## The *d*- and *f*-Block Elements

CHAPTER

## A TRY YOURSELF

## ANSWERS

**1.** This is because Ag (Z = 47) can show +2 oxidation state in which silver will have partially filled *d* orbitals.

**2.** The five *d*-orbitals when combined make a complete sphere. Thus, if the *d*-subshell is half-filled or completely filled, the distribution of electron density will be symmetrical as compared to an assymmetrical distribution of electron density for a partially filled *d*-orbital. This symmetrical distribution of electrons make the half-filled or completely filled *d*-subshell more stable (lower energy) as compared to partially filled *d*-subshell (more energy).

**3.** No unpaired electron is present in *d*-orbital of Hg, this no possibility of *d*-*d* transition. Here, weak metallic bond is present between atoms.

**4.** Enthalpy of atomisation reaches maximum upto middle, due to increased number of unpaired electrons at middle in the series. Greater is the *d-d* transition, stronger is metallic bond but after middle decrease in the number of unpaired electrons occurs that leads to weak metallic bonds. As a result, enthalpy of atomisation decreases.

**5.** Since standard reduction potential of  $Co^{3+}$  is larger than that of  $Co^{2+}$ , it suggests that  $Co^{3+}$  can be more easily reduced. Therefore, it is better oxidising agent than  $Co^{2+}$ .

**6.** Magnetic properties of a substance is determined by the number of unpaired electrons in the ion. In Cu(II) ion the outermost orbital configuration is  $3d^{9}$ . Thus, there is one unpaired electron. Hence, it is paramagnetic. In Cu(I), the outermost orbital configuration is  $3d^{10}$ . There is no unpaired electron and therefore, Cu (I) is diamagnetic.

7. In the solution, the following equilibria exists :  $Cr_2O_7^{2-} + H_2O \implies 2CrO_4^{2-} + 2H^+$  On adding an alkali, the concentration of  $H^+$  ions decreases, this equilibrium shifts in the forward direction. According to Lechatelier principle on adding acid, an increase in the  $H^+$  ions occurs causing the equilibrium to shift in backward direction.

8. (i) 
$$Cr_2O_7^{2-} + 6Fe^{2+} + 14H^+ \xrightarrow{\Delta} 2Cr^{3+} + 6Fe^{3+} + 7H_2O^{3+}$$

(ii) 
$$Cr_2O_7^{2-} + 3H_2S + 8H^+ \xrightarrow{\Delta} 2Cr^{3+} + 3S + 7H_2O$$

(iii) 
$$2KMnO_4 \xrightarrow{\Delta} K_2MnO_4 + MnO_2 + O_2$$

(iv) 
$$MnO_4^- + \frac{5Fe^{2+}}{5Fe^{2+}} + \frac{8H^+}{5Fe^{2+}} \rightarrow \frac{Mn^{2+}}{5Fe^{3+}} + \frac{4H_2O}{5Fe^{3+}}$$

**9.** The decreasing order of reducing strength is  $Sm^{2+} > Eu^{2+} > Yb^{2+}$ .

**10.** Due to lanthanoid contraction, Hf and Zr have almost similar size and therefore, their properties are similar.

**11.** 5*f* electrons of actinoids are less effectively shielded which results in quenching of orbital contribution. Therefore, magnetic moments of actinoid ions are less than the theoretically predicted values.

**12.** Chemistry of actinoids is more complicated than lanthanoids because

(i) actinoids show greater number of oxidation states due to the comparable energies of 5f, 6d and 7s-orbitals.

(ii) most of the actinoids are radioactive and the study of their chemistry in the laboratory is difficult.

- **13.** MnO<sub>2</sub>
- **14.** V<sub>2</sub>O<sub>5</sub>

**15.** TiCl<sub>4</sub> and Al(CH<sub>3</sub>)<sub>3</sub> (Zieglar Natta) are useful in the polymerisation of alkene.

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