# **CO-ORDINATION COMPOUNDS**

#### **QUESTIONS**

### **VSA QUESTIONS (1 - MARK QUESTIONS)**

- 1. Define the term coordination compound?
- 2. Write the names of counter ions in
  - (i) Hg [Co (SCN)<sub>4</sub>] and (ii)  $[Pt(NH_3)_4]$  Cl<sub>2</sub>.
- 3. Write the oxidation state of nickel in [Ni(CO)<sub>4</sub>]
- \*4. What is the coordination number of central atom in  $[Co(C_2O_4)_3]^{3-}$ ?

[**Ans.** : 6]

- 5. What is the coordination number of iron in [Fe (EDTA)]<sup>-</sup> ? [Ans.: 6]
- 6. Write the name of a complex compound used in chemotherapy.

[Ans.: Cis-Platin.  $[Pt(NH_3)_2 Cl_2]$ 

- 7. Name the compound used to estimate the hardness of water volumetrically.
- 8. Give the IUPAC name of [Pt Cl<sub>2</sub> (NH<sub>2</sub>CH<sub>3</sub>) (NH<sub>3</sub>)<sub>2</sub>] Cl.
- \*9. How many geometrical isomers are possible for the tetrahedral complex [Ni(CO)<sub>4</sub>].

[Ans.: No isomer, as the relative positions of the unidentate ligands attached to the central metal atom are same with respect to each other].

- 10. Arrange the following in the increasing order of conductivity in solution.  $[Ni(NH_3)_6]Cl_2$ ;  $[Co(NH_3)_6]Cl_3$  and  $[CoCl_2(en)_2]$  CI
- 11. Arrange the following ligands in increasing order of  $\Delta_0$  (Crystal field splitting energy) for octahedral complexes with a particular metal ion.

 $Cl^-$ ,  $NH_3$ ,  $I^-$ , CO, en.

- 12. Write I.U.P.A.C. name of Tollens' reagent.
- 13. Which is more stable?  $K_3[Fe(CN)_6]$  or  $K_4[Fe(CN)_6]$
- 14. Calculate the overall dissociation equilibrium constant for the  $[Cu(NH_3)_4]^{2+}$  ion. Given that overall stability constant ( $\beta_4$ ) for this complex is 2.1 × 10<sup>13</sup>. [Ans.: 4.7 × 10<sup>-14</sup>]
- \*15. What is a chelate ligand? Give one example.
- Write the I.U.P.A.C. name of Li[AlH<sub>4</sub>].
- 17. Name one homogeneous catalyst used in hydrogenation of alkenes.
- \*18. Name the types of isomerism shown by coordination entity : [CrCl<sub>2</sub>(Ox)<sub>2</sub>]<sup>3-</sup>
- \*19. [Ti(H<sub>2</sub>O)<sub>6</sub>]Cl<sub>3</sub> is coloured but on heating becomes colourless. Why?
- \*20. Write the IUPAC name of ionization isomer of [Co(NH<sub>3</sub>)<sub>5</sub>(SO<sub>4</sub>)] Br
- \*21. Write the formula and the name of the coordinate isomer of  $[Co(en)_3]$   $[Cr(CN)_6]$ .

[Ans.:  $[Cr(en)_3]$   $[Co(CN)_6]$  Tris- (ethane -1, 2, diammine) chromium (III) hexacyanocobaltate (III)]

## SA(I) TYPE QUESTIONS (2-MARK QUESTIONS)

- \*22. Write two differences between a double salt and a coordination compound with the help of an example of each.
- 23. Mention the main postulates of Werner's Theory.
- 24. Define (a) Homoleptic and (b) Heteroleptic complexes with the help of one example of each.
- 25. In the following coordination entity: [Cu(en)<sub>2</sub>]<sup>2+</sup>
  - (a) Identify the ligand involved and
  - (b) Oxidation state of copper metal.
- 27. Calculate the magnetic moments of the following complexes:
  - (i)  $[Fe(CN)_6]^{4-}$  (ii)  $[CoF_6]^{3-}$
- 28. Explain the following:
  - (a)  $[Fe(CN)_6]^{3-}$  is an inner orbital complex whereas  $[FeF_6]^{3-}$  is an outer orbital complex.
  - (b) NH<sub>3</sub> acts as complexing agent but NH<sub>4</sub><sup>+</sup> does not.

- 29. What type of structural isomerism is represented by the following complexes:
  - (a)  $[Mn(CO)_5(SCN)]$  and  $[Mn(CO)_5(NCS)]$
  - (b)  $[Co(NH_3)_5(NO_3)] SO_4$
- 30. How are complex compounds applicable in (a) electroplating of silver, gold or other noble metals (b) in photography.
- 31. Explain on the basis of Valance Bond Theory that diamagnetic  $[Ni(CN)_4]^{2-}$  has square planar structure and paramagnetic  $[NiCl_4]^{2-}$  ion has tetrahedal geometry.
- 23. Explain as to how the two complexes of nickel  $[Ni(CN)_4]^{2-}$  and  $Ni(CO)_4$  have different structures but do not differ in their magnetic behaviours. (At. no. of Ni = 28).
- 34. Draw the structures of geometrical isomers of the coordination complexes— $[Co(NH_3)_3Cl_3]$  and  $[CoCl_2(en)_2]^+$
- 35. Write the IUPAC name of the complexes :
  - (a)  $[NiCl_2 (PPh_3)_2]$  (b)  $[Co(NH_3)_4 Cl(NO_2)]$  Cl
  - (c)  $K[Cr(H_2O)_2 (C_2O_4)_2]$

[Hint.: (a) Dichloridobis(triphenylphosphine)nickel (II);

- 36. Using IUPAC norms write the formulae for the following :
  - (a) Terabromidocuprate (II)
  - (b) Pentaamminenitrito-O- Cobalt (III)
- \*37. How does EDTA help as a cure for lead poisoning?

[Ans.: Calcium in Ca-EDTA complex is replaced by lead in the body. The more soluble complex lead-EDTA is eliminated in urine].

37. A complex is prepared by mixing CoCl<sub>3</sub> and NH<sub>3</sub> in the molar ratio of 1:4. 0.1 m solution of this complex was found to freeze at -0.372°C. What is the formula of the complex?

 $K_f$  of water = 1.86°C/m

[Hint:  $\Delta T_f = i K_f \times m = i \times 1.86 \times 0.1$ 

 $\Delta T_f(obs) = 0.373$ °C

This means each molecule of complex dissociates into two ions. Hence the formula is i = 2

\*38. The [Mn(H<sub>2</sub>O)<sub>6</sub>]<sup>2+</sup> ion contains five unpaired electrons while [Mn(CN)<sub>6</sub>]<sup>4-</sup> ion contains only one unpaired electron. Explain using Crystal Field Theory:

#### SA (II) TYPE QUESTIONS (3 - MARK QUESTIONS)

- 39. Account for the following -
  - (i) [NiCl<sub>4</sub>]<sup>2-</sup> is paramagnetic while [Ni(CO)<sub>4</sub>] is diamagnetic though both are tetrahedral.
  - (ii)  $[Fe(H_2O)_6]^{3+}$  is strongly paramagnetic whereas  $[Ni(NH_3)_6]^{2+}$  weakly paramagnetic.
  - (iii)  $[\text{Co(NH}_3)_6]^{3+}$  is an inner orbital complex whereas  $[\text{Ni(MH}_3)_6]^{2+}$  is in outer orbital complex.
- 40. Compare the following complexes with respect to their shape, magnetic behaviours and the hybrid orbitals involved.
  - (a)  $[CoF_6]^{3-}$
  - (b)  $[Cr(NH_3)_6]^{3+}$
  - (c)  $[Fe(CN)_6]^{4-}$

[Atomic Number : Co = 27, Cr = 24, Fe = 26]

- 41. Draw the structure of
  - (a) cis-dichloridotetracyanochromate (II) ion
  - (b) mer-triamminetrichloridocobalt (III)
  - (c) fac-triaquatrinitrito-N-cobalt (III)
- 42. Name the central metal atom/ion present in (a) Chlorophyll (b) Haemoglobin (c) Vitamin B-12. [Ans.: (a) Mg (b) Fe; (c) Co.]
- 43. A metal complex having composition  $Cr(NH_3)_4$   $Cl_2Br$  has been isolated in two forms 'A' and 'B'. The form 'A' reacts with  $AgNO_3$  solution to give white precipitate which is readily soluble in dilute aqueous ammonia, whereas 'B' gives a pale yellow precipitate which is soluble in concentrated ammonia solution. Write the formula of 'A' and 'B'. Also mention the isomerism which arises among 'A' and 'B'.

[Hint: A = [Cr(NH<sub>3</sub>)<sub>4</sub> BrCl] Cl; B = [Cr(NH<sub>3</sub>)<sub>4</sub>Cl<sub>2</sub>] Br are ionisation isomers].

- 44. Write the limitations of Valence Bond Theory.
- 45. Draw a sketch to show the splitting of d-orbitals in an octahedral crystal field state for a d<sup>4</sup> ion. How the actual electronic configuration of the split

d-orbitals in an octahedral crystal field is decided by the relative values of  $\Delta_0$  and pairing energy (P)?

- \*46. For the complex [Fe(en)<sub>2</sub>Cl<sub>2</sub>]Cl identify
  - (a) the oxidation number of iron.
  - (b) the hybrid orbitals and the shape of the complex.
  - (c) the magnetic behaviour of the complex.
  - (d) the number of geometrical isomers.
  - (e) whether there is an optical isomer also?
  - (f) name of the complex. [At. no. of Fe = 26]
- 48. A chloride of fourth group cation in qualitative analysis gives a green coloured complex [A] in aqueous solution which when treated with ethane -1, 2-diamine (en) gives pale yellow solution [B] which on subsequent addition of ethane -1, 2-diamine turns to blue/purple [C] and finally to violet [D]. Identify [A], [B], [C] and [D] complexes,

$$[\textbf{Hint.} : \mathsf{Nickel}, \ [\mathsf{A}] = [\mathsf{Ni}(\mathsf{H}_2\mathsf{O})_6]^{2+} \ ; \qquad [\mathsf{B}] = [\mathsf{Ni}(\mathsf{H}_2\mathsf{O})_4 \ (\mathsf{en})]^{2+} \ ;$$

$$[C] = [Ni(H_2O)_2 (en)_2]^{2+}; [D] = [Ni(en)_3]^{2+}.$$