

## APPLICATION OF COORDINATION

### COMPOUNDS

Application of formation in Qualitative analysis :-

a) In detecting metal ions:

Formation of Complexes is for detecting metal ions in solutions. E.g.

i). If an ammoniacal solution of the substance gives a scarlet red precipitate with DMG, it indicates the presence of  $\text{Ni}^{2+}$ .

ii). If a solution of the substance gives a blood red colour with potassium thiocyanate ( $\text{KCNs}$ ) it indicates, the presence of  $\text{Fe}^{3+}$ . The blood colour is due to the formation of the complex  $[\text{Fe}(\text{CNS})_6]^{3-}$ .

iii). If a solution of the substance gives a deep blue colour with ammonia it indicates the presence of  $\text{Cu}^{2+}$ . The deep blue colour is due to the formation of the complex  $[\text{Cu}(\text{NH}_3)_4]^{2+}$ .

iv). Oxine (8-hydroxy quinoline) is used for the detection of Mg, Al, Zn etc.



b) In separating metal ions:

i). Separation of  $\text{Ag}^+$  and  $\text{Hg}_2^{2+}$  in first group

analysis is done by adding aqueous ammonia.

$\text{AgCl}$  dissolves forming a complex  $[\text{Ag}(\text{NH}_3)_2]\text{Cl}$ .

$\text{HgCl}$  does not dissolve. It can be separated by filtering it.

ii). Separation of  $\text{Cu}^{2+}$  and  $\text{Cd}^{2+}$  in second group

analysis is done by adding  $\text{KCN}$  solution. The sulphides of both metals forms complexes and go into solution.

Now  $\text{H}_2\text{S}$  is passed  $\text{K}_2[\text{Cu}(\text{CN})_4]$  is more stable than

$\text{K}_3[\text{Cd}(\text{CN})_4]$ . So  $\text{Cd}^{2+}$  ion is precipitated on passing

$\text{H}_2\text{S}$ . It can be separated by filtration.

iii). In second group  $\text{Hg}^{2+}$ ,  $\text{Cu}^{2+}$ ,  $\text{Cd}^{2+}$ ,  $\text{Bi}^{2+}$ , arsenic,

antimony and tin are precipitated as their sulphides.

To the precipitate yellow ammonium sulphide solution

is added. Arsenic, antimony and tin alone form

complex sulphide  $[\text{AsS}_4]^{3-}$ ,  $[\text{SbS}_4]^{3-}$  and  $[\text{SnS}_4]^{4-}$

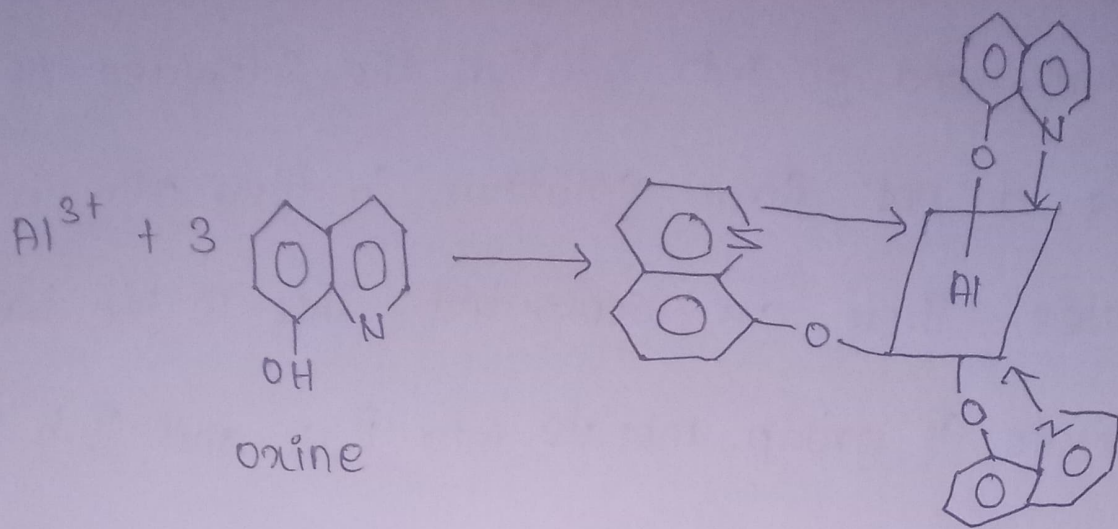
respectively, and go into solution the sulphides of elements do not form solution. So they remain as precipitates. They are separated. This is the basis of separate II group metals into II a and II b sub groups.

Quantitative Analysis:-

i).  $\text{Ni}^{2+}$  forms a complex with dimethyl glyoxime. The complex is bis (dimethyl glyoximate) nickel (II). It is scarlet red in colour. The formation of this complex is used for the gravimetric estimation of  $\text{Ni}^{2+}$  in solutions.

ii).  $\text{Al}^{3+}$  forms a complex with oxine (8-hydroxy quinoline  $\text{C}_9\text{H}_7\text{ON}$ ). The complex is trisoxynato aluminium (III) -  $\text{Al}(\text{C}_9\text{H}_7\text{ON})_3$ . The formation of this complex is used for the gravimetric estimation of  $\text{Al}^{3+}$  in solutions. The N and  $\text{O}^-$  are the donor sites.





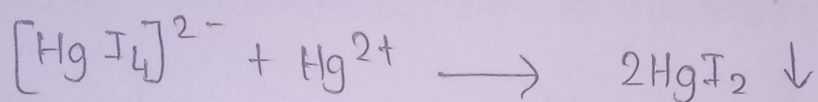
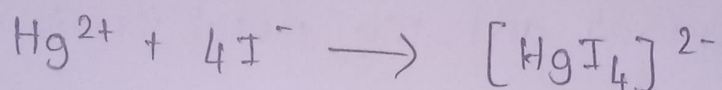
iii).  $K^+$  forms a complex with sodium hexanitrito Cobaltate (III). The complex is potassium hexanitrito Cobaltate (III)  $K_3[Co(NO)_2)_6]$ . The formation of this complex is used for the gravimetric estimation of  $K^+$  in solutions.

iv). Hardness of water is estimated by complexometric titration with standard EDTA using EBT (eriochrome black T) or murexide as indicator. The indicator metal complex and the EDTA metal complex have different colours. So the end point in such titrations are indicated by a colour change.

v). Metal ions like  $Fe^{2+}$ ,  $Ni^{2+}$ ,  $Cu^{2+}$  etc. which are present in traces are estimated spectrometrically

using suitable complexing agents.

vi). Iodine can be estimated by titrating it against  $\text{Hg}^{2+}$  in solutions. The formation of a red precipitate of  $\text{HgI}_2$ , indicates the end point.



(At the end point)

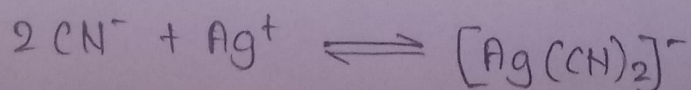
(Red precipitate)

Complexometric titration:

In these titrations ions (other  $\text{H}^+$  and  $\text{OH}^-$  ions) are converted into a soluble, slightly dissociated ions or compounds.

Description / Explanation with example:

When  $\text{AgNO}_3$  is titrated against  $\text{KCN}$ ,  $\text{Ag}^+$  is converted into the soluble complex  $\text{Ag}[\text{CN}]_2^-$  ion which dissociates slightly.





## Principle:

\* Metal ions can be determined by titrating them with a reagent that complexes them in solution. The solution to be titrated is buffered at a suitable pH and an indicator is added and the metal ion is titrated with a standard solution of the complexing agent. A sharp colour change marks the end point of the titration.

\* Complexometric titrations are convenient and accurate. They have replaced time consuming gravimetric procedures. Except for the alkali metals most metal cations can be determined by titration with a suitable complexing agent.

\* A suitable complexing agent is EDTA. It forms stable chelates with a large number of metal ions. It is used as a primary standard in complexometric titrations.