


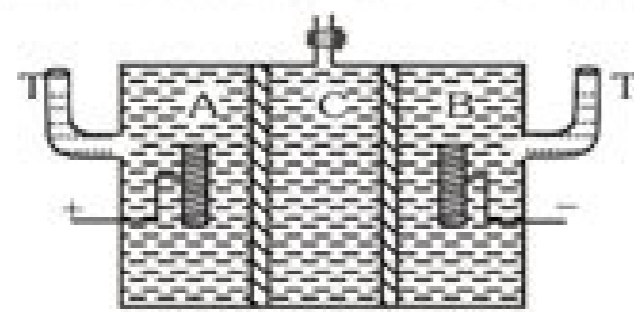
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Since the current in the colloidal solution must be carried by both positive and negative particles, ions of the diffused layer must be moving in a direction opposite to the direction of the movement of colloid particles. In a  $\text{Fe}(\text{OH})_3$  sol which is positively charged, the sol particles move to the negative electrode where their charge is neutralised and they aggregate and finally precipitate out. Thus, the entire colloidal matter settles down at the bottom.

◆ **Importance** : This phenomenon of electrophoresis is made use of in the following ways :

- (i) Determining the charge on the colloidal particles : direction of movement of the colloidal particles in the electric field shows the charge on them.
  - (ii) It can also be used to determine the rate at which colloidal particles migrate under the influence of an electric field.
  - (iii) It is also used in the identification and determination of homogeneity.
  - (iv) It is of great importance for the preparative separations of the colloidal substances.
- (ii) **Electro osmosis** : It is also known as electro-endosmosis, in the above experiment, a partition is made by animal membrane or parchment paper in between two electrodes, so that only the dispersion medium can move through it and not the colloidal particles. When potential difference is set up between the electrodes, then the dispersion medium is seen to move in a direction opposite to the direction of movement of the colloidal particles. This movement of the dispersion medium relative to the dispersed phase under the influence of the electric field is known as electro-osmosis. This is indicated by the rise of water level in one limb of the U-tube.



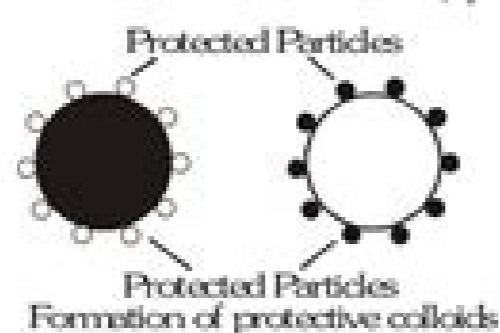
Measurement of electro-osmosis

#### □ **ELECTRIC DOUBLE LAYER:**

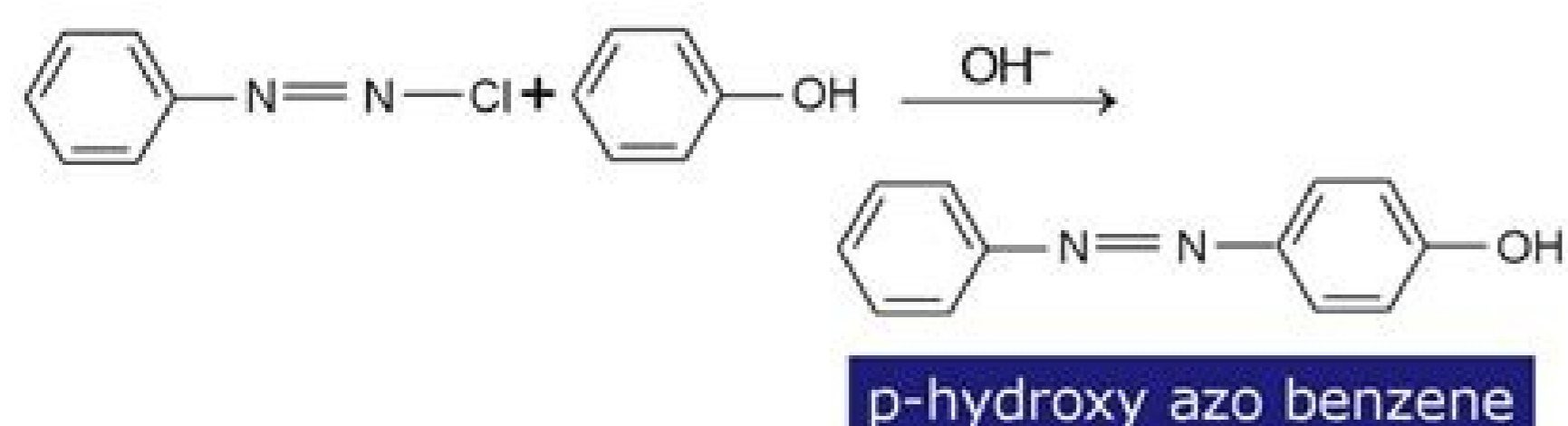
The surface of a colloidal particle acquires a positive or a negative charge by selective adsorption of ions carrying +ve or -ve charges respectively. The charges layer attracts counter ions from the medium which forms a second layer. Thus, an electrical double layer is formed on the surface of the particles i.e., one due to adsorbed ions and the other due to oppositely charged ions forming a diffused layer. This layer consists of ion of both the signs, but its net charge is equal and opposite to those adsorbed by the colloidal particles. The existence of charges of opposite signs on the fixed and diffused parts of the double layer creates a potential difference between these layers. This potential difference between the fixed charge layer and diffused layer of opposite charge is called electrokinetic potential or zeta potential.

#### □ **PROTECTION OF COLLOIDS :**

**Protection** : When certain hydrophilic colloids such as gum, gelatin, agar-agar etc. are added to a hydrophobic colloid, the stability of the latter is markedly increased. Now the addition of the small amounts of electrolytes does not cause the precipitation of the hydrophobic colloid. This action of the hydrophilic colloids to prevent precipitation of the hydrophobic colloid by the electrolytes is called protection and the hydrophilic colloid is called protective colloid. It is further observed that the protective colloid not only increase the stability of the hydrophobic colloid but the latter can be evaporated to dryness and the dry mass peptised by simply shaking with water. Thus the protective colloid converse an irreversible (hydrophobic) colloid into a reversible colloid.



## Coupling Reaction



### Abnormal Properties of HF

- (a) Liquid at room temperature due to H bonding.
- (b) Most thermal stable than other halogen hydrides.
- (c) It is weak acid but is extremely stable. It is not oxidised by strong oxidising agents. On the other hand all other halogen acid is oxidised.
- (d) HF form two type of salt ( $\text{NaHF}_2$ ,  $\text{Na}_2\text{F}_2$ )
- (e) This acid attacks on silica of glass & formation of following compounds.



### 7. OXY ACID OF HALOGEN FAMILY

Name of the acids and their salts	Oxidation state of the halogen in the acid	Oxoacids			Structure	Stability	Acidic strength
		Cl	Br	I			
Hypochlorous Hypochlorite	+1	HClO	HBrO	HOI	H-O-X	Decreases	Increases
Chlorous, Chlorites	+3	HClO <sub>2</sub>			H-O-X→O	Decreases	Increases
Chloric, Chlorates	+5	HClO <sub>3</sub>	HBrO <sub>3</sub>	HOI <sub>3</sub>	$\begin{matrix} \text{O} \\ \uparrow \\ \text{H-O-X} \end{matrix}$ →O	Decreases	Increases
Perchloric, Perchlorates	+7	HClO <sub>4</sub>	HBrO <sub>4</sub>	HOI <sub>4</sub>	$\begin{matrix} \text{O} \\ \uparrow \\ \text{HO-X} \\ \downarrow \\ \text{O} \end{matrix}$ →O	Decreases	Increases

Acidity order : HOX < HXO<sub>2</sub> < HXO<sub>3</sub> < HXO<sub>4</sub>

Oxidising power :HOX > HXO<sub>2</sub> > HXO<sub>3</sub> > HXO<sub>4</sub>

Thermal stability : HOX < HXO<sub>2</sub> < HXO<sub>3</sub> < HXO<sub>4</sub>

- (a) **Fluoric (I) acid (HOF)** : It is obtained when F<sub>2</sub> is passed over ice at 273 K.

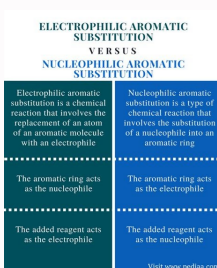
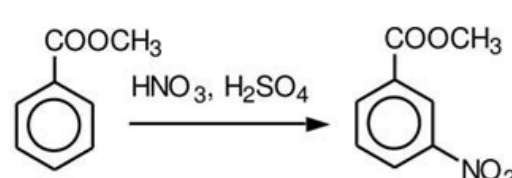
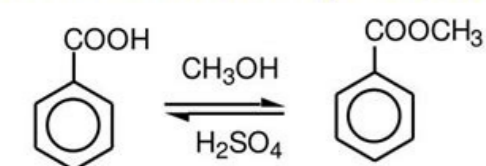


It is colourless unstable gas which decomposes to HF and O<sub>2</sub>. It is strong oxidising agent and oxidises H<sub>2</sub>O to H<sub>2</sub>O<sub>2</sub>.

- (b) **Hypochlorous acid or chloric (I) acid (HOCl)** : It can be prepared by following reactions.



### Nitration of Methyl Benzoate



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