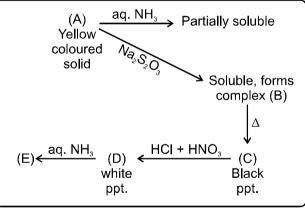
Exercise-1

> Marked questions are recommended for Revision.

PART - I : SUBJECTIVE QUESTIONS

Section (A) : Ist Group

- A-1. Name of one chloride which is soluble in hot water as well as in excess of HCI.
- A-2. Why do lead salts turn black on keeping for a long time in the laboratory ?
- A-3. A metal salt forms a yellow precipitate (P) with potassium iodide solution and black precipitate (Q) with acidified H₂S. Yellow precipitate (P) dissolve in excess of concentrated solution of KI and ppt.reappears on dilution. Then metal salt could be and also write the chemical equation.
- A-4. When calomel reacts with ammonia solution, a black precipitate is formed. Write the chemical equation and also name the reaction nature.
- **A-5.** What products are formed when precipitate formed by the reaction of Hg₂²⁺ ions and excess of sodium hydroxide solution is boiled ?
- A-6. A metal salt forms a green precipitate with KI solutiion and white ppt with KCI. This green precipitate under goes disproportionation reaction in excess of reagent which results in a soluble compound and black precipitate then write the chemical equations and identify the cation in the original sample.
- A-7. Why bright yellow precipitate of AgI is soluble in KCN and Na₂S₂O₃ solution.
- A-8. Solid (A) is used in photographic film, Identify (A) to (E) ?



Section (B) : II A Group

B-4.

- B-1. Why Na₂S cannot be used in place of H₂S (in presence of HCI) as a reagent for IInd group cations ?
- B-2. Does sodium hydroxide solution can be used to differentiate Hg(I) from Hg(II) ?
- B-3. What happens when white precipitate of Bi(OH)₃ is boiled ?

Identify A to F. (F) (A) K₄[Fe(CN)₆] BaCl₂ (B) (C) Chocolate ← White ppt. Blue solution brown ΚI (D) (E) Hypo White ppt. Brown ppt. Solution

Section (C) : IIB Group

- C-1. An original solution is prepared in conc. HCl when diluted a white ppt. is formed. What does it indicate ?
- C-2.∞ Why HNO₃ can not be used in place of HCl as a reagent (H₂S in presence of HCl) for IInd group cations?
- **C-3.** A chloride of an element (X) forms yellow precipitate with H₂S in acidic medium. This yellow precipitate is soluble in Conc. HNO₃ due to formation of compound (Y). Compound (Y) gives ammonium molybdate test then find the metal salt (X).
- C-4.> Identify (A) based on the following facts :
 - (i) (A) reduces $HgCl_2$ solution to white ppt. changing to grey.
 - (ii) (A) turns FeCl₃ yellow coloured solution to green.
 - (iii) (A) gives white ppt. with NaOH soluble in excess of NaOH.
 - (iv) (A) gives yellow dirty ppt. on passing H₂S gas, soluble in yellow ammonium sulphide (YAS).
 - (v) (A) gives chromyl chloride test.

Section (D) : IIIrd Group

- D-1. Why is the original solution boiled with conc. HNO₃ in group III ?
- D-2. Do Fe(III) salts and Fe(II) salts both give red colouration with dimethylglyoxime in ammonical solution. If not then which iron salt gives red colouration with dimethylglyoxime ?
- D-3. Which colour precipitate is formed by Fe(II) salt with potassium ferrocyanide, (i) in complete absence of air and (ii) under ordinary atmospheric conditions ?

Section (E) : IVth Group

- **E-1.** Partial precipitation of Mn²⁺ as Mn(OH)₂ occurs with ammonia solution but the precipitate is soluble in ammonium salts. Explain ?
- E-2. Why Zn(II) salt is not precipitated as Zn(OH)₂ by ammonia solution in the presence of excess of ammonium chloride ?
- E-3. In the absence of copper ion, zinc ion form a white precipitate with the ammonium tetrathiocyanatomercurate(II) How the presence of Cu²⁺ ion affect this test ?
- E-4. A Identify (A) to (D).

(A) $\xrightarrow{\Delta}$ (B) + (C) (On cooling (C) turns white) Colourless water soluble compound $| NH_4OH |$ and H_2S

Section (F) : Vth, VIth and Zero Group

F-1. What is the formula of iodide of Millon's base ?

- F-2. What happens when ammonia gas is passed into a solution of sodium cobaltinitrite ?
- **F-3.** What will happen if the precipitation of Vth group cation by ammonium carbonate is replaced by Na₂CO₃?

(D) (White ppt.)

- **F-4.** What happens when ammonium sulphate solution is added to a solution containing both Sr²⁺ and Ca²⁺ ions?
- **F-5.** Which colour precipitate is obtained when a solution of Ca²⁺ ions reacts with potassium ferrocyanide.

PART - II : ONLY ONE OPTION CORRECT TYPE

Section (A) : Ist Group

	A metal nitrate reacts with KI solution concentrated solution (6 M) of KI dissol (A) Hg2 ²⁺ (B) Ag ⁺		
A-2.	Three separate samples of a solution o with excess ammonia solution, one forr a black precipitate with H_2S . The salt co (A) AgNO ₃ (B) Pb(NO ₃) ₂	ned a white precipitate with dil	
A-3.১	Consider the following observation :		
	$\begin{array}{ll} M^{n+} + HCI \mbox{ (dilute)} & \longrightarrow \mbox{ white precipitat} \\ The metal ion \ M^{n+} \mbox{ will be :} \\ (A) \ Hg^{2+} & (B) \ Ag^{+} \end{array}$	$e \xrightarrow{a \to a} water soluble \xrightarrow{a \to a}$ (C) Pb ²⁺	(D) Sn ²⁺
A-4.	When calomel reacts with NH ₄ OH solut (A) Hg ₂ Cl ₂ (B) Hg(NH ₂)Cl	•	(D) HgCl ₂ .NH ₃
A-5.১	Consider the following equilibrium : AgCl \downarrow + 2NH ₃ \implies [Ag(NH ₃) ₂] ⁺ + Cl ⁻ White ppt of AgCl appears on adding (A) NH ₃ (B) aq. NaBr	(C) aq. HNO₃	(D) aq. NH₄I
A-6.	AgCI with NH₃ forms a complex : (A) [Ag(NH₃)₂]Cl (B) AgNO₃	(C) [Ag(NH ₂) ₂]Cl	(D) Ag mirror
	on (B) : IIA Group Sometimes yellow turbidity appears we absence of II group radicals. This is been (A) sulphur is present in the mixture as (B) IV group radicals are precipitated as (C) of the oxidation of H ₂ S gas by some (D) III group radicals are precipitated as	cause : impurity. s sulphides. acid radicals.	in slightly acidic medium in the
B-2.	 H₂S in the presence of HCl precipitates (A) HCl activates H₂S (B) HCl increases concentration of Cl⁻ (C) HCl decreases concentration of S²⁻ (D) HCl lowers the solubility of H₂S in second secon		ause :
В-3.	When small amount of SnCl ₂ is added The silky white precipitate is due to the (A) Hg ₂ Cl ₂ (B) SnCl ₄		silky white precipitate is obtained. (D) Hg
B-4.	When excess of dilute NH ₄ OH is add colour is developed. This is due to the f (A) $[Cu(NH_3)_6]^{2+}$ (B) $Cu(OH)_2$		copper sulphate an intense blue (D) (NH ₄) ₂ SO ₄
B-5.æ	A black sulphide is formed by the action (A) cupric chloride (B) cadmium cl		(D) ferric chloride.
B-6.	Which one of the following salts will pro (A) Ag ₂ CO ₃ (B) Pb(CO ₃)	duce clear and transparent ori (C) Hg ₂ CO ₃	ginal solution in 2M HCl ? (D) CuCO₃
B-7.	When bismuth chloride is poured into a (A) BiO.OH (B) Bi ₂ O ₃	large volume of water the whit (C) BiOCI	e precipitate produced is of : (D) Bi(OH)₃
B-8.24	In which of the following pairs the p precipitates are soluble in excess KI so (A) Hgl ₂ , Hg ₂ l ₂ (B) Hgl ₂ , Bil ₃		(D) Cdl ₂ , Pbl ₂

B-9.æ		nal solution (i.e. O.S) c lium hydroxide. The me (B) iron		solution gives a yellow precipitat (D) lead
Sectio C-1.	on (C) : IIB Group Which of the following (A) HgS	metal sulphide is solub (B) PbS	le in YAS (yellow ammc (C) Bi ₂ S ₃	onium sulphide) : (D) Sb2S3
C-2.			sium ammonium arsen s formed. The colour of (C) White	ate is treated with acidified silve precipitate is : (D) Brownish black
	on (D) : III rd Group When NH ₄ Cl is added (A) the dissociation of (B) the concentration of (C) the concentrations (D) the concentration of	NH₄OH increases. f OH⁻ increases. of both OH⁻ an NH₄⁺ ir		
D-2.æ	To avoid the precipitat the third group solution (A) Concentrated HNC (C) Concentrated H ₂ S0	n should be : ₀ is added	n ²⁺ , Mn ²⁺ and Ni ²⁺ alon (B) Treated with exc (D) Treated with exc	
D-3.	NH₄OH is brought ne	ar to it. An acidic sol olves in NaOH solutio	ution of (X) on addition	increases when and rod dipped i on of NH ₄ Cl and NH ₄ OH gives oes not give precipitate with H ₂ (D) ZnCl ₂
D-4.১	An original solution of	of an inorganic salt in	dilute HCI gives a b	prown colouration with potassiur acetate solution. The cation of th (D) none
D-5.	Fe(OH) ₃ and Cr(OH) ₃ (A) Aq. NH ₃	precipitates can be con (B) HCl	npletely separated by : (C) NaOH/H ₂ O ₂	(D) H ₂ SO ₄
Sectio E-1.	(i) It gives white precip			own on exposure to air. ammonium salts. (D) Ni ²⁺
E-2.	A metal salt form prec acetic acid then metal (A) ZnS		ence of (NH₄OH + NH₄O (C) MnS	Cl) and this precipitate is soluble i (D) NiS
E-3.24		ly the concentration of f H₃(aq) [Zn(NH		on of the complex ion $[Zn(NH_3)_4]^{2+}$
	add to the solution son $(A) H_2O$	ne : (B) HCI (aq)	(C) NH ₃ (aq)	(D) NH₄Cl (aq)
-4. æ		IH₄OH. On passing H₂		precipitate with NH₄OH. This wa ution a white precipitate is formed (D) Zn
	. ,			
<u>5.2</u>	A metal salt solution v metal is : (A) Ni	vhen treated with dime (B) Zn	thyl glyoxime and NH₄C (C) Co	OH gives a rose red complex. Th (D) Mn.

E-6.	The ion that can not be (A) Pb ²⁺	e precipited by H ₂ S in (B) Bi ³⁺	presence of dil. HCl. (C) Cu ²⁺	(D) Ni ²⁺
	on (F) : V th , VI th and		cipitate with :	
F-1.234	Aqueous Solution of B (A) K ₂ CrO ₄	(B) AgNO ₃	(C) (CH ₃ COO) ₂ Pb	(D) (A) and (B) both
F-2.	but that one which doe	es not produce precipi	tate is :	oduce a precipitate in every case
	(A) BaCl ₂ (aq)	(B) CaBr ₂ (aq)	(C) Na ₂ SO ₄ (aq)	(D) Pb(NO ₃) ₂ (aq)
F-3.	may be	. .		on as well as with dilute H ₂ SO ₄ . It
	(A) Pb(NO ₃) ₂	(B) Ba(NO ₃) ₂	(C) BaCl ₂	(D) CuCl ₂
F-4.æ	Mg is not precipitated (A) MgCO ₃ is soluble i (C) MgCO ₃ is soluble i	n water.	(B) K₅p of MgCO₃ is h (D) None.	igh.
F-5.			n potassium chromate in a chloride or iodide it is : (B) Basic lead carbor (D) Strontium nitrate	cetic acid, a white ppt with dilute nate
F-6.	obtained. The black	precipitate dissolves		his solution, a black precipitate is D ₃ . On adding a few drops of that of : (D) CdSO4
F-7.a	The yellow precipitate (A) Hgl₄ ^{2−} (C) NH2–Hg–O–Hg–I	formed by passing ar	nmonia into Nessler's reag (B) NH₂O–Hg–HgI (D) NH₃–Hg–I	ent in due to the formation of

1. Match the precipitates listed in column-I with their suitable solvents listed in column-II.

	Column I		Column II
	(precipitate)		(solvent)
(A)	AgCl↓ (white)	(p)	Concentrated HCI.
(B)	CuS↓ (black)	(q)	Dilute ammonia solution (excess).
(C)	Zn(OH)₂↓ (white)	(r)	Potassium cyanide solution.
(D)	BaCO₃↓ (white)	(s)	Hot 50% nitric acid.
		(t)	Sodium hydroxide solution.

2. Match the basic radicals listed in column-I with the properties listed in column-II.

	Column-I		Column-II
	(Basic radical)		(Properties)
(A)	Mn ²⁺	(p)	Forms coloured metaborate in oxidising flame in borax bead test.
(B)	Cr ³⁺	(q)	Forms white precipitate with sodium hydroxide but on exposure to air turns rapidly brown.
(C)	Al ³⁺	(r)	With both potassium cyanide (not in excess) and ammonia solution separately forms reddish brown precipitate.
(D)	Fe ³⁺	(s)	With excess of sodium hydroxide forms soluble complex but on adding acid to soluble complex, a precipitate is obtained which redissolves on adding excess of acid.

Qu	alitative Analysis (Cation	ns)		
	Exercise- 2	2		
🔈 Mar	ked questions are recom			
	PART - I	: ONLY ONE O	PTION CORREC	ТТҮРЕ
1.১	Which of the following con (A) AgNO ₃ (mpound does not gives B) Pb(NO ₃) ₂	s ppt with dil. HCl ? (C) Hg ₂ (NO ₃) ₂	(D) Hg(NO ₃) ₂
2.	KI gives precipitate with a (A) Ag ⁺ , Hg ₂ ²⁺ , Pb ²⁺ ((C) Na+, Ca ²⁺ , Mg ²⁺	(D) Ag ⁺ , Ca ²⁺ , Sr ²⁺
3.	Three test tubes A, B, C added in excess). Followi A : Black ppt, B : Brown p A, B and C contain respe (A) Pb ²⁺ , Hg ₂ ²⁺ , Ag ⁺ (C) Ag ⁺ , Pb ²⁺ , Hg ₂ ²⁺	ing changes occur. opt, which disolve in NH		ach, aqueous solution NaOH is colves in excess of NaOH
4.æ	Salt mixture $\xrightarrow{dil. HCI}$ wh	hite ppt. <u>Heated</u> and filtered under hot condition	→ Filtrate <u>Cooled</u>	like crystal
	salt mixture contains cation (A) Pb^{2+} and Hg^{2+} (ons of : B) Pb ²⁺ and Hg ₂ ²⁺	(C) Pb ²⁺ and Ag ⁺	
5.	A compound (X) reacts in Bright ye	(X)(aq) KI(aq) NaOH(aq) NaOH(aq)	NH₃(aq) Colourless	
	The compound (X) is likel (A) Pb(NO ₃) ₂ (ly to be B) CaCrO₄	solution (C) Hg(NO ₃) ₂	(D) AgNO ₃
6.2	colour is due to the forma		f HgCl ₂ , a white ppt tur (C) Sn	ning grey is obtained. The grey (D) Hg
7.	heating. When hydrogen subtance is a:	sulphide is passed thro	ough the hot acidic solut	vith dil HCl, which dissolves on ion, a black ppt is obtained.The
8.海	Three seperate sample c with excess of ammonia a black precipitate with H	solution, one formed a	-	 (D) Pb²⁺ salt one formed a white precipitate I. NaCl solution and one formed (D) MnSO₄
9.	addition of aqueous Nac medium. An aqueous so solution. The metal in the	DH. The metal salt sol plution of the metal s	ution also gives a black	tanding it gives a brown ppt on k ppt on bubbling H ₂ S in basic k colour of the permanganate (D) iron
10.১	K₄[Fe(CN)₀] can be used (A) only Fe²+, Fe³+ (to precipitate one or m B) only Fe ³⁺ ,Zn ²⁺ ,Cu ²⁺		²⁺ , Cu ²⁺ , Ca ²⁺ : (D) all of these.

- 11.2 Ferric alum gives deep red colour with NH₄SCN due to the formation of : (A) AI(SCN)₃ (B) [Fe(SCN)₃]-(C) Fe(SCN)₃ (D) None of these. 12. Nessler's reagent is used to detect. (A) CrO₄²⁻ (B) PO₄^{3−} (C) MnO₄-(D) NH₄+ 13. On the addition of a solution containing CrO₄²⁻ & CH₃COOH acid to the solution of Ba²⁺, Sr²⁺ and Ca²⁺ ions, the ppt obtained first will be of : (A) CaCrO₄ (B) SrCrO₄ (C) BaCrO₄ (D) a mixture of all the three 14. A salt on treatment with dil. HCl gives a pungent smelling gas and a yellow precipitate. The salt gives green flame when tested. The solution gives a vellow precipitate with potassium chromate. The salt is : (A) NiSO₄ (B) BaS_2O_3 (C) PbS_2O_3 (D) CuSO₄ A chloride dissolves appreciably in cold water. When placed on a platinum wire in Bunsen flame, no 15.🕰 distinctive colour is noticed. The cation of chloride is : (A) Ma²⁺ (B) Ba²⁺ (C) Pb2+ (D) Ca2+ In fifth group, (NH₄)₂CO₃ is added to precipitate out the carbonates. We do not add Na₂CO₃ because : 16. (A) CaCO₃ is soluble in Na₂CO₃ (B) Na₂CO₃ increases the solubility of fifth group carbonates (C) MqCO₃ will be precipitated out in fifth group (D) none
- A metal salt solution forms a yellow precipitate with potassium chromate in acetic acid, a white precipitate with dilute sulphuric acid, but gives no precipitate with sodium chloride or iodide, it is :

 (A) lead salt
 (B) silver salt
 (C) barium salt
 (D) strontium salt

PART - II : SINGLE AND DOUBLE VALUE INTEGER TYPE

- **1.** An aqueous solution contains Hg²⁺, Hg₂²⁺, Pb²⁺, Ag⁺, Bi³⁺ and Cd²⁺. Out of these, how many ions will produce white precipitate with dilute HCl ?
- **2.** A solution of Hg²⁺ ion on treatment with a solution of cobalt(II) thiocyanate gives rise to a deep blue crystalline precipitate. Then the coordination number of mercury in the deep blue coloured compound is:

3. FeSO₄ + KCN
$$\xrightarrow{\Delta}$$
 (X) $\xrightarrow{\text{Conc.H}_2SO_4}$ (Y) \uparrow
(excess) (Z) $\xrightarrow{\text{dil.H}_2SO_4}$ (Z) \uparrow
(as)

The sum of number of $p\pi$ - $p\pi$ bonds present in gas (Y) & (Z) ?

- 4. An alcoholic solution of dimethylglyoxime is added to an aqueous solution of nickel(II) chloride. Slow addition of ammonium hydroxide led to the precipitation of a bright-red coloured metal complex. Find out the number of hydrogen bonds present in the structure of the complex.
- 5. In how many of the following reactions, one of the product is obtained as a yellow precipitate :
 - (a) $Ba^{2+} + CrO_4^{2-} \longrightarrow product$
 - (b) $NH_{4^+} + [PtCl_6]^{2-} \longrightarrow product$
 - (c) $NH_{4^+} + [Co(NO_2)_6]^{3-} \longrightarrow product$
 - (d) $Ca^{2+} + [Fe(CN)_6]^{4-} + K^+ \longrightarrow product$
 - (e) $Sr^{2+} + CO_3^{2-} \longrightarrow product$

Q	ualitative Analysis (Cations)
6.	Total number of metal nitrates given below, which give the following reaction, (i) AgNO ₃ (ii) Pb(NO ₃) ₂ (iii) Cu(NO ₃) ₂ (iv) Cd(NO ₃) ₂ (v) Zn(NO ₃) ₂ (v) Ni(NO ₃) ₂ (vi) Al(NO ₃) ₃
7.	Aqueous CuSO ₄ decolorizes on addition of excess KCN due to formation of complex (A). In complex "A". If (i) Number of d orbitals participating in hybridisation is / are 'a' (ii) Coordination number of Cu is "c". then find 8a + 5c.
8.24	$\begin{array}{l} Co^{2+} + CN^{-} \longrightarrow \ "A" \ (reddish \ brown) \ ppt. \\ \ "A" + CN^{-} \ (excess) \longrightarrow \ "B" \ (brown \ solution) \\ \ "B" + O_{2} + 2H_{2}O \longrightarrow \ "C" \ (yellow \ solution) \\ \ then \ find \ (a + b) \ for \ complex \ "C" \ if \\ \ b = charge \ on \ complex \ (consider \ the \ magnitude) \end{array}$
9.2	How many of the following reactions give yellow ppt.(i)NaBr + AgNO3 \longrightarrow (ii)NaI + AgNO3 \longrightarrow (iii)NaBr + Pb(NO3)2 \longrightarrow (iv)NaI + Pb(NO3)2 \longrightarrow (v)Na2S + Cd(CH3COO)2 \longrightarrow (vi)K2CrO4 + Pb(CH3COO)2 \longrightarrow (vii)K2CrO4 + AgNO3 \longrightarrow (viii)NaBr + Chlorine water (excess) \longrightarrow PART - III : ONE OR MORE THAN ONE OPTIONS CORRECT TYPE
1.	Which of the following is/are correctly matched ?(A) $Bil_3 \downarrow \longrightarrow Black precipitate(B) CuI \downarrow \longrightarrow White precipitate(C) Pbl_2 \downarrow \longrightarrow Yellow precipitate(D) Hgl_2 \downarrow \longrightarrow Red precipitate$
2.2	Which of the following are completely soluble in concentrated ammonia solution?(A) AgCl(B) AgBr(C) Ag2CrO4(D) AgI
3.24	 Hg₂I₂↓ (green) → boiled with H₂O products Which of the following statement is / are correct with respect to the products ? (A) Black precipitate of mercury(I) oxide is formed. (B) Violet colour gas is evolved. (C) Red precipitate of HgI₂ is formed. (D) Mercury is obtained
4.	Which of the following cations form coloured (not white) precipitates with aqueous solution of KI and this precipitate does not dissolve in excess of reagent ? (A) Hg^{2+} (B) Hg^{2+}_{2} (C) Ag^{+} (D) Cu^{2+}
5.	 Which of the following statements is/are true ? (A) Ag⁺ ions do not give white precipitate with excess of concentrated HCI. (B) Cu²⁺ ions produce a white precipitate when KCN solution is added in a small quantity, and allowed to stand. (C) Hg²⁺ ions give deep blue precipitate with cobalt acetate and ammonium thiocyanate. (D) Black precipitate of Bil₃ turns orange when heated with water.

Qu	alitative Analysis (Cat	ions)		
6.24	KI solution is the reage (A) Hg²+	ent for the analysis of : (B) Pb ²⁺	(C) Ag⁺	(D) Cu ²⁺
7.	Which of the following (A) HgS	metal sulphide are in bro (B) PbS	wn/balck in colour ? (C) Bi ₂ S ₃	(D) Sb ₂ S ₃
8.	Which of the following (A) Cu ²⁺	cations form(s) black pre (B) Sb ³⁺	cipitate(s) with H ₂ S (g) ? (C) Pb ²⁺	(D) Bi ³⁺
9.	Which of the following (A) Zn ²⁺ , Mn ²⁺	mixture of cations can be (B) Zn ²⁺ , Cd ²⁺	e separated by adding ex (C) Zn ²⁺ , Pb ²⁺	ccess NH ₃ solution ? (D) Zn ²⁺ , Cu ²⁺
10.24	(Unbalanced equation) Which is / are correct for (A) X is a yellow crysta (B) X is a green coloure (C) IUPAC name of X i		ble in water. Rinman's green. -N– cobaltate (II)	
11.24	(A) It gives a brown pred(B) It gives a white pred(C) It in excess gives a	is/are correct for potassiu ecipitate with Cu ²⁺ ions. cipitate of mixed salt with bluish white/white precip red colouration with Fe ³⁺	$n Ca^{2+}$ ions. Ditate with Zn^{2+} .	

PART - IV : COMPREHENSION

Read the following passage carefully and answer the questions.

Comprehension #1

Aqueous solution of 'A' $\xrightarrow{H_2S (g)}$ Black precipitate 'B', soluble in 50% HNO₃ forming 'C'.

NH₃ solution

Alkaline Na₂SnO₂

Black precipitate 'D'

(D) CuO.OH

White precipitate dissolves in

hydrochloric acid but on dilution with water again white turbidity appears 'E'.

Moreover, the salt 'A' on heating with solid $K_2Cr_2O_7$ and concentrated H_2SO_4 produces deep red vapours which dissolve in sodium hydroxide solution forming a yellow solution. This yellow solution gives yellow precipitate with Ba(NO₃)₂ solution.

On the basis of the aforesaid characteristic informations answer the following questions :

- Acidified solution of 'A', on treatment with KI gives black precipitate 'F' which dissolves in excess of reagent forming the coloured compound 'G'. The chemical composition of 'F' and 'G' are respectively :

 (A) Hgl₂ and [Hgl₄]²⁻
 (B) Pbl₂ and [Pbl₄]²⁻
 (C) Bil₃ and [Bil₄]⁻
 (D) Cul and Cul₂.
- The black precipitate 'F' on heating with water produces :
 (A) Hg(OH)₂
 (B) BiOI
 (C) BiO.OH
- **3.** Select the correct statement.

(A) Aqueous solution of 'A' reacts with $AgNO_3$ solution to give white precipitate which turns into yellow on treatment with sodium arsenite.

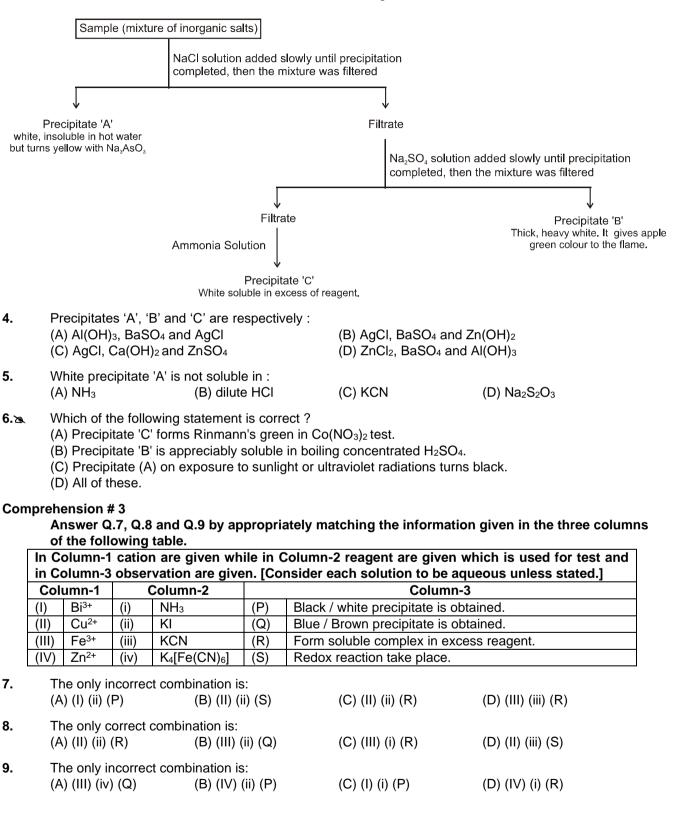
(B) Aqueous solution of 'A' produces white precipitate with sodium hydroxide which turns into yellowishwhite on boiling.

(C) White turbidity 'E' is soluble in dilute mineral acids.

(D) All of these.

Comprehension # 2

A student was given a sample of colourless solution containing three cations and was asked to identify the cations. Student carried out a series of reactions as given below.



Oualitative Analysis (Cations) Exercise-3 * Marked Questions may have more than one correct option. PART - I : JEE (ADVANCED) / IIT-JEE PROBLEMS (PREVIOUS YEARS) A metal nitrate reacts with KI to give a black precipitate which on addition of excess of KI is converted 1. into orange colour solution. The cation of the metal nitrate is : [JEE - 2005, 3/84] (A) Hg²⁺ (B) Bi3+ (C) Pb2+ (D) Cu+ 2. In the given reaction sequence, Identify (A) and (B). (Excess) Blood red F^{-} (excess) \rightarrow colourless(B) Fe³⁺ -Write the IUPAC name of (A) and (B). (a) (b) Find out the spin only magnetic moment of B. [JEE 2005, 4/144] 3. A white precipitate is obtained when a solution is diluted with H₂O and boiled. On addition of excess NH₄Cl/NH₄OH, the volume of precipitate decreases leaving behind a white gelatinous precipitate. Identify the precipitate which dissolves in ammonia solution or NH₄Cl. [JEE 2006, 3/184] (A) $AI(OH)_3$ (D) Ca(OH)2 (B) $Zn(OH)_2$ (C) $Mg(OH)_2$ In blue solution of copper sulphate excess of KCN is added then solution becomes colourless due to 4. the formation of : [JEE 2006, 3/184] (B) Cu²⁺ get reduced to form [Cu(CN)₄]³⁻ (A) [Cu(CN)₄]²⁻ (C) $Cu(CN)_2$ (D) CuCN MgSO₄ + NH₄OH + Na₂HPO₄ \longrightarrow white crystalline precipitate. The formula of crystalline precipitate 5. [JEE 2006, 3/184] is: (A) MgCl₂. MgSO₄ (B) MgSO₄ (C) Mg(NH₄)PO₄ (D) Mg(PO₄)₂ A solution of a metal ion when treated with KI gives a red precipitate which dissolves in excess KI to 6. give a colourless solution. Moreover, the solution of metal ion on treatment with a solution of cobalt(II) thiocyanate gives rise to a deep blue crystalline precipitate. The metal ion is : [JEE - 2007, 3/162] (D) Co2+ (A) Pb2+ (B) Hg²⁺ (C) Cu2+ 7.* A solution of colourless salt H on boiling with excess NaOH produces a non-flammable gas. The gas evolution ceases after some time. On addition of Zn dust to the same solution, the gas evolution restarts. The colourless salt(s) **H** is (are) : [JEE 2008, 4/163] (A) NH₄NO₃ (B) NH₄NO₂ (C) NH₄Cl (D) (NH₄)₂SO₄ Paragraph for Question Nos. 8 to 10 p-Amino-N, N-dimethylaniline is added to a strongly acidic solution of X. The resulting solution is treated with a few drops of aqueous solution of Y to yield blue coloration due to the formation of methylene blue. Treatment of the aqueous solution of Y with the reagent potassium hexacyanoferrate(II) leads to the formation of an intense blue precipitate. The precipitate dissolves on excess addition of the reagent. Similarly, treatment of the solution of Y with the solution of potassium hexacyanoferrate(III) leads to a brown coloration due to the formation of Z. The compound X is : [JEE 2009, 4/160] 8. (A) NaNO₃ (B) NaCl (D) Na₂S (C) Na_2SO_4 The compound Y is : 9. [JEE 2009, 4/160] (A) MgCl₂ (B) FeCl₂ (C) FeCl₃ (D) ZnCl₂

10. The compound **Z** is : (A) Mg₂[Fe(CN)₆] (B) Fe[Fe(CN)₆] (C) Fe₄[Fe(CN)₆]₃ [JEE 2009, 4/160] (D) K₂Zn₃[Fe(CN)₆]₂

Paragraph for Question Nos. 11 to 13

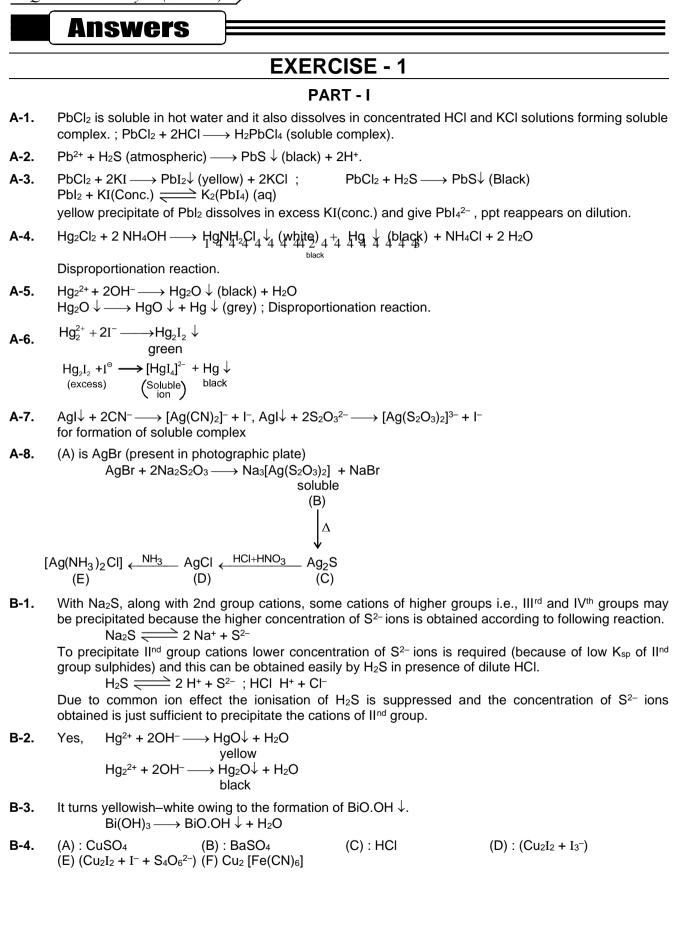
When a metal rod M is dipped into an aqueous colourless concentrated solution of compound N the solution turns light blue. Addition of aqueous NaCl to the blue solution gives a white precipitate O. Addition of aqueous NH_3 dissolves O and gives an intense blue solution.

11. The metal rod **M** is : [JEE 2011, 3/180] (A) Fe (B) Cu (D) Co (C) Ni [JEE 2011, 3/180] The compound **N** is : 12. (A) AgNO₃ (B) $Zn(NO_3)_2$ (C) $AI(NO_3)_3$ (D) Pb(NO₃)₂ [JEE 2011, 3/180] 13. The final solution contains (A) [Pb(NH₃)₄]²⁺ and [CoCl₄]²⁻ (B) [AI(NH₃)₄]³⁺ and [Cu(NH₃)₄]²⁺ (C) $[Ag(NH_3)_2]^+$ and $[Cu(NH_3)_4]^{2+}$ (D) [Ag(NH₃)₂]⁺ and [Ni(NH₃)₆]²⁺ Passing H₂S gas into a mixture of Mn²⁺, Ni²⁺, Cu²⁺ and Hg²⁺ ions in an acidified agueous solution 14. precipitates: [JEE 2011, 3/180] (A) CuS and HgS (B) MnS and CuS (C) MnS and NiS (D) NiS and HgS 15.* 2Cu^I \implies Cu⁰ + Cu^{II} The equilibrium, in aqueous medium at 25° C shifts towards the left in the presence of : [JEE 2011, 4/180] (C) SCN⁻ (D) CN-(A) NO₃-(B) CI-16.* For the given aqueous reaction which of the statement(s) is (are) true ? [JEE 2012, 4/136] dilute H₂SO₄ brownish-yellow solution excess KI + K₃[Fe(CN)₆] ZnSO₄ (white precipitate + brownish- yellow filtrate) Na₂S₂O₃ colourless solution (A) The first reaction is a redox reaction (B) White precipitate is $Zn_3[Fe(CN)_6]_2$ (C) Addition of filtrate to starch solution gives blue colour. (D) White precipitate is soluble in NaOH solution 17. Upon treatment with ammoniacal H₂S, the metal ion that precipitates as a sulphide is : [JEE(Advanced) 2013, 2/120] (A) Fe(III) (B) AI (III) (C) Mg(II) (D) Zn(II) Paragraph for Question 18 and 19 An aqueous solution of a mixture of two inorganic salts, when treated with dilute HCI, gave a precipitate (P) and a filtrate (Q). The precipitate P was found to dissolve in hot water. The filtrate (Q) remained unchanged, when treated with H_2S in a dilute mineral acid medium. However, it gave a precipitate (R) with H_2S in an ammoniacal medium. The precipitate **R** gave a coloured solution (S), when treated with H₂O₂ in an aqueous NaOH medium. [JEE(Advanced) 2013, 3/120] 18. The precipitate P contains (A) Pb²⁺ (B) $Hg_{2^{2+}}$ (C) Ag⁺ (D) Hg²⁺ 19. The coloured solution **S** contains [JEE(Advanced) 2013, 3/120] (A) Fe₂(SO₄)₃ (B) CuSO₄ (C) ZnSO₄ (D) Na₂CrO₄ Among PbS, CuS, HgS, MnS, Ag₂S, NiS, CoS, Bi₂S₃ and SnS₂, the total number of **BLACK** coloured 20. sulphides is : [JEE(Advanced) 2014, 3/120]

21.*	• • • •	where BOTH the ion	s are precipitated upon pa	
	HCI, is(are) (A) Ba ²⁺ , Zn ²⁺	(B) Bi ³⁺ , Fe ³⁺	(C) Cu ²⁺ , Pb ²⁺	[JEE(Advnaced) 2015, 4/16 (D) Hg ²⁺ , Bi ³⁺
22.	In the following reac	tion sequence in aqu	eous solution, the species	X, Y and Z, respectively, are [JEE(Advanced) 2016, 4/12
	$S_2O_3^{2-}$	$\xrightarrow{g^+}$ x $\xrightarrow{Ag^+}$	→ Y with time	Y
		Clear solution	white precipitate	black precipitate
	(A) [Ag(S₂O₃)₂]³-, Ag (C) [Ag(SO₃)₂]³-, Ag		(B) [Ag(S₂O₃)₃]⁵-, (D) [Ag(SO₃)₃]³-, A	
23.*	The correct option(s) to distinguish nitrate	e salts of Mn ²⁺ and Cu ²⁺ ta	ken separately is (are) IEE(Advanced) 2018, 4/120]
	(B) Only Cu ²⁺ show (C) Only Mn ²⁺ show	the formation of prec the formation of prec	blour in the flame test ipitate by passing H ₂ S in a sipitate by passing H ₂ S in f	acidic medium
	PART - II : JE	E (MAIN) / AIE	EE PROBLEMS (PREVIOUS YEARS)
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1.	(1) From a mixed pression (2) Ferric ions gave(3) On boiling a solution	lowing statement is c ecipitate of AgCl and a deep green precipi ition having K ⁺ , Ca ²⁺		[AIEEE 2003, 3/225 ssolves only AgCl. ferrocyanide solution. precipitate of K ₂ Ca(CO ₃) ₂ .
1. 2.	 (1) From a mixed present (2) Ferric ions gave (3) On boiling a solution (4) Manganese salts A red solid is insolution red solid in a test tube 	lowing statement is c ecipitate of AgCl and a deep green precipi ition having K ⁺ , Ca ²⁺ a s give a violet borax b ble in water. Howeve	correct ? AgI, ammonia solution dis tate on adding potassium and HCO3 ⁻ ions we get a p pead test in the reducing fi er it becomes soluble if som n of some violet coloured f	[AIEEE 2003, 3/225 ssolves only AgCl. ferrocyanide solution. precipitate of K ₂ Ca(CO ₃) ₂ .
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2.	 (1) From a mixed predict (2) Ferric ions gave (3) On boiling a solution (4) Manganese salts A red solid is insolution red solid in a test tube on the cooler parts of (1) (NH4)2 Cr2O7 Which of the following (1) Zn2[Fe(CN)6] (3) (NH4)3[As(Mo3Offer Content of the following content of the following content of the following (1) Zn2[Fe(CN)6] (3) (NH4)3[As(Mo3Offer Content of the following content of the following content of the following content of the following for the following content of th	lowing statement is c ecipitate of AgCl and a deep green precipi ition having K ⁺ , Ca ²⁺ a give a violet borax b ble in water. Howeve be results in liberation of the test tube. The r (2) HgI ₂ ng compounds is not 10)4] organic acid 'X' pro	correct ? Agl, ammonia solution dis tate on adding potassium f and HCO3 ⁻ ions we get a p bead test in the reducing fla er it becomes soluble if som n of some violet coloured f red solid is : (3) HgO colored yellow ? (2) K3[Co(NO2)6] (4) BaCrO4	[AIEEE 2003, 3/225 ssolves only AgCl. ferrocyanide solution. precipitate of K ₂ Ca(CO ₃) ₂ . ame . me KI is added to water. Heating the umes and droplets of a metal appe [AIEEE 2003, 3/225 (4) Pb ₃ O ₄ .
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3.		ted HCl is added to an aqui hich complex ion gives blue	e colour in this reaction				
	(1) [CoCl ₄] ²⁻	(2) [CoCl ₆] ^{3–}	[JEE(M a (3) [CoCl ₆] ^{4–}	hin) 2015 Online (11-04-15) (4) [Co(H ₂ O) ₆] ²⁺	, 4/120]		
4.	A pink coloured s	salt turns blue on heating.	•	cation is most likely ? /ain) 2015 Online (11-04-1	5). 4/1201		
	(1) Co ²⁺	(2) Cu ²⁺	(3) Zn ²⁺	(4) Fe ²⁺	,,		
5.	in dil. HCI produ		with NaOH solution a	sing H ₂ S. A solution of this nd bluish-white precipitate Main) 2017 Online (08-04-1 (4) Ni ²⁺	with basic		
6.	 (1) Mn²⁺ (2) Zn²⁺ (3) Co²⁺ (4) Ni²⁺ The incorrect statement is : [JEE(Main) 2018 Online (16-04-18), 4/120] (1) Cu²⁺ ion gives chocolate coloured precipitate with potassium ferrocyanide solution. (2) Cu²⁺ and Ni²⁺ ions give black precipitate with H₂S in presence of HCl solution. 						

- (3) Ferric ion gives blood red colour with potassium thiocyanate.
- (4) Cu²⁺ salts give red coloured borax bead test in reducing flame.



				EXER	CISE	- 2			
1.	(A - p,q,r ; I	B - r,s ; C -	p,q,r,s,t ; D -	· ·	2.	(A - p,q ; B -	p,s ; C - s ;	D - p,r)	
		_			RT - III				
F-3.	(C)	F-4.	(B)	F-5.	(C)	F-6.	(C)	F-7.	(C)
E-4.	(D)	E-5.	(A)	E-6.	(D)	F-1.	(D)	F-2.	(C)
D-4.	(B)	D-5.	(C)	E-1.	(A)	E-2.	(C)	E-3.	(B)
C-1.	(D)	C-2.	(B)	D-1.	(D)	D-2.	(B)	D-3.	(B)
B-5.	(A)	B-6.	(D)	B-7.	(C)	B-8.	(B)	B-9.	(D)
A-6.	(A)	B-1.	(C)	B-2.	(C)	B-3.	(A)	B-4.	(C)
A-1.	(C)	A-2.	(B)	A-3.	(C)	A-4.	(B)	A-5.	(C)
			, <u>, , , , 2</u> 0u	, -	RT - II				
F-5.	White preci Ca ²⁺ + 2K ⁺		tained. ∣ ^{4–} —→ K₂Ca	[Fe(CN)₅] J					
-			$2^{2-} \longrightarrow (NH_4)$,		e complex).			
F-4.	-		\rightarrow SrSO ₄ \downarrow (white) + N⊦	4+				
F-3.		- 、 /	recipitated.		I + (Yenc				
F-2.			ammonium c ₅] —→ (NH₄)₃						
F-1.			NH ² or Hg						
E-4.	(A) : Zn (N0	D 3)2	(B) : NO ₂	(C) : Z	ZnO	(D) : ZnS			
		sence of th	(whit $N)_4]^{2-} \longrightarrow C$ e copper ions	u[Hg(SCN).		lex Co-precipita	ated with tha	t of zinc com	plex and
E-3.	→ Zn ²⁺ + [⊦	lg(SCN)₄]²⁻	→Zn[Hg(S						
E-2.	It is due to	the lowering	$-2 H_2O$ g of OH ⁻ ion of attained	concentratio		l₄⁺ use of common	ion effect of	⁻ NH₄⁺ to such	n a value
			•			in backward di	ection.		
E-1.		-	→ Mn(O		-	-			
D-3.	Only Fe(II) salts give soluble red iron(II) dimethylglyoxime in ammonical solution. (i) White precipitate, K ₂ Fe[Fe(CN) ₆] and (ii) Pale blue precipitate.								
D-2.	otherwise F	e ²⁺ would r	not be precipi	tated in gro	up III.		-		
C-4. D-1.	(A) is SnCl ₂ Fe ²⁺ salt a		to Fe ³⁺ sal	t by boiling	a with c	onc. HNO3, be	fore adding	NH₄CL and	NH₄OH
C-3. C-4.	3As ₂ S ₃ + 28 H ₃ AsO ₄ (y)	8HNO₃+4H₂ + 12 NH₄M		.sO ₄ (y) + 18	3H + 9S	O₄ ^{2–} + 28 NO↑ 12MoO₃↓(yellov	w) +21 NH4N	1O3+12H2O	
C-2.	solution car	n not be filte	ered causing	unnecessa	ry troubl		AS_2S_3 (both	yellow ppt).	Colloidal
C-1.	 Presence of Sb³⁺ or Bi³⁺, their chloride hydrolyse to oxychlorides in presence of excess of water. BiCl₃ + H₂O → BiOCl + 2HCl SbCl₃ + H₂O → SbOCl + 2HCl HNO₃ is a powerful oxidising solution. Causing confusion with CdS, AS₂S₃ (both yellow ppt). Colloidal solution can not be filtered causing unnecessary trouble. 								

				PA	RT - I				
1.	(D)	2.	(A)	3.	(B)	4.	(C)	5.	(D)
6.	(D)	7.	(D)	8.	(B)	9.	(D)	10.	(D)
11.	(C)	12.	(D)	13.	(C)	14.	(B)	15.	(A)
16.	(C)	17.	(C)						
				PA	RT - II				
1.	3 (Hg ₂ ²⁺ , Pb	²⁺ , Ag ⁺)		2.	4	3.	4	4.	2
5.	3 (a,b,c) 6. 3 (I, IV, V) 7. 20 8. a = 6 ; b = 3 (a + b = 9						= 9)		
9.	5 (i), (ii), (iv)), (v), (vi)							ŗ
				Þ۸	RT - III				
1.	(ABCD)	2.	(ABC)	3.	(CD)	4.	(BC)	5.	(ABCD)
6.	(ABCD)	 7.	(ABC)	8.	(ACD)	9.	(AC)	10.	(AD)
11.	(ABC)		(7120)	0.	(/(02)	0.	(/(0)		(7.2)
•••	(7.20)								
				PAF	RT - IV				
1.	(C)	2.	(B)	3.	(D)	4.	(B)	5.	(B)
6.	(D)	7.	(C)	8.	(D)	9.	(B)		
			E	EXER	CISE - 3				
				PA	RT - I				
1.	(B)								
2.	(a) (A) = Pe	ntaaquath	iocyanato-S-iroi	n(III) ; (B)	= Hexafluorio	oferrate(III) (b) 5.93 B.M.		
3.	(B)	4.	(B)	5.	(C)	6.	(B)	7.*	(AB)
8.	(D)	9.	(C)	10.	(B)	11.	(B)	12.	(A)
13.	(C)	14.	(A)	15.*	(BCD)	16.*	(ACD)	17.	(D)
18.	(A)	19.	(D)	20.	7	21.*	(CD)	22.	(A)
23.*	(BD)								
				PA	RT - II				
			JEE(MAI	N) OFF	LINE PRO	BLEMS			

1.	(1)	2.	(2)	3.	(1)	4.	(3)		
JEE(MAIN) ONLINE PROBLEMS									
1.	(3)	2.	(3)	3.	(1)	4.	(1)	5.	(2)
6.	(2)								