

HINTS & SOLUTIONS

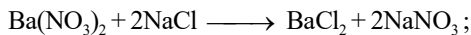
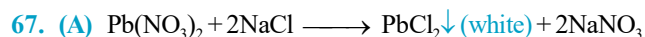
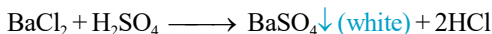
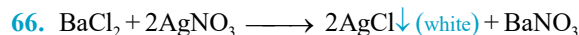
EXERCISE - 1

Single Choice

- $\text{SO}_3^{2-} + \text{Zn} + 8\text{H}^+ \longrightarrow \text{H}_2\text{S} \uparrow + 3\text{Zn}^{2+} + 3\text{H}_2\text{O}$
 $\text{Pb}^{2+} + \text{S}^{2-} \longrightarrow \text{PbS} \downarrow (\text{black})$
 $\text{Ag}^+ + \text{S}^{2-} \longrightarrow \text{Ag}_2\text{S} \downarrow (\text{black})$
 $2\text{MnO}_4^- + 5\text{H}_2\text{S} + 6\text{H}^+ \longrightarrow 2\text{Mn}^{2+} + 5\text{S} + 8\text{H}_2\text{O}$
- $\text{SO}_3^{2-} + \text{Ba}(\text{OH})_2 \longrightarrow \text{BaSO}_3 \downarrow (\text{white}) + 2\text{OH}^-$
 $3\text{SO}_2 + \text{Cr}_2\text{O}_7^{2-} + 2\text{H}^+ \longrightarrow 2\text{Cr}^{3+} (\text{green colour solution}) + 3\text{SO}_4^{2-} + \text{H}_2\text{O}$
- (i) $\text{Cr}_2\text{O}_7^{2-} + 3\text{SO}_2 + 2\text{H}^+ \longrightarrow 2\text{Cr}^{3+} (\text{green}) + 3\text{SO}_4^{2-} + \text{H}_2\text{O}$; CO_2 and O_2 do not give reactions with acidified $\text{K}_2\text{Cr}_2\text{O}_7$
 (ii) $\text{Ca}(\text{OH})_2 + \text{CO}_2 \longrightarrow \text{CaCO}_3 \downarrow (\text{white}) + \text{H}_2\text{O}$; O_2 does not give reaction with acidified $\text{K}_2\text{Cr}_2\text{O}_7$.
 (iii) O_2 dissolves in pyrogallol.
- (A) $\text{Cd}^{2+} (\text{aq}) + \text{H}_2\text{S} (\text{g}) \longrightarrow \text{CdS} \downarrow (\text{yellow}) + 2\text{H}^+ (\text{aq})$
 (B) $\text{Co}_3^{2-} + 4\text{Hg}^{2+} + 3\text{H}_2\text{O} \longrightarrow \text{Hg}_4\text{O}_3\text{CO}_3 \downarrow (\text{reddish-brown}) + 6\text{H}^+$
 $\text{HCO}_3^- (\text{aq})$ does not give precipitate.
 (C) $\text{SO}_3^{2-} + 3\text{Zn} + 8\text{H}^+ \longrightarrow \text{H}_2\text{S} \uparrow + 3\text{Zn}^{2+} + 3\text{H}_2\text{O}$
 (D) $5\text{SO}_2 + 2\text{IO}_3^- + 4\text{H}_2\text{O} \longrightarrow \text{I}_2 + 5\text{SO}_4^{2-} + 8\text{H}^+$
- $\text{Br}^- + \text{H}_2\text{SO}_4 \longrightarrow \text{HBr} + \text{HSO}_4^-$;
 $\text{HBr} + \text{H}_2\text{SO}_4 \longrightarrow \text{Br}_2 + \text{SO}_2 + 2\text{H}_2\text{O}$
- $\text{Ag}^+ + \text{Br}^- \longrightarrow \text{AgBr} \downarrow (\text{pale yellow})$;
 $\text{NaBr} + \text{Na}_2\text{CO}_3 \longrightarrow \text{No reaction.}$
 $2\text{NaBr} + 2\text{H}_2\text{SO}_4 \longrightarrow \text{Br}_2 \uparrow (\text{reddish brown}) + \text{SO}_2 + \text{SO}_4^{2-} + 2\text{Na}^+ + 2\text{H}_2\text{O}$
- $3\text{AgCl} + \text{Na}_3\text{AsO}_3 \longrightarrow \text{Ag}_3\text{AsO}_3 \downarrow (\text{yellow}) + 3\text{Cl}^- + 3\text{Na}^+$
 $\text{AsO}_4^{3-} + 3\text{Ag}^+ \longrightarrow \text{Ag}_3\text{AsO}_4 \downarrow (\text{brownish-red})$
- $2\text{NO}_3^- + 4\text{H}_2\text{SO}_4 + 6\text{Fe}^{2+} \longrightarrow 6\text{Fe}^{3+} + 2\text{NO} \uparrow + 4\text{SO}_4^{2-} + 4\text{H}_2\text{O}$
 $\text{Fe}^{2+} + \text{NO} \uparrow + \text{SO}_4^{2-} \longrightarrow [\text{Fe}(\text{NO})]^{2+}\text{SO}_4^{2-}$
- (A) $3\text{AgCl} \downarrow + \text{AsO}_3^{3-} \longrightarrow \text{Ag}_3\text{AsO}_3 \downarrow (\text{yellow}) + 3\text{Cl}^-$
 AgI is unaffected by this treatment.
 (B) $\text{AgCl} + 2\text{NH}_3 \longrightarrow [\text{Ag}(\text{NH}_3)_2]\text{Cl}$
 AgI is not soluble in dilute ammonia solution.
 (C) Both soluble in potassium cyanide, forming soluble complexes.
 (D) Both insoluble in dilute HNO_3 .
- $\text{OCl}^- + 3\text{I}^- + \text{H}_2\text{O} \longrightarrow \text{I}_3^- + 2\text{OH}^- + \text{Cl}^-$
 $\text{I}_3^- + \text{starch} \longrightarrow \text{blue-black spot on starch paper due to the formation of iodine-starch adsorption complex.}$
- $2\text{I}^- + \text{Cl}_2 \longrightarrow \text{I}_2 (\text{violet}) + 2\text{Cl}^-$
 $5\text{Cl}_2 (\text{excess}) + \text{I}_2 + 6\text{H}_2\text{O} \longrightarrow 2\text{HIO}_3 (\text{colourless}) + 10\text{HCl}$
- $\text{Ba}^{2+} + \text{CrO}_4^{2-} \longrightarrow \text{BaCrO}_4 \downarrow (\text{yellow})$;
 $\text{Ag}^+ + \text{Cl}^- \longrightarrow \text{AgCl} \downarrow (\text{white})$
- $\text{NH}_4^+ + [\text{PtCl}_6]^{2-} \longrightarrow (\text{NH}_4)_2[\text{PtCl}_6] \downarrow (\text{yellow})$
 $[\text{Co}(\text{NO}_2)_6]^{3-} + 3\text{NH}_4^+ \longrightarrow (\text{NH}_4)_3[\text{Co}(\text{NO}_2)_6] \downarrow (\text{yellow})$
- $\text{NH}_4^+ + \text{OH}^- \xrightarrow{\Delta} \text{NH}_3 \uparrow + \text{H}_2\text{O}$
 (A) NH_3 , alkaline in nature turns red litmus blue;
 $\text{NH}_3 + \text{HCl} \longrightarrow \text{NH}_4\text{Cl} (\text{white fumes})$
 (B) $2\text{HgNO}_3 + 2\text{NH}_3 \longrightarrow \underset{\text{black}}{\text{Hg}_4(\text{NH}_2)_2\text{NO}_3} + \underset{\text{black}}{\text{Hg}_4\text{NH}_2} + \text{NH}_4\text{NO}_3$
 $\text{CuSO}_4 + 4\text{NH}_3 \longrightarrow [\text{Cu}(\text{NH}_3)_4]\text{SO}_4 (\text{intense blue})$
 (C) $2\text{K}_2(\text{HgI}_4) + \text{NH}_3 + 3\text{KOH} \longrightarrow \text{HgOHgNH}_2\text{I} \downarrow (\text{brown}) + 7\text{KI} + 2\text{H}_2\text{O}$
- $\text{Pb}^{2+} + \text{H}_2\text{S} \longrightarrow \text{PbS} \downarrow (\text{black}) + 2\text{H}^+$;
 $3\text{PbS} + 8\text{HNO}_3 \longrightarrow 3\text{Pb}(\text{NO}_3)_2 + 2\text{NO} \uparrow + 3\text{S} \downarrow + 4\text{H}_2\text{O}$
 $\text{Pb}^{2+} + \text{SO}_4^{2-} \longrightarrow \text{PbSO}_4 \downarrow (\text{white})$
- $\text{Pb}(\text{NO}_3)_2 + 2\text{NH}_4\text{OH} \longrightarrow \text{Pb}(\text{OH})_2 \downarrow (\text{white}) + 2\text{NH}_4\text{NO}_3$
 $\text{Pb}(\text{NO}_3)_2 + 2\text{NaCl} \longrightarrow \text{PbCl}_2 \downarrow (\text{white}) + 2\text{NaNO}_3$
 $\text{Pb}(\text{NO}_3)_2 + \text{H}_2\text{S} \longrightarrow \text{PbS} \downarrow (\text{black}) + 2\text{NaNO}_3$
- $\text{AgCl} + 2\text{CN}^- \longrightarrow [\text{Ag}(\text{CN})_2]^- + \text{Cl}^-$;
 $\text{AgCl} + 2\text{S}_2\text{O}_3^{2-} \longrightarrow [\text{Ag}(\text{S}_2\text{O}_3)_2]^{3-} + \text{Cl}^-$
 $\text{AgCl} + 2\text{NH}_3 \longrightarrow [\text{Ag}(\text{NH}_3)_2]^+ + \text{Cl}^-$
 AgCl is soluble in concentrated solution of KCl .
 $\text{AgCl} + \text{Cl}^- \longrightarrow [\text{AgCl}_2]^- (\text{soluble complex})$

32. $\text{HgI}_2 \downarrow (\text{red}) + 2\text{I}^- \rightarrow [\text{HgI}_4]^{2-}$ (soluble colourless complex);
 $\text{BiI}_3 \downarrow (\text{black}) + \text{I}^- \rightarrow [\text{BiI}_4]^-$ (soluble orange complex).
34. $\text{Pb}^{2+} + \text{CrO}_4^{2-} \longrightarrow \text{PbCrO}_4 \downarrow (\text{yellow});$
 $\text{PbCrO}_4 \downarrow + 4\text{OH}^- \longrightarrow [\text{Pb}(\text{OH})_4]^{2-} + \text{CrO}_4^{2-}.$
36. $\text{H}_2\text{S} \rightleftharpoons 2\text{H}^+ + \text{S}^{2-}; \text{HCl} \rightleftharpoons \text{H}_2^+ + \text{Cl}^-.$
 Due to common ion effect, the ionisation of H_2S is suppressed and thus low concentration of S^{2-} ions is obtained. This much of S^{2-} ions concentration is enough to precipitate only IInd group cations (because of the low K_{sp} of IInd group sulphides).
40. (A) $\text{Hg}^{2+} + \text{Co}^{2+} + 4\text{SCN}^- \longrightarrow \text{Co}[\text{Hg}(\text{SCN})_4] \downarrow (\text{deep blue});$
 (B) $2\text{Hg}^{2+} + \text{Sn}^{2+} + 2\text{Cl}^- \longrightarrow \text{Hg}_2\text{Cl}_2 \downarrow (\text{white}) + \text{Sn}^{4+};$
 $\text{Hg}_2\text{Cl}_2 + \text{Sn}^{2+} \longrightarrow \text{Hg} \downarrow (\text{black}) + \text{Sn}^{4+} + 2\text{Cl}^-$
 (C) $2\text{Hg}^{2+} + \text{NO}_3^- + 4\text{NH}_3 + \text{H}_2\text{O} \longrightarrow \text{HgO} \cdot \text{Hg}(\text{NH}_2)\text{NO}_3 \downarrow (\text{white})$
 (D) KCN no effect i.e. no reaction.
42. (A) $\text{Bi}^{3+} + 3\text{I}^- \longrightarrow \text{BiI}_3 \downarrow (\text{black})$
 (B) $2\text{Cu}^{2+} + 5\text{I}^- \longrightarrow \text{Cu}_2\text{I}_2 \downarrow (\text{white}) + \text{I}_3^- (\text{brown solution})$
 (C) $\text{Pb}^{2+} + 2\text{I}^- \longrightarrow \text{PbI}_2 \downarrow (\text{yellow})$
 (D) $\text{Hg}_2^{2+} + 2\text{I}^- \longrightarrow \text{HgI}_2 (\text{red/scarlet})$
44. (A) $\text{Cu}^{2+} + \text{S}^{2-} \longrightarrow \text{CuS} \downarrow (\text{black}),$
 (B) $\text{Cd}^{2+} + \text{S}^{2-} \longrightarrow \text{CdS} \downarrow (\text{yellow}),$
 (C) $\text{Zn}^{2+} + \text{S}^{2-} \longrightarrow \text{ZnS} \downarrow (\text{white}),$
 (D) $2\text{Fe}^{3+} + \text{H}_2\text{S} \longrightarrow 2\text{Fe}^{2+} + 2\text{H}^+ + \text{S} \downarrow (\text{milky white})$
45. $2\text{CuSO}_4 + 4\text{KI} \longrightarrow 2\text{CuI} \downarrow (\text{white}) + 2\text{K}_2\text{SO}_4 + \text{I}_2$
 $3\text{SCN}^- + \text{Cu}^{2+} \longrightarrow \text{CuSCN} \downarrow (\text{white}) + (\text{SCN})_2$
 $6\text{CN}^- + \text{Cu}^{2+} \longrightarrow [\text{Cu}(\text{CN})_4]^{3-} + (\text{CN})_2$
47. Both dissolves in acids. $\text{Cr}(\text{OH})_3$ is partially soluble while $\text{Fe}(\text{OH})_3$ is not soluble in aqueous NH_3 . Only $\text{Cr}(\text{OH})_3$ not $\text{Fe}(\text{OH})_3$ is soluble by $\text{NaOH}/\text{H}_2\text{O}_2$ according to the reaction.
 $\text{Cr}(\text{OH})_3 \downarrow (\text{green}) \xrightarrow{\text{NaOH}/\text{H}_2\text{O}_2} \text{CrO}_4^{2-} (\text{yellow solution}) + \text{H}_2\text{O}.$
50. $\text{Fe}^{3+} + [\text{Fe}(\text{CN})_6]^{3-} \longrightarrow \text{Fe}[\text{Fe}(\text{CN})_6] \text{ brown colouration}$
 $3\text{Fe}^{3+} + 6\text{CH}_3\text{COO}^- + 2\text{H}_2\text{O} \rightleftharpoons [\text{Fe}_3(\text{OH})_2(\text{CH}_3\text{COO})_6]^{+} + 2\text{H}^+.$
52. $\text{Fe}^{3+} + 3\text{SCN}^- \longrightarrow \text{Fe}(\text{SCN})_3 (\text{deep red colouration})$
 $\text{Co}^{2+} + 4\text{SCN}^- \xrightarrow{\text{amyl alcohol}} [\text{Co}(\text{SCN})_4]^{2-}$ (blue colouration)
 $\text{Cu}^{2+} + 2\text{SCN}^- \longrightarrow \text{Cu}(\text{SCN})_2 \downarrow (\text{black});$
 $2\text{Cu}(\text{SCN})_2 \longrightarrow 2\text{CuSCN} \downarrow (\text{white}) + (\text{SCN})_2.$
53. $\text{Fe}^{2+} + [\text{Fe}(\text{CN})_6]^{4-} \rightarrow$ white precipitate ;
 $\text{Fe}^{3+} + [\text{Fe}(\text{CN})_6]^{4-} \rightarrow$ Prussian blue precipitate.
 $\text{Zn}^{2+} + [\text{Fe}(\text{CN})_6]^{4-} \rightarrow$ Bluish white/white precipitate ;
 $\text{Cu}^{2+} + [\text{Fe}(\text{CN})_6]^{4-} \rightarrow$ chocolate brown precipitate
 $\text{Ag}^+ + [\text{Fe}(\text{CN})_6]^{4-} \rightarrow$ white precipitate ;
 $\text{Ca}^{2+} + \text{K}_4[\text{Fe}(\text{CN})_6] \rightarrow$ white precipitate.
54. $\text{Zn}^{2+} + 2\text{NH}_4\text{OH} \longrightarrow \text{Zn}(\text{OH})_2 \downarrow (\text{white}) + 2\text{NH}_4^+ ;$
 $\text{Zn}(\text{OH})_2 + 4\text{NH}_4\text{OH} \longrightarrow \text{Zn}(\text{NH}_3)_4 + 6\text{H}_2\text{O}$
 $\text{Zn}^{2+} + \text{S}^{2-} \longrightarrow \text{ZnS} (\text{white})$
56. $\text{Na}_2\text{B}_4\text{O}_7 \cdot 10\text{H}_2\text{O} \xrightarrow{\Delta} \text{Na}_2\text{B}_4\text{O}_7 + 10\text{H}_2\text{O} ;$
 $\text{Na}_2\text{B}_4\text{O}_7 \xrightarrow{\Delta} 2\text{NaBO}_2 + \text{B}_2\text{O}_3$
 $\text{CuO} + \text{B}_2\text{O}_3 \xrightarrow{\Delta} \text{Cu}(\text{BO}_2)_2 (\text{copper (II) metaborate}) -$
 blue bead in oxidising flame.
58. $2\text{CoS} + 6\text{HCl} + 2\text{HNO}_3 \longrightarrow 3\text{CoCl}_2 + 2\text{NO} + 3\text{S} + 4\text{H}_2\text{O}$
 $\text{CoCl}_2 + 6\text{NaHCO}_3 \longrightarrow \text{Na}_4[\text{Co}(\text{CO}_3)_3] + 2\text{NaCl} + 3\text{H}_2\text{O} + 3\text{CO}_2$
 $2\text{Na}_4[\text{Co}(\text{CO}_3)_3] + \text{Br}_2 \longrightarrow 2\text{Na}_3[\text{Co}(\text{CO}_3)_3] + 2\text{NaBr}$ (Green)
60. (A) $\text{Zn}(\text{OH})_2 \downarrow + 2\text{OH}^- \longrightarrow [\text{Zn}(\text{OH})_4]^{2-}$
 (B) $\text{Zn}(\text{OH})_2 \downarrow + 2\text{H}^+ \rightleftharpoons \text{Zn}^{2+} + 2\text{H}_2\text{O}$
 (C) & (D) White precipitate of $\text{Zn}(\text{OH})_2$ dissolves in excess of ammonia solution and in solutions of ammonium salts owing to the production of tetraamminezincate(II).
 $\text{Zn}(\text{OH})_2 \downarrow + 4\text{NH}_3 \rightleftharpoons [\text{Zn}(\text{NH}_3)_4]^{2+} + 2\text{OH}^-$
61. Strontium chloride gives crimson colour flame in Bunsen burner.
 BaCl_2 -apple green, CaCl_2 -brick red, KCl -lilac(violet).
63. (A) titan yellow is absorbed by magnesium hydroxide producing a deep-red colour or precipitate.
 (B) $\text{Mg}^{2+} + \text{NH}_3 + \text{HPO}_4^{2-} \longrightarrow \text{Mg}(\text{NH}_4)\text{PO}_4 \downarrow (\text{white}).$
 (C) Blue lake is formed by the adsorption of reagent on $\text{Mg}(\text{OH})_2.$
64. $\text{Ba}^{2+} + \text{CrO}_4^{2-} \longrightarrow \text{BaCrO}_4 \downarrow (\text{yellow});$
 $\text{Ag}^+ + \text{Br}^- \longrightarrow \text{AgBr} \downarrow (\text{pale yellow}).$

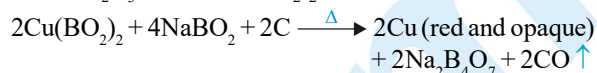
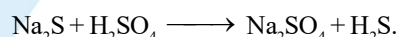
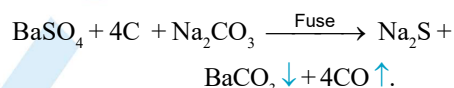
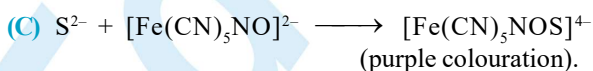
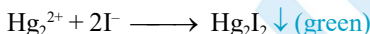
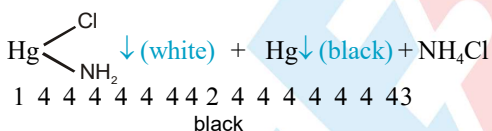
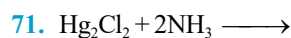
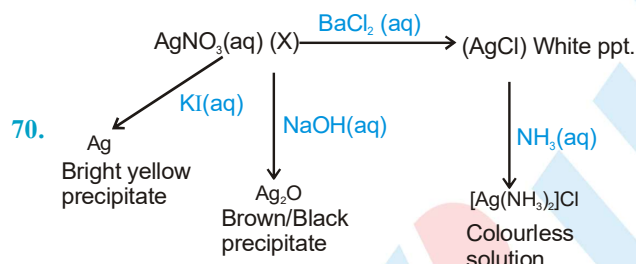
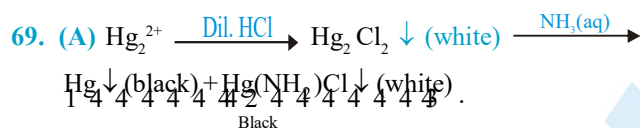
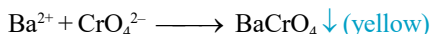
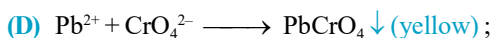
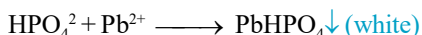
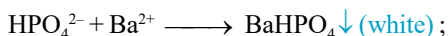
SALT ANALYSIS AND QUALITATIVE ANALYSIS



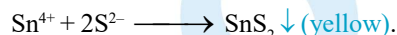
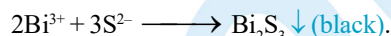
BaCl_2 is soluble in water

(B) Na_2SO_4 gives white precipitate of PbSO_4 and BaSO_4 with $\text{Pb(NO}_3)_2$ and $\text{Ba(NO}_3)_2$ respectively.

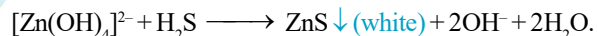
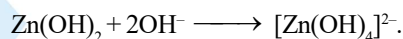
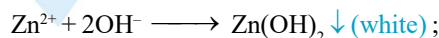
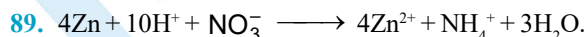
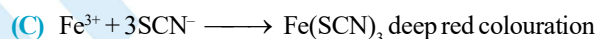
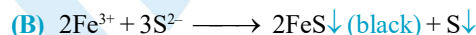
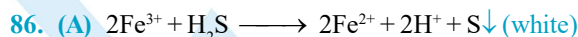
(C) Disodium hydrogen phosphate gives white precipitate with both salts.



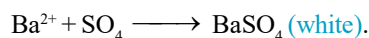
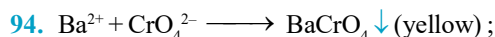
77. Both belong to same group i.e. IInd group and their K_{sp} values are low ; so both are precipitated according to the following reactions.



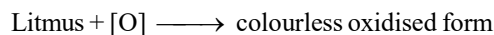
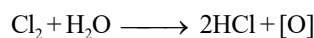
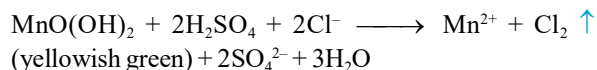
83. As it provides NH_4^+ ions which suppresses the ionisation of NH_4OH so that only the group 3rd cations are precipitated as hydroxides because of their low solubility products and NO_3^- ions do not produce precipitate with the cations of IVth, Vth and VIth groups.



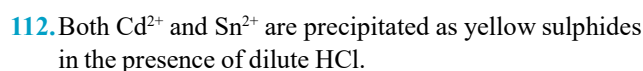
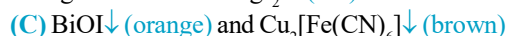
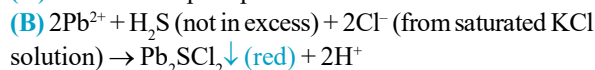
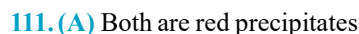
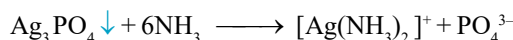
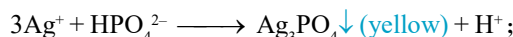
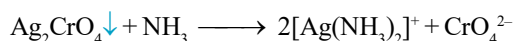
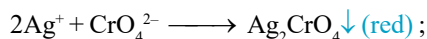
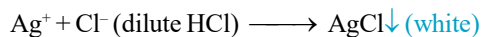
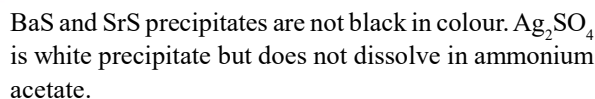
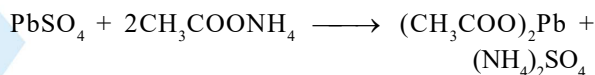
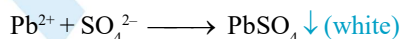
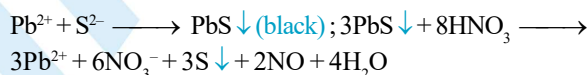
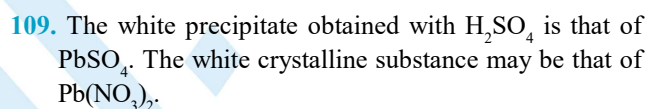
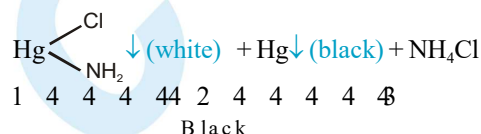
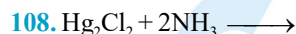
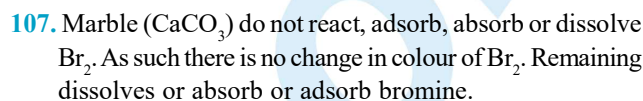
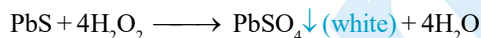
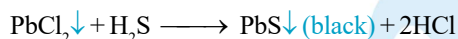
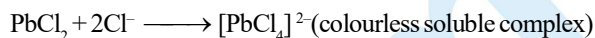
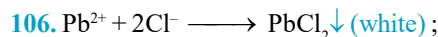
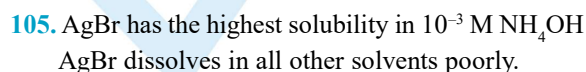
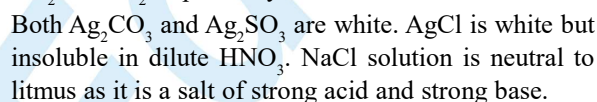
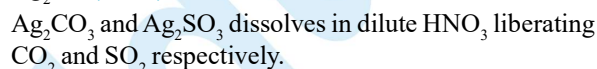
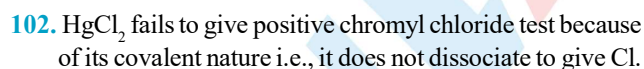
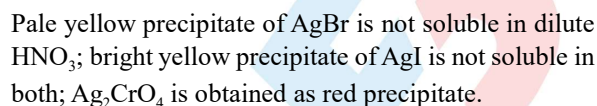
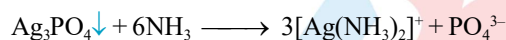
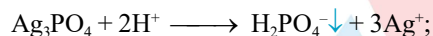
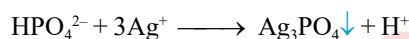
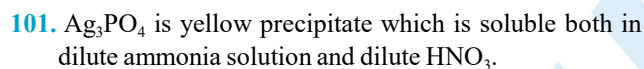
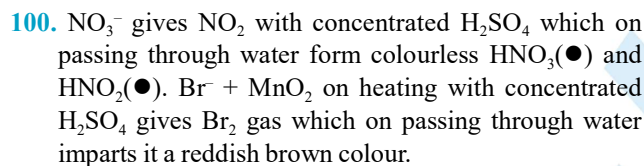
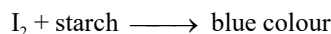
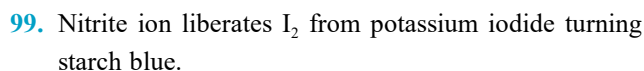
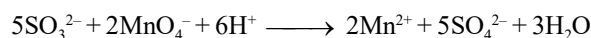
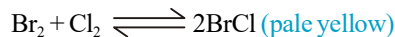
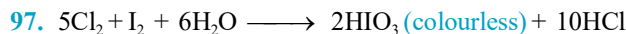
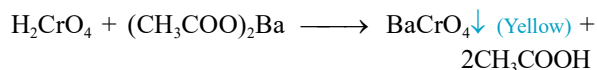
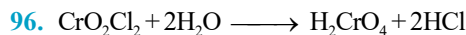
92. Ni^{2+} and Fe^{2+} both on reaction with alkaline solution of dimethyl glyoxime give red precipitate and red solution respectively but not zinc.

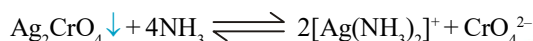
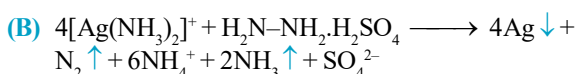
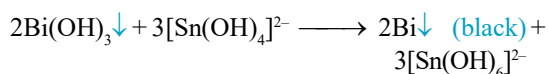
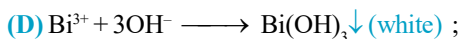
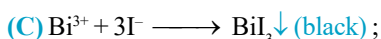
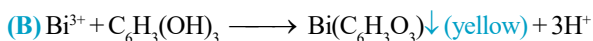
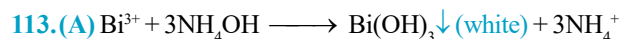


K_{sp} of SrCrO_4 is high in acetic acid, so no precipitate is formed. Lead carbonate and basic lead carbonate both gives precipitate with K_2CrO_4 and NaCl .



Cl_2 is a yellowish green gas which bleaches litmus paper by oxidation.

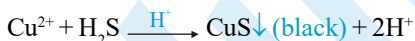
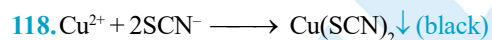
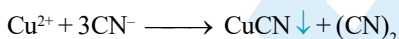
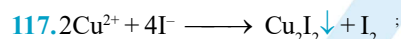




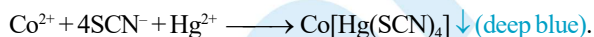
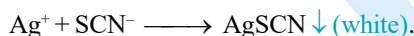
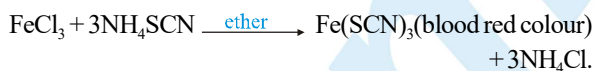
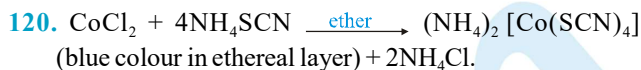
So, all statements are correct.

115. The filter paper ash test is substitute for cobalt nitrate charcoal cavity test. Double oxide $\text{ZnO} \cdot \text{CoO}$ formed is green in colour. It is called Rinmann's green.

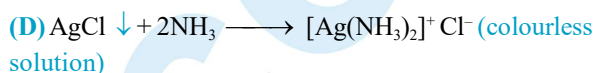
116. $\text{Cd}^{2+} + \text{H}_2\text{S} \longrightarrow \text{CdS} \downarrow + 2\text{H}^+$, reaction is reversible; if the concentration of strong acid in the solution is above 0.5 M, precipitation is incomplete. Concentrated acid dissolves the precipitate for the same reason.



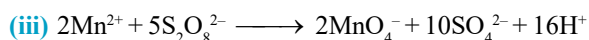
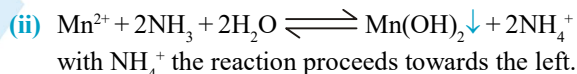
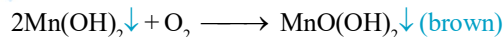
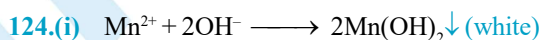
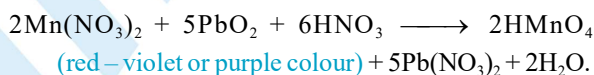
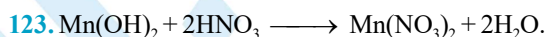
119. Function of strong electrolyte NH_4Cl is to suppress the ionisation of NH_4OH so that the concentration of OH^- ions in the solution is decreased but it is sufficient to precipitate the third group basic radicals because the solubility product of group III hydroxides is lower than IV, V and VI group hydroxides. The $\text{Cr}(\text{OH})_3 \downarrow$ is slightly soluble in excess of precipitant, upon boiling the solution, $\text{Cr}(\text{OH})_3$ is precipitated.



121. (A), (B) and (C) all gives blue colouration in solution or blue precipitate.



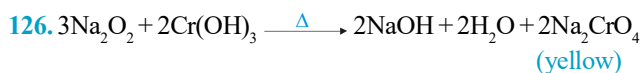
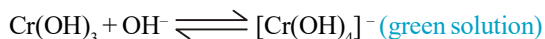
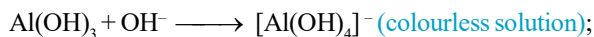
It in etherial layer develops blue colouration.



125. In first step with H_2S gas in acidic medium Cd^{2+} and Cu^{2+} will precipitate as CdS and CuS respectively. Thus the filtrate will now contain chromium, iron and aluminium.

In second step with NH_4Cl and NH_4OH , Cr^{3+} , Fe^{3+} and Al^{3+} will precipitate out as their hydroxides.

In third step on addition of excess of NaOH , $\text{Al}(\text{OH})_3$ and $\text{Cr}(\text{OH})_3$ precipitates will dissolve forming soluble complexes but $\text{Fe}(\text{OH})_3$ precipitate is unaffected. Hence the filtrate will contain $\text{Na}[\text{Al}(\text{OH})_4]$ and $\text{Na}[\text{Cr}(\text{OH})_4]$.



127. On adding H_2O_2 in alkaline medium or SnCl_2 solution in acidic medium, the $[\text{Fe}(\text{CN})_6]^{3-}$ part of the compound is reduced and prussian blue is precipitated.

128. (A) $\text{Zn(OH)}_2 \downarrow + 2\text{OH}^- \rightleftharpoons [\text{Zn(OH)}_4]^{2-}$
 (B) and (C) $\text{Zn(OH)}_2 \downarrow + 4\text{NH}_3 \rightleftharpoons [\text{Zn(NH}_3)_4]^{2+} + 2\text{OH}^-$ or NH_4^+
129. Ba^{2+} salts gives yellow precipitate with K_2CrO_4 solution and this precipitate is not soluble in CH_3COOH . Ba^{2+} ions gives apple green colour in the flame test.
130. (B) $\text{BaCO}_3 + \text{ZnS}$ mixture dissolves in HCl but is insoluble in water. Further the solution in HCl will be colourless due to the formation of soluble BaCl_2 and ZnCl_2 .
131. (A) No precipitate with K_2CrO_4 in acetic acid as its k_{sp} is high.
 (B) $\text{Ca}^{2+} + 2\text{K}^+ + [\text{Fe(CN)}_6]^{4-} \longrightarrow \text{K}_2\text{Ca}[\text{Fe(CN)}_6] \downarrow$ (white)
 (C) It imparts brick red colour to Bunsen flame.
 (D) $\text{Ca(HCO}_3)_2$ is formed which is water soluble.

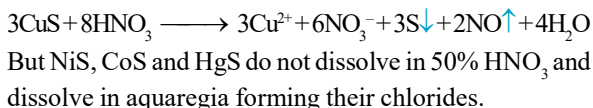
EXERCISE - 2

Part # 1 : Multiple Choice

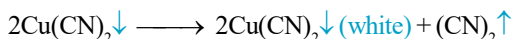
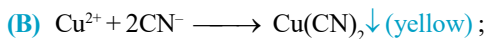
4. (A) $\text{Na}_2\text{S} + \text{H}^+ \longrightarrow \text{H}_2\text{S} \uparrow + 2\text{Na}^+$;
 Na_2SO_4 gives no reaction with H_2SO_4 .
 (B) 2MnO_4^- (pink solution) + $5\text{H}_2\text{S} + 6\text{H}^+ \longrightarrow 2\text{Mn}^{2+}$ (colourless) + $5\text{S} \downarrow + 8\text{H}_2\text{O}$.
 No colour change is observed with Na_2SO_4 .
 (C) $\text{S}^{2-} + [\text{Fe(CN)}_5\text{NO}]^{2-} \longrightarrow [\text{Fe(CN)}_5\text{NOS}]^{4-}$ (purple or violet colouration).
 No colour change is observed with Na_2SO_4 .
 (D) $\text{S}^{2-} + \text{Cd}^{2+} \longrightarrow \text{CdS} \downarrow$ (yellow).
 Na_2SO_4 forms CdSO_4 which is water soluble.
5. (A) $\text{Br}^- + \text{H}^+ \longrightarrow \text{HBr} \uparrow$;
 $2\text{HBr} + 2\text{H}_2\text{SO}_4 \longrightarrow \text{Br}_2 \uparrow$ (reddish brown) + $\text{SO}_2 \uparrow$ + $\text{SO}_4^{2-} + \text{H}_2\text{O}$.
 (B) $4\text{NO}_3^- + 2\text{H}_2\text{SO}_4 \longrightarrow 4\text{NO}_2 \uparrow$ (reddish brown) + $\text{O}_2 \uparrow$ + $2\text{SO}_4^{2-} + 2\text{H}_2\text{O}$.
 (C) $\text{SO}_3^{2-} + 2\text{H}^+ \longrightarrow \text{SO}_2 \uparrow$ (colourless) + H_2O .
 (D) $3\text{I}^- + 2\text{H}_2\text{SO}_4 \longrightarrow \text{I}_3^- \uparrow$ (violet/purple) + $\text{SO}_4^{2-} + 2\text{H}_2\text{O} + \text{SO}_2$.

9. (A) AgCl dissolves completely forming $[\text{Ag(NH}_3)_2]\text{Cl}$;
 $\text{AgCl} + 2\text{NH}_4\text{OH} \longrightarrow [\text{Ag(NH}_3)_2]\text{Cl} + 2\text{H}_2\text{O}$.
 (B) AgBr dissolves completely forming $[\text{Ag(NH}_3)_2]\text{Br}$ soluble complex.
 $\text{AgBr} + 2\text{NH}_4\text{OH} \longrightarrow [\text{Ag(NH}_3)_2]\text{Br} + 2\text{H}_2\text{O}$.
 (C) $\text{Ag}_2\text{CrO}_4 + 4\text{NH}_3 \longrightarrow 2[\text{Ag(NH}_3)_2]^+ + \text{CrO}_4^{2-}$
 (D) AgI is insoluble in concentrated aqueous NH_3 .
11. (A) $\text{Ag}^+ + 2\text{CrO}_4^{2-} \longrightarrow \text{Ag}_2\text{CrO}_4 \downarrow$ (red) [X].
 (B) $\text{Ag}_2\text{CrO}_4 \downarrow + 4\text{NH}_3 \longrightarrow 2[\text{Ag(NH}_3)_2]^+ + \text{CrO}_4^{2-}$.
 (C) $\text{Cr}^{3+} + 3\text{OH}^- \longrightarrow \text{Cr(OH)}_3 \downarrow$ (green).
 $\text{Cr(OH)}_3 + \text{OH}^- \longrightarrow [\text{Cr(OH)}_4]^-$ (green coloured soluble complex).
 (D) $3\text{SO}_2 + \text{Cr}_2\text{O}_7^{2-} + 2\text{H}^+ \longrightarrow 2\text{Cr}^{3+}[\text{Y}] + 3\text{SO}_4^{2-} + \text{H}_2\text{O}$.
13. (A) Fe^{2+} responds to this test but not Fe^{3+} ; Fe(II) gives soluble red iron(II) dimethylglyoxime in alkaline solution.
 (D) $\text{Ag}_2\text{O} \downarrow + 4\text{NH}_3 + \text{H}_2\text{O} \longrightarrow 2[\text{Ag(NH}_3)_2]^+ + 2\text{OH}^-$
 $\text{Ag}_2\text{O} \downarrow + 2\text{H}^+ \longrightarrow 2\text{Ag}^+ + \text{H}_2\text{O}$
 (B) and (C) are correct statements.
14. (A) Red colour solution or precipitate is produced when reagent reacts in alkaline solution.
 (B) $\text{HO.SO}_2.\text{NH}_2 + \text{HNO}_2 \longrightarrow \text{N}_2 + 2\text{H}^+ + \text{SO}_4^{2-} + \text{H}_2\text{O}$.
 (C) $\text{Fe}^{2+} + [\text{Fe(CN)}_6]^{3-} \longrightarrow \text{Fe}^{3+} + [\text{Fe(CN)}_6]^{4-}$.
 $4\text{Fe}^{3+} + [\text{Fe(CN)}_6]^{4-} \longrightarrow \text{Fe}_4[\text{Fe(CN)}_6]_3 \downarrow$ (Turnbull's blue).
 (D) $4\text{Cr(OH)}_3 \downarrow + 5\text{O}_2^{2-} \longrightarrow 4\text{CrO}_4^{2-}$ (yellow solution) + $6\text{H}_2\text{O}$.
15. (A) $\text{PbSO}_4 \downarrow + \text{H}_2\text{SO}_4$ (hot and concentrated) $\longrightarrow \text{Pb}^{2+} + 2\text{HSO}_4^-$ (soluble)
 (B) It dissolves forming $\text{Na}_2[\text{Pb(OH)}_4]$ soluble complex.
16. $\text{B}_4\text{O}_7^{2-} + 4\text{Ag}^+ + \text{H}_2\text{O} \longrightarrow 4\text{AgBO}_2 \downarrow$ (white) + 2H^+
 $2\text{AgBO}_2 \downarrow + 3\text{H}_2\text{O} \xrightarrow[\Delta/\text{H}_2\text{O}]{\text{Hydrolysis}} \text{Ag}_2\text{O} \downarrow$ (brown) + $2\text{H}_3\text{BO}_3$

17. CuS dissolves in 50% HNO₃ ;



21. (A) Correct.

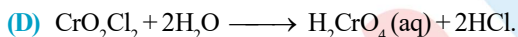


22. (NH₄)₂ CO₃ and (NH₄)₂ SO₄ can not be used as they would also precipitate the IV and Vth group cations.

23. (A) $\text{Cr}^{3+} + 3\text{NH}_3 + 3\text{H}_2\text{O} \longrightarrow \text{Cr}(\text{OH})_3\downarrow + 3\text{NH}_4^+$
The reaction is reversible; on addition of NH₄⁺, shifts to backward direction. Thus if excess of NH₄⁺ salt is added, then precipitation of Cr(OH)₃ will not take place. However, because of very small k_{sp} of iron (III) hydroxide complete precipitation will take place even in the presence of ammonium salts. (k_{sp} = 3.8 × 10⁻³⁸)

(B) Concentration of CO₃²⁻ provided by Na₂CO₃ in aqueous solution is just sufficient to precipitate Mg²⁺ ion as MgCO₃ along with Ba²⁺, Ca²⁺ and Sr²⁺ as their carbonates.

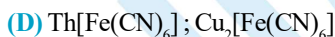
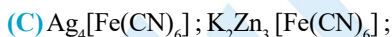
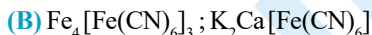
(C) The oxidising anions like MnO₄⁻, Cr₂O₇²⁻, ClO₄⁻, etc., also respond to this test.



24. $\text{Co}^{2+} + 2\text{CN}^- \longrightarrow \text{Co}(\text{CN})_2\downarrow$ (reddish - brown)

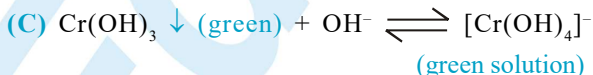


25. (A) Cu₂[Fe(CN)₆] ; K₂Zn₃[Fe(CN)₆]₂ ;



26. (A) $\text{Hg}^{2+} + \text{Co}^{2+} + 4\text{SCN}^- \longrightarrow \text{Co}[\text{Hg}(\text{SCN})_4]\downarrow$ (deep blue)

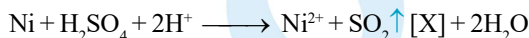
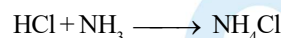
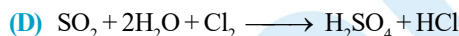
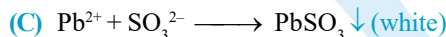
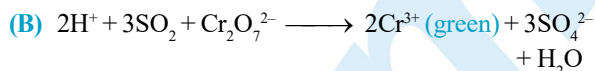
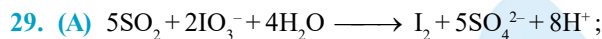
(B) Soluble in NaOH forming [Al(OH)₄]⁻, not in NH₃ (aq)



(D) Correct statement.

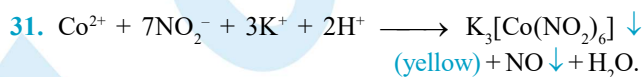
27. Iron and NO exist as Fe(II) and NO⁺ respectively.

28. Cr = Green ; Co = Blue.



With dilute H₂SO₄, hydrogen gas is liberated.

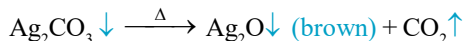
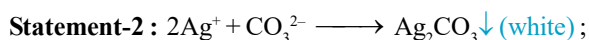
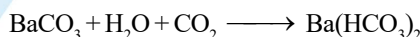
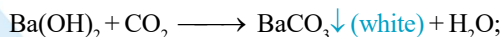
30. BF₃ colour the flame green ; B(OC₂H₅)₃ burns with green edged flame ; Barium chloride (volatile) gives apple green colour to flame.



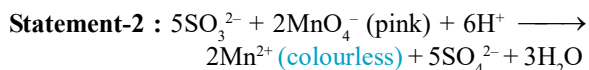
So [X] is K₃[Co(NO₂)₆]. It is called Fischer's reagent ; [Co(III)(NO₂)₆]³⁻ has d²sp³ hybridisation and is diamagnetic. It's IUPAC name is potassium hexanitrito – N – cobaltate(III).

Part # II : Assertion & Reason

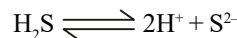
2. Statement-1 :



3. Statement-1 : $\text{Ag}_2\text{SO}_3\downarrow + \text{SO}_3^{2-} \longrightarrow 2[\text{Ag}(\text{SO}_3)]^-$ (soluble complex)



5. Statement-1 : $\text{Zn}^{2+} + \text{H}_2\text{S} \rightleftharpoons \text{ZnS} + 2\text{H}^+$;

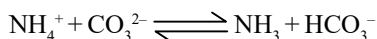


Sulphide ion concentration obtained from the H₂S is depressed so much by the hydrogen-ion concentration from acid that it is too low to exceed the solubility product of ZnS and consequently precipitation ceases.

Statement-2 : ZnS is insoluble in caustic alkali solution, but dissolves according to the following reaction in dilute HCl.

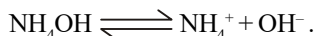


7. **Statement-1** : In the presence of NH_4^+ salts, no precipitation of Mg^{2+} occurs because the equilibrium

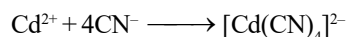


is shifted towards the formation of HCO_3^- ions. K_{sp} of the precipitate being high (K_{sp} of pure MgCO_3 is 1×10^{-5}), the concentration of carbonate ions necessary to produce a precipitate is not attained.

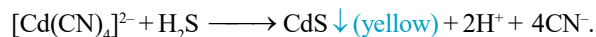
Statement-2 : NH_4OH turns red litmus blue.



9. **Statement-1** : $\text{Cu}^{2+} + 6\text{CN}^- \longrightarrow [\text{Cu}(\text{CN})_4]^{3-} + (\text{CN})_2$;



Statement-2 : $[\text{Cu}(\text{CN})_4]^{3-}$ being stable complex is not effected by H_2S gas but $[\text{Cd}(\text{CN})_4]^{2-}$ is not so stable and gives yellow precipitate with H_2S .



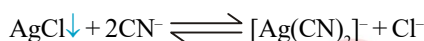
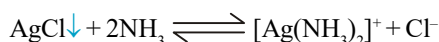
12. **Statement-1** : In presence of ammonium salts the reaction proceeds in backward direction forming ammonia gas.

Statement-2 : $\text{Mg}(\text{OH})_2 \downarrow$ is insoluble in water.

EXERCISE - 3

Part # I : Matrix Match Type

1. (A) $\text{AgCl} \downarrow + \text{Cl}^- \longrightarrow [\text{AgCl}_2]^-$ soluble complex.



AgCl is insoluble in dilute HNO_3 .

- (B) $2\text{CuS} \downarrow + 8\text{CN}^- \longrightarrow 2[\text{Cu}(\text{CN})_4]^{3-} + \text{S}_2^{2-}$.



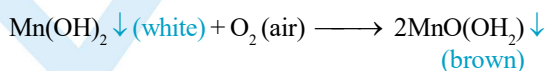
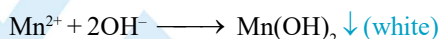
- (C) $\text{Zn}(\text{OH})_2 \downarrow + 2\text{H}^+ \rightleftharpoons \text{Zn}^{2+} + 2\text{H}_2\text{O}$



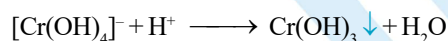
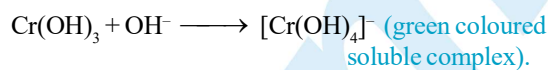
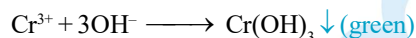
- (D) $\text{BaCO}_3 + 2\text{H}^+ \rightleftharpoons \text{Ba}^{2+} + \text{CO}_2 + \text{H}_2\text{O}$

BaCO_3 is slightly soluble in solution of ammonium salts of strong acids only.

3. (A) Mn^{2+} meta borate $\longrightarrow$ Amethyst red



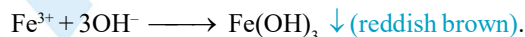
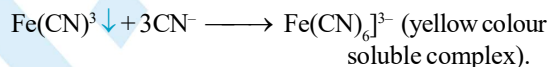
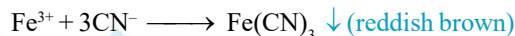
- (B) Cr^{3+} metaborate $\longrightarrow$ Green



- (C) $\text{Al}^{3+} + 3\text{OH}^- \longrightarrow \text{Al}(\text{OH})_3 \downarrow (\text{white})$



- (D) Fe^{3+} metaborate $\longrightarrow$ Pale yellow



Part # II : Comprehension

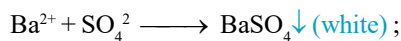
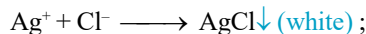
Comprehension # 1 :

1. Sample $\xrightarrow{\text{NaCl}}$ Precipitate 'A' insoluble chloride (AgCl) + Filtrate (Ba^{2+} and Zn^{2+})

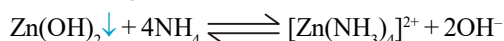
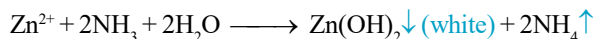


ppt. 'C' insoluble hydroxide ($\text{Zn}(\text{OH})_2$) $\xleftarrow[\text{NH}_3]{\text{aqueous}}$

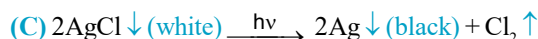
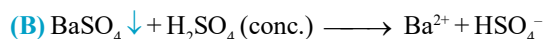
Filtrate (Zn^{2+}) + ppt. 'B' insoluble sulphate (BaSO_4)



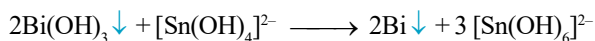
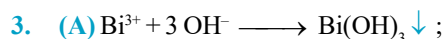
Ba^{2+} salts give apple green colour to the Bunsen flame.



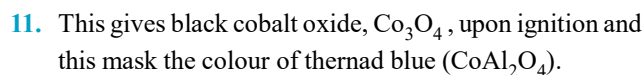
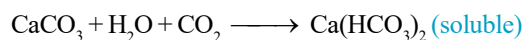
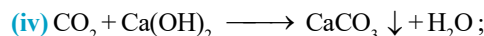
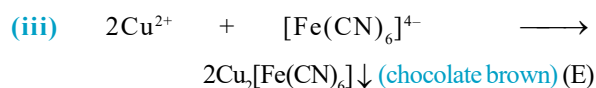
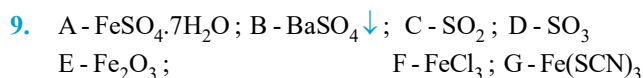
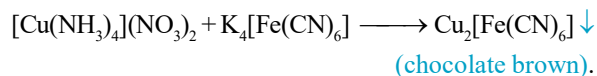
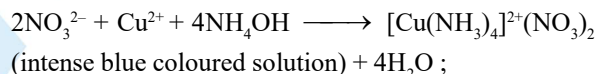
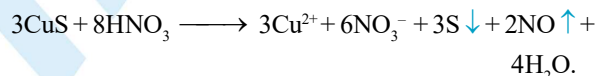
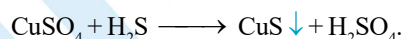
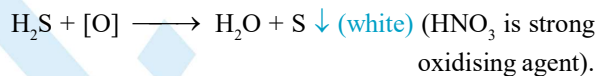
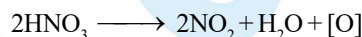
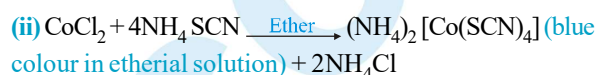
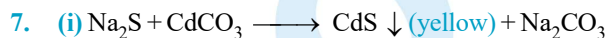
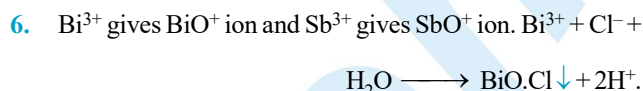
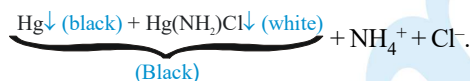
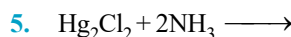
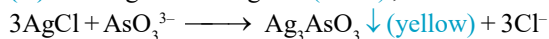
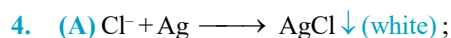
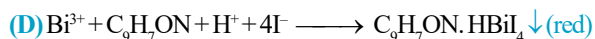
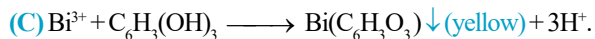
3. (A) $\text{CoO} \cdot \text{ZnO}$ = Rinmann's green reagent used as green pigment.



Comprehension # 2 :

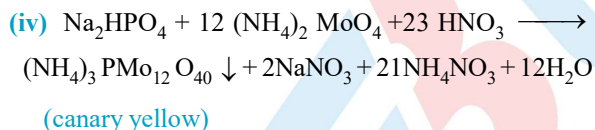
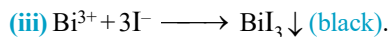
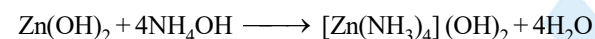
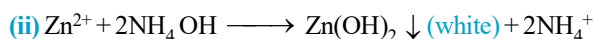
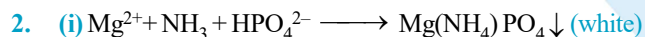
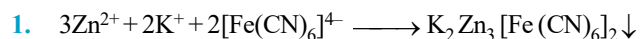


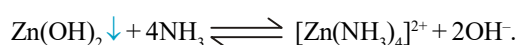
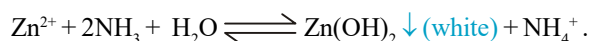
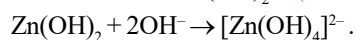
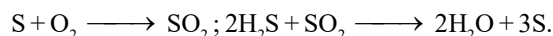
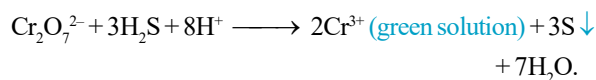
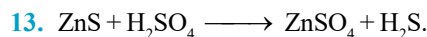
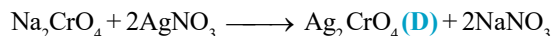
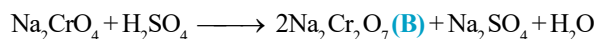
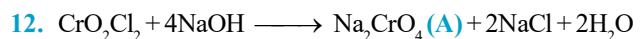
(B) is not correct as 'D' is precipitate of bismuth.



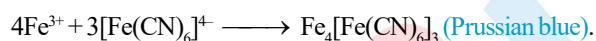
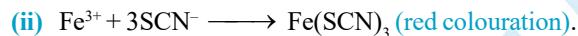
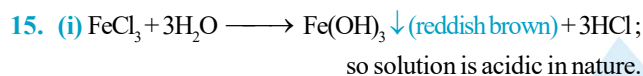
EXERCISE - 4

Subjective Type

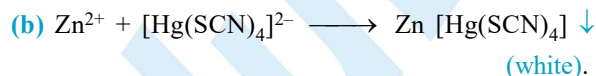




14. The given unknown mixture contains NH_4^+ , Fe^{2+} and Cl^- ions or NH_4Cl and FeCl_2

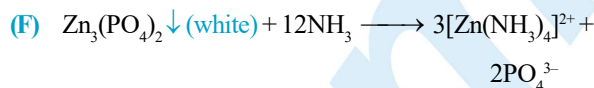
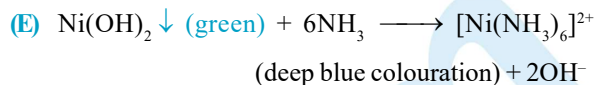


16. (a) Violet (or blackish-purple) precipitate consisting of $\text{Zn}[\text{Hg(SCN)}_4]$ and $\text{Cu}[\text{Hg(SCN)}_4]$ is formed.



Soluble in sodium hydroxide not in NH_3 .

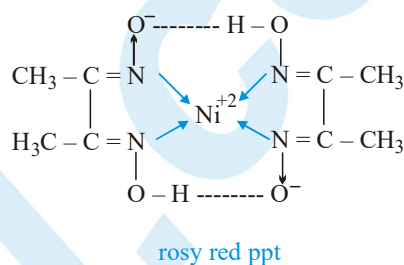
(D) Soluble in sodium hydroxide not in NH_3 .



(G) Insoluble in NH_3 , soluble appreciably in boiling concentrated H_2SO_4 .

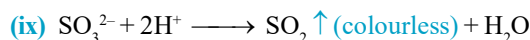
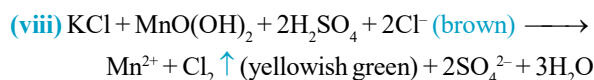
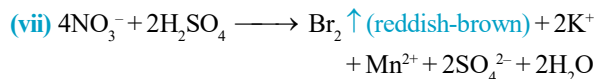
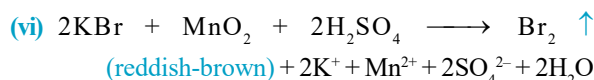
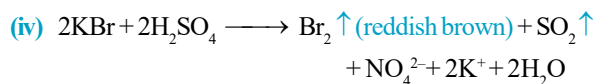
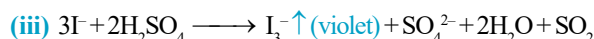
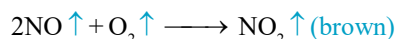
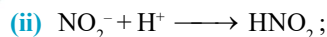
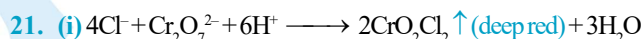
(H) $\text{Bi(OH)}_2\text{NO}_3$ is insoluble in NH_3 .

(I) Insoluble in NH_3 but soluble in ammonium salts liberating NH_3 .



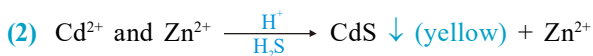
19. Here NO is NO^+ ; so in $[\text{Fe(H}_2\text{O)}_5\text{NO}]^{2+}$, the oxidation state of iron is +1.

20. NaCl-Golden yellow; KCl-Lilac; CuCl_2 -bluish-green, BaCl_2 -Apple green; SrCl_2 -Crimson; CaCl_2 -Brick red.





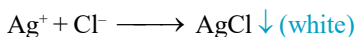
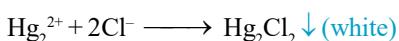
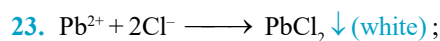
(in solution)



(in solution)

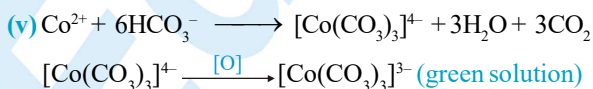
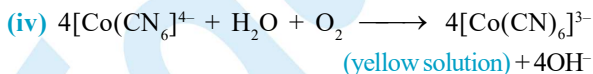
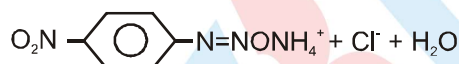
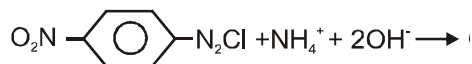


(in solution).



K_{sp} of chlorides of Pb^{2+} and Hg_2^{2+} is low as compared to K_{sp} of Hg^{2+} and Cd^{2+} . Chloride ion concentration provided by dilute HCl is just enough to exceed the K_{sp} of PbCl_2 and Hg_2Cl_2 . Thus Pb^{2+} and Hg_2^{2+} are precipitated as their chlorides.

24. A red colouration is obtained.



EXERCISE - 5

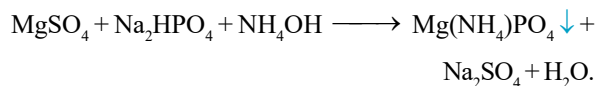
Part # I : AIEEE/JEE-MAIN

1. Methyl orange shows Red (pinkish) color in Acidic medium & yellow color in basic medium since original solution is basic so
 initial color $\Rightarrow$ yellow
 & Titrated with acid so
 Final color $\Rightarrow$ pinkish (red)

Part # II : IIT-JEE ADVANCED

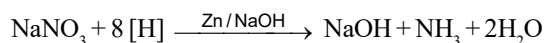
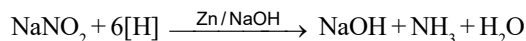
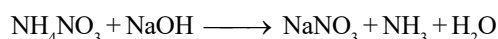
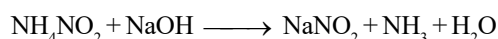
1. $\text{Mg}^{2+} + 2\text{HCO}_3^- (\text{from sodium salt}) \longrightarrow \text{Mg}(\text{HCO}_3)_2 (\text{soluble}) \xrightarrow{\Delta} \text{MgCO}_3 \downarrow (\text{white}) + \text{H}_2\text{O} + \text{CO}_2$
2. Hg^{2+} gives red precipitate of HgI_2 which dissolves in excess KI forming colourless $[\text{HgI}_4]^{2-}$ ions.
 $\text{Bi}^{3+} + 3\text{I}^- \longrightarrow \text{BiI}_3 \downarrow (\text{black})$;
 $\text{BiI}_3 + \text{I}^- \longrightarrow [\text{BiI}_4]^- (\text{orange solution})$
 Pb^{2+} gives yellow precipitate of PbI_2 . Cu^{2+} gives white precipitate of Cu_2I_2 with evolution of iodine.
3. $[\text{Zn}(\text{OH})_4]^{2-} \xrightarrow{\text{H}_2\text{O-boiled}} \text{Zn}(\text{OH})_2 (\text{white}) + 2\text{OH}^-$
 $\text{Zn}(\text{OH})_2$ precipitate is readily soluble in excess of ammonia and in solutions of ammonium salts due to the formation of tetraamminezinc(II).
 $\text{Zn}(\text{OH})_2 + 4\text{NH}_3 \longrightarrow [\text{Zn}(\text{NH}_3)_4]^{2+} (\text{soluble complex}) + 2\text{OH}^-$
 or
 $\text{Zn}(\text{OH})_2 + 4\text{NH}_4\text{OH} \xrightarrow{\text{NH}_4\text{Cl}} [\text{Zn}(\text{NH}_3)_4]^{2+} + 4\text{H}_2\text{O} + 2\text{OH}^-$
 But $\text{Al}(\text{OH})_3$, $\text{Mg}(\text{OH})_2$ and $\text{Ca}(\text{OH})_2$ don't dissolve in excess of NH_3 solution.
4. $\text{Cu}^{2+} + 6\text{CN}^- \longrightarrow [\text{Cu}(\text{CN})_4]^{3-} + (\text{CN})_2$
 $\text{Cu}^{2+} + 2\text{CN}^- \longrightarrow \text{Cu}(\text{CN})_2 \downarrow (\text{yellow}); 2\text{Cu}(\text{CN})_2 \downarrow \longrightarrow 2\text{CuCN} \downarrow (\text{white}) + (\text{CN})_2 \downarrow$
 $\text{CuCN} \downarrow + \text{CN}^- \longrightarrow [\text{Cu}(\text{CN})_4]^{3-} (\text{colourless soluble complex}).$

5. MgSO_4 on reaction with Na_2HPO_4 in presence of NH_4OH gives white precipitate of $\text{Mg}(\text{NH}_4)\text{PO}_4$.

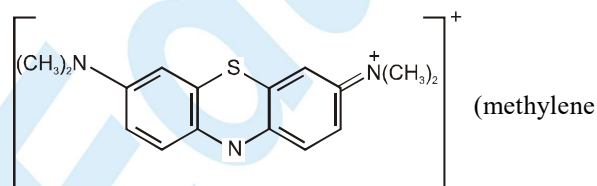
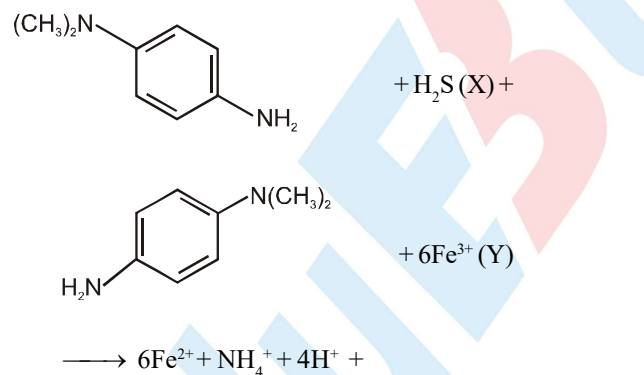
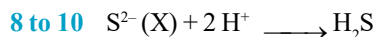


6. $\text{Hg}^{2+} + 2\text{I}^- \longrightarrow \text{HgI}_2 \downarrow$ (scarlet / red)
 $\text{HgI}_2 + 2\text{I}^-$ (excess) $\longrightarrow [\text{HgI}_4]^{2-}$ (soluble complex)
 $\text{Hg}^{2+} + \text{Co}(\text{SCN})_2 \longrightarrow \text{Co}[\text{Hg}(\text{SCN})_4] \downarrow$ (deep blue)

7. All ammonium salts on reaction with alkali produce ammonia. The nitrate and nitrite also on reduction with nascent hydrogen (produced by the reaction of zinc and sodium hydroxide) produce the ammonia gas according to the following reactions.



So options (A) and (B) are correct.



blue).

The compound X is Na_2S and Y is FeCl_3 .

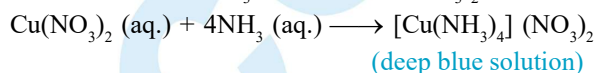
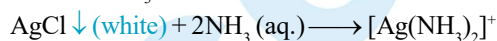
11, 12 & 13

$\text{Cu}(\text{M}) + \text{AgNO}_3(\text{N})$ (aqueous colourless solution) $\longrightarrow$
 Resultant solution contains $\text{Cu}(\text{NO}_3)_2$ (blue solution) and
 AgNO_3 (colourless solution)

(N)

Note : Here it is considered that complete AgNO_3 is not utilized in the reaction.

$\text{AgNO}_3(\text{aq}) + \text{NaCl}(\text{aq}) \longrightarrow \text{AgCl} \downarrow$ (white) (O) + NaNO_3
 Solution containing white ppt. of AgCl also contains
 $\text{Cu}(\text{NO}_3)_2$ which developed deep blue colouration with
 aqueous NH_3 solution

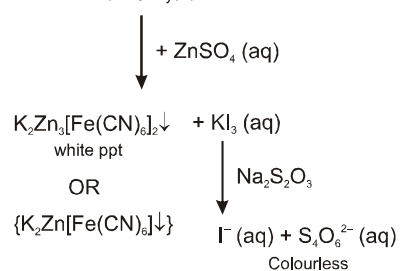
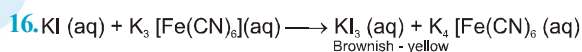


So, Metal rod M is Cu.

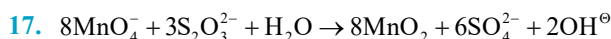
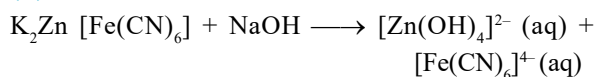
The compound N is AgNO_3 and the final solution contains $[\text{Ag}(\text{NH}_3)_2]^+$ and $[\text{Cu}(\text{NH}_3)_4]^{2+}$

14. In presence of acidic medium, ionisation of H_2S is suppressed so less number of S^{2-} ions are produced. So only those sulphides are precipitated which have low solubility product (K_{sp}) value, For example CuS and HgS .

15. Precipitation of insoluble cuprous salts pushes the equilibrium backwards (i.e. towards the left)



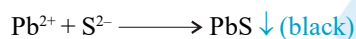
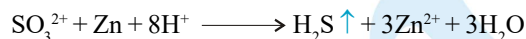
(D) with NaOH



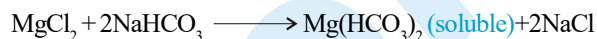
22. (A) Cu^{2+} show characteristic green colour in the flame test.
- (B) Only Cu^{2+} can give precipitate in acidic medium on passing H_2S
- (C) Both Cu^{2+} and Mn^{2+} show the formation of precipitate by passing H_2S in faintly basic medium.
- (D) $E^\circ_{\text{Cu}^{2+}/\text{Cu}} > E^\circ_{\text{Mn}^{2+}/\text{Mn}}$, as per electrochemical series.

MOCK TEST

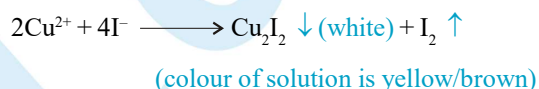
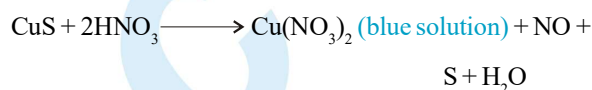
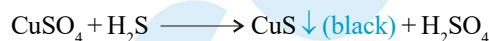
1. (A)



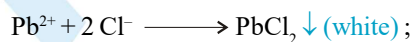
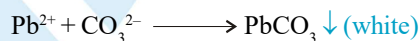
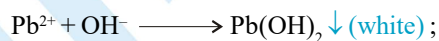
2. (A)



3. (B)

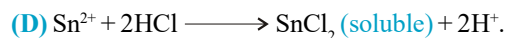


4. (D)



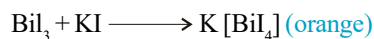
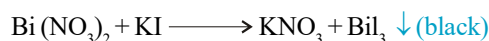
5. (D)

(A) (B) (C) can be precipitated by HCl as well as H_2S as their insoluble chlorides and sulphides respectively.

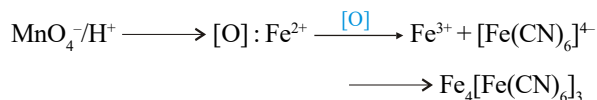


It can be precipitated only by H_2S as its insoluble sulphide (brown).

6. (B)



7. (A)



8. (B)

On adding HCl the equilibrium will shift in backward direction.

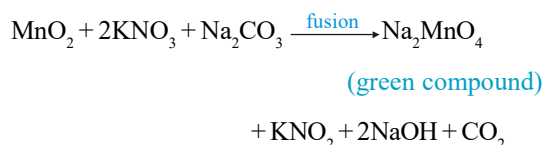
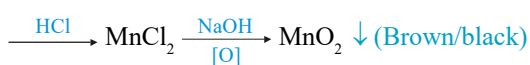
9. (A)

ZnS & MnS are soluble in conc. HCl. Ni^{2+} & Co^{2+} are insoluble in conc. HCl but dissolves when KClO_3 is added, forming corresponding chlorides.

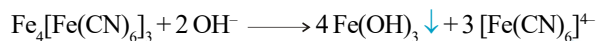


Ni^{2+} forms white precipitate which is soluble in excess KCN.

10. (D)



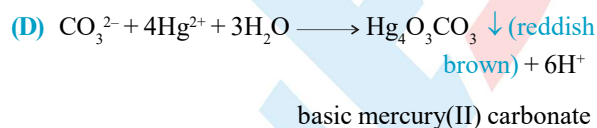
11. (B)



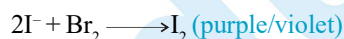
12. (B, C, D)



(C) True statement

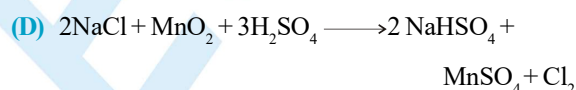


13. (C,D)

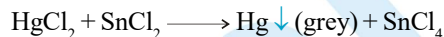
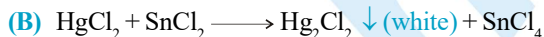
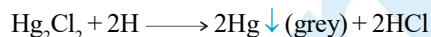
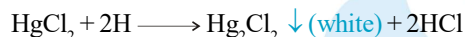
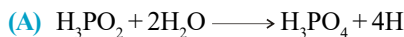


14. (C, D)

(C) Heavy metal chlorides like AgCl does not give chromyl chloride.



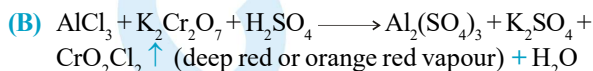
15. (A,B)



(C) Red precipitate is formed which dissolves in excess of KI.

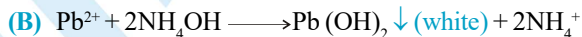
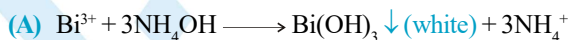
(D) White precipitate of $\text{Hg}(\text{NH}_3)\text{Cl}$ is formed.

16. (A, B, D)

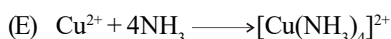
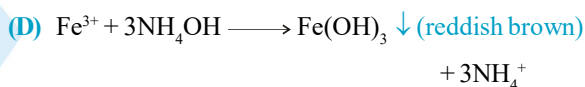


(D) Gives blue bead of $\text{Al}_2\text{O}_3 \cdot \text{CoO}$

17. (A, B, D)

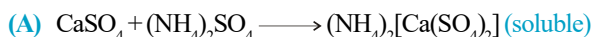


(C) Due to common ion effect of NH_4^+ , the OH^- ion concentration is not excess enough to precipitate Mg^{2+} as $\text{Mg}(\text{OH})_2$ because of its high solubility product.



In presence of ammonium salts, precipitation does not occur at all, but blue colour solution is formed right away.

18. (A,B,C,D)



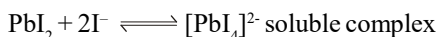
(B) BaCrO_4 is insoluble in dilute acetic acid & thus gets precipitated with K_2CrO_4 (low K_{sp} in K_2CrO_4 solution in acetic acid).

(C) $\text{Cr}(\text{OH})_3$ is soluble in NaOH & Br_2 water forming sodium chromate while $\text{Fe}(\text{OH})_3$ is insoluble.

(D) $[\text{Cu}(\text{CN})_4]^{3-}$ is more stable than $[\text{Cd}(\text{CN})_4]^{2-}$ gives yellow precipitate with H_2S .

19. (A)

20. (B)



Reaction is reversible and shifts in backward direction on dilution with water.

21. (D)

Dilute solution contains more of water thus precipitate formed gets dissolved.

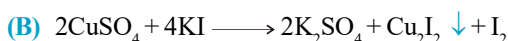
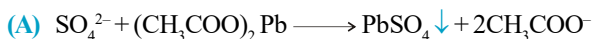
22. (A)

23. (D)

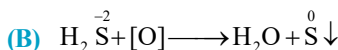
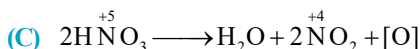
All are black and gives gas with dil. H_2SO_4



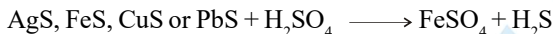
24. (D)



25. (D)

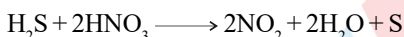


26. (A)



(A)

(B)

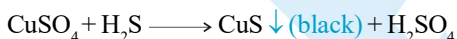


(C)

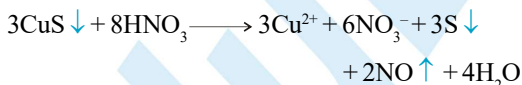
(D)



(E)



(F)



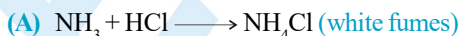
(G)

complex



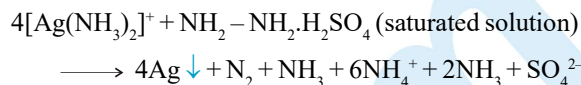
(chocolate brown)

27. (D)



(C) Nitrogen gas is obtained

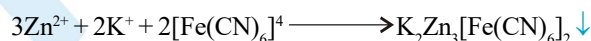
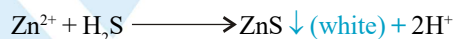
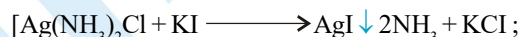
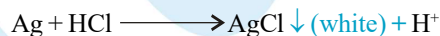
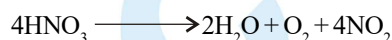
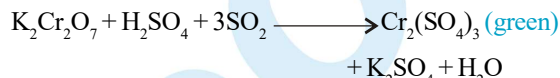
28. (C)



This is called silver mirror test. Finely divided silver metal adhere to the glass walls forming an attractive mirror.

29. (D)

Silver is precipitated.



31. $[\text{A}] = \text{K}^+$ and I^- , $[\text{B}] = \text{Hg}^{2+}$ and Cl^-

As black sulphide is only soluble in aquaregia and further it gives greyish black ppt with SnCl_2 the one of the cation may be Hg^{2+} . As salt mixture gives lilac colour in flame, the other cation is K^+ .

