

#### **CHEMISTRY**

# **BOOKS - CHHAYA CHEMISTRY (BENGALI ENGLISH)**

### COORDINATION COMPOUNDS OR COMPLEX COMPOUNDS

# Example

1. Calculate the overall complex dissociation equilibrium constant for

 $\left[Ag(NH_3)_2\right]^+$  ion, given that  $eta_2$  for this complex is  $1.6 imes 10^7$ .



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Warm Up Exercise

1. What are first order and higher order compounds? Give examples.

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2. Write four differences between double salts and complex salts.
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3. How will you identify the ions present in Mohr salt?
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<b>4.</b> How will you prove that $\left[Fe(CN)_2.4KCN ight]$ is a complex salt?
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<b>5.</b> Classify double salts with examples .
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<b>6.</b> What is primary valency? Why is it known as ionisable valency?
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7. What is secondary valency? What is it known as non-ionisable valency?
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<b>8.</b> Determine the coordination number of Ni and Co in the complexes $\left[Ni(CO)_4 ight]$ and $\left[Co(NH_3)_6 ight]^{3+}$ .
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<b>9.</b> Give an example each of cationic, anionic and electrically neutral coordination entity:
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<b>10.</b> Explain the following terms :
ligand
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11. Explain the following terms :
coordination number
Watch Video Solution
<b>12.</b> Explain the following terms :
central metal atom or ion
Watch Video Solution
<b>13.</b> Explain the following terms : counter ion
Watch Video Solution

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<b>14.</b> Explain the following terms :
coordination sphere.
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<b>15.</b> Mention the coordination entity, ligand, counter ion, oxidation
number of the central metal ion of the coordination compound
[C(NH)]CO
$ig[Cu(NH_3)_4ig]SO_4.$
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46 D. C. C. L. L. C. L. L. C.
<b>16.</b> Distiguish between homoleptic and heteroleptic complexes with one
example of each.
example of each.
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17. Determine the net charge of the coordination entity, comparising of a Co (III)-ion,  $5NH_3$  molecules and  $1Cl^-$  ion.



**18.** Mention the denticity and structure of the following lignads : (i)  $C_5H_5\overset{...}{N}$  (ii)  $\mathrm{acac}^-$  (iii)  $\mathrm{EDTA}^{4-}$  (iv) trien



- 19. Define the following with an example of each.
- (i) Ambident ligand



**20.** Define the following with an example of each.

(ii) Flexidenate ligand



**21.** Define the following with an example of each.

(iii) Chelating ligand



**22.** What is a  $\pi$  - acid ligand ? Give an example.



**23.** What do you mean by Effective Atomic Number (EAN)?

Calculate the EAN of Pt and Fe in the complexes

 $\left[PtCl_{6}\right]^{2-} \ \ {
m and} \ \left[Fe(CO)_{5}\right]$  respectively.



**24.** Give the IUPAC names of the following complexes.

$$K_3ig[Cr(CN)_6ig]$$



**25.** Give the IUPAC names of the following complexes.

$$Na_2igl[Co(NO_2)_6igr]$$



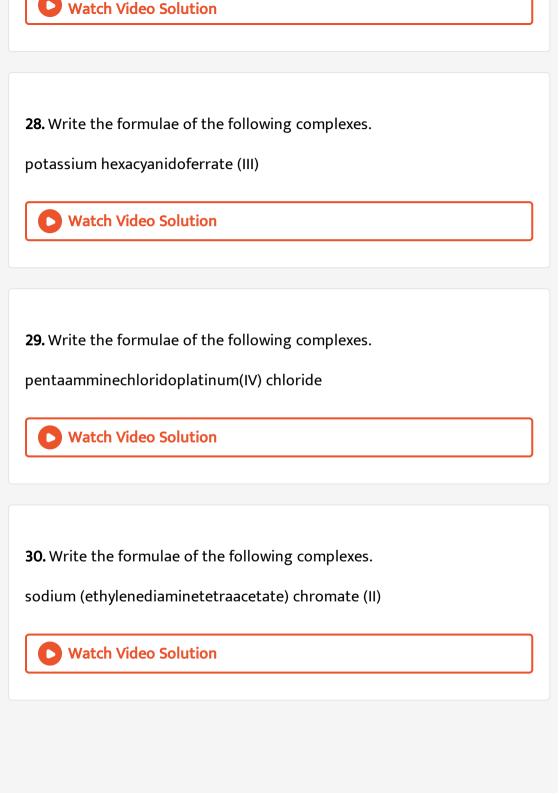
**26.** Give the IUPAC names of the following complexes.

$$\left[ Cr(NH_3)_6 \right] \left[ Co(C_2O_4)_3 \right]$$



**27.** Give the IUPAC names of the following complexes.

 $\big[Co(en)_3\big]Cl_3$ 



31. Write the formulae of the following complexes.

hexakis(methylisocyanide) iron (II) bromide



**32.** Write the formulae of the following complexes.

hexacyanidoplatinic (IV) acid



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Identify the isomerism exhibited by the complexes, 33. (i)  $[Co(NH_3)_5(ONO)]Cl$  and  $[Co(NH_3)_5NO_2]Cl$ . Suggest a method to distinguish between the two compounds.



**34.** Write the ionisation of  $\left[Co(NH_3)_5SO_4\right]Br$ . Why does the compound exhibit ionisation isomerism?



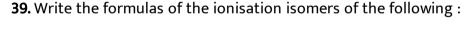
**35.** The tetrahedral compounds of the type  $MA_4, MA_3B, MA_2B_2, MA_2BC$ , do not exhibit geometrical isomerism - justify the statement (M = central metal atom or ion, A, B, C, D = monodenatate ligands).



**36.** What are facial and meridional isomers? Explain with the help of an  $MA_3B_3$  type compound with (A, B = monodenate ligands).



37. Identify the type of isomerism in the following compounds: (i)  $[Co(NH_3)_4Cl_2]6^+$ Watch Video Solution 38. Identify the type of isomerism in the following compounds: (ii)  $\left[Zn(NH_3)_2Cl_2\right]$  (tetrahedral) **Watch Video Solution** 



- (i)  $\left[Ni(en)_2(NO_2)_2\right]Cl_2$ 
  - Watch Video Solution

- **40.** Write the formulas of the ionisation isomers of the following :
- (ii)  $\left[Fe(NH_3)_5CN\right]SO_4$



41. What do you mean by inner orbital and outer orbital complex?



**42.** How do you determine the hybridisation of the central metal ion of a complex from its magnetic moment?



**43.** Discuss the formation of  $\left[Ni(NH_3)_4
ight]^{2+}$  with the help of valence bond theory.



**44.** Discuss the formation of  $\left[FeF_6\right]^{3-}$  with the help of valence bond theory.



**45.** Write the limitations of valence bond theory.



**46.** Why do  $Co^{2+}$ ,  $Ni^{2+}$ ,  $Cu^{2+}$ ,  $Zn^{2+}$  always form outer orbital octahedral complexes?



**47.** What is crystal field splitting? How do the d-orbitals split under the influence of an octahedral field?



**48.** What do you mean by crystal field stabilisation energy (CFSE)? Calculate the value of CFSE for a  $d^4$  metal ion in presnce of weak - field ligand in octahedral complex.



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49. How d-orbitals undergo splitting in a tetrahedral crystal field?

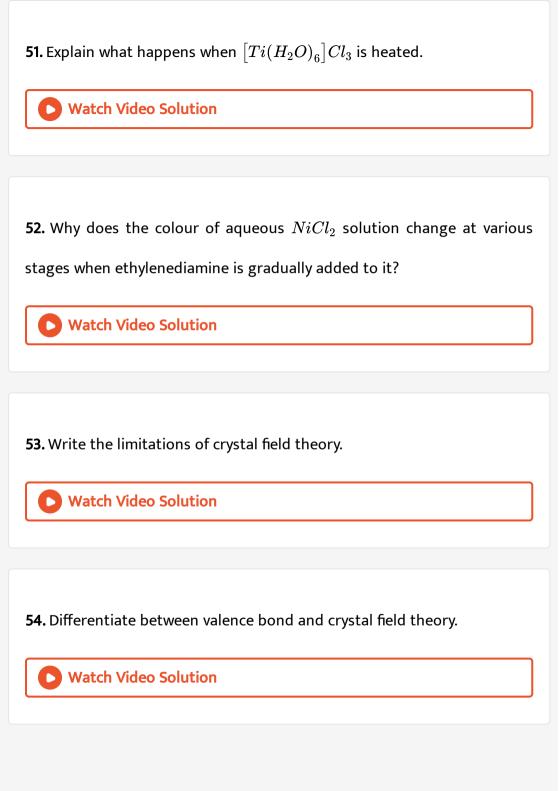


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**50.** What do you mean by spectrochemical series? Arrange the ligands of the following complexes in order of their crystal field spiltting ability:  $[CrCl_6]^{3-}$ ,  $[Cr(H_2O)_6]^{3+}$ ,  $[Cr(CN)_6]^{3-}$  and  $[Cr(NH_3)_6]^{3+}$ 



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**55.** Write the relation between overall and stepwise stability constants of coordination compounds formed through 1, 2, 3 and 4 steps.



**56.** Why is the stability of  $\left[Ni(en)_3
ight]^{2+}$  more than that of  $\left[Ni(NH_3)_6
ight]^{2+}$ ?

57. Arrange the following in increasing order of conductivity.



 $[Co(NH_3)_3Cl_3], [Co(NH_3)_5C, ]Cl_2, [Co(NH_3)_5]Cl_2 \text{ and } [Co(NH_3)_4Cl_2]$ 

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**58.** Name bidentate lignads with (i) two neutral donor groups and (ii) two acid groups.

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<b>59.</b> What are inner metallic complexes? Give examples.
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<b>60.</b> Mention four applications of chelate formation.
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<b>61.</b> Explain the role of coordination compounds in analytical chemistry,
with the help of examples.
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<b>62.</b> Name the metal ions present in chlorophyll, haemoglobin, vitamin ${\cal B}_{12}$ and cis-platin.

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<b>63.</b> What is calgon? Water is it used?
Watch Video Solution
<b>64.</b> Give two examples of metal carbonyls.
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<b>65.</b> What do you mean by synerigic effect?
Watch Video Solution
<b>66.</b> How does hypo remove unreacted $AgBr$ from photographic films?
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- **1.** Give the coordination number of Fe in  $\left[Fe(EDTA)\right]^-$ .
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- **2.** Give an example of a neutral coordination compound in which the central metal atom is  $sp^3d^2$  hybridised.
  - Watch Video Solution

- - Watch Video Solution

**3.** Name the isomer(s) of  $[Pt(SCN)(NH_3)_3]$  SCN.

- **4.** Write the IUPAC name for the coordination isomer o  $\left[ Cr(NH_3)_6 \right] \left[ Co(CN)_6 \right].$ 
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<b>5.</b> Calculate the oxidation number of cobalt in the complex, $\left[Co(NH_3)_2(NO_2)Cl\right]\left[Au(CN)_2\right].$
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<b>6.</b> Which of the following cannot act as a ligand? $NH_3,CH_4,CO,H_2O$ Watch Video Solution
7. Write the metals present in the following compounds:  (1) Chlorophyll
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8. Write the metals present in the following compounds:

(2) cis- platin

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- $\boldsymbol{9.}$  Write the metals present in the following compounds :
- (3) Haemoglobin



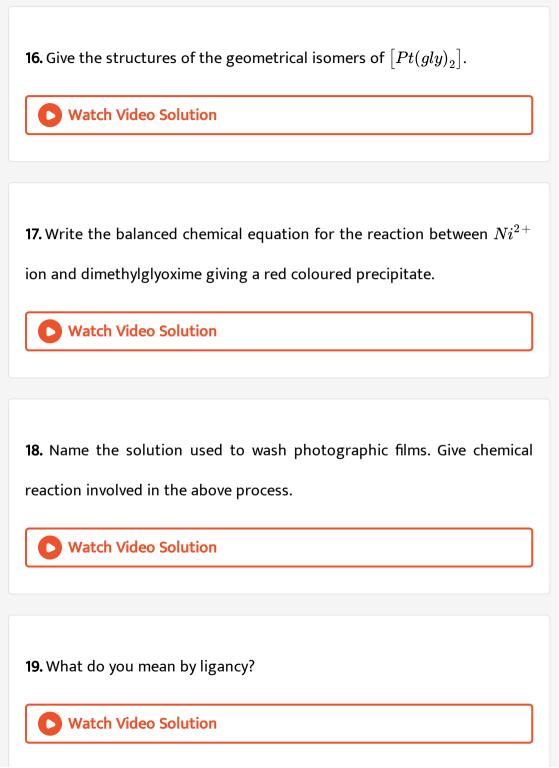
- 10. Write the metals present in the following compounds:
- (4) Vit.  $B_{12}$ .



11. What is the coordination number of the metal ion in an octahedral complex ? Why does  $NH_3$  take part in complex formation but not  $NH_4^+$ ?



12. Is the complex formation an exothermic or an endothermic process? Give reasons. **Watch Video Solution** tempertaure affect the stability of coordination **13.** How does compounds? **Watch Video Solution** 14. Mention any two important conditions that help in the formation of a metal - complex. **Watch Video Solution** 15. Give the name and structure of an unsymmetrical bidentate ligand. **Watch Video Solution** 



- **20.** In the inner orbital complex ion,  $[Ni(CN_4)]^{2-}$ , give:
- (1) the type of hybridisation in  $Ni^{2\,+}$
- (2) the orbitals involved in hybridisation.



**21.** What do you mean by crystal field splitting energy in tetrahedral and octahedral complexes?



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- **22.**  $\left[NiCl_4\right]^2$  is an outer orbital complex. Give:
- (1) the type of hybridisation in  $Ni^{2+}$
- (2) the orbitals involved in hybridisation.



23. Write the name and structure of a the ligand that is used to determine the hardness of water (i.e., in the qualitative analysis of  $Ca^{2+}$ and  $Mq^{2+}$  ).



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**24.**  $\left[Ni(NH)_4\right]^{2-}$  is diamagnetic but  $\left[NiCl_4\right]^{2-}$  is paramagnetic. Give the hybridisation of  $Ni^{2+}$  in the given complex ions.



25. Arrange the given ligands in order of their ligand field strength.

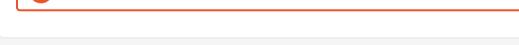
$$\left[CrCl_{6}
ight]^{3-},\left[Cr(H_{2}O_{6}]^{3+},\left[Cr(CN_{6})
ight]^{3-},\left[Cr(NH_{3})_{6}
ight]^{3+}$$



26. Which of the following exhibits geometrical isomerism? (1) linear (2) square planar (3) tetrahedral (4) octahedral

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27. What is the basis of spectrochemical series?



**28.** Write the names of the coordination entities that are used as electrolytes in electroplating with gold & silver.



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**29.** Why are most of the coordination compounds of transition elements paramagnetic in nature?



**30.** A transition metal M and ligand L forms a coordination compound  $ML_4$ . Express the overall stability constant of the system,  $\beta_4$ , in terms of concentration of its various components.



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31. What do you mean by labile complex?



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**32.** The reaction of sulphide ion with nitroprusside ion gives a violet complex. Write the balanced equation for the given reaction.



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**33.** How can you identify cis -  $\left[Pt(NH_3)_2Cl_2\right]$  from trans -  $\left[Pt(NH_3)_2Cl_2\right]$ .



**34.** Give an example each of symmetrical and unsymmetrical bidentate ligand.



**35.** Consider the following equilibria along with the respective stability constants:

$$Aq^{+} + NH_{3} \Leftrightarrow [Aq(NH_{2})]^{+}, K_{1} = 3.5 \times 10^{3}$$

$$\left[Ag(NH_3)
ight]^+ + NH_3 \Leftrightarrow \left\lceil Ag(NH_3)_2 
ight
ceil^+,$$

$$K_2 = 1.7 \times 10^3$$

Calculate the overal formiation constant of  $\left[Ag(NH_3)_2
ight]^+$  .



## Question Answer Zone For Board Examination Short Answer Type

1. Why do the transition elements form complexes?



**2.** The square planar complexes with coordination number 4  $(MA_2B_2 \ {
m or} \ MA_2BC)$  show geometrical isomerism but tetrahedral complexes do not. Explain.



**3.** With the help of suitable tests, distinguish between  $\left[Co(NH_3)_5Br\right]SO_4$  and  $\left[Co(NH_3)_5SO_4\right]Br$ . What type of isomerism is shown by these compounds?



**4.** Though the salts of s - and p - block elements are colourless, why are the salts of transition elements coloured?



**5.** Calculate the oxidation number of the central metal atoms or ions : (any 4): (1)  $\left[Co(NH_3)_4Cl(NO_2)\right]^+$ 



**6.** Calculate the oxidation number of the central metal atoms or ions :





**7.** Calculate the oxidation number of the central metal atoms or ions : (any 4): (3)  $\left[Cr(C_2O_4)_3\right]^{3-}$ 

- **8.** Calculate the oxidation number of the central metal atoms or ions :
- (any 4): (4)  $\left[CoCl_2(en)_2
  ight]^+$ 
  - View Text Solution

- **9.** Calculate the oxidation number of the central metal atoms or ions :
- (any 4): (5)  $\left[Co(NH_3)_5(ONO)\right]^{2+}$ 
  - Watch Video Solution

- 10. Write the type of hybridisation in  $\left[Cr(CO)_6\right]$ . Also explain its structure and magnetic property.
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11. Give the geometrical isomers of  $[Pt(NH_3)_2Cl_2]$ .

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12. Write the formula of potassiumtrioxalatoferrate (III).

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**13.** Calculate the Effective Atomic number of Mn is  $Mn_2(CO)_{10}$ .



**14.** If a coordination compound,  $CoCl_3.4NH_3$  is heated, it does not give out  $NH_3$ , while if the same reacts with  $AgNO_3$  solution, gives a white precipitate of AgCl. Write the structure and IUPAC name of the compound.



**15.**  $\left[M(\mathrm{AA})_2X_2\right]^{n+}$  is an optically active complex ion. What is its geometry? Give an example.



**16.** How do the transition elements form  $\pi$  – complexes ?



17. Why can't there be any low spin tetrahedral complex?



**18.** Calculate the volume of 0.1M  $AgNO_3$  solution required for complete precipitation of chloride ions present in 30 mL of 0.01M solution of  $\left[Cr(H_2O)_6\right]Cl_3$ .

A. Millimoles of  $igl[Cr(H_2O)_6igr]Cl_3$ 

 $= ext{volume (in mL)} imes ext{molarity} = 30 imes 0.01 = 0.3$ 

 $\therefore$  Millimoles of  $Cl^-$  in

 $ig[ {\it Cr}(H_2O)_6 ig] {\it Cl}_3 = 3 imes ext{millimoles of } ig[ {\it Cr}(H_2O)_6 ig] {\it Cl}_3$ 

=3 imes0.3=0.9

So, millimoles of  $Ag^{\,+}\,$  ion required = millimoles of  $Cl^{\,-}\,$  ion present

in  $\left[Cr(H_2O)_6\right]Cl_3=0.9$ 

But millimoles of  $Ag^+$  ion = volume of

 $AgNO_3$  solution (in mL) imes molarity of  $AgNO_3$  solution

 $\therefore 0.9 = ext{volume of} \ AgNO_3 \ ext{Soluion (in mL)} imes 0.1$ 

Hence volume of  $AgNO_3$  solution (in mL)  $= rac{0.9}{0.1} = 9$ 

В.

C.

D.

Answer:

**19.** Why do the following transition metal ions appear colourless ?  $Cu^+, Ag^+, Zn^{2+}, Hg^{2+}$  and  $Cd^{2+}.$ 

A. From the electronic configurations of the metal atoms,  $Cu\big(3d^{10}4s^1\big),\,Ag\big(4d^{10}5s^1\big),\,Zn\big(3d^{10}4s^2\big),\,Hg\big(5d^{10}6s^2\big)\qquad\text{and}$   $Cd\big(4d^{10}5s^2\big),\,\text{we can see that the given metal ions do not have}$  unpaired electrons in their (n-1)d orbital and the orbitals are completely filled  $(d^{10})$  with electrons :

Hence, these electrons do not undergo d-d transition by absorbing light from visible spectrum and therefore the given metal ions appear colourless.

 $Cu^{+}\left(3d^{10}
ight), Ag^{+}\left(4d^{10}
ight), Zn^{2+}\left(3d^{10}
ight), Hg^{2+}\left(5d^{10}
ight) ext{ and } Cd^{2+}\left(4d^{10}
ight)$ 

В.

C.

_	
U.	

### **Answer:**



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**20.** Explain which of the following compounds has effective atomic number of central metal atom in accordance with Sidwick theory?  $(1) \ K_3[Fe(CN_6)] \ (2) \ K_4[Fe(CN)_6]$ 



**21.** Why does the paramagnetic nature of the 3d transition series increase till Cr and then decrease regularly?



22. Calculate the 'spin-only' magnetic moment of the tetrahedral complex,

 $H_g^n[Co(SCN)_4].$ 

 $\therefore$  Electronic configuration of  $Co^{2+}$  ion :



 $Co^{2+}$  undergoes  $sp^{3}$  – hybridisation and forms a tetrahedral complex.

In this case, no pairing of electrons takes place in the 3d orbitals. Hence, the given complex has 3 unpaired electrons.

$$\therefore$$
 Magnetic moment,  $\mu = \sqrt{n(n+2)}$ 

$$=\sqrt{3(3+2)}=\sqrt{15}=3.87BM$$





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**24.** The stability (or formation) constant  $\left(K_f\right)$  for  $\left\lceil Ag(NH_3)_2 \right
ceil^+$  is  $1.6 imes10^7$ . Calculate the ratio of the concentrations of  $\left[Ag(NH_3)_2
ight]^+$ 

**23.** Show the structures of the isomers of  $\lceil Co(NH_3)_3Cl_3 \rceil$ .

and  $Ag^{\,+}$  in 0.12M  $NH_3$  solution.



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### Solved Wbchse Scanner 2015

**1.** Which of the following complex ions has no 'd' electron(s) in the central metal atom -

A. 
$$\left[Cr(H_2O)_6
ight]^{3+}$$

B. 
$$\left[Co(NH_3)_5Cl
ight]^{2+}$$

C. 
$$\left[Fe(CN)_6
ight]^{3}$$

D. 
$$[MnO_4]^-$$

### Answer: d



**2.** Which of the following is the correct electronic configuration of Ni in  $Ni(CO)_{\mbox{\tiny A}} ext{-}$ 

A.  $[Ar]3d^84s^2$ 

B.  $[Ar]3d^{10}4s^0$ 

C.  $[Ar]3d^84s^0$ 

D.  $[Ar]3d^94s^1$ 

### Answer:



### Solved Wbchse Scanner 2016

1. An aqueous solution of a complex compound of formula  $Co(NH_3)_5 Br(SO_4)$  reacts readily with aqueous  $AgNO_3$  to give a yellowish white precipitate. Write down the structural formula of the complex and mention the reaction involved.



### **Solved Wbchse Scanner 2017**

1. When tetraamminechloridonitrito	cobalt (	III) nitrate	is	dissolved	in	
water, how many ions will be formed from one molecule -						

- A. 4
- B. 3
- C. 2
- D. 0

### **Answer: C**



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**2.** Explain why  $\left[Cu(NH_3)_4
ight]^{2+}$  is coloured but  $\left[Cu(CN)_4
ight]^{3-}$  is colourless.

## **Solved Wbchse Scanner 2018**

- is the oxidation number of the central metal  $\left[Cr(NH_3)_4(NO_2)Cl
  ight]$  (atomic no. of Cr = 24 ) -
  - A. 0
  - B. + 1
  - D. + 2

 $\mathsf{C.} + 3$ 

Answer: d



- **2.** How many isomers are possible for  $\left[Co(NH_3)_4Cl_2\right]Cl$ ?

# Solved Cbse Scanner Delhi 2014

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- 1. Give the formula of the following coordination entities:
- (a)  $Co^{3\,+}$  ion is bound to one  $Cl^{\,-}$  , one  $NH_3$  molecule and two bidentate ethylene diamine (en) molecules.



- 2. Give the formula of the following coordination entities:
- (b)  $Ni^{2\,+}$  ion is bound to two water molecules and two oxalate ions.

Write the name and magnetic behaviour of each of the above coordination entities.



**1.** Write IUPAC name of the complex  $\left[Cr(NH_3)_4Cl_2\right]Cl$ . What type of isomerism is exhibited by the complex  $\left[Co(en)_3\right]^{3+}$  (en = ethane - 1, 2 - diamine) ? Why is  $\left[NiCl_4\right]^{2-}$  paramagnetic but  $\left[Ni(CO)_4\right]$  is diamagnetic ? (Atomic no : Cr = 24, Co = 27, Ni = 28)



### Solved Cbse Scanner Delhi 2015

**1.** What is the IUPAC name of  $[Ni(NH_3)_6]Cl_2$ ?



2. Indicate the types of isomerism exhibited by the following complexes :

(i) 
$$\left[Co(NH_3)_5(NO_2)\right]^{2+}$$



- 3. Indicate the types of isomerism exhibited by the following complexes : (ii)  $\lceil Co(en)_3 \rceil Cl_3$  (en = ethylene diamine)
  - Watch Video Solution
- **4.** Indicate the types of isomerism exhibited by the following complexes : (iii)  $\lceil Pt(NH_3)_2Cl_2 \rceil$ 
  - Watch Video Solution

# Solved Cbse Scanner Outside Delhi 2015

- **1.** What is meant by chelate effect ?
  - Watch Video Solution

- **2.** Write the name of the following : (i)  $\left[Co(NH_3)_6
  ight]Cl_3$ 
  - Watch Video Solution

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3. Write the IUPAC name of the following:

- (i)  $\left[Co(NH_3)_6\right]Cl_3$
- (ii)  $\left[NiCl_4
  ight]^{2-}$
- (iii)  $K_3igl[Fe(CN)_6igr]$



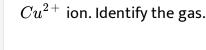
**4.** Write the IUPAC names of the following coordination compounds :

$$K_3[Fe(CN)_6]$$



### Solved Cbse Scanner Delhi 2016

1. On adding NaOH to ammonium sulphate, a colourless gas with pungent odour is evolved which forms a blue coloured complex with





**2.** When a coordination compound  $CrCl_3.6H_2O$  is mixed with  $AgNO_3$ , 2 moles of AgCl are precipitated per mole of the compound. Write - (i) structural formula of the complex, (ii) IUPAC name of complex.



**3.** For the complex  $\left[Fe(CN)_6\right]^{3-}$ , write the hybridization type, magnetic character and spin nature of the complex. (At. Number : Fe = 26)



**4.** Draw one of the geometrical isomers of the complex  $\left[Pt(en)_2Cl_2\right]^{2+}$  which is optically active.



### Solved Cbse Scanner East Zone 2016

**1.** When a coordination compound  $PtCl_4.6NH_3$  is mixed with  $AgNO_3$ , 4 moles of AgCl are precipitated per mole of the compound. Write - (i) structural formula of the complex



**2.** When a coordination compound  $PtCl_4.6NH_3$  is mixed with  $AgNO_3$ , 4 moles of AgCl are precipitated per mole of the compound. Write - (ii) IUPAC name of the complex.



**3.** For the complex  $[CoF_6]^{3-}$ , write the hybridization type, magnetic character and spin nature of the complex (At. Number : Co = 27)



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**4.** Why is the complex  $\left[Co(en)_3
ight]^{3+}$  more stable than the complex

 $[CoF_6]^{3-}$ ?



### Solved Cbse Scanner Delhi 2017

1. What type of isomerism is shwon by the complex  $\lceil Co(NH_3)_6 \rceil \lceil Cr(CN)_6 \rceil$ ?

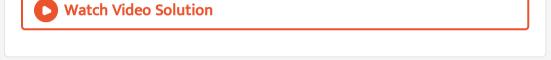


**2.** Why a solution of  $\left[Ni(H_2O)_6\right]^{2+}$  is green while a solution of  $\left[Ni(CN)_4\right]^{2-}$  is colourless? (At. No. of Ni = 28)



<b>3.</b> Write IUPAC name of the following complex : $\left[Co(NH_3)_5(CO_3)\right]Cl.$	
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**4.** What type of isomerism is shown by the complex  $\left[Co(en)_3\right]Cl$ ?



**5.** Write the hybridisation and magnetic character of  $\left[Co(C_2O_4)_3
ight]^3-$  . (At.

No. of Co = 27)



**6.** Write IUPAC name of the following complex :  $\left[Cr(NH_3)_3Cl_3\right]$ .



<b>1.</b> Using IUPAC norms write the formulae for the following :
--

- (i) Sodium dicyanidoaurate (I)
  - Watch Video Solution

- **2.** Using IUPAC norms write the formulae for the following:
- (ii) Tetraamminechloridonitrito N platinum (IV) sulphate
  - Watch Video Solution

- **3.** What type of isomerism is shown by the complex  $Co(NH_3)_5(SCN)^{2+}$  ?
  - Watch Video Solution

<b>4.</b> Why is $\left[NiCl_4 ight]^{2-}$ paramagnetic while $\left[Ni(CN_4) ight]^{2-}$ is diamagnetic ?
(At. No. of Ni = 28)
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5. Using IUPAC norms write the formulae for the following :
(i) Tri(ethane - 1, 2 - diamine) chromium (III) chloride
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<b>6.</b> Using IUPAC norms write the formulae for the following :
(ii) Potassium tetrahydroxozincate (II)
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7. Using IUPAC norms write the formulae for the following :
(i) Potassium trioxalatoaluminate (III)



- **8.** Using IUPAC norms write the formulae for the following :
- (i) Dichloridobis (ethane 1, 2 diamine) cobalt (III)
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### Solved Cbse Scanner All India 2018

**1.** Write the coordination number and oxidation state of platinum in the complex  $\lceil Pt(en)_2Cl_2 \rceil$ ?



- 2. Write the formulas for the following compounds:
- (6) Iron (III) hexacyanoferrate (II)
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3. What type of isomerism is exhibited by the complex

 $[Co(NH_3)_5Cl]SO_4$ ?



**4.** Write the hybridisation and number of unpaired electrons in the complex  $\left[CoF_6\right]^{3-}$ . (At. No. of Co = 27)



### Solved Ncert Textbook Problems Ncert Intext Questions

**1.** On the basis of the following observation made with aqueous solution, assign secondary valencies to metals in the following compounds :





2. Write the formulas for the following compounds -(1) Tetraammineaquachloridocobalt (III) chloride **Watch Video Solution** 3. Write the formulas for the following compounds -(2) Potassium tetrahedroxozincate (II) **Watch Video Solution** 4. Write the formulas for the following compounds -(3) Potassium trioxalatoaluminate (III) **Watch Video Solution** 5. Write the formulas for the following compounds -(4) Dichloridobis (ethane - 1, 2 - diamine) cobalt (III) Watch Video Solutio

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<b>6.</b> Write the formulas for the following compounds -
(5) Tetracarbonylnickel (0)
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7. Write the IUPAC names of the following coordination compounds:
(1) $\left[Pt(NH_3)_2Cl(NO_2) ight]$
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 ${\bf 8.}$  Write the IUPAC names of the following coordination compounds :

(2)  $K_3ig[Cr(C_2O_4)_3ig]$ 

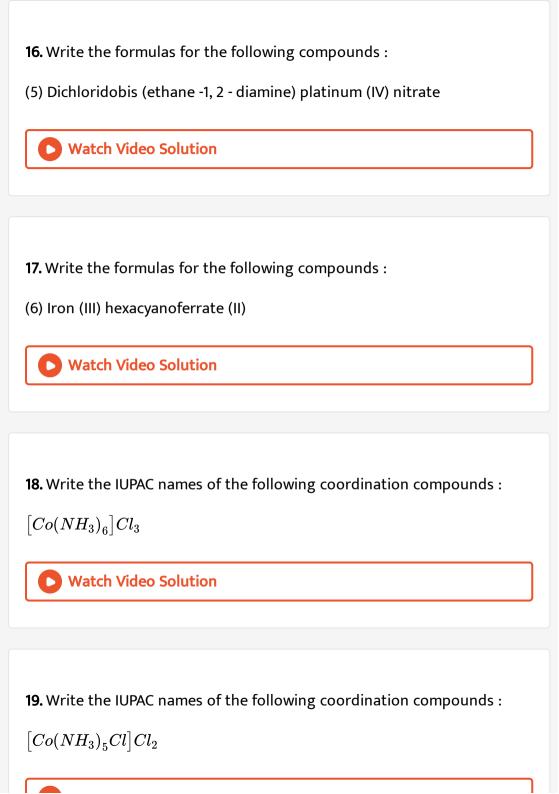
- - **11.** Write the IUPAC names of the following coordination compounds :

$$Hgigl[Co(SCN)_4igr]$$



- **12.** Write the formulas for the following compounds :
- (1) Tetraamminediaquacobalt (III) chloride

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<b>13.</b> Write the formulas for the following compounds:
(2) Potassium tetracyanonickelate (II)
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14. Write the formulas for the following compounds :
(3) Tris (ethane - 1, 2 - diamine) chromium (III) chloride
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<b>15.</b> Write the formulas for the following compounds:
(4) Amminebromidochloridonitrito - N - platinate (II)
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**20.** Write the IUPAC names of the following coordination compounds :

$$K_3ig[Fe(CN)_6ig]$$



21. Write the IUPAC names of the following coordination compounds:

 $K_3[Fe(C_2O_4)_3]$ 



**22.** Write the IUPAC names of the following coordination compounds :

$$K_2[PdCl_4]$$



23. Write the IUPAC names of the following coordination compounds:

 $\lceil Pt(CN_3)_2Cl(NH_2CH_3) \rceil Cl$  itrate

(6) Iron (III) hexacyanoferrate (II)



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24. Why is geometrical isomerism not possible in tetrahedral complexes having two different types of uni - dentate ligands coordinated with the central metal ion?



**25.** Draw structure of geometrical isomers of  $\lceil Fe(NH_3)_2(CN)_4 \rceil \rceil^-$ 



26. Out of the following two coordination entities which is chiral (1)  $cis - \left[CrCl_2(ox)_2\right]^{3-}$ (optically active)? (2)

 $trans - \left[ CrCl_2(ox)_2 \right]^{3-}$ 



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27. Indicate the types of isomerism exhibited by the following complexes and draw the structures for these isomers:

 $K[Cr(H_2O)_2(C_2O_4)_2]$ 



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28. Indicate the types of isomerism exhibited by the following complexes and draw the structures for these isomers:

 $[Co(en)_3]Cl_3$ 



**29.** Indicate the types of isomerism exhibited by the following complexes and draw the structures for these isomers :

 $[Co(NH_3)_5(NO_2)](NO_3)_2$ 



**30.** Indicate the types of isomerism exhibited by the following complexes and draw the structures for these isomers :

 $[Pt(NH_3)(H_2O)Cl_2]$ 



**31.** Give evidence that  $\left[Co(NH_3)_5Cl\right]SO_4$  &  $\left[Co(NH_3)_5SO_5\right]Cl$  are ionisation isomers.



**32.** The spin only magnetic moment of  $\left[MnBr_4\right]^{2-}$  is 5.9 BM. Predict the geometry of the complex ion.



**33.** Explain on the basis of valence bond theory that  $\left[Ni(CN)_4\right]^{2-}$  ion with square planar structure is diamagnetic and the  $\left[NiCl_4\right]^{2-}$  ion with tetrahedral geometry is paramagnetic.



**34.**  $\left[Fe(H_2O)_6\right]^{3+}$  is strongly paramagnetic whereas  $\left[Fe(CN)_6\right]^{3-}$  is weakly paramagnetic. Explain.



**35.**  $[NiCl_4]^{2-}$  is paramagnetic while  $\left[Ni(CO)_4\right]$  is diamagnetic though both are tetrahedral. Why?



**36.** Explain  $\left[Co(NH_3)_6\right]^{3+}$  is an inner orbital complex whereas  $\left[Ni(NH_3)_6\right]^{2+}$  is an outer oubital complex.



**37.** Predict the number of unpaired electrons in the square planar  $\left[Pt(CN)_4\right]^{2-}$  ion.



**38.** Hexaaquamanganese (II) ion contains five unpaired electrons, while the hexacyano ion contains only one unpaired electron. Explain using

Crystal Field Theory.



**39.** Calculate the overall complex dissociation equilibrium constant for the  $\left[Cu(NH_3)_4\right]^{2+}$  ion, given that  $eta_4$  for this complex is  $2.1 imes 10^{13}$ .

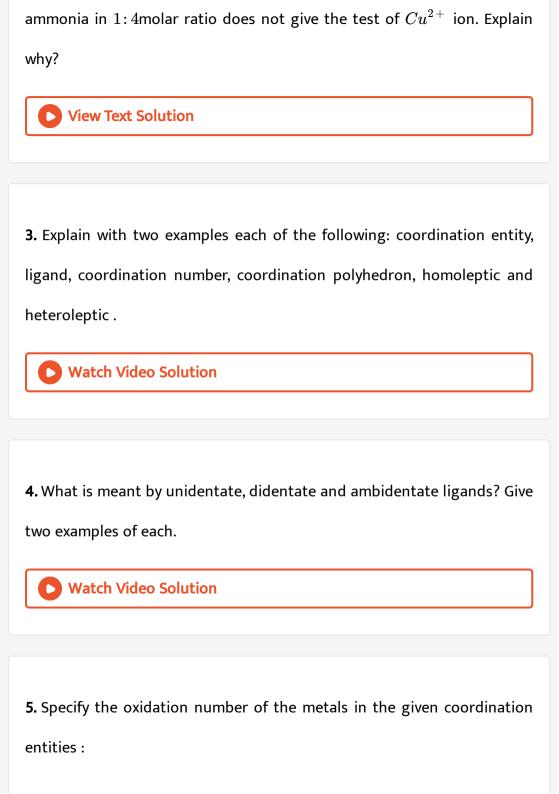


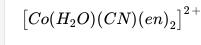
# Solved Ncert Textbook Problems Ncert Exercise Questions

**1.** Explain the bonding in coordination compounds in terms of Werner's postulates.



**2.**  $FeSO_4$  solution mixed with  $(NH_4)_2SO_4$  solution in  $1\colon 1$  molar ratio gives the test of  $Fe^{2+}$  ion but  $CuSO_4$  solution mixed with aqueous







6. Specify the oxidation number of the metals in the given coordination entities:

$$\left[ CoBr(En)_2 \right]^{2+}$$



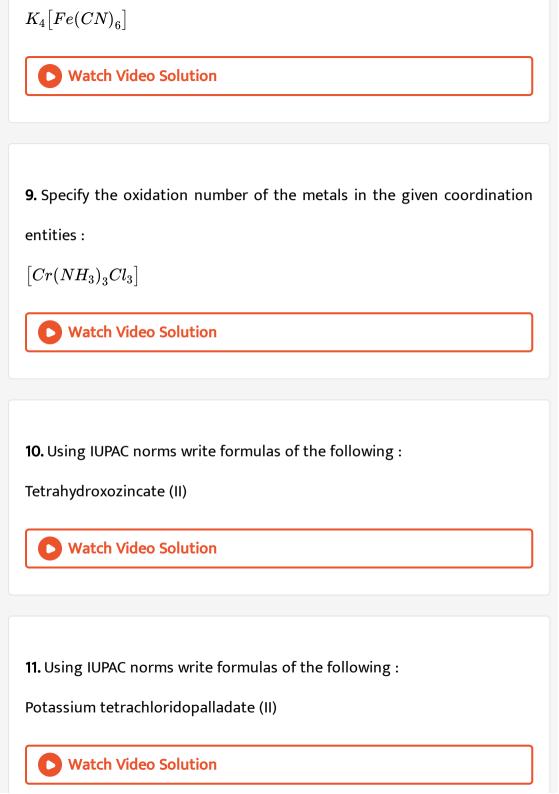
7. Specify the oxidation number of the metals in the given coordination entities:

 $[PtCl_4]^{2-}$ 



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8. Specify the oxidation number of the metals in the given coordination entities:



12. Using IUPAC norms write formulas of the following: Diamminedichloridoplatinum (II) **Watch Video Solution** 13. Using IUPAC norms write formulas of the following: Potassium tetracyanonickelate (II) **Watch Video Solution** 14. Using IUPAC norms write formulas of the following: Pentaamminenitrito - O - cobalt (III)

15. Using IUPAC norms write formulas of the following: Hexaamminecobalt (III) sulphate **Watch Video Solution** 16. Using IUPAC norms write formulas of the following: Potassium tri(oxalato) chromate (III) **Watch Video Solution** 17. Using IUPAC norms write formulas of the following: Hexaammineplatinum (IV) **Watch Video Solution** 18. Using IUPAC norms write formulas of the following: Tetrabromidocuprate (II)

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19. Using IUPAC norms write formulas of the following:

Pentaamminenitrito - N - cobalt (III)



20. Using IUPAC norms write systematic names of:

 $\left[ Co(NH_3)_6 \right] Cl_3$ 



21. Using IUPAC norms write systematic names of:

 $[Pt(NH_3)_2Cl(NH_2CH_3)]Cl$ 



**22.** Using IUPAC norms write systematic names of :

$$\big[Ti(H_2O)_6\big]^{3\,+}$$



**23.** Using IUPAC norms write systematic names of :

$$\left[Co(NH_3)_4Cl(NH_2)\right]Cl$$



**24.** Using IUPAC norms write systematic names of :

$$\big[Mn(H_2O)_6\big]^{2\,+}$$



**25.** Using IUPAC norms write systematic names of :

 $[NiCl_4]^{2\,-}$ 



**26.** Using IUPAC norms write systematic names of :

$$\big[Ni(NH_3)_6\big]Cl_2$$



# **27.** Using IUPAC norms write systematic names of : $\left[Co(en)_3 ight]^{3+}$



**28.** Using IUPAC norms write systematic names of :

 $\left[Ni(CO)_4\right]$ 



29. List various types of isomerism possible for coordination compounds, giving an example of each.



30. How many geometrical isomers are possible in the following coordination entities?

$$\big[Cr(C_2O_4)_3\big]^{3\,-}$$



31. How many geometrical isomers are possible in the following coordination entities?

 $[Co(NH_3)_3Cl_3]$ 



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32. Draw the structures of optical isomers of:

$$\left[Cr(C_2O_4)_3\right]^{3\,-}$$



33. Draw the structures of optical isomers of:

 $\big[PtCl_2(en)_2\big]^{2\,+}$ 



34. Draw the structures of optical isomers of:  $\left[ Cr(NH_3)_2 Cl_2(en) 
ight]^+$ 



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35. Draw all the isomers (geometrical and optical) of:

 $ig[ CoCl_2(en)_2 ig]^+$ 

**36.** Draw all the isomers (geometrical and optical ) of :

$$\big[Co(NH_3)Cl(en)_2\big]^{2\,+}$$



**37.** Draw all the isomers (geometrical and optical ) of :

$$\left[ Co(NH_3)_2 Cl_2(en) \right]^+$$



**38.** Write all the geometrical isomers of  $[Pt(NH_3)(Br)(Cl)(py)]$  and how many of these will exhibit optical isomerism?



**39.** Aqueous copper sulphate solution (blue) gives - (1) a green precipitate with aqueous potassium fluoride



**40.** Aqueous copper sulphate solution (blue) gives - (2) a bright green solution with aqueous potassium chloride. Explain these experimental results.



**41.** What is the coordination entity formed when excess of aqueous KCN is added to aqueous solution of  $CuSO_4$ ? Why is it that no precipitate of copper sulphide is obtained when  $H_2S(g)$  is passed through this solution?



**42.** Discuss the nature of bonding in the given coordination entities on the basis of valence bond theory:

 $\left\lceil Fe(CN)_6 
ight
ceil^{4-}$ 



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**43.** Discuss the nature of bonding in the given coordination entities on the basis of valence bond theory:

 $\left[FeF_{6}
ight]^{3}$ 



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**44.** Discuss the nature of bonding in the given coordination entities on the basis of valence bond theory:

 $\big[ Co(C_2O_4)_3 \big]^{3\,-}$ 



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**45.** Discuss the nature of bonding in the given coordination entities on the basis of valence bond theory:

 $[CoF_6]^{3}$ 



**46.** Draw figure to show the splitting of d - orbitals in an octahedral crystal field.



- **47.** What is spectrochemical series? Explain the difference between a weak
- field ligand and a strong field ligand.



**48.** What is crystal field splitting energy ? How does the magnitude of  $\Delta_0$  decide the actual configuration of d orbitals in a coordination entity?



**49.**  $\left[Cr(NH_3)_6
ight]^{3+}$  is paramagnetic while  $\left[Ni(CN)_4
ight]^{2-}$  is diamagnetic.

Explain why?



**50.** A solution of  $\left[Ni(H_2O)_6\right]^{2+}$  is green but a solution of  $\left[Ni(CN)_4\right]^{2-}$  is colourless. Explain.



**51.**  $\left[Fe(CN)_6\right]^{4-}$  and  $\left[Fe(H_2O)_6\right]^{2+}$  are of different coloures in dillute solutions. Why ?

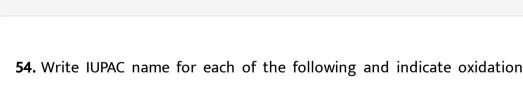
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**52.** Discuss the nature of bonding in metal carbonyls.



53. Write IUPAC name for each of the following and indicate oxidation state, electronic configuration and coordination number. Also give stereochemistry, magnetic moment:

 $K[Cr(H_2O)_2(C_2O_4)_2].3H_2O$ 



state, electronic configuration and coordination number. Also give

 $[Co(NH_3)_5Cl]Cl_2$ 



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stereochemistry, magnetic moment:

**55.** Write IUPAC name for each of the following and indicate oxidation state, electronic configuration and coordination number. Also give stereochemistry, magnetic moment :

 $CrCl_3(py)_3$ 



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**56.** Write IUPAC name for each of the following and indicate oxidation state, electronic configuration and coordination number. Also give stereochemistry, magnetic moment :

 $Cs[FeCl_4]$ 



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**57.** Write IUPAC name for each of the following and indicate oxidation state, electronic configuration and coordination number. Also give

 $stere ochem is try, \, magnetic \, \, moment: \, \,$ 

 $K_4[Mn(CN)_6]$ 



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**58.** Give oxidation state, d - orbital occupation and coordination number of the central metal ion in the given complexes :

- (1)  $K_3igl[Co(C_2O_4)_3igr]$
- (2)  $cis-\left[Cr(en)_2Cl_2\right]Cl$
- (3)  $(NH_4)_2[CoF_4]$
- (4)  $[Mn(H_2O)_6]SO_4$



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**59.** What is meant by stability of a coordination compound in solution? State the factors which govern stability of complexes.



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60. What is meant by chelate effect? Given example. Watch Video Solution 61. Discuss briefly giving an example in each case the role of coordination compounds in: (1) biological systems **Watch Video Solution 62.** Discuss briefly giving an example in each case the role of coordination compounds in: (2) medicinal chemistry **Watch Video Solution** 63. Discuss briefly giving an example in each case the role of coordination compounds in: (3) analytical chemistry **Watch Video Solution** 

**64.** Discuss briefly giving an example in each case the role of coordination compounds in : (4) extraction/metallurgy of metals.



**65.** How many ions are produced from the complex  $\left[Co(NH_3)_6\right]Cl_2$  in solution : 6, 4, 3 or 2



**66.** Amongest the following ions which one has the highest magnetic value ?

$$ig[Zn(H_2O)_6ig]^{2+}$$
, $ig[Fe(H_2O)_6ig]^{2+}$ , $ig[Cr(H_2O)_6ig]^{3+}$ 



67. Amongest the following ions which one has the highest magnetic

value?

 $igl[Zn(H_2O)_6igr]^{2+}$  ,  $igl[Fe(H_2O)_6igr]^{2+}$  ,  $igl[Cr(H_2O)_6igr]^{3+}$ 



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**68.** Amongest the following ions which one has the highest magnetic

value ?

 $\left[Zn(H_2O)_6\right]^{2+}$ ,  $\left[Fe(H_2O)_6\right]^{2+}$ ,  $\left[Cr(H_2O)_6\right]^{3+}$ ,



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**69.** The oxidation number of cobalt in  $Kigl[Co(CO)_4igr]$  is -

A. + 1

B.+3

 $\mathsf{C.}-1$ 

#### **Answer: C**



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**70.** What will be the correct order for the wavelengths of absorption in the visible region for the following :

$$ig[ Ni(NO_2)_6 ig]^{4-}, ig[ Ni(NH_3)_6 ig]^{2+} < ig[ Ni(H_2O)_6 ig]^{2+} ?$$



71. Amongst the following, the most stable complex is -

A. 
$$\left[Fe(H_2O)_6
ight]^{3+}$$

B. 
$$\left[Fe(NH_3)_6
ight]^{3+}$$

C. 
$$\left[Fe(C_2O_4)_3
ight]^{3}$$

D. 
$$\left[FeCl_6\right]^{3}$$

### Answer:



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# **Higher Order Thinking Skill Hots Questions**

**1.** The molecular formula of a complex id  $PtCl_4.2KCl$ . From electrical conductance measurements, the molecule was found to contain three ions. The compound does not give any precipitate with  $AqNO_3$  solution. Identify the compound.



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**2.** Addition of iodine to KI solution does not bring down its freezing point. But, addition of mercuric iodide to KI solution increases its freezing point. Give reasons.



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**4.** Octahedral complexes of which transition metal ions have zero CFSE.



Explain.

**5.** Explain the principle of removal of rust stains from clothes using oxalic acid solution.



**6.** Explain the following : (1) Nickel does not form low spin octahedral complexes.

- **7.** Explain the following : (2) The  $\pi-$  complexes are known for the transition metals only.
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- **8.** Explain the following : (3) CO is considered to be a stronger ligand than  $NH_3$  for some metals.
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**9.**  $Co^{2+}$  is easily oxidised to  $Co^{3+}$  in the presence of a strong ligand.

Explain.



10. Which of the following is the most stable complex?

$$\left[Fe(NH_3)_6
ight]^{3+}, \left[Fe(C_2O_4)_3
ight]^{3-}, \left[FeCl_6
ight]^{3-}$$



**11.** What do you mean by macrocyclic effect in relation to the stability of coordination compounds?



**12.** Why do the metals of the second and third transition series have a greater tendency to form low-spin complexes as compared to the first transition series?



**13.** How many moles of AgCl will be precipitated if an excess of  $AgNO_3$  solution is added to one litre 1 (M) solution of  $\left[CrCl(H_2O)_5\right]Cl_2$ ?



**14.** Give an example of a coordination compound in which the flexidentate sulphate ion behaves as a monodentate ligand.



**15.**  $\left[CuCl_4\right]^{2-}$  exists but  $\left[CuI_4\right]^{2-}$  does not. Give reason.



16. Can the metal carbonyl compounds be called organometallics?



17. Compare the values of formation constant,  $K_f$  of the following coordination entities with their stabilities.

$$igl[Ni(NH_3)_6igr]^{2+}, k_f = 5.31 imes 10^8, \quad igl[Ni(en)_3igr]^{2+}, k_f = 1.12 imes 10^{18}$$



**18.** A coordination compound,  $CrCl_3.4H_2O$  gives a precipitate of AgCl with silver nitrate solution. From the value of molar conductivity, the aqueous solution of the compound was found to contain 2 moles of ions. Write the name and structural formula of the compound.



**19.** Show all stereoisomers of  $\left[Co(NH_3)_2Cl_2(en)\right]^+$ . Mention their inter relationship.



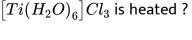
**20.** Give the structures of the stereo isomers of  $\left[Co(en)_2Br_2\right]^+$  and mention their inter relationships.



21. A metal complex having composition  $Cr(NH_3)_4Cl_2Br$  has been isolated in two forms (P) and (Q). The form (P) reacts with  $AgNO_3$  to give a white precipitate. Readily soluble in dilute aqueous ammonia, whereas (Q) gives a pale yellow precipitate soluble in concentrated ammonia. Write the formula of (P) and (Q) and state the hybridisation of Cr in each. Calculate the magnetic moments (spin only value).



22. Will there be any change in colour of the compound formed, if





23. If a coordination compound,  $CoCl_3.4NH_3$  is heated, it does not give out  $NH_3$ , while if the same reacts with  $AgNO_3$  solution, gives a white precipitate of AgCl. Write the structure and IUPAC name of the compound.



**24.** Arrange the following complexes in decreasing order of their electrical conductance :

$$\left[\operatorname{CoCl}(NH_3)_5\right]\operatorname{Cl}_2,\left[\operatorname{CoCl}_2(NH_3)_4\right]\operatorname{Cl},\left[\operatorname{Co}(NH_3)_6\right]\operatorname{Cl}_3,\left[\operatorname{CoCl}_3(NH_3)_3\right]$$



**25.** Octahedral and tetrahedral complexes made of the same metal ion have different coloures. Explain.



# Entrance Questions Engineering And Medical Archive Wbjee

1. Amongst  $\left[NiCl_4\right]^{2-}, \left[Ni(H_2O)_6\right]^{2+}, \left[Ni(PPh_3)_2Cl_2\right], \left[Ni(CO)_4\right] \& \left[Ni(CN)_4\right]^{2-}$ 

A. 
$$\left[NiCl_4
ight]^{2-}, \left[Ni(H_2O)_6
ight]^{2+}, \left[Ni(PPh_3)_2Cl_2
ight]$$

B. 
$$ig[Ni(CO)_4ig], ig[Ni(PPh_3)_2Cl_2ig], ig[NiCl_4ig]^2$$

D. 
$$igl[Ni(PPh_3)_2Cl_2igr], igl[Ni(CO)_4igr], igl[Ni(CN)_4igr]^{2-}$$

C.  $\left[Ni(CN)_4\right]^{2-}, \left[Ni(H_2O)_6\right]^{2+}, \left[NiCl_4\right]^{2-}$ 

Answer: A

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2. Optical isomerism is exhibited by (ox = oxalate anion, en = ethylenediamine)-

A. 
$$cis - igl[ CrCl_2(ox)_2 igr]^{3-}$$

B. 
$$\left[Co(en)_3\right]^{3+}$$

C. 
$$trans - \left[ CrCl_2(ox)_2 
ight]^{3-}$$

D. 
$$\left[Co(\otimes)(en)_2\right]^+$$

# Answer: A::B::D



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**3.** Addition of sodium thiosulphate solution to a solution of silver nitrate given 'X' as white precipitate, insoluble in water but soluble in excess thiosullphate solution to give 'Y'. On boiling in water, 'Y' gives 'Z'. 'X', 'Y' and 'Z' respectively are -

A. 
$$Ag_2S_2O_3,$$
  $Na_3ig[Ag(S_2O_3)_2ig],$   $Ag_2S$ 

B. 
$$Ag_2SO_4$$
,  $Naig[Ag(S_2O_3)_2ig]$ ,  $Ag_2S_2$ 

$$\mathsf{C.}\,Ag_2S_2O_3,Na_5\big[Ag(S_2O_3)_3\big],AgS$$

D. 
$$Ag_2SO_3, Na_3ig[Ag(S_2O_3)_2ig], Ag_2O$$

## **Answer: A**



**4.** Cold ferrous sulphate solution on absorption of NO develops brown colour due to the formation of -

A. paramagnetic  $\left[Fe(H_2O)_5NO\right]SO_4$ 

B. diamagnetic  $\left[Fe(H_2O)_5(N_3)\right]SO_4$ 

C. paramagnetic  $igl[Fe(H_2O)_5(NO_3)igr](SO_4)_2$ 

D. diamagnetic  $\left[Fe(H_2O)_4(SO_4)\right]NO_3$ 

### Answer: A



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**5.**  $PbCl_2$  is insoluble in cold water. Addition of HCl increases its solubility due to -

A. formation of soluble complex anions lik  $[PbCl_3]^-$ 

B. oxidation of Pb (II) to Pb(IV)

C. formation of  $igl[Pb(H_2O)_6igr]^{2+}$ 

D. formation of polymeric lead complexes

# Answer: A



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- **6.** The number of unpaired electrons in  $\left[CoCl_4\right]^{2-}$  ,  $\left[Ni(CO)_4\right]$  and  $igl[ Cu(NH_3)_6 igr]^{2+}$  respectively are -
  - A. 2,2,1
  - B. 3,0,1
  - C. 0,2,1
  - D. 2,2,0

**Answer: B** 

7. Ferric ion forms a prussian blue precipitate due to the formation of -

A. 
$$K_4igl[Fe(CN)_6igr]$$

$$\mathsf{B.}\,K_3\big[Fe(CN)_6\big]$$

C. 
$$Fe(CNS)_3$$

D. 
$$Fe_4igl[Fe(CN)_6igr]_3$$

### **Answer: D**



**8.** Silver chloride dissolves in excess of ammonium hydroxide solution. The cation present in the resulting solution is -

A. 
$$\left[Ag(NH_3)_6
ight]^+$$

B. 
$$\left[Ag(NH_3)_4
ight]^+$$

C.  $Ag^+$ 

D.  $\left[Ag(NH_3)_2
ight]^+$ 

### **Answer: D**



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# **Entrance Questions Engineering And Medical Archive Jee Main**

**1.** Which of the following facts about the complex  $\left[Cr(NH_3)_6\right]Cl_3$  is incorrect -

A. the complex involves  $d^2sp^3$  hybridisation and is octahedral in shape

B. the complex is paramagnetic

C. the complex is an outer orbital complex

D. the complex gives white pricipitate with silver nitrate solution

# Answer: C

**2.** Magnetic moment (spin only) of  $\left[NiCl_4\right]^2$  -

B. 5.46 BM

A. 1.82 BM

C. 2.82 BM

D. 1.41 BM

## **Answer: C**



**3.** Which among the following will be named as dibromidobis (ethylenediamine) chrominium (III) bromide -

A.  $[Cr(en)Br_2]Br$ 

B.  $\left[Cr(en)_3\right]Br_3$ 

$\left[Cr(en)_2Br_2\right]$	Br
	$\left[ Cr{(en)}_2Br_2 \right]$

D.  $[Cr(en)Br_4]$ 

### **Answer: C**



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**4.** Which of the following complex species is not expected to exhibit optical isomerism-

A. 
$$\left[Co(en)_3
ight]^{3+}$$

$$\mathsf{B.}\left[ Co(en)_{2}Cl_{2}\right] ^{+}$$

C. 
$$\left[Co(NH_3)_3Cl_3\right]$$

D. 
$$igl[ {Co(en)(NH_3)}_2 Cl_2 igr]^+$$

## **Answer: C**



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**5.** The octahedral complex of a metal ion  $M^{3\,+}$  with four monodentate lignads  $L_1,\,L_2,\,L_3$  and  $L_4$  absorb wavelengths in the region of red, green, yellow and blue respectively. The increasing order of ligand strength of the four ligands is -

A. 
$$L_1 < L_2 < L_4 < L_3$$

B. 
$$L_4 < L_3 < L_2 < L_1$$

C. 
$$L_1 < L_3 < L_2 < L_4$$

D. 
$$L_3 < L_2 < L_4 < L_1$$

### **Answer: C**



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**6.** Number of geometric isomers that can exist for square planar  $\left[Pt(Cl)(py)(NH_3)(NH_2OH)\right]^+$  is -

A. 4

- B. 6
- C. 2
- D. 3

# **Answer: D**



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- 7. Which of the following compounds is not yellow coloured -
  - A.  $(NH_4)_3ig[As(Mo_3O_{10})_4ig]$
  - B.  $BaCrO_4$
  - $\mathsf{C}.\,Zn_2ig[\mathit{Fe}(\mathit{CN})_6ig]$
  - D.  $K_3igl[Co(NO_2)_6igr]$

# **Answer: C**



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8. Which one of the following complexes shows optical isomerism -

A. 
$$\left[Co(NH_3)_3Cl_3\right]$$

$$\mathsf{B.}\, cis - \big[\mathit{Co}(en)_2\mathit{Cl}_2\big]\mathit{Cl}$$

C. 
$$trans - \lceil Co(en)_2 Cl_2 \rceil Cl$$
 (en = ethylenediamine)

D. 
$$\left[Co(NH_3)_4Cl_2\right]Cl$$

#### **Answer: B**



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**9.** The pair having the same magnetic moment is (At. No. : Cr = 24, Mn = 25,

A. 
$$\left[Cr(H_2O)_6\right]^{2+}$$
 and  $\left[CoCl_4\right]^{2-}$ 

B. 
$$\left[Cr(H_2O)_6\right]^{2+}$$
 and  $\left[Fe(H_2O)_6\right]^{2+}$ 

C. 
$$\left[Mn(H_2O)_6\right]^{2+}$$
 and  $\left[Cr(H_2O)_6\right]^{2+}$ 

D. 
$$\left[CoCl_4\right]^{2-}$$
 and  $\left[Fe(H_2O)_6\right]^{2+}$ 

## **Answer: B**



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**10.** On treatment of 100 mL of 0.1(M) solution of  $CoCl_3.6H_2O$  with excess

 $AgNO_3,\, 1.2 imes 10^{22}$  ions are precipitated. The complex is -

A. 
$$igl[ {Co(H_2O)}_6 igr] Cl_3$$

B. 
$$\left[Co(H_2O)_5Cl\right]Cl_2$$
.  $H_2O$ 

C. 
$$\left[Co(H_2O)_4Cl_2\right]Cl.2H_2O$$

D. 
$$\left[Co(H_2O)_3Cl_3\right].3H_2O$$

### **Answer: B**



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11. The oxidation states of Cr  $\left[Cr(H_2O)_6\right]Cl_3, \left[Cr(C_6H_6)_2\right] \text{ and } K_2\left[Cr(CN)_2(O_2)(O_2)(NH_3)\right]$ 

in

respectively are -

A. +3, +4 and +6

B. +3, +2 and +4

C. +3, 0 and +6

D. +3, 0 and +4

#### **Answer: C**



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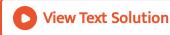
**12.** For 1 molal aqueous solution of the following compounds, which one will show the highest freezing point -

A.  $\left[Co(H_2O)_6\right]Cl_3$ 

B.  $\left[Co(H_2O)_5Cl\right]Cl_2$ .  $H_2O$ 

C.  $\left[Co(H_2O)_4
ight]Cl_2.2H_2O$ 

D.  $[Co(H_2O_3Cl_3].3H_2O$ 



13. Consider the following reaction and statements:

$$\left[ {Co(NH_3)}_4 Br_2 
ight]^+ + Br^- 
ightarrow \left[ {Co(NH_3)}_3 Br_3 
ight] + NH_3$$

(I) two isomers are produced if the reactant complex ion is a cis - isomers

(II) two isomers are produced if the reactant complex ion is a trans - isomer

(III) only one isomer is produced if the reactant complex ion is trans - isomer

(IV) only one isomer is produced if the reactant complex ion is a cis - isomer

The correct statements are -

A. (III) and (IV)

B. (II) and (IV)

C. (I) and (II)

D. (I) and (III)

#### Answer: D



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# **Entrance Questions Engineering And Medical Archive Neet**

**1.** The d-electron configurations of  $Cr^{2+}, Mn^{2+}, Fe^{2+}$  and  $Co^{2+}$  are  $d^4, d^5, d^6$  and  $d^7$  respectively. Which one of the following will exhibit minimum paramagnetic behaviour -

A. 
$$igl[Mn(H_2O_6]^{2\,+}igr]$$

B. 
$$\left[Fe(H_2O)_6
ight]^{2+}$$

C. 
$$\left[ Co(H_2O)_6 
ight]^{2+}$$

D. 
$$\left[Cr(H_2O)_6
ight]^{2+}$$

### **Answer: C**



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**2.** The complex,  $[Pt(py)(NH_3)BrCl]$  will have how many geometrical isomers -

A. 3

B. 4

C. 0

D. 2

### **Answer: A**



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**3.** The complex,  $\left[Co(NH_3)_6\right]\left[Cr(CN)_6\right]\left[Cr(NH_3)_6\right]\left[Co(CN)_6\right]$ the examples of which type of isomerism -

A. linkage isomerism

B. ionization isomerism

	C.	coordination	isomerism
--	----	--------------	-----------

D. geometrical isomerism

### **Answer: C**



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- 4. Of the following complex ions, which is diamagnetic in nature -
  - A.  $\left[NiCl_4
    ight]^{2}$  –
  - B.  $\left[Ni(CN)_4
    ight]^{2-}$
  - C.  $\left[CuCl_4
    ight]^{2-}$
  - D.  $[CoF_6]^{3-}$

## Answer: B



5. Which of the following complex compounds will exhibit highest paramagnetic behaviour -

**6.** Which of the following carbonyls will have the strongest C-O bond -

A. 
$$\left[Ti(NH_3)_6
ight]^{3+}$$

B. 
$$\left[Cr(NH_3)_6
ight]^{3+}$$

C.  $\left[ Co(NH_3)_6 \right]^{3+}$ 

D. 
$$igl[Zn(NH_3)_6igr]^{2+}$$

## **Answer: B**



- A.  $Mn(CO)_6^+$ 
  - B.  $Cr(CO)_6$
  - $C.V(CO)_6^-$
  - D.  $Fe(CO)_5$

#### Answer: A



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**7.** Which one of the following is an outer orbital complex and exhibits paramagnetic behaviour -

- A.  $\left[Cr(NH_3)_6
  ight]^{3+}$
- B.  $\left[ Co(NH_3)_6 
  ight]^{3+}$
- C.  $\left[Ni(NH_3)_6\right]^{2+}$
- D.  $\left[Zn(NH_3)_6
  ight]^{2+}$

#### Answer: C



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**8.** Which among the following will be named as dibromidobis (ethylenediamine) chromium (III) bromide -

A.  $[Cr(en)_2Br]Br_2$ 

 $\operatorname{B.}\left[Cr(en)_{2}Br_{2}\right]Br$ 

- C.  $[Cr(en)Br_2]Br$
- D.  $\left[Cr(en)_3\right]Br_3$

# **Answer: B**



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- 9. A magnetic moment of 1.73 BM will be shown by one among the folllowing -
- A.  $\left[CoCl_{6}\right]^{4-}$ 
  - $\mathsf{B.}\left[Cu(NH_3)_4\right]^{2\,+}$
  - C.  $\left[Ni(CN)_4\right]^{2-}$

# **Answer: B**

D.  $TiCl_4$ 

10. Which of the following complexes is used to be as an anticancer agent-

A. 
$$mer - \left[Co(NH_3)_3Cl_3\right]$$

$$\mathsf{B.}\, cis - \left[ PtCl_2(NH_3)_2 \right]$$

C. 
$$cis-K_2[PtCl_2Br_2]$$

$$\mathsf{D.}\, Na_{2}CoCl_{4}$$

#### **Answer: B**



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**11.** Among the following complexes, the one which shown zero crystal field stabilization energy (CFSE)-

A. 
$$igl[Mn(H_2O)_6igr]^{3\,+}$$

C. 3 D. 4

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# **Answer: C**

= ethylenedisamine)-

B.  $\left[Fe(H_2O)_6\right]^{3\,+}$ 

C.  $\left[ Co(H_2O)_6 
ight]^{2+}$ 

D.  $igl[ {Co(H_2O)}_6 igr]^{3\,+}$ 

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**12.** Number of possible isomers for the complex  $\left[Co(en)_2Cl_2
ight]$  will be (en

# B. 1

**Answer: B** 







**13.** The hybridisation involved in complex  $\left[Ni(CN)_4\right]^{2-}$  is

 $({\rm At.no.~of~Ni}=28)$ 

A.  $dsp^2$ 

 $\mathsf{B.}\,sp^3$ 

 $\mathsf{C.}\,d^2sp^2$ 

D.  $d^2sp^3$ 

#### **Answer: A**



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**14.** The name of complex ion,  $\left[Fe(CN)_6\right]^{3-}$  is -

A. hexacyanoiron (III) ion

B. hexacyanitoferrate (III) ion

C. tricyanoferrate (III) ion

D. hexacyanidoferrate (III) ion

#### **Answer: D**



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15. The sum of coordination number and oxidation number of the metal

M in the complex  $igl[M(en)_2(C_2O_4)igr]Cl$  is -

- A. 9
- B. 6
- C. 7
- D. 8

## Answer: A



16. The correct increasing order of trans - effect of the following species is

A. 
$$CN^{-} > Br^{-} > C_{6}H_{5}^{-} > NH_{3}$$

B. 
$$NH_3 > CN^- > Br^- > C_6H_5^-$$

C. 
$$CN^{\,-} > C_6 H_5^{\,-} > Br^{\,-} > N H_3$$

D. 
$$Br^- > CN^- > NH_3 > C_6H_5^-$$

### **Answer: C**



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17. Jahn-Teller effect is not found in high spin complex of -

A.  $d^9$ 

 $B.d^7$ 

 $\mathsf{C}.\,d^8$ 

 $D.d^4$ 



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**18.** Which of the following has longest C-O bond length ? (Fee C-O bond length in CO is  $1.128 {\rm \AA}$ )

- A.  $igl[Mn(CO)_6igr]^+$
- B.  $Ni(CO_4$
- C.  $\left[Co(CO)_4\right]^-$
- D.  $\left\lceil Fe(CO)_4 \right\rceil^{2-}$

#### Answer: D



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**19.** Pick out the correct statement with respect to  $igl[Mn(CN)_6igr]^3$  -

- A. it is  $sp^3d^2$  hybridised and octahedral
- B. it is  $sp^3d^2$  hybridised and tetrahedral
- C. it is  $d^2sp^3$  hybridised and octahedral
- D. it is  $dsp^2$  hybridised and square planar.



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**20.** The correct order of the stoichiometry of AgCl formed when  $AgNO_3$  in excess in treated with the complexes :

 $CoCl_3.6NH_3, CoCl_3.5NH_3, CoCl_3.4NH_3$  respectively is -

- A.  $1AgCl,\,3AgCl,\,2AgCl$
- $\mathsf{B.}\, 3AgCl,\, AgCl,\, 2AgCl$
- $\mathsf{C.}\ 3AgCl,\ 2AgCl,\ 1AgCl$
- ${\tt D.}\ 2AgCl,\ 3AgCl,\ 1AgCl$



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**21.** Correct increasing order for the wavelengths of absorption in the visible region for the complexes of  $Co^{3+}$  is -

A. 
$$\left[Co(en)_3
ight]^{3+}, \left[Co(NH_3)_6
ight]^{3+}, \left[Co(H_2O)_6
ight]^{3+}$$

B. 
$$\left[Co(H_2O)_6\right]^{3+}, \left[Co(en)_3\right]^{3+}, \left[Co(en)_3\right]^{3+}$$

C. 
$$\left[Co(H_2O)_6
ight]^{3+}, \left[Co(NH_3)_6
ight]^{3+}, \left[Co(en)_3
ight]^{3+}$$

D. 
$$\left[Co(NH_3)_6
ight]^{3+}, \left[Co(en)_3
ight]^{3+}, \left[Co(H_2O)_6
ight]^{3+}$$

#### Answer: A



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**22.** Iron carbonyl,  $Fe(CO)_5$  is -

C. trinuclear D. mononuclear Answer: D **Watch Video Solution 23.** The type of isomerism shown by the complex  $\lceil CoCl_2(en)_2 \rceil$  is -A. linkage isomerism B. geometrical isomerism C. ionisation isomerism D. coordination isomerism Answer: B Watch Video Solution

A. dinuclear

B. tetranuclear

**24.** The geometry and magnetic behaviour of the complex  $\left[Ni(CO)_4\right]$  are

A. tetrahedral geometry and paramagnetic

B. square planar geometry and diamagnetic

C. square planar geometry and paramagnetic

D. tetrahedral geometry and diamagnetic

#### **Answer: D**



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# Entrance Questions Engineering And Medical Archive Aiims

1. Which of the following is diamagnetic -

A.  $\left[Cu(NH_3)_4
ight]^{2+}$ 

B.  $\left[NiCl_4
ight]^{2}$  –

C.  $[PtCl_4]^{2-}$ 

D.  $\left[Cu(H_2O)_4
ight]^{2+}$ 

#### **Answer: C**



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- 2. The wavelength of light absorbed is highest in -
  - A.  $\left[Co(NH_3)_5Cl
    ight]^{2+}$
  - B.  $\left[Co(NH_3)_5H_2O\right]^{3+}$
  - C.  $\left[Co(NH_3)_6
    ight]^{3+}$
  - D.  $\left[Co(en)_3\right]^{2+}$

#### **Answer: A**



<b>3.</b> Which of the following metal ion forms unstable complex with $CN^-$ -		
A. $Ag(I)$		
B. $Zn(II)$		
C. $Cu(II)$		
D. $Cr(II)$		
Answer: A		
Watch Video Solution		
<b>4.</b> Least coordination number is shown by -		
A. $Co_2(CO)_8$		
B. $Mn_2(CO)_{10}$		
<del></del>		
C. $\left[Fe(en)_2NH_3 ight]$		

#### **Answer: A**



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- 5. The diamagnetic species is -
  - A.  $\left[Ni(CN)_4
    ight]^{2}$
  - B.  $\left[NiCl_4
    ight]^2$  -
  - C.  $\left[CoCl_4
    ight]^{2-}$
  - D.  $[CoF_6]^{2-}$

#### **Answer: A**



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**6.** In which of the following pairs both the complexes show optical isomerism -

B. 
$$igl[ Co(en)_3 igr] Cl_3, cis - igl[ Co(en)_2 Cl_2 igr] Cl$$

C.  $[PtCl \text{ (dien)}]Cl, [NiCl_2Br_2]^{2-}$ 

D.  $[Co(NO_3)_3(NH_3)_3]$ ,  $cis - [Pt(en)_2Cl_2]$ 

7. Amongst the following, the most stable complex is -

A.  $cis - \left[ Cr(C_2O_4)_2Cl_2 
ight]^{3-}, cis - \left[ Co(NH_3)_4Cl_2 
ight]$ 



**Answer: B** 

B.  $igl[Fe(NH_3)_6igr]^{3\,+}$ 

A.  $\left[Fe(H_2O)_6\right]^{3+}$ 

C.  $\left[Fe(C_2O_4)_3
ight]^{3-}$ 

D.  $\left[FeCl_{6}
ight]^{3}$  –

**Answer: C** 

8. Select the correct statement -

A. geometrical isomer may differ in dipole moment and visible/UV spectra.

B. complexes of the type  $[MA_3B_3]$  can also have facial (fac) and meridional (mer) isomer.

C. no optical isomer exists for the complex trans-  $igl[Co(en)_2Cl_2igr]^+$  .

D. all of these.

#### **Answer: D**



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**9.** Calculate the overall complex dissociation equilibrium constant for the  $\left[Cu(NH_3)_4\right]^{2+}$  ions, given that stability constant  $(eta_4)$  for this complex is  $2.1 imes 10^{13}$ -

A.  $8.27 \times 10^{-13}$ 

B.  $4.76 \times 10^{-14}$ 

 $\text{C.}\ 2.39 \times 10^{-7}$ 

D.  $1.83 \times 10^{14}$ 

### **Answer: B**



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10. Which of the following is a pair of diamagnetic complexes -

A. 
$$\left[ {Co(NH_3)}_6 
ight]^{3+}, \left[ {Fe(CN)}_6 
ight]^{4-}$$

B. 
$$\left[Co(ox)_3
ight]^{3-}, \left[FeF_6
ight]^{3-}$$

C. 
$$\left[Fe(ox)_3
ight]^{3-}, \left[FeF_6
ight]^{3-}$$

D. 
$$\left[Fe(CN)_6
ight]^{3-}, \left[CoF_6
ight]^{3-}$$

# **Answer: A**



<b>11.</b> Trien is -			
A. hexadentate, mono anionic			
B. tetradentate, neutral			
C. tetradenatate, dianion			
D. monodentate, anion			
Answer: B  Watch Video Solution			
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C.  $\left[Co(CO)_4\right]^-$ 

D. 
$$Fe(CO)_5$$

**Answer: A** 



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## **Solved Ncert Exemplar Problems**

**1.** Which of the following complexes formed by  $Cu^{2\,+}$  ions is most stable -

A. 
$$Cu^{2+} + 4NH_3 \Leftrightarrow \left[Cu(NH_3)_4\right]^{2+}, \log K = 11.6$$

B. 
$$Cu^{2+} + 4CN^- \Leftrightarrow \left[Cu(CN)_4
ight]^{2-}, \log K = 27.3$$

C. 
$$Cu^{2+} + 2en \Leftrightarrow \left[Cu(en)_3\right]^{2+}, \log K = 15.4$$

D. 
$$Cu^{2+} + 4H_2O \Leftrightarrow \left\lceil Cu(H_2O)_4 
ight
ceil^{2+}, \log K = 8.9$$

#### **Answer: B**



**2.** Colour of the coordination compounds depends on the crystal field splitting. What will be the correct order of absorption of wavelength of light in the visible region, for  $\left[Cu(NH_3)_6\right]^{3+}$ ,  $\left[Co(CN)_6\right]^{3-}$ ,  $\left[Co(H_2O)_6\right]^{3+}$ -

A. 
$$\left[ Co(CN)_6 \right]^{3-} > \left[ Co(NH_3)_6 \right]^{3+} > \left[ Co(H_2O)_6 \right]^{3+}$$

$$\mathsf{B.} \left[ Co(NH_3)_6 \right]^{3+} > \left[ Co(H_2O)_6 \right]^{3+} > \left[ Co(CN)_6 \right]^{3-}$$

C. 
$$\left[ Co(H_2O)_6 \right]^{3+} > \left[ Co(NH_3)_6 \right]^{3+} > \left[ Co(CN)_6 \right]^{3-}$$

D. 
$$\left[ {Co(CN)_6 } \right]^{3-} > \left[ {Co(NH_3)_6 } \right]^{3+} > \left[ {Co(H_2O)_6 } \right]^{3+}$$

#### **Answer: C**



- **3.** When 0.1 mol  $CoCl_3(NH_3)_5$  is treated with excess of  $AgNO_3$ , 0.2 mol of AgCl are obtained. The conductivity of solution will correspound to -
  - A. 1:3 electrolyte
  - B. 1:2 electrolyte

C. 1:1 electrolyte

D. 3:1 electrolyte

#### **Answer: B**



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**4.** When 1 mole  $CrCl_3.6H_2O$  is treated with excess of  $AgNO_3,\,\,$  3 mol

AgCl are obtained. Formula of the complex :

A.  $\left[CrCl_3(H_2O)_3\right].3H_2O$ 

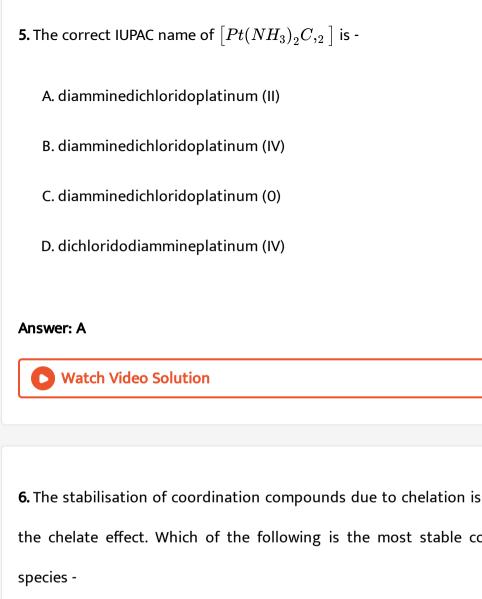
B.  $\left[CrCl_2(H_2O)_4\right]Cl.\ 2H_2O$ 

C.  $\left[CrCl(H_2O)_5\right]Cl_2$ .  $H_2O$ 

D.  $\left[Cr(H_2O)_6\right]Cl_3$ 

#### Answer: D





6. The stabilisation of coordination compounds due to chelation is called the chelate effect. Which of the following is the most stable complex

A. 
$$\left[Fe(CO)_5\right]$$

$$\operatorname{B.}\left[Fe(CN)_{6}\right]^{3}-$$

C. 
$$\left[Fe(C_2O_4)_3
ight]^{3-}$$

D. 
$$\left[Fe(H_2O)_6
ight]^{3+}$$



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- 7. Indicate the complex ion which shows geometrical isomerism -
  - A.  $\left[Cr(H_2O)_4Cl_2
    ight]^+$
  - $\mathsf{B.}\left[Pt(NH)_3\right)_3Cl\right]$
  - C.  $\left[ {Co(NH_3)_6} 
    ight]^{3+}$
  - D.  $\left[ Co(CN)_5(NC) \right]^{3-}$

**Answer: A** 



**8.** The CFSE for octahedral  $[CoCl_6]^{4-}$  is  $18000cm^{-1}$ . The CFSE for tetrahedral  $[CoCl_4]^{2-}$  will be -

A. 
$$18000cm^{-1}$$

B.  $16000cm^{-1}$ 

C.  $8000cm^{-1}$ 

D.  $20000cm^{-1}$ 

#### Answer: C



- **9.** Due to the presence of ambidentate ligands coordination compounds show isomerism. Palladium complexes of the type  $\left[Pt(C_6H_5)_2(SNC)_2\right]$  and  $\left[Pt(C_6H_5)_2(NSC)_2\right]$  are -
  - A. linkage isomers
  - B. coordination isomers

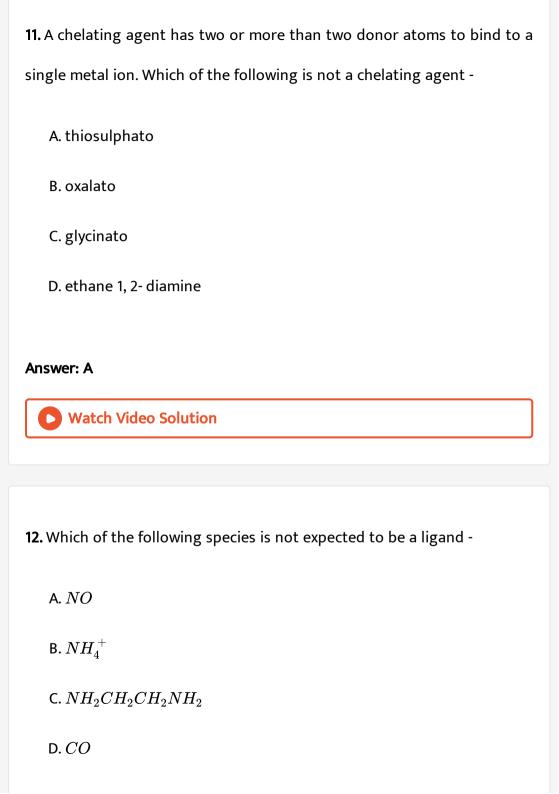
C. ionisation isomers	
D. geometrical isomers	
Answer: A	
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<b>10.</b> The compounds $igl[Co(SO_4)(NH_3)_5igr]Br$ and represent -	$\big[Co(SO_4)(NH_3)_5\big]Cl$
A. linkage isomerism	
B. ionisation isomerism	

C. coordination isomerism

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D. no isomerism

**Answer: D** 



#### **Answer: B**



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- **13.** What kind of isomerism exists between  $\left[Cr(H_2O)_6\right]Cl_3$  (violet) and  $\left[Cr(H_2O)_5cl\right]Cl_2$ .  $H_2O$  (greyish green)-
  - A. linkage isomerism
  - B. solvate isomerism
  - C. ionisation isomerism
  - D. coordination isomerism

#### **Answer: B**



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**14.** IUPAC name of  $igl[Pt(NH_3)_2Cl(NO_2)igr]$  is -

A. platinum diaminechloronitrite

B. chloronitritio - N - ammineplatinum (II)

C. diamminechloridonitrito - N - platinum (II)

D. diamminechloronitrito - N - platinate (II)

#### **Answer: C**



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15. Atomic number of Mn, Fe and Co are 25, 26 and 27 respectively. Which of the following inner orbital octahedral complex ions are diamagnetic -

A. 
$$\left[Co(NH_3)_6
ight]^{3+}$$

B. 
$$ig[Mn(CN)_6ig]^{4-}$$

C. 
$$\left[Fe(CN)_6\right]^{4-}$$

D. 
$$\left[Fe(CN)_6
ight]^{3-}$$

## Answer: A::C

**16.** Atomic number of Mn, Fe and Co an dNi are 25, 26 27 and 28 respectively. Which of the following outer orbital octahedral complexes have same number of unpaired electrons -

A. 
$$\left[MnCl_{6}
ight]^{3}$$
 –

B. 
$$\left[FeF_6\right]^{3}$$

$$\mathsf{C.}\left[CoF
ight]^{3}$$

D. 
$$\left[Ni(NH_3)_6
ight]^{2+}$$

#### Answer: A::C



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**17.** Which of the following options are correct for  $\left[Fe(CN)_6\right]^{3-}$  -complex -

- A.  $d^2sp^3$  hybridisation
- B.  $sp^3d^2$  hybridisation
- C. paramagnetic
- D. diamagnetic

### Answer: A::C



- **18.** An aqueous pink solution of cobalt(II) chloride changes to deep blue on addition of excess of HCl. This is because -
  - A.  $\left[Co(H_2O)_6
    ight]^{2+}$  is transformed into  $\left[CoCl_6
    ight]^{4-}$
  - B.  $\left[Co(H_2O)_6
    ight]^{2+}$  is transformed into  $\left[CoCl_4
    ight]^{2-}$
  - C. tetrahedral complexes have smaller crystal field splitting than octahedral complexes.

D. tetrahedral complexes have larger. Crystal field splitting than octahedral complexes.

### **Answer: B::C**



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19. Which of the following complexes are homoleptic -

- A.  $\left[ Co(NH_3)_6 
  ight]^{3\,+}$
- B.  $\left[Co(NH_3)_4Cl_2\right]^+$
- C.  $\left\lceil Ni(CN)_4 
  ight
  ceil^{2-}$
- D.  $\left[Ni(NH_3)_4Cl_2
  ight]$

### Answer: A:C



20. Which of the following complexes are heteroleptic -

A. 
$$\left[Cr(NH_3)_6
ight]^{3\,+}$$

B. 
$$\left[Fe(NH_3)_3Cl_2\right]^+$$

C. 
$$\left[Mn(CN)_6\right]^{4-}$$

D. 
$$\left[Co(NH_3)_4Cl_2\right]$$

### **Answer: B::D**



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21. Identify the optically active compounds from the following -

A. 
$$\left[Co(en)_3
ight]^{3+}$$

$$\text{B.} \, trans - \left[ Co(en)_2 Cl_2 \right]^+$$

$$\mathsf{C.}\,cis - \left[Co(en)_2Cl_2\right]^+$$

D. 
$$\left[Cr(NH_3)_5Cl\right]$$

### Answer: A::C



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**22.** Identify the correct statements for the behaviour of ethane -1, 2 - diamine as a ligand -

A. it is a neutral ligand.

B. it is a didentate ligand.

C. it is a chelating ligand

D. it is a unidentate lignad.

### Answer: A::B::C



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23. Which of the following show linkage isomerism -

A.  $\left[Co(NH_3)_5(NO_2)
ight]^{2+}$ B.  $\left[Co(H_2O)_5CO
ight]^{3+}$ 

D.  $\left\lceil Fe(en)_2Cl_2
ight
ceil^+$ 

C.  $\left[Cr(NH_3)_5SCN\right]^{2+}$ 

### Answer: A::C



**24.** Arrange the given complexes in the increasing order of conductivity of their solution:  $[Co(NH_3)_3Cl_3], [Co(NH_3)_4Cl_2]Cl, [Co(NH_3)_6]Cl_3, [Cr(NH_3)_5Cl]Cl_2$ 



**25.** A coordination compound  $CrCl_3.4H_2O$  precipitates AgCl when treated with  $AgNO_3$ . Molar conductance of its solution corresponds to a total of two ions. Write structural formula of the compound and name it.

**26.** A complex of the type  $\left[M(\mathrm{AA})_2X_2\right]^{n+}$  is known to be optically active. What does this indicate about the structure of the complex ? Give one example.



**27.** Magnetic moment of  $\left[MnCl_4
ight]^{2-}$  is 5.92 BM. Explain.



**28.** On the basis of crystal field theory explain why Co(III) forms paramagnetic octahedral complex with weak - field ligands whereas it forms diamagnetic octahedral complex with strong - field ligands.



**29.** Why are low spin tetrahedral complexes not formed?



**30.** Give the electronic configuration of the following complexes on the basis of Crystal Field Splitting theory.

$$\left[CoF_{6}
ight]^{3-}, \left[Fe(CN)_{6}
ight]^{4-} ext{ and } \left[Cu(NH_{3})_{6}
ight]^{2+}$$



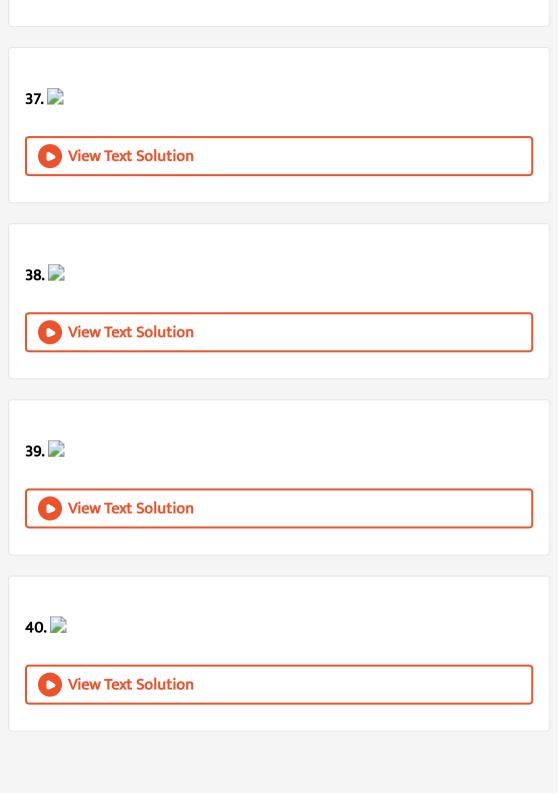
**31.** Explain why  $\left[Fe(H_2O)_6\right]^{3+}$  has magnetic moment value of 5.92 BM whereas  $\left[Fe(CN)_6\right]^{3-}$  has a value of only 1.74 BM.



**32.** Arrange the following complex ions in increasing order of crystal field splitting energy  $\left(\Delta_0\colon \left[Cr(Cl)_6\right]^{3-}, \left[Cr(CN)_6\right]^{3-}, \left[Cr(NH_3)_6\right]^{3+}\right)$ .



33. Why do compounds having similar geometry have different magnetic moment? **Watch Video Solution 34.**  $CuSO_4.5H_2O$  is blue in colour while  $CuSO_4$  is colourless. Why? **Watch Video Solution** 35. Name the type of isomerism when ambidentate ligands are attached to central metal ion. Give two examples of ambidentate ligands. **Watch Video Solution** 36. **View Text Solution** 



**41.** Assertion (A): Toxic metal ions are removed by chelating ligands.

Reason (R): Chelate complexes tend to be more stable.

A. Both (A) and (R) are true and (R) is the correct explanation of (A)

B. Both (A) and (R) are true but (R) is not the correct explanation of (A)

C. (A) is true but (R) is false

D. (A) is false but (R) is true

### Answer: A



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**42.** Assertion (A) :  $\left[Cr(H_2O)_6\right]Cl_2$  and  $\left[Fe(H_2O)_6\right]Cl_2$  are reducing in nature.

Reason (R): Unpaired electrons are present in their d - orbitals.

A. Both (A) and (R) are true and (R) is the correct explanation of (A)

B. Both (A) and (R) are true but (R) is not the correct explanation of (A)

C. (A) is true but (R) is false

D. (A) is false but (R) is true

### **Answer: B**



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**43.** Assertion (A): Linkage isomerism arises in coordination compounds containing ambidentate lignad.

Reason (R): Ambidentate ligand has 2 different donor.

A. Both (A) and (R) are true and (R) is the correct explanation of (A)

B. Both (A) and (R) are true but (R) is not the correct explanation of (A)

C. (A) is true but (R) is false

D. (A) is false but (R) is true

### **Answer: A**



**44.** Assertion (A) : Complexes of  $MX_6$  and  $MX_5L$  type (X and L are unidentate) do not show geometrical isomerism.

Reason (R): Geometrical isomerism is not shown by complexes of coordination number 6.

A. Both (A) and (R) are true and (R) is the correct explanation of (A)

B. Both (A) and (R) are true but (R) is not the correct explanation of (A)

C. (A) is true but (R) is false

D. (A) is false but (R) is true

### **Answer: C**



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**45.** Assertion (A) :  $\left[Fe(CN)_6\right]^{3-}$  ion shows magnetic moment corresponding to two unpaired electrons.

Reason (R) : Because it has  $d^2sp^3$  type hybridisation.

A. Both (A) and (R) are true and (R) is the correct explanation of (A)

B. Both (A) and (R) are true but (R) is not the correct explanation of (A)

C. (A) is true but (R) is false

D. (A) is false but (R) is true

### Answer: D



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**46.** Using crystal field theory, draw energy level diagram, write electronic configuration of central metal atom/ion, determine magnetic moment value in the following:

$$\left[CoF_{6}
ight]^{3-},\left[Co(H_{2}O)_{6}
ight]^{2+},\left[Co(CN)_{6}
ight]^{3-}$$



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47. Using crystal field theory, draw energy level diagram, write electronic configuration of central metal atom/ion, determine magnetic moment

value in the following:

$$\left[FeF_{6}
ight]^{3-},\left[Fe(H_{2}O)_{6}
ight]^{2+},\left[Fe(CN)_{6}
ight]^{4-}$$



**48.** Using valence bond theory, explain the following in relation to the complexes given below :

$$\left[Mn(CN)_6\right]^{3-}, \left[Co(NH_3)_6\right]^{3+}, \left[Cr(H_2O)_6\right]^{3+}, \left[FeCl_6\right]^{4-}$$
 (i) Type of hbridisation. (ii) Inner or outer orbital complex. (iii) Magnetic

\_\_\_\_

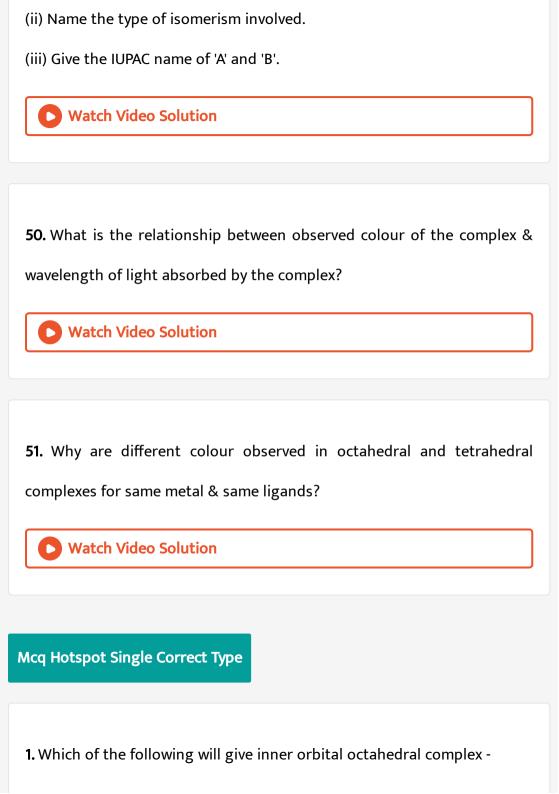
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Answer the following questions.

behaviour. (iv) Spin only magnetic moment value.

**49.**  $CoSO_4Cl.5NH_3$  exists in two isomeric forms 'A' and 'B'. Isomer 'A' reacts with  $AgNO_3$  to give white precipitate, but does not react with  $BaCl_2$ . Isomer 'B' gives white precipitate with  $BaCl_2$  but does not react with  $AgNO_3$ .

(i) Identify 'A' & 'B' and write their structural formulas.



 $\mathsf{C}.\,d^6$ D. All of these

### **Answer: C**

A.  $d^7$ 

 $B.d^8$ 



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2. Which one of the following is paramagnetic -

A.  $\left[Ni(H_2O)_6
ight]^{2\,+}$ 

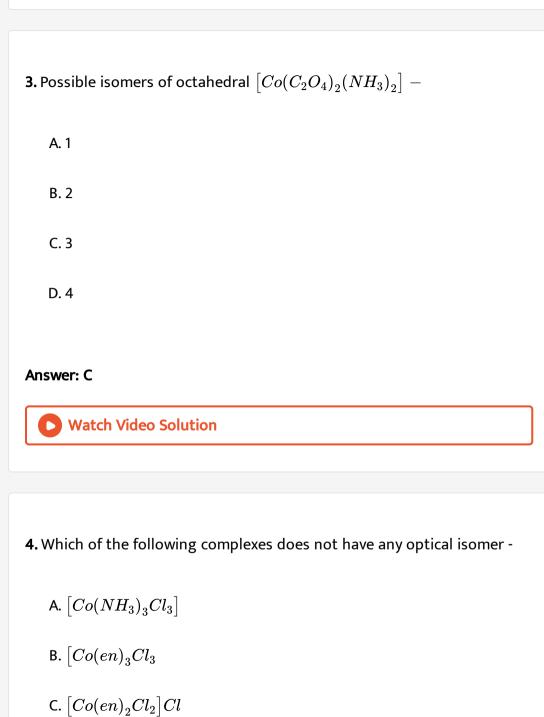
 $\operatorname{B.}\left[Ni(CO)_4\right]$ 

C.  $\left[Zn(NH_3)_4
ight]^{2+}$ 

D.  $\left[ {Co(NH_3)}_6 
ight]^{3\,+}$ 

# **Answer: A**





D. 
$$\left[Co(en)(NH_3)_2Cl_2\right]Cl$$

### Answer: A



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- **5.** The central metal atom of which of the following species does not have any d-electron -
  - A.  $[MnO_4]^{\,-}$
  - B.  $\left[Co(NH_3)_6
    ight]^{3+}$
  - C.  $\left[Fe(CN)_6
    ight]^{3}$
  - D.  $igl[Cr(H_2O)_6igr]^{3\,+}$

### Answer: A



<b>6.</b> Effective atomic number of Cr in $igl[ Cr(NH_3)_6 igr] Cl_3$ is -
A. 32
B. 33
C. 34
D. 35
Answer: B
Watch Video Solution
7. Which of the following $den^2$ hybridized complex ions is square plan

**7.** Which of the following  $dsp^2$  hybridised complex ions is square planar-

A.  $\left[Ni(CN)_4
ight]^{2-}$ 

B.  $\left[Cu(NH_3)_4
ight]^{2+}$ 

C.  $[PtCl_4]^{2-}$ 

D. All of these

### Answer: D



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8. Which of the following does not form an amine even in presence of excess ammonia -

- A.  $Al^{3\,+}$
- B.  $Ag^+$
- C.  $Cu^{2+}$
- D.  $Cd^{2+}$

### Answer: A



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9. Which of the following is in accordance with spectrochemical series -

A. 
$$Cl^- < F^- < [C_2 O_4]^{2-} < NO_2^- < CN^-$$

B.  $CN^- < \left[ C_2 O_4 \right]^{2-} < Cl^- < NO_2^- < F^-$ 

C.  $[C_2O_4]^{2-} < F^- < Cl^- > NO_2^- < CN^-$ 

D. 
$$F^{\,-} < C l^{\,-} < N O_2^{\,-} < C N^{\,-} < [C_2 O_4]^{2\,-}$$

## Answer: A



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10. Which one of the following is a high spin complex -

B.  $[FeF_6]^{3}$ 

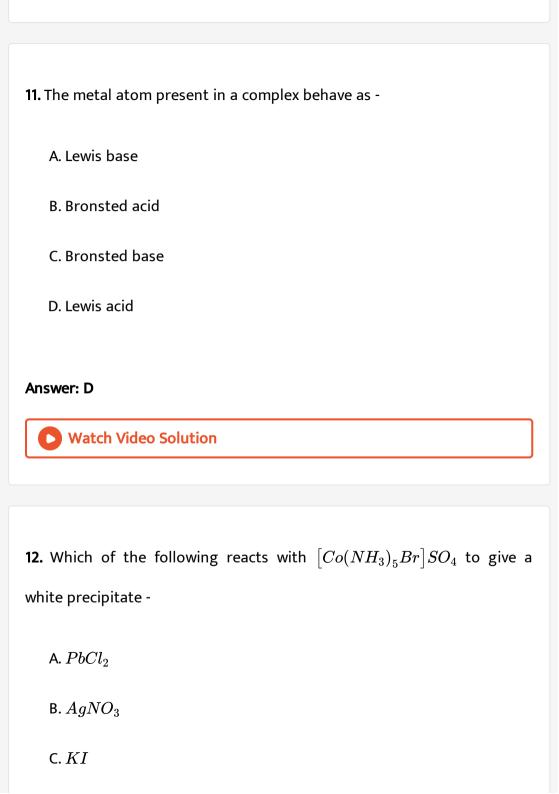
A.  $[CoCl_6]^{3}$ 

C.  $\left[Mn(H_2O)_6\right]^{2+}$ 

D. all of these



Answer: D



### **Answer: A**



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- 13. Which of the following ions exhibit coordination number 4 -
  - A.  $Pt^{2+}$
  - B.  $Cr^{3+}$
  - C.  $Fe^{3+}$
  - D.  $Pt^{4+}$

### **Answer: A**



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14. Platinum reacts with aquaregia to produce -

maximum -

- C.  $PtCl_4$ 
  - D.  $PtCl_2$

A.  $Pt(NO_3)_4$ 

B.  $H_2[PtCl_6]$ 

## **Answer: B**



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- 15. Molar conductivity of the aqueous solution of which of the following is
  - A.  $\left[Pt(NH_3)_6\right]Cl_4$ 

    - C.  $\left[Pt(NH_3)_4Cl_2\right]Cl_2$

B.  $[Pt(NH_3)_5Cl]Cl_3$ 

D.  $\left[Pt(NH_3)_3Cl_3\right]Cl$ 

## Answer: A

16. The reagent used for identifying nickel ion is -

A. potassium ferrocyanide

B. phenolphthalein

C. dimethylglyoxime

D. edta

### **Answer: C**



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**17.** As per stability constants (imaginery) given, which of the following is the strongest ligand -

(i) 
$$Cu^{2+}+4NH_2\Leftrightarrow \left[Cu(NH_3)_4
ight]^{2+},\left(K=4.5 imes 10^{11}
ight)$$

(ii) 
$$Cu^{2+} + 4CN^- \Leftrightarrow \left[Cu(CN)_4
ight]^{2+}, \left(K = 2.0 imes 10^{27}
ight)$$

**Answer: B** 

A.  $NH_3$ 

B.  $CN^-$ 

 $\mathsf{C}.\,en$ 

D.  $H_2O$ 

(iii)  $Cu^{2+} + 2en \Leftrightarrow \left[Cu(en)_2
ight]^{2+}, \left(K = 3.0 imes 10^{15}
ight)$ 

(iv)  $Cu^{2+} + 4H_2O \Leftrightarrow \left \lceil Cu(H_2O)_4 
ight 
ceil^{2+}, \left (K=9.5 imes 10^8 
ight )$ 

18. The reason behind two different colours of the complex entity

 $igl[Co(NH_3)_4Cl_2igr]^+$  is -

- A. ionisation isomerism
- B. optical isomerism
- C. geometrical isomerism

D. linkage isomerism

### **Answer: C**



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**19.** In which of the following pairs of complex ions, spin - only magnetic moment of both are same -

A. 
$$\left[Cr(H_2O)_6\right]^{2+}, \left[CoCl_4\right]^{2-}$$

B. 
$$\left[Cr(H_2O)_6\right]^{2+}, \left[Fe(H_2O)_6\right]^{2+}$$

C. 
$$\left[Mn(H_2O)_6
ight]^{2+}, \left[Cr(H_2O)_6
ight]^{2+}$$

D. 
$$\left[CoCl_4\right]^{2-}, \left[Fe(H_2O)_6\right]^{2+}$$

### **Answer: B**



**20.** The Fe-C bond in  $Fe(CO)_{ar{5}}$  displays-

A.  $\pi$  character

 $\operatorname{B.}\sigma\operatorname{character}$ 

C. ionic character

D. both  $\pi$  and  $\sigma$  character

### **Answer: D**



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21. The formula of tris(ethylenediammine) cobalt (III) sulphate is -

- A.  $\left[Co(en)_2SO_4\right]$
- B.  $\left[Co(en)_3SO_4\right]$
- C.  $igl[Co(en)_3igr]_2SO_4$
- D.  $\left[Co(en)_3
  ight]_2(SO_4)_3$

### **Answer: D**



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- **22.** In aqueous solution, which gives  $Fe^{3\,+}$  ion -
  - A.  $\left[Fe(CN)_6
    ight]^{3}$  –
  - B.  $\left[Fe(CN)_6\right]^{4-}$
  - C.  $Fe_2(SO_4)_3$
  - D.  $NH_4(SO_4)_2$ .  $FeSO_4.6H_2O$

### **Answer: C**



- 23. Which of the groups act as ambident ligands -
  - A.  $CO_3^{2-}$

B.  $CN^{\,-}$ 

 $\mathsf{C.}\,NO_2^-$ 

D. ethylenediamine

### Answer: B::C



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### 24. Identify complexes which are expected to be coloured -

A.  $Ti(NO_3)_4$ 

B.  $\left[Cu(\mathrm{NCCH_3})_4\right]^+BF_4^-$ 

C.  $\left[Cr(NH_3)_6
ight]^{3+}3Cl^-$ 

D.  $K_3[VF_6]$ 

### Answer: C::D



25. Which of the following statement(s) is/are correct -

A. 
$$\left[Ni(Cl)_{\scriptscriptstyle 4}\right]^{2-}$$
 &  $\left[Ni(CN)_{\scriptscriptstyle 4}\right]^{2-}$  differ in magnetic property

B. 
$$[NiCl_4]^{2-}$$
 &  $\left[Ni(CN)_4
ight]^{2-}$  differ in their geometry.

C.

$$\left[NiCl_{4}
ight]^{2-}$$
 &  $\left[Ni(CN)_{4}
ight]^{2-}$  differ in primary valencies of nickel.

D.

Answer: A::B::D

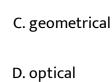
$$\left[NiCl_4
ight]^2$$
 &  $\left[Ni(CN)_4
ight]^2$  differ in hybridisation state of nickel

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- 26. Which of the following isomerisms are shown by the complex
  - A. ionisation

 $[CoCl_2(OH)_2(NH_3)_2]Br$  -

- B. linkage



Answer: A::C::D



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**27.** In test of  $NO_3^-$  ion, the dark brown ring complex is formed. Which is true of this complex -

- A. the colour is due to charge transfer spectra.
- B. iron and NO both have +1 charge.
- C. the complex species can be represented as  $\left\lceil \stackrel{I}{Fe}(H_2O)_5 NO 
  ight
  ceil^{2+}$
- D. iron has +2 oxidation state and NO is neutral.

Answer: A::B::C



28. Which of the following are outer orbital complexes -

A.  $\left[Co(NH_3)_6
ight]^{3+}$ 

 $\mathrm{B.}\left[Mn(CN)_{6}\right]^{3}-$ 

C.  $\left[Ni(NH_3)_6
ight]^{2+}$ 

D.  $\left[FeF_6
ight]^{3}$  -

### Answer: C::D



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## **Exercise Very Short Answer Type Questions Answer In One Two Sentences**

1. What is coordination polyhedron? Give an example.



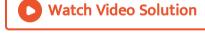
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**2.** Give one example each of perfect and imperfect complexes.

**3.** What is  $\pi$  — acid lignad? Give example?



**4.** Write the IUPAC name of the ionisation isomer of  $\lceil Co(NH_3)_5 SO_4 \rceil Br$ .



 $[Co(en)_3][Cr(CN)_6].$ 

the

IUPAC name

**6.** How many isomers are possible for  $\lceil Co(NH_3)_4 Cl_2 \rceil Cl$ ?

of

5.

Write



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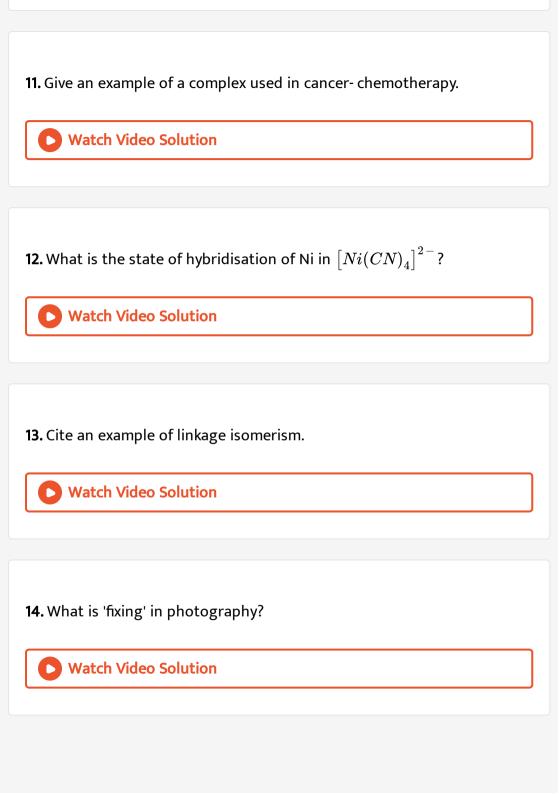


the coordination

isomer

of

<b>7.</b> How will you identify cis- and trans- isomers of $\left[Pt(NH_3)_2Cl_2 ight]$ ?
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8. Why do most transition elements and their compounds exhibit paramagnetism?
Watch Video Solution
<b>9.</b> Why do the tetrahedral compounds of the type $MA_2B_2$ not display geometrical isomerism?
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10. Name the ligand used in the treatment of lead toxicity.
Watch Video Solution



<b>15.</b> Give example of symmetrical & unsymmetrical bidentate lignads.
Watch Video Solution
<b>16.</b> Give two examples of complexes having biological importance.
Watch Video Solution
17. Which coordination compounds removes hardness of water?
Watch Video Solution
<b>18.</b> Which coordination number(s) is/are most common in coordination compounds?
Watch Video Solution

<b>19.</b> What is indicated by ' $\Delta_0 < P$ ' in an octahedral crystal field?			
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**21.** How many isomers are possible for the compound  $\left[Cu(NH_3)_4\right][PtCl_4]$ ?



22. Name a organometallic compound used as homogeneous catalyst.



<b>23.</b> How many unpaired electrons are there in $\left[Pt(CN)_4\right]^{2-}$ ion (square planar)?			
Watch Video Solution			
Exercise Fill In The Blanks			
<b>1.</b> Due to $dsp^2$ -hybridisation, shape of a compound becomes			
Watch Video Solution			
2. cis-/ trans- isomerism is not possible of structure.			
Watch Video Solution			
3. In metal carbonyls, metal is present in oxidation state.			
Watch Video Solution			

<b>4.</b> As perVBT, hybridisation state of $Cu$ in $\left[CuCl_4\right]^{2-}$ is Watch Video Solution
<b>5.</b> Magnetic moment of a high - spin complex is than of a low-spin complex.
Watch Video Solution
<b>6.</b> CFSE of an octahedral $d^4$ high-spin complex is
Watch Video Solution
<b>7.</b> $\left[Cr(NH_3)_6 ight]^{3+}$ is an orbit complex and in nature.
Watch Video Solution

<b>8.</b> Chemical name of $\left[C_4H_4Fe(CO)_3 ight]$ is
Watch Video Solution
<b>9.</b> Coordination compound containing ligand exhibit linkage isomerism only.
Watch Video Solution
Exercise Short Answer Type Questions
1. What do you mean by denticity of a ligand? Give example each of monodentate lignads.
Watch Video Solution
2. What is chelating ligand and ambident ligand? Give example.

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**3.** Which of the following categories of complexes display geometrical isomerism: (i) linear (ii) square planar (iii) tetrahedral (iv) octahedral.





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**6.** Write geometrical shapes and hybridisation state of the central metal atom for following:

**5.** Will there be any change in colour if  $\lceil Ti(H_2O)_6 \rceil Cl_3$  is heated?

4. What do you mean by crystal field splitting and CFSE?

(i)  $\left[ZnCl_4
ight]^2$ 

**7.** Write geometrical shapes and hybridisation state of the central metal atom for following:

(ii) 
$$\left[Ni(CN)_4
ight]^{2-}$$



**8.**  $FeSO_4$  solution mixed with  $(NH_4)_2SO_4$  solution in 1:1 molar ratio gives the test of  $Fe^{2+}$  ion, but  $CuSO_4$  solution mixed with aqueous  $NH_3$  in 1:4 molar ratio does not give the test of  $Cu^{2+}$  ion. Explain.



- **9.** Calculate the oxidation number of the central metal atom or ion for the following :
- (i)  $\left[Cr(C_2O_4)_3\right]^{3-}$



**10.** Calculate the oxidation number of the central metal atom or ion for the following :

(ii) 
$$igl[CoCl_2(en)_2igr]^+$$



**11.** Calculate the oxidation number of the central metal atom or ion for the following :

(iii) 
$$\left[PtClNO_2(NH_3)_4\right]^{2+}$$
.



**12.** What are  $t_{2g}$  and  $e_g$  orbitals?



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**13.** A solution of  $\left[Ni(H_2O)_6\right]^{2+}$  is green but a solution of  $\left[Ni(CN)_4\right]^{2-}$  is colourless. Explain.



**14.** Explain the nature of bonding in  $\left[Fe(CN)_6\right]^{3-}$  and  $\left[FeF_6\right]^{3-}$  on the basic of valence bond theory.



**15.** Give the IUPAC names with an example, for each of (i) cationic complex (ii) anionic complex and (iii) neutral complex.



**16.** Why is  $Co^{2+}$  easily oxidised to  $Co^{3+}$  in presence of a strong-field ligand?



**17.** Why is CO a stronger complexing agent than  $NH_3$ ?



**18.** Why do metal ions never form low-spin tetrahedral complex?



**19.** Calculate the effective atomic number (EAN) of the central metal for the following:

- (i)  $\left[Ag(NH_3)_2
  ight]^{2+}$ 
  - Watch Video Solution

**20.** Calculate the effective atomic number (EAN) of the central metal for the following:

- (ii)  $\left[Fe(CN)_6
  ight]^{3}$ 
  - Watch Video Solution

- **21.** Calculate the effective atomic number (EAN) of the central metal for the following:
- (iii)  $igl[Co(NH_3)_5Cligr]^{2+}$ 
  - Watch Video Solution

- **22.** Discuss the importance of chelate formation.
  - Watch Video Solution

1. Which of the following will give inner orbital octahedral complex -
A. $d^7$
B. $d^8$
$C.d^6$
D. all the given
Answer: C
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Practice Set 10
<b>1.</b> The $Fe-C$ bond in $Fe(CO)_5$ displays-
A. $\pi$ character
B. $\sigma$ character
C. ionic character

D. both $\pi$ and $\sigma$ character
Answer: D
Watch Video Solution
ractice Set 11
<b>1.</b> The metal atom present in a complex as -
A. Lewis base

B. Bronsted acid

C. Bronsted base

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D. Lewis acid

**Answer: D** 

**1.** For the ion  $\left[Cr(NH_3)_4Cl_2
ight]^+$  what is the oxidation state or Cr?

A. + 3

B. + 2

C. + 1

D. 0

#### **Answer: A**



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### **Practice Set 13**

A.  $\left[Fe(CN)_6\right]^{3}$ 

$$(V)_6]^{3}$$

**1.** In aqueous solution, which gives  $Fe^{3\,+}$  ion -

 $\mathsf{C.}\, Fe_2(SO_4)_3$ D.  $(NH_4)_2SO_4$ .  $FeSO_4.6H_2O$ 

B.  $\left[Fe(CN)_6\right]^{4-}$ 

# **Answer: C**



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# Practice Set 14

1. Which of the following is optically active -

A.  $\left[Zn(en)_2
ight]^{2+}$ 

B.  $\left[Zn(en)(NH_3)_2\right]^{2+}$ 

D.  $igl[Co(H_2O)_4(en)igr]^{3\,+}$ 

C.  $\left[Co(en)_3\right]^{3+}$ 

**Answer: C** 



**Practice Set 15** 

1. What is chelating ligand and ambident ligand? Give example.



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Practice Set 16

**1.** Why is CO a stronger complexing agent than  $NH_3$ ?



**Watch Video Solution** 

**Practice Set 17** 

**1.** Why is  $Co^{2+}$  easily oxidised to  $Co^{3+}$  in presence of a strong-field ligand?



## Practice Set 18

**1.** A solution of  $\left[Ni(H_2O)_6\right]^{2+}$  is green but a solution of  $\left[Ni(CN)_4\right]^{2-}$  is colourless. Explain.



## Practice Set 19

**1.** Will there be any change in colour if  $\lceil Ti(H_2O)_6 \rceil Cl_3$  is heated?



**1.** (a) Write down the IUPAC name of  $igl[Fe(C_2O_4)(H_2O)_4igr]_2SO_4.$ 



#### **Practice Set 21**

1. (b) Determine the hybridisation of the central metal ion.

 $\big[Fe(C_2O_4)(H_2O)_4\big]_2SO_4$ 



2. How does complex salt differ from double salt?

