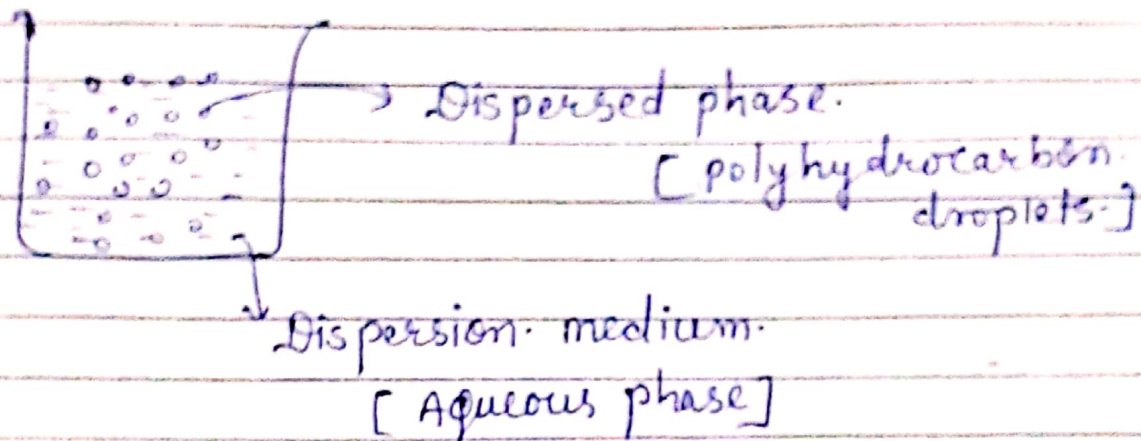


UNIT - I

- Latex :- Latex is emulsion of polyhydro-
Carbon droplets in aqueous solution/medium.
• Latex define as a stable colloidal dis-
-persion of a polymeric particle in an esse-
-ntially aqueous medium.



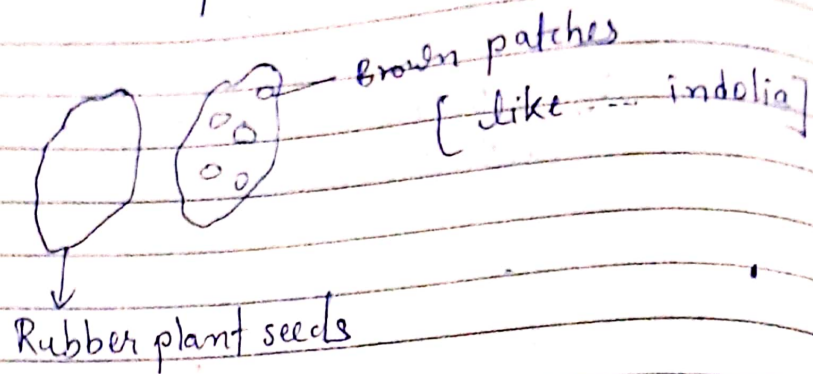
- Like Colloidal Solution.
- It is Colloidal dispersion of negatively charge particle

Natural latex is obtained from - Rubber plant [*Hevea Brasiliensis*]

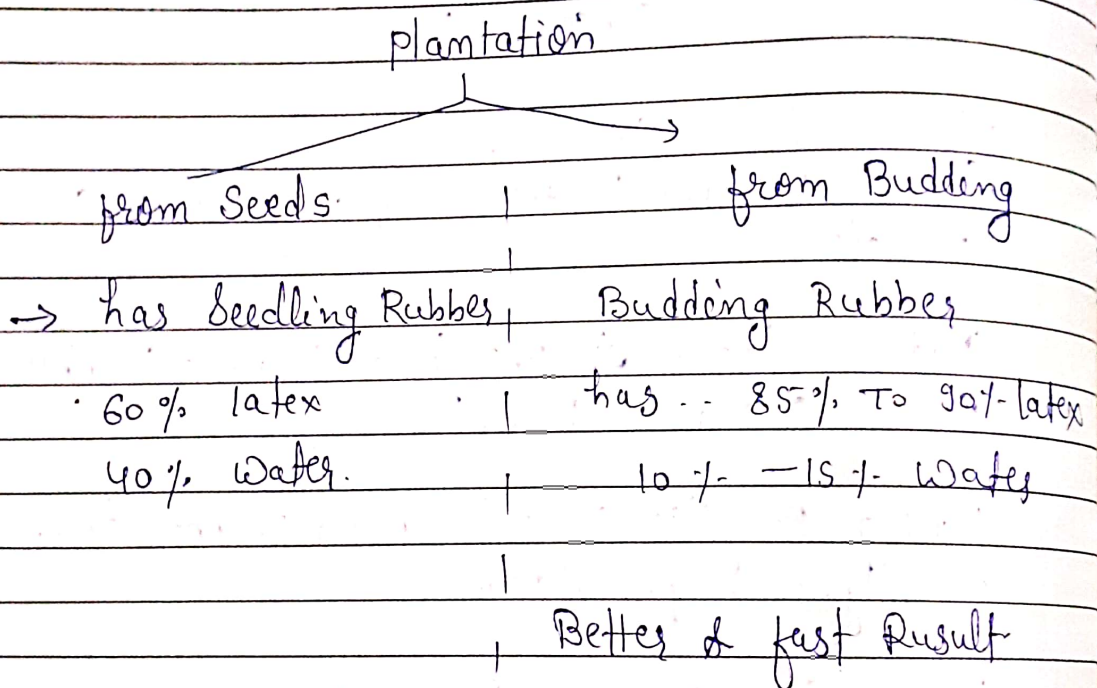
Hevea Brasiliensis → is para rubber tree.
Family - Euphorbiaceae.

Latex, collected from Tree, process is known as Tapping and its sap like extract known as latex, it is primary source of Natural latex, Natural Rubber -

Seeds of plant → like cornucule and usually are globose in shape.



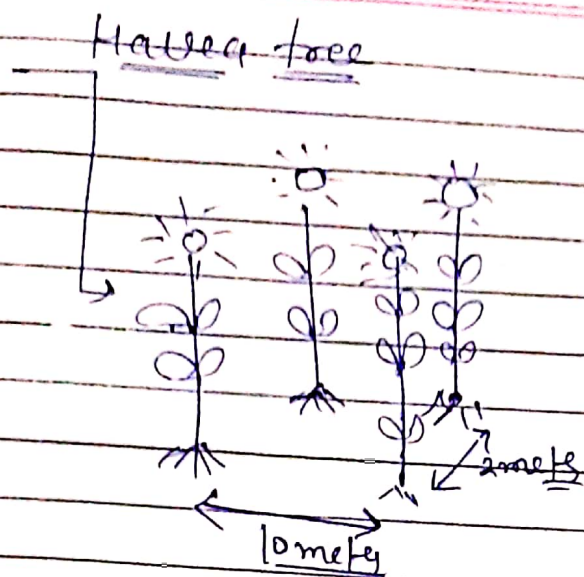
Rubber Tree seeds Complyte germination in 10-15 days,



Rubber tree →

Havea / Guayule / ficus elastation,
Castilloa Elastica, Parthenium argentatum,
Landolphia.

Havea - is most popular Rubber tree



Tapping processing time --- 5 AM to 9 AM

Is right.

- * Fresh latex pH, approximately is 7, The latex coagulates when pH is lowered to around 4.2-5, fresh latex coagulates spontaneously within few hours of leaving the tree, Because Reduction of pH due to degradation of carbohydrates present in latex by Bacteria and hydrolysis of the lipid substance to fatty acid and its interaction with metal ions are the chief reason for this coagulation,

Composition of fresh latex
as follows

	%
(TSC) Total Solid Content	36 %
(DRC) Dry Rubber Content	33 %
Proteins	1-1.5 %
Resins	1-2.5 %
Ash	1 %
Sugars	1 %
Water	Add to 100% (Generally 30-35 %)

Advantage of Latex :-

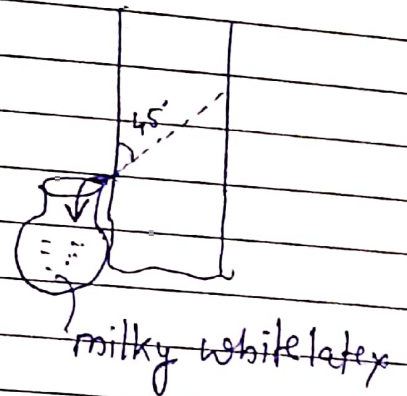
- ① Low Viscosity so Easy to shape
- ② light equipment of Simple Construction for processing
- ③ low energy Requirements,
- ④ No - Heat generation during processing
[ultra - accelerators - low temp. curing
- ⑤ unbroken rubber -- Better physical & Ageing properties

Flow chart of Field Latex to RSS sheet...



Havea Brasiliensis

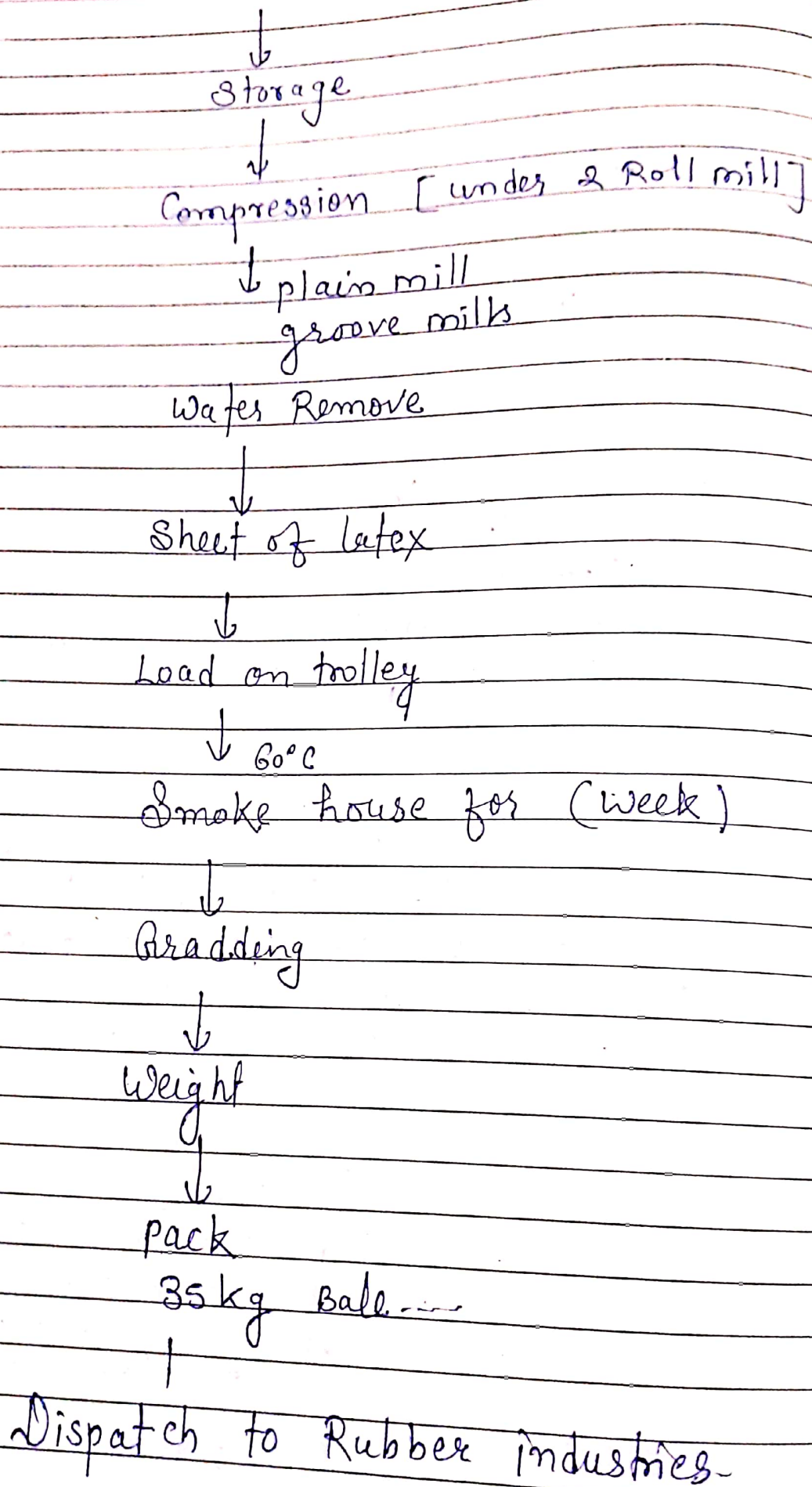
Collect latex
[process known as
Tapping]



Preservation of field latex
Ammonia, Na sulphate,
formalins etc...

Sieving (60 mesh)

Coag Coagulated.
By HCOOH



preservation :-

The latex is to be preserved immediately after collection, against rise in acidity by bacterial putrefaction.

Among the preservatives used, Ammonia is most commonly used owing to certain advantages like ... removal by blowing air or reaction with HCHO

Recently Sodium pentachlorophenate is being used along with small amount of Ammonia...

Common preservatives

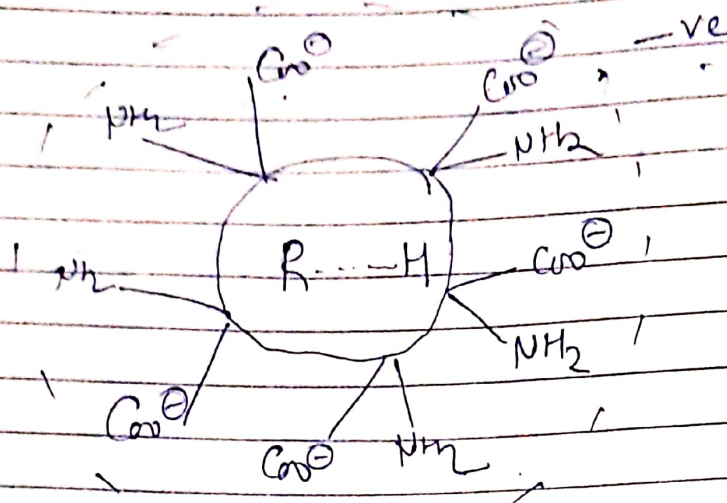
→ NH_3 , potash, formalin, Sodium Sulphite, Boric acid,

Stability of NR Latex :-

Latex consist of fine particles of size 20-20000 nm, and covered by electrical charged.
- Repulsion Between the particle and layer of adsorbed water giving a mechanical barrier,

Reason for instability :-

Mechanical agitation Impingement of two particle - Coalescence (झटका-में मिलना)
Reduction in electrical charge due to pH.
formation of insoluble soap by reaction with metal ions.



Stability of NR latex

- Reduction of free surface energy between the polymer particle interface
- By imparting charge of same polarity.
- By putting some heavy SEETHING layer of tightly bound water molecules. (ଅବଳାହୁଆ)

→ Concentration of latex

In this process eliminates most of the Non Rubber Constituents, This is done by 4 following methods,

- creaming
- centrifugation
- electrodecantation
- evaporation.

(a) Centrifugation :- The Rotating Bowl of the Centrifuge is generally fitted inside

With a number of metallic discs. As the Bowl Rotate at high speed 6000 rpm, The mass of latex is broken up into thin conical shells. The latex is fed to the machine from a constant head to ensure uniform flow. The Cream Centrifuge towards the axis of Rotation.

In which $\rightarrow$ Centrifugal force are used. Centrifugal Latexes are commercially available as high Ammonia (HA; minimum 0.6% Ammonia) and low and low Ammonia type 0.3%.

(b) Evaporation $\rightarrow$

Evaporation latex concentrate is made by heating suitably preserved latex without skin formation on the surface. The stabilizers used are generally KOH soap and/or glue.

A suitable evaporator consist of a jacketed drum rotating along the axis, steam is admitted in the jacket and hot air inside over the latex surface, a hollow cylinder rotating along the drum surface avoid skin formation.

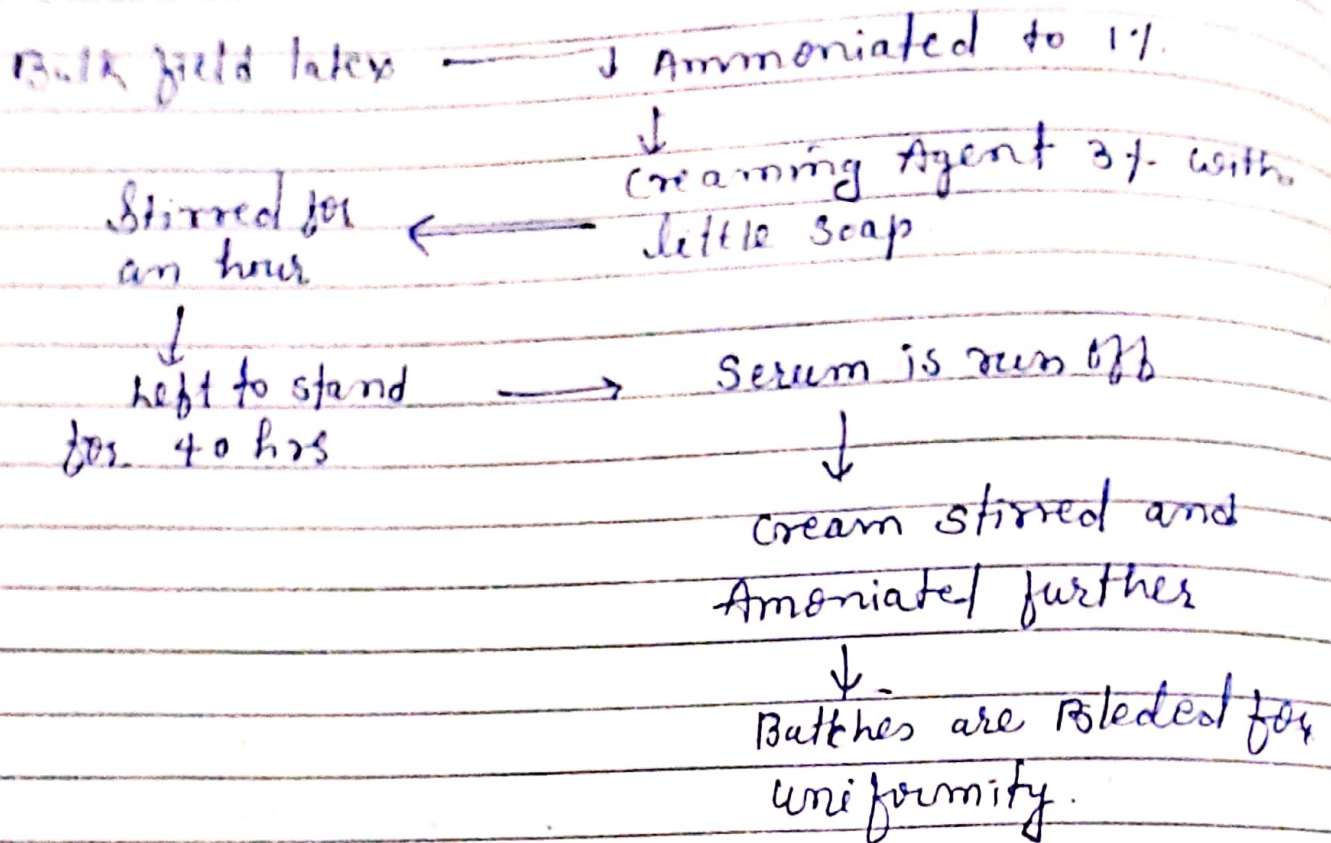
Evaporated latex has high stability & used for adhesive and latex/cement composites.

(c) Creaming

Latex particle are lighter than serum in which they are suspended.

Have tendency to go up on the surface.

Creaming agent Ammonium alginate, tamarind seed gum, Irish moss and locust bean gum, make the particles to form loose aggregates



⑤ Electrodécantation :-

Depends of (-ve) charge on surface of Rubber particle, electric current is passed, Rubber particle move towards the Anode.

Difference between Natural latex and Synthetic latex.

Natural latex	Synthetic latex
→ is more durable	→ is durable
→ is ecofriendly	→ is cheaper
→ is hypoallergenic	→ is hypoallergenic
→ is dust mite resistant	→ is dust mite resistant
→ stay fresh and cool at night	→ stay fresh & cool at night
	→ Support back & joint

Latex grades

Type	Source
① Ribbed Smoke Sheet	Coagulated field latex, [finalization in smoke house]
② pale crepe	Coagulum of Natural latex liquid.
③ Estate Brown Crepe.	Estate cuplumb tree latex
④ Thin Brown Crepe	Cuplumb, wet slab, tree latex unsmoked sheet,
⑤ Thick Blanket crepe	— I —
⑥ plate book crepe	— II —
⑦ pure smoked blanket crepe	Remilled RSS & RSS Cutting.

Natural Rubber :-

Rubber tapping the milky white liquid latex is collected from rubber trees in a cup by making a slight cut in the tree bark. The collected latex is washed, filtered and reacted with acid to coagulate (coagulate) the rubber particles.

It is created by enzymatic process by many plants.

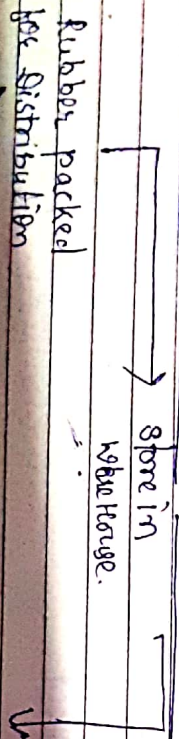
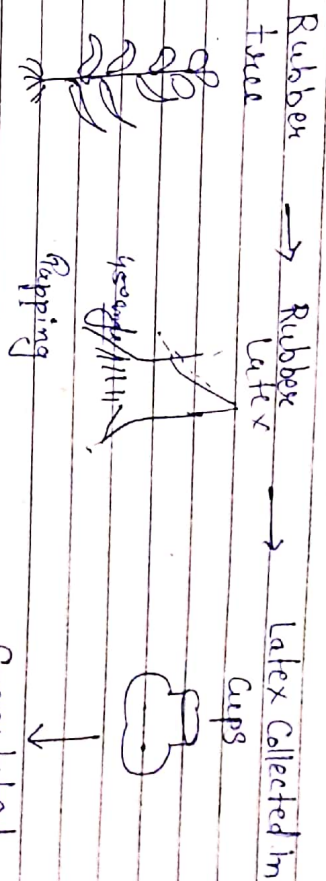
Belonging family → Euphorbiaceae

Climatic Condition →

Avg temp. $25^{\circ}\text{C} - 28^{\circ}\text{C}$

@ 80% Humidity

Outline Manufacturing Diagram of N.R.



Pattetised Rubber Black

Rubber used by factories around world wide.

Lab test



Drying

(over sun)

Shaping

Creeping

Washing

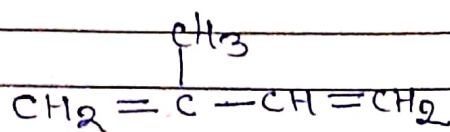
Shredding

(machines)

Area @ World wise $\rightarrow$ [South eastern Asia]

Country $\rightarrow$ Malaysia, Indonesia, China, Sri Lanka, Vietnam, India etc.

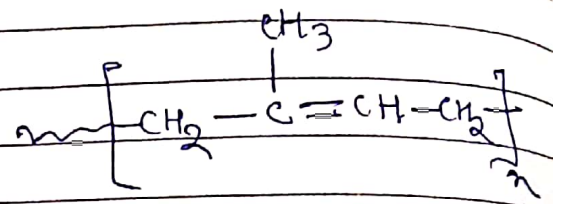
NR - [chemical structure.]



[Isoprene]

2, Methyl 1,3 Butadiene

polymerisation $\rightarrow$



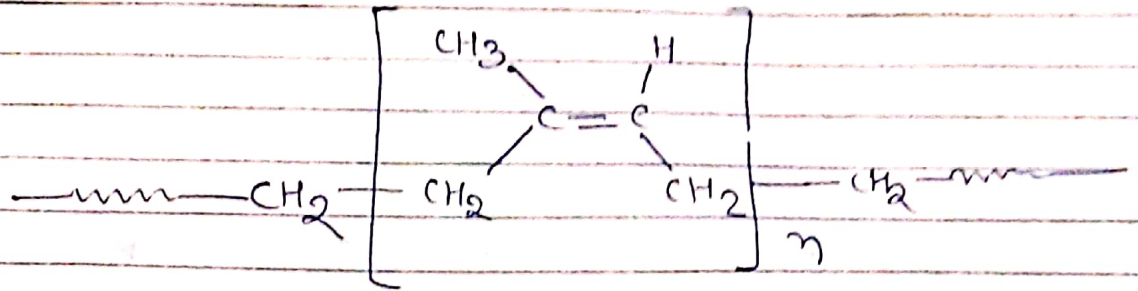
poly isoprene

Importance $\rightarrow$

- (i) unsaturation
- (ii) Non polar
- (iii) Methyl bulky group
- (iv) 99% < Content [poly cis-1,4 isoprene] & some may be bonded also non isoprene structural unit proteins, Amino Acids, phospholipids in Macromolecules Backbone.
- (v) Final packing - sheet form
- Block form.

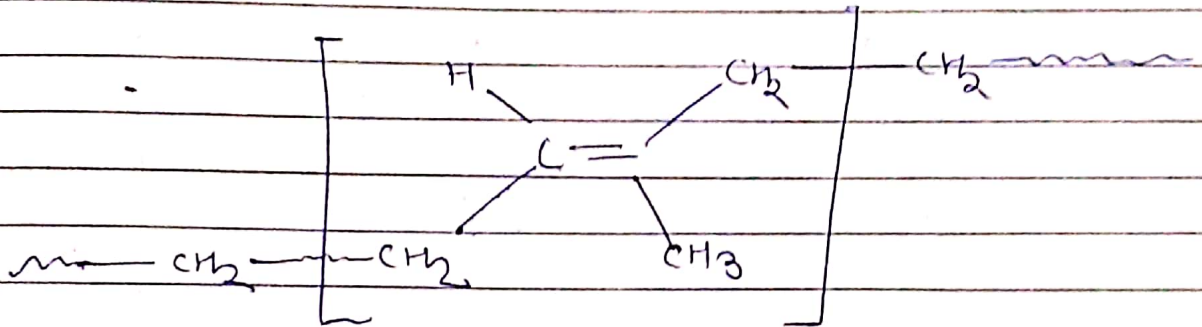
In polyisoprene $\rightarrow$ Trans 1,4 polyisoprene is.
Dominant isomer in Gutta-percha or Balata,

* Poly - cis - 1,4 isoprene



(poly - cis - 1,4 isoprene.)

* Trans - 1,4 isoprene.



(poly Trans 1,4 isoprene.)

⇒ Physical properties of NR

Density = 0.92 gm/cc

Refractive index 1.52 @ 20°C

Molecular Weight 2 lac - 5 lac

[200000 - 500000]

Dielectric Constant = 2.37

Volume Resistivity = 10^{15} ohms/cc

$T_g = -75^\circ\text{C}$

⇒ Advantages : →

poor resistance to cut,

Better , Good Tensile strength.

Technically Specified Rubber (TSR)

TSR was started by Malaysia in 1965 with the introduction of SmR (Standard Malaysian Rubber)

TSR Grade →

SmR L	SmR GP	TSNR-10
SmR WF	SmR 10	TSNR-20
SmR CV	SmR 20	TSNR-50
SmR LV	SmR 50	
SmR S		

Properties	SmR L, WF, CV, LV,	SmR 5	SmR 10,	SmR 20,	SmR 50,
Dist Content @ 40 mesh.	0.03 %	0.05 %	0.10 %	0.20 %	0.50 %
Ash Content	0.50 %	0.60	0.75	1.00	1.50 %
Nitrogen Content (%)	0.60 %	0.60 %	0.60 %	0.60 %	0.60 %
Volatile matter	0.80 %	0.80 %	0.80 %	0.80 %	0.80 %

OTHER TYPES OF NATURAL RUBBER

* Hevea plus MG Rubber →

These are Chemically modified grade of Natural Rubber, prepared by graft polymerization of methyl methacrylate in NR-Rubber Concentrate.

This is available in latex or dry rubber form.

Two Grade are prepared viz, MG30 and MG49 contains 30% and 49% of PMMA respectively.

→ Dry MG Rubber used in Adhesive

→ Latex MG is blended with normal latex to Achieve films with improved modulus & tear,

* ENR [Epoxidized Natural Rubber]

it is produced by reaction of latex with perfo-
mic acid formed in situ.

ENR gives rise to improved air impermeability and resistance to oil

ENR also has good adhesive properties useful for Rubber to metal bonding.

* Thermoplastic Natural Rubber [TPNR.]

it is prepared by Blending of NR and polypropylene / polyethylene & different proportions to give Rubber with different stiffness properties.

Potential application includes footwear, sport goods, seals hoses, bumpers, spoilers and Car body protection strips.

Natural Rubbers are different from Synthetic

Due to is obtained from Natural source ... (tree)
have high gum ; green tack and highest T.S.

Created by Enzymatic process in plant.

SBR

STYRENE BUTADIENE RUBBER.

SBR is World's oldest Synthetic Rubber, it is developed in 1930's,

it is copolymer of styrene and butadiene normally contain 23% bound styrene

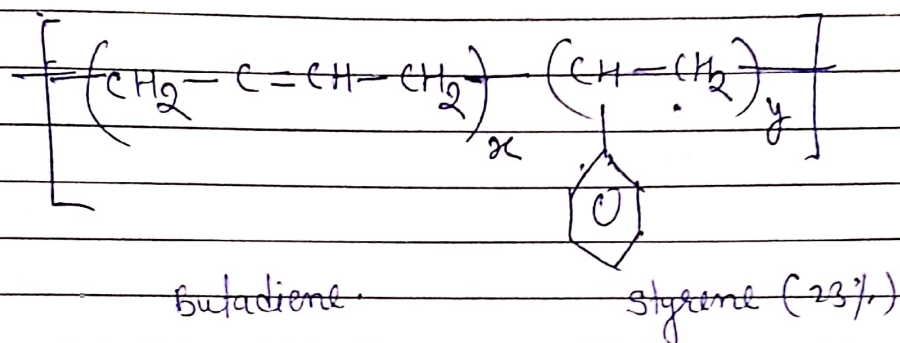
it is prepared by emulsion & solution polymerisation

it is not hardened with storage

It is a Slow Curing Rubber

good Abrasion resistance. (due to hard)

Structure :-

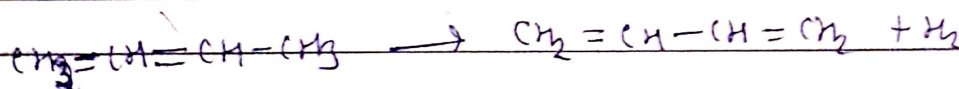
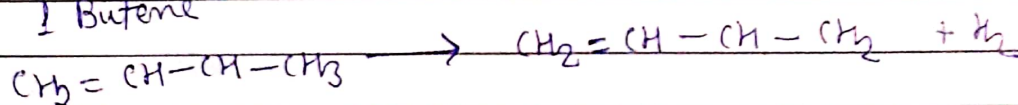


Butadiene production

(1) from n-Butanes.

The n-Butanes are obtained from C₃-C₄ streams from the cracking of petroleum crude propane & isobutane are separated by fractionation and isobutane is removed by absorption in 65% Sulphuric acid

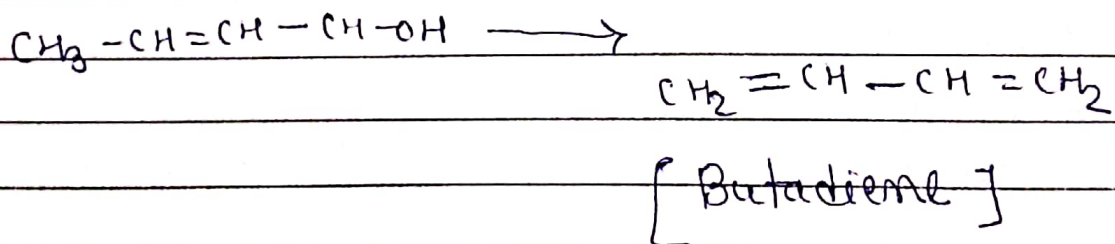
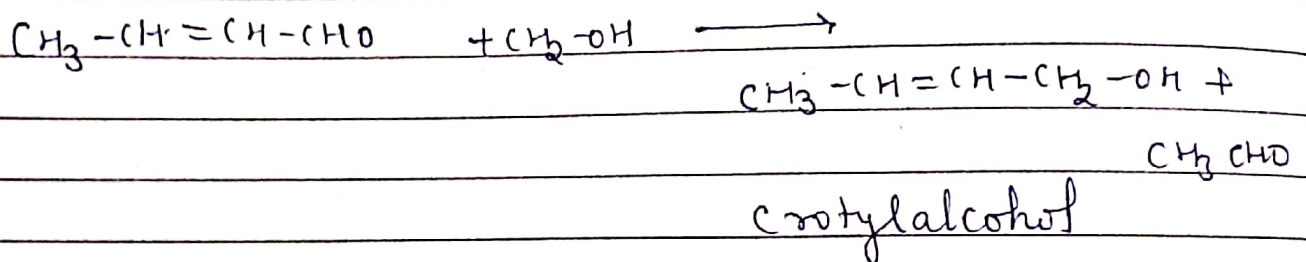
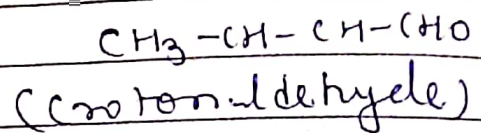
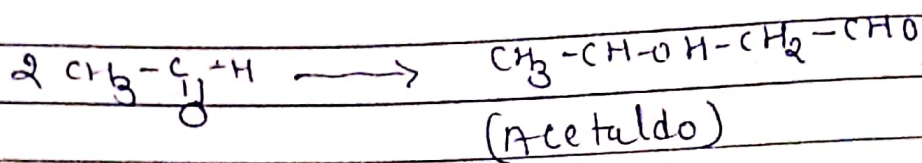
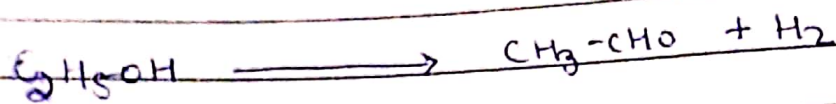
1 Butene



2 Butene

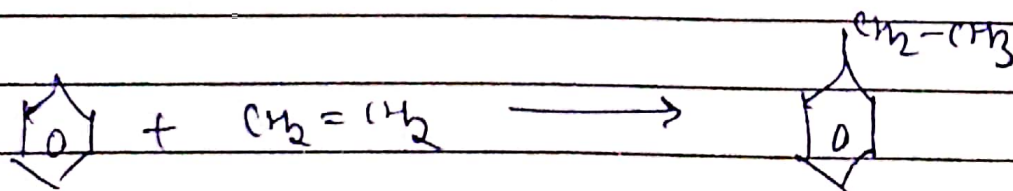
Typically a mixed stream containing 1 Butane & 2 Butene is preheated to 650°C & mixed with Superheated Steam... and finally made. Butadiene

2. production of Butadiene from Ethyl Alcohol.

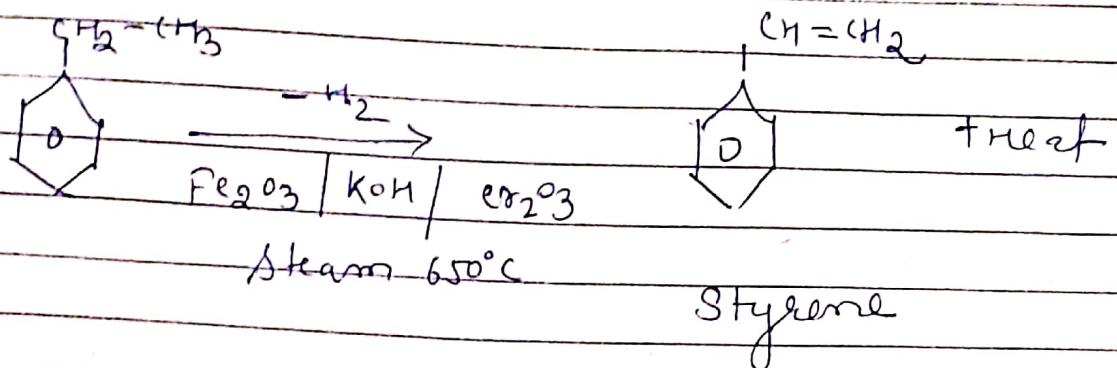


Styrene :-

Styrene is produced from ethyl benzene, which is produced by alkylating Benzene with ethylene



↓
Dehydrogenation of ethyl benzene

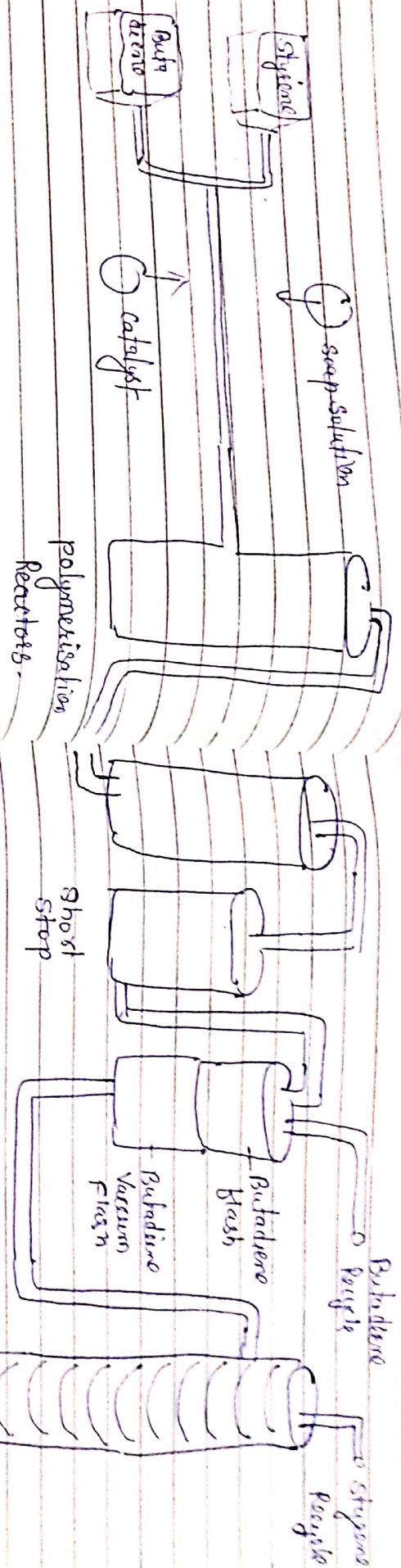


Emulsion polymerisation of SBR.

Emulsion polymerisation is done at two-temperature 50°C and 5°C . The polymer obtained at 50°C is called a hot polymer and the other one is called the cold polymer.

General purpose SBR is manufactured by reaction of aqueous emulsion of butadiene and styrene in the presence of soap solution, polymerisation initiator and regulators to form a high molecular weight copolymer with elastic properties. This can be prepared with technique also, but the properties of a solution SBR slightly varies from that of an emulsion SBR.

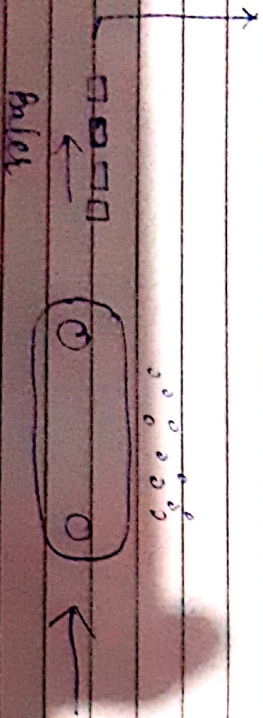
outline manufacturing process --- (P.T.O.)



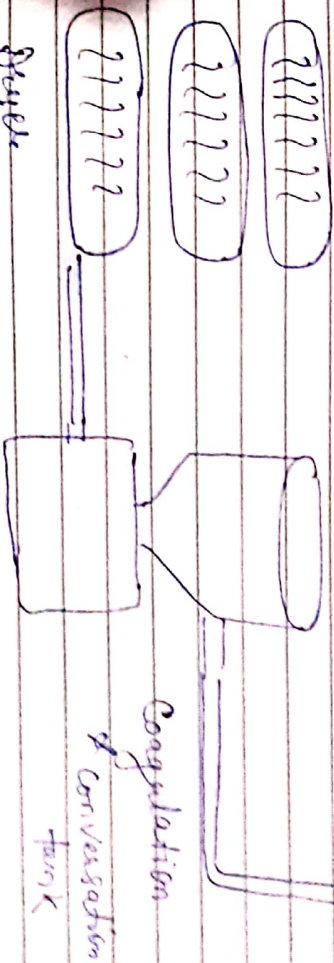
STYRENE
BUTADIENE
RUBBER



Bales



later
Blond



properties of SBR

- (i) T_g of SBR is $\approx -59^{\circ}\text{C}$
- (ii) Density = 0.94
- (iii) Mooney viscosity = 35-55

Advantage of SBR

- (i) Better resistance to reversion
- (ii) higher resistance to abrasion at high speed.
- (iii) Better resistance to ageing specially at high temperature
- (iv) Better resistance to flex fatigue
- (v) Excellent impact strength

Disadvantage of SBR

- (i) Necessity to use reinforcing filler.
- (ii) poor resistance to ozone, sun light.
- (iii) Very little resistance to oils, gasoline, hydrocarbon solvents.

GRADE of SBR Rubber

All Hot & Cold polymerised SBR, with or without oil and or C-Black are classified by IISRP (International Institute of Synthetic Rubber producers.) under the following series,

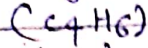
1000 - Hot polymerised non pigmented Rubbers

- 1500 - Cold polymerised non pigmented rubber.
- 1600 - Cold polymerised Carbon Black - oil Rubber master Batches. 14 phr or less of oil.
- 1700 - Cold polymerised oil Rubber master Batches
- 1800 - Cold polymerised C-Black - oil rubber master Batches containing more than 14 phr of oil
- 1900 - miscellaneous dry polymer master Batches.

Not → Solution SBR provide low - Rolling resistance →

3. PBR [POLY BUTADIENE-RUBBER]

→ polybutadiene is a homopolymer of Butadiene



→ It is prepared either by solution or emulsion polymerisation.

→ $T_g = -100^\circ C$

→ Density = 0.91 to 0.92

→ it is crystalline because of the stereoregularity in microstructure.

→ Good Wet Traction

→ poor Behaviour on mills.

→ Mostly used with NR & SBR (blend)

→ This blend, shows higher extrusion rate

→ poor tack,

→ poor Road grip of tyre tread

(A feature Related to the high Resilience)

→ poor tear & tensile strength compare with NR & SBR

→ BR has tendency to cold flow

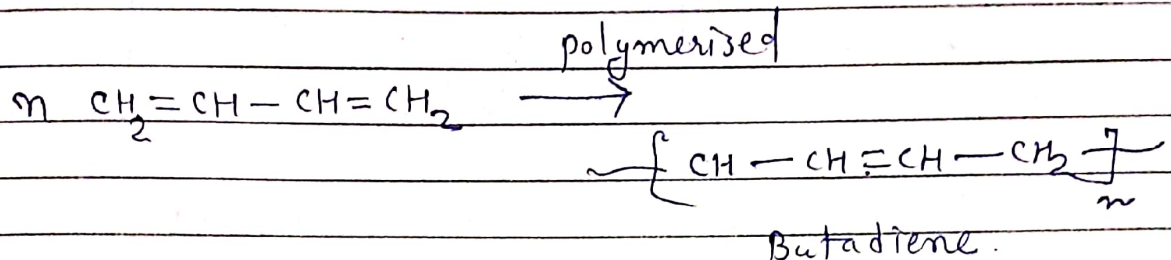


slow during storage.

Advantages

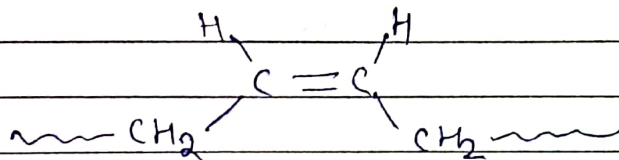
- Low Temperature flexibility
- Heat Ageing resistance
- High resilience at low deformation.
- ozone resistance
- Ability to accept higher level of filler and oil with less deterioration of its properties.
- good Abrasion resistance

Chemical structure $\rightarrow$

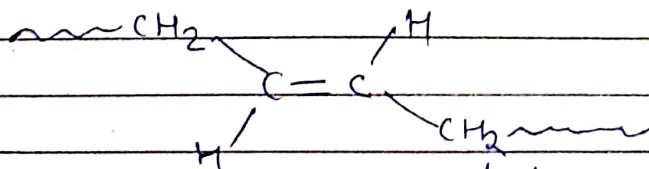


Structure

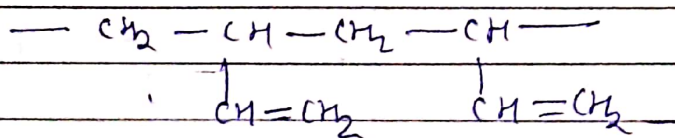
(i) 1,4 Cis - poly Butadiene



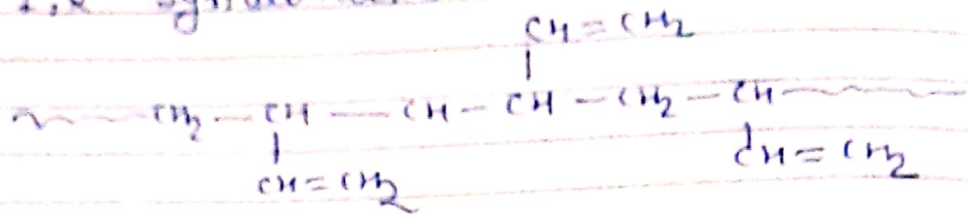
(ii) 1,4 Trans - poly Butadiene -



(iii) 1,2-isotactic poly Butadiene



(b) 1,2 - syndiotactic



→ Manufacturing process

The monomer Solvent purified & Dried the Catalyst is prepared Separately and added to reaction Vessels Containing monomer & Pyre Solvents, when the desired degree of polymerisation is reached The Catalyst, inactivated., The unreacted Solvent is Removed & polymer is recovered & Washed, dried & Balled--

GRADE →

Dry polymer	--	1200 - 1249
oil extended	---	1250 - 1299
Black master Batch..	-	1300 - 1349
oil Black master Batch-		1350 - 1399
latex...		1400 - 1449
miscellaneous ~		1449 - 1499

Disadvantages

- poor processibility if used alone
- Necessity to used Reinforcing fillers
- very little resistance to heat and ozone
- very little resistance to oil and gasoline hydrocarbon Solvents--

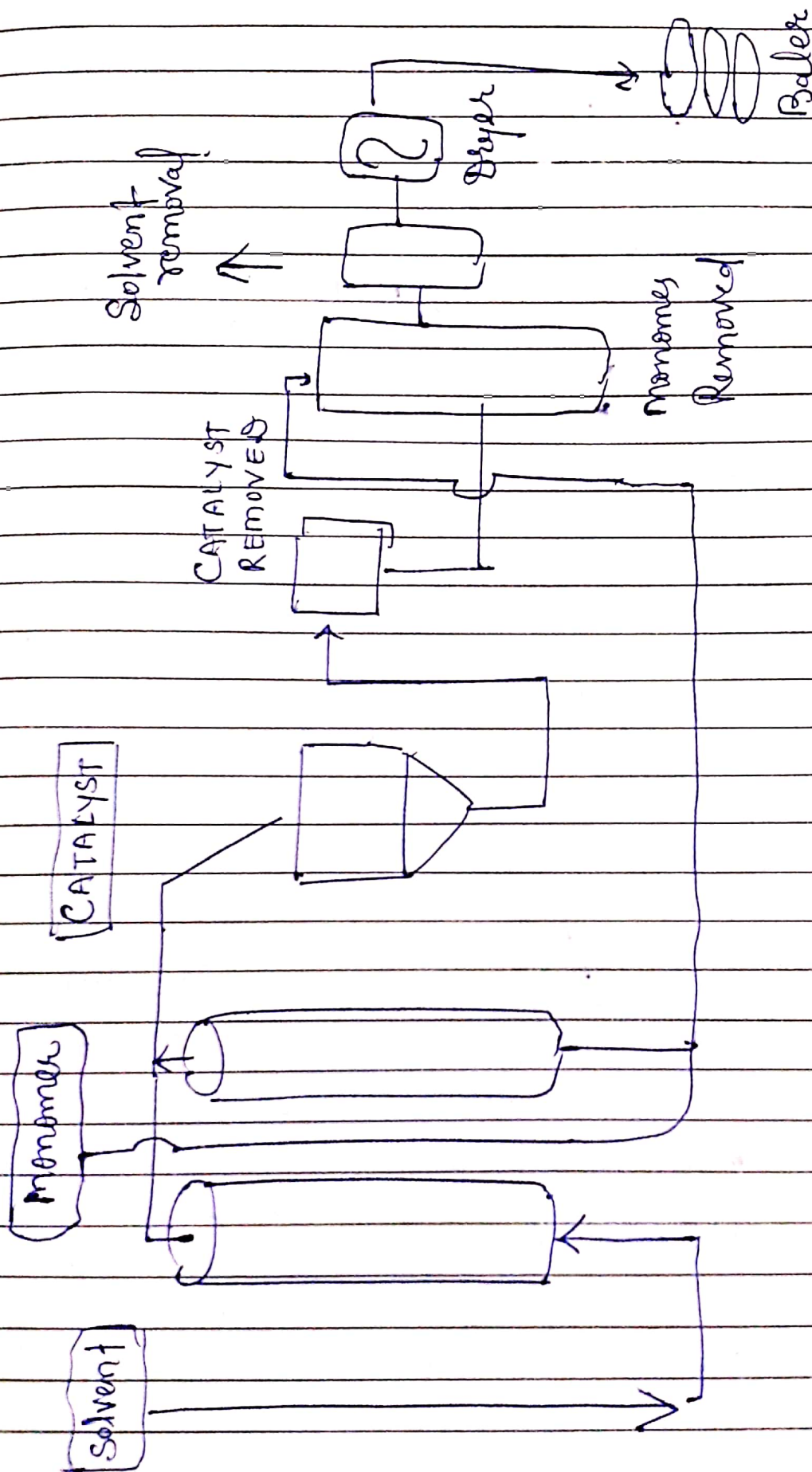
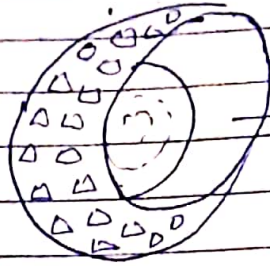


Fig → Flow Diagram for polybutadiene rubber

Application of polymer blend



major Tyre sectors

①

NR / BR
Blend

- Truck Tyre Tread (RIB)
- Steel Cord Coating

Truck Tyre Side Wall

②

SBR / BR
Blend

- Car Tyre tread
- Light Tyre (Truck tyre Tread)

③

NR / SBR
Blend

- Car tyre tread Compound
- LCV Tread Compound
- Bead Compound

Special purpose Rubber / other Rubbers.

- ACM → Acrylic Rubber
- EPDM → Ethylene propylene Rubber
- IIR → Butyle rubber
- NBR → Nitrile Rubber
- CR → Chloroprene Rubber
- CIIR → Chlorinated Butyle Rubber
- BIIR → Brominated Butyle Rubber
- AEM → Ethylene - Acrylate Rubber
- HNBR → Hydrogenated Nitrile Rubber.
- CSM → chlorosulphonated polyolefins.
- T → polysulphide Rubbers,

[MQ, VMQ]
[PMQ, PVMQ] → Silicone Rubber

FKM, CFM → Fluoro / fluoro Rubber ✓

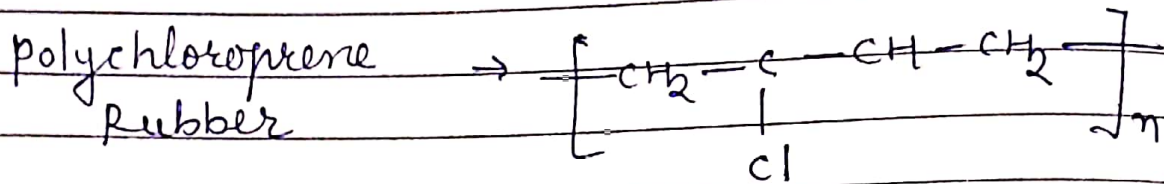
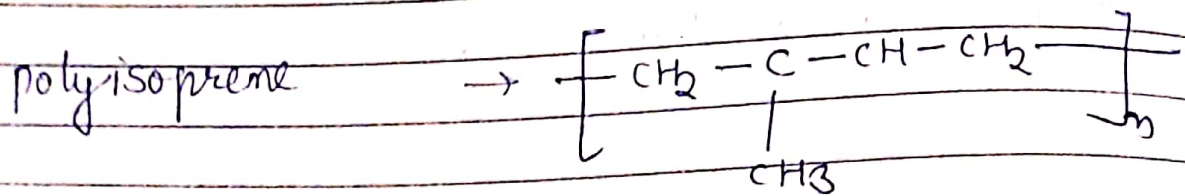
FVMQ → Fluorosilicone Rubber,

① CR

[Neoprene]

POLY CHLORO PRENE RUBBER,

polychloroprene Rubber is first synthetic Special purpose Rubber.



- polychloroprene rubber is manufactured by emulsion polymerisation technique.
- it is also known as chloroprene.

properties $\rightarrow$

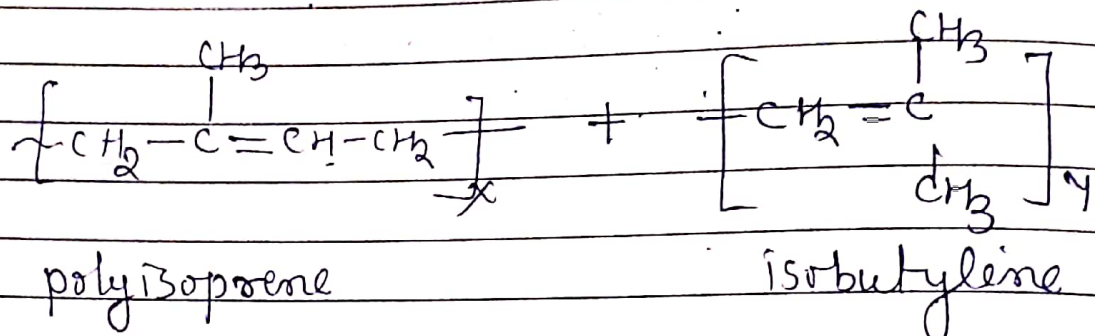
- CR rubber has good resistance to O_2 & O_3
- good physical properties over a wide range of curing temp.
- polychloroprene rubber has "heat memory" (minimum during processing stages avoid scorch)
- resistance to flame (flamoresistance)
- The Rubber is resistance to bacterial attack
- it has lower gas permeability than NR,
- CR has poorer electrical resistance than NR. But resistance to Corona crack.

uses of CR :-

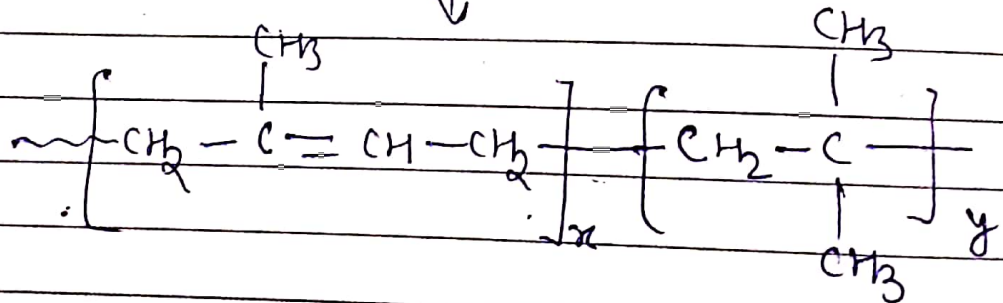
- Hoses of chemical & Cover for Hose (oil.)
- gaskets.
- O-rings (Heat resist. properties)
- Diaphragms.
- Special purpose conveyors in coal mines.)
- transmission V or timing Belts.

② IIR
(Isobutylene - isoprene Rubber)
Butyle Rubber.

- Butyle Rubber is produced by Co-polymerising isobutylene with small amount of isoprene.
- it has extremely low permeability to gases and moisture. Because very few (low) double bonds.



$$\begin{array}{l} x = 1\% \\ y = 99\% \end{array}$$



[IIR]

- The small amount of unsaturation is included to furnish the necessary sites of Sulfur Vulcanisation.

Major suppliers of this elastomers in the united states are Exxon chemical company and cities service company.

flow Diagram of Butyle Rubber production:-

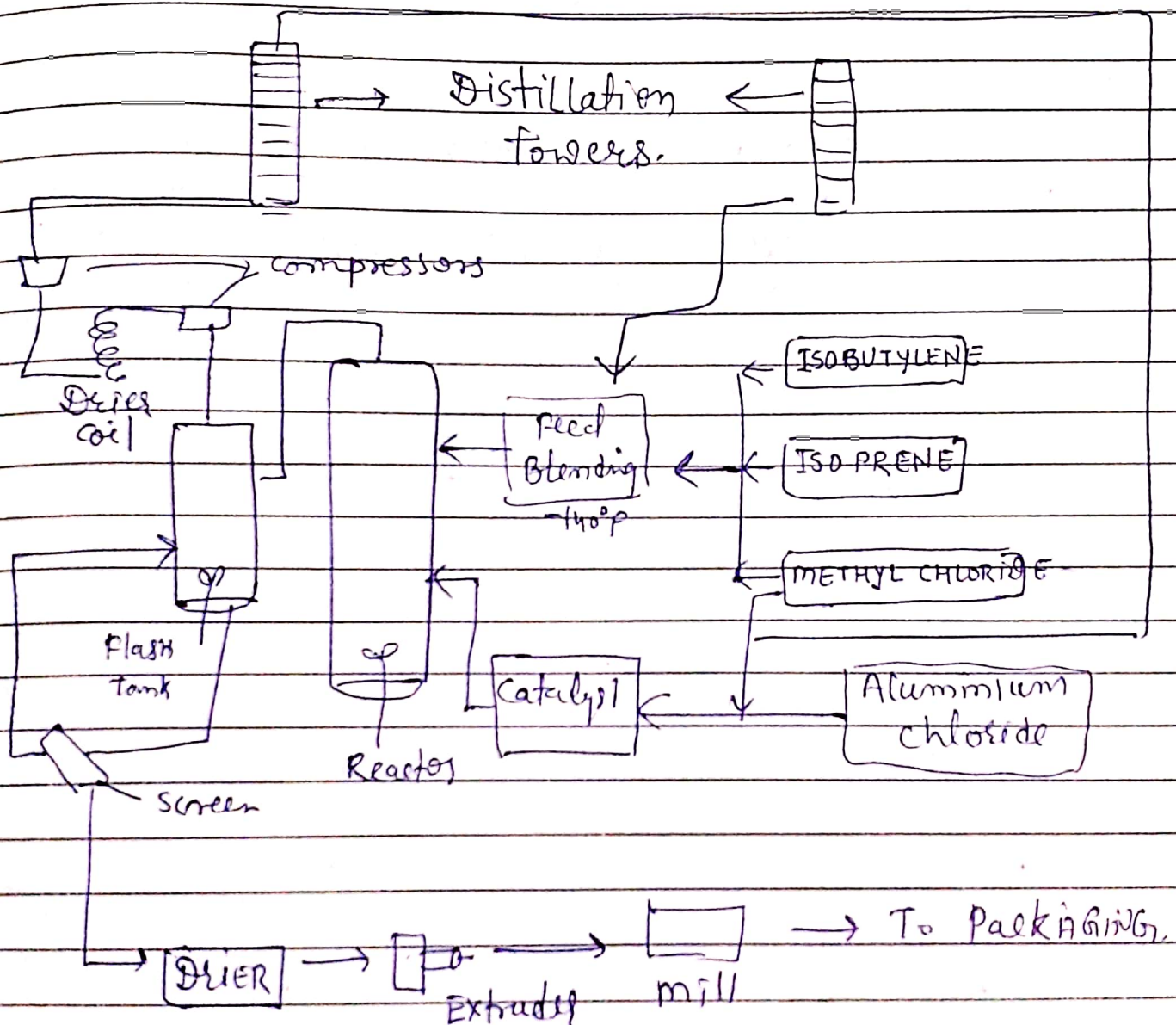


fig → flow Diagram for Butyle Rubber production.

GRADE →

(i) at low unsaturation (0.6 - 1.27%)

used where max. resistance to ozone, weather flexing and chemical required. But is slow curing grade

ex → irrigation tubing, tank lining, roof coverings, cable insulation, gaskets

(ii) medium unsaturation (1.5 - 2 mole %)

This is the largest consumption grade. Higher unsaturation gives faster g-curing and higher mod gives higher filler loading and greater compounding latitude

Ex - inner tubes,

Tyre curing bladders

Shock absorbers,

hoses,

pedal pads, weather strips, bushing

radiator hoses, floor mats,

(iii) high unsaturation (above 2% mole)

This is faster curing grade. It gives very good heat resistance, used for

Rain coats seals,

conveyor belts; Footwear

gaskets

process aids →

Butyl is highly saturated rubber so plasticizers are also highly saturated. & oil - paraffin process oil used

and with resin cure $\rightarrow$ Rubber,

Typical formulation, (exa.)

Butyle - 100 phr
GPR Black - 70 phr
paraffin oil - 25 phr
zinc oxide - 5 phr
St. acid - 2 phr
TMTD - 1 phr
MBT - 0.5 phr
Sulphur - 1.5 phr

The polyisobutylene portion provide a high degree of impermeability to Butyle, which leads to an exclusive use in inner tubes. The CH_3 group clustering -

- around the backbone packs. The molecules tightly to prevent any seepage of air.

Types

- Bromobutyle
chlorobutyle --

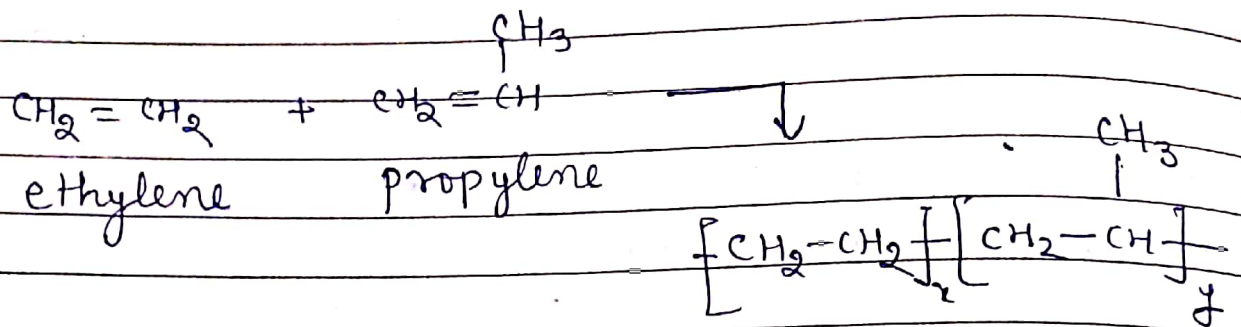
properties

- Excellent gas impermeability.
- very good electrical insulation properties.
- Good Gum Tensile strength
- Good heat resistance...

③ EPDM

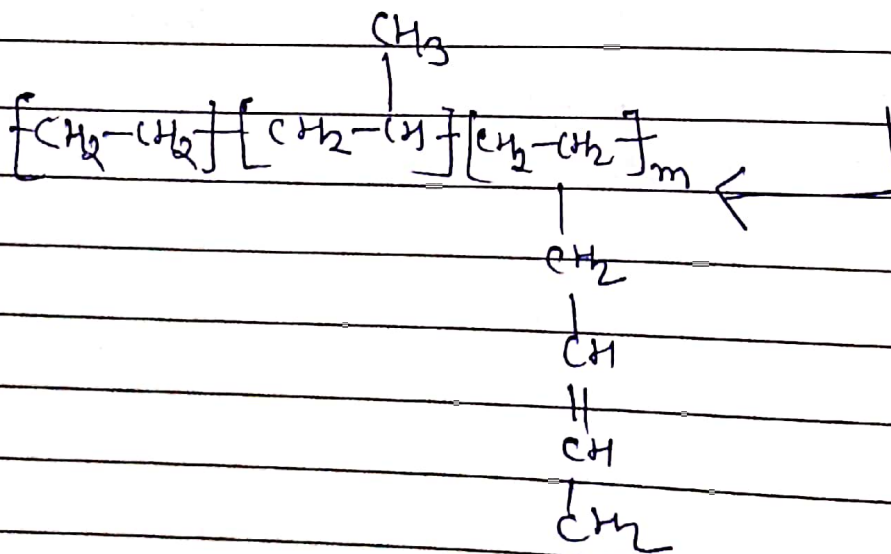
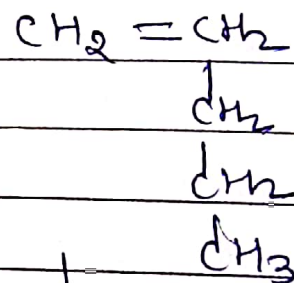
Ethylene propylene terpolymer
 Crackless Rubber,

These are copolymers of Ethylene and propylene. Containing in addition a small amount of a diene for unsaturation, just enough for unsaturation, just sulphur vulcanisation, EPDM can also be vulcanized with peroxide vulcanizing agent.



Ethylene propylene
 Rubber -

+



EPDM (2 to 10% mole unsaturation)

The Double bond is not in main polymer but pendent to it, This accounts for it EPDMs. good weather and ozone resistance, if and when oxygen react with the double bond, there is no chain scission of the polymer backbone, hence no cracking of Rubber.

Because the Various chain unit are Randomly arranged in the chain, EPDMs do not crystallize on stretching and require a reinforcing filler to develop high strength,

EPDMs. are very hard at room temperature,
EPDM Applications.

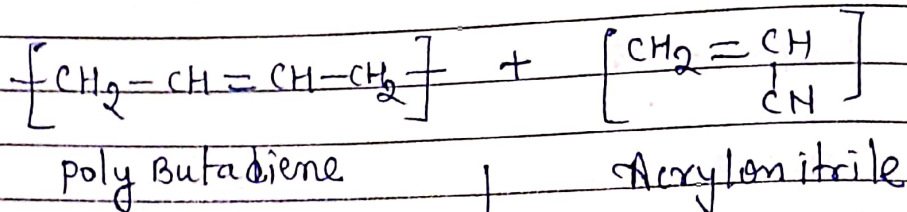
- use in - Cable insulation & jacketing
- Radiator hose,
- electrical cable,
- washing machine hose,
- Seals,
- door fenders,
- window profile & door,
- white sidewall of tyre,

(4) NBR

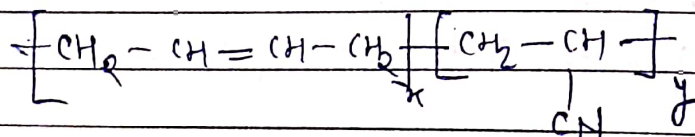
Nitrile Rubber

Nitrile Rubber is a copolymer of two monomers Butadiene and Acrylonitrile.

The Acrylonitrile content varies 20 to 50% range depending on the grade, The ACN content only makes the polymer polar and hence resistance to non polar mineral oils, grease and petrols,



↓
Emulsion polymⁿ
at 5°C (known cold polymⁿ)



NBR

$x = 50\% - 80\%$ range,

$y = 20\% - 50\%$ range,

properties :-

⇒ In NBR, The ACN contents only makes the polymer polar. So Excellent oil resistance

→ Good fuel resistance,

- Abrasion resistance
- poor tear resistance
- Solvent resistance

⑤ HNBR (HSN or ENM)
Hydrogenated Nitrile - Butadiene Rubber

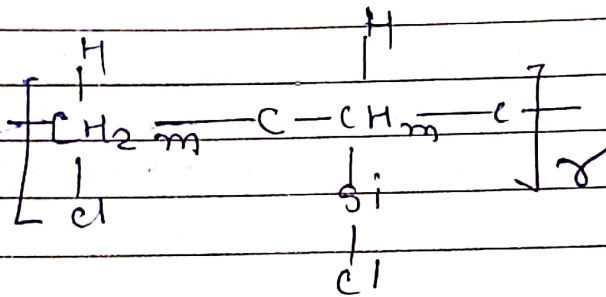
Completely hydrogenated NBRs have a fully saturated polymer chain & can be cross-linked with peroxides, the vulcanizates gives the highest resistance to hot air, and hot oils that can be achieved with NBRs.

(6) CSM

chloro sulphonated polyethylene.

CSM is a synthetic Rubber prepared by chloro-sulphonation of polyethylene.

- The resulting polymer is extremely stable chemically, and its vulcanizates possess -
- outstanding resistance to ozone, as well as excellent flame ret, heat & weather & abrasion resistance.



Curative - metallic oxide / Epoxy resin

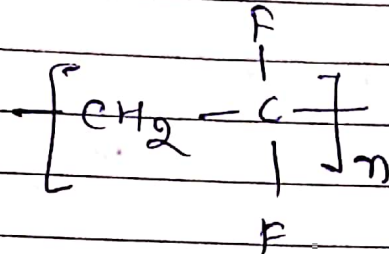
properties → resistance to Burning chemicals, oil
Ageing

uses → cables, hoses, molded goods.

(7) FKM / FFKM

fluorocarbon rubber / elastomer.

Fluoroelastomer are highly specialized products, that as class show the best resistance to all Rubber to attack heat, & chemicals and solvents & Excellent steam resistance.



Heat & ageing resistance 1000hr @ 220°C =

Shaft seals in cars, Components in aircrafts & rockets.

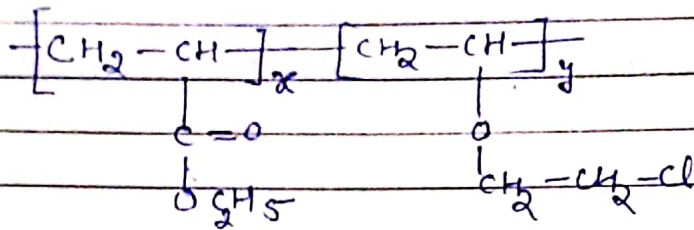
properties

- steam resist.
- high temp. compression set resistance.
- resistance to heat, light, ozone, solvents,

(8) ACM

polyacrylic rubbers

polyacrylics are high heat & oil resistance - specially elastomers.



$$x/y = 20 \text{ or more}$$

polyacrylic rubbers have two major component.

(i) 95 to 99 % polymer in backbone.

(ii) reactive cure-site comprising 1 to 5 % of the polymer.

The backbone is made of monomeric acid esters.

Primary curatives $\rightarrow$ Thiourea, Di or polyamine + Acid acceptors.

properties $\rightarrow$ Good heat oil & ozone resist looks
@ 160-170°C

use $\rightarrow$ oil seals, O-rings, oil hoses.

exg Acrylic rubbers.

(9) Silicone Rubbers

Silicone Rubber is considered as an organo-mineral high polymer called poly-siloxane, having the general formula $[RR'SiO]$ and consisting of alternating silicon and oxygen atoms in the main chain.

Silicone rubber is available form →

(i) silicone rubber is available form.

(a) Solid elastomer

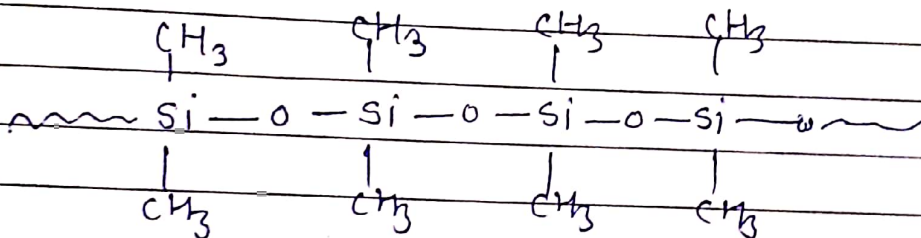
(b) Sponge

(c) potting resin

(d) Sealant

(e) adhesive

Rubber has structure is



The Si-atom in the polymer chain can carry other side groups resulting different types of properties. A side group is vinyl.

properties -

(1) Low Temp. flexibility

(2) Radiation Resistance

(3) Heat oil & solvent resistance.

• Curing Agent / curative. / Vulcanisation

Vulcanisation is the process by which mainly plastic Rubber is converted into elastic Rubber or hard rubber.

The process which involves chemical linking of polymer chain is called "cross-linking".

The cross linking of S-atoms, C-C-bond, polyvalent organic radical or polyvalent-metal ions.

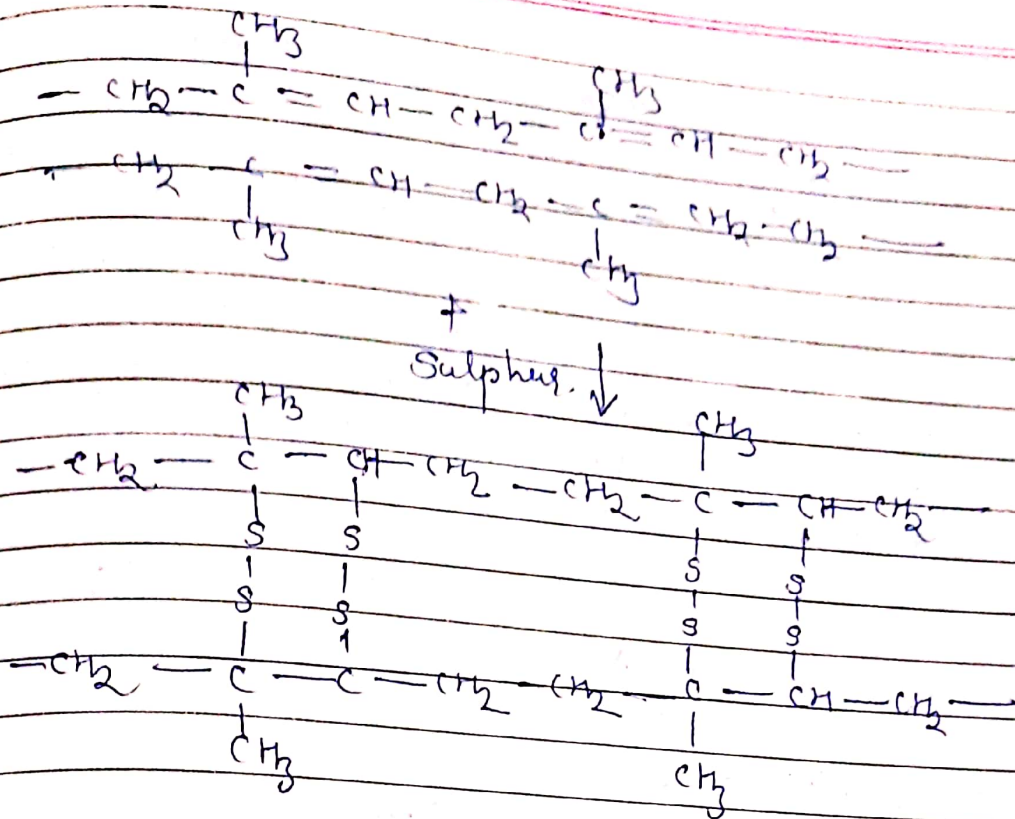
Due to vulcanization the long rubber molecule become crosslinked and 3 dimensional network is formed.

Due to increase the properties

- (i) Rubber loses its tackiness,
- (ii) Become insoluble in solvents,
- (iii) more resistant to deterioration caused by heat, light.

Increase in

- P.S.T
- Toughness,
- hardness increase
- permanent set decrease
- it has excellent resilience,
- it possesses low water absorption tendency.
- it has higher resistance to oxidation & wear & tear abrasion,
- it is better electrical insulator.



Types of Sulphur

- (i) Rhombic / soluble
- (ii) Amorphous / insoluble

(i) Rhombic (soluble)

This is an allotropic form of Sulphur, which occurs as an 8₈

This is soluble in CS₂ and also in Rubber

Temp. ↑
NR at R. Temp.

80°C

120°C

Solubility ↑

1%

4.5% ↑

10% ↑

of mixing temp. is high the proportion of Sulphur dissolved at high temperature tends to crystallize out and diffuse to surface when cooled know blooming

(Reduction in building tack during processing)
cheaper & widely available.

(ii) Amorphous sulphur. / Insoluble sulphur: →
Allotropic form, occur in polymeric form with
m.wt 100,000 to 300,000 (black to blue) range
formed by rapid cooling of molten sulphur or
precipitation from aqueous solution of sulphur
derived acids,

Insoluble in CS_2 & Rubber also therefore
has no tendency to bloom.

Sulphur Vulcanization system.

- (i) Conventional system (CV)
- (ii) Semiefficient system (S-EV)
- (iii) Efficient vulcanization system (EV)

(i) Conventional system: →

in the case →

- high proportion of sulphur - (1.5-3 phr)
- low amount of accelerator (1-1.5 phr)
- The network formed mainly polysulphide bond

→ In these systems, these cross link will
decompose at elevated temp. leading to loss
of cross linking and network containing
modification.

The network inefficiently crosslinked.

(ii) Semi-efficient crosslinked systems,

Having efficient cross-linking (monosulphide links) in between EV system & CV systems.

In this - S or sulphur donors and accelerator phr at levels intermediate between those of EV and CV systems,

→ cross linking have thermal stability

(iii) Efficient systems.

In which -

Small amount of sulphur (0-0.5 phr)
high proportion of accelerator (2-6 phr)
have the final network crosslinked with
mainly monosulphide bonds.

Such network have excellent thermal stability

	Typical CV system	Semi EV	EV systems
polysulphide bond %	65	40	0
Disulphide % bond	25	25	25
monosulphide bond %	10	35	75