

→ Cross-linked polymers are insoluble and infusible. Depending on nature & frequency of cross-links, such polymers may show different orders of swelling in solvent.

Ex: PF Resin, epoxy resins, vulcanised rubber etc.

Q → why branch polymer is soluble but linear polymer is not soluble.

Ans → Branched polymers are readily soluble & fusible than linear polymers of comparable chain length or m.wt

(5) ON THE BASIS OF APPLICATION and physical properties.

① Rubber → These polymers are characterized by long-chain elasticity. They are rather weak, dimensionally unstable and undergo high elongation even on application of low stress.

They exhibit tensile strengths in the range 300-3000 Psi.

Exa: Natural Rubber, and synthetic rubbers.

→ They are also known as elastomers. The cross linking of rubber is commonly known as vulcanization.

② Plastics : These are polymers which can be shaped into hard and tough utility articles by the application of heat & pressure.

* These are usually much stronger than rubber. Some of them are hard, horny rigid, stiff. Plastics exhibits tensile strength ranging b/w 4,000 - 15,000 psi.

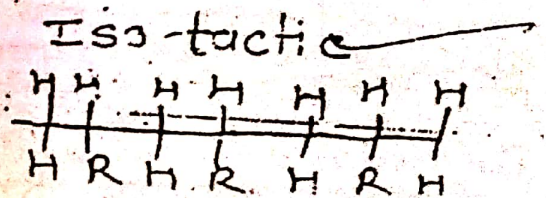
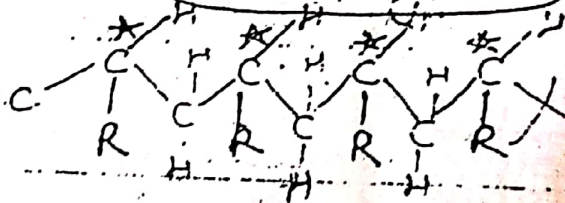
Exo: Polyethylene, polypropylene, polystyrene. PVC, PF & VF resins, polycarbonates.

(c) fibre: They are strongest of the three diff. types of polymers.

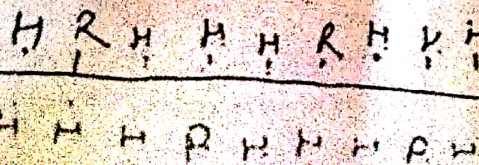
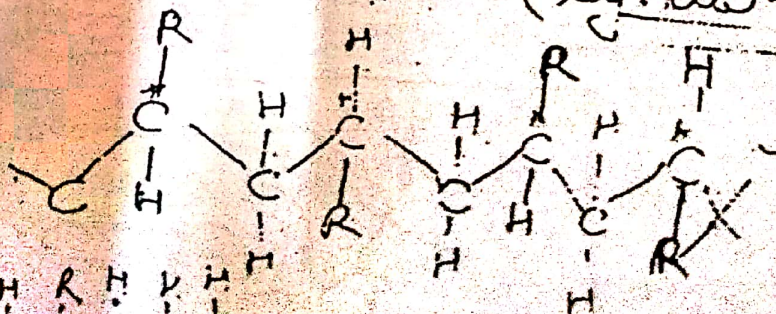
Exo: Rayon, Polyester, silk.

6. ON THE BASIS OF TACTICITY ⇒

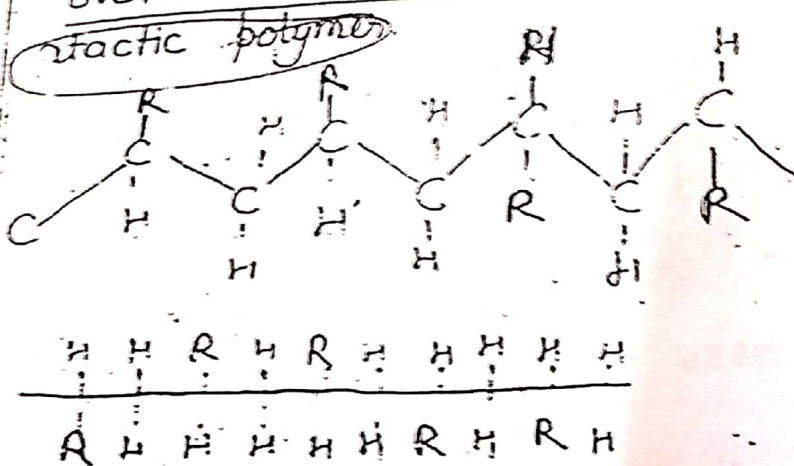
(i) Iso-tactic: Polymers where in all adjacent asymmetric carbon atoms over the chain length of the polymer chain have the same space configuration are known as Isotactic polymers.



(ii) Syndio-tactic ⇒ Polymers in which chain is made up of units having opposite space configuration of each symmetric carbon atom are known as Syndio-tactic polymers.



~~(ii) Syndiotactic~~
 (iii) Atactic polymer $\Rightarrow$ Polymer in which
 gps are arranged randomly in space
 over the chain are known as



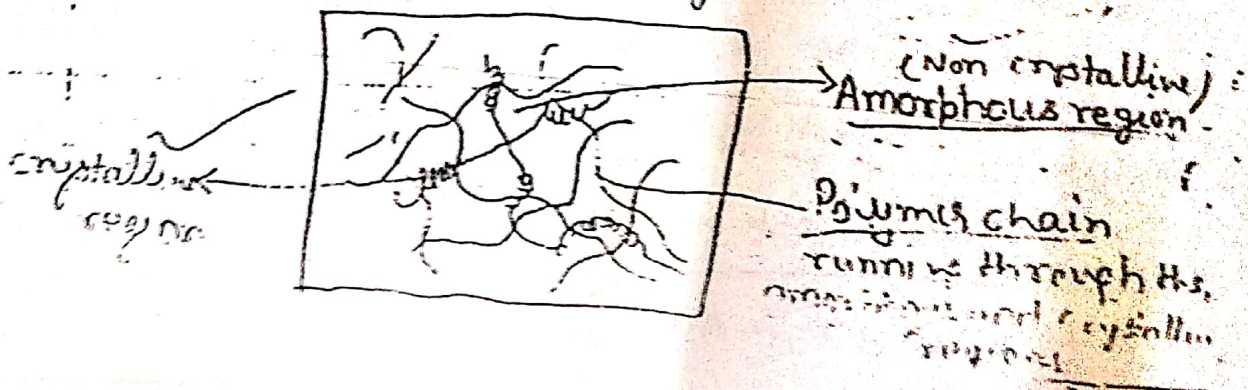
⑦ ON THE BASIS OF CRYSTALLINITY $\Rightarrow$

When polymers cooled from molten state they exhibit diff. tendency to crystallize at different rates depending on many factors.

① Crystalline $\Rightarrow$ Polymer showing a high extent of crystallinity more than 60% are commonly known as crystalline polymers.

② Semicrystalline $\Rightarrow$ Those showing a significant but relatively lower degree of crystallinity are termed as semicrystalline polymers.

③ Non crystalline $\Rightarrow$ No crystal formation



2008 (2/12) - Chemical
 (8) ON THE BASIS OF composition $\Rightarrow$ On this basis the polymers can be classified as-

(i) Homo-polymer $\Rightarrow$ When a polymer is obtained from a single monomer unit it is called homopolymer e.g. polystyrene which is formed from monomer styrene.

$\rightarrow$ Some other examples of homopolymers are polyethylene, polypropylene.

With some exceptions, polymers made in chain reactions often contain only carbon atoms in main chain (homochain-polymer).

(ii) Co-polymers $\Rightarrow$ When a polymer contains two and more monomer units in the chain then it is known as a Co polymer.

It is possible to designate copolymers as

(a) Random copolymers $\Rightarrow$ when monomer units are arranged in an irregular manner, the co-polymer molecule is called Random co-polymer.
 Ex: $-A-B-B-A-B-A-B-A-A-B-$

(b) Alternate Co-polymers $\Rightarrow$ when monomer units in a copolymer molecule are arranged in the chain in regular manner, the molecule is called alternate - copolymers.

Ex: $-A-B-A-B-A-B-A-B-A-$

(c) Block-Copolymers $\Rightarrow$ Linear copolymers in which the like monomeric units occur in relatively long sequences are called block co-polymers.

Exo: -A-A-A-B-B-B-B-A-A-A-B-B-B-B-

Block copolymers

Block copolymers may be produced by the following four tech.

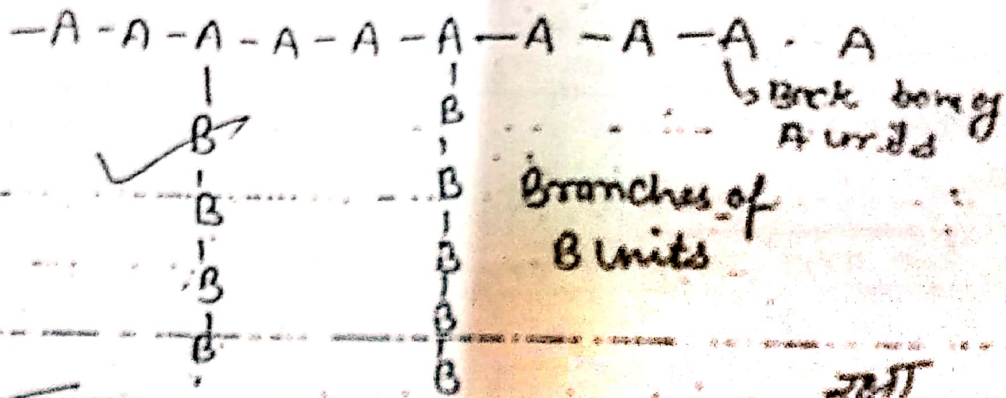
1. Utilise reactive end groups

2. Activate end groups

3. Fragment a chain to produce active end groups

4. Transport reactive end groups into a different media

* (d) Graft Copolymer $\Rightarrow$ These are branched copolymers in which the branches have different chemical structure to that of main chain.



* The importance of the block and graft copolymer is that the resultant material tends to exhibit the properties of each homopolymer.

* They also have same unique property arising from the chemical linkage b/w the homopolymer sequence.

* Ex: Pure polystyrene is quite brittle, whereas polymerisation of about 5% of Rubber produced material which is strong and tough.

$\rightarrow$ 90 part of vinyl chloride & 10 part of vinyl acetate, the random co-polymerisation