

Colloidal dispersions - are defined as those polyphasic system (heterogeneous), where at least one dimension of the dispersed phase measure b/w one nm and one micrometer.

↳ Colloid ⇒ glue-like (in Greek) substances.  
eg - glue, gelatin, starch.

Heterogeneous  
↳ but not visible

Classification of dispersed system

① Classification According to physical state of two phases:

S.No.	Dispersed phase	Dispersion medium	Name system	Examples
1.	Solid	Solid	Solid solution	Colloidal gold
2.	Liquid	Solid	Crystals	wax, gels
3.	Gas	Solid	Solid foam	foam, rubber
4.	Solid	Liquid	Sols	most paints, proteins
5.	Liquid	Liquid	emulsion	milk
6.	Gas	Liquid	foam (liquid)	Soap lather
7.	Solid	Gas	Aerosol of solid	Soda water Smoke
8.	Liquid	Gas	Aerosol of liquid	dust storm Fogs, clouds

# ② Classification According to particle diameter -

① Molecular dispersions  
 ↳ dispersed particles range ~~from~~ less than 1 nm  
 ↳ invisible in electron microscope  
 ↳ pass through semi-permeable membranes  
 eg. oxygen gas, glucose etc.

② Colloidal dispersions  
 ↳ Range from 1 nm to 1  $\mu$ m  
 ↳ visible in only electron-microscope (not ordinary microscope)  
 ↳ pass through filter paper but not pass through semi-permeable membrane  
 ↳ eg. colloidal silver sol, natural & synthetic polymers

③ Coarse dispersion  
 ↳ more than 1  $\mu$ m  
 ↳ visible under ordinary microscope.  
 ↳ Do not pass ~~semi~~ filter paper, retained on the filter paper.

↳ Rapid diffusion  
 ↳ not settle  
 ↳ Homogeneous  
 ↳ Tyndall effect don't show  
 ↳ Brownian motion negligible  
 ↳ Separation not possible

↳ very slow diffusion  
 ↳ under centrifugation settle down  
 ↳ Heterogeneous  
 ↳ show  
 ↳ show (occured)  
 ↳ separate through semipermeable membrane

↳ not diffusion  
 ↳ under gravity settle down  
 ↳ Heterogeneous.  
 ↳ show  
 ↳ may be / may not  
 ↳ separate through filter paper.

⇒



# Size and shape of Colloidal particles

Particle Size: Particle size influences the colour of a dispersion. The wavelength of light absorbed by a particles is approximately related to its density. radius.

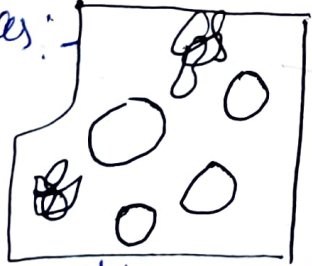
⇒ The larger the particle, the shorter the wavelength of light transmitted, i.e. the colour approaches the violet end of the spectrum.

eg Colloidal gold has red colour (650 to 750 nm) while intermediate size is in violet colour.

## Particle shape:-

The shape of particles of colloidal is depends upon method by which they are prepared and the type of interaction between dispersed phase with dispersion medium.

- They may be exists in cubical, ~~spher~~ spherical, spiral thread, cylindrical disc & rod shaped,
- It visible in electron microscope and look like as:-



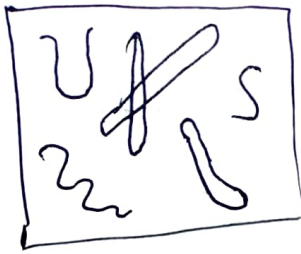
Spheres and globules  
surfactants, poliomyelites



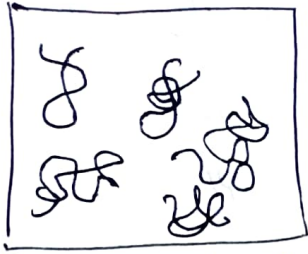
Short rods and prolate ellipsoids



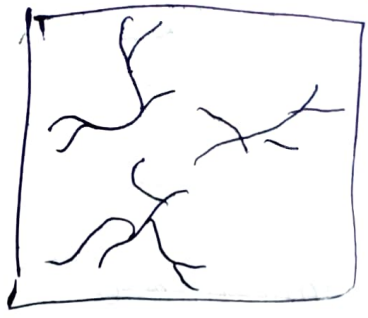
oblate ellipsoids and flakes



• Long rods & threads



• Loosely coiled threads



• Branched thread.

⇒ It also influence the ~~colour~~ <sup>viscosity</sup> of particles.

eg Spherical particles dispersion of low viscosity while linear particles generally produce more viscous dispersion.

⇒ Classification of Colloids:-

• Based on nature of interaction (affinity) between dispersed phase and dispersion medium

three types -

- ① Lyophobic Colloids ✓
- ② Lyophobic Colloids ✓
- ③ Association Colloids ✓

④ Lyophilic Colloids: Lyo → solvent, philic → loving -  
- Also called as solvent loving colloids.

• These are those solution, in which the dispersed phase has great attraction for the dispersion medium.

Types

Hydrophilic

- When dispersion medium is water

eg - Acacia, Albumin & gelatin indize in

Lipophilic

→ when dispersion medium is oil (other than water)

eg - Rubber, Poly-styrene in non-aqueous (Benzene)



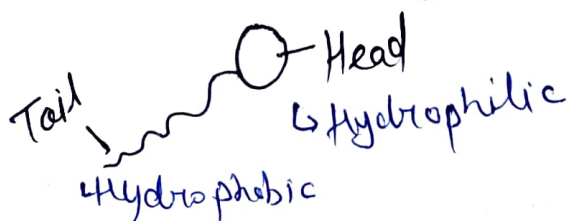
- ⇒ These are thermodynamically stable.
- ⇒ Viscosity of these solution generally increase on addition of dispersed phase.

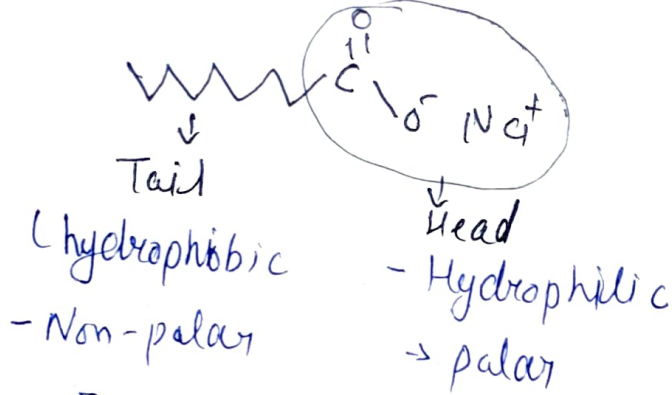
### ② Lyophobic Colloids: Lyo → Solvent, Phobic → Hating.

- Also termed as Solvent hating colloids.
- In which the dispersed phase has very less interaction [no affinity] for the dispersion medium.
- They are thermodynamically unstable (low stability).
- Viscosity of these solution does not increase on addition of dispersed phase.
- If the dispersion medium is water then they also called as Hydrophobic colloids.
- eg. metal such as gold, silver in water.

### ③ Association Colloids:

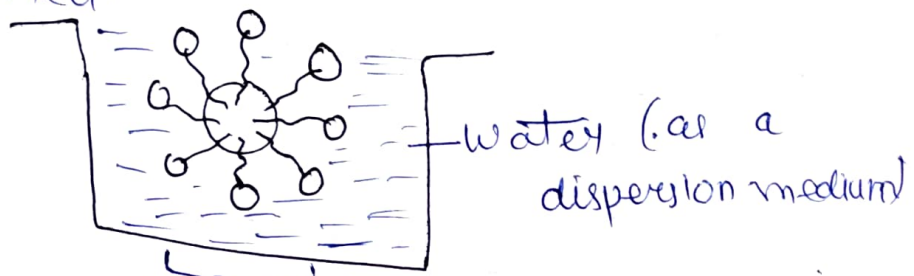
- ↳ Also called as Amphiphilic colloids.
- Molecules / Ions have both polar and non-polar group.
- They exist separately at low concentration.
- They associate at cmc (critical micelle concentration) to form micelle of colloidal size.
- They are also thermodynamically stable.





CMC → [Critical micelle Concentration]

The ~~the~~ minimum concentration at which micelle are formed.



↳ micelle (Associated Colloids)

- these amphiphiles also known as surfactants -
- Viscosity increases with addition of amphiphiles.
- ⇒ Comparative account of their general properties

Characteristic features/properties	Lyophilic colloids	Lyophobic colloids	Association Colloids
1. Nature of Interaction	Strong, high affinity	Little affinity (very low)	Depend on type of dispersion medium
2. Dispersed phase	mostly organic molecules	mostly inorganic molecules	Aggregation of surfactants
3. Size	Small ✓	Large ✓	Small ✓
4. Viscosity ✓	Increased	Same as dispersion m.	increases
5. Reversibility profil.	Reversible	Irrversible	Reversible
6. Stability	Stable high	Less stable	stable
7. Method of preparation	Simple & Easy	special method required	Depends on CMC



# ⇒ Optical, Kinetic & Electrical properties

## ① Optical properties of Colloids -

These help to know about size, shape, structure of & molecular weight of colloids.

⇒ these are following -

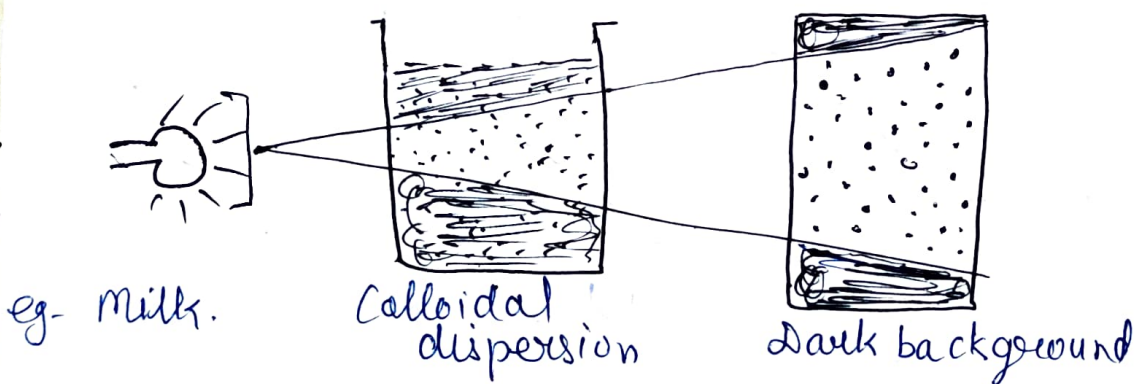
- ① Tyndall Effect (Light scattering)
- ② Ultramicroscopy
- ③ Electron microscopy
- ④ Turbidity

### ① Tyndall Effect :- When a beam of light is passed

through a colloidal solution (dispersion) kept in dark, the path of the beam get illuminated with blue colour.

This phenomena is known as Tyndall effect and the path is known as Tyndall cone.

⇒ The Tyndall effect is due to the scattering of light by colloidal particles.



• True (Homogeneous) solution does not show this, they have small particle size.

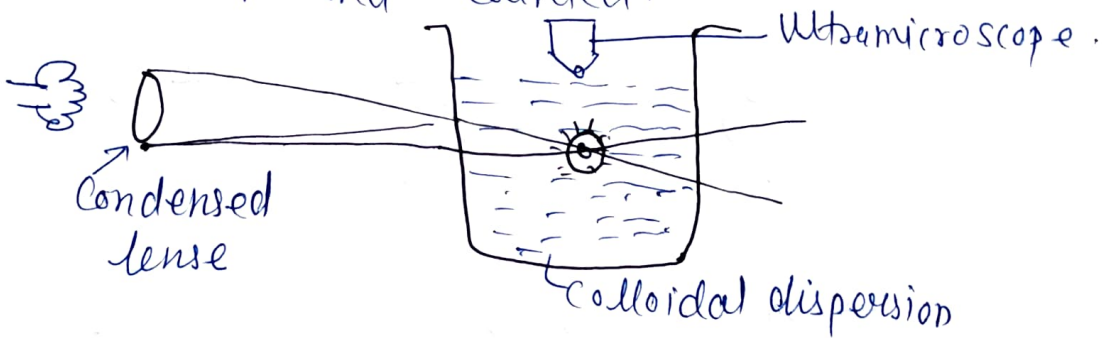
⇒ Heterogeneous (colloidal) dispersion show this, -  
 Lyophobic → show more  
 Lyophilic → show less (no effect)

(ii) Ultramicroscope :-

Ultramicroscope has been used to observe the Tyndall beam.

When a intense light beam is passed through the solution (colloidal dispersion) against a dark background at right angles to the plane of observation.

⇒ The particles will appear as the bright spot which can be observed and counted.



(iii) Electron Microscope :-

It give the actual picture of the colloidal particles.

- High energy electron beam are passed, it used to observe the size, shape and structure of colloidal particles.
- Useful of in lyophilic.

(iv) Turbidity :- All colloidal dispersion show turbidity according to molecular weight of colloids particles, spectrophotometer are used to check this.

Turbidity  $\propto$  molecular weight



## ② Kinetic Properties of Colloids:-

These properties help to know about the motion of colloidal particles in colloidal dispersion.

### Ⓐ Brownian Motion—

(i) Diffusion —

(ii) Sedimentation —

(iii) Viscosity —

### Ⓐ Brownian Motion:-

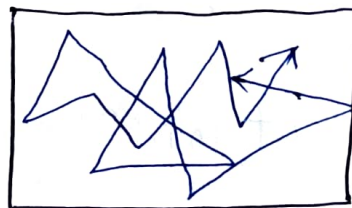
It is the zig-zag motion of colloidal particles in colloidal dispersion in continuously random manner.

→ It is given by scientist Robert Brown.

→ Particles continuously strike to each other and to the wall of container.

→ Velocity  $\uparrow$  → particle size  $\downarrow$

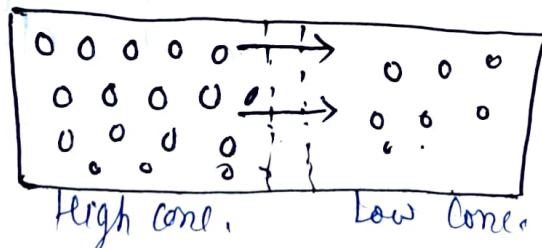
→ Brownian movement  $\uparrow$  → stability  $\uparrow$



### (ii) Diffusion:-

It is the movement of particles from an area of higher concentration to the area of lower concentration.

→ It is based on Fick's first law, that particles diffuse spontaneously (continuously) until the diffusion equilibrium is reached (is attained).



(iii) Sedimentation :-

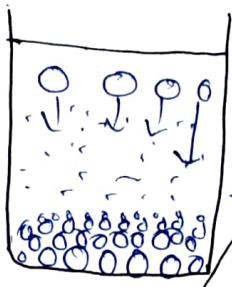
It is the settling down of dispersed phase particles into dispersion medium due to gravity.

⇒ It is depends upon molecular weight of colloidal particles (mol. weight ↑ = sedimentation ↑).

⇒ It is also depends upon the density difference of dispersed phase to the dispersion medium.

⇒ Brownian motion ↑  
↓  
Sedimentation ↓

⇒ Sedimentation ↑.  
↓  
stability ↓



→ stoke's law used to determine sedimentation.

⇒  $\eta = \eta_0 (1 + 2.5 \phi)$

Where,  $\eta$  = viscosity of dispersion Pa.s.  
 $\eta_0$  = viscosity of dispersion medium, Pa.s  
 $\phi$  = Volume fraction of particles

(iv) Viscosity :-

It is the resistance to fluid to flow under an applied stress.

⇒ It is depend upon -

- shape, size, molecular weight
- interaction between dispers phase and dispersion medium.

⇒ molecular weight & viscosity.

• Einstein describe on equation of flow to dilute colloidal dispersions of spherical particles -

⇒  $\eta = \eta_0 (1 + 2.5 \phi)$



### ③ Electrical Properties of Colloids -

These properties help to know about the charge on colloidal particles in dispersion (colloidal dispersion).

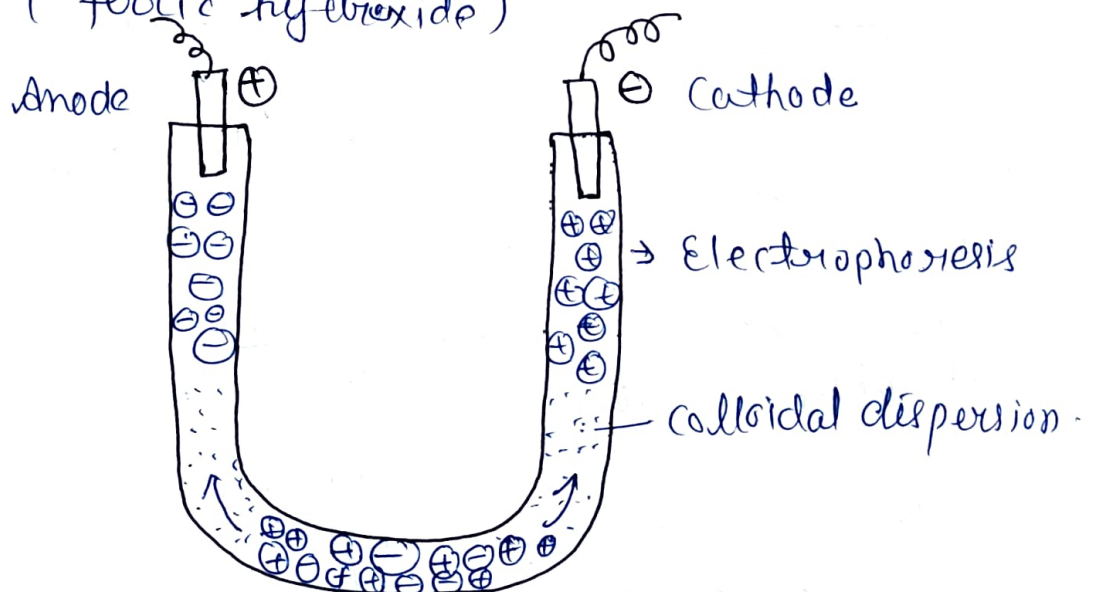
- (i) Electrophoresis
- (ii) Electrical double layer

#### (i) Electrophoresis :-

When an electric field is applied on colloidal dispersion, then the particles carrying charges move towards opposite charge electrode. i.e. -

→ Negative charged particles move towards anode (Kaoline, Sulphur).

→ Positive (+ve) charged particles move towards cathode (ferric hydroxide)



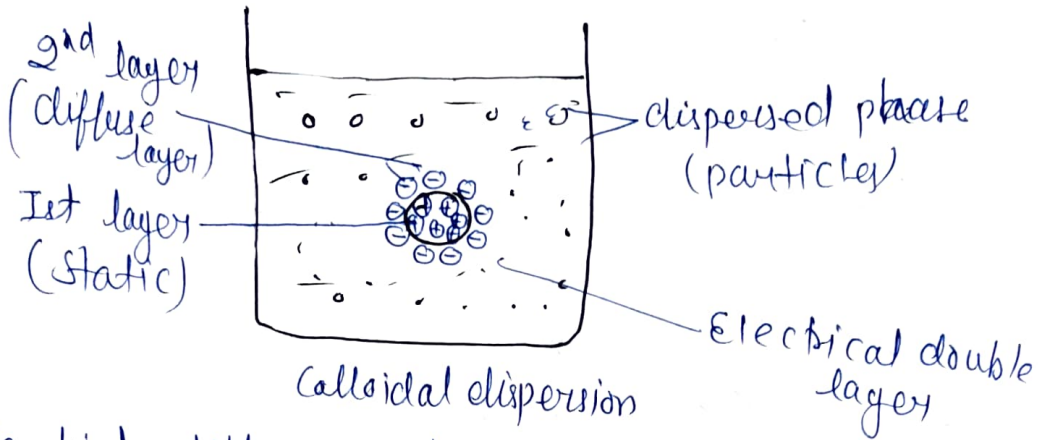
#### (ii) Electrical double layer :-

Helmholtz explained it in 1879.

→ In this theory, at the first layer charge is imparted to the surface of particles which is immovable also known as static layer (or Helmholtz layer)

and the second layer consists of diffused mobile ions (according to first layer).

⇒ The charge develop (present) on both the layers are equal

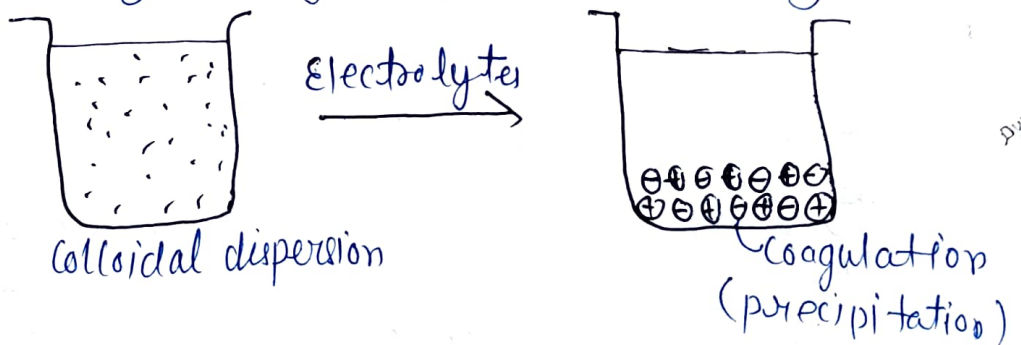


⇒ The potential difference between two layers i.e. static and diffuse layer is called as Zeta potential or Electrokinetic potential.

### ⇒ Effect of Electrolytes:

On addition or removal of electrolyte in colloidal dispersion may affect the stability of colloids.

⇒ On addition of excess of electrolytes, particles of colloidal dispersion precipitate due to accumulation of oppositely charged particles (stability decreases)

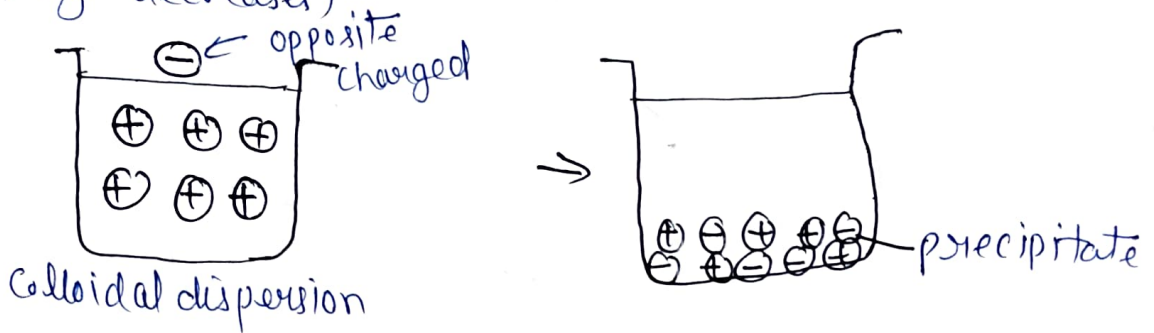




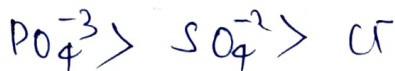
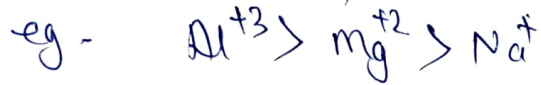
## Hardy Schulze law (rule):-

The phenomenon of deposition of colloid particles when oppositely charged electrolyte on it.  
(stability decreases)

eg



⇒ Charge on Cation/Anion of Coagulation/precipitation.



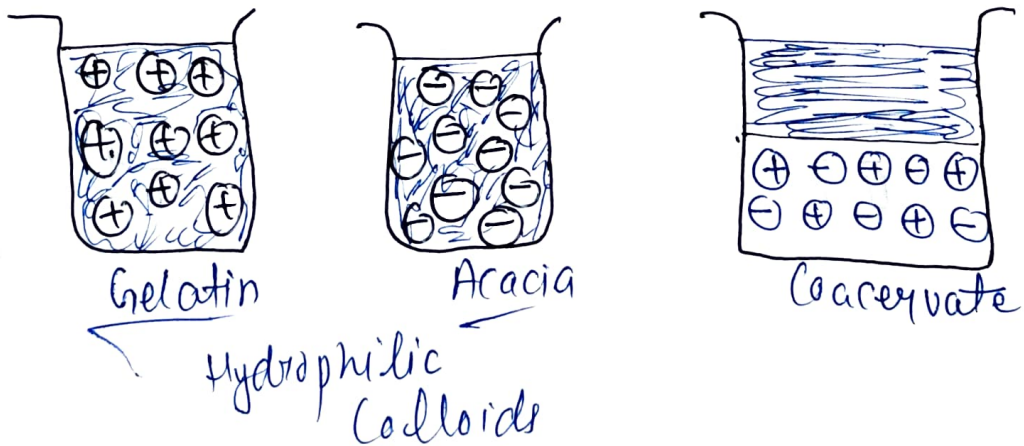
⇒ But if we add same charged particles then particles and electrolytes repel each other and increase stability of colloidal dispersion.

## ⇒ Coacervation:-

When two opposite charged hydrophilic colloids are mixed then there will be separation of the colloid rich layer.

⇒ The colloid-rich layer is known as Coacervate. The phenomenon is called as Coacervation.

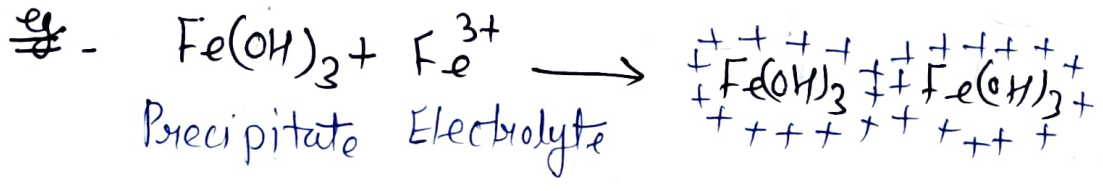
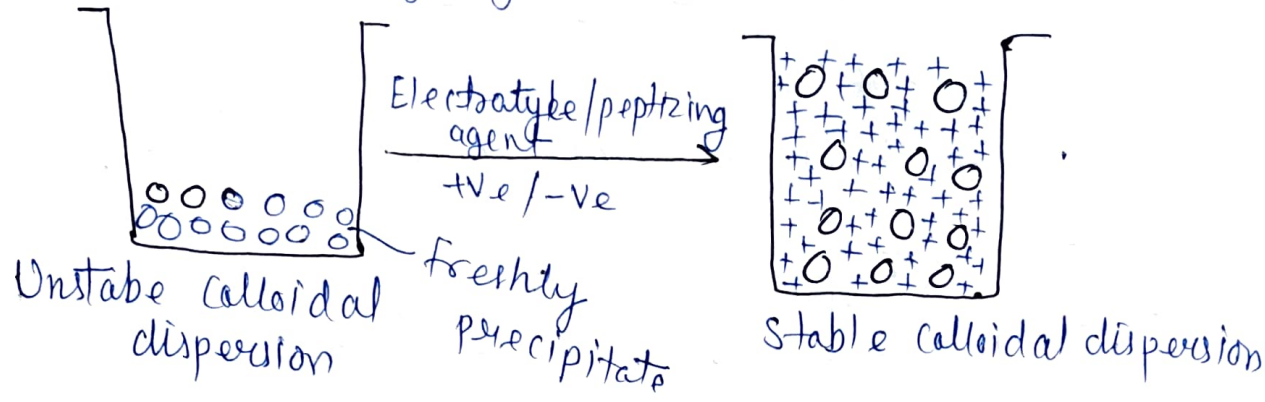
eg



### ⇒ Peptization :-

It is used for the formation of stable colloidal dispersion.

"It is the process of converting a precipitate into colloidal dispersion by shaking it with dispersion medium in the presence of small amount of electrolyte/peptizing agent."



### ⇒ Protective Action :-

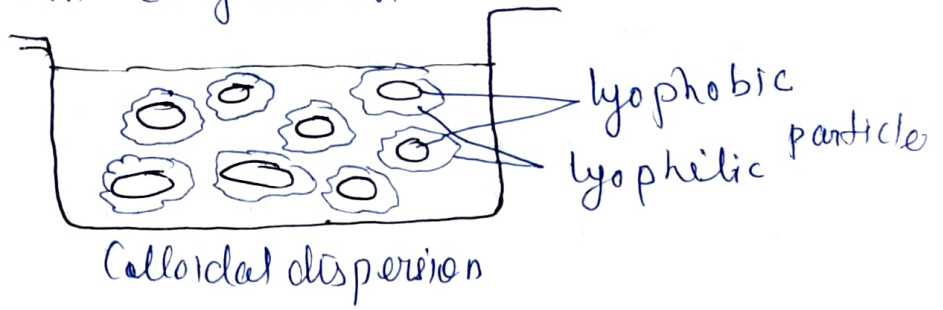
The addition of lyophilic colloidal solution into a lyophobic colloidal solution prevents a lyophobic sol from coagulation.

- This phenomenon is known as protective action and the lyophilic colloid used for this purpose is known as protective colloids.

mechanism- when lyophilic colloid added to a lyophobic colloid, the particles of lyophilic cover the surface of lyophobic particle and behave as protective colloids - and the particles of lyophobic behave as lyophilic colloids and



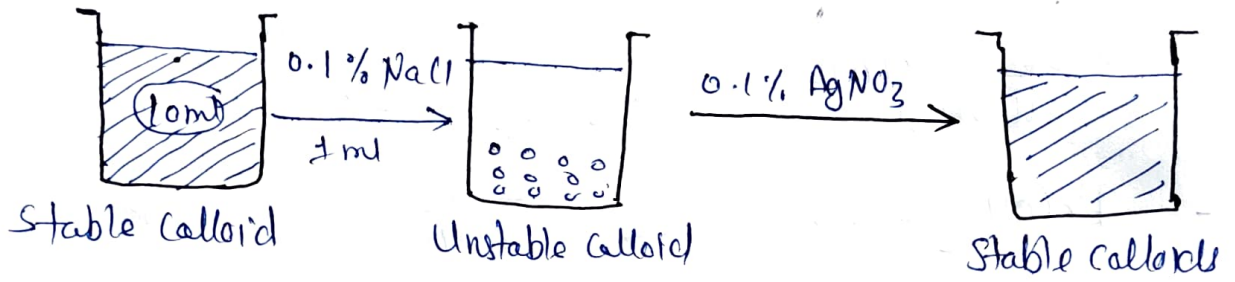
and prevents from coagulation.



### Gold number -

The amount of protective colloids in mg. which prevents the coagulation of gold solution (10 ml) when 1 ml of NaCl added to it.

eg-



→ Amount of 0.1% AgNO<sub>3</sub> to make it stable.