DCAM classes

# SOLVED EXAMPLES

Ex. 1 Amongst  $[TiF_6]^{2-}$ ,  $[CoF_6]^{3-}$ ,  $Cu_2Cl_2$  and  $[NiCl_4]^{2-}$  [Atomic number ; Ti = 22, Co = 27, Cu = 29, Ni = 28] the colourless species are :

(A)  $[TiF_6]^{2-}$  and  $[Cu_2Cl_2]$  (B)  $Cu_2Cl_2$  and  $[NiCl_4]^{2-}$  (C)  $[TiF_6]^{2-}$  and  $[CoF_6]^{3-}$  (D)  $[CoF_6]^{3-}$  and  $[NiCl_4]^{2-}$ 

Ans. (A)

Ans.

Sol.

Sol. In  $[TiF_6]^{2-}$  the titanium is in +4 oxidation state having the electronic configuration  $[Ar]^{18} 3d^0 4s^0$ . Similarly in  $Cu_2Cl_2$  the copper is in +1 oxidation state having the electronic configuration  $[Ar]^{18} 3d^{10} 4s^0$ . As they do not have any unpaired electrons for d-d transition, they are therefore colourless.

In  $[NiCl_4]^{2-}$  the nickel is in +2 oxidation state and electronic configuration is  $[Ar]^{18} 3d^8 4s^0$ . As it has two unpaired electrons, so the complex is coloured.

In  $[CoF_6]^{3-}$ , the cobalt is in +3 oxidation state having electron configuration [Ar]  $3d^6 4s^0$ . As it has four unpaired electrons, so the complex is coloured.

- Ex.2 On the basis of trends in the properties of the 3d-series elements, suggests possible  $M^{2+}$  aqua ions for use as reducing agents, and write a balanced chemical equation for the reaction of one of these ions with  $O_2$  in acidic solution.
- **Sol.** Because oxidation state +2 is most stable for the later elements of 3d-series elements, strong reducing agents include ions of the metals on the left of the series: such ions include  $V^{2+}$  (aq) and  $Cr^{2+}$  (aq) The Fe<sup>2+</sup> (aq) ion is only weakly reducing. The Co<sup>2+</sup> (aq), Ni<sup>2+</sup> (aq), and Cu<sup>2+</sup> (aq) ions are not oxidized in water.

 $Fe^{3+} \xrightarrow{+0.77} Fe^{2+} \xrightarrow{-0.44} Fe$ 

The chemical equation for the oxidation is then

 $4 \operatorname{Fe}^{2+}(aq) + \operatorname{O}_{2}(g) + 4\operatorname{H}^{+}(aq) \longrightarrow 4\operatorname{Fe}^{3+}(aq) + 2\operatorname{H}_{2}\operatorname{O}(\ell).$ 

**Ex.3** Match the reactions given in column-I with the characteristic(s) of the reaction products given in column-II.

	Column-I	Column-II			
(A)	$\text{TiCl}_4$ Zn	(p) One of the products is bright orange coloured but diamagnetic.			
<b>(B)</b>	FeCl <sub>3</sub> 573K	(q) One of the products is green coloured and paramagnetic.			
(C)	$KMnO_4 \xrightarrow{750K}$	(r) One of the products is violet and paramagnetic.			
<b>(D)</b>	$K_2Cr_2O_7 + H_2SO_4$ (cold & conc.)	(s) One of the products exists as dimer.			
[A-r];	[B-s; [C-q]; [D-p].				
(A) $\text{TiCl}_4 \xrightarrow{\text{Zn}} \text{TiCl}_3$ , violet (one unpaired electron so d-d transition is possible).					
( <b>B</b> ) 2Fe		≻CI Scl gas dimer.			

(C) 
$$2KMnO_4 \xrightarrow{750K} K_2MnO_4$$
 green (one unpaired electron so d-d transition is possible) +  $MnO_2 + O_2$ .

**(D)**  $K_2Cr_2O_7 + 2H_2SO_4 \longrightarrow 2CrO_3$  bright orange (diamagnetic) + 2KHSO\_4 + H\_2O.

- **Ex.4** Among the following statements choose the true or false statement(s).
  - (a)  $K_2Cr_2O_7$  on heating with charcoal gives metallic potassium and  $Cr_2O_3$ .
  - (b) On heating in current of  $H_2$  the crystalline KMnO<sub>4</sub> is converted into KOH and Mn<sub>3</sub>O<sub>4</sub>.
  - (c) Hydrated ferric chloride on treatment with 2, 2-dimethoxypropane gives anhydrous ferric chloride.
- Ans. (a) False (b) False (c) True
- Sol. (a)  $K_2Cr_2O_7 + 2C$  (charcoal)  $\xrightarrow{\Delta} Cr_2O_3 + K_2CO_3 + CO\uparrow$ .

**(b)**  $2KMnO_4 + 5H_2 \xrightarrow{\Delta} 2KOH + 2MnO + 4H_2O.$ 

(c) FeCl<sub>3</sub>. 
$$6H_2O + 6CH_3 \longrightarrow FeCl_3$$
 (anhydrous) + 12CH<sub>3</sub>OH + 6CH<sub>3</sub>COCH<sub>3</sub>.

- **Ex.5** A compound (A) is used in paints instead of salts of lead. Compound (A) is obtained when a white compound (B) is strongly heated. Compound (B) is insoluble in water but dissolves in sodium hydroxide forming a solution of compound (C). The compound (A) on heating with coke gives a metal (D) and a gas (E) which burns with blue flame. (B) also dissolves in ammonium sulphate solution mixed with ammonium hydroxide. Solution of compound (A) in dilute HCl gives a bluish white / white precipitate (F) with excess of  $K_4$ [Fe(CN)<sub>6</sub>]. Identify (A) to (F) and explain the reactions.
- Ans. (A) ZnO, (B)  $Zn(OH)_2$ , (C)  $Na_2ZnO_2$ , (D) Zn, (E) CO, (F)  $K_2Zn_3$  [Fe(CN)<sub>6</sub>]<sub>2</sub>

Sol. 
$$Zn(OH)_2$$
 (B)  $\longrightarrow$   $ZnO$  (A) + H<sub>2</sub>O

 $Zn(OH)_{2}(\mathbf{B}) \downarrow + 2OH^{-} \longrightarrow [Zn(OH)_{4}]^{2-}(\mathbf{C})$  (soluble complex).

 $ZnO(\mathbf{A}) + C \xrightarrow{\Delta} Zn(\mathbf{D}) + CO(E).$ 

 $Zn(OH)_2$  (B) + 4NH<sub>3</sub>  $\longrightarrow$  [Zn(NH<sub>3</sub>)<sub>4</sub>]<sup>2+</sup> (soluble complex) + 2OH<sup>-</sup>.

 $ZnO + 2HCl \longrightarrow ZnCl_2 + H_2O.$ 

 $3\text{ZnCl}_2 + 2\text{K}_4[\text{Fe}(\text{CN})_6] \longrightarrow \text{K}_2\text{Zn}_3[\text{Fe}(\text{CN})_6]_2 \downarrow \text{(bluish white/white)}(\text{F}) + 6\text{KCl}.$ 

- **Ex.6** An unknown inorganic compound (X) gave the following reactions:
  - (i) The compound (X) on heating gave a residue, oxygen and oxide of nitrogen.
  - (ii) An aqueous solution of compound (X) on addition to tap water gave a turbidity which did not dissolve in HNO<sub>3</sub>.
  - (iii) The turbidity dissolves in  $NH_4OH$ .

Identify the compound (X) and give equations for the reactions (i), (ii) & (iii).

**Ans.** 
$$X = AgNO_3$$

Sol.  $2 \text{AgNO}_3(X) \xrightarrow{\Delta} 2 \text{Ag} + 2 \text{NO}_2 + \text{O}_2$ .

 $AgNO_3(aq.) + Cl^- \longrightarrow AgCl \downarrow (white) + NO_3^-.$ 

 $AgCl + 2NH_3 \longrightarrow [Ag(NH_3)_2]^+$  (soluble complex).

<b>Ex.</b> 7	Which of the following is true for the species have	ing 3d <sup>4</sup> configuration ?
	(A) $Cr^{2+}$ is reducing in nature.	<b>(B)</b> Mn <sup>3+</sup> is oxidising in nature.
	(C) Both (A) and (B)	(D) None of these
Sol.	$Cr^{2+}$ is reducing as its configuration changes from d in octahedral crystal field spliting. On the other ha configuration which has extra stability. Therefore, ( <b>C</b> ) option is correct.	$^4$ to d <sup>3</sup> , the latter having a half-filled t <sup>3</sup> $_{2g}$ energy level of 3d orbitals and, the change from Mn <sup>3+</sup> to Mn <sup>2+</sup> results in the half-filled (d <sup>5</sup> )
<b>Ex. 8</b>	Which of the following increasing order of oxidisi	ng power is correct for the following species ?
	$VO_{2}^{+}, MnO_{4}^{-}, Cr_{2}O_{2}^{2-}$	Cr
	(A) $VO_2^+ < Cr_2O_2^{-2} < MnO_4^{-2}$	<b>(B)</b> $VO_2^+ < MnO_4^- < Cr_2O_7^{2-}$
	(C) $Cr_{2}O_{7}^{2-} < VO_{7}^{+} < MnO_{4}^{-}$	( <b>D</b> ) $\operatorname{Cr}_{2}O_{7}^{2-} < \operatorname{MnO}_{4}^{-} < \operatorname{VO}_{2}^{+}$
Sol.	This is attributed to the increasing stability of the	lower species to which they are reduced.
	$MnO_4^{-}$ is reduced to $Mn^{2+}$ which has stable half fill	led valence shell electron configuration [3d <sup>5</sup> ].
	$Cr_2O_7^{2-}$ is reduced to $Cr^{3+}$ which has half filled $t^{3}_{2a}$	energy level of 3d orbitals in octahedral crystal field spliting
	$VO_{3}^{+}$ is reduced to $V^{3+}$ which has electronic config	guration $[Ar]^{18} 3d^2 4s^0$ .
	So the order of increasing stability of the reduced of oxidising power is $VO_2^+ < Cr_2O_7^{-2} < MnO_4^{-1}$ .	species is $Mn^{2+} > Cr^{3+} > V^{3+}$ and, therefore, the increasing order
	Therefore, (A) option is correct.	
Ex. 9.	Which of the following statement(s) is/are corr	ect?
	(A) Transition metals and many of their compo	unds show paramagnetic behaviour.
	(B) The enthalpies of atomisation of the transit	ion metals are high
	(C) The transition metals generally form colour	red compounds
	(D) Transition metals and their many compoun	ds act as good catalyst.
Sol.	<ul> <li>(A) As metal ions generally contain one or mo generally paramagnetic</li> </ul>	ore unpaired electrons in them & hence their complexes are
	(B) Because of having larger number of unpair interaction and hence stronger bonding be	red electrons in their atoms, they have stronger inter atomic tween the atoms.
	(C) According to CFT, in presence of ligands the electrons.	he colour of the compound is due to the d-d transition of the
	(D) This activity is ascribed to their ability to ac	dopt multiple oxidation state and to form complexes.
	Therefore, (A,B,C,D) options are correct.	
Ex. 10	Statement-1 : The number of unpaired electrons i	n the following gaseous ions :
	$Mn^{3+}$ , $Cr^{3+}$ , $V^{3+}$ and $Ti^{3+}$ are 4, 3, 2 a	nd 1 respectively.
	Statement-2 : Cr <sup>3+</sup> is most stable in aqueous solut	tion among these ions.
	(A) Statement-1 is True, Statement-2 is True; State	ement-2 is a correct explanation for Statement-1.
	(B) Statement-1 is True, Statement-2 is True; State	ment-2 is NOT a correct explanation for Statement-1
	(C) Statement-1 is True, Statement-2 is False	
	(D) Statement-1 is False, Statement-2 is True	
Sol.	<b>(B)</b> $Mn^{3+} = [Ar]^{18} 3d^4$ , $Cr^{+3} = [Ar]^{18} 3d^3$ , $V^{3+} = [Ar]^{18} 3d^3$	$^{18} 3d^2$ , $Ti^{3+} = [Ar]^{18} 3d^1$
	Cr <sup>3+</sup> is most stable in aqueous solution because it h	has half filled $t_{2}^3$ energy level of 3d orbitals in octahedral crystal

field spliting and according to crystal field theory (CFT) it has highest value of CFSE i.e.  $1.2 \Delta_0$ .

S1 : Interstitial compounds have high melting points, higher than those of pure metals. Ex. 11 S2 : Permanganate titrations in presence of hydrochloric acid are unsatisfactory. **S3**: KMnO<sub>4</sub> does not act as an oxidising agent in strong alkaline medium. S4 :  $KMnO_4$  on heating in a current of  $H_2$  gives MnO. (A) T T F T (B) T F F T (C) T F T T (D) F F T F Sol. **S1**: Due to strong interatomic forces. S2 : Some of the hydrochloric acid is oxidised to chlorine and thus we get less volume of  $KMnO_4$  than the actual one. **S3**: MnO<sub>4</sub><sup>-</sup> + e<sup>-</sup>  $\longrightarrow$  MnO<sub>4</sub><sup>2-</sup> **S4**:  $2KMnO_4 + 5H_2 \longrightarrow 2KOH + 2MnO + H_2O.$ Therefore, (A) option is correct. Ex. 12 Match the reactions in Column I with nature of the reactions/type of the products in Column II. **Column II** Column I (A)  $AgNO_{2}(aq) + I_{2}(excess) + H_{2}O \longrightarrow$ (**p**) Disproportionation (B)  $K_2MnO_4(aq) + CO_2(g) \longrightarrow$ (q) Comproportionation (C) Na<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>+C  $\longrightarrow$ (r) Redox (D)  $CuCl_{aq}(aq) + Cu(s) \longrightarrow$ (s) One of the products is insoluble in water  $(A \rightarrow p, r, s)$ ;  $(B \rightarrow p, r, s)$ ;  $(C \rightarrow r, s)$ :  $(D \rightarrow q, r, s)$ Ans. (A)  $5\text{AgNO}_3(aq) + 3\overset{0}{I}_2(\text{excess}) + 3\text{H}_2\text{O} \longrightarrow \overset{+\text{V}}{\text{HIO}_3} + 5\overset{-1}{\text{SAgI}} + 5\text{HNO}_3$ Sol. So it is redox and disproportionation reaction. AgI insoluble in water. **(B)**  $3K_2^{+VI}$   $MnO_4(aq) + 2CO_2(g) \longrightarrow 2KMnO_4 + MnO_2 + 2K_2CO_3$ So it is redox and disproportionation reaction. MnO, insoluble in water. (C)  $2C^{0} + Na_{2}Cr_{2}O_{7} \xrightarrow{\Delta} Cr_{2}O_{3} + Na_{2}CO_{3} + CO^{+III} + Na_{2}O_{3} + CO^{+III} + CO^{+III} + CO^{+III} + CO^{-1}$ So it is redox reaction. Cr<sub>2</sub>O<sub>3</sub> (green pigment) is insoluble in water. (**b**)  $\overset{\text{+II}}{\text{CuCl}_2}$  (aq) +  $\overset{0}{\text{Cu}}$  (s)  $\longrightarrow \overset{\text{+I}}{\text{Cu}_2\text{Cl}_2}$  (s) So it is redox and comproportionation reaction. Cu<sub>2</sub>Cl<sub>2</sub> is insoluble in water. **Ex.13** When CO<sub>2</sub> is passed into aqueous : (A) Na,  $CrO_4$  solution, its yellow colour changes to orange. (B) K<sub>2</sub>MnO<sub>4</sub> solution, it disproportionates to KMnO<sub>4</sub> and MnO<sub>2</sub> (C) Na<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> solution, its orange colour changes to green (D) KMnO<sub>4</sub> solution, its pink colour changes to green. Sol. (A) Na<sub>2</sub>CrO<sub>4</sub>  $H^{+}$  Na<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> (orange colour) (B)  $MnO_4^{2-}$  H<sup>+</sup>  $MnO_4^{-}$  +  $MnO_2$ , in neutral or acidic medium (C) False - In acidic medium no colour change takes place. (**D**)  $MnO_4^- + e^- \xrightarrow{OH} MnO_4^{2-}$ ; in strong alkaline medium pink colour of  $KMnO_4$  changes to green. Therefore, (A,B) options are correct.

- Ex. 14 Which of the following statement(s) is (are) not correct with reference to ferrous and ferric ions
  - (A) Fe<sup>3+</sup> gives brown colour with potassium ferricyanide
  - (B) Fe<sup>2+</sup> gives blue precipitate with potassium ferricyanide
  - (C) Fe<sup>3+</sup> gives red colour with potassium sulphocyanide
  - (D)  $Fe^{2+}$  gives brown colour with potassium sulphocyanide
- **Sol.** Fe<sup>3+</sup> produces red colouration with KSCN but Fe<sup>2+</sup> does not give brown colour with KSCN. Therefore, (**D**) option is correct.
- Ex. 15 Statement-1 : Ammonical silver nitrate converts glucose to gluconic acid and metallic silver is precipitated. Statement-2 : Glucose acts as a weak reducing agent.
  - (A) Statement-1 is True, Statement-2 is True; Statement-2 is a correct explanation for Statement-1.
  - (B) Statement-1 is True, Statement-2 is True; Statement-2 is NOT a correct explanation for Statement-1
  - (C) Statement-1 is True, Statement-2 is False
  - (D) Statement-1 is False, Statement-2 is True
- **Sol.** (A)  $Ag_2O + C_6H_{12}O_6 \rightarrow 2Ag + C_6H_{12}O_7$ .
- **Ex. 16** What is the composition of mischmetal alloy and what are it's uses ?
- Ans. Mischmetal consists of lanthanoid metal (~95%) and iron (~5%) and traces of S,C,Ca and Al. Mischmetal is used in Mg based alloy to produce bullets, shell and lighter flint.



9.	Magnetic moment of $Cr^{+2}$	$(Z=24), Mn^{+2}(Z=25) and$	Fe <sup>2+</sup> (Z=26) are x,y,z. They (C) $z \le x = y$	are in order : (D) $y = 7 \le y$	
	$(\mathbf{A}) \mathbf{X} < \mathbf{y} < \mathbf{Z}$	$(\mathbf{D}) \mathbf{X} \ge \mathbf{y} \ge \mathbf{Z}$	(C)Z > X - y	$(\mathbf{D})\mathbf{X} - \mathbf{Z} > \mathbf{y}$	
10.	The magnetic moment $c$	of <sub>25</sub> Mn in ionic state is $\sqrt{1}$	5 B.M, then Mn is in : $(C) \downarrow 4$ state	$(\mathbf{D}) + 5$ state	
11	$(\mathbf{A}) + 2$ state	( <b>B</b> ) +5 state	(C) +4 state	$(\mathbf{D})$ +5 state	
11.	(A) $Cu^+$ , $Zn^{2+}$ , $Sc^{3+}$	(B) $Mn^{2+}$ , Fe <sup>3+</sup> , Ni <sup>2+</sup>	(C) $Cr^{2+}$ , $Mn^{3+}$ , $Sc^{3+}$	<b>(D)</b> $Cu^{2+}$ , $Ni^{2+}$ , $Ti^{4+}$	
12.	Which of the following h (A) Zn <sup>2+</sup>	as the maximum number of <b>(B)</b> Fe <sup>2+</sup>	unpaired d-electron? (C) Ni <sup>2+</sup>	<b>(D)</b> Cu <sup>2+</sup>	
13.	The highest magnetic mo (A) 3d <sup>5</sup>	ment is shown by the transi (B) 3d <sup>2</sup>	tion metal ion with the oute (C) 3d <sup>7</sup>	ermost electronic configuration is : (D) 3d <sup>9</sup>	
14.	A metal ion from the firs electrons are expected to (A) 1	t transition series has a mag be present in the ion? (B) 2	(C) 3	of 3.87 B.M. How many unpaired (D)4	
15.	The colour of transition metal ions is attributed to :(A) exceptionally small size of cations(C) incomplete (n - 1) d - subshell(D) absorption of infrared radiations				
16.	Which one of the ionic sp (A) Ti <sup>4+</sup>	pecies will impart colour to (B) Cu <sup>+</sup>	an aqueous solution ? (C) Zn <sup>2+</sup>	<b>(D)</b> Cr <sup>3+</sup>	
17.	<ul> <li>MnO<sub>4</sub> - is of intense pink colour, though Mn is in (+7) oxidation state. It is due to :</li> <li>(A) oxygen gives colour to it</li> <li>(B) charge transfer when Mn gives its electron to oxygen</li> <li>(C) charge transfer when oxygen gives its electron to Mn making it Mn(+VI) hence coloured</li> <li>(D) none is correct</li> </ul>				
18.	CuSO <sub>4</sub> .5H <sub>2</sub> O is blue in co (A) It contains water of co (B) SO <sub>4</sub> <sup>2-</sup> ions absorb red (C) Cu <sup>2+</sup> ions absorb red (D) Cu <sup>2+</sup> ions absorb all co	plour because rystallization. l light. light. volours except red from the v	white light.		
19.	The yellow colour of chr (A) Cr <sup>3+</sup>	omates changes to orange o <b>(B)</b> $Cr_2O_3$	on acidification due to formation $(C) \operatorname{Cr}_2 O_7^{2}$	ation of : (D) $\operatorname{CrO}_4^-$	
20.	Cementite is : (A) interstitial compound (C) a compound resembli	l of iron and carbon ing cement	(B) an alloy of Fe and Cr (D) an ore of iron		
21.	<ul> <li>The catalytic activity of t</li> <li>(A) their chemical reactiv</li> <li>(B) their magnetic behavi</li> <li>(C) their filled d-orbitals.</li> <li>(D) their ability to adopt their</li> </ul>	the transition metals and the ity. Tour. Mourle oxidation state and	eir compounds is ascribed to their complexing ability.	o :	
22.	Which forms interstitial (A) Fe	compounds? (B) Co	( <b>C</b> ) Ni	(D) All	
23.	A compound is yellow w $(A) Al_2O_3$	hen hot and white when col (B) PbO	d. The compound is : (C) CaO	(D) ZnO	

24.	On heating $ZnCl_2.2H_2O$ , (A) $ZnCl_2$	the compound obtained is : (B) Zn(OH)Cl	(C) Zn(OH) <sub>2</sub>	<b>(D)</b> Zn	
25.	When copper is placed in of the green crust is :	the atmosphere for sufficie	nt time, a green crust is forn	ned on its surface. The composition	
	(A) $Cu(OH)_2$	(B) CuO	(C) $CuCO_3$	<b>(D)</b> $CuCO_3$ . $Cu(OH)_2$	
26.	$KMnO_4$ is the oxo salt of (A) $MnO_2$	: <b>(B)</b> Mn <sub>2</sub> O <sub>7</sub>	(C) $MnO_3$	<b>(D)</b> $Mn_2O_3$	
27.	When $K_4[Fe(CN)_6]$ is add (A) $Fe_3[Fe(CN)_6]_4$	ded to $\text{FeCl}_3$ , the complex co ( <b>B</b> ) $\text{Fe}_4[\text{Fe}(\text{CN})_6]_3$	ompound formed is : (C) K <sub>2</sub> Fe[Fe(CN) <sub>6</sub> ]	<b>(D)</b> $K_2Fe_3[Fe(CN)_6]_2$	
28.	FeCl <sub>3</sub> .6H <sub>2</sub> O is actually : (A) [Fe(H <sub>2</sub> O) <sub>6</sub> ]Cl <sub>3</sub> (C) [Fe(H <sub>2</sub> O) <sub>4</sub> Cl <sub>2</sub> ]Cl.2H <sub>2</sub> O		(B) [Fe(H <sub>2</sub> O) <sub>5</sub> Cl]Cl <sub>2</sub> .H <sub>2</sub> O (D) [Fe(H <sub>2</sub> O) <sub>3</sub> Cl <sub>3</sub> ].3H <sub>2</sub> O		
29.	The solubility of silver be $(A) Ag_2SO_3$	romide in hypo solution (ex <b>(B)</b> $Ag_2S_2O_3$	(C) $[Ag(S_2O_3)]^-$	on of : <b>(D)</b> $[Ag(S_2O_3)_2]^{3-}$	
30	In dilute alkaline solution (A) $MnO_4^{2-}$	n, $MnO_4^-$ changes to : (B) $MnO_2$	(C) $Mn_2O_3$	(D) MnO	
31.	$Cl_2$ gas is obtained by va (A) KMnO <sub>4</sub> (s)+conc l	$\begin{array}{c} \text{trious reactions but not by :} \\ \text{HCl} \xrightarrow{\Delta} \end{array}$	( <b>B</b> ) KCl(s)+K <sub>2</sub> Cr <sub>2</sub> O <sub>7</sub> (s)	(B) KCl(s)+K <sub>2</sub> Cr <sub>2</sub> O <sub>7</sub> (s)+conc H <sub>2</sub> SO <sub>4</sub> $\longrightarrow$	
	(C) $MnO_2(s) + conc HC$	$1 \xrightarrow{\Delta}$	( <b>D</b> ) KCl(s)+ $F_2(\ell) \longrightarrow$		
32.	FeCl, dissolves in :				
	(A) water	(B) ether	(C) ammonia	<b>(D) (A) and (B) both</b>	
33.	Which of the following compounds is used as the starting material for the preparation of potassium dichroma(A) $K_2SO_4.Cr_2(SO_4)_3.24H_2O$ (chrome alum)(B) PbCrO_4 (chrome yellow)(C) FeCr.O_4 (chromite)(D) PbCrO_4.PbO (chrome red)				
34.	$\operatorname{CrO}_3$ dissolves in aqueor (A) $\operatorname{CrO}_4^{2-}$	us NaOH to give : (B) Cr(OH) <sub>3</sub>	(C) $Cr_2O_7^{2-}$	<b>(D)</b> Cr(OH) <sub>2</sub>	
35.	CuSO <sub>4</sub> solution + lime is (A) Lucas reagent (C) Fehling solution A	called :	<ul><li>(B) Fenton's reagent</li><li>(D) Bordeaux mixture</li></ul>		
36.	The developer used in pl (A) hydroquinone	hotography is an alkaline so (B) glycerol	olution of : (C) phenol	(D) picric acid	
37.	When acidified solution of $K_2Cr_2O_7$ is shaken with aqueous solution of $FeSO_4$ , then :(A) $Cr_2O_7^{2-}$ ion is reduced to $Cr^{3+}$ ions(B) $Cr_2O_7^{2-}$ ion is converted to $CrO_4^{2-}$ ions(C) $Cr_2O_7^{2-}$ ion is reduced to $Cr$ (D) $Cr_2O_7^{2-}$ ion is converted to $CrO_3$				
38.	The final products obtain	ned for the following reaction	on is :		
	$\operatorname{KWINO}_4 (\operatorname{excess})$ (A) Mn <sub>2</sub> O <sub>2</sub>	$P + H_2 SO_4$ (concentrated ar (B) MnO	$(C) \operatorname{Mn}_{3}O_{4}$	<b>(D)</b> $MnO_3^+$	
39.	When $AgNO_3$ (aq) reacts (A) $AgIO_3$	with excess of iodine, we g ( <b>B</b> ) HIO <sub>3</sub>	get : (C) AgO	( <b>D</b> ) HI	

40.	$ZnO + CoO \xrightarrow{\Delta} X$ ; P	roduct 'X' colour is :			
	(A) Green	(B) Blue	(C) Pink	(D) Bluish green	
41.	The compound that gets $(A) \operatorname{Co}_2(\operatorname{SO}_4)_3$	(B) NiSO <sub>4</sub>	to atmosphere is : (C) KMnO <sub>4</sub>	(D) FeSO <sub>4</sub>	
42.	<ul> <li>The f-block of the periodic table contains those elements in which :</li> <li>(A) only 4f orbitals are progressively filled in 6th period.</li> <li>(B) only 5f orbitals are progressively filled in 7th period.</li> <li>(C) 4f and 5f orbitals are progressively filled in 6th and 7th periods respectively.</li> <li>(D) none</li> </ul>				
43.	Among the lanthanoides (A) Lu	the one obtained by synthe (B) Pm	tic method is : (C) Pr	<b>(D)</b> Gd	
44.	The most common lanthat (A) lanthanum	noide is : (B) cerium	(C) samarium	<b>(D)</b> plutonium	
45.	<ul> <li>The lanthonoide contraction is reponsible for the fact that</li> <li>(A) Zr and Y have about the same radius</li> <li>(B) Zr and Nb have similar oxidation state</li> <li>(D) Zr and Ce have the same oxidation state</li> </ul>				
46.	Lanthanoid and actinides resemble in :(A) electronic configuration(C) ionization energy(D) formation of complexes				
47. 48.	The separation of lanthanoids by ion exchange method is based on(A) sizes of the ions(B) oxidation state of the ions(C) the solubility of their nitrates(D) basicity of hydroxides of lanthanidesAcross the lanthanide series, the basicity of the lanthanoide hydroxides :(A) increases(A) increases(B) decreases(C) first increases and then decreases(D) does not change			ions s of lanthanides	
49.	The +3 ion of which one of (A) La	of the following has half fill (B) Lu	ed 4f subshell ? (C) Gd	(D) Ac	
50.	Actinides : (A) are all synthetic eleme (C) have only short lived	ents isotopes	<ul><li>(B) includes element 104</li><li>(D) have variable valency</li></ul>		
51.	<ul> <li>Atoms of the transition elements are smaller than those of the s-block elements, because :</li> <li>(A) there is increase in the nuclear charge along the period.</li> <li>(B) orbital electrons are added to the penultimate d-subshell rather than to the outer shell of the atom.</li> <li>(C) the shielding effect of d-electrons is small.</li> <li>(D) All of these</li> </ul>				
52.	<ul> <li>Which of the following fa</li> <li>(A) Poor shielding of 4 f-</li> <li>(B) Effective shielding of 6</li> <li>(C) Poorer shielding of 5</li> <li>(D) Greater shielding of 5</li> </ul>	actor may be regarded as the electrons in compare to othe one of the 4 f-electrons by d electron by 4 f- electrons. d electron by 4 f- electron.	e main cause of Lanthanide her electrons in the sub-shell another in the sub-shell.	contraction ?	
53.	Which of the following tr (A) Copper	ansition metal ions has the (B) Nickel	lowest density ? (C) Scandium	(D) Zinc	

54.	The correct ground state electronic configuration of chromium atom (Z = 24) is : (A) [Ar] $4d^5 4s^1$ (B) [Ar] $3d^4 4s^2$ (C) [Ar] $3d^6 4s^0$ (D) [Ar] $3d^5 4s^1$
55.	Transition metals :(A) exhibit only diamagnetism(B) undergo inert pair effect(C) do not form alloys(D) show variable oxidation states
56.	<ul> <li>The radii (metallic) of Fe,Co and Ni are nearly same.</li> <li>This is due to : <ul> <li>(A) lanthanide contraction</li> <li>(B) the fact that successive addition of d–electrons screen the outer electrons (4s) from the inward pull of the nucleus.</li> <li>(C) increase in radii due to increase in 'n' is compensated by decrease in radii due to increase in effective nuclear charge (Z).</li> <li>(D) atomic radii do not remain constant but decrease in a normal gradation.</li> </ul> </li> </ul>
57.	<ul> <li>Of the ions Zn<sup>2+</sup>, Ni<sup>2+</sup> and Cr<sup>3+</sup> (atomic number Zn = 30, Ni = 28, Cr = 24) :</li> <li>(A) only Zn<sup>2+</sup> is colourless and Ni<sup>2+</sup> and Cr<sup>3+</sup> are coloured.</li> <li>(B) all three are colourless.</li> <li>(C) all three are coloured.</li> <li>(D) only Ni<sup>2+</sup> is coloured and Zn<sup>2+</sup> and Cr<sup>3+</sup> are colourless.</li> </ul>
58.	Among the following pairs of ions, the lower oxidation state in aqueous solution is more stable than the other, in: (A) $Zn^{2+}$ , $Zn^{3+}$ (B) $Cu^+$ , $Cu^{2+}$ (C) $Cr^{2+}$ , $Cr^{3+}$ (D) $V^{2+}$ , $VO^{2+}$
<b>59.</b> <b>60.</b>	$\begin{array}{c c c c c c c c c c c c c c c c c c c $
61.	Which one of the following shows highest magnetic moment ?(A) $V^{3+}$ (B) $Cr^{3+}$ (C) $Fe^{3+}$ (D) $Co^{3+}$
62. 63.	Amongst the following the lowest degree of paramagnetism per mole of the compound at 298 K will be shown by:(A) $MnSO_4.4H_2O$ (B) $CuSO_4.5H_2O$ (C) $FeSO_4.6H_2O$ (D) $NiSO_4.6H_2O$ Match the compounds of column X with oxidation state of column Y.Column XColumn YI[Cr(H_2O)_6]Cl_35IIColumn YIIColumn YIIII K_3CrO86IV(NH_3)_3CrO43II <th< td=""></th<>
64.	Which of the following transition element shows the highest oxidation state :(A) Mn(B) Fe(C) V(D) Cr

65.	<ul> <li>Standard reduction electrode potential of Zn<sup>2+</sup> / Zn is – 0.76 V. This means :</li> <li>(A) ZnO is reduced to Zn by H<sub>2</sub></li> <li>(B) Zn can't liberates H<sub>2</sub> with concentrated acids</li> <li>(C) Zn is generally the anode in an electrochemical cell</li> <li>(D) Zn is generally the cathode in an electrochemical cell</li> </ul>					
66.	Which one of the transiti (A) Cu <sup>+</sup>	on metal ions is coloured in (B) Zn <sup>2+</sup>	aqueous solution ? (C) Sc <sup>3+</sup>	<b>(D)</b> V <sup>4+</sup>		
67.	Compound that is both pa (A) $K_2Cr_2O_7$	aramagnetic and coloured is <b>(B)</b> $(NH_4)_2$ [TiCl <sub>6</sub> ]	s : (C) VOSO <sub>4</sub>	<b>(D)</b> $K_3[Cu(CN)_4]$		
68.	Which of the following c (A) $Ag_2SO_4$	ompounds is expected to be <b>(B)</b> $\operatorname{CuF}_2$	e coloured ? (C) $MgF_2$	(D) CuCl		
69.	German silver is an alloy (A) Zn + Ni	of copper and : (B) Al + Ag	(C) $Zn + Ag$	(D) $\operatorname{Sn} + \operatorname{Zn}$		
70.	Ferric sulphate on heating $(A)$ SO <sub>2</sub> and SO <sub>3</sub>	g gives : (B) SO <sub>2</sub> only	(C) SO <sub>3</sub> only	<b>(D)</b> S		
71.	<ul> <li>Anhydrous ferric chloride is prepared by :</li> <li>(A) dissolving ferric hydroxide in dilute HCl.</li> <li>(B) dissolving ferric hydroxide in concentrated HCl.</li> <li>(C) by passing dry Cl<sub>2</sub> gas over heated scrap iron</li> <li>(D) by dissolving iron (III) oxide in concentrated HCl.</li> </ul>					
72.	Most transition metals : I : form sets of compound II : form coloured ions in III : burn vigorously in ox IV : form complex comport of these : (A) I, II, III are correct	ls which display different o solution. xygen. und. (B) II, III, IV are correct	xidation states of the metal. (C) I, II are correct	<b>(D)</b> all are correct.		
73.	The aqueous solution of t (A) $Zn (NO_3)_2$	the following salts will be c (B) LiNO <sub>3</sub>	oloured in the case of : (C) Co(NO <sub>3</sub> ),	(D) Potash alum		
74.	<ul> <li>Which one of the followi</li> <li>(A) Colour of hydrated ic</li> <li>(B) Variable oxidation state</li> <li>(C) High enthalpy of atom</li> <li>(D) Paramagnetic behavior</li> </ul>	ng characteristics of the tra ons. tes. nization. our.	nsition metals is associated	with their catalytic activity?		
75.	At 300°C, FeCl <sub>3</sub> : (A) decomposes into FeC (C) sublimes to give liqui	$Cl_2$ and $Cl_2$ d FeCl_3	<ul><li>(B) decomposes into Fe a</li><li>(D) sublimes to give gase</li></ul>	nd $Cl_2$ ous dimer (FeCl_3) <sub>2</sub>		
76.	Iron is rendered passive I (A) HCl	by treatment with concentration $(\mathbf{B}) \mathrm{H}_2 \mathrm{SO}_4$	tted : (C) $H_3PO_4$	(D) HNO <sub>3</sub>		
77.	Lucas reagent is : (A) Anhydrous $ZnCl_2 + H$ (C) $H_2SO_4 + HCl$	Cl (conc.)	<b>(B)</b> $MnO_2 + H_2O$ <b>(D)</b> $NO + H_2O$			

<b>78.</b>	Sodium thiosulphate is us	ed in photography because	of its :		
	<ul><li>(A) oxidising behaviour</li><li>(C) complexing behaviour</li></ul>		<ul><li>(B) reducing behaviour</li><li>(D) photochemical behaviour</li></ul>	our	
79.	MnO <sub>4</sub> <sup>2-</sup> (1 mole) in neutra (A) 2/3 mole of MnO <sub>4</sub> <sup>-</sup> and (C) 1/3 mole of Mn <sub>2</sub> O <sub>7</sub> and	e) in neutral aqueous medium disproportionates to : $MnO_4^-$ and 1/3 mole of $MnO_2^-$ (B) 1/3 mole of $MnO_4^-$ and 2/3 mole of $MnO_2^-$ $Mn_2O_7$ and 1/3 mole of $MnO_2^-$ (D) 2/3 mole of $MnO_7^-$ and 1/3 mole of $MnO_2^-$			
80.	<ul> <li>When H<sub>2</sub>O<sub>2</sub> is added to an</li> <li>(A) solution turns green di</li> <li>(B) solution turns yellow di</li> <li>(C) a blue coloured composition</li> <li>(D) solution gives green pairs</li> </ul>	acidified solution of $K_2Cr_2$ ue to formation of $Cr_2O_3$ due to formation of $K_2CrO_4$ bund $CrO(O_2)_2$ is formed pt of $Cr(OH)_3$	0 <sub>7</sub> :		
81.	Which one of the followin (A) Fe	ng dissolves in hot concentr (B) Zn	ated NaOH ? (C) Cu	<b>(D)</b> Ag	
82.	The compound used for gr (A) $Cu_2(SCN)_2$	ravimetric estimation of Cu( (B) Cu <sub>2</sub> O	II) is : (C) $Cu_2I_2$	<b>(D)</b> $Cu_2CO_3$	
83.	In the reaction, $2CuCl_2$ (A) $Cu_2Cl_2$	$+2H_2O + SO_2 \rightarrow A + H_2SO_4$ <b>(B)</b> Cu	+ 2HCl ; A is (C) CuSO <sub>4</sub>	(D) CuS	
84.	The number of moles of K (A) 2/5	$MnO_4$ that will be needed to <b>(B)</b> 3/5	o react with one mole of sulp (C) 4/5	phite ion in acidic medium is (D) 1	
<b>85.</b>	Ammonium dichromate is (A) $CrO_3$	used in some fire works. The <b>(B)</b> $Cr_2O_3$	ne green coloured powder b (C) Cr	lown in the air is (D) $CrO(O_2)$	
00.	(A) p-block elements	(B) lanthanoides	(C) actinoides	(D) transition elements	
87.	The number of moles of K solution is :	$MnO_4$ that will be needed to	o react completely with one	mole of ferrous oxalate in acidic	
	(A) 3/5	<b>(B)</b> 2/5	<b>(C)</b> 4/5	<b>(D)</b> 1	
88.	<ul><li>Which of the following is</li><li>(A) Low melting points</li><li>(C) Show variable oxidation</li></ul>	the most suitable description on states.	<ul> <li>on of transition elements ?</li> <li>(B) No catalytic activity.</li> <li>(D) Exhibit inert pair effect</li> </ul>		
89.	<ul> <li>The tendency of the transition elements to form coloured compound is attributed to :</li> <li>(A) transition of electrons from one atoms to the other.</li> <li>(B) transition of electrons from s-orbitals of the outer shells to p-orbitals.</li> <li>(C) d-d-transition of electron in last but one shell.</li> <li>(D) none of the reason is correct.</li> </ul>				
90.	The ions from among the $i$ (i) Ti <sup>4+</sup> , (ii) Cu <sup>+1</sup> (A) (i) and (ii) only	following which are colourl , (iii) Co <sup>3+</sup> , ( <b>B</b> ) (i), (ii) and (iii)	ess are : (iv) $Fe^{2+}$ . (C) (iii) and (iv)	<b>(D)</b> (ii) and (iii).	
91.	Which of the following do (A) Titanium	bes not belong to 3d series (B) Iron	of transition elements ? (C) Palladium	(D) Vanadium.	
92.	For the process $Cu(g) \rightarrow$ (A) 3d sub-shell	$Cu^+(g) + e^-$ , the electron is (B) 4s sub-shell	to be removed from (C) 3p sub-shell	<b>(D)</b> any of the above.	

93.	<ul> <li>In general, the melting and boiling points of transition metals :</li> <li>(A) increase gradually across the period from left to right.</li> <li>(B) decrease gradually across the period from left to right.</li> <li>(C) first increase till the middle of the period and then decrease towards the end.</li> <li>(D) first decrease regularly till the middle of the period and then increase towards the end.</li> </ul>				
94.	The transition elements are more metallic than the representative elements because they have : (A) the electrons in d-orbitals (B) electron pairs in d-orbitals (D) empty metallic orbitals				
95.	Which oxide of manganes (A) MnO	e is most acidic in nature ? (B) Mn <sub>2</sub> O <sub>7</sub>	(C) $Mn_2O_3$	<b>(D)</b> MnO <sub>2</sub> .	
96.	Which of the following tra (A) Co <sup>3+</sup>	ansition metal ions has least (B) Fe <sup>3+</sup>	magnetic moment ?	<b>(D)</b> V <sup>3+</sup>	
97.	<ul> <li>Within each transition series, the oxidation states :</li> <li>(A) regularly decrease from left to right.</li> <li>(C) first decrease upto middle and then increase.</li> <li>(D) none of these.</li> </ul>				
98.	The maximum oxidation st (A) $+5, +6, +3, +3$	tate shown by $V(Z=23)$ , Cr( (B) + 3, +4, +5, +2	(Z = 24), Co(Z = 27), Sc(Z = (C) + 5, +3, +2, +1)	<ul><li>21) are respectively :</li><li>(D) + 4 in each case.</li></ul>	
99.	The stability of particular oxidation state of a metal in aqueous solution is determined by :(A) enthalpy of sublimation of the metal(B) ionisation energy(C) enthalpy of hydration of the metal ion(D) all of these.				
100.	Which of the following sh	all have the highest value of	f magnetic moment ?		
101.	(A) Zn(II) ion The highest oxidation stat	(B) Mn(IV) ion e is exhibited by the transit	(C) Fe(II) ion tion metals with configuration	(D) Ti(III) ion.	
1010	(A) $(n-1) d^3ns^2$	<b>(B)</b> $(n-1) d^5 ns^1$	(C) $(n-1) d^5 ns^2$	<b>(D)</b> $(n-1) d^8 ns^2$ .	
102.	In general, the transition e (A) C	lements exhibit their highes (B) S	st oxidation states in their co (C) S and P	ompounds with elements like : (D) F and O.	
103.	Transition elements are frequently used as catalyst because :(A) of variable oxidation state(B) of high ionic charge(C) large surface area of reactants(D) of their specific nature.				
104.	The first ionisation energies of the elements of the first transition series : (A) increase as the atomic number increase. (B) decrease as the atomic number increase.				

(C) do not show any change as the addition of electrons takes place in the inner (n - 1) d-orbitals. (D) increase from Ti to Mn and then decrease from Mn to Cu.

Exercise # 2 [Multiple Correct Choice Type Questions] Part # I 1. The colour of the transition metal ions is/are due to : (A) d-d transition of electrons in presence of ligands (B) charge transfer from ligand to metal ion. (C) change in the geometry (D) polarisation of anion by cation Which of the following statement(s) is/are not correct ? 2. (A) The blue colour of aqueous CuCl<sub>2</sub> is due to  $[Cu(H_2O)_4]^{2+1}$ (B) The yellow colour of aqueous  $CuCl_{3}$  is due to  $[CuCl_{4}]^{2-1}$ (C) The green colour of aqueous CuCl, is due to the presence of both  $[Cu(H,O)_{4}]^{2+}$  and  $[CuCl_{4}]^{2-}$ (D) The blue colour of aqueous CuCl<sub>2</sub> is due to  $[CuCl_4]^{2-}$ 3. Select correct statement (s). (A)  $MnO_4^{-}$  is intense pink colour due to d-d transition of electron. (B) Cu(I) is diamagnetic while Cu(II) is paramagnetic. (C) CrO<sub>2</sub> is amphoteric oxide. (**D**)  $[Ti(H_2O)_{\ell}]^{3+}$  and  $[Sc(H_2O)_{\ell}]^{3+}$  both are coloured in aqueous solution. 4 In photography, quinol is used as developer according to following reaction.  $\langle \bigcirc \rangle$ -OH +2AgBr +2OH<sup>-</sup>  $\longrightarrow$  O= $\langle$  $\rangle = O + 2Ag + 2H_2O + 2Br$ HO Which of the following describe(s) the role of quinol in this reaction ? (A) It acts as an acid. (B) It acts as a weak base. (C) It acts as an oxidising agent. (D) It acts as a reducing agent. 5. Which of the following statement (s) is/are correct? (A)  $S_2 O_2^{2-}$  oxidise Ag<sup>+</sup> in presence of pyridine and give red colour compound. (B)  $MnO_4^{2-}$  disproportionates to yield  $MnO_4^{-}$  and  $MnO_2$  in presence of H<sup>+</sup> ions. (C) In  $Cr_2O_7^{2-}$  each Cr is linked to four oxygen atoms. (D) Ti<sup>3+</sup> is purple while Ti<sup>4+</sup> is colourless. Following reaction(s) is/are involved in the iodometric estimation. 6. (A)  $\operatorname{Cr}_2 \operatorname{O}_7^{2-} + \operatorname{H}^+ + \operatorname{I}^- \longrightarrow 2\operatorname{Cr}^{3+} + \operatorname{I}_2; \operatorname{I}_2 + \operatorname{S}_2 \operatorname{O}_3^{2-} \longrightarrow \operatorname{S}_4 \operatorname{O}_6^{2-} + \operatorname{I}^-$ (B)  $\operatorname{MnO}_4^- + \operatorname{H}^+ + \operatorname{I}^- \longrightarrow \operatorname{Mn}^{2+} + \operatorname{I}_2; \operatorname{I}_2 + \operatorname{S}_2 \operatorname{O}_3^{2-} \longrightarrow \operatorname{S}_4 \operatorname{O}_6^{2-} + \operatorname{I}^-$ (C)  $MnO_4^- + OH^- + I^- \longrightarrow MnO_2 + I_2; I_2 + S_2O_3^{2-} \longrightarrow S_4O_6^{2-} + I^-$ (D)  $Cr_2O_7^{2-} + OH^- + I^- \longrightarrow 2 Cr^{3+} + I_2; I_2 + S_2O_3^{2-} \longrightarrow S_4O_6^{2-} + I^-$ 7. Correct statements about transition metals are that they : (A) form complex (B) show variable oxidation states (C) show magnetic properties (D) do not form coloured compounds 8. Transition elements have greater tendency to form complexes because they have : (A) vacant d-orbitals (B) small size (C) higher nuclear charge (D) variable oxidation states Which of the following statements are correct when a mixture of NaCl and K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> is gently warmed with conc. H<sub>2</sub>SO<sub>4</sub>? 9. (A) Deep red vapours are liberated (B) Deep red vapours dissolve in NaOH (aq.) forming a yellow solution. (C) Greenish yellow gas is liberated

(D) Deep red vapours dissolve in water forming yellow solution

- 10. Which of the following chemical reaction(s) is/are involved in developing of photographic plate ? (A)  $C_6H_4(OH)_2 + 2AgBr \longrightarrow 2Ag + C_2H_3O_2 + 2HBr$ **(B)** AgBr + 2Na<sub>2</sub>S<sub>2</sub>O<sub>2</sub> $\longrightarrow$  Na<sub>2</sub> [Ag(S<sub>2</sub>O<sub>2</sub>)<sub>2</sub>] + NaBr (C)  $2AgNO_3 + Na_2S_2O_3 \longrightarrow Ag_2S_2O_3 + 2NaNO_3$ (D)  $AgNO_3 + KCN \longrightarrow AgCN + KNO_3$ 11. Pyrolusite is MnO<sub>2</sub> used to prepare KMnO<sub>4</sub>. Steps are :  $MnO_{2} \xrightarrow{I} MnO_{4}^{2-} \xrightarrow{II} MnO_{4}^{-}$ Steps I and II are respectively : (A) fuse with KOH / air, electrolytic oxidation (B) fuse with KOH / KNO<sub>2</sub>, electrolytic oxidation (C) fuse with concentrated HNO<sub>2</sub> / air, electrolytic reduction (D) dissolve in H<sub>2</sub>O, oxidation 12. Which are correct statements ? (A) In less acidic solution  $K_2Cr_2O_2$  and  $H_2O_2$  gives violet coloured diamagnetic  $[CrO(O_2)(OH)]^-$  ion. (B) In alkaline  $H_2O_2$ ,  $K_2CrO_2$  (with tetraperoxo species  $[Cr(O_2)_4]^{3-}$ ) is formed with  $K_2Cr_2O_2$ . (C) In ammonical solution of  $H_2O_2$ ,  $(NH_3)_3CrO_4$  is formed with  $K_2Cr_2O_7$ . (**D**)  $\operatorname{CrO}_{4}^{2-}$  changes to  $\operatorname{Cr}_{2}\operatorname{O}_{7}^{2-}$  by oxidation. 13. Which of the following reaction(s) is/are incorrect for silver nitrate? (A)  $6 \text{ AgNO}_3 + 3I_2 (\text{excess}) + 3 \text{ H}_2\text{O} \longrightarrow \text{AgIO}_3 + 5 \text{ AgI} + 6 \text{HNO}_3$ **(B)** AgNO<sub>3</sub> (excess) + 2 KCN  $\longrightarrow$  K[Ag(CN)<sub>2</sub>] + KNO<sub>3</sub> (C)  $2AgNO_3 + 4Na_2S_2O_3(excess) \longrightarrow 2Na_3[Ag(S_2O_3)_2] + 2NaNO_3$ (D)  $PH_3 + 6AgNO_3 + 3H_2O \longrightarrow 6Ag + 6HNO_3 + H_3PO_3$ 14. Which of the following reaction(s) is/are used for the preparation of anhydrous FeCl, ? (A)  $FeCl_3 6H_2O + 6SOCl_2 \longrightarrow FeCl_3 + 12HCl + 6SO_2$ (B)  $Fe(OH)_3 \downarrow + 3HCl \longrightarrow FeCl_3 + 3H_2O$ (C)  $2Fe + 4HCl (aq.) + Cl_2 \longrightarrow 2FeCl_3 + 2H_2$ (D)  $2Fe + 3Cl_{2}(dry) \rightarrow 2FeCl_{2}$ 15. Which of the following is/are false? (A) Na<sub>2</sub>Cr<sub>2</sub>O<sub>2</sub> is used as a primary standard in volumetric analysis. (B) Potassium permanganate in excess on treatment with conc. H<sub>2</sub>SO<sub>4</sub> forms manganese heptoxide (C) Phosphine, arsine and stibine all precipitates silver from silver nitrate (D) From kipp's apparatus waste ferric sulphate and ferrous sulphate mixture is obtained when air or oxygen is passed for longer time. 16. When CO<sub>2</sub> is passed into aqueous : (A) Na<sub>2</sub>CrO<sub>4</sub> solution, its yellow colour changes to orange.
  - (B)  $K_2$ MnO<sub>4</sub> solution, it disproportionates to KMnO<sub>4</sub> and MnO<sub>2</sub>
  - (C)  $Na_2Cr_2O_7$  solution, its orange colour changes to green
  - **(D)** KMnO<sub>4</sub> solution, its pink colour changes to green.
  - 17. Which of the following statement(s) is/are correct ?
    - (A) Transition metals and many of their compounds show paramagnetic behaviour.
    - (B) The enthalpies of atomisation of the transition metals are high.
    - (C) The transition metals generally form coloured compounds.
    - (D) Transition metals and their many compounds act as good catalyst.

18.	<ul> <li>Potassium manganate (K<sub>2</sub>MnO<sub>4</sub>) is formed when :</li> <li>(A) potassium permanganate reacts with formaldehyde in acidic medium.</li> <li>(B) manganese dioxide is fused with potassium hydroxide in air.</li> <li>(C) manganese hydroxide reacts with K<sub>2</sub>O<sub>2</sub> in alkaline medium.</li> <li>(D) potassium permanganate is heated at 750K.</li> </ul>					
19.	Select the incorrect st (A) In $K_2MnO_4$ and C (B) Both sodium and (C) Potassium dichro (D) Potassium perman	atement(s) : $rO_2Cl_2$ , the central transit potassium dichromate ca mate on strong heating enganate on heating with s	tion metals/ions have same of in be used as primary standa volves oxygen gas and form solid KOH evolves oxygen	oxidation state. ard in volumetric estimations. ns green powder. gas and forms a black powder.		
20.	<ul> <li>Select the correct statement(s) with respect to oxides and oxoanions of transition metals.</li> <li>(A) Among oxides of chromium, CrO is basic, Cr<sub>2</sub>O<sub>3</sub> is amphoteric and CrO<sub>3</sub> is acidic.</li> <li>(B) No higher oxides of iron above Fe<sub>2</sub>O<sub>3</sub> are found.</li> <li>(C)Ti,V, Cr and Mn form oxides MO and their correct increasing order of acidic character is MnO<cro<vo<tio.< li=""> <li>(D) Vandium(V) oxide does not react with acids but reacts with alkalies only.</li> </cro<vo<tio.<></li></ul>					
21.	In which of the follo (A) $KMnO_4$	wing compound(s), the (B) CrO <sub>3</sub>	colour is due to the charge (C) CuCl <sub>2</sub>	ge transfer spectra ? (D) Cu <sub>2</sub> O.		
22.	Which of the followities $(A) \operatorname{Cr}_2 O_3$	ng oxide(s) of transition (B) $V_2O_5$	n metals is/are amphoteric (C) Mn <sub>2</sub> O <sub>7</sub>	in nature. (D) ZnO.		
23.	<ul> <li>The hydrated cupric chloride is strongly heated. Which of the following statement(s) is (are) correct for this?</li> <li>(A) It is reduced to Cu<sub>2</sub>Cl<sub>2</sub>.</li> <li>(B) Cupric oxide is formed along with Cu<sub>2</sub>Cl<sub>2</sub>.</li> <li>(C) Cl<sub>2</sub> is liberated only but it is not liberated along with HCl.</li> <li>(D) Cl. and HCl both are liberated.</li> </ul>					
24.	<ul> <li>Cuprous chloride can be prepared :</li> <li>(A) by passing SO<sub>2</sub> through the solution containing CuSO<sub>4</sub> and NaCl.</li> <li>(B) by heating excess of copper with concentrated HCl in presence of a little KCIO<sub>3</sub>.</li> <li>(C) by boiling copper sulphate solution with excess of copper turnings in presence of hydrochloric acid.</li> <li>(D) by dissolving cupric oxide or copper carbonate in concentrated HCI.</li> </ul>					
State W 25.	ate Weather the Questions is True or False : . S1 : Acidic dichromate solutions on treatment with $H_2O_2$ gives deep blue $CrO(O_2)_2$ . S2 : A deep red liquid, $CrO_2Cl_2$ is formed by the reaction of chromium(III) oxide with HCl in presence of conc. $H_2SO_4$ . S3 : $(NH_4)_2Cr_2O_7$ on heating yields green chromium(III) oxide and nitrogen gas. S4 : $K_2Cr_2O_7$ on heating with charcoal produces $K_2CO_3$ . and arrange in the order of true/false.					
	(A) T T T T	<b>(B)</b> T F T T	(C) T F T F	<b>(D)</b> F F T T		
26.	<b>S1</b> : $Mn^{II}$ ions in solution is oxidised to $MnO_4^-$ by $PbO_2$ or $NaBiO_3$ . <b>S2</b> : $MnO_4^{2-}$ ions in neutral, acid or slightly basic solutions readily disproportionates. <b>S3</b> : $KMnO_4$ gives MnO on heating in current of hydrogen. and arrange in the order of true/ false.					
	(A) I I I	(B) TFT	(C) T F F	(D) F I F.		

**S1**: Interstitial compounds have high melting points, higher than those of pure metals. 27. **S2**: Permanganate titrations in presence of hydrochloric acid are unsatisfactory. **S3**: KMnO<sub>4</sub> does not act as an oxidising agent in strong alkaline medium. **S4** :  $KMnO_4$  on heating in a current of H, gives MnO. (A) T T F T **(B)** T F F T (C) T F T T (D) F F T F 28. S1 :  $Mn^{2+}$  compounds are more stable than  $Fe^{2+}$  towards oxidation to their +3 state. S2: Titanium and copper both in the first series of transition metals exhibits +1 oxidation state most frequently. **S3**: Cu<sup>+</sup> ion is stable in aqueous solutions. S4 : The  $E^{\theta}$  value for the  $Mn^{3+}/Mn^{2+}$  couple is much more positive than that for  $Cr^{3+}/Cr^{2+}$  or  $Fe^{3+}/Fe^{2+}$ . (A) T T F T (B) T F F T (C) T F T T (D) F F T F 29. S1 : Covalent and ionic radii of Nb and Ta are almost the same. **S2**: Ionisation energies of transition elements decrease with increase in atomic number in a given group. **S3**: Iodide of Millon's base is believed to have the structure. Ι S4 : Yellow colour of  $CrO_4^{2-}$  is due to d-d transition. (A) T T F T **(B)** T F F T (C) T F T F (D) F F T F **30**. If a non metal is added to the interstitial sites of a metal, then the metal becomes : **(B)** less tensile (C) less malleable (D) more ductile. (A) softer 31. The reaction  $MnO_4^- + e^- \implies MnO_4^{2-}$  takes place in : (B) acidic medium (A) a basic medium (strong) (D) both acidic and basic medium. (C) neutral medium 32. Reaction of KMnO<sub>4</sub> in neutral or very weakly alkaline solution can be represented as : (A)MnO<sub>4</sub><sup>-+</sup>+2H<sub>2</sub>O+3e<sup>-</sup> $\rightarrow$  MnO<sub>2</sub>+4OH<sup>-</sup>. **(B)**  $2MnO_4^- + 2OH^- \rightarrow 2MnO_4^{2-} + 1/2O_2 + H_2O.$ (C)  $MnO_4^- + 8H^+ + 5e^- \rightarrow Mn^{2+} + 4H_2O_4$ . (D)  $MnO_4^- + e^- \rightarrow MnO_4^{2-}$ . 33. On heating KMnO<sub>4</sub>, one among the following is not formed : (C) MnO<sub>2</sub> (D) MnO. (A) K<sub>2</sub>MnO<sub>4</sub>  $(\mathbf{B})O_{2}$ 34. Among the following series of transition metal ions, the one where all metal ions have  $3d^2$  electronic configuration is (A)  $Ti^{3+}$ ,  $V^{2+}$ ,  $Cr^{3+}$ ,  $Mn^{4+}$ **(B)** Ti<sup>+</sup>, V<sup>4+</sup>, Cr<sup>6+</sup>, Mn<sup>7+</sup> (C)  $Ti^{2+}$ ,  $V^{3+}$ ,  $Cr^{2+}$ ,  $Mn^{3+}$ (**D**)  $Ti^{2+}$ ,  $V^{3+}$ ,  $Cr^{4+}$ ,  $Mn^{5+}$ . 35. The pair of the compounds in which both the metals are in the highest possible oxidation state is, (A)  $[Fe(CN)_{6}]^{3-}$ ,  $[Co(CN)_{6}]^{3-}$ **(B)**  $CrO_{2}Cl_{2}$ ,  $MnO_{4}^{-}$ . (**D**)  $[Co(CN)_{6}]^{3-}$ , Mn<sub>2</sub>O<sub>7</sub>. (C) TiO<sub>2</sub>, MnO<sub>2</sub> Which of the following can be used for the conversion of potassium manganate to potassium permanganate? 36. (B) Cl<sub>2</sub> (C) CO, (D) All.  $(\mathbf{A}) \mathbf{O}_{2}$ One of the products formed due to the reaction between KMnO<sub>4</sub> and HCl is : 37. (A) red liquid, CrO<sub>2</sub>Cl<sub>2</sub> (B) black powder, MnO, (C) greenish yellow gas, Cl, (**D**) colourless liquid,  $HClO_4$ .

38.	Which one of the follow (A) Sulphur dioxide	ing compounds does not de (B) Ferric chloride	colourise an acidified a (C) Hydrogen peroxi	equeous solution of KMnO <sub>4</sub> ? de <b>(D)</b> Ferrous sulphate.
39.	Which of the following it (A) Only $MnO_2$ . (C) Both $MnO_2$ and KMm	s used in the preparation of $nO_4$ .	chlorine ? (B) Only KMnO <sub>4</sub> . (D) None.	
40.	The yellow colour soluti	on of Na <sub>2</sub> CrO <sub>4</sub> changes to o	range red on passing C	O, gas due to the formation of :
	(A) $CrO_5$	<b>(B)</b> $CrO_3$	(C) $Na_2Cr_2O_7$	( <b>D</b> ) $\operatorname{Na_3CrO_8}$ .
41.	Reaction of potassium cl (A) CuCrO <sub>4</sub> only (C) CuCr <sub>2</sub> O <sub>7</sub> only	nromate and CuSO <sub>4</sub> in aque	ous solution produces : <b>(B)</b> $Cu_{s}[CrO_{4}]_{2}$ only <b>(D)</b> both $CuCrO_{4}$ and	Cu <sub>2</sub> Cr <sub>2</sub> O <sub>7</sub> .
42.	$\operatorname{FeCr}_2O_4 + \operatorname{Na}_2CO_3 + O_2 -$	$\xrightarrow{\text{Fusion}} [X] \xrightarrow{H^+} [Y]$	$\xrightarrow{H^{+}}_{H_2O_2} [Z]$	
	Which of the following	statement is true for the com	pounds [X], [Y] and [Z	]?
	(A) In all three compoun	ds, the chromium is in $+6$ or	xidation state.	
	(B) [Z] is a deep blue–v dioxygen.	violet coloured compound v	which decomposes rapi	dly in aqueous solution into Cr <sup>3+</sup> and
	<ul><li>(C) Saturated solution of</li><li>(D) All of these.</li></ul>	[Y] gives bright orange com	pound, chromic anhydr	ide, with cold and concentrated $H_2SO_4$ .
43.	$KMnO_4$ in excess on trea heating forming (Y). The	tment with concentrated $H_2^{2}$ e (X) and (Y) are respectively	$SO_4$ forms a compound y :	(X) which decomposes explosively on
	(A) $Mn_2O_7$ , $MnO_2$		<b>(B)</b> $\operatorname{Mn_2O_7}$ , $\operatorname{Mn_2O_3}$	
	(C) $MnSO_4$ , $Mn_2O_3$		<b>(D)</b> $\operatorname{Mn}_2O_3$ , $\operatorname{MnO}_2$	
44.	Potassium dichromate in	alkaline solution, with 30%	H <sub>2</sub> O <sub>2</sub> produces :	
	(A) $K_{3}CrO_{8}$	<b>(B)</b> $\operatorname{CrO}(O_2)_2$	$(\mathbf{C}) \operatorname{CrO}_{3} \qquad (\mathbf{D})$	$K_2 CrO_4$ .
45.	<ul> <li>Which of the following s</li> <li>(A) An acidified solution</li> <li>(B) In acidic solution did</li> <li>(C) Ammonium dichrom</li> <li>(D) Potassium dichroma</li> </ul>	statement is wrong ? of $K_2Cr_2O_7$ liberates iodine chromate ions are converted ate on heating undergo exoth te is used as a titrant for Fe <sup>2</sup>	from iodides. to chromate ions. hermic decomposition t <sup>+</sup> .	o give $Cr_2O_3$ .
46.	Oxygen is absorbed by m is known as :	olten Ag, which is evolved or	n cooling and the silver p	particles are scattered ; the phenomenon
	(A) silvering of mirror.	<b>(B)</b> spitting of silver.	(C) frosting of silver.	<b>(D)</b> hairing of silver.
47.	The image on an expose	d and developed photograph	nic film is due to :	
	(A) AgBr	<b>(B)</b> $[Ag(C_2O_3)_2]^{3+}$	(C) Ag	<b>(D)</b> $Ag_2O$ .
48.	Which of the following of	loes not react with AgCl?		
	$(\mathbf{A}) \operatorname{Na}_2 \operatorname{S}_2 \operatorname{O}_3$	(B) NH <sub>4</sub> OH	(C) NaNO <sub>3</sub>	<b>(D)</b> Na <sub>3</sub> AsO <sub>3</sub> .
49.	Addition of iron fillings	to CuSO, solution caused m	recipitation of Cu owing	z to the :
	(A) reduction of $Cu^{2+}$	(B) oxidation of Cu <sup>2+</sup>	(C) reduction of Fe	(D) None of these

50.	Identify the incorre (A) $CuSO_4$ reacts w (B) $CuSO_4$ reacts w (C) $CuSO_4$ reacts w (D) $CuSO_4$ on stron	ct statement. ith KCl in aqueous solution to g ith KI in aqueous solution to g ith NaOH and glucose in aqueo g heating gives CuO.	give $Cu_2Cl_2$ . ive $Cu_2I_2$ . ous medium to give $Cu_2O$ .		
51.	Solution of CuCl in (A) CO <sub>2</sub>	NH <sub>4</sub> OH absorbs : ( <b>B</b> ) SO <sub>2</sub>	(C) $H_2SO_4$	<b>(D)</b> CO.	
52.	Which of the following reactions represents "developing" in photography ? (A) $AgNO_3 + NaBr \rightarrow AgBr + NaNO_3$ . (B) $AgBr + 2Na_2S_2O_3 \rightarrow Na_3[Ag(S_2O_3)_2] + NaBr$ . (C) $AgBr + hv \rightarrow AgBr$ . (D) $C_6H_4(OH)_2 + 2AgBr \rightarrow C_6H_4O_2 + 2HBr + 2Ag$ . (E) $AgBr + 2NH \rightarrow [Ag(NH)]Br$				
53.	Transition metal wite (A) a base	th low oxidation state will act as (B) an acid	s : (C) both (A) and (B)	(D) none of these.	
54.	Boiling CuCl <sub>2</sub> with (A) CuCl	Cu in concentrated HCl gives : (B) CuCl <sub>2</sub>	(C) CuCl <sub>3</sub>	<b>(D)</b> $Cu_2Cl$ .	
55.	CuFeS <sub>2</sub> + $O_2$ (excess <b>Note :</b> Correspondi Which is correct che (A) (X) liberates iod (B) (Y) on heating I (C) (X) forms bluist (D) (Y) on reaction	s) $\xrightarrow{\Delta} X(s) + Y(s)$ ngs sulphates of copper and iro pice for (X) and (Y) ? line from soluble iodides like K iberates SO <sub>3</sub> only h white precipitate with sodium with potassium ferricyanide give	on are formed respectively. I hydroxide which redissolve ves brown precipitate.	es in excess of sodium hydroxide.	
56.	An extremely hot co (A) Cu <sub>2</sub> O	opper wire reacts with steam to $(B)$ CuO <sub>2</sub>	produce : (C) Cu <sub>2</sub> O <sub>2</sub>	<b>(D)</b> CuO.	
57.	A compound of iron acidic solution. The (A) Fe <sub>2</sub> O <sub>4</sub>	exists as a dimer in vapour stat compound is : (B) FeSO	e. It is hygroscopic in nature (C) FeCl,	and dissolves in water giving brown (D) FeCl <sub>2</sub> .	
58.	<ul> <li>Which of the following statements is not correct ?</li> <li>(A) La(OH)<sub>3</sub> is less basic than Lu(OH)<sub>3</sub></li> <li>(B) In lanthanide series ionic radius of Ln<sup>3+</sup> ions decreases</li> <li>(C) La is actually an element of transition series rather than lanthanide series</li> <li>(D) Atomic radii of Zr and Hf are same because of lanthanide contraction</li> </ul>				
59.	Transuranic elemen (A) Np	ts begin with (B) Cm	(C) Pu	<b>(D)</b> U	
60.	Lanthanide contract (A) Gd	tion is observed in : (B) At	( <b>C</b> ) Xe	<b>(D)</b> Ac	
61.	The names transition and inner transition metals are used to indicate the elements of :(A) d-block elements only(C) p- and d-blocks elements respectively(D) d- and f-blocks elements respectively				

62.	The correct statemet (i) all the d and f-ble (ii) all the d and f-b (iii) all the d- and f- (A) (i) only	ent(s) from among the followi ock elements are metals lock elements form coloured block elements form paramag (B) (i) and (ii)	ing is/are : ions gnetic ions (C) (ii) and (iii)	(D) All	
63.	The actinides show	ing +7 oxidation state are :	(C) Nn Pu	(D) Am Cm	
64.	(A) O, NP Which of the follow (A) Curium	ving is not an actinoide ? (B) Californium	(C) uranium	(D) terbium	
65.	Lanthanide contrac (A) shielding by 4f	tion is due to increase in : electrons	(B) atomic number	s	
66.	<ul> <li>(b) size of 41 orbitals</li> <li>(c) effective indicial charge</li> <li>(d) size of 41 orbitals</li> <li>(e) size of 41 orbitals</li> <li>(f) size of 3d series because :</li> <li>(g) size of 2n atom is smaller</li> </ul>				
67.	The less stable oxic (A) $Cr^{2+}$	lation states of Cr are : (B) $Cr^{3+}$	<b>(C)</b> Cr <sup>4+</sup>	<b>(D)</b> Cr <sup>6+</sup>	
68.	Which of the follow (A) Transition elem (B) $IE_2$ of $_{23}V <_{24}Cr$ (C) Ni(II) compour (D) The elements w series.	ving statements are correct? nents exhibit higher enthalpie $>_{25}$ Mn and $_{28}$ Ni $<_{29}$ Cu $>_{30}$ Zn ads are more stable than Pt(II) which gives the greatest numb	s of atomization as they hat where as Pt(IV) compoun- per of oxidation states doe	we stronger interatomic interaction ds are more stable than nickel (IV). s not occur in or near the middle of the	
69.	Which of the follow (A) Ni <sup>2+</sup>	ving ions give(s) colourled a (B) Fe <sup>2+</sup>	queous solution? (C) Cu <sup>2+</sup>	<b>(D)</b> Cu <sup>+</sup>	
70.	<ul> <li>Which of the following statement is/are correct ?</li> <li>(A) Transition metals and their many compounds act as good catalyst.</li> <li>(B) The enthalpies of atomistation of the transition metals are high.</li> <li>(C) The transition metals generally form interstitial compounds with small atoms like C, B, H etc.</li> <li>(D) All transition metal compounds are not paramagnetic.</li> </ul>				
71.	Select correct stater (A) PH <sub>3</sub> reduces Ag (B) Organic tissues	ment (s). NO <sub>3</sub> to metallic Ag. turn AgNO <sub>3</sub> black by reduci	ng it to Ag.		

- (C) AgCN is soluble in KCN.(D) Zr and Ta have almost similar size due to lanthanide contraction.

	Part # II	>>	[Assertion & Reason Type Questions]
	Each questie (A) Statemen (B) Statemen (C) Statemen (D) Statemen (E) Both Sta	on has 5 ch nt-1 is true, nt-1 is true, nt-1 is true, nt-1 is false itements ar	<b>toices (A), (B), (C), (D) and (E) out of which only one is correct.</b> Statement-2 is true and Statement-2 is correct explanation for Statement-1 Statement-2 is true and Statement-2 is not correct explanation for Statement-1, Statement-2 is false b, Statement-2 is true e false
1.	Statement-1 Statement-2	: Mn aton : The effe	n loses ns electrons first during ionisation as compared to $(n - 1)$ d-electrons. Exclive nuclear charge experienced by $(n - 1)$ d electrons is greater than that by ns electron.
2.	Statement-1 Statement-2	: CuSO <sub>4</sub> .5 : All the f	${}^{5}\mathrm{H}_{2}\mathrm{O}$ on heating to 250°C loses all the five $\mathrm{H}_{2}\mathrm{O}$ molecules and becomes anhydrous. ive $\mathrm{H}_{2}\mathrm{O}$ molecules are co-ordinated to the central $\mathrm{Cu}^{2+}$ ion.
3.	Statement-1 Statement-2	: Silver cl : AgCl for	noride dissolves in excess ammonia. rms a soluble complex, $[Ag(NH_3)_2]$ Cl with ammonia.
4.	Statement-1 Statement-2	<ul> <li>The order</li> <li>The atom radii of 4</li> </ul>	er of atomic radii of Cu, Ag and Au is Cu < Ag $\approx$ Au. nic radii of 4d series elements are larger than those of 3d series elements but generally the 4d and 5d series elements are almost identical.
5.	Statement-1 Statement-2	1 :4d & 5d 2 :Lanthar	series elements have nearly same atomic radius.
6. 7.	Statement-1 Statement-2 Statement-1	<ul><li>Tungste</li><li>Tungste</li><li>CrO, is a</li></ul>	n has very high melting point n is a covalent compound an acid anhydride.
	Statement-2	: $CrO_3^3$ is o	obtained as bright orange crystals by the reaction of $K_2 Cr_2 O_7$ with cold concentrated $H_2 SO_4$ .
8.	Statement-1	: Solid po ammoni	tassium dichromate gives greenish yellow vapours with concentrated $H_2SO_4$ and solid um chloride.
	Statement-2	2 : The read chloride	ction of ammonium chloride with solid $K_2 Cr_2 O_7$ and concentrated $H_2 SO_4$ produces chromyl .
9.	Statement-1 Statement-2	: Permang : Hydroch	anate titrations is not carried out in presence of hydrochloric acid.
10.	Statement-1 Statement-2	: Copper	metal is turned green when exposed to atmospheric $CO_2$ and moisture. gets covered with a green layer of basic copper carbonate.
11.	Statement-1 Statement-2	: Ammon : Glucose	iacal silver nitrate converts glucose to gluconic acid and metallic silver is precipitated. acts as a weak reducing agent.
12.	Statement-1 Statement-2	: 4d & 5d : Lanthar	series elements have nearly same atomic radius.
13.	Statement-1 Statement-2	: The free : Half fille	gaseous chromium atom has six unpaired electrons. ed orbital has greater stability than fully filled.
14.	Statement-1 Statement-2	: $K_2 CrO_4$ : $CrO_4^{2-}$ io	has yellow colour due to charge transfer. on is tetrahedral in shape.

- **15. Statement-1 :** The green manganate is paramagnetic but the purple permanganate is diamagnetic in nature. **Statement-2 :**  $MnO_4^{2-}$  contains one unpaired electron while in  $MnO_4^{-}$  all electrons are paired.
- 16. Statement-1 : The highest manganese fluoride is  $MnF_4$  and the highest oxide is  $Mn_2O_7$ . Statement-2 : In  $Mn_2O_7$ , each Mn is tetrahedrally surrounded by O's including Mn–O–Mn bridge.
- 17. Statement-1: The lowest oxide of a transition metal (say, chromium, atomic number 24) is basic where as the highest oxide is usually acidic.

**Statement-2** :  $Cr_2O_3$  is amphoteric in nature.

- Statement-1: In acid solution permanganate is reduced to Mn<sup>2+</sup> by an excess of reducing agent.
   Statement-2: MnO<sub>4</sub><sup>-</sup> redused in Mn<sup>2+</sup> in acidic medium and the product in the presence of an excess of permanganate is MnO<sub>2</sub>.
- **19. Statement-1 :** The number of unpaired electrons in the following gaseous ions ;  $Mn^{3+}$ ,  $Cr^{3+}$ ,  $V^{3+}$  and  $Ti^{3+}$  are 4, 3, 2 and 1 respectively.

Statement-2: Cr<sup>3+</sup> is most stable in aqueous solution amongst these ions .

- Statement-1: The value of enthalpy of atomisation is maximum at about the middle of each series.
   Statement-2: There is one unpaired electron per d-orbital and this results in stronger interaction.
- **21. Statement-1** : The spin only magnetic moment of  $Sc^{3+}$  is 1.73 B.M.
  - Statement-2: The spin only magnetic moment of an ion is equal to  $\sqrt{n(n+2)}$ ; where n is the number of unpaired electrons in the ion.
- 22. Statement-1: Hydrochloric acid is not used to acidify a  $KMnO_4$  solution in volumetric analysis of Fe<sup>2+</sup> and C<sub>2</sub>O<sub>4</sub><sup>2-</sup> because.

Statement-2: Part of the oxygen produced from KMnO<sub>4</sub> and HCl is used up in oxidising HCl to Cl<sub>2</sub>.

- Statement-1: Potassium dichromates gives deep red vapours with concentrated H<sub>2</sub>SO<sub>4</sub> and sodium chloride.
   Statement-2: The reaction of sodium chloride with solid K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> and concentrated H<sub>2</sub>SO<sub>4</sub> produces chromyl chloride.
- 24. Statement-1 : Solution of  $Na_2CrO_4$  in water is intensely coloured. Statement-2 : Oxidation state of Cr in  $Na_2CrO_4$  is (+VI).
- 25. Statement-1: Silver nitrate is reduced to silver by the hydrides of 15th group elements (except NH<sub>3</sub>) because Statement-2: They act as strong reducing agents.
- 26. Statement-1:  $Ag_2 S + 4 KCN \xrightarrow{O_2} 2K [Ag(CN)_2] + K_2 S$ Statement-2: The reaction is carried out in presence of air or  $O_2$  so that  $K_2S$  is oxidised to  $K_2SO_4$  thereby shifting the equilibrium in forward direction.
- Statement-1: Reaction of thionyl chloride with hydrated ferric chloride yields anhydrous ferric chloride.
   Statement-2: Water of crystallisation present with ferric chloride reacts with thionyl chloride to liberate HCl and SO<sub>2</sub> gases.
- Statement-1: Hydroquinone is used as a developer for developing black and white photographic film.
   Statement-2: Hydroquinone reduces silver bromide to black silver particles and an inverted image of the object is produced on a celluloid film.

	Exercise # 3	Part # I [Matrix Match Type Questions]				
1.	Match the reactions listed Column (II).	in Column (I) with the characteristic(s) of the products/type of reactions listed in				
	Column - I (A) $MnO_4^{2-}+CO_2 \longrightarrow$ (B) $CrO_4^{2-}+H^+ \longrightarrow$	<ul> <li>Column - II</li> <li>(p) Two pungent smelling gases are liberated.</li> <li>(q) Show disproportionation reaction.</li> </ul>				
	(C) FeSO <sub>4</sub> $\xrightarrow{\Delta}$	(r) Dimeric bridged tetrahedral metal ion.				
	<b>(D)</b> CuCl <sub>2</sub> 2H <sub>2</sub> O $\xrightarrow{\Delta}$	(s) One of the products has central metal in its highest stable oxidation state.				
2.	Match the salts/mixtures list Column - I	ed in Column (I) with their respective name listed in Column (II). Column - II				
	(A) $ZnS + BaSO_4$ mixture	(p) Lunar caustic				
	(B) $FeSO_4$ (NH <sub>4</sub> ), SO <sub>4</sub> . 6H,O	(q) Schwitzer's regent.				
	(C) AgNO <sub>3</sub>	(r) Lithopone				
	<b>(D)</b> $[Cu(NH_3)_4]SO_4$	(s) Mohr's salt				
3.	Match the pairs of complex listed in Column (II).	es/compounds listed in Column (I) with the characteristic(s) of the reaction products				
	Column - I	Column - II				
	(A) Cu(I) and Zn (II) comple	(p) Pair of compounds having similar colour and some magnetic moment but equal.				
	( <b>B</b> ) KMnO <sub>4</sub> and $K_2Cr_2O_7$	(q) Pair of compounds which are diamagnetic but coloured.				
	(C) $Cu_2O$ and $HgI_2$	(r) Pair of compounds having metals in the highest stable oxidation states.				
	<b>(D)</b> $VOCI_2$ and $CuCI_2$	(s) Pair of compounds which show diamagnetism and are colourless.				
4.	Column-I (Metals)	Column-II (Ores)				
	(A) Zn	(p) Cyanide process				
	<b>(B)</b> Cu	(q) hydrometallurgical process				
	(C) Ag	(r) roasting				
	( <b>D</b> ) Pt	(s) brass.				
5.	Column-I (Alloys)	Column-II (Constituents)				
	(A) $\operatorname{TiCl}_4$	(p) Adams catalyst in reduction				
	<b>(B)</b> $PdCl_2$	(q) In preparation of $(CH_3)_2$ SiCl <sub>2</sub>				
	(C) Pt/PtO	(r) Used as the Natta catalyst in polythene production				
	(D) Cu	(s) Wake process for converting $C_2H_4$ to $CH_3CHO$				
6.	Match the reactions in Colu	mn I with the types of products / the use of products in Column II.				
	Column - I	Column - II				
	(A) HO-OH + Age	$Br(s) \rightarrow$ (p) Turnbull's blue pigment				
	<b>(B)</b> $\operatorname{BaCl}_2 + \operatorname{K}_2\operatorname{Cr}_2\operatorname{O}_7 + \operatorname{H}_2\operatorname{SO}_7$	$_4 \rightarrow$ (q) Schwitzer's reagent				
	$(C) \operatorname{FeSO}_4 + \operatorname{K}_3[\operatorname{Fe}(CN)_6] \rightarrow$	(r) Rinmann's green pigment				
( <b>D</b> ) $Cu(OH)_2 + NH_4OH + (NH_4)_2SO_4 \rightarrow$ (s) Chromyl chloride test						

- (s) Chromyl chloride test
- (E)  $\operatorname{ZnO} + \operatorname{Co(NO_3)_2} \xrightarrow{\Delta}$
- (t) Photography

7. Match the reactions in Column I with nature of the reactions/type of the products in Column II.

**Column I** Column II (A)  $\operatorname{FeSO}_4 \xrightarrow{\Delta}$ (p) One of the products is coloured due to charge transfer (B)  $Mn^{2+} + S_2O_8^{2-} + H_2O \longrightarrow$ (q) One of the products is in + VI oxidation state (r) Redox reaction (C) Na<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> + H<sub>2</sub>SO<sub>4</sub>(conc.)  $\rightarrow$ (saturated solution) (D)  $N_2H_4 + CuSO_4 \longrightarrow$ (s) One of the products is acidic oxide Match the properties given in Column I with the transition elements given in Column II. Column I **Column II** (Property) (Transition elements) (A) Highest oxidation state. **(p)**Cr. (B) Highest density. (q) Os. (C) Element with maximum unpaired electrons. (r) Tc. (D) Radioactive transition element. (s) Ru. Match the transition element ions given in Column I with the characteristic(s) of products given in Column II.

Column I	Column II
(A) $Cu^{2+}$ .	(p) Form amphoteric oxide.
<b>(B)</b> $Zn^{2+}$ .	(q) Diamagnetic and colourless compounds.
(C) $Cr^{3+}$ .	(r)) Coloured hydrated transition metal ion.
<b>(D)</b> $Ni^{2+}$ .	(s) Paramagnetic.

8.

9.

[Comprehension Type Questions]

### **Comprehension #1**

(i) A powdered substance (A) on treatment with fusion mixture gives a green coloured compound (B).

(ii) The solution of (B) in boiling water on acidification with dilute  $H_2SO_4$  gives a pink coloured compound (C) and brown colour compound (D).

(iii) The aqueous solution of (A) on treatment with NaOH and  $Br_2$ -water gives the compound (D).

(iv) A solution of (D) in concentrated  $HNO_3$  on treatment with lead peroxide at boiling temperature produced a compound (E) which was of the same colour at that of (C).

(v) A solution of (A) on treatment with a solution of barium chloride gave a white precipitate of compound (F) which was insoluble in concentrated HNO<sub>3</sub> and concentrated HCl.

- 1. Which of the following is true for compound (C)?
  - (A) It oxidises ammonia to nitrogen dioxide in neutral medium.
  - (B) It's pink colour is due to d-d transition.
  - (C) It can be oxidised by ozone.

(D) It is obtained by alkaline fusion of pyrolusite followed by electrolytic oxidation.

2. The oxidation state of central metal ions of (A), (B) and (C) compounds are respectively : (A)  $\downarrow$  H  $\downarrow$  VI and  $\downarrow$  VII

(A) $+11$ , $+V1$ and $+V11$	$(\mathbf{B})$ +II, + VI and + VI
$(\mathbf{C})$ +II, + VII and + VII	(D) +VI, +VII and +VII

- **3.** Consider the following statements ;
  - (I) anions of both (B) and (C) are diamagnetic and have tetrahedral geometry.
  - (II) anions of both (B) and (C) are paramagnetic and have tetrahedral geometry.
  - (III) anion of (B) is paramagnetic and that of (C) is diamagnetic but both have tetrahedral geometry.
  - (IV) green coloured compound (B) in a neutral or acidic medium disproportionates to give (C) and (D). of these select the correct one from the codes given :
  - (A) I and III only (B) II and III only (C) II and IV only (D) III and IV only

### **Comprehension #2**

Paramagnetism is a property due to the presence of unpaired electrons. In case of transition metals, as they contain unpaired electrons in the (n-1) d orbitals, most of the transition metal ions and their compounds are paramagnetic. Paramagnetism increases with increases in number of unpaired electrons. Magnetic moment is calculated from 'Spin only formula' viz.

 $\mu = \sqrt{n(n+2)}$  B.M n = number of unpaired electrons

Similarly the colour of the compounds of transition metals may be attributed to the presence of incomplete (n-1)d sub-shell. When an electron from a lower energy of d-orbital is excited to a higher energy d-orbital, the energy of excitation corresponds to the frequency of light absorbed. This frequency generally lies in the visible region. The colour observed corresponds to complementary colour of the light absorbed. The frequency of the light absorbed is determined by the nature of the ligand.

- 1. Which of the following pair of compounds is expected to exhibit same colour in aqueous solution. (A) FeCl<sub>2</sub>, CuCl<sub>2</sub> (B) VOCl<sub>2</sub>, CuCl<sub>2</sub> (C) VOCl<sub>2</sub>, FeCl<sub>2</sub> (D) FeCl<sub>2</sub>, MnCl<sub>2</sub>
- Titanium shows magnetic moment of 1.73 BM in its compound. What is the oxidation state of titanium in the compound ?
   (A)+2
   (B)+1
   (C)+3
   (D)+4
- 3. The colourless species is : (A) VCl<sub>3</sub> (B) VOSO<sub>4</sub> (C) Na<sub>3</sub>VO<sub>4</sub> (D)  $[V(H_2O)_5SO_4]2H_2O$
- 4. Identify the incorrect statement.
  - (A) Mn<sup>2+</sup> has the highest paramagnetism amongst the bivalent cations of the I<sup>st</sup> transition series.
  - (B) The coloured ions or compounds of transition elements are due to d-d transition and charge transfer spectrum.
  - (C) In 3 d series the paramagnetic character first increase to maximum & then starts decreasing.
  - (D) None of these

## **Comprehension #3**

Transition metals usually form coloured complexes and d - d transitions ( $t_{2g} \leftrightarrow e_g$ ) are responsible for colour as the energy difference between  $t_{2g}$  and  $e_g$  lies in visible region. But all the coloured ions are not due to d-d transition but charge transfer bands also play important roles. Charge transfer bands may be of two types. (i) ligand to metal (CTLM) (ii) metal to ligand (CTML).

Charge transfer transition always produces intense colour as compared to d-d transition.

**1.** Select the incorrect statement :

(A) d-block metal ions are usually coloured.

(B) Colour of the most of d-block metal ions is generally due to d-d transition.

(C) All the complexes of Cu<sup>+</sup> are colourless on account of diamagnetic nature i.e. d<sup>10</sup> configuration.
 (D) CrO, is bright orange due to CTLM.

2.  $MnO_4^-$  is dark purple coloured although Mn is in (+ VII) oxidation state with 3d° configuration :

(A) due to d-d transition.

(B) due to CTML spectra.

(C) due to CTLM spectra

(D) none of these.

- **3.** Select the correct statement.
  - (A) In CTML, no net reduction-oxidation takes place because of the short life time of excited state.
  - **(B)** Cu<sub>2</sub>O is a red coloured salt.
  - (C) Vermilon (HgS) is a red coloured compound.
  - (D) All of these.

### **Comprehension #4**

Read the following graph and gives the answer for the following questions.



#### Trends in enthalpies of atomisation of transition elements

- The dip in the boiling point of manganese in 3d series is due to :

   (A) less delocalisation of valence electrons owing to extra stable half filled configuration.
   (B) smaller size of atom.
   (C) less electronegativity of atom.
  - (D) all of these
- 2. Consider the following statements :
  - (I) The transition elements (except the 12<sup>th</sup> group elements) are very much hard and have low volatility.
  - (II) Generally in a given series the melting points of the transition elements rises to a maximum and then fall as the atomic number increases.
  - (III) Transition metals have high thermal and electrical conductivity and metallic luster.

Select the correct from the codes given.

(A) I and III only (B) II and III only (C) All of these (D) None of these

### Comprehension # 5

Pyrolusite ore on oxidation with  $\text{KClO}_3/\text{KNO}_3$  in basic medium produces dark green coloured compound (A), which on electrolysis produces a purple coloured compound (B). The purple coloured compound can be crystallised to deep purple rhombic prisms. It shows different reactions in different mediums. Excess of compound (B) on heating with concentrated H<sub>2</sub>SO<sub>4</sub> gives an explosive oil (C), which on heating

(C) neutral

(D) amphoteric

- decomposes to gives another compound (D) along with oxygen.1. The nature of compound (C) is :
- 2. On passing CO<sub>2</sub> in the solution of (A), a purple coloured solution along with brown precipitate is observed. Select the correct statement.
  - (A) This is a disproportionation reaction where oxidation number changes from +6 to +7, +4.
  - (B) This is a comproportionation reaction where oxidation number changes from +3 to +2.

**(B)** acidic

- (C) both (A) and (B)
- (D) none

(A) basic

- 3. Identify (D) . (A) Mn<sub>2</sub>O<sub>2</sub> (B) M
  - (A) Mn<sub>2</sub>O<sub>7</sub>
    (B) MnO<sub>2</sub>
    (C) MnSO<sub>4</sub>
    (D) Mn<sub>2</sub>O<sub>3</sub>
    (D) Mn<sub>2</sub>O<sub>3</sub>
    (D) Mn<sub>2</sub>O<sub>3</sub>
- 4. When (D) reacts with PbO<sub>2</sub>/HNO<sub>3</sub> it produces a purple/pink coloured compound (E). Identify (E)
   (A) NaMnO<sub>4</sub> (B) Mn(OH)<sub>2</sub> (C) HMnO<sub>4</sub> (D) Na<sub>2</sub>MnO<sub>4</sub>
- 5. Red coloured oxide (M) whose cation has pseudo inert electronic configuration reacts with compound (B) in acidic medium gives another oxide (N) which is black in colour. When sulphate of its metal is reacted with  $K_4[Fe(CN)_6]$  it produces brown precipitate. Select the correct statement.
  - (A) The brown precipitate formed is  $Cu_2[Fe(CN)_6]$
  - **(B)** The red coloured oxide is  $Cu_2O$
  - (C) The black coloured oxide is CuO

**(D)** All of these

### **Comprehension #6**

Photography is based on the nature of silver halides. Except AgF, the silver halides are photosensitive. These undergo decomposition in light and turn black due to formation of free silver.

 $2AgBr \longrightarrow 2Ag + Br_2$ 

The photographic plates or films are prepared in a dark and dust free room by applying a light sensitive mixture called emulsion which is prepared by adding 20 per cent aqueous solution of silver nitrate to ammonium bromide solution containing gelatin.

When such a film is exposed, the emulsion gets affected. Since different parts of the body reflect light of varied intensity, the film or plate is affected proportionately. An actual but inverted image of the object is thus formed on the film or plate which is not visible to the eye. It is, therefore, called the latent image.

When this exposed film or plate is dipped in a developer which contains a reducing agent, the parts affected most during exposure are reduced to maximum. The image becomes visible. It is called a negative. The remaining sensitive emulsion on the negative is removed by dissolving it in hypo solution (fixer). Finally, a positive of the negative already prepared is made on silver bromide paper.

**1.** The compound formed on the unexposed photographic film or plate is :

(A) silver nitrate.	(B) ammonium bromide.
(C) diamminesilver bromide.	<b>(D)</b> silver bromide.

The exposed part of the film or plate after developing contains :
 (A) silver metal
 (B) silver oxide
 (C) silver bromide(D) silver nitrate.

3.	The solution of the developer consists :	
	(A) alkaline solution of pyrogallol	(B) alkaline solution of quinol
	(C) either (A) or (B)	(D) neither (A) nor (B).

- 4. Silver halides are used in photography because these compounds :
  - (A) are insoluble in water
  - (B) are affected by light
  - (C) are soluble in ammonia solution
  - **(D)** easily stick on the surface of photographic plate or film
- 5. Silver bromide dissolves in hypo solution forming : (A)  $Ag_2S_2O_3$  (B)  $Ag_2S$  (C)  $Na_3[Ag(S_2O_3)_2]$  (D)  $NaAgS_2O_3$

	Exercise # 4 [Subjective Type Questions]
1	Why are $Mn^{2+}$ compounds more stable than $Fe^{2+}$ towards oxidation to their +3 state?
2	How is the variability in oxidation states of transition metals different from that of the non transition Illustrate with examples.

metals?

- 3. Which metal in the first series of transition metals exhibits +1 oxidation state most frequently and why?
- 4. Between Na<sup>+</sup> and Ag<sup>+</sup>, which is stronger Lewis acid and why?
- 5. Why the highest oxidation state of a metal is exhibited in its oxide or fluoride only?
- 6. What may be the stable oxidation state of the transition element with the following d electron configurations in the ground state of their atoms:  $3d^3$ ,  $3d^5$ ,  $3d^8$  and  $3d^4$ ?
- 7. The  $E^{\theta}$  (M<sup>2+</sup>/M) value for copper is positive (+0.34V). What is possibly the Statement-2 for this?
- 8. Why is the  $E^{\theta}$  value for the  $Mn^{3+}/Mn^{2+}$  couple much more positive than that for  $Cr^{3+}/Cr^{2+}$  or  $Fe^{3+}/Fe^{2+}$ ? Explain.
- 9. Which is a stronger reducing agent  $Cr^{2+}$  or  $Fe^{2+}$  and why?
- **10.** Copper dissolves in dilute nitric acid but not in dilute HCl. Why?
- 11. Blue colour of  $CuSO_4$  solution is discharged slowly when an iron rod is dipped into it. Why?
- **12.** Copper (I) salts are not known in aqueous solutions.
- **13.** Explain why  $Cu^+$  ion is not stable in aqueous solutions?
- 14. The aqueous solution of FeCl<sub>3</sub> is acidic. Why?
- **15.** Ferric iodide is very unstable but ferric chloride is not.
- 16. Calculate the number of unpaired electrons in the following gaseous ions:  $Mn^{3+}$ ,  $Cr^{3+}$ ,  $V^{3+}$  and  $Ti^{3+}$ . Which one of these is the most stable in aqueous solution?
- 17. Mercurous ion is written as  $Hg_2^{2+}$  whereas cuprous ion is written as  $Cu^+$ . Explain.
- **18.** Copper sulphate dissolves in  $NH_4OH$  solution but  $FeSO_4$  does not.
- A hydrated metallic salt (A), light green in colour, on careful heating gives a white anhydrous residue (B), (B) is soluble in water and its aqueous solution gives a dark blue precipitate (C) with potassium hexacyanidoferrate(III). (B) on strong heating gives a brown residue (D) and a mixture of two gases (E) and (F). The gaseous mixture when passed through acidified potassium dichromate, produces green colour solution and when passed through lead acetate solution gave a white precipitate. Out of two gases (E) can act as both reducing as well as oxidising agent.
  - (a) Identify (A), (B), (C), (D), (E) and (F) and give the reactions involved.
  - (b) Explain why salt (A) becomes white on heating.
- 20. A white substance (A) reacts with dilute  $H_2SO_4$  to produce a colourless suffocating gas (B) and a colourless solution (C). The reaction of gas (B) with potassium iodate and starch solution produces a blue colour solution. Aqueous solution of (A) gives a white precipitate with BaCl<sub>2</sub> solution which is soluble in dilute HCl. Addition of aqueous NH<sub>3</sub> or NaOH to (C) produces first a precipitate which dissolves in excess of the respective reagent to produce a clear solution. Similarly addition of excess of potassium ferrocyanide to (C) produces a precipitate (D) which also dissolves in aqueous NaOH giving a clear solution. Identify (A), (B), (C) and (D). Write the equations of the reactions involved.

- 21. Write the balanced chemical equation for the following reaction :-Nitrogen is obtained in the reaction of aqueous ammonia with potassium permanganate
- 22. An aqueous solution of inorganic compound (X) gives following reactions.
  - (i) With an aqueous solution of barium chloride a precipitate insoluble in dilute HCl is obtained.
  - (ii) Addition of excess of KI gives a brown precipitate which turns white on addition of excess of hyposolution.
  - (iii) With an aqueous solution of  $K_4$ [Fe(CN)<sub>6</sub>], a brown coloured precipitate is obtained.

Identify (X) and give equations for the reaction for (i), (ii) and (iii) observations.

- 23.  $H_2S$  gas is passed through an acidic solution of  $K_2Cr_2O_7$ . The solution turns milky, why?
- 24. What happens when (i) a small amount of  $KMnO_4$  is added to concentrated  $H_2SO_4$  (ii) an excess amount of  $KMnO_4$  is added to concentrated  $H_2SO_4$  solution.
- 25. Complete and / balance the following equation :  $Ag_2S + CuCl_2 + Hg$  .....+ S + 2Ag
- 26. Which out of the two,  $La(OH)_3$  and  $Lu(OH)_3$ , is more basic and why?
- 27. One among the lanthanoides, Ce(III), can be easily oxidized to Ce(IV) (At. No. of Ce = 58). Explain why?
- 28. Transition metals forms a large number of interstitial compounds. Explain.
- **29.** Explain the large difference in melting point of Cr(1920°C) and Zn(420°C).
- 30. Although  $Cu^+$  does not exist in solution state, but  $CuCl_{(s)}$  is formed in presence of  $Cl^-$  ions in aqueous solution of  $Cu_{(s)}$  and  $Cu^{2+}$ .
- 31. Atomic radius in a transition series does not vary much while that of s and p-block does. Why ?
- 32. The paramagnetic character in 3d transition series increases upto Cr and then decreases. Explain.
- **33.** The ionisation energies of first five members of 3d-series increase with increase in atomic number and then become constant or irregular for next five members. Explain.
- 34. Pyrolusite on heating with KOH in the presence of air gives a dark green compound (A). The solution of (A) on treatment with H<sub>2</sub>SO<sub>4</sub> gives a purple coloured compound (B), which gives following reactions :
  - (i) KI on reaction with alkaline solution of (B) changes into a compound (C).
  - (ii) The colour of the compound (B) disappears on treatment with the acidic solution of FeSO<sub>4</sub>.

(iii) With concentrated H<sub>2</sub>SO<sub>4</sub> compound (B) gives (D) which can decompose to yield (E) and oxygen on heating.

Identify (A) to (E) and write balanced chemical equations for the formation of (A) and (B) and for the steps (i) to (iii).

- **35.** A black compound (A) in solid state is fused with KOH and KClO<sub>3</sub>. The aqueous extract of fused mass is green colour solution (B). On passing CO<sub>2</sub> gas through it pink colour of (C) is noticed along with some black insoluble mass of (A). The pink coloured solution is decolourised by  $Fe^{2+}$  in acidic medium. What are (A) (B) and (C).
- 36. (i) An ore (A) on roasting with sodium carbonate and lime in the presence of air gives two compounds, (B) and (C).
  - (ii) The solution of (B) in conc. HCl on treatment with potassium ferrocyanide gives a blue colour or precipitate of compound (D).
  - (iii) The aqueous solution of (C) on treatment with conc.  $H_2SO_4$  gives a yellow coloured compound (E).
  - (iv) Compound (E) when treated with KCl gives an orange-red compound (F) which is used as an oxidising agent.
  - (v) The solution of (F) on treatment with oxalic acid and then with an excess of potassium oxalate gives blue/violet crystals of compound (G).

Identify (A) to (G) and give balanced chemical equations for reactions at steps (i) to (v).

- 37. Compound (A) is a light green crystalline solid. It gives the following tests.
  - (i) Compound (A) is heated strongly. Gases (B) and (C) with pungent smell come out. A brown residue (D) is left behind.
  - (ii) It dissolves in dilute sulphuric acid. No gas is produced.
  - (iii) A drop of  $KMnO_4$  is added to the above solution. The pink colour disappears.
  - (iv) The gas mixture (B and C) is passed into a dichromate solution. The solution turns green.
  - (v) The green solution from step (iv) gives a white precipitate (E) with a solution of barium nitrate.
  - (vi) Residue (D) from step (i) is heated on charcoal in a reducing flame. It gives a magnetic substance. Name the compounds (A), (B), (C), (D) and (E).
- 38. Cuprous chloride is insoluble in water and dilute HCl but dissolves in concentrated HCl.
- **39.** (i) A blue coloured compound (A) on heating gives two products, (B) and (C).
  - (ii) A metal (D) is deposited on passing hydrogen through heated (B).
  - (iii) The solution of (B) in HCl on treatment with  $K_4[Fe(CN)_6]$  gives a chocolate brown coloured precipitate of compound (E).

(iv) (C) turns lime water milky which disappears on continuous passage of (C) forming a compound (F). Identify (A) to (F) and give chemical equations for the reactions at steps (i) to (iv).

- 40. The  $Cr_2O_7^{2-}$  ion is a powerful oxidant in acidic medium but is weak oxidant in basic medium.
- 41. Anhydrous FeCl<sub>3</sub> cannot be obtained by heating hydrated ferric chloride. Why ?
- **42.**  $FeCl_3(aq.)$  gives  $CO_2$  with NaHCO\_3(aq.). Explain.
- **43.** Why is  $AgNO_3$  also called lunar caustic ?
- 44. Hydrated zinc chloride cannot be dehydrated on heating.
- 45. Compete the following by identifying (A) to (H)

(i)  $\operatorname{CuSO}_4:\operatorname{5H}_2O \xrightarrow{100^\circ C}$  (A)  $\xrightarrow{230^\circ C}$  (B)  $\xrightarrow{800^\circ C}$  (C) + (D).

(ii) AgNO<sub>3</sub>  $\longrightarrow$  (E) + (F) + O<sub>2</sub>.

**46.** Why is AgNO<sub>3</sub> kept in brown coloured bottles ?

I	Exercise # 5	Part # I [	Previous Year Ques	tions] [AIEEE/JEE-N	MAIN]
1.	Cerium (Z = 58) is an incorrect? (1) The common oxida (2) The +3 oxidation s (3) The +4 oxidation s (4) Cerium (IV) acts as	n important member of ation state of cerium are tate of cerium is more st tate of cerium is not know an oxidizing agent.	the lanthanoide. Which +3 and +4. able than +4 oxidation s own in solution.	of the following statement	nt about cerium is [AIEEE 2004]
2.	The lanthanide contra (1) Zr and Y have abo (3) Zr and Hf have ab Which of the followin	ction is responsible for ut the same radius out the same radius	the fact that (2) Zr and Nb h (4) Zr and Zn h	ave similar oxidation state ave same oxidation state.	[ <b>AIEEE 2005]</b>
5.	<ol> <li>Greater shielding of</li> <li>Poorer shielding of</li> <li>Effective shielding</li> <li>Poor shielding of</li> </ol>	of 5d electrons by 4f elect f 5d electron by 4f elect g of one of 4f electrons by one of 4f electron by and	ctrons cons by another in the sub-she other in the sub-shell.		[AIEEE 2005]
4.	The "spin-only" magnumber of $Ni = 28$ ) (1) 2.84	netic moment [in units o (2) 4.90	f Bohr magneton, $(\mu_B)$ o (3) 0	f Ni <sup>2+</sup> in aqueous solution (4) 1.73	would be (atomic [AIEEE 2006]
5.	Lanthanoid contractio (1) the appreciable shi (2) the appreciable shi (3) the same effective (4) the imperfect shiel	n is caused due to : elding on outer electron elding on outer electror nuclear charge from Ce ding on outer electrons	ns by 4 <i>f</i> electrons from t ns by 5 <i>f</i> electrons from t to Lu by 4f electrons from the	he nuclear charge he nuclear charge nuclear charge	[AIEEE 2006]
6.	Identify the incorrect (1) The chemistry of v (2) 4f and 5f orbitals a (3) d-block elements s (4) La and Lu have par	statement among the fol arious lanthanoids is ve re equally shielded. how irregular and erration tially filled d orbitals an	lowing. ry similar. c chemical properties an d no other partially filled	nong themselves. d orbitals.	[AIEEE 2007]
7.	The actinoids exhibit (1) The actinoids are a (2) The 5f orbitals ext (3) The 5f orbitals are (4) There is a similarit	more number of oxidation nore reactive than the latend farther from the nuc more buried than the 4f y between 4f and 5f orb	on states in general than anthanoids. leus than the 4f orbitals. orbitals itals in their angular part	the lanthanoids. This is b t of the wave function	ecause [AIEEE 2007]
8.	Larger number of oxi being. (1) lesser energy diffe (2) more energy differ (3) more reactive natu	dation states are exhibit rence between 5f and 6c ence between 5f and 6d re of the actinoids than	ted by the actinoids that I than between 4f and 5c than between 4f and 5d the lanthanoids	n those by the lanthanoids I orbitals orbitals	s, the main reason [AIEEE 2008]

(4) 4f orbitals more diffused than the 5f orbitals

9.	In context with the tra (1) In the highest oxid (2) In the highest oxid for bonding	ansition elements, which of the dation states, the transition mo- lation states of the first five tra	ne following statements is etal show basic character ansition elements (Sc to N	incorrect ? and form cationic com In), all the 4s and 3d ele	[AIEEE 2009] nplexes. ectrons are used
	<ul> <li>(3) Once the d<sup>5</sup> config</li> <li>(4) In addition to the r</li> </ul>	guration is exceeded, the tend normal oxidation states, the ze	lency to involve all the 3 ro oxidation state is also s	d electrons in bonding shown by these element	decreases. ts in complexes.
10.	Knowing that the Ch statement is incorrect (1) The ionic sizes of (2) Ln (III) compound (3) Ln (III) hydroxide (4) Because of the lar	emistry of lanthanoids (Ln) i ? Ln (III) decrease in general w ds are generally colourless. s are mainly basic in character ge size of the Ln (III) ions the	is dominated by its +3 or with increasing atomic num r e bonding in its compound	xidation state, which on the constant of the c	of the following [AIEEE 2009] nic in character.
11.	The correct order of	$E^{\circ}_{M^{2^+}/M}$ values with negative :	sign for the four successi	ve elements Cr, Mn, Fe	e and Co is
12.	<ul> <li>[AIEEE 2010]</li> <li>(1) Mn &gt; Cr &gt; Fe &gt; Co</li> <li>(2) Cr &gt; Fe &gt; Mn &gt; Co</li> <li>(3) Fe &gt; Mn &gt; Cr &gt; Co</li> <li>(4) Cr &gt; Mn &gt; Fe &gt; Co</li> <li>(5) In context of the lanthanoids, which of the following statement is not correct?</li> <li>(1) There is a gradual decrease in the radii of the members with increasing atomic number in the series.</li> <li>(2) All the member exhibit +3 oxidation state.</li> <li>(3) Because of similar properties the separation of lanthanoids is not easy.</li> <li>(4) Availability of 4f electrons results in the formation of compounds in +4 state for all the members of the series.</li> </ul>				
13.	The outer electron cor (1) $4f^3 5d^5 6s^2$	nfiguration of Lu (Atomic No (2) $4f^8 5d^0 6s^2$	: 71) is : (3) $4f^4 5d^4 6s^2$	(4) $4f^{14}5d^16s^2$	[AIEEE 2011]
14.	Iron exhibits +2 and + 3 oxidation states. Which of the following statements about iron is incorrect ?(1) Ferrous oxide is more basic in nature than the ferric oxide.[AIEEE 2012](2) Ferrous compounds are relatively more ionic than the corresponding ferric compounds[AIEEE 2012](3) Ferrous compounds are less volatile than the corresponding ferric compounds[AIEEE 2012](4) Ferrous compounds are more easily hydrolysed than the corresponding ferric compounds.[AIEEE 2012]				
15.	Which of the following arrangements does not represent the correct order of the property stated against it ? (1) $V^{2+} < Cr^{2+} < Mn^{2+} < Fe^{2+}$ : paramagnetic behaviour [JEE(Mains) 2013] (2) $Ni^{2+} < Co^{2+} < Fe^{2+} < Mn^{2+}$ : ionic size (3) $Co^{3+} < Fe^{3+} < Cr^{3+} < Sc^{3+}$ : stability in aqueous solution (4) $Sc < Ti < Cr < Mn$ : number of oxidation states				
16.	Four successive members of the first row transition elements are listed below with atomic numbers. Which one of				
	them is expected to h	ave the highest $E_{M^{3+}/M^{2+}}^{0}$ val	lue ?	[JEF	E(Mains) 2013]
	(1) $Cr(Z=24)$	(2) $Mn(Z=25)$	(3) $Fe(Z=26)$	(4) $Co(Z=27)$	

	Part # I		[Previous Year Q	uestions][IIT-JEE A	ADVANCED]	
1.	The proc	luct of oxidation	$1 \text{ of } I^- \text{ with } MnO_4^- \text{ in alk}$	caline medium is :		[ <b>JEE 2004</b> ]
2.	(A) $IO_3^-$ The pair (A) MnC	of compounds $D_2$ , FeCl <sub>3</sub>	<b>(B)</b> $I_2$ having metals in their h	(C) IO <sup>-</sup> nighest oxidation state is (B) [MnO <sub>4</sub> ] <sup>-</sup> , CrO	( <b>D</b> ) $IO_4^-$ : $D_2Cl_2$	[ <b>JEE 2004</b> ]
3.	(C) [Fe(C) Which o (A) FeCl	$(CN)_6]^{2-}$ , $[Co(CN)_6]^{2-}$ , $[Co(CN)_3]_3$ , $CuCl_2$	$[b_6]^{3-}$ pair of compounds is ex (B) VOCl <sub>2</sub> , CuCl <sub>2</sub>	( <b>D</b> ) [NiCl <sub>4</sub> ] <sup>2–</sup> , [Ni xpected to exhibit same of ( <b>C</b> ) VOCl <sub>2</sub> , FeCl <sub>2</sub>	(CO) <sub>4</sub> ]. colour in aqueous solution ? ( <b>D</b> ) FeCl <sub>2</sub> , MnCl <sub>2</sub>	[JEE 2005]
4.	Give equ thiosulpl process.	uations and deso hate solution is t	cribe the process for th reated with acidic solut	e developing of black an tion turns milky white. Gi	d white photographic film. ve the half reaction of the ab	When sodium pove described [JEE 2005]
5.	MC (A Colou liquid a temper transition White t	$Cl_4 \xrightarrow{Zn} (E)$ rless t room ature, n metal moist Air fumes c)	) purple colour comp	bound		
	Identify	(A), (B) and (C)	). Also explain colour di	ifference between MCl₄ a	nd <b>(B)</b> .	[ <b>JEE 2005</b> ]
6.	Match th	ne reactions in C Column I	Column I with nature of	f the reactions/type of the Column II	products in Column II.	[JEE 2007]
	(A)	$O_2^- \rightarrow O_2 + O_2$	$D_2^{2-}$	(p) Redox reaction	n	
	<b>(B)</b>	$CrO_{4}^{2-} + H^{+}$ -	$\rightarrow$	(q) One of the pr	oducts has trigonal planar s	tructure
	(C)	$MnO_4^- + NO_2^-$	$+ H^+ \rightarrow$	(r) Dimeric bridg	ed tetrahedral metal ion	
	<b>(D</b> )	$NO_3^- + H_2SO_2$	$_{4} + Fe^{2+} \rightarrow$	(s) Disproportion	ation	
7.	Among t (A) CuCl	the following, the <b>(B)</b> K	ne coloured compound [Cu(CN) <sub>4</sub> ]	is : ( <b>C</b> ) CuF <sub>2</sub>	<b>(D)</b> [Cu(CH <sub>3</sub> CN) <sub>4</sub> ]BF <sub>4</sub>	[ <b>JEE 2008</b> ]
8.	The oxic	lation number o	f Mn in the product of a	alkaline oxidative fusion	of MnO <sub>2</sub> is.	[ <b>JEE 2009</b> ]
9.	Reduction (A) 3 ele (C) 3 ele	on of the metal of the trons in neutral ctrons in alkalin	centre in aqueous perm l medium e medium	anganate ion involves : (B) 5 electrons in (D) 5 electrons in	neutral medium acidic medium	[ <b>JEE 2011</b> ]

10.	The colour of light absorbed by an aqueous solution of CuSO <sub>4</sub> is :				
	(A) organge-red	(B) blue-green	(C) yellow	(D) violet	
11.	Which of the following halides react(s) with $AgNO_3(aq)$ to give a precipitate that dissolves in $Na_2S_2O_3(aq)$ ?				
	(A) HCl	(B) HF	(C) HBr	(D) HI	[ <b>JEE 2012</b> ]

		моск	TEST	
	SE	CTION - I : STRAIGH	HT OBJECTIVE TYPE	
1.	In which compound does (A) $NH_4 VO_2$	vanadium have an oxidation (B) $K_4[V(CN)_6]$	n number of +4? (C) $VSO_4$	(D) VOSO <sub>4</sub>
2.	In which of the following operiodic table? (A) $MnO_4^-$	oxoanion the oxidation state (B) $Cr_2O_7^{-2}$	e of central atom is not same as that o (C) $VO_4^{3-}$	f its group number in the <b>(D)</b> $\text{FeO}_4^{2-}$
3.	Which of the following st (A) Transition elements er (B) $IE_2$ of ${}_{23}V < {}_{24}Cr > {}_{25}M$ (C) Ni(II) compounds are (D) The elements which give	atements is incorrect? xhibit higher enthalpies of a n and $_{28}$ Ni $<_{29}$ Cu $>_{30}$ Zn. more stable than Pt(II) whe ive the greatest number of or	ntomization as they have stonger interer as Pt (IV) compounds are more statistication states do not occur in or near	eratomic interaction. able than Ni(IV). r the middle of the series.
4.	At 300°C, FeCl <sub>3</sub> : (A) Decomposes into FeC (C) Sublimes to give liquid	$l_2$ and $Cl_2$ d FeCl <sub>3</sub>	<ul> <li>(B) Decomposes into Fe and Cl<sub>2</sub></li> <li>(D) Sublimes to give gaseous dimensional dimensiona</li></ul>	r (FeCl <sub>3</sub> ) <sub>2</sub>
5.	Fe $Cr_2O_4$ + Na $_2CO_3$ + $O_2$ — Which of the following sta (A) In all three compound (B) [Z] is a deep-blue viole (C) Saturated solution of (D) All of the above	$\xrightarrow{\text{Fusion}} [X] \xrightarrow{H^+}_{H_2O} [Y] \xrightarrow{H^+}_{H_2O}$ atements are true for composed of the chromium is in +6 oxistic to coloured compound which of [Y] gives bright orange composed of the chromited of the chromit	$\xrightarrow{H^{+}}{H_2O_2} [Z]$ unds [X], [Y] and [Z] ? dation state. decomposes rapidly in aqueous solution apound, chromic anhydride, with correct	on into $Cr^{+3}$ and dioxygen. nc. $H_2SO_4$
6.	FeCl <sub>3</sub> .6H <sub>2</sub> O + C(CH <sub>3</sub> ) <sub>2</sub> (CH Reaction product are : (A) FeCl <sub>3</sub> , CH <sub>3</sub> OH and CH <sub>3</sub> (C) FeCl <sub>2</sub> , HCl and CH <sub>3</sub> CC	$(1_3O)_2 \rightarrow \text{Products}$ $_3\text{COCH}_3$ $(OCH_3)$	(B) $(CH_3O)_3Fe$ , HCl and $H_2O$ (D) $Fe(OH)_3$ , $FeCl_3$ and $CH_3COCH_3$	
7	SECTIC Purclusite is MpO, used t	DN - II : MULTIPLE (	NnO I MnO <sup>2-</sup> II MN	<u>)</u> -
1.	<ul> <li>Stesp I and II are respective</li> <li>(A) fuse with KOH/ air and</li> <li>(B) fuse with KOH/KNO<sub>3</sub></li> <li>(C) fuse with conc. HNO<sub>3</sub></li> <li>(D) dissolve in H<sub>2</sub>O and or</li> </ul>	vely : d electrolytic oxidation and electrolytic oxidation /air and electrolytic reductio xidation	$2, \text{ who}_2 \xrightarrow{} \text{ who}_4 \xrightarrow{} \text{ who}_4$	· <sub>4</sub> .
8.	Which of the following ch (A) $C_6H_4(OH)_2 + 2AgBr \rightarrow (C) AgBr + 2NH_3(aq) \rightarrow [A]$	hemical reaction(s) is(are) in $Ag(NH_3)_2]Br$	woled in the developing of photogra (B) $AgBr + 2Na_2S_2O_3 \rightarrow Na_3[Ag(S_2) \rightarrow 2AgBr + Na_2S_2O_3 \rightarrow Ag_2S_2O_3 + Ag_2S_2O_$	nphic plate? O <sub>3</sub> ) <sub>2</sub> ]+NaBr 2NaBr

9. The hydrated cupric chloride is strongly heated. Which of the following statement(s) is (are) correct for this :
 (A) It is reduced to Cu<sub>2</sub>Cl<sub>2</sub>
 (B) Cupric oxide is formed along wiht Cu<sub>2</sub>Cl<sub>2</sub>
 (C) Only Cl<sub>2</sub> is liberated
 (D) Cl<sub>2</sub> and HCl both are liberated

- **10** Potassium manganate  $(K_2 MnO_4)$  is formed. When :
  - (A) cuprous oxide reacts iwth alkaline  $KMnO_4$  solution
  - (B) manganese dioxide is fused with  $K_2CO_3$  in presence of KNO<sub>3</sub>.
  - (C) formaldehyde reacts with potassium permanganate in presence of concentrated potassium hydroxide solution.
  - (D) potassium permanganate is heated with potassium hydroxide.

## **SECTION - III : ASSERTION AND REASON TYPE**

- **11. Statement -1** : The value of enthalpy of atomisation is generally maximum at about the middle of each series.
  - **Statement 2**: There is one unpaired electron per d-orbital and this results in stronger interatomic interaction.
  - (A) Statement 1 is True, Statement-2 is True, Statemen-2 is a correct explanatio for Statement-1.
  - (B) Statement-2 is True, Statement-2 is True, Statement-2 is NOT a correct explanation for Statement-1.
  - (C) Statement-1 is True, Statement-2 is False
  - (D) Statement-1 is False, Statement-2 is True
- 12. Statement-1: To a solution of potassium chromate if a strong acid is added it changes its colour from yellow to organce.
  - **Statement 2** : The colour change is due to hte oxidation of potassium chromate.
  - (A) Statemen-1 is True, Statement-2 is True, Statement-2 is a correct explanation for Statement-1.
  - (B) Statement-1 is True, Statement-2 is True, Statement-2 is NOT a correct explanation for Statement-1.
  - (C) Statement-1 is True, Statement-2 is False.
  - **(D)** Statement-1 is False, Statement-2 is True

Consider the following statements and arrange in the order of true/false as given in the codes.

- **S1** : Interstitial compounds have high melting points, higher than thoses of pure metals.
  - **S2** : Permangnate titrations in presence of hydrochloric acid are unsatisfactory.
  - **S3** :  $KMnO_4$  does not act as an oxidising agent in strong alkaline medium.

**S4** :  $KMnO_4$  on heating in current of H<sub>2</sub> gives MnO.

- 14. S1 :  $Mn^{2+}$  compounds more stable than  $Fe^{2+}$  towards oxidation to their +3 state.
  - S2 : Titanium and copper both in the first series of transition metals exhibits +1 oxidation state most frequently.
  - **S3** : Cu<sup>+</sup> ion is stable in aqueous solutions.
  - S4 : The E° value for the  $Mn^{3+}/Mn^{2+}$  couple much more positive than that for  $Cr^{3+}/Cr^{2+}$  or  $Fe^{3+}/Fe^{2+}$
  - (A) TTFT (B) TFFT (C) TFTT (D) FFTF

## **SECTION - IV : COMPREHENSION TYPE**

Read the following comprehensions carefully and answer the questions.

#### **Comprehension #1**

13.

The (n-1) d shel of electrons in d-block elements is expanding and, therefore, they have many physical and chemical properties in common. They show variable oxidation state and all are metals. The transition elements i.e. d-block elements show an unparalled tendency to form coordination compounds with those groups while are able to donate an electron pair (i.e. lewis base).

- **15.** Which oneis most acidic?
  - **(A)**  $Cr_2O_3$  **(B)**  $V_2O_5$  **(C)**  $Mn_2O_7$  **(D)**  $Fe_2O_3$

- **16.** Which of the following statement is incorrect?
  - (A) Across a period from Sc toCu the densities increase with increasing atomic number
  - (B) The melting point of transition elements rise to a maximum from Sc to Cr and then decreases from Fe to Zn.
  - (C) Transition elements have high entablpies of atomization and in 3d series increases regularly from Sc to Cu.
  - (D) On going down a group from 3d to 6d series the stability of higher oxidation state increases with increasing atomic number.

17. Which of the following products are obtained when Na<sub>2</sub>CO<sub>3</sub> is added to a solution of copper sulphate?

- (A) Basic coper carbonate  $[CuCO_3Cu(OH)_2]$ , sodium sulphate and  $CO_2$ .
- **(B)** Copper hydroxide, sodium sulphate and  $CO_2$ .
- (C) Copper carbonate sodium sulphate and  $CO_2$ .
- (D) Copper carbonate and sodium sulphate.

#### Comprehension # 2

Paramagnetism is a property due to the presence of unpaired electrons. In case of transition metals, as they contain unpaired electrons in the (n-1) d orbitals, most of the transition metal ions and their compounds are paramagnetic. Paramagnetism increases with increases in number of unpaired electrons. Magtnetic moment is calculated from 'Spin only formula' viz.

 $\mu = \sqrt{n(n+2)}$  B.M. = n = no. of unpaired electrons

Similarly the colour of the compounds of transitions metals may be attributed to the presence of incomplete (n-1) d subshell. When an electron from lower energy of d-orbital is excited to a higher energy d-orbital, the energy of excitation corresponds to the frequency of light absorbed. This frequence generally lies in the visible region. The colour observed corresponds to complementary colour of the light absorbed. The frequency of the light absorbed is determined by the nature of the ligand.

- 18. Which of the following of compounds is expected to exhibit same colour in aqueous solution.
  (A) FeCl<sub>2</sub>, CuCl<sub>2</sub>
  (B) VOCl<sub>2</sub>, CuCl<sub>2</sub>
  (C) VOCl<sub>2</sub>, FeCl<sub>2</sub>
  (D) FeCl<sub>2</sub>, MnCl<sub>2</sub>
- 19. Titanium shows magnetic moment of 1.73 BM in its compound. What is the oxidation state of titanium in the compound?
   (A)+2
   (B)+1
   (C)+3
   (D)+4
- **20.** Identify the incorrect statement.
  - (A)  $Mn^{2+}$  has the highest paramagnetism amongst the bivalent cations of the I<sup>st</sup> transition series.
  - (B) The coloured ions or compounds of transition elements are due to d-d transition and charge transfer spectrum.
  - (C) In 3 d series the paramagnetic character first increase to maximum & then starts decreasing
  - **(D)** None of these

#### Comprehension #3

- (I) A powdered substanced (A) on treatment with fusion gives a green coloured compound (B).
- (II) The solution of (B) in boiling water on acidification with dilute  $H_2SO_4$  gives a pink coloured compound (C)
- (III) The aqueous solution of (A) on treatment iwth NaOH and Br<sub>2</sub>– water gives a compound (D).
- (IV) A solution of (D) in conc.  $HNO_3$  on treatment with lead peroxide at boiling temperature produced a compound (E) which was of the same colour at that of (C).
- (V) A solution of (A) on treatment iwht a solution of barium chloride gave a white precipitate of compound (F) which was insoluble in conc. HNO<sub>3</sub> and conc. HCl.

- **21.** Which of the following is true for compound (C).
  - (A) It oxidises ammonia to nitrogen dioxide in neutral medium
  - (B) It's pink colour is due to d-d transition.
  - (C) It can be oxidised by ozone.

(D) It is obtained by alkaline fusion of pyrolusite in presence of air followed by electrolytic oxidation.

- 22. The oxidation state of centre metal ions of (A), (B) and (C) compounds are respectively:
   (A) + II, + VI and + VII
   (B) + II, + VI and + VI
   (C) + II, + VII and VII
   (D) + VI, + VII and + VII
- 23. Consider the following statements :
  - (I) Anions of both (B) and (C) are diamagnetic and have tetradedral geometry.
  - (II) Anions of both (B) and (C) are paramagnetic and have tetrahedral geometry.
  - (III) Anions of (B) and (C) are paramagnetic and have tetrahedral geometry.
  - (IV) Green coloured compound (B) in a neutral or acidic medium disproportionated to give (C) and (D). Of these select the correct one from the codes given :
  - (A) I and III only (B) II and III only (C) II and IV only (D) III and IV only

### **SECTION - V : MATRIX - MATCH TYPE**

24Match the reactions in Column I with nature of the reactions/type of the products in Columns II.Column IColumn II(A)  $MnO_4^{2-} + H^+ \rightarrow$ (p) One of the products of transition elementis paramagnetic(B)  $Cu^+(aq) \rightarrow$ (p) One of the products of transition elementis paramagnetic(C)  $Cr_2O_7^{-2}(s) + H^+(conc.) + Cl^-(s) \rightarrow$ (r) One of the products is liberated as coloured vapours.(D)  $Fe_2(SO_4)_3 + l^- \rightarrow$ (s) In one of the products central atom exhibits its highest oxidation state.

25 Match the reaction in Column I with type of the products/applications in Column II. Column I

(A)  $\operatorname{FeSO}_4 \xrightarrow{\Delta}$ (B)  $\operatorname{Mn}^{2+} + \operatorname{S_2O}_8^{2-} + \operatorname{H_2O} \rightarrow$ (C)  $\operatorname{Na_2Cr_2O_7(saturated salt)} + \operatorname{H_2SO}_4(\operatorname{conc.}) \rightarrow$ (D)  $\operatorname{N_2H_4} + \operatorname{CuSO}_4 \rightarrow$ 

- (p) One of the product is coloured due to charge transfer
- (q) One of the product is in + VI oxidation state
- (r) Redox reaction
- (s) One of the product is acidic oxide
- 26 Match the reaction in Column I with type of the products/applications in Column II. Column I Column II
  - (A) HO- $\bigcirc$ -OH(aq)+AgBr(s)  $\rightarrow$ (B) BaCl<sub>2</sub>(s)+K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>(s)+H<sub>2</sub>SO<sub>4</sub>(conc.)  $\rightarrow$ (C) FeSO<sub>4</sub>(aq)+K<sub>3</sub>[Fe(CN)<sub>6</sub>](aq)  $\rightarrow$ (D) Cu(OH)<sub>2</sub>(s)+NH<sub>4</sub>OH(aq)+(NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>(s) $\rightarrow$ (E) ZnO (s)+Co(NO<sub>3</sub>)<sub>2</sub>(s)  $\xrightarrow{\Delta}$

Column

- (q) Schwitzer's reagent
- (r) Rinmann's green pigment

(p) Turnbull's blue pigment

(s) Chromyl chloride test

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(t) Photography
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### **SECTION - VI : SUBJECTIVE TYPE**

- 27. Write two compounds in which compound does vanadium have an oxidation number of +4?
- 28. How many elements are present in each transition series? Why this number can not be less or more?
- **29.** Why do transition element exhibit higher enthalpies of atomisation ?
- 30. The second and third member in each group of transition elements have similar atomic radii. Explain why?
- 31.  $E^{\circ}$  for Mn<sup>+3</sup> / Mn<sup>+2</sup> is more positive then for Fe<sup>+3</sup> / Fe<sup>+2</sup>. Account for this?

## **ANSWER KEY**

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#### **EXERCISE - 1**

1.	С	2.	С	<b>3.</b> B	<b>4.</b> C	5. B	6. A	<b>7.</b> C	<b>8.</b> A	9. D	10. C	11. B	12. B	13. A
14.	С	15.	С	16. D	17. C	18. C	19. C	<b>20.</b> A	<b>21.</b> D	<b>22.</b> D	23. D	<b>24.</b> B	<b>25.</b> D	<b>26.</b> B
27.	В	28.	С	<b>29.</b> D	<b>30</b> B	<b>31.</b> B	32. D	<b>33.</b> C	<b>34.</b> A	35. D	<b>36.</b> A	<b>37.</b> A	<b>38.</b> A	<b>39.</b> B
40.	А	41.	D	<b>42.</b> C	<b>43.</b> B	<b>44.</b> B	<b>45.</b> C	<b>46.</b> A	<b>47.</b> A	<b>48.</b> B	<b>49.</b> C	50. D	<b>51.</b> D	52. A
53.	С	54.	D	55. D	56. B	<b>57.</b> A	58. A	<b>59.</b> A	<b>60.</b> A	61. C	<b>62.</b> B	<b>63.</b> A	<b>64.</b> A	<b>65.</b> C
66.	D	67.	С	<b>68.</b> B	<b>69.</b> A	<b>70.</b> C	<b>71.</b> C	<b>72.</b> C	<b>73.</b> C	74. B	<b>75.</b> D	76. D	77. A	<b>78.</b> C
79.	А	80.	С	<b>81.</b> B	82. A	<b>83.</b> A	<b>84.</b> A	<b>85.</b> B	<b>86.</b> A	<b>87.</b> A	<b>88.</b> C	<b>89.</b> C	90. A	91. C
92.	В	93.	С	94. C	95. B	96. D	<b>97.</b> B	98. A	99. D	<b>100.</b> C	<b>101.</b> C	102. D	103. A	<b>104.</b> A

#### EXERCISE - 2 : PART # I

 1. A, B, D
 2. D
 3. B
 4. A, D
 5. A, B, C, D
 6. A, B
 7. A, B, C
 8. A, B, C
 9. A, B, D

 10. A, B
 11. A, B
 12. A, B, C
 13. A, B
 14. A, D
 15. A
 16. A, B
 17. A, B, C, D
 18. B, C, D

 19. B, D
 20. A, B
 21. A, B, D
 22. A, B
 23. A, B, D
 24. A, B
 25. B
 26. A
 27. A
 28. B

 29. C
 30. C
 31. A
 32. A
 33. D
 34. D
 35. B
 36. D
 37. C
 38. B
 39. C
 40. C
 41. D

 42. D
 43. A
 44. A
 45. B
 46. B
 47. C
 48. C
 49. A
 50. A
 51. D
 52. D
 53. B
 54. A

 55. A
 56. D
 57. C
 58. A
 59. A
 60. A
 61. D
 62. A
 63. C
 64. D
 65. C
 66. A, C

 67. A, C, D
 68. A, B, C
 69. A, B, C, D
 71. A, B, C
 71. A, B, C
 71. A, B, C

#### PART # II

 1. A
 2. C
 3. A
 4. B
 5. A
 6. C
 7. B
 8. D
 9. A
 10. A
 11. A
 12. A
 13. C

 14. B
 15. A
 16. B
 17. B
 18. B
 19. B
 20. A
 21. D
 22. A
 23. A
 24. B
 25. A
 26. A

 27. A
 28. A

#### **EXERCISE - 3 : PART # I**

1.	$A \rightarrow q, s; B \rightarrow r, s; C \rightarrow p, s; D \rightarrow p, s$	2.	$A \rightarrow r, B \rightarrow s; C \rightarrow p; D \rightarrow q$
3.	$A \rightarrow s$ ; $B \rightarrow q, r$ ; $C \rightarrow q, D \rightarrow p$	4.	$A \rightarrow r, s; B \rightarrow r, s; C \rightarrow p, q; D \rightarrow p$
5.	$A \rightarrow r$ ; $B \rightarrow s$ ; $C \rightarrow p$ ; $D \rightarrow q$	6.	$A \rightarrow t$ ; $B \rightarrow s$ ; $C \rightarrow p$ ; $D \rightarrow q$ ; $E \rightarrow r$
7.	$A \rightarrow q, r, s; B \rightarrow p, q, r; C \rightarrow p, q, s; D \rightarrow q, r$	8.	$A \rightarrow q, s; B \rightarrow q; C \rightarrow p; D \rightarrow r.$
9.	$A \rightarrow r, s; B \rightarrow p, q; C \rightarrow p, r, s; D \rightarrow r, s.$		

#### PART # II

Comprehension #1:	1.	D	2.	А	3.	В				
Comprehension # 2 :	1.	В	2.	С	3.	С	4.	D		
Comprehension # 3 :	1.	С	2.	С	3.	D				
Comprehension #4:	1.	А	2.	С						
Comprehension # 5 :	1.	В	2.	А	3.	В	4.	С	5.	D
Comprehension # 6 :	1.	D	2.	А	3.	С	4.	В	5.	С

#### **EXERCISE - 5 : PART # I**

 1. 3
 2. 3
 3. 4
 4. 1
 5. 4
 6. 2
 7. 2
 8. 1
 9. 1
 10. 2
 11. 1
 12. 4
 13. 4

 14. 4
 15. 1
 16. 4

#### PART # II

#### **1.** A **2.** B **3.** B

4. Developer is usually a weak reducing agent like potassium ferrous oxalate, an alkaline solution of pyrogallol or an alkaline solution of quinol.

In the process of development of the photographic film, the exposed/activated AgBr grains are preferentially reduced by one of the reducing agent described above.



 $AgBr + e^{-} \longrightarrow Ags + Br^{-}$ 

The photographic film is then fixed by washing with hypo solution to remove the unreduced AgBr grains from the film.

$$\begin{split} & AgBr + 2Na_2S_2O_3 \longrightarrow Na_3[Ag(S_2O_3)_2] + NaBr. \\ & S_2O_3^{2-} + H_2O \longrightarrow 2SO_2 + 2H^+ + 4e^-. \\ & S_2O_3^{2-} + 6H^+ \longrightarrow 2S \downarrow \text{ (white milky)} + 3H_2O. \end{split}$$

(A) is TiCl<sub>4</sub> as it has no unpaired electron and is liguid at room temperature on account of covalent character because of high polarising power of Ti<sup>+4</sup>. TiCl<sub>4</sub> being covalent gets hydrolysed forming TiO<sub>2</sub>(H<sub>2</sub>O)<sub>2</sub> and HCl (B) which fumes in air. In [Ti(H<sub>2</sub>O)<sub>6</sub>]Cl<sub>3</sub> complex Ti(III) has one unpaired electron(3d<sup>1</sup>) which gives violet / purple colour due to d-d transition.

$$\begin{array}{c} \text{TiCl}_{4} \xrightarrow{Zn} \text{TiCl}_{3} \xrightarrow{\text{Hydrolysis}}_{\text{H}_{2}\text{O}} \xrightarrow{[\text{Ti} (\text{H}_{2}\text{O})_{6}]\text{Cl}_{3}}_{\text{purple or violet}}\\ \text{Colour less} \qquad (B) \\ & \downarrow \\ \text{moist air} \\ \text{TiO}_{2} \cdot (\text{H}_{2}\text{O})_{2} + \text{HCl} \text{ (white fumes)} \end{array}$$

6.  $A \rightarrow p, s; B \rightarrow r; C \rightarrow p, q; D \rightarrow p.$  7. (C) 8. 6 9. ACD 10. A 11. ACD

#### **MOCK-TEST**

 1. D
 2. D
 3. D
 4. D
 5. D
 6. A
 7. A,B
 8. A,B
 9. A,B,D
 10. A,B,C,D
 11. A

 12. C
 13. A
 14. B
 15. C
 16. C
 17. A
 18. B
 19. C
 20. D
 21. D
 22. A
 23. D

 24.  $A \rightarrow p, s; B \rightarrow r; C \rightarrow p, q; D \rightarrow p$  25.  $A \rightarrow q, r, s; B - p, q, r; C - p, q, s 4 - q, r$  

 26.  $A \rightarrow t; B \rightarrow s; C \rightarrow p; D \rightarrow q; E \rightarrow r$ 

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