









Essentials of

Nutrition and Biochemistry

for BSc Nursing Students

As per the syllabus of INC for BSc Nursing

Special Features

- Thoroughly Updated and Revised Edition
- Perfect Amalgamation of Theoretical and Clinical Aspects
- 30+ Clinical Correlation Boxes
- 300+ Figures, Tables and Flowcharts
- A Special Chapter on Sample Collection and Normal Values of Biochemical Parameters
- Covers an Exclusive Section on Nutrition for the first time

3rd Edition





CBS Publishers & Distributors Pvt. Ltd.





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Third Edition

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Preface to Third Edition

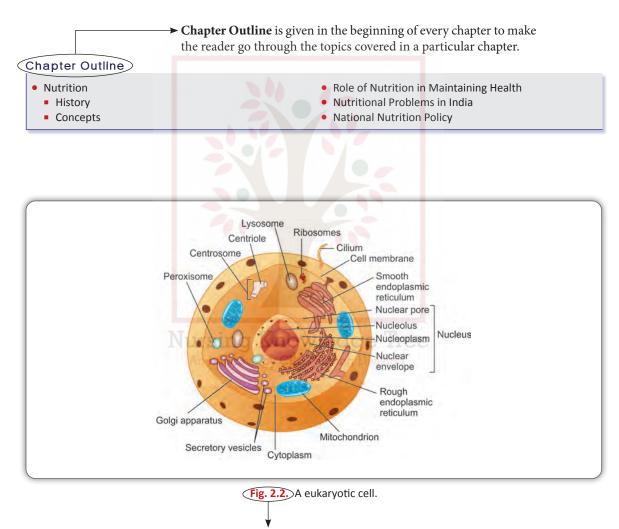
Lots of encouragement and support of the faculty members and the students of several Nursing Colleges motived and inspired me to write this book. Extraordinary success of my books entitled "Food & Nutrition" and "Essentials of Biochemistry for BSc Nursing Students" encouraged me to present the new edition of Essentials of Nutrition and Biochemistry for BSc Nursing Students.

A major goal is to provide the basics of Nutrition as well as Biochemistry to the BSc Nursing students in a concise and interesting manner. All the chapters which have been divided under various Units have been written in a lucid manner. The text in each chapter has been supplemented with suitable tables, outlined flowcharts and figures. Some important questions have also been included at the end of each chapter under the segment Assess Yourself.

HARBANS LAL

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Special Features of the Book



Studded with 200+ fully colored Images and Illustrations for easy grasp of the relevant topics.

Numerous Tables are used to clarify the concepts and make the reading enjoyable and informative.

TABLE 7.4: Biological importance of proteins

Biological Importance of protein	Examples
As structural proteins	Collagen, keratins, etc.
As enzymes	Pepsin, amylase, etc.
As hormones	Insulin
As transporters	Hemoglobin
As receptors	Hormone receptors
As storage molecules	Ferritin
In immune response	γ-globulin

A number of Clinical Correlation boxes have been included in-between the text from the clinical point of view.

Clinical Correlation

Zellweger Syndrome

Absence of the functional peroxisomes results in a rare autosomal recessive disease referred to as Zellweger syndrome. It is characterized by abnormalities of the several organs due to the decreased levels of plasmalogens.



 At the end of every chapter Assess Yourself covering subjective and objective Qs provided for self assessment.

Long and Short Answer Questions

- Define carbohydrate. Classify carbohydrates.
- 2. Write short notes on:

 - b. Homopolysaccharides Knowledge Tree

Multiple Choice Questions

- 1. Which of the following is a non-reducing sugar?
 - a. Sucrose c. Glucose

b. Maltose

- d. Lactose
- 2. Sucrose has glucose and
 - a. Maltose

b. Fructose

c. Glycogen

d. Galactose

Syllabus

NUTRITION

Placement: First Year Time: Theory 60 hours

Course description: The Course is designed to assist the students to acquire knowledge of nutrition for maintenance of optimum health at different stages of life and its application for practice of nursing.

Unit	Time	(Hrs)	Learning	Content	Teaching Learning	Evaluation
	Th.	Pr.	Objectives		Activities	
I	4		Describe the relationship between nutrition and health Nurs: An In	Introduction Nutrition: History Concepts Role of nutrition in maintaining health Nutritional problems in India National nutritional policy Factors affecting food and nutrition: socioeconomic, cultural, tradition, production, system of distribution, life style and food habits, etc. Role of food and its medicinal value Classifications of foods Food standards Elements of nutrition: macro and micro Calorie, BMR	 Lecture discussion Explaining using charts Panel discussion 	Short answersObjective type
II	2		 Describe the classification, functions, sources and recommended daily allowances (RDA) of carbohydrates 	Carbohydrates Classification Caloric value Recommended daily allowances Dietary sources Functions Digestion, absorption and storage, metabolism of carbohydrates Malnutrition: Deficiencies and over consumption	 Lecture discussion Explaining using charts 	Short answersObjective type

Unit	Time	(Hrs)	Learning Objectives	Content	Teaching Learning Activities	Evaluation
	Th.	Pr.	Objectives		Activities	
111	2		Describe the classification, functions, sources and recommended daily allowances (RDA) of fats	 Fats Classification Caloric value Recommended daily allowances Dietary sources Functions Digestion, absorption and storage, metabolism Malnutrition: Deficiencies and over consumption 	 Lecture discussion Explaining using charts 	Short answersObjective type
IV	2		 Describe the classification, functions, sources and recommended daily allowances (RDA) of proteins 	Proteins Classification Caloric value Recommended daily allowances Dietary sources Functions Digestion, absorption, metabolism and storage Malnutrition: Deficiencies and over consumption	 Lecture discussion Explaining using charts 	Short answersObjective type
V	3		Describe the daily calorie requirement for different categories of people An In	 Energy Unit of energy–Kcal Energy requirements of different categories of people Measurements of energy Body mass index (BMI) and basic metabolism Basal metabolic rate (BMR)– determination and factors affecting 	 Lecture discussion Explaining using charts Exercise Demonstration 	Short answersObjective type
VI	4		 Describe the classification, functions, sources and recommended daily allowances (RDA) of vitamins 	Vitamins Classification Recommended daily allowances Dietary sources Functions Absorption, synthesis, metabolism storage and excretion Deficiencies Hypervitaminosis	 Lecture discussion Explaining using charts 	Short answersObjective type

Unit	Time	(Hrs)	Learning	Content	Teaching Learning	Evaluation
	Th.	Pr.	Objectives		Activities	
VII	4		 Describe the classification, functions, sources and recommended daily allowances (RDA) of minerals 	Minerals Classification Recommended daily allowances Dietary sources Functions Absorption, synthesis, metabolism storage and excretion Deficiencies Over consumption and toxicity	 Lecture discussion Explaining using charts 	Short answersObjective type
VIII	3		Describe the sources, functions and requirements of water and electrolytes	 Water and Electrolytes Water: Daily requirement, regulation of water metabolism, distribution of body water, Electrolytes: Types, sources, composition of body fluids Maintenance of fluid and electrolyte balance Over hydration, dehydration and water intoxication Electrolyte imbalances 	 Lecture discussion Explaining using charts 	Short answersObjective type
IX	5	15	 Describe the cookery rules and preservation of nutrients Prepare and serve simple beverages and different types of foods 	Cookery Rules and Preservation of Nutrients Principles, methods of cooking and serving Preservation of nutrients Safe food handling-toxicity Storage of food Food preservation, food additives and its principles Prevention of Food Adulteration Act (PFA) Food standards Preparation of simple beverages and different types of food		 Short answers Objective type Assessment of practice sessions
х	7	5	 Describe the balanced diet for different categories of people 	Balanced Diet Elements Food groups Recommended daily allowance Nutritive value of foods Calculation of balanced diet for different categories of people Planning menu Budgeting of food Introduction to therapeutic diets: naturopathy—diet	 Lecture discussion Explaining using charts Practice session Meal planning 	Exercise on menu planning

Unit	Time	(Hrs)		Content	Teaching Learning Activities	Evaluation
	Th.	Pr.	Objectives		Activities	
XI	4		 Describe various national programs related to nutrition Describe the role of nurse in assessment of nutritional status and nutrition eduction 	 Role of Nurse in Nutritional Programs National programs related to nutrition Vitamin A deficiency program National iodine deficiency disorders (IDD) program Midday meal program Integrated child development scheme (ICDS) National and International agencies working towards food/nutrition NIPCCD, CARE, FAO, NIN, CFTRI (Central food technology and research institute) etc. Assessment of nutritional status Nutrition educational and role of nurse 	 Lecture discussion Explaining with slide/films shows Demonstration of assessment of nutritional status 	Short answersObjective type

Nursing Knowledge Tree

BIOCHEMISTRY

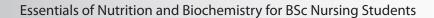
Placement: First Year Time: Theory-30 Hours

Course descriptions: The Course is designed to assist the students to acquire knowledge of the normal biochemical composition and functioning of human body and understand the alterations in biochemistry in diseases for practice of nursing.

Unit	Time	Learning	Content	Teaching Learning	Assessment Methods
	(Hrs)	Objective		Activities	
ı	3	 Describe the structure, composition and functions of cell Differentiate between prokaryote and eukaryote cell Identify techniques of microscopy 	Introduction Definition and significance in nursing Review of structure, composition and functions of cell Prokaryote and eukaryote cell organization Microscopy	 Lecture discussion using charts, slides Demonstrate use of microscope 	 Short answer questions Objective type
II	6	Describe the structure and function of cell Nurs An I	Structure and Functions of Cell Membrane • Fluid mosaic model tight junction, cytoskeleton • Transport mechanism: diffusion, osmosis, filtration, active channel, sodium pump • Acid base balance- maintenance and diagnostic tests • PH buffers	• Lecture discussion	 Short answer questions Objective type
Ш	6	Explain the metabolism of carbohydrates	Composition and Metabolism of Carbohydrates • Types, structure, composition and uses • Monosaccharides, disaccharides, polysaccharides, oligosaccharides	 Lecture discussion Demonstration of blood glucose monitoring 	Short answer questionsObjective type

Unit	Time (Hrs)	Learning Objective	Content	Teaching Learning Activities	Assessment Methods
			 Metabolism Pathways of glucose: Glycolysis Gluconeogenesis: Cori's cycle, tricarboxylic acid (TCA) cycle Pentose phosphate pathways (Hexose mono phosphate) Regulation of blood glucose level Investigations and their interpretations 		
IV	4	Explain the metabolism of lipids Nurs An I	Composition and Metabolism of Lipids Types, structure, composition and uses of fatty acids Nomenclature, roles and prostaglandins Metabolism of fatty acid Breakdown Synthesis Metabolism of triacylglycerols Cholesterol metabolism Biosynthesis and its regulation Bile salts and bilirubin Vitamin D Steroid hormones Lipoproteins and their functions: VLDLs-IDLs, LDs and HDSs Transport of lipids Athrosclerosis, Investigations and their interpretations	 Lecture discussion using charts Demonstration of laboratory tests 	 Short answer questions Objective type

Unit	Time (Hrs)	Learning Objective	Content	Teaching Learning Activities	Assessment Methods
V	6		Composition and Metabolism of Amino Acids and Proteins Types, structure, composition and uses of amino acids and proteins Metabolism of amino acids and proteins Protein synthesis, targeting and glycosylation Chromatography Electrophoresis Sequencing Metabolism of nitrogen Fixation and assimilation Urea cycle Hemes and chlorophylls Enzymes and co- enzymes Classification Properties Kinetics and inhibition Control Investigations and their interpretations		Short answer questions Objective type
VI	2	Describe types, composition and utilization of vitamins and minerals	Composition of Vitamins and Minerals Vitamins and minerals: Structure Classification Properties Absorption Storage and transportation Normal concentration Investigations and their interpretations	 Lecture discussion using charts Demonstration of laboratory tests 	 Short answer questions Objective type



vvii	١
VVII	1

Unit	Time (Hrs)	Learning Objective	Content	Teaching Learning Activities	Assessment Methods
VII	3	Describe immunochemistry	 Immunochemistry Immune response, Structure and classification of immunoglobins Mechanism of antibody production Antigens: HLA typing. Free radical and antioxidants Specialised protein: collagen, elastin, keratin, myosin, lens protein. Electrophoretic and quantitative determination of immunoglobins—ELISA etc. Investigations and their interpretations 	Lecture discussion Demonstrate laboratory tests	 Short answer questions Objective type

Nursing Knowledge Tree



Carbohydrates

LEARNING OBJECTIVE

After going through this unit, you will be able to:

Describe the classification, functions, sources and recommended daily allowances (RDA) of carbohydrates

UNIT OUTLINE

Chapter 2 Carbohydrates and their Significance

Carbohydrates and their Significance

Chapter Outline

- Carbohydrates
- Classification
- Calorie Value
- Recommended Daily Allowances
- Dietary Sources
- Functions

- Digestion, Absorption and Storage
- Metabolism
- Malnutrition
 - Deficiencies
 - Overconsumption

Carbohydrates are macronutrients and are one of the three main sources of energy. They are called carbohydrates, because at the chemical level, they contain carbon, hydrogen and oxygen, where the last two are present in the ratio of 2:1. For example, glucose $(C_6H_{12}O_6)$ contains 6 carbons, 12 hydrogens and 6 oxygen $(CH_2O)_6$.

CLASSIFICATION OF CARBOHYDRATES

Carbohydrates can be classified into different types, according to the size of the molecules and their structure as well as according to their nutritional significance.

According to the Size of the Molecule and their Structure

According to the size of the molecule and their structure, carbohydrates are classified into four groups, i.e., as monosaccharide, disaccharide, oligosaccharides and polysaccharide.

- Monosaccharides: Monosaccharides are also called simple sugars. Common monosaccharides include
 glucose, galactose and fructose. They share the same molecular formula, i.e., as C₆H₁₂O₆. Because of their
 six carbon atoms, each is a hexose:
 - The most important being glucose, as blood sugar. It is the immediate source of energy for cellular respiration.
 - Galactose, a sugar in milk (and yogurt)
 - Fructose, a sugar found in honey
 Monosaccharides, in a closed-chain form, can form glycosidic bonds with other monosaccharides,
 creating disaccharides (such as sucrose) and polysaccharides (such as starch). Monosaccharides may

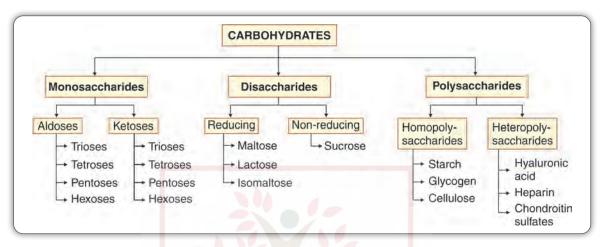


Fig. 2.1: Classification of carbohydrates

be further converted into structural polysaccharides such as cellulose and pectin for cell wall construction, or into energy reserves in the form of storage polysaccharides, such as starch and glycogen.

- **Disaccharides:** Disaccharides are formed by linking two monosaccharides with the removal of a molecule of water (Fig. 2.1). Commonly available disaccharides include lactose, maltose and sucrose.
 - Lactose is formed when a molecule of glucose is linked to galactose. It is also called milk sugar, since it is found naturally in milk. Lactose is synthesized in the mammary glands during lactation and is secreted into the milk. During digestion, lactose is hydrolyzed to its monosaccharide units, i.e., glucose and galactose by the intestinal enzyme referred to as lactase.
 - Maltose is formed when a molecule of glucose is linked to another molecule of glucose. It is obtained
 during hydrolysis of starch. During digestion, maltose is hydrolyzed to its monosaccharide units, i.e.,
 glucose and glucose by the intestinal enzyme referred to as maltase.
 - Sucrose is formed when a molecule of glucose is linked to fructose. It is also referred to as table sugar since it is commonly used on the table for routine use in home-made drinks. Sucrose is also referred to as cane sugar since it is obtained from the stems of sugarcane and roots of sugar beet. During digestion, sucrose is split into its constituent monosaccharides glucose and fructose by the enzyme sucrase (also called invertase).
 - Sucrose is used by people as a sweetener for foods (e.g., toast and cereal) and beverages (e.g., coffee and tea).
- Polysaccharides: Polysaccharides are made up of many monosaccharide molecules. Examples of
 polysaccharides are starch, glycogen (the form in which glucose is stored in the body), and cellulose and
 pectin (components classed as dietary fiber), etc.
 - Starches: Starch or amylum is a polymeric carbohydrate consisting of numerous glucose units joined by glycosidic bonds. This polysaccharide is produced by most green plants as energy storage. Starch molecules arrange themselves in the plant in semi-crystalline granules. Each plant species has a unique starch granular size, e.g., rice starch is relatively small while potato starches have larger granules. Starch content of different foods varies, e.g., potatoes 15%, wheat 55%, corn 65% and rice 75%.

Starch is a mixture of two polymers, i.e., amylose and amylopectin. Natural starches consist of about 10–30% amylose and 70–90% amylopectin.

- Amylose is a linear polysaccharide, composed entirely of D-glucose units joined by the α -1,4-glycosidic linkages.
- Amylopectin is a branched-chain polysaccharide, composed of glucose units linked primarily by α -1,4-glycosidic bonds but with occasionally α -1,6-glycosidic bonds, which are responsible for the branching. A molecule of amylopectin may contain many thousands of glucose units with branch points occurring after about every 25–30 units.

In the human body, several enzymes known collectively as **amylases degrade starch** sequentially into usable glucose units. Alpha-amylases are found in plants and animals. **Human saliva is rich in amylase** and the **pancreas also secretes the enzyme**.

The complete hydrolysis of starch yields, in successive stages, glucose:

$$Starch \rightarrow Dextrin \rightarrow Maltose \rightarrow Glucose$$

Starch is the *most common carbohydrate in the human diet* and is contained in many staple foods. The major sources of starch intake worldwide are the cereals (**rice, wheat and maize**) and the root vegetables (**potatoes and cassava**). Widely used prepared foods containing starch are bread, pancakes, cereals, noodles and pasta.

- **Glycogen**: Glycogen is the energy reserve carbohydrate of animals including humans, hence is also referred to as **animal starch**. Like starch in plants, glycogen is found as granules **in liver** (4–8% by weight of tissue) **and in skeletal muscle cells** (0.5–1.0%). When fasting, humans draw on these glycogen reserves during the first day without food to obtain the glucose needed to maintain metabolic balance.
 - Glycogen is **structurally quite similar to amylopectin**, however glycogen is more branched (8–12 glucose units between branches) and the branches are shorter. Enzyme **phosphorylase catalyzes the breakdown of glycogen to glucose**, when energy is needed, by a process called glycogenolysis.
- Cellulose: Cellulose is a polysaccharide (homopolysaccharide) like amylose, but it differs from amylose in the way that glucose molecules, in cellulose are linked together by β-1,4-glycosidic linkages instead of α-1,4-glycosidic linkages present in amylose. Cellulose is found in plants, as a structural component of their cell wall. We are, however, unable to digest cellulose due to the reason that human beings do not contain enzyme β-amylase, which is required for the digestion of cellulose.

Classification of Carbohydrates According to Nutritional Significance

According to their significance in nutrition, carbohydrates are classified as simple sugars and complex carbohydrates including fiber.

Simple Sugars

- Intrinsic sugars, which are incorporated into the cellular structure of foods, e.g., glucose, fructose, pentoses, etc., i.e., the sugars found in whole fruits and vegetables.
- Extrinsic sugars, which are not bound into a cellular structure, e.g., lactose (milk sugar) found in dairy products, sucrose (table sugar) found in confectionery, etc.

Complex Carbohydrates

- **Starch:** It is found in potatoes, bread, rice and pasta.
- **Dietary fiber:** These are the carbohydrate polymers, which are neither digested nor absorbed in the small intestine, e.g., cellulose, pectin, etc.

DIETARY FIBER

Dietary fiber refers to indigestible carbohydrates, such as cellulose and hemicellulose, pectin, lignin, gums and mucilage. Pectin, gums, mucilages and some of the hemicelluloses are soluble in water and hence, are also called **soluble fibers**. On the other hand, cellulose, various hemicelluloses and lignin are insoluble in water and are referred to as **insoluble fibers**.

Fiber cannot be digested by human beings due to the absence of the required digestive enzymes but is essential for digestion. Fiber promotes healthy bowel movement and decreases the risk of chronic diseases, such as coronary heart disease and diabetes. However, unlike sugars and starches, fiber is not absorbed in the small intestine and is not converted to glucose. Instead, fiber passes into the large intestine, relatively intact, where it is converted to hydrogen, carbon dioxide and fatty acids. Sources of fiber include fruits, grains and vegetables, especially, legumes.

Significance of Dietary Fiber

High intake of dietary fiber has many **health benefits**. For example, diets rich in fiber decreases risk of obesity, type 2 diabetes, colon cancer, cardiovascular disease and gastrointestinal disorders:

- Diets rich in fiber are usually, low in fat and provide lesser calories since such a diet forms bulk of the
 food, which takes longer to eat, and gives a feeling of fullness. Dietary fiber not only regulates body
 weight but also lowers the risk of obesity.
- Higher intake of soluble fiber also delays stomach emptying and thus, regulates blood glucose level. This in turn also reduces the risk of type 2 diabetes.
- Diet rich in fiber also lowers the risk of certain types of cancer. It is due to the reason that food with large
 amounts of fruits and vegetables, provides antioxidants, which protect our body against cell damage and
 cancer.
- Soluble fiber also lowers serum cholesterol, thus, reduces the risk of a cardiovascular disease.
- Diet rich in insoluble fiber also adds bulk to feces and softens stool for its easy passage. Thus, dietary fiber helps in **prevention of constipation and hemorrhoids**, particularly, when fluid intake is high.
- Excess of fiber in the diet may, however, cause deficiency of some minerals, such as zinc, calcium, magnesium and iron. It is due to the reason that dietary fiber binds these minerals and thus reduces their absorption.

Good Carbohydrates versus Bad Carbohydrates

Carbohydrates form the basis of most diets, making up half of total energy (calorie) intake. Dietary carbohydrates can be separated into two types, as complex (good) carbohydrates and simple (bad) carbohydrates.

Good carbohydrates: Carbohydrates, usually, considered to be good includes complex carbohydrates
such as those obtained from whole grains, fruits, vegetables, beans and legumes. They contain larger





Fig. 2.2: Sources of good carbohydrates

Fig. 2.3: Sources of bad carbohydrates

chains of sugar molecules and take longer to digest than processed grains. They are not only processed slowly, but also contain a bounty of other nutrients (Fig. 2.2).

Good carbohydrates are:

- Low or moderate in calories
- High in nutrients
- Devoid of refined grains and sugar
- High in naturally occurring fiber
- Low in sodium
- Low in saturated fat
- Very low in, or devoid of, cholesterol and trans fat
- Bad carbohydrates: Carbohydrates, usually, considered bad include refined sugars, such as those obtained
 from pastries, white sugar, honey, fruit juices and soda, highly processed foods, white rice, white bread and
 other white-flour foods (Fig. 2.3).

They are digested more quickly by the body. This makes them a faster source of energy. Also, these types of carbohydrates do not offer as many additional nutrients.

Bad carbohydrates are:

- High in calories
- Full of refined sugars
- Low in many nutrients

- Low in fiber
- High in sodium
- High in saturated fat, cholesterol and trans fat

Calorific Value of Carbohydrates

Calorific value or caloric value of the food is the amount of heat produced by combustion of one gram of fuel. It is expressed in terms of calories per gram or joules per kilogram. Calorific value of carbohydrate is 4 kcal/g.

Recommended Daily Allowances

Carbohydrates intake, for most people, should be such that it provides nearly 45 – 65% of total calories. Since, one gram of carbohydrate provides 4 calories, so a diet to provide 1,800 calories per day, should have between 200 and 300 grams of carbohydrates. However, people with diabetes should not eat more than 200 grams,

while pregnant women need at least 175 grams of carbohydrates per day.

Dietary Sources

Main dietary sources of carbohydrates include grains, vegetables and fruits. Complex carbohydrates are present in whole grains (such as cereals and rice), bran, beans, bread and pasta. Milk and milk products provide lactose, while beverages, jams, jellies and candies provide refined sugar, such as glucose and fructose (Fig. 2.4).

Fig. 2.4: Dietary sources of carbohydrates

Foods which are rich in carbohydrates include:

- Dairy: Milk, yogurt and ice cream
- Fruits: Whole fruits and fruit juice
- Grains: Cereal grains, bread and rice
- Legumes: Beans and pulses
- Starchy vegetables: Potatoes and corn
- Sugary sweets: Soda (when fluid intake is high), candy, cookies and other desserts.

Functions of Carbohydrates

Carbohydrates are necessary for good health and give a variety of benefits:

- **Brain health:** Carbohydrates are important for brain functions. They are a quick source of energy and have influence on mood and memory.
- Weight loss: Although, carbohydrates are often blamed for weight gain, the right kind of carbohydrate can actually help to maintain a healthy weight. This can happen due to the consumption of many good carbohydrates, especially, whole grains and vegetables with skin, which contain fiber.
- Good sources of nutrients: Whole, unprocessed fruits and vegetables are well known for their nutrients contents. Due the same reason, some of the food stuffs are even considered as super foods, such as the green leafy vegetables, sweet potatoes, berries, citruses and apples. Whole grains also have significantly higher amounts of fiber, energy and polyunsaturated fats, as well as all micronutrients (except vitamin B₁₂ and sodium). Further, whole grains have also been shown to contain antioxidants.
- Heart health: Fiber also helps to lower cholesterol. This is due to the reason that the digestive process
 requires bile acids, which are made, partly, from cholesterol. As the digestion improves, liver pulls
 cholesterol from blood, to form more bile acid, thereby, reducing the amount of LDL (the bad cholesterol).

DIGESTION, ABSORPTION AND STORAGE

DIGESTION OF CARBOHYDRATES

- Digestion of carbohydrates begins in the **mouth**, where **salivary amylase** hydrolyses starch into small polysaccharides (dextrins) and maltose.
- Chewing stimulates production of saliva and mixes the enzyme, salivary amylase with food. Only about 5% of the dietary starch is hydrolyzed, in the mouth.

- When dietary contents enter the stomach, Hcl (present in the gastric juice) stops the action of salivary amylase.
- After dietary contents enter small intestine, pancreas starts secreting pancreatic amylase, which further
 hydrolyses starch, and converts it to maltose, isomaltose, oligosaccharides and dextrins.
- Thereafter, brush border disaccharidases ridases hydrolyze disaccharides monosaccharides (Table 2.1).

TABLE 2.1: Effect of variou	s digestive enzymes o	on dietary carbohydrates
------------------------------------	-----------------------	--------------------------

Enzyme	Site of action	Catalytic action
Salivary amylase	Mouth	${\sf Starch/glycogen} \rightarrow {\sf Partially\ hydrolyzed\ dextrins,\ oligosaccharides}.$
		isomaltose and maltose
Pancreatic amylase	Small intestine	Partially hydrolyzed dextrins/oligosaccharides → Dextrins, maltose, isomaltose and maltotriose
α-Dextrinase	Small intestine	α -Limit dextrins \rightarrow maltotriose and glucose
Disaccharidases	Small intestine	
Sucrase		Sucrose → Glucose and fructose
Lactase		Lactose → Glucose and galactose
Maltase		Maltose/maltotriose → Glucose
Isomaltase		Isomaltose → Glucose

ABSORPTION OF MONOSACCHARIDES

All the dietary carbohydrates are absorbed into mucosal cell linings the small intestine, as monosaccharides, by two mechanism as fructose is absorbed by **facilitated diffusion**, glucose and galactose are absorbed by **secondary active transport**, by **sodium-linked glucose transporter-1** (SLGT-1).

After their absorption, all the mono-saccharides pass through the intestinal mucosal cells in the villi, and enter liver, via portal vein. In the liver, most of the galactose and fructose are further converted to glucose. Thereafter glucose is either stored in the liver or is oxidized for the generation of energy.

STORAGE OF GLUCOSE

Carbohydrates, mainly **glucose**, are **stored as glycogen in the liver and muscle**. Liver glycogen is hydrolyzed to glucose during fasting, whereas muscle glycogen is used as a source of energy during exercise. **Excess of glucose is also converted to fat and is stored in the adipose tissue as triacylglycerols (fat).**

METABOLISM OF CARBOHYDRATES

Metabolic Fates of Glucose

Glucose is metabolized in the liver and is utilized in various organs of the body, by different processes.

 Glucose is oxidized to produce energy, where it is converted to pyruvate, by the process of glycolysis. Thereafter, pyruvate is converted to acetyl CoA, which enters citric acid cycle, and produces energy (ATP) through electron transport chain and oxidative phosphorylation.

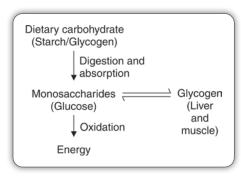


Fig. 2.5: Metabolic fates of glucose

• Glucose is also converted to glycogen, and is stored in the liver and muscle, by the process of **glycogenesis**. When blood glucose level is low, glycogen is converted to glucose, by the process of **glycogenolysis** (Fig. 2.5).

Glycolysis

Glycolysis is also called Embden Meyerhof glycolysis). Various reactions of glycolysis are outlined in Figure 2.6. In this process, firstly, glucose is converted to glucose-6-phosphate.

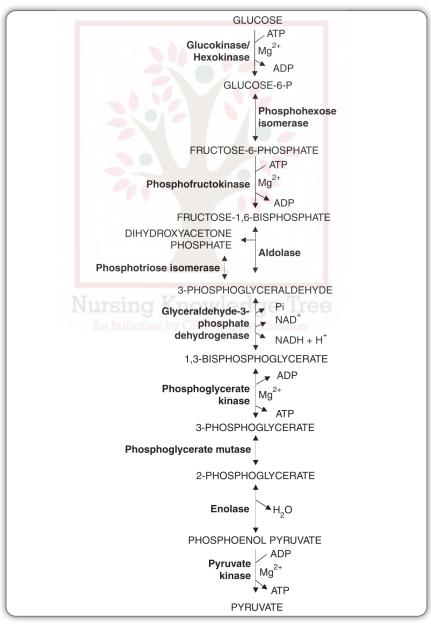


Fig. 2.6: Outline of glycolysis

Thereafter, via fructose-6-phosphate, it is converted to fructose-I, 6-bisphosphate. In the next step, fructose-I, 6-bisphosphate (a hexose) is cleaved to two trioses, i.e., 3-phosphoglyceraldehyde and dihydroxyacetonephosphate. Finally, these are converted, via phosphoenolpyruvate, to pyruvate, which has two fates:

Under aerobic conditions, pyruvate is transported into mitochondria, and is oxidatively decarboxylated to acetyl CoA. Via Krebs cycle and oxidative phosphorylation, a molecule of glucose provides **38** ATP.

Under anaerobic conditions, such as in the muscle during exercise, a molecule of glucose, however, provides only **2ATP**.

Glycogenesis

Glycogenesis, also referred to as **glycogen synthesis**, is the process of the conversion of glucose to glycogen, mainly in the liver and muscle. Various reactions of glycogenesis are outlined in Figure 2.7.

Regulation of Blood Glucose Level

Blood glucose level is maintained within the **normal** physiological **range of 60–90 mg/100 mL**, even in the fasting (post-absorptive)

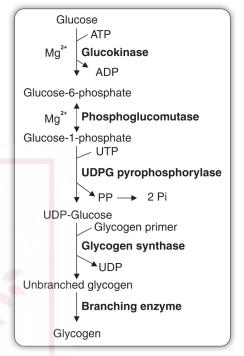


Fig. 2.7: Glycogenesis

state. If blood glucose level falls below the normal range, it is referred to as **hypoglycemia**. If blood glucose level rises above the normal range, it is referred to as **hyperglycemia**.

Blood glucose level is regulated by several hormones. Besides, some of the food stuffs also help in this process.

Regulation of Blood Glucose Level by Hormones

Insulin lowers blood glucose level, while several others, like glucagon, epinephrine (adrenaline), glucocorticoids, growth hormone and thyroxine increase blood glucose level (Fig. 2.8).

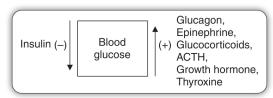


Fig. 2.8: Effects of hormones on blood glucose

During hyperglycemia (when blood glucose level is increased, such as after a meal), β -cells release insulin, which, in turn, increases glucose uptake by the cells and stimulates glycogen synthesis.

During fasting, when blood level falls, α -cells release glucagon, which, in turn, stimulates the breakdown of liver glycogen and forms glucose (Fig. 2.9).

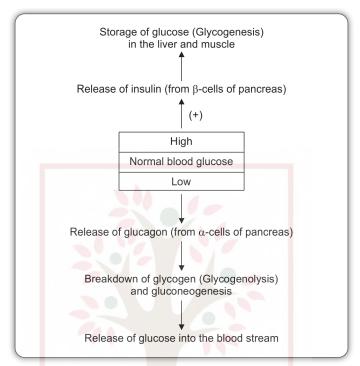


Fig. 2.9: Regulation of blood glucose by insulin and glucagon

During stress, epinephrine is released by the adrenal glands. It exhibits effects similar to glucagon and ensures that all the cells have adequate energy during emergency. Thus, epinephrine is also called **fight-or-flight** hormone.

Effect of The Type of Food String Washing Division

Type of the food also exhibits its effect on blood glucose level. Foods which are rich in simple carbohydrates or starch, but low in fiber and fat, are digested and absorbed rapidly. As a result of it, they cause large and rapid rise in blood glucose concentration. Body reacts to the rising blood glucose level and releases insulin, which, in turn, lowers blood glucose level and brings it to normal. Other foods, especially, those, which are rich in dietary fiber, resistant starch or fat, cause lesser response to blood glucose with small alterations.

Glycemic Index

Glycemic index (GT) refers to the effect of a particular type of food stuff on blood glucose level, i.e., it is a measure of how quickly the glucose is released into the bloodstream, after eating it. Glycemic index, thus, measures how quickly, and how much, a carbohydrate raises blood sugar. High-glycemic foods, like pastries, raise blood sugar rapidly, whereas low-glycemic foods raise it gently and to a lesser degree. Low GI foods, that are rich in fiber, should form an important component of a healthy diet.

MALNUTRITION

DEFICIENCY OF CARBOHYDRATES

When a person is on a carbohydrate-free diet, protein and fat are used as alternate sources of energy. Excessive breakdown of these substances, in turn, leads to the excessive formation of ketone bodies. Such individuals exhibit symptoms, similar to those seen during starvation, i.e., abnormal fat metabolism, breakdown of tissue proteins, increased sodium excretion, loss of energy and fatigue. Small amount of carbohydrates (50–100 g) prevent these symptoms. Not getting enough carbohydrate can cause problems, such as:

- Without sufficient fuel, body gets no energy.
- Additionally, without sufficient glucose, central nervous system suffers, which may cause dizziness, and mental and physical weakness.
- If body has insufficient carbohydrates intake, or stores, it will consume protein, as a source of fuel. This is problematic, because the body needs protein to make muscles. Using protein for fuel, instead of carbohydrates, also puts stress on the kidneys leading to the passage of painful by-products, such as urea, in the urine.
- People, who do not consume enough carbohydrates, may also suffer from deficiency of dietary fiber, which can cause digestive problems and constipation.

Overconsumption of Carbohydrates

Overconsumption of carbohydrates, in turn, leads to increased calories intake and causes obesity. This can also lead to diabetes mellitus and a cardiac disorder.

- Weight gain and obesity: Carbohydrates are the preferred source of energy for the body. When we eat any type of carbohydrate, our body breaks it to glucose, which is released into the blood stream and causes a rise in blood glucose level. If we eat moderate amount of carbohydrates, body's cells use, what they need for energy and store excess of it as glycogen. However, glycogen stores are limited, and once stores are saturated, excess of glucose is stored as fat. Thus, over consumption of carbohydrates, for a longer period, results in weight gain and obesity.
- Poor appetite control: Eating too much of carbohydrates, in one sitting, causes large increase in blood glucose level. This, in turn, causes pancreas to release a large amount of insulin, which leads to an equally large decrease in blood glucose level. When blood glucose level suddenly falls, brain sends hunger signals, meaning that one demands more food. Thus, the overall result is poor appetite control, as one is constantly craving for food, despite eating regularly.
- Fatigue: As overconsumption of carbohydrates causes a rapid surge and thereafter, a rapid fall in blood
 glucose level. This has a direct effect on overall energy levels. As a result of it, one will suddenly feel
 full of energy, and thereafter, shortly the energy level will crash. Fluctuating energy levels affect proper
 functioning of the body, since one will be constantly switching between the feelings of too much energy
 and not enough energy.
- Dental caries: Dental caries is the progressive destruction (demineralisation) of teeth, by the acids
 produced by bacteria acting on sugars and other fermentable carbohydrates on the tooth's surfaces.
 Longer a carbohydrate remains in the mouth, more likely it will promote dental caries. Foods that stick to
 the teeth, such as candies, cookies and beverages like cola, are more likely to cause dental caries. The most

- important way to protect dental caries is regular brushing of teeth with a fluoride containing toothpaste (twice daily). Fluoride strengthens tooth enamel and provides resistance to decay.
- Risk of type 2 diabetes: Level of glucose in blood is carefully monitored and maintained within the narrow limits, by the action of the hormone insulin. Over consumption of carbohydrates cause pancreas to produce, and release, increased amount of insulin. Pushing the pancreas to over exert, ultimately, interferes with its ability to produce insulin. Both, insulin resistance and pancreas failing to produce enough insulin, are characteristics of type 2 diabetes. This, in turn, also causes permanent damage to the body and affects all major organs and vessels including heart, kidneys, eyes, nerves and blood vessels.
- Risk of a cardiovascular disease (CVD): A high blood cholesterol level is also associated with increased risk of CVD. Some forms of complex carbohydrates, such as pectin in fruits and beta-glucan in oats, may, slightly, reduce the level of cholesterol in blood. These are the forms of soluble fiber. There is also some evidence that resistant starch may have a slight beneficial effect on the risk CVD. Regular consumption of wholegrain foods has also been linked with heart health, possibly, as a result of the breakdown products (e.g., short chain fatty acids), which are formed and absorbed in large bowel.
- **Risk of cancer:** Some studies suggest that population with a high intake of fiber-rich foods, experiences lower incidence of large bowel cancer than the population with low intakes of these foods.

Assess Yourself

Long and Short Answer Questions

- 1. Define carbohydrate. Describe the classification of carbohydrates.
- 2. Describe calorific value, RDA, dietary sources and functions of carbohydrates.
- 3. Describe various disorders which may occur as a result of overconsumption of carbohydrates.
- 4. Define glycolysis. Outline the process of glycolysis.
- 5. Define glycogenolysis. Outline reactions of this pathway.
- 6. What is glycogenesis? Give reactions of glycogenesis.
- 7. Differentiate between
 - a. Simple sugars and complex carbohydrates
 - b. Good carbohydrates and bad carbohydrates
 - c. Anaerobic and aerobic glycolysis
- 8. Discuss briefly:
 - a. Digestion of carbohydrates
 - b. Absorption of monosaccharides
 - c. Hormonal regulation of blood glucose
 - d. Gluconeogenesis
- 9. Write notes on:
 - a. Dietary fiber
 - b. Starches
 - c. Cellulose
 - d. Functions of carbohydrates

- e. Calorific value of carbohydrates
- f. Dental caries
- g. Glycemic index

Cell and its Structure

Chapter Outline

- Prokaryotic Cells
- Eukaryotic Cells

Structure and Functions of Intracellular Components

- The Nucleus
- The Endoplasmic Reticulum
- The Ribosomes

- The Golgi Apparatus
- The Mitochondria
- The Lysosomes
- The Peroxisomes
- The Cytosol
- The Plasma Membrane

The cell is a structural and functional unit of life. All animals and plants are made up of a large number of such units, in a manner to the utilization of bricks in the construction of a building.

Living cells are divided into two groups, i.e., the **prokaryotic cells** and the **eukaryotic cells**. As their name suggests (pro = prior to; karyot = nucleus; eu = true), the fundamental difference between them is the absence or presence of a true nucleus.

PROKARYOTIC CELLS

The simplest form of the cell is a prokaryotic cell. Prokaryotes, e.g., bacteria are **unicellular** and have one of the three basic shapes, viz. spheroidal (Cocci), rodlike (Bacilli) and helically-coiled (Spirilla).

- A prokaryotic cell is small in size (1 to 10 nm), relatively simple in structure and has only a single membrane, called cell membrane, which is usually surrounded by a rigid cell wall of characteristic structure.
- There may or may not be a surrounding **capsule**.
- Besides, there is a **single chromosome** comprised of a molecule of double helical DNA which is densely coiled to form a **nuclear zone**.
- Reproduction is by **asexual division**. The best characterized prokaryotic cell is *Escherichia coli*.
- Some prokaryotes **possess pili** and **flagella** for adhesion and movement, respectively (Fig. 2.1).

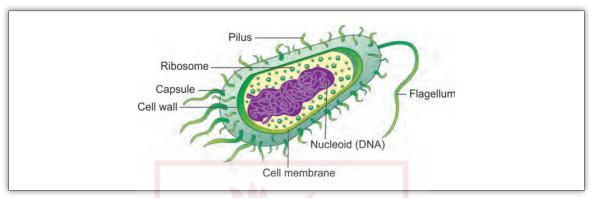


Fig. 2.1. A prokaryotic cell.

Organelles of Prokaryotic Cell

Cell Wall

- Found only in prokaryotic cells
- Surrounds the plasma membrane
- Gives the cell its shape
- Prevents bursting when turgid pressure is high
- Helps anchor appendages like pili and flagella
- Composed of cellulose microfibers which form a thick wall
- Grows with the cell
- Helps the cell maintain osmotic balance.

Plasma Membrane

- Found in both prokaryotic and eukaryotic cells
- Encloses the interior of the cell
- Regulates the flow of material in and out of the cell.

Ribosomes

- 70S ribosome is made up of 50S and 30S components
- Translates the genetic code from the DNA to make proteins
- They can be either free or attached to endoplasmic reticulum.

Nucleoid

- There is no clear nuclear cytoplasmic difference
- Not a membrane bound nucleus
- An area of the cytoplasm where the strands of DNA are found.

Organelles not found in Prokaryotic Cells

- Endoplasmic reticulum
- Golgi apparatus
- Mitochondria
- Nucleus

Pili

- Small hairlike projections emerging from the outside cell surface
- Assists in the cell attaching to other cells and surfaces. Thus helps in pathogenic organisms to stick to the surface.

Flagella

- Flagella are hairlike structures that allow the cell to move by beating in a propeller-like motion
- They help bacterium to move towards nutrients, away from toxic chemicals, etc.

EUKARYOTIC CELLS

Animals, plant, fungi and protozoa are called eukaryotes which may be unicellular or multicellular.

- Eukaryotic cells are one to ten thousand times **large in size** and are more **complex in structure** than the prokaryotic cells.
- They may **vary from one tissue to another** with respect to their functions, e.g., the liver parenchymal cell, adipose cell, nerve cell, renal tubular cell, white blood cell, etc.
- Generally, a eukaryotic cell has a well-defined membrane-bound nucleus containing several chromosomes.
- Their chromosomes undergo replication of DNA during mitosis and get separated into daughter chromosomes, i.e., these cells reproduce by cell division.
- A typical eukaryotic cell contains various organelles such as nucleus, endoplasmic reticulum, Golgi apparatus, mitochondria, etc. (Fig. 2.2).

Differences between Prokaryotic and Eukaryotic Cell Structures

Major differences between prokaryotic and eukaryotic cell structures are given in Table 2.1.

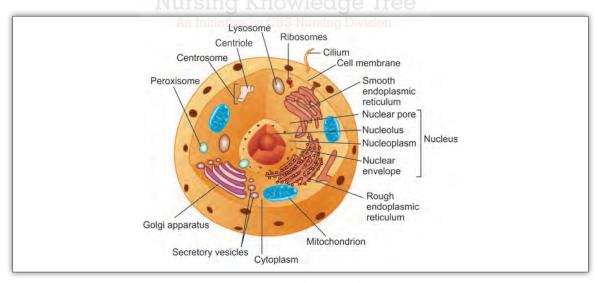


Fig. 2.2. A eukaryotic cell.

Parameters Prokaryotic cells Eukaryotic cells Cell size Small Large **Overall organization** Simple Complex Cell membrane **Boundary** Cell membrane and cell wall, sometimes surrounded by a capsule Subcellular entities Few Many Nucleus They have a single nucleus which contains Well defined nucleus with a double helix DNA that is present in a poorly membrane and multiple DNA double defined region called nucleoid helices organized into chromatin Reproduction Asexual Sexual **Examples** Bacteria and blue-green algae Plant and animal cells

TABLE 2.1. Major differences between prokaryotic and eukaryotic cell structures

STRUCTURE AND FUNCTIONS OF INTRACELLULAR COMPONENTS

THE NUCLEUS

- The nucleus is the largest component of the cell, containing DNA organized into separate chromosomes and is surrounded by a membrane called nuclear membrane.
- The **nuclear membrane** consists of two layers which are separated by an intermembrane space termed as perinuclear space (**cisterns**). The **outer membrane** though is continuous with the endoplasmic reticulum but the two layers of the nuclear membrane are fused together at several places producing nuclear pores for the exchange of materials between the nucleus and the cytoplasm.
- The nucleus is filled with the **nucleoplasm** which has a discrete body called nucleolus and a thread-like structure called chromatin (Fig. 2.3).

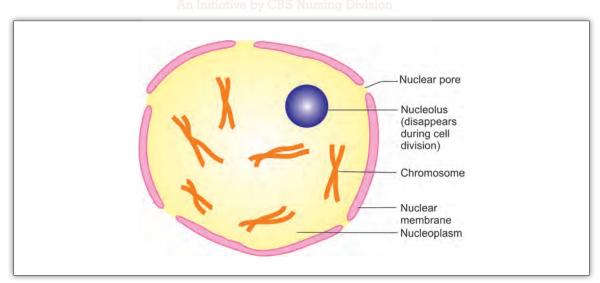


Fig. 2.3. The nucleus.

- Nucleolus: The number of nucleoli may vary from one cell type to another. The genes for three of the
 four ribosomal RNA molecules are located in the nucleolus. Nucleoli are rich in RNA and disappear
 during cell division.
- **Chromatin:** It contains most of the cellular DNA in association with basic proteins, termed histones. At the time of cell division, the chromatin is organized into small thread-like structures called **chromosomes**. Human somatic cell contains 23 pairs of chromosomes.

Important Functions of the Nucleus

Important functions of the nucleus include:

- Control of cell division (DNA replication)
- **Protein synthesis** (by controlling the synthesis of RNA)

The regulation of DNA synthesis and other functions of the nucleus are severely disturbed in some pathological conditions such as cancer.

DNA and **DNA** polymerase are the markers of the nucleus.

THE ENDOPLASMIC RETICULUM

Endoplasmic reticulum is a system of membranes (lipid bilayer structures) with a network of vesicular spaces. This network is present throughout the cytoplasmic matrix and grows by its own synthesis. These membranes run parallel to each other creating channels which are called **cisternae**. The interior of the endoplasmic reticulum thus is well connected with perinuclear spaces and through pores on the cell surface, with the extracellular space. Cisternae have a role in the exchange of materials between the cell and the extracellular fluid. The surface of the endoplasmic reticulum may or may not bear ribosomes. Accordingly, endoplasmic reticulum is of two types:

- 1. **Rough endoplasmic reticulum (RER):** It is also called the granular type of endoplasmic reticulum since it has small granules attached to it. These granules are termed as ribosomes.
- 2. **Smooth endoplasmic reticulum (SER):** It is also called the agranular type of endoplasmic reticulum since it consists of the membranous structure only and does not contain ribosomes on its outer surface. The SER has enzymes for the biosynthesis of lipids and glycoproteins. Further, SER are very important in hepatocytes where these are primarily concerned with oxidative metabolism and for the detoxification of many drugs and other toxic organic molecules.

Glucose-6-phosphatase is a marker enzyme for the endoplasmic reticulum.

THE RIBOSOMES

- Ribosomes consist of ribonucleoprotein particles of two sizes, i.e., 50S and 30S in prokaryotes or 60S and 40S in eukaryotes. Because of their high RNA content, ribosomes are the site of protein synthesis.
- Ribosomes on the RER are associated with the synthesis of proteins for export from the cell.
- Free ribosomes, on the other hand, are present in the cytoplasm and synthesize proteins for use within the cell.
- **RNA** is used as a marker for the ribosomes.

THE GOLGI APPARATUS

- The Golgi apparatus is a smooth membrane system with **vacuoles**. It is rich in lipids and is considered to be the site where secretions from other organelles are brought and assembled.
- The **newly synthesized proteins** are also transferred from RER and **stored in the Golgi apparatus**, temporarily.
- Some of the synthesized proteins also **undergo post-translational modifications** within the Golgi apparatus and thereafter, are transported to different destinations. The Golgi apparatus is thus, especially active in cells which produce proteins for export. They form secretory granules for the proteins after their synthesis on the ribosomes.
- Galactosyl transferase is a marker enzyme for the Golgi apparatus.

THE MITOCHONDRIA

The mitochondria are the major organelle of a eukaryotic cell lacking any direct structural relationship with other organelles and contain its own DNA (Fig. 2.4).

A mitochondrion produces energy in the form of ATP (adenosine triphosphate) for the cellular functions and is thus, called **power house of the cell**. Thus, depending upon energy requirement of the cell, mitochondria may vary in size, shape and number, from cell to cell.

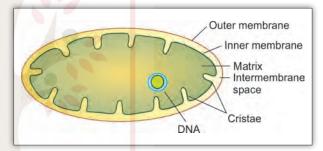


Fig. 2.4. A mitochondrion.

- Besides producing energy, mitochondria also help to **control the level of calcium** in the cytoplasm.
- Most of the cells contain several hundred mitochondria.

Mitochondrial Membrane

It is a double-layered structure where the two layers are separated from each other by 50–100 Å intermembrane space. Several enzymes especially those involved in the nucleotide metabolism are located here.

- The **outer membrane** of the mitochondria has a smooth structure. It is composed of both lipids and proteins and is freely permeable to most of the small molecules. Several enzymes involved in lipid metabolism, such as the enzymes for fatty acid elongation, glycerol phosphate acyltransferase and phospholipase A, are associated with the outer membrane of the mitochondria.
- The **inner membrane** of the mitochondria has a denser structure. It has more proteins than lipids. The inner membrane has extensive irregular foldings, called **cristae**. Cytochromes, the enzymes of electron transport chain and flavoproteins are localized within the inner membrane of the mitochondria.

Matrix

The intra-mitochondrial space (enclosed within the inner membrane) is called mitochondrial matrix. This chamber contains enzymes for β -oxidation of fatty acids, citric acid cycle and glutamate dehydrogenase. In addition, mitochondria also have DNA, referred to as mtDNA (mitochondrial DNA).

Parameters	Nuclear DNA	mtDNA
Location	Nucleoplasm	Mitochondrial matrix
Inheritance	Both paternal and maternal	Maternal
Shape	Linear	Circular
Base pairs	3 x 10 ⁹	16 x 10 ³
Introns	Present	Absent
Histones	Present	Absent
Encoded proteins	Used within the cell as well as exported	Used within mitochondria
Repair mechanisms	Well developed	Poorly developed
Mutation rate	Low	High

TABLE 2.2. Comparison between nuclear DNA and mitochondrial DNA (mtDNA)

Mitochondrial DNA: Mitochondria are the only cellular organelles that contain their own chromosomal DNA (mtDNA), which is maternally inherited. Human mtDNA is a small double stranded circular molecule (about 16,000 base pairs), encoding 13 polypeptides that are integrated into the inner mitochondrial membrane along with other polypeptides encoded by nuclear genes. In addition, it encodes 2 rRNAs and 22 tRNAs that are used in protein synthesis within the organelle. mtDNA differs from nuclear DNA in several aspects.

Comparison between Nuclear DNA and mtDNA

Comparison between nuclear DNA and mtDNA is given in Table 2.2.

In contrast to nuclear DNA, the mtDNA is exposed to high levels of mutagenic free radicals and is not protected by the usual DNA repair mechanisms. This results in mutations in mtDNA that affect mitochondrial structure and function, leading to **various muscular** and **neurological disorders**, such as Leber's hereditary optic neuropathy, mitochondrial encephalopathy, chronic progressive external opthalmoplegia, mitochondrial encephalopathy lactic acidosis and stroke like episode (MELAS), etc.

Succinate dehydrogenase and glutamate dehydrogenase are marker enzymes for mitochondria.

THE LYSOSOMES

Lysosomes (derived from the Greek words lysis, meaning **to loosen** and soma, **body**) are the membrane-bound cell organelle found in animal cells. They are absent in red blood cells. Lysosomes are structurally and chemically spherical vesicles containing hydrolytic enzymes which are capable of breaking down all kinds of biomolecules including proteins, lipids, carbohydrates, nucleic acids and cellular debris.

- Lysosomes are known to contain more than fifty different enzymes which are all active at an acidic
 environment of about pH 5. Thus, lysosomes act as waste disposal system of the cell by digesting
 unwanted materials in the cytoplasm both from outside of the cell and obsolete components inside the
 cell.
- Further, lysosomes are responsible for **cellular homeostasis** for their involvements in secretion, plasma membrane repair, cell signaling and energy metabolism, which are related to health and diseases.
- Depending on their functional activity their sizes can vary.
- Acid phosphatase is a marker enzyme of the lysosomes.

Suicide Bags

Lysosome is a bag (oval, rounded) that is loaded with digestive enzymes which, when activated, act upon cellular structures and demolish them leading to cell death. They are referred to as suicide-bags due to their role in **autolysis**, a process by which a biological cell self-destructs, i.e., during the period of starvation it engulfs or eats the whole cell.

Residual Bodies

Cytoplasmic vacuole containing the leftover products of digestion (e.g., membrane fragments or ferritin-like particles) after fusion with the contents of a lysosome is referred to as a residual body. Residual bodies are either secreted by the cell via exocytosis (generally in macrophages) or become lipofuscin granules that remain in the cytosol indefinitely. Longer-living cells like neurons and muscle cells usually have a higher concentration of lipofuscin than other more rapidly proliferating cells.

THE PEROXISOMES

- Peroxisomes, also called microbodies, contain certain oxidative enzymes, e.g., uric acid oxidase, D-amino acid oxidase and catalase, etc.
- The major function of the peroxisomes is in hepatocytes where they are involved in the oxidation of fatty acids by a modified α-oxidation pathway.

Clinical Correlation

Zellweger Syndrome

Absence of the functional peroxisomes results in a rare autosomal recessive disease referred to as Zellweger syndrome. It is characterized by abnormalities of the several organs due to the decreased levels of plasmalogens.

- Besides, peroxisomes also represent a very primitive cellular organelle which provides protection to the cell from the toxicity of hydrogen peroxide (H₂O₂).
- Catalase and uric acid oxidase are marker enzymes of the peroxisomes.

THE CYTOSOL

- The cytosol or **cell-sap** is a structureless material filling the cell (aqueous matrix) in which all the cellular organelles float.
- It is a colloidal solution of proteins containing nearly 70% water.

Clinical Correlation

Lysosomal Storage Diseases

Lysosomes are responsible for a group of genetically inherited disorders called lysosomal storage diseases (LSD). They are a type of inborn errors of metabolism caused by malfunction of one of the enzymes. The rate of incidence is estimated to be 1 in 5,000 live births.

The primary cause is deficiency of an acidic hydrolase while there are conditions of defects in lysosomal membrane proteins that fail to transport the enzyme or non-enzymatic soluble lysosomal proteins. The initial effect is accumulation of specific macromolecules or monomeric compounds inside the endosomal—autophagic—lysosomal system. This results in abnormal signaling pathways, calcium homeostasis, lipid biosynthesis and degradation and intracellular trafficking, ultimately leading to pathogenic disorders. The organs most affected are brain, viscera, bone and cartilage. There is no direct medical treatment to cure LSDs.

Gaucher's disease: Gaucher's disease is due to deficiency of the enzyme glucocerebrosidase. Consequently, the enzyme substrate, the fatty acid glucosylceramide accumulates, particularly in white blood cells. This in turn affects spleen, liver, kidneys, lungs, brain and bone marrow. The disease is characterized by bruises, fatigue, anemia, low blood platelets, osteoporosis and enlargement of the liver and spleen.

- Besides proteins, the cytosolic fraction also contains various enzymes for glycolysis, gluconeogenesis and HMP shunt (Hexomono phosphate shunt), and a variety of organic as well as inorganic substances such as glucose, potassium and magnesium.
- The cytosol is in contact with all the cellular organelles and an important vehicle for the transport of metabolites from one organelle to the other.
- The cytosol of all eukaryotic cells also contains a network of fibers, collectively called **cytoskeleton**, which includes microtubules, intermediate filaments and microfilaments.
- Lactate dehydrogenase is a marker enzyme of the cytosol fraction of the cell.

THE PLASMA MEMBRANE

Plasma membrane or cell membrane is the outer membrane of the cell. It is in contact with the extracellular matrix.



Assess Yourself

Long and Short Answer Questions

- Describe structure, functions and biochemical markers of various subcellular entities of a eukaryotic cell.
- 2. Explain:
 - a. Mitochondria as a power house of the cell.
 - b. Functions of the rough endoplasmic reticulum.
 - c. Differences between nuclear and mitochondrial DNA.
- 3. Write short notes on:
 - a. The nucleus
 - c. The Golgi apparatus
 - e. Suicide bags
 - g. Lysosomal storage diseases

- b. Ribosomes
- d. Matrix
- f. Age-pigments
- h. Lysosomes and peroxisomes

Multiple Choice Questions

- 1. Protein synthesis takes place mainly in:
 - a. Ribosomes
 - c. Centrosome

- b. Mitochondria
- d. Golgi apparatus
- 2. In Lysosomal storage disorders, true is:
 - a. The lysosomes are deficient in enzyme hydrolase
 - b. There is a defect in fusion of lysosomes and phagosomes
 - c. There is a defect in lysosomal membrane
 - d. There is increased degradation of heteroglycans
- 3. False about Mitochondria is: no Knowledge Tree
 - a. Absent in RBCs
 - b. Powerhouse of the cell
 - c. Electron transport chain enzymes are present in outer membrane
 - d. Cytochrome P450 in inner membrane
- 4. Mitochondrial DNA is:
 - a. Closed circular
 - c. Linear

- b. Nicked circular
- d. Open circular

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- Clinical correlation boxes have been added at various places in the text to help the students understand the clinical relevance of the subject.
- · Chapters are studded with frequently asked questions under Assess Yourself Section for self-evaluation.
- · Important terms and points have been highlighted throughout the chapters.

About the Author

Harbans Lal *PhD, FACBI, FSOBSI, FIAO,* joined Medical College, Rohtak (Haryana), presently Postgraduate Institute of Medical Sciences, Pt BD Sharma University of Health Sciences, Rohtak (Haryana) as a Lecturer in Biochemistry and after nearly 34 years of service, retired in 2009 from the same institution as Senior Professor. Thereafter, he joined the Department of Biochemistry at Maharaja Agrasen Medical College, Agroha, Hisar (Haryana) as a Senior Professor and Head, from where he retired in July 2017. During this period, he supervised a large number of PhD and MD students, wrote 9 textbooks and published more than 150 research papers in various National and International Journals. He is also a peer reviewer of several National and International Research Journals.

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