POLYMER BLENDS or more different polymers or copolymers that are not dinked by covalent bonds having better properties than individual component. Ineparation of Polymer Blends! - following methods can be used! -9 - For amorphous and semicrystalline polymers, mixing of Pops polymers is done at temp above Tg or Tm resp. This mixing gives pyechamical Blends. b- Polymers are dissolved in common solvents & their solvent is removed, trques solution-cast Blands. c-fine dispersion of polymere in water (laters) are mixed. I then mixed polymers are coagulated, it gives later Blands. d-Cross-linked polymer is swollen with different monomer, then monomer is polymerised & cross-linked. It gives Inter-Penetrating Polymer Networks (IPN) or chemical blends. For making a blend of polymers, it is essential that they are compatible with each other i.e. both the polymers should be triendly in it. be friendly in structure. Ex: (i) Iwo hydrocarbon polymers are blended (polyethique + prolypropylene) (ii) Wood (polyamide) + Jerylene (polyester) as they have O' atoms & similar structural parts (coo. & -cont)

(iii) Cellulose nitrale & polyalyeofs are possible to blend. (iv) Polyestei + cotton (cellulose fibre) can be blended. classification of Polymer Blends! Ta classification can be on the basis of preparation of folyblands: - nechanical blunds, Chemical Blands, Later Blands and Solution-cast Blands. (b) or on-the basis of compatibility.

Miscible Bands

- usually clear Dumiscible Blends -> usually opaque -> Exhibit a single glasstranstron -> Exhibit-seperate Tgs of individual components no phese characteristics of each component under stress. solid becomes soft upon heating - temp. at which an amorphous solid becomes soft uponto or brittle upon cooling. The G.T. temp is lower than MP. of its crystalline form it it has one )

Properties of Polymen blends! -A polymer blend components as seperate phasewhen Viewedunder microscope. Blw polymeric chains, only rander waals forces, dipole-dipole into the chains. dipole-dipole interactions of H-bonding exist. Insperties of a polymer blend are closely related to the properties of individual components A polymer blend can be be synergistic for any property l'e better than the weighted average) or Non-syneigistic because of unfavourable dipole-dipole attractions or disruption of such attractions between components -> Blending usually improve properties like processes ability, Impact strength, abrasion resistance etc. Degradation of us sensitive polymers can be protected by blending e.g. PMMA is degraded by gamma radiations but when it blends with strene-acrylonihile copolymer, -> Polyester (high fensile strength, crease resistant, durable but does not absorb sweat easily & gets heated by sunlight) is Hended with cotton to make it cooler. Blend of natural subber & synthetic compatible subber. Natural kubber - weak, too much clastic water absorbing, 2 tacky Butadiene subber - proprides better properties with faulty properties minimized. A crylonitride sysème subbent Butadiene name plates other such things where reflecting surface is required. (Homopogner + copogner) - for making sports equipments & transport entrainers. (Homopolymen+ Homogodymen)

Mechanical / physical properties of homogenous polymer-blends can be predicted using the following semi- empirical P -> particular properly V- volume fraction in the P=P,V,+P,V2+IV,V2 nu'xture, I > Interaction term (may bet ve, -ve or zero) When ID=0 P = P1 V1 + P2 V2 i.e-Properties are strictly additine when I = tre (positive term) The property in this ease is better than the weighted on average and the blend is paid to be Synergistic for average and the blend is paid to be Synergistic for that properly. Results in very favourable dipole-dipole attraction blo polimer components. attraction bjw polymer components! When I = -ve (Negative term) p=P,V,+P2V2-2V,V2, The property is worse than the weighted average (Non-syneigistie). Graph for property Vs volume fraction Non-syneigisin arise when a favourable intérmolecular interaction is disrupted. volume paction -) The Tg's of miscible blends simple rule of mixtures equationi- Typ Tg = w, Tg, +w2 Tg 2 ) in K.

w -> wt practions, Tg -> in K.

polymer % of polymer B

hiokols (T) Polysulphides or Polythioethers -Preparation! - They are typically the liquid polymers that can be crosslinked by converting—the terminal thiol groups (-SH) of the polymer to disulphide (-S-S-) links: n cley cycl + n Nas -s Na -cy-cy-is-stn Sodium Polysulphide 1, 2-dich loro ethane Thiokof Rubber ) Typical temp. range = -45°C-105°C. > The elastomer has a high sufflux content, approximately 80% by wto, making it a high density material with a high density material resistance to swelling by hydrocarbon site. hydrocarbon oils. on the other hand, causes a strong tendency to relax & flow under messure. & flow under pressure. Applications: The polymer is most used as a low molecular wholiqued that cures in place to form an elastomeric sealant. elastomeric sealant. (2) Primarily used in oil-resistant & weather-resistant seals and gaskets. (3) Also used in gasoline hoses & other applications.

Kevlark (Poly (p. phenylene te & phthalamiole)) reparation -+ HAN-(0)-NH2 1,4-benzeue dicarbocylie acid 1,4-benzeue d'annive (p-phenylene d'annive) et Le poly merisations Copolymer Condentation polymer f 6 20 2-NH-(0)-NH)n Properties -→ Benzene eings are linked to the amide linkage; therefore. Kevlar is a very sicial polimer. As it is an aromater polyamide, il is alsolenewmen as Kevlar has abbility to be sprin into fileres having 5-times the tensile strength of steel. Applications: (a) for making Tyses, brakes, clutch lining & other can parts
(b) Bullet proof vests (c) Safety helinets (d) Aérospace à aircraft industries.