Polymers



Macromolecules

• Individual molecule of very high molecular weight. e.g., Heamoglobin, Chlorophyll etc.

Monomer

 Monomer is the molecule that forms the basic unit for Polymer. e.g., Propene, Styrene, Vinyl chloride etc.

Polymer

 Natural, semisynthetic and synthetic compounds consisting of number of molecules (monomers) called Polymer. e.g., Polypropene, Polystyrene, PVC, Nylon-6 etc.

Definition

Polymers which are obtained from animal and plants are known as natural polymers.

Concept Ladder

- S - C	7
	4
	4
- 5	Т
	1
	8

The number of monomers which are joined together in a polymer constitutes the degree of polymerization.



I. Classification Based upon source

(1) Natural polymers

Polymers which are found in nature. eg. Starch, cellulose and natural rubber, Silk, Proteins, Nucleic acid. ...etc.

(2) Semisynthetic polymers

It is obtained from naturally occuring polymers by chemical treatment. Most of the semisynthetic polymers are prepared

Rack you				
Why macromo macromo polymers	lecul	les	ymers but are	are all not

from cellulose. eg. cellulose acetate, cellulose nitrate, cellulose xanthate and rayon.

(3) Synthetic polymers

Man maid polymers are known as synthetic polymer. eg. PVC, polyethylene, polystyrene, nylon-6, nylon-6,6; nylon-6,10; terylene, synthetic rubbers etc.

Classification Based upon structure

1. Linear polymers

Monomers are joined together to form long straight chains. The various linear polymeric chains are stacked over one another to give a well packed structure close packed in nature, having high densities, high melting point and high tensile (pulling) strength.

Note:

All fibers are linear polymers. eg. cellulose, silk, nylon, terylene etc.

2. Branched chain polymers

Polymers in which the monomeric units constitute a branched chain. Branched chain polymers have lower melting point low densities and tensile strength as compared to linear polymers.

Examples are amylopectin, glycogen, low density polyethylene and all vulcanized rubbers.

3. Cross linked or Three-Dimensional network polymers

When linear polymeric chains are joined together to form a three dimensional network structure. These polymers are hard,

Concept Ladder

Polymers in which repeating units are derived from two or more types of monomers units are called copolymers. eg. nylon-6,6; Buna-S; Buna-N; Polyesters.

Definition

Man-made polymers, i.e. polymers prepared in laboratory are known as synthetic polymers.

Previous Year's Question

Which compound forms linear polymer due to H-bond?

	[AIPMT]
(1) H ₂ O	(2) NH ₃
(3) HF	(4) HCl

rigid and brittle. Cross linked polymers are always condensation polymers. Resins are cross linked polymers.

Classification Based upon molecular force

1. Elastomers

Polymers in which the intermolecular foreces of attraction between the polymer chains are the weakest (weak van der Waals forces of attraction) these polymers consist of randomly coiled molecular chains of irregular shape having a few cross links. Examples are natural rubber, Buna-S, Buna-N etc.

2. Fibres

Those polymers in which teh intermolecular forces of attraction are the strongest are called fibers. These polymers held together by H-bonding or dipole-dipole interaction. Fibres have high tensile strenght, least elasticity having high melting point and low solubility.

3. Thermoplastics

In thermoplastics intermolecualr forces of attraction are in between those of elastomers and fibres. Thermoplastics become soft and viscous on heating and rigid on cooling.

Examples are polythene, nylon-6, nylon-6,6 etc.

4. Thermosetting polymers

These polymers have low molecular masses and are semi-fluid substances. These polymers are hard and infusible.

Examples are melamine-formaldehyde, bakelite (phenol-formaldehyde) etc.

Concept Ladder

Ethene (Monomer)

Polymers whose repeating structural units are derived from only one type of monomer units are called homopolymers. $nCH_2 = CH_2 \longrightarrow [-CH_2 - CH_2 -]_n$

Rack your Brain



Polythene (polymer)

Why branched chain polymers do not pack well?

Concept Ladder



Linear polymers may be condensation as well as addition polymers. Examples are cellulose, polypeptide, nucleic acid, nylon, terylene etc.

Classification based upon polymerisation

1. Addition polymerization

- (a) Addition polymers are those in which addition reaction takes place
- (b) If monomer is ethylene, then addition polymer may be either linear polymer of branched chain polymer. Examples are polystyrene, polytetrafluoroethylene, polyacrylonitrile etc.
- (c) If monomer is 1,3-butadiene or 2substituted-1, 3-butadiene

 $\begin{pmatrix} CH_2 = C - CH = CH_2 \\ I \\ G \end{pmatrix}$, then polymer is always

branched chain polymer.

$$n CH_{2} = C - CH = CH_{2} \longrightarrow \begin{pmatrix} G \\ -CH = C - CH = CH - \\ (Monomer) & (Polymer) \end{pmatrix}_{n}$$

Monomer Polymer G = H: (i) Polybutadiene (a) 1.3-Butadiene G-CH₃: 2-Methyl-1, (ii) Polyisoprene (b) 3-butadiene or isoprene G=Cl; 2-Chloro-1, (iii) Polychloroprene 3-butadiene or (c) (Neoprene) chloroprene

(d) Addition polymers are also known as chain growth polymers.

Concept Ladder

Addition polymerization takes place in three steps: Initiation, chain propagation and chain termination.

Previous Year's Question

Regarding cross-linked or network polymers, which of the following statements is incorrect? [NEET]

(1) They conttain covalent bonds between various linear polymer chains.

(2) They are formed from bi-and tri-functional monomers.

(3) Examples are bakelite and melamine.

(4) They contain strong covalent bonds in their polymer chains.

Types of Addition Polymerisation

(a) Free Radical Polymerisation

 Free radical polymerisation takes in the presence of radical initiators. Such as dioxygen, benzoyl peroxide, acetyl peroxide etc.

> For example, polymerization of ethene is carried out at high temperatures (350-570K) and at high pressure (1000-2000 atm) in presence of dioxygen or a small amount of benzoyl peroxide as radical initiator.

Mechanism

Chain initiating steps:





 $\dot{C}_{6}H_{5} + CH_{2}=CH_{2} \longrightarrow C_{6}H_{5}-CH_{2}-\dot{C}H_{2}$ Ethene Free radical (I) (Monomer)

Chain propagating steps:

$$C_{6}H_{5}-CH_{2}-\dot{C}H_{2} + CH_{2}=\dot{C}H_{2} \longrightarrow C_{6}H_{5}-CH_{2}-CH_{2}-CH_{2}-\dot{C}H_{2} \longrightarrow C_{6}H_{5}-CH_{2}-CH_{2}-\dot{C}H_{2} \longrightarrow C_{6}H_{5}-(CH_{2}-CH_{2})_{n}-CH_{2}-\dot{C}H_{2}$$

Free radical (II)

Previous Year's Question

stick to wound?

Which can absorb over 90% of its

own mass of water and does not

[AIPMT]

Chain terminating steps:

Step 1: By combination of free radicals (II)

$$C_{6}H_{5}-(CH_{2}-CH_{2})_{n}-CH_{2}-\dot{C}H_{2} + \dot{C}H_{2}-CH_{2}-(CH_{2}-CH_{2})_{n}-C_{6}H_{5} \longrightarrow$$

$$C_{6}H_{5}-(CH_{2}-CH_{2})_{n}-CH_{2}CH_{2}CH_{2}CH_{2}-(CH_{2}-CH_{2})_{n}-C_{6}H_{5}$$
Polythene (Polymer)



Step 2: By disproportionation of free radicals (II)

$$C_{6}H_{5}-(CH_{2}-CH_{2})_{n}-CH_{2}-\dot{C}H_{2} + \dot{C}H_{2}-CH_{2}-(CH_{2}-CH_{2})_{n}-C_{6}H_{5} \xrightarrow{\text{Disproportionation}} \\C_{6}H_{5}-(CH_{2}-CH_{2})_{n}-CH_{2}CH_{3} + C_{6}H_{5}-(CH_{2}-CH_{2})_{n}-CH_{2}=CH_{2}$$

$$_{5}$$
—(CH₂—CH₂)_n—CH₂=

(b) Cationic Polymerisation

Cationic polymerisation takes place in the presence of strong protonic acids such as H₂SO₄, AlCl₃, BF₃ etc.



(c) Anionic Polymerization

In anionic polymerisation, it takes place in the presence of strong bases such as KNH2, NaOH, KOH, some organometallic compounds etc.



2. Condensation polymerization

(a) They are formed due to condensation reactions.

- (b) Condensation polymerization is also known as step growth polymerization.
- (c) For condensation polymerization, monomers may should have at least two functional groups. (functional groups may be same or different).

For example,

Previous Year's Question



Which one of the following is not a condensation polymer?

[AIPMT]

- (1) Melamine (2) Glyptal
- (3) Dacron (4) Neoprene

$$nNH_{3} - R - NH_{3} + nHOOC - R' - COOH \xrightarrow{Condensation} \left[-NH - R - NH - C - R' - C - \right]_{n}^{+} (n-1)H_{2}O$$
Polyamide

$$nHO - R - OH + nHO - C - R' - C - OH \xrightarrow{Condensation} - O - R - O - C - R' - C - n + (n-1)H_2O$$
Polyester

$$nNH_2 - R - COOH \xrightarrow{Condensation} \left[-NH - R - C - \\ Polyamide - NH - R - C - \\ n \right] n$$

 (d) During condensation elimination of small molecules like ammonia, alcohol, water, HCl takes place.

$$n \text{ NH}_{2} - R - \text{COOH} \xrightarrow{\text{Condensation}}_{\text{reaction}} \left[\begin{array}{c} O \\ -NH - R - C - \\ Polyamide \end{array} \right]_{n} + \left(\begin{array}{c} n \\ 2 - 1 \end{array} \right)$$

 Monomer having three functional groups always gives cross linked polymer. Examples are : Urea formaldehyde resin, phenolformaldehyde resin.

Concept Ladder

Monomers having only two functional group always give linear polymer.

Natural Rubber

The main source of natural rubber is braziliensis tree. Natural rubber is obtained from latex. Again latex is coagulated with acetic acid and formic acid and coagulated mass is then squeezed.

Natural rubber is a sticky, gummy & soft and insoluble in water, alkalies & dilute acids. Natural rubber soluble in non-polar solvents. It has low elasticity and low tensile strength. Example : Polyisoprene.

Concept Ladder

Natural rubber is a polymer of 2-methyl-1, 3-butadiene (isoprene). On average, molecules of rubber contain 5000 isoprene units held together by head to tail. All the double bonds in rubber are cis, hence natural rubber is cis-polyisoprene.

n $CH_2 = C - CH = CH_2 \xrightarrow{Polymerization} [-CH_2 - C - CH - CH_2 -]_n$ Isoprene Polyisoprene

Structure of Natural Rubber (cis-polyisoprene)



Synthetic Rubber or Polymerisation of Dienes

Polymers of 1,3-butadienes are called synthetic rubbers because they have some properties of natural rubber. Synthetic rubber are water proof and great elasticity.

1. Homopolymers

Monomer of this class is 2-substituted G $CH_2 = C - CH = CH_2$ where G = H, Cl or CH₃.

2 2 3

Zieglar-Natta catalyst is used for polymerisation which gives stereo regular polymers.

Previous Year's Q	uestion 😗
natural rubber is a	polymer of [AIPMT]
(1) Styrene (3) Butadiene	(2) Ethyne (4) Isoprene



Dieprene (Neoprene) was the first synthetic rubber manufactured on large scale. It is the monomer of chloroprene (2-chlorobutadiene).

2. Copolymers

Copolymers are derived from two or more types of monomer units. Example of copolymers are :

1	Buna-S, SBR (Styrene- Butadiene rubber)	$C_{6}H_{5} - CH = CH_{2} + CH_{2} = CH - CH = CH_{2}$
2	Buna-N, NBR (Nitrile- Butadiene rubber)	$CH_2 = CH - CN + CH_2 = CH - CH = CH_2$ (25%)
3	Butyl rubber	$CH_{3} - C = CH_{2} + Butadiene$ (2%) CH_{3} (98%)
4	ABS; Acrylonitrile, Butadiene, Styrene	$\label{eq:CH2} \begin{split} CH_2 &= CH - CN + CH_2 = CH_2 = CH - CH = CH_2 \\ &+ C_6 H_5 CH = CH_2 \end{split}$

(a) Buna-S (SSR : Styrene-butadiene rubber)

Buna-S rubber is a copolymer of three moles of butadiene and one mole styrene. Buna-S is generally compounded with

Rack your Brain

Which polymer is used to make 'non-stick' cookware?

Synthetic rubbers have

Concept Ladder

some improved properties. They are more flexible, tougher and more durable than natural rubber.

9.

carbon black and vulcanised with sulpur. It is extremely resistant to tear & wear and therefore used in the manufacture of tyres and other mechanical rubber goods. It is obtained as a result of free radical copolymerisation of its monomers.

(b) Buna -N

Buna-N is obtained by copolymerisation of butadiene and acrylonitrile (General purpose Rubber acrylonitrile or GRA)

Concept Ladder

In Buna-S, 'Bu' stands for butadiene, 'na' for symbol of sodium (Na) which is a polymerizing agent and 'S' stands for styrene. It is an elastomer (General purpose styrene Rubber or GRS).



Buna-N is rigid and resistant to the action of organic solvents, lubricating oil and petrol. It is also used for making fuel tanks.

$$nCH_{2}=CH-CH=CH_{2} + nCH_{2}=CH \longrightarrow \begin{bmatrix} CH_{2}-CH=CH-CH_{2}-CH_{2}-CH_{2} \end{bmatrix}_{n}^{L}$$

Buna-S is also called?	
(1) Polyurethane	(2) SBR
(3) Teflon	(4) Bakelite

Sol. (2)

SBR is known as styrene butadiene.

Vulcanisation of rubber

Natural rubber soft & sticky and becomes even more so at high temperatures and brittle at low temperanures. Therefore, rubber is generally used in the temperature range 283-335 K where its elasticity is maintained. It has a large water absorption capacity, has low tensile strength and low resistance to abrasion. It is also not-resistant to the action of organic solvents and is also easily attacked by axidising agents. These properties can be improved markedily by a process called vulcanization.



It consists of heating raw rubber with sulphur at 373-415 K. Since this process is slow, therefore, additives like zinc oxide, are used to acoelerate the rate of vulcanization The vulcanized rubber thus obtained has excellent elasticity, low water absorption tendency and is resistant to the action of organic solvents and oxidising agents During vulcanization, sulphur bridges or crosslinks between polymer chans are introduced either reactive allylic positions or at the sites of the double bonds.

Nvlon

Nylons are prepared by the condensation polymerisation of dibasic acids with diamines. Nylon contain amide linkages having a protein like structure.

Concept Ladder

Synthetic rubbers are either homopolymers 3-butadiene of 1, its derivatives or or copolymers are of 3-butadiene or 1. its derivatives with another unsaturated monomer.





(1) Nylon-6,6 (Nylon-six, six)

It is obtained by condensation polymerisation of a diamine with six carbon atoms (hexamethylenediamine) and a dibasic acid having 6 carbon atoms (adipic acid).

n HOOC(CH₂)₄COOH + n H₂N(CH₂)₆NH₂ Adipic acid Hexamethylene -(n-1)H₂O Nylon-6, 6 diamine

(2) Nylon-6,10 (Nylon-six, ten)

It is obtained by condensation polymerisation of six carbon atoms (hexamethylenediamine) and a dibasic acid with 10 carbon atoms (sebacic acid).

These fibres are light, very strong, flexible and elastic having retain creases and drip dry property. These polyemrs are inert towards biological and chemicals agents. These polymers can be blended with wool to make carpets, garments, tyre cords, ropes, etc.

(3) Nylon-6 (Perlon L)

A polyamide closely related to nylon is known as perlon L (Germany) or Nylon–6 (USA). During prolonged heating of caprolactum at 260–270°C. It is formed by self condensation of a large number of molecules of amino caproic acid. Since, caprolactum is more easily available, it is used for polymerization, with is carried out in the presence of H_2O that first hydrolysis the lactam to amino acid. Subsequently, the amino acid can react with the lactam and the process goes on and onto formed by polyamide polymer.

Caprolactam is obtained by Backman

Concept Ladder

Nylon fibres are stronger than natural fibres and so are used in making cords and ropes.

Previous Year's Question

Nylon-6, 6 is a polyamide obtained by the reaction of [AIPMT] (1) COOH(CH₂)₄COOH + $H_2NC_6H_4NH_2$ (p) (2) COOH(CH₂)₄COOH + $NH_2(CH_2)_6NH_2$ (3) COOH(CH₂)₄COOH + $NH_2(CH_2)_4NH_2$ (4) COOHC₆H₄COOH (p) + $NH_2(CH_2)_6NH_2$

rearrangement of cyclohexanone oxime.

Polytetrafluoroethene (PTEE) or Teflon

Teflon is manufactured by heating tetrafluoroethene in presence of peroxides or ammonium persulphate catalyst at high pressures.

n $F_2C = CF_2 \xrightarrow{(NH_4)_2S_2O_8} [-F_2C - CF_2 -]_n$ Tetrafluoroethene

Polyacrylonitrile (PAN) or Orlon

During Polymerisation of acrylonitrile in the presence of a peroxide catalyst gives polyacrylonitrile.

$$n \operatorname{CH}_{2} = \operatorname{CH} \xrightarrow{\operatorname{Peroxides}} \left(\operatorname{-CH}_{2} - \operatorname{CH}_{-} \right)$$

Melamine-formaldehyde resin

This resin is formed by condensation polymerization of melamine and formaldehyde.

It is a quite hard polymer and is used widely for making plastic crockery under the name melamine. The articles made from this polymer do not break even when dropped from considerable height.

Rack your Brain

Why the articles made from melamine do not break even if they fall from several heightes?

Concept Ladder



PAN is used as a substitute for wool in the manufacture of orlon and acrilan fibres which are used for making clothes, carpets and blankets.



Sol. Bakelite is a polymer formed by the monomers of phenol and formaldehyde. This phenol-formaldehyde resin is a thermosetting polymer.

Bakelite

Phenol-formaldehyde resins are obtained by the reaction of phenol and formaldehyde in the presence of either an acid or a basic catalyst. It starts with the initial formation of ortho and para hydroxymethyl phenol derivatives, which further react with phenol to form compounds where rings are joined to each other with –CH2 groups. This reaction involves the formation of methylene bridges in ortho, para or both ortho and para positions. Linear or cross- linked materials are obtained depending on the condition of the reaction.



The bakelite is prepared by the reaction between

[AIPMT]

- (1) Phenol and formaldehyde
- (2) Tetramethylene glycol
- (3) Urea and formaldehyde
- (4) Ethylene glycol



Polyesters

Dacron is a common polyester, prepared using ethylene glycol and terephthalic acid. The reaction is carried out at 140°C to 180°C in the presence of zinc acetate and Sb₂O₃ as catalyst.

The dacron is crease resistant and has low moisture absorption. It has high tensile strength. It is mainly used in making wash and wear gamets, in blending with wood to provide better crease and wrinkle resistance.

Molecular mass of polymer.

Normally, a polymer contains chains of varying lengths and therefore, its molecular mass is always expressed as an average. In contrast, natural polymers such as protein contains chain of identical length and hence, have definite molecular mass.

Number average molecular mass (\overline{M}_n)

$$\overline{M}_{n} = \frac{\sum N_{i}M_{i}}{\sum_{i}N_{i}}$$

Here $N_{\rm i}$ is the number of molecules of molecular mass $M_{\rm i}$



Definition

The ratio of the weight and number average molecular masses (\overline{M}_w / \overline{M}_n) is called Poly Dispersity Index (PDI).

The number average molecular mass and mass average molecular mass of a polymer are 30,000 and 40,000 respectively. the PDI of the polymer is :

Sol. PDI (Polydispersity Index) is the ratio of the mass average molecular mass to the number average molecular mass.

Therefore, PDI =
$$\frac{40000}{30000}$$
 = 1.33

Thus, PDI of the polymer is >1.

Here N_i is the number of molecules of molecular mass M_i . Methods such as light scattering and ultracentrifuge depend on the mass of the individual molecules and yield weight as average molecular masses. \overline{M}_n is determined by osmotic pressure measurement (colligative properties).

Some natural polymers, which are generally monodispersed, the PDI is unity (i.e. $\overline{M}_w = \overline{M}_n$).

In synthetic polymers, which are always polydisperse, PDI > 1 because \overline{M}_w is always higher than \overline{M}_n .

Biodegradable Polymers:

Many polymers are quite resistant to the environmental degradation processes and are thus responsible for the accumulation of polymeric solid waste materials (environmental polution). These solid wastes cause acute environmental and health problems and remain undegraded for quite a long time (100 — 1000 years). For general awareness and concern, the problems created by the polymeric solid wastes, certain new biodegradable synthetic polymers have been designed and developed. These polymers contain functional groups similar to the functional groups present in biopolymers.





Which of the following is a biodegradable polymer?

- [AIIMS]
- (1) Cellulose
- (2) Polythene
- (3) Polyvinyl chloride
- (4) Nylon-6

4 Identify the type of polymer. —A—A—A—A—A—A—A—

Sol. Here the monomer is –A-. Since the same or identical monomer is being repeated to form this above polymer, so this is a homopolymer. ('Homo' means same or identical)

By far the most important class of biodegradable polymers are aliphatic polyesters and polyamides. Some of these are discussed below :

(i) Nylon-2-Nylon-6

It is an alternating polyamide of glycine (containing two carbon atoms) and ϵ -aminocaproic acid or 6-aminohexanoic acid. (containing six carbon atoms)



Nylon-2-Nylon-6

(ii) PHBV

It is a thermoplastic co-polymer of 3-hydroxy-butanoic acid and 3-hydroxypentanoic acid in which the two monomer units are connected by ester linkages.

n HO-CH-CH₂-COOH + n HO-CH-CH₂-COOH $\xrightarrow{\text{Polymerization}}$ \downarrow CH₃ CH₂CH₃

3-Hydroxybutanoic acid 3-Hydroxypentanoic acid

$$\begin{bmatrix} O-CH-CH_2-C-O-CH-CH_2-C\\ I\\ CH_3 & O\\ CH_2CH_3 & O \end{bmatrix}_n + (2n-1) H_2O$$



Identify the type of polymer. —A—B—B—A—A—A—B—A—

- **Sol.** Monomers 'A' and 'B' are being joined in a random manner in the same polymer chain. It is the example of a copolymer. Copolymers are formed by the mixture of two or more unsaturated monomers reacts to each other.
 -) 6 Why are rubbers called elastomers?
- **Sol.** Rubbers can be stretched by applying mechanical forces and come back to its original position after the removal of applied force. So they are called elastomers.
 - 7 Can the enzyme be called a polymer?
- **Sol.** The enzyme are the catalysts which enhances the biological (bio-chemical) reactions. Enzymes are made up of proteins (monomeric units are amino acids). Thus, enzymes are also polymers.
 - Can nucleic acids, proteins and starch be considered as step growth polymers?
- **Sol.** Step growth polymers are condensation polymers. Proteins, starch and nucleic acids can be considered as step-growth polymers. Those polymers in which monomers possess bifunctional or trifunctional functional groups can combine by the loss of a simple molecule.
 - 9 To have practical applications why are cross-links required in rubber?
- **Sol.** In order to increase the quality of rubber, cross-linking is done which helps to increasing elastomeric properties and thermal stability.



- **Sol.** cis-polyisoprene has a coiled structure and does not have a straight chain. It can be streched like a spring. On stretching, the molecules become partially aligned with respect to each other and on withdrawing the force, the chains come back to their original coild state.
 - Q11 What is the structural difference between HDP and LDP? How does the structure account for different behaviour and nature, hence the use of a polymer?
- Sol. Low density polymer is a low density structure which has branched structure and high density polymer is a high-density polymer which has a linear structure. Low density polymer is a lower melting point and flexible whereas high density polymer is chemically inert and has a higher melting point.
 - Q12 Assertion: Rayon is a semi-synthetic polymer and is taken as a better choice than cotton fabric.
 Reason: Mechanical and aesthetic properties of cellulose can be improved by acetylation.
- Sol. Assertion and reason both are correct statements and reason explain the assertion.

Q13 Which factor imparts crystalline nature to a polymer like nylon?

Sol. One molecule of Nylon can join another by intermolecular H –bonding which is present between two terminal groups –C=O and –NH. Since intermolecular H – bonding is relatively strong, it can lead to close packing of chains that impacts the crystalline character of Nylon.

14 Name the polymers used in laminated sheets and give the name of monomeric units involved in its formation.

Sol. Urea-formaldehyde resin is used in laminated sheets. For the formation of laminated sheets, urea and formaldehyde are the monomeric units.

Q15 Which of the follownig polymer is stored in liver of animals? (1) Amylose (2) Cellulose (3) Amylopectin (4) Glycogen

Sol. (4)

Glycogen is a multibranched polysaccharide of glucose that serves as a form of energy storage in animals.

Glycogen is the analogue of starch, a glucose polymer in plants, having a siimilar structure of amylopectin, but more extensively branched and compact than starch. Starch is polymer of α -glucose and consist of two components anylose and amylopectin.

Amylose is water soluble component which constitutes about 15-20% of starch. Amylose and amylopectin is involuble in water and constitutes about 80-85% of starch.

Amylose and amylopectin are the main storage polysaccharide of plants. Cellulose occurs exclusively in plants and it is the most abundant organic substance in plant kingdom. it is a predominant constituent of cell wall of plants cells.

Q16 Natural ruberr is a polymer of (1) cis-isoprene (3) both (1) and (2)

(2) trans-isoprene(4) None of the above

Sol. (1)

Natural rubber is a polymer of cis-isoprene, [2-methyl-1, 3-butadiene].

(1) teflon (2) polystyrene	017 A con	densation polyme	r among t	he following poly	/mers is
	(1) tet	flon		(2) polystyrene	
(3) PVC (4) dacron	(3) PV	/C		(4) dacron	

Sol. (4)

The polymer formed by the condensation polymerisation is known as condensation polymer. Dacron (terylene) is a condensation polymer. It is formed by the condensation polymerisation of terephthalic acid and ethylene glycol.

- 1. Have high molecular mass and formed by union of monomers.
- 2. Chain growth polymers or addition polymers are formed by successive addition of monomers without loss of simple molecules like H₂O, NH₃ etc.
- 3. Step growth or condensation polymers :

these are formed through series of independent steps. Each step involves condensation between two monomers leading to formation of polymers. More than one monomer unit is involved.

4. Homopolymers :

Chapter Summary

Polymers made of same monomer.

5. Copolymers :

Polymers made of different types of monomers.

6. Elastomers :

The strands of polymer are held together by weak inter molecular forces (van der wall's). e.g. - **Vulcanised rubber.**

- 7. Fibres are held together by hydrogen bonds. e.g. nylon, polyester, polyamide.
- 8. Thermoplastics can be easily moulded on heating. They don't have cross-links, e.g.- polyethene.
- **9.** Thermosetting polymers have cross links, cannot be remolded on heating. e.g. Bakelite.
- **10. Plastisizer** are high boiling esters which are added to plastic to make it soft and rubber like.

SOME COMMON ADDITION POLYMERS/ CHAIN GROWTH POLYMER (HOMOPOLYMERS)				
S. No.	Name(s)	Formula	Monomer	Uses
1.	Polyethylene (low density (LDPE))	-(CH ₂ -CH ₂) _n -	CH ₂ =CH ₂ (ethylene)	film wrap, plastic bags
2.	Polyethylene (high density (HDPE))	-(CH ₂ -CH ₂) _n -	CH ₂ =CH ₂ (ethylene)	electrical insulation bottles, toys
3.	Polypropylene (PP) different grades	СН ₃ [-СН—СН ₂]_	CH ₂ =CHCH ₃ (propylene)	similar to LDPE carpet, upholstery
4.	Poly vinyl chloride (PVC)	СІ [-СН—СН ₂]_	CH ₂ =CHCl (vinyl chloride)	pipes, siding, flooring
5.	Poly(vinylidene chloride) (Saran A)	Cl -CH-CH ₂ -CH Cl	CH ₂ =CCl ₂ (vinylidene chloride)	seat covers, films
6.	Polystyrene (PS) (Styron)	[−CH−CH ₂] _¬	CH ₂ =CHC ₆ H ₅ (styrene)	toys, cabinets packaging (foamed)
7.	Polyacrylonitrile (PAN, Orlon, Acrilan)	N C I I-CH−CH₂]_	CH ₂ =CHCN (acrylonitrile)	rugs, blankets clothing
8.	Polytetrafluoroethyl- ene (PTFE, Teflon)	-(CF ₂ -CF ₂) _n -	CF ₂ =CF ₂ (tetrafluoroethyl- ene)	non-stick surfaces electrical insulation

OME COMMON ADDITION POLYMERS/ CHAIN GROWTH POLYMER (HOMOPOLYMERS)

9.	Poly(methyl methacrylate) (PMMA, Lucite, Plexiglas, perspex)	-[CH ₂ C(CH ₃) CO ₂ CH ₃] _n -	CH ₂ =C(CH ₃) CO ₂ CH ₃ (methyl methacrylate)	lighting covers, signs skylights
10.	Poly(vinyl acetate) (PVAc)	-(CH ₂ - CHOCOCH ₃) _n -	CH ₂ =CHOCOCH ₃ (vinyl acetate)	latex paints, adhesives
11.	Polyisoprene cis form Trans-form (Natural rubber) gutta-purcha (1,1-addition) $-H_2C$ CH_2 H H C H	-[CH ₂ - CH=C(CH ₃)- CH ₂] _n -	CH ₂ =CH- C(CH ₃)=CH ₂ (isoprene)	requires vulcaniza- tion for practical
12.	Polychloroprene (cis + trans) (Neoprene)	-[CH ₂ - CH=CCl-CH ₂] _n -	CH ₂ =CH-CCl=CH ₂ (chloroprene)	synthetic rubber oil resistant

	Some Condensation POLYMERS (Step Growth Polymers)					
S. No.	Name(s)	Formula	Monomer	Uses		
1.	Polyester/ Dacron/ Terylene/Mylar		HO ₂ C C ₆ H ₄ CO ₂ H (Terephthalic acid) HO– CH ₂ CH ₂ –OH Ethylene glycol	Copolymer Step growth, Linear Polymer		
2.	Glyptal or Alkyds resin		HO ₂ C–C ₆ H ₄ –CO ₂ H (Phthalic acid) HO–CH ₂ CH ₂ –OH	Copolymer Step growth, Linear Polymer		

3.	Polycarbonate Lexan		$(HO-C_6H_4-)_2$ $C(CH_3)_2$ (Bisphenol A) $X_2C=O (X = OCH_3$ or Cl)	Copolymer Step growth, linear Polymer
4.	Polyamide (Nylon 6,6)	~[CO(CH ₂) ₄ CO-NH(CH ₂) ₆ NH] _n ~	$HO_{2}C-(CH_{2})_{4}-CO_{2}HH_{2}N-(CH_{2})_{6}-NH_{2}$	Copolymer Step growth, Linear Polymer
5.	Nylon 6,10	O -(-C-(CH ₂) ₆)-C-NH-(CH ₂) ₆ -NH-)-	$HOOC-(CH_2)_8-$ COOH $H_2N-(CH_2)_6-NH_2$	Copolymer Step growth, Linear Polymer
6.	Bakelite	OH OH CH ₂ -CH ₂ -CH	PhOH + HCHO in (excess)	Copolymer Cross-linked polymer Step growth
7.	Urea- formaldehyde resin	(-NH-CO-NH-CH ₂ -) _n	H ₂ N–CO–NH ₂ (Urea) HCHO (Formaldehyde)	Copolymer Step growth Cross-linked polymer
8.	Melamine formaldehyde resin	$ \begin{array}{c} \begin{array}{c} \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\$	Melamine + Formaldehyde	Cross-linked polymer

Some Useful Copolymers				
Monomer A	Monomer B	Copolymer	Uses	
H ₂ C=CHCl	H ₂ C=CCl ₂	Saran	films & fibers	
H ₂ =CH-CH=CH ₂ 1, 3-Butdiene		Buna Rubber (Homo-polymer)	In manufacture of tyres, hoses, etc.	
H ₂ C=CHC ₆ H ₅	H ₂ C=CH- CH=CH ₂	SBR styrene butadiene rubber (Buna-S)	Tires	
H ₂ C=CHCN	H ₂ C=CH- CH=CH ₂	Nitrile Rubber (Buna-N)	Adhesives hoses	
H ₂ C=CHCN	$H_2C=CH-CH=CH_2 + H_2C=CHC_6H_5$	ABS rubber	Fuel tanks	