## The d-and f-Block Elements

## **Practice Questions**

**1**. Which of the following element does not have  $(n-1) d^{10}ns^2$  electronic configuration?

(a) Zn	<i>(b)</i> Cd
(c) Hg	<i>(d)</i> Cu

2. The ground state electronic configuration of neutral titanium atom is

(a) [Ar] $4s^2 4 p^2$	(b) [Ar] $3d^2 4s^2$
(c) [Ar] $4s^2 p_x^1 p_y^1$	(d) [Ar] $3d^5$

- **3.** The third ionisation enthalpy is minimum form (a) Mn (b) Ni (c) Co (d) Fe
- **4.** Which element can have oxidation state from 4 to 6? (a) Fe (b) Mg (c) Co (*d*) Cr
- 5. Magnetic moment of a transition metal ion is found to be 3.87 BM. The number of unpaired electrons present in it is (a) 2*(b)* 3

<i>(c)</i> 4		(d) 5
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6. Which of the following aqueous solutions will be coloured?

(a) $\operatorname{Zn}(\operatorname{NO}_3)_2$	(b) $LiNO_3$
(c) $CoNO_3$	(d) $HgCl_2$

- 7. Which of the following alloys contain Cu and Zn? (a) Brass (b) Bronze (d) All of these (c) Bell metal
- **8.** Acidified potassium dichromate oxidises (a) iodides to iodine (b) sulphides to sulphur (c) tin (IV) to tin (II) (d) Both (a) and (b)
- **9.** Which of the following can react with  $K_2Cr_2O_7$ ?

(a) $SO_3^{-2}$	(b) $CO_3^{-2}$
(c) $SO_4^{-2}$	( <i>d</i> ) $NO_{3}^{-}$

**10.** Dichromates are generally prepared by the fusion of chromite ore with (a) sodium carbonate (b) potassium carbonate

(c) Both (a) and	(b) <i>(a</i>	l) Neither (a	a) nor (b	)

- **11.** Permanganate ion  $(MnO_{4})$  is dark purple coloured though Mn is in + 7 oxidation state with  $d^0$ configuration. This is due to (a) *d*-*d* transition (b) charge transfer from metal to ligand
  - (c) charge transfer from ligand to metal
  - (d) All of the above

**12.** Name the gas that can readily decolourise acidified KMnO<sub>4</sub> solution.

(a) CO <sub>2</sub> (c) NO <sub>2</sub>		$\begin{array}{c} (b) \text{ SO}_2\\ (d) \text{ P}_2 \text{ O}_5 \end{array}$

**13.** The product of following reaction is

 $K_2CrO_4 + dil. HNO_3 (excess) \longrightarrow$ (a)  $\operatorname{Cr}^{3+}_{2}$  and  $\operatorname{Cr}_{2}O_{2}^{2-}$ (b)  $\operatorname{Cr}_2\operatorname{O}_7^{2-}$ ,  $\operatorname{NO}_3^-$  and  $\operatorname{H}_2\operatorname{O}$ (c) Only  $Cr^{3+}$ (d) Only  $Cr^{7+}$ 

- 14. The green manganate and purple permanganate are respectively (a) paramagnetic, diamagnetic
  - (b) diamagnetic, paramagnetic
  - (c) paramagnetic, paramagnetic
  - (d) diamagnetic, diamagnetic
- **15.** The most common lanthanoid among the following is (a) lanthanum (b) cerium (c) promethium (d) plutonium
- **16.** Which of the following pairs has the same size? (a)  $Zn^{2+}$ ,  $Hf^{4+}$ (b)  $Fe^{2+}$ ,  $Ni^{2+}$ (d)  $Zr^{4+}$ ,  $Hf^{4+}$ (c)  $Zr^{4+}$ ,  $Ti^{4+}$
- 17. The correct order of ionic radii of Ce, Pm, Gd and Dy in +3 oxidation state is (a)  $Ce^{3+} < Gd^{3+} < Pm^{3+} < Dy^{3+}$ (b)  $Ce^{3+} < Pm^{3+} < Gd^{3+} < Dy^{3+}$ (c)  $Dy^{3+} < Gd^{3+} < Pm^{3+} < Ce^{3+}$ (d)  $Pm^{3+} < Ce^{3+} < Dv^{3+} < Gd^{3+}$
- **18.** A man made white silvery metal, radioactive in nature, has strong tendency to form oxocations and complexes. It is used as a nuclear fuel in atomic reactor. This metal is a
  - (a) actinide
  - (b) lanthanide
  - (c) representative element
  - (*d*) transition metal
- **19.** All the actinoids are believed to have the electronic configuration of

(a) $6s^2$	(b) $7s^2$
(c) $5f^{14}$	(d) $6d^{10}$

**20.** What will be the most common oxidation state shown by the actinoids?

(a) -3	<i>(b)</i> +3
<i>(c)</i> –4	<i>(d)</i> +4

21.	Which of the follow	ving elements shows maximum	
	number of different	oxidation states in its compounds	?
	(a) Eu	(b) I o	

( <i>a</i> ) Eu	( <i>b</i> ) La
(c) Gd	<i>(d)</i> Am

22.	The actinoids resemb	le the lanthanoids in having
	more compounds in	
	(a) +3 state	<i>(b)</i> +4 state
	A 1	

(c) +5 state	(d) +2 state

23. Compound(s) useful in the battery industries is/are(a) MnO<sub>2</sub>(b) Zn

(c)	Ni/Cd	(d)	All of these

- **24.** Catalyst used in the oxidation of  $SO_2$  in the manufacture of  $H_2SO_4$  is
  - (a)  $CuCl_2$
  - (b)  $V_2O_5$
  - (c)  $MnO_2$
  - (d) None of thesee
- **25.** Which of the following compounds form the basis, if Ziegler-Natta catalysts is used to manufacture of polythene?

(a) TiCl<sub>4</sub>
(b) Al(CH<sub>3</sub>)<sub>3</sub>
(c) TiCl<sub>4</sub> with Al(CH<sub>3</sub>)<sub>3</sub>
(d) None of these

## ANSWERS

1	. (d)	2.	(b)	3.	(d)	4.	(d)	5.	(b)	6.	(C)	7.	(a)	8.	(d)	9.	(a)	10.	(C)
11	. (c)	12.	(b)	13.	(b)	14.	(a)	15.	(b)	16.	(d)	17.	(C)	18.	(a)	19.	(b)	20.	(b)
21	. (d)	22.	(a)	23.	(d)	24.	(b)	25.	(C)										

## **Hints & Solutions**

**1.** (*d*) The electronic configurations of Zn, Cd and Hg are represented by the general formula  $(n - 1) d^{10}ns^2$ . The orbitals in these elements are completely filled in the ground state as well as in their common oxidation states.

Cu has electronic configuration  $3d^{10}4s^1$ . Thus, Cu does not have  $(n-1)d^{10}ns^2$  electronic configuration.

**3.** (*d*) Fe has minimum value of third ionisation enthalpy. Ground state electronic configuration of Fe is  $[Ar]3d^{6}4s^{2}$ .

 $Fe^{2+}$  has [Ar]3 $d^6$  configuration, whereas  $Fe^{3+}$  has [Ar]3 $d^5$ . The latter is a stable configuration and easier to ionise  $Fe^{2+}$  to  $Fe^{3+}$  than expected.

Hence possess least value.

Ni has highest value of third ionisation enthalpy due to its greater nuclear charge and smaller size.

Due to the same reason, Co > Fe > Mn should be the order for the remaining elements, but the anomalous order is due to greater stability of  $\text{Mn}^{2+}$  having [Ar] $3d^5$  configuration than  $\text{Mn}^{3+}$  with [Ar] $3d^4$ .

Whereas for  $\text{Co}^+$ , the electronic configuration is [Ar] $3d^7$ , Thus, ionisation enthalpy needed to remove third electron is less as compared to that of Mn<sup>2+</sup>.

**4.** (*d*) Oxidation state of alkaline earth metal (i.e. Mg) is fixed and equal to (+)2.

Oxidation state of *d*-block elements can vary,

i.e. for Fe and Co it is as follows :

Oxidation state of Co = (+) 2 to (+) 4 and oxidation state of Fe = (+) 2 to (+) 6, but (+) 6 is less stable.

Oxidation state of Cr = (+) 2 to (+) 6, where (+) 6 state is more stable.

2)

**5.** (b) Magnetic moment of a transition metal ion

$$(\mu) = \sqrt{n(n+1)}$$

 $3.87 = \sqrt{n(n+2)}$  (Given,  $\mu = 3.87$ )

On solving, n = 3

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Thus, number of unpaired electrons are 3.

- **6.** (*c*) Only Co<sup>+</sup>, because of the presence of unpaired electrons in *d*-orbitals show *d*-*d* transition and, hence it is coloured.
- **8.** (*d*) Acidified potassium dichromate oxidises iodides to iodine, sulphides to sulphur, tin(II) to tin(IV), iron(II) salts to iron (III).
- **9.** (*a*) Oxidation state of central atoms are as follows :

Sulphur in  $SO_3^{-2} = +4$ Carbon in  $CO_3^{-2} = +4$ Sulphur in  $SO_4^{-2} = +6$ 

Nitrogen in 
$$NO_3^- = +5$$

Since,  $K_2Cr_2O_7$  is a strong oxidising agent it can oxidise the species, which is not in its most possible positive oxidising state. Thus,  $SO_3^{-2}$  can react with  $K_2Cr_2O_7$  as follows :

 $\operatorname{Cr}_2O_7^{-2}(aq) + 3\operatorname{SO}_3^{-2}(aq) + 8\operatorname{H}^+(aq) \longrightarrow 2\operatorname{Cr}^{3+}(aq)$ 

 $+ 3SO_4^{2-}(aq) + 4H_2O(l)$ 

**10.** (c) Dichromates are generally prepared from chromate which in turn are obtained by the fusion of chromite ore (FeCr<sub>2</sub>O<sub>4</sub>) with sodium or potassium carbonate in free access of air.

 $\begin{array}{l} 4FeCr_{2}O_{4}+8Na_{2}CO_{3}+7O_{2}\longrightarrow\\ \\ 8Na_{2}CrO_{4}+2Fe_{2}O_{3}+8CO_{2}\\ \\ 2Na_{2}CrO_{4}+2H^{+}\longrightarrow Na_{2}Cr_{2}O_{7}+2Na^{+}H_{2}O \end{array}$ 

**11** (c) In  $MnO_4^{-}$ ,

x + (-2)4 = -1

 $\Rightarrow$  x = +7

 $_{25}$  Mn<sup>+7</sup> = [Ar], no unpaired electrons.

Thus, it will not show *d-d* transition. It is dark purple coloured due to charge transfer from ligand to metal.

**12.** (*b*) SO<sub>2</sub> gas can readily decolourise acidified KMnO<sub>4</sub> solution because KMnO<sub>4</sub> is an oxidising agent that easily oxidises SO<sub>2</sub>,

$$2MnO_4^- + 5SO_2 + 2H_2O \longrightarrow 2Mn^{2+} + 5SO_4^{2-} + 4H^+$$

while other options such as  $NO_2$  (strong oxidising agent),  $CO_2$  (neither oxidising agent nor reducing agent) cannot decolourise acidified KMnO<sub>4</sub> solution.

**13.** (b) The product of given reaction is  $Cr_2O_7^{2-}$ ,  $NO_3^-$  and  $H_2O_7$ ,

$$\begin{array}{ccc} 2CrO_4^{2-} + 2HNO_3 \longrightarrow & Cr_2O_7^{2-} + 2NO_3^- + H_2O\\ Chromate & (Dil.) & Dichromate \\ (yellow) & (orange) \end{array}$$

- **14.** (*a*) The manganate and permanganate ions are tetrahedral. Here, the  $\pi$ -bonding takes place by overlap of *p*-orbitals of oxygen with *d*-orbitals of manganese. The green manganate is paramagnetic because of one unpaired electron but the purple permanganate is diamagnetic due to charge transfer.
- **15.** (*b*) Lanthanum is a *d*-block element, whereas plutonium is an actinoid. Both cerium and promethium are lanthanoids. But cerium is a common lanthanoid because it occurs naturally, whereas Pm does not occur naturally and is radioactive.
- **16.** (d)  $Zr^{4+}$  and  $HF^{4+}$  have same size due to lanthanoid contraction.
- **18.** (*a*) Actinide is a man-made white silvery metal, radioactive in nature has a strong tendency to form oxocations and complexes.
- **19.** (*b*) All the actinoids are believed to have the electronic configuration of  $7s^2$  and variable occupancy of 5f and 6d-subshell.
- **25** (c) TiCl<sub>4</sub> with Al(CH<sub>3</sub>)<sub>3</sub> forms the basis of *Ziegler-Natta* catalysts that is used to manufacture polythene.