during festivals and processions in our society.

Exposure to extremely high sound level (150 decibels or more) generated during a take-off of a jet plane or rocket, may damage ear drums and cause permanent hearing loss.

Noise also can cause sleeplessness, increased heartbeat, altered breathing pattern and psychological stress. Noise may negatively interfere with child's learning and behaviour pattern. The common sources of noise pollution are machines, transportation, construction sites, industries, Public functions, festivals, family functions etc.



Activity :

Find out different sources of noise pollution in your surrounding.

Reduction of noise in our industries can be brought about by use of sound absorbent materials or by muffling the noise. Laws which prohibit blowing horn in the areas of schools and hospitals, need to be implemented strictly to curb decibel levels.

Govt. of India has rules and regulations against firecrackers and loudspeakers. Supreme court of India has banned loudspeakers at public gatherings after 10pm.



Internet my friend

We have studied about health effects of noise pollution on humans. How does this pollution affect birds? Does it affect marine life? Find out.



Can you recall?

1. Define water pollution.
2. When will you call a water body polluted?

c. Water Pollution and its Control:

Water has been a controlling factor in distribution and density of living beings in an area. Human water consumption increased many folds with increased industrialisation.

Increased utilisation and discharge of harmful waste water in water bodies has caused severe pollution.

Most of the water pollution is manmade. Polluted water may be turbid, foul smelling, and may contain number of pathogens, heavy metals, oils etc.

Realising the importance of maintaining the cleanliness of the water bodies, Government of India has passed the **Water (prevention and control of pollution) Act 1974** to safe guard our water resources.



Can you recall?

1. Where does domestic waste water go in urban and rural areas?
2. What is importance of sewage treatment plant?

Domestic sewage and Industrial Effluents:

Even a small quantity of about 0.1% impurities in water, can make it unfit for human consumption and also affect aquatic plants and animals. Solids are relatively easy to separate but dissolved salts such as nitrates, phosphates, other nutrients and toxic metal ions as well as organic compounds, are difficult to remove.

Domestic sewage is one of the most common source of water pollution. It contains biodegradable organic matter. It readily gets decomposed by bacteria and other microorganisms. They use organic matter as substrate and utilise some amount of sewage.

It is possible to estimate biodegradable organic matter in sewage water by measuring **Biochemical Oxygen Demand (BOD)**. It is the amount of dissolved oxygen required by microorganisms for decomposing the organic matter present in water. It is expressed in milligram of oxygen per litre (mg/L) of water. High BOD indicates intense level of microbial pollution.

Microorganisms involved in biodegradation of organic matter in water body consume lot of dissolved oxygen. This results in sharp decline in oxygen level of water which leads to mortality of fish and other aquatic creatures.

Presence of large amount of nutrients in water causes excessive growth of **planktonic** free-floating algae specially, blue green algae. This is called **algal bloom** which gives colour to the water bodies. Algal bloom often releases toxins in water. So, fish die due to toxicity. Quality of water deteriorates and becomes toxic for human beings and other animals.

Another threat to aquatic ecosystem, is water hyacinth (*Eichhornia crassipes*). It is an aquatic plant, native of amazon basin, highly problematic invasive species. It was first introduced in India for its lovely purple coloured flowers. But, now it is a nuisance as it grows excessively and covers entire water body. This plant grows faster than our ability to remove it. It is commonly called 'Terror of Bengal'.

Natural Eutrophication is aging of a lake due to nutrient enrichment of water. Depending on the size of the lake, climatic conditions and other factors, natural aging of lakes require thousands of years. However, due to pollutants from human activities like effluents from agricultural lands, industries and homes (house hold) have accelerated this aging process. This phenomenon is called **Cultural or Accelerated Eutrophication**. Observe the flow chart and understand the process of eutrophication.

This results in non-availability of dissolved oxygen for aquatic organisms. This leads to death of fish and other aquatic organisms. Its decomposition further depletes the dissolved oxygen. So, a lake can literally get choked to death.

A few substances usually present in industrial waste waters can undergo biological magnification. The phenomenon through which

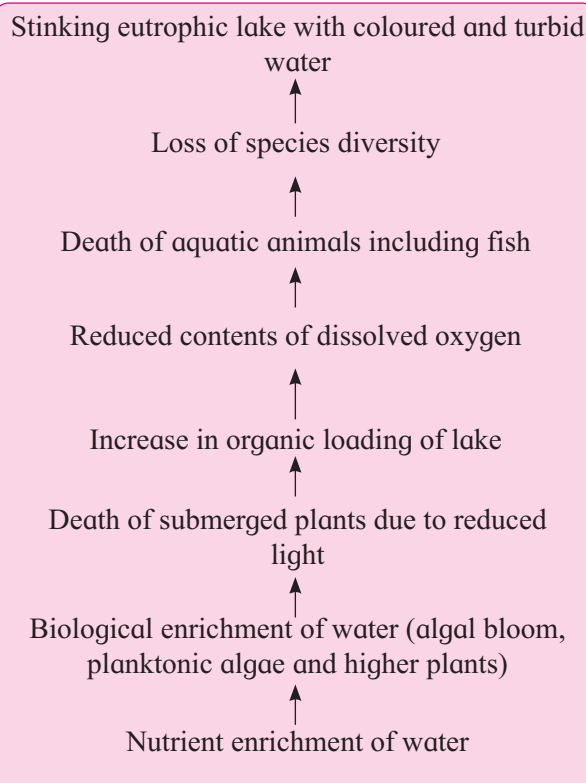


Fig. 15.8 : Eutrophication

certain pollutants get accumulated in tissues in increasing concentration along the food chains (successive trophic levels) is called **Biological Magnification (Biomagnification)**

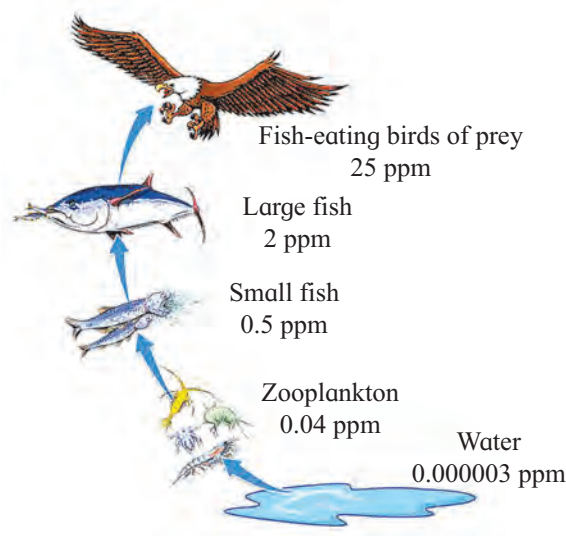


Fig. 15.9 : Biomagnification of DDT

Many of the pesticides are non-degradable. So, they get accumulated in the tissues of organisms. They neither get metabolised nor are excreted.

They are passed on to the next trophic level. This is commonly seen in case of DDT and Mercury. Observe the food chain in the given figure. It shows how biomagnification of DDT takes place.



Use your brain power

Why do you think the amount of DDT is maximum in birds?

Thermal pollution of water is caused due to rise in temperature of water. The main source of thermal pollution are the thermal and nuclear power plants. The power plants use water as coolant and release hot water. As many organisms are sensitive to temperature, sudden rise in temperature leads to loss of flora and fauna.



Internet my friend

1. Find out biomagnification of diclofenac and DDT in birds.
2. Find out various types of agrochemicals which have ill effects on ecosystem.

Measures to reduce sewage water:

In order to conserve water and prevent creation of sewage, ecosan is a sustainable system for handling human excreta using dry composting toilets. This is a practical, efficient and cost-effective solution for human waste disposal.

Ecological sanitation (Ecosan) is an approach to sanitation provision which safely reuses excreta in agriculture. It reduces the need for chemical fertilizers.

Ecosan toilet is a closed system that does not need water. It is an alternative to leach pit toilets in place where water is scarce or where there is risk of ground water contamination.

It is based on the principle of recovery and recycling of nutrients from excreta to create a valuable resource for agriculture.

When the pit of an ecosan toilet fills up, it is closed and sealed. After about 8-9 months, the faeces are completely composted to organic manure. There are working EcoSan toilets in many areas of Gujrat, Kerala, Tamilnadu and Sri Lanka.

Recycling of sewage water by reverse osmosis will help to solve the problem of not only scarcity of water but also the disposal of sewage water. At Tirumala hills, millions of pilgrims visit. Every building here has R.O. system which solves the problem of huge water demand. Recycling of sewage water is seen in many townships in Mumbai. Rain water harvesting is encouraged and made mandatory for new constructions by Municipal Corporation.



Can you tell?

1. Explain BOD and its effects on aquatic ecosystem.
2. Ecological sanitation is the need of the day. Justify.
3. Explain the phenomenon of biological magnification.

Solid Waste Management :

You have already studied about solid waste management in ninth standard.



Can you recall?

1. Why is it necessary to separate wet and dry garbage?
2. What are the six R's to combat solid waste?

Solid waste is everything that goes to trash. It includes wastes from home, offices, stores, schools, hospitals etc. Wastes are collected and disposed by municipality. Municipal solid waste generally consists of paper, food, plastic, glass, metals, rubber, leather, textile etc;

Burning reduces volume of the waste. Generally, it is not burnt to completion. So open dumps become breeding ground for rats and flies.

Sanitary landfills are substitute for open burning dumps. Here wastes are dumped in depression or trench. Everyday wastes are added to this pit. Landfills get filled very soon especially in metro cities. The amount of garbage generated is too high. In addition to this, there is a danger of chemicals percolating and reaching down to ground water and contaminate this water source.

All wastes can be categorised in three types-

1. Bio degradable
2. Recyclable
3. Non bio degradable

Our rag pickers and *kabadiwallahs* do a great job of separation of materials for recycling. Primary goal of all citizens should be to reduce generation of waste.

The biodegradable materials can be put into deep pits in the ground and left for natural degradation. Non biodegradable wastes have to be disposed off.

Maharashtra government in a notification on 23rd June 2018 banned use of plastic to fight pollution caused due to extensive use of plastic. Ban on the use, sale, distribution and storage of plastic material.

Mission-Plastic Free Maharashtra :

The union environment ministry has amended rules in 2016 in order to strengthen the implementation of environmentally sound management of hazardous waste in the country.

It also includes ban on use of Thermocol.

Large amount of use of plastic can be avoided, if we inculcate the habit of carrying cloth bags when we go for shopping and strictly refusing plastic bags from the vendors. Government should charge high penalty/fine for the individuals not abiding by this rule.

Hospitals generate harmful waste that contains disinfectants, harmful chemicals and also pathogenic microorganisms.

Such wastes require careful treatment and disposal. You are aware of colour code for disposal of biomedical waste.

Irreparable computers and other electronic goods as well as electrical waste are known as **e-wastes**. They are buried in landfills or are completely burnt. Major part of e-waste generated by developed countries are exported to developing countries like China, India and Pakistan. During recycling process of this waste, metals like copper, iron, silicon, nickel and gold are recovered. Developed countries have facilities for recycling of e-waste. In developing countries, manual participation is involved. So, workers are exposed to toxic substances from e-waste. Treatment of e-waste should be carried out in environment friendly manner.

You have already studied about radioactive pollution.



Can you recall?

1. What is greenhouse effect?
2. Enlist greenhouse gases.
3. How do you correlate green house effect and global warming?

15.8 Greenhouse effect and Global warming:

Greenhouse effect : It is responsible for heating of earth's surface and atmosphere. Without greenhouse effect, the average temperature of Earth would have been -18°C rather than average of 15°C .

Of the solar radiation that reaches earth, Clouds and gases reflect about $1/4^{\text{th}}$ and absorb some of it. But half of total incoming radiations reach the earth's surface and heat it. Small portion of it, is reflected back. Earth's surface re emits the heat in the form of infrared radiations. Part of these radiations do not escape into the space because atmospheric gases (CO_2 , CH_4) absorb a major portion.

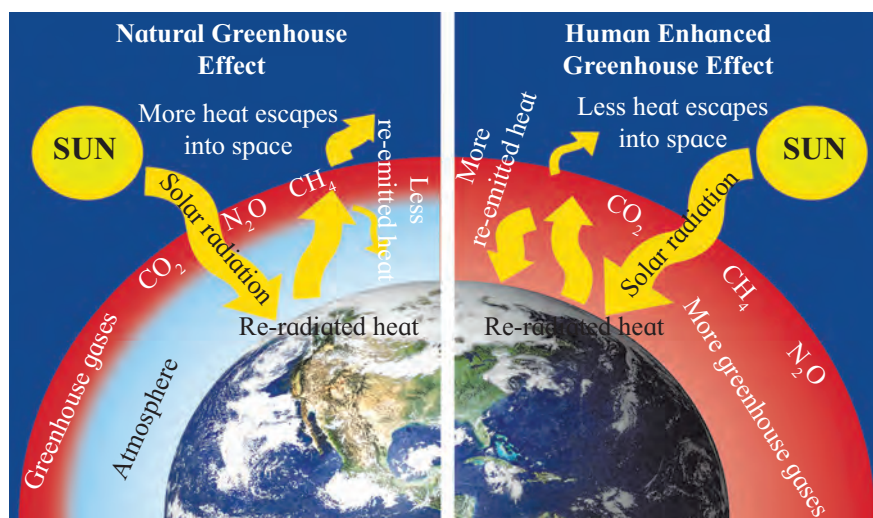


Fig. 15.10 : Greenhouse effect

The molecules of these gases radiate heat energy and a major part of it comes back to earth's surface. Thus reheating the earth. This cycle is repeated many times. Similar phenomenon prevails under green house/glass house. Hence, it is called **green house effect**. Gases CO₂ and CH₄ involved in the process are called **green house gases**.

The atmosphere around the earth acts as glass wall of a greenhouse. It absorbs solar radiations from the sun and radiates it to earth. Atmosphere prevents infrared radiations emitted by the earth to escape into the space.

Gases responsible for this effect are carbon di-oxide (CO₂), Methane (CH₄), chloroflorocarbons (CFC), Nitrous oxide (N₂O) and water vapours.

Because of burning of fossil fuels in industries, by automobiles, burning of agricultural wastes, levels of CO₂ are increasing. Biogas plants, paddy fields, cattle sheds add methane to atmosphere. Chloroflorocarbons are emitted by fire extinguishers and air conditioners.

Global warming :

Increase in atmospheric concentration of green house gases, has resulted in the heating of Earth or rise in the temperature. During past century, the temperature of the Earth has increased by 0.6°C, most of it during last three decades.

This rise in temperature leads to unfavourable changes in environment and resulting in odd climatic changes.(eg; El Nino effect). El Nino effect results in melting of polar ice caps and Himalayan snow caps which may be the cause for submerging of the coastal areas.



Do you know ?

In order to overcome the problem of global warming, **Chewang Norphel, Ice Man of India** has built 13 artificial glaciers. He is an Indian civil engineer from Ladakh.

Norphel noticed a small stream had frozen solid, under the shade of a group of poplar trees, though it flowed freely elsewhere in his yard.

The reason for this phenomenon :

The flowing water was moving quickly to freeze while the sluggish trickle of water beneath the trees, was slow enough to freeze. Based on this, he created artificial glaciers by diverting a river into a valley, slowing the stream by constructing checks.

The artificial glaciers increase the ground water recharge, rejuvenating the spring and providing water for irrigation. He constructed them at lower elevations, so that they melt earlier expanding the growing season.

Global warming can be controlled by reducing use of fossil fuel, improving efficiency of energy usage, reducing deforestation, planting trees and checking of human population growth.



Do you know ?

El Nino effect is a climate cycle in Pacific ocean with global impact on weather pattern. The cycle begins when warm water in the western tropical Pacific ocean shifts eastwards along the equator towards the coast of America. Normally, this warm water pools near Indonesia and Philippines.

La Nina is a climatic pattern that describes the cooling of surface ocean waters along the tropical west coast of South America.

15.9 Ozone depletion :

Ozone is a form of oxygen. In the stratosphere, ozone is photo-dissociated and is generated by absorption of short wave length UV radiations.



Generation and dissociation of ozone are in equilibrium leading to steady concentration of ozone in the stratosphere (12 to 15 kilometers from Earth's surface in the atmosphere).

This ozone layer acts as shield that absorbs UV radiations from the sun and protects all types of life on earth. Absorption of UV radiations by ozone blanket is proportional to its thickness. Thickness is more above the poles than at the equator. UV rays are highly injurious to living organisms since DNA and proteins of living organisms absorb UV rays and its high energy breaks the chemical bonds within these molecules.

Thickness of the ozone in a column of air from the ground to the top of the atmosphere is measured as **Dobson units** (DU).

The ozone shield has been disturbed due to increased rate of ozone degradation by **Chlorofluorocarbon** (CFC). CFCs move upwards and reach stratosphere. UV rays act on them and release Cl atoms. Cl degrades ozone releasing molecular oxygen. Cl atoms act as catalyst. So they remain in the stratosphere and continue the effect of ozone degradation. This results in ozone depletion. Although it occurs widely in the stratosphere, the depletion is particularly marked over the Antarctic region. This has resulted in formation of large area of thinned ozone layer, commonly called **ozone hole**.

UV radiations of wave length shorter than UV-B i.e. 100-280nm are almost completely absorbed by earth's atmosphere, given that the ozone layer is intact. UV-B (wavelength 280-322nm) damages DNA and mutation may occur. It causes aging of skin, damage to skin cells and various types of skin cancers. In human eye, cornea absorbs UV-B radiations and a high dose of UV-B causes inflammation of cornea called **snow blindness**, cataract etc; Such exposure may permanently damage cornea.

Recognising the harmful effects of ozone depletion, an international treaty, known as the **Montreal Protocol** was signed at Montreal (Canada) in 1987 to control emission of ozone depleting substances.

Later many more efforts have been made and protocols have laid down definite roadmaps separately for developing and developed countries for reducing emission of CFCs and other ozone depleting chemicals.



Can you tell?

1. How pollution by domestic garbage can be controlled?
2. Give effect of CFC on ozone shield.
3. Give an account of possible effects of global warming.
4. Plastic ban in Maharashtra is an essential step. Give reason.



Internet my friend

1. Find out more about artificial glaciers.
2. Collect information about colour code used for biomedical waste.

15.10 Deforestation :

Deforestation is conversion of forest area into non-forest area. According to an estimate, almost 40% forests are lost in the tropics and only 1% in temperate region. The scenario of deforestation is grim in India. At the beginning of 20th century, forest cover was about 30% in India. By the end of the century, it got reduced to 19.4 %. The National Forest Policy 1988 of India has recommended 33% forest cover for the plains and 67% for the hills.

Deforestation takes place by conversion of forest to agricultural land so as to feed the growing human population. Trees are cut for timber, firewood, for keeping cattle in farm and for other purposes.

Slash and burn agriculture commonly called **Jhum Cultivation** in north eastern parts of India, where farmers cut down trees of the forest and burn the plant remains. The ash is used as fertilizer and the land is used for farming and cattle grazing. After cultivation, the area is left for several years so as to allow its recovery.

Major Effect of Deforestation is the increased concentration of CO₂ in the atmosphere because trees hold lot of carbon in their biomass, are lost with deforestation.

It leads to the loss of biodiversity due to habitat destruction, disturbs hydrologic (= hydrological) cycle, causes soil erosion and desertification in extreme cases.



Can you tell?

1. How Jhum cultivation has led to deforestation in recent years?
2. Comment on deforestation status of the world and its major effects.

Reforestation is the process of restoring a forest that once existed but was destroyed or removed at some time in past. Reforestation occurs naturally in a deforested area. We can speed up this by plantation of tree with due consideration to biodiversity that existed before.

Best example of peoples participation in reforestation is **Saalumara Thimmakka**, an Indian environmentalist from state of Karnataka noted for her work in planting and tending to 385 banyan trees along a 4km stretch of highway between Hulikal and Kudur. She has also planted nearly 8000 other trees. Her work has been honoured with the National Citizens Award of India. She was also conferred with Padma Shri in 2019.

Moirangthem Loiya from Manipur dedicated 17 years of his life to restore Punshilok forest. He left his job and took over the task of bringing back the lost glory of 300 acres forest land. He planted a variety of trees like, bamboo, oak, ficus, teak, jackfruit and magnolia. Today the forest has over 250 varieties of plants including 25 varieties of bamboo. It is now selected as home by great diversity of animals too.



Can you recall?

What is Chipko movement?

Case study of people's participation in conservation of forests.

History of people's participation in India can be traced back to 1731. The ministers of the king of Jodhpur in Rajasthan tried to cut forest to procure wood for a new palace. Local Bishnoi community known for peaceful co-existence with nature, opposed king's men.

A Bishnoi woman **Amrita Devi** hugged the trees and lost her life in an attempt for protecting the forest. Her three daughters and hundreds of other Bishnois too lost their lives.

The Government of India has recently instituted **Amrita Devi Bishnoi Wildlife protection Award for** individuals or community from rural areas that have shown extraordinary courage and dedication in protecting wildlife.

In 1980s, realising the significance of participation of local communities, Government of India introduced the concept of **Joint Forest Management (JFM)** so as to work closely with local communities for protecting and managing forests. In return, for their services to the forest, the communities get benefit of various forest products (Fruits, gum, rubber, medicine etc.) and thus the forest can be conserved in a sustainable manner.

15.11 Mission Harit Maharashtra :

An ambitious project of planting 50 crore trees in four years was taken up by Government of Maharashtra in the year 2016. Under this project yearly targets were given to each district. The plantation drive is in line with National Forest Policy (NFP) which aims at maintaining 33% forest cover in the country. A 24-hour toll free helpline number 1926 called 'Hello Forest' has been set up to provide information regarding plantation, protection and for mass awareness.

The Forest Department has created a mobile application called 'My Plants' to record details of the plantation such as numbers, species and location. Authorities are expected to take care of saplings in the first year i.e., year 2016, 2.87 crore saplings were planted. In 2017, 5.17 crore and in 2018, 15.17 crore plantation count was achieved! In the year 2019, the government aimed at a phenomenal 33 crore sapling plantation which was launched at Anandwan, Warora.

The government has decided to adapt Japanese Miyawaki method of plantation for this project. State Forest Department and Social Forestry Department have run successful pilot plantation programmes using Miyawaki pattern in various districts like, Beed, Hingoli, Pune, Jalgaon, Aurangabad etc.



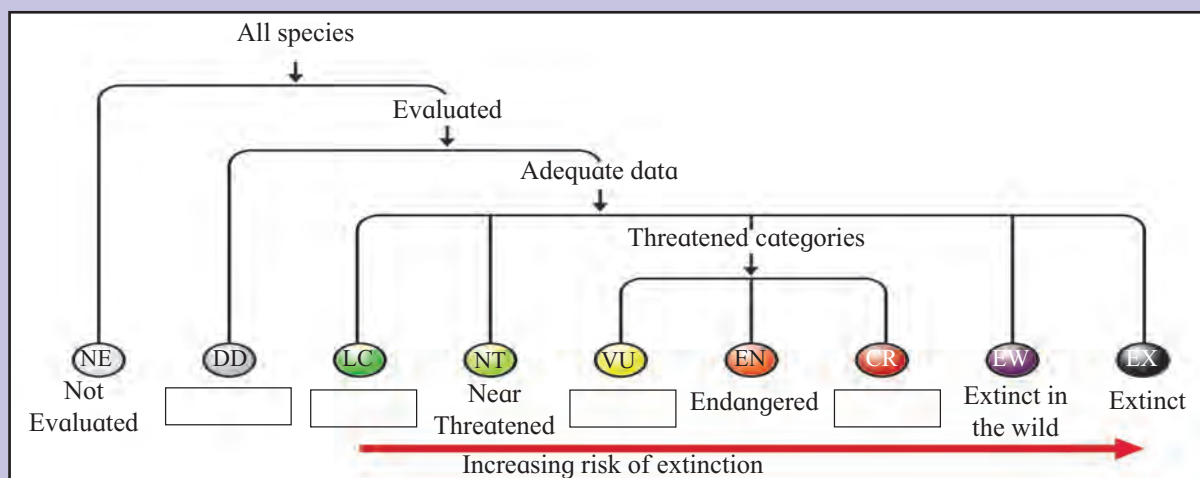
Use your brain power

Floods in Sangli and Kolhapur in August 2019, were responsible for many problems during and after the floods. Think and enlist different types of problems faced by flood affected areas.



Activity :

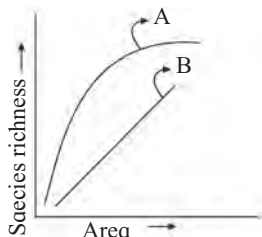
Complete the following chart :



Exercise

Q. 1 Choose the correct option

- Observe the graph and select correct option.



- Line A represents, $S=CA^2$
 - Line B represents, $\log C= \log A + Z \log S$
 - Line A represents, $S=CA^Z$
 - Line B represents, $\log S= \log Z + C \log A$
- Select odd one out on the basis of *Ex situ* conservation.
 - Zoological park
 - Tissue culture
 - Sacred groves
 - Cryopreservation
 - Which of the following factors will favour species diversity?
 - Invasive species
 - Glaciation
 - Forest canopy
 - Coextinction
 - The term “terror of Bengal’ is used for _____.
 - algal bloom
 - water hyacinth
 - increased BOD
 - eutrophication
 - CFC are air polluting agents which are produced by _____.
 - Diesel trucks
 - Jet planes
 - Rice fields
 - Industries

Q. 2 Very short answer type questions.

- Give two examples of biodegradable materials released from sugar industry.
- Name any 2 modern techniques of

protection of endangered species.

- Where was ozone hole discovered?
- Give one example of natural pollutant.
- What do you understand by EW category of living being?

Q. 3 Short answer type questions.

- Dandiya raas* is not allowed after 10.00 pm. Why?
- Tropical regions exhibit species richness as compared to polar regions. Justify.
- How does genetic diversity affect sustenance of a species?
- Is green house effect boon or bane? Give your opinion.
- How does CO cause giddiness and exhaustion?
- Name two types of particulate pollutants found in air. Add a note on ill effects of the same on human health.

Q. 4 Long answer type questions.

- Montreal protocol is an essential step. Why is it so?
- Name any 2 personalities who have contributed to control deforestation in our country. Elaborate on importance of their work.
- How BS emission standards changed over time? Why is it essential?
- During large public gatherings like *Pandharpur vari* mobile toilets are deployed by the government. Explain how this organic waste is disposed.
- How Indian culture and traditions helped in bio-diversity conservation? Give importance of conservation in terms of utilitarian reasons.

Project : Find out at least 2 plant and animal varieties native to Maharashtra which are endangered. Find out their IUCN status and reasons for the same. Suggest conservation measures.

Notes

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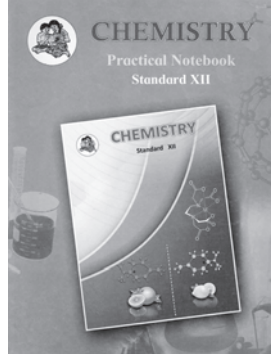
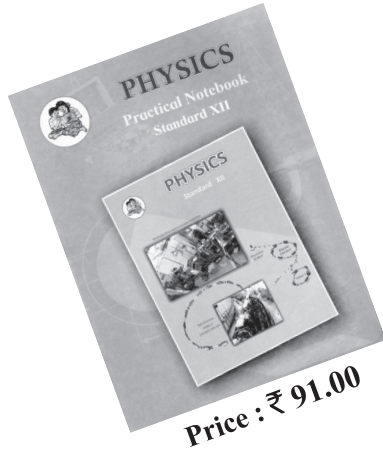


Notes

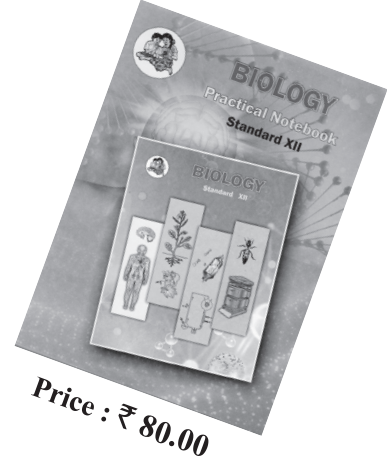
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