

### **CHEMISTRY**

### FOR IIT JEE ASPIRANTS OF CLASS 12 FOR CHEMISTRY

### **COORDINATION COMPLEXES**

### W.E.

**1.** When excess of silver nitrate solution is added to aqueous solution containing 0.1 molar  $CoCl_3$ .  $xNH_3$ . If 43.05 g of silver chloride precipitated, then x value is



**2.** The secondary valence of  $Co^{-3}$  is 6. Calculate the number of moles of AgCl precipitated, when excess of  $AgNO_3$  solution is added to 1.5 lit of 0.2 M.  $CoCl_3$ .  $5NH_3$  solution



3. The number of stereo isomers of the compound is

$$\left[ Co \left( C_2 O_4 \right) \left( NH_3 \right)_2 \left( NO_2 \right)_2 \right]$$



**4.** If  $\left[Co\left(NH_3\right)_6\right]^{y^+}$  follows EAN rule, the oxidation state of Co is (At. No. of CO = 27 and At. No. of Kr = 36).



**5.** Which of the following is correct value of x in  $Cr(CO)_{\chi}$ ?



**6.** The spin only magnetic moment of  $\left[FeBr_4\right]^-$  is 5.92 BM. Predict the geometry of complex ion.



**7.** Arrange the following in the increasing order of stabilization energy of following in presence of strong field ligands.



**8.** Give the stabilization energy of  $d^7$  electrons in presence of strong and weak field ligands.



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**9.** In  $\left[Ti\left(H_2O\right)_6\right]^{3+}$  complex  $Ti^{3+}$  has only one electron in 3d-orbital. Its spectrum shows a single absorption peak of 5000Å then correcponding energy is



**10.** 
$$Zn^{+2} + 2NH_3 \Leftrightarrow \left[ Zn \left( NH_3 \right)_2 \right]^{+2}, K_1 = 2 \times 10^{-3}$$

$$\left[Zn(NH_3)_2\right]^{+2} + 2NH_3 \Leftrightarrow \left[Zn(NH_3)_4\right]^{2+}, K_2 = 1.5 \times 10^{-3}$$

Find out the instability constant?



C.U.Q

1. The following does not give a precipitate either with  $AgNO_3$  or  $BaCL_2$ 

A. 
$$\left[ Co(NH_3)_5 Cl \right] SO_4$$

$$B. \left[ Co \left( NH_3 \right)_3 Cl_3 \right]$$

$$\mathsf{C.} \left[ \mathsf{Co} \left( \mathsf{NH}_3 \right)_4 \mathsf{Cl}_2 \right] \mathsf{Cl}$$

D. 
$$\left[Co(NH_3)_2Cl_4\right]Cl$$

### Answer: 2



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**2.** Which of the following has the highest molar conductivity in solution?

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

B. 
$$\left[Co(NH_3)_5Cl\right]Cl_2$$

C. 
$$\left[ Co(NH_3)_4 Cl_2 \right] Cl$$

$$\mathsf{D.}\left[\mathit{Co}\!\left(\mathit{NH}_3\right)_6\!\mathit{Cl}_3\right]$$



- **3.** The transition metals have a strong tendency to form complaexes because of
- (i) smaller sizes of the metal ions
- (ii) variable oxidation states
- (iii) high ionic charges of metal ions
- (iv) availability of vacant d-orbitals for bond formation.
  - A. I only
  - B. ii only

C. I & ii	
D. I,ii,iii	
Answer: 4	
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<b>4.</b> The ionizable valency of Ni in $Ni(CO)_4$ is	
<b>A.</b> 2	
B. 4	
C. 0	

**D.** 1

**Answer: 3** 

5. According to Werner's theory transition metals possesses
A. only one type of valency
B. two types of valencies
C. three types of valencies
D. four types of valencies
Answer: 2
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6. The primary valency of the metal ion is satisfied by
6. The primary valency of the metal ion is satisfied by
6. The primary valency of the metal ion is satisfied by  A. neutral molecules

D. all	the	above
D. all	the	above



**7.** No of ionizable & non-ionizable  $Cl^-$  ions in  $CoCl_35NH_3$  representively are

**A**. 3, 0

B. 2, 1

**C**. 1, 2

D. 0, 3

### Answer: 2



8. Central metal ion in complex compound acts as
A. Lewis acid
B. Lewis base
C. Arrhenius acid
D. Arrhenius base
Answer: 1
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9. Which one of the following acts as a Lewis base in complexes
A. <i>CO</i> <sub>2</sub>
B. $BF_3$
C. <i>NH</i> <sub>3</sub>



- 10. Potassium ferrocyanide is a
  - A. Complex salts
  - B. Normal salts
  - C. Double salts
  - D. Basic salts

### **Answer: 1**



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11. Example for a coordination compound is

B. 
$$K_2SO_4$$
.  $Al_2(SO_4)_3$ .24 $H_2O$ 

C. 
$$CoCl_3.6NH_3$$

D. 
$$FeSO_4$$
.  $\left(NH_4\right)_2SO_4.6H_2O$ 



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12. in which of the following transition metal complexes does the metal exhibits zero oxidation state.

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

$$\mathsf{B.}\left[\mathit{Fe}\big(H_2O\big)_6\right]\!\!\mathsf{SO}_4$$

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

$$\mathsf{D.}\left[\mathit{Fe}\Big(H_2O\Big)_6\right]\!Cl_3$$

# Answer: 3 **Watch Video Solution** 13. The number of ions formed form a formula unit of potassium ferricyanide in solution is **A.** 2 B. 4



**C**. 5

D. 6

A. M-SCN B. M-NCS C. M-CNS D. M-CSN Answer: 2 **Watch Video Solution 15.** Number of chlorides satisifying secondary valency in  $CoCl_34NH_3$ **A.** 2 **B**. 3 **C**. 4 D. 1 **Answer: 1** 

**16.** Which of the following is cationic complex?

A. 
$$K_4$$
 [ $Fe(CN)_6$ ]

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

$$\mathsf{C.}\left[\mathit{Co}\left(\mathit{NH}_3\right)_3 \mathit{Cl}_3\right]$$

D. 
$$\left[ Cu(NH_3)_4 \right] SO_4$$

### Answer: 4



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**17.** The no. of moles of AgCl obtained when excess  $AgNO_3$  is added to one mole of  $\left[Cr\Big(NH_3\Big)_4Cl_2\right]Cl$ 

**A.** 1 **B**. 2 **C**. 3 D. 4 **Answer: 1** 



**18.** Positive ligand in sodium nitroprusside complex is

- - **A.** *Na* <sup>+</sup>
  - B. *NO* +
  - C. *CN* +
  - $\mathrm{D.}\,H_3O^{\,+}$



**19.** The no. of moles of AgCl precipitated when excess of  $AgNO_3$  is mixed with one mole of  $\left[ Cr(NH_3)_3 Cl_3 \right]$  is

**A.** 0

B. 1

**C**. 2

**D**. 3

**Answer: 1** 



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A. Cobalt trichloride penta amonium

**20.** IUPAC name of the complex  $CoCl_35NH_3$  is

- B. Penta amine carbonyl chloride
- C. Trichloro penta amino cabalt
- D. Pentaaminechlorocobalt (III) chloride



**21.** The property of possessing atleast one atom that is attached to four non-identical groups in tetrahedral geomentry is called

- A. polarisation
- B. chirality
- C. enantiomerism
- D. meridionity

### Answer: 2



### 22. A racemic mixture has a net rotation

A. to right of chirality

B. to left of original plane

C. to right of original plane

D. zero

### **Answer: 4**



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23. Optical isomer have

A. property of chirality

B. almost identical chemist properties

C. almost identical physical properties

D. all the above

### **Answer: 1**



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### **24.** The effective atomic number of iron in $\left[Fe(CN)_6\right]^{3-}$ is

**A.** 34

**B.** 36

**C**. 37

**D.** 35

### Answer: 4



25. Which does not obey EAN rule?

A. 
$$Fe(CO)_5$$

$$B. K_4 \Big[ Fe(CN)_6 \Big]$$

$$C. \left[ Cu(NH_3)_4 \right] SO_4$$

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

### **Answer: 3**



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**26.** The effective atomic number of central metal ion is wrongly calculated in the following complex?

A. In  $\left[Ni(CO)_4\right]$  the EAN of Ni is 36

B. In  $K_2$   $Ni(CO)_4$  the EAN of Ni is 36

D. In  $\left[ Cr(NH_3)_6 \right] Cl_3$  the EAN of Cr is 33

C. In  $K_3$   $Fe(CO)_6$  the EAN of Fe is 35

### Answer: 1



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27. According to effective atomic number rule the central metal acquires:

A. inert gas configuration

C. duplet

B. octet

D. quartet

### Answer: 4



28. The shape of the complex species will be square planar if its coordination number is

- **A.** 2
- B. 6
- **C**. 5
- D. 4

### Answer: 4



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**29.** Which of the following is outer orbital complex?

- A.  $\left[ CoF_6 \right]^{-3}$  B.  $\left[ Cu \left( H_2O \right)_6 \right]^{+2}$

$$\mathsf{C.} \left[ \mathsf{Co} \left( \mathsf{NH}_3 \right)_6 \right]^{+2}$$

D. Both 1 and 2

### **Answer: 1**



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### **30.** $sp^3d^2$ hybridisation is present in

A. 
$$\left[ CoF_6 \right]^{-3}$$

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

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

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

### Answer: 2



**31.** The type of hybridisation present in  $\left[Cu(NH_3)_4\right]^{+2}$  ion is

A.  $sp^3$ 

B.  $dsp^2$ 

 $C. sp^3d$ 

D.  $sp^3d^2$ 

### Answer: 3



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**32.** The shape of  $\left[CoF_6\right]^{-3}$  is

A. Square planner

B. Trigonal bipyramidal

C. Octahedral

D. Tetrahedral

### **Answer: 1**



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33. The hybridisation of metal atom & geometry of complex in

$$\left[Ag\left(NH_3\right)_2\right]^{-3}$$
 are

- A. sp, linear
- B.  $sp^2$ , linear
- C.  $sp^2$ , trigonal planner
- D. sp, angular

### **Answer: 1**



**34.** The magnetic moment of  $\left[Ni(Cl)_4\right]^{2-}$  is

A. 2.85*BM* 

B. 1.83*BM* 

C. 4.86*BM* 

D. 5.95*BM* 

### **Answer: 1**



**35.** Which of the following system has maximum number of the unpaired electrons in an inner octahedral complex?

A.  $d^4$ 

B. *d*<sup>9</sup>

 $C. d^7$ 

D.  $d^5$ 

### **Answer: 1**



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### 36. In an octahedral crystal field, the correct set of low orbitals are

A. 
$$d_{xy}$$
,  $d_{xz}$ ,  $d_{x^2-y^2}$ 

B. 
$$d_{x^2-y^2}$$
,  $d_{z^2}$ 

C. 
$$d_{xy}$$
,  $d_{xz}$ ,  $d_{yz}$ 

D. 
$$d_{xy}$$
,  $d_{x^2-y^2}$ 

### Answer: 3



**37.** For the same metal, the stabiliing energies of tetrahedral and octahedral complex are related as

A. 
$$\Delta_t = \Delta_0$$

$$B. \Delta_t \times 4 = \Delta_0 \times 6$$

$$\mathsf{C.}\,\Delta_t\times 9=\Delta_0\times 4$$

D. 
$$\Delta_t \times 6 = \Delta_0 \times 4$$

#### **Answer: 3**



**38.** The orbitals having lower energy in tetrahedral complexs according to CFT are

$$A. d_{xy}, d_{yz}, d_{z^2}$$

B. 
$$d_{xy}$$
,  $d_{yz}$ ,  $d_{x^2-y^2}$ 

- $\mathsf{C.}\,d_{\mathit{xy}},\,d_{\mathit{yz}},\,d_{\mathit{zx}}$
- D.  $d_{x^2-y^2}$ ,  $d_{z^2}$



- 39. The metal which does not form poly nuclear carbonly is
  - A. Mn
  - B. Fe
  - C. Cr
  - D. Co

### **Answer: 3**



**40.** Nessler's reagent is

A.  $K_2$  [ $HgI_4$ ]

 $B. K_2 \Big[ HgI_2 \Big]$ 

 $C. K_2[HgCl_4]$ 

D.  $HgI_2$ 

### Answer: 1



**41.** Among the following metal carbonyls the C - O bond order is lowest in .

A. 
$$\left[V(CO)_6\right]^{-1}$$

B.  $Fe(CO)_5$ 

 $\mathsf{C.}\left[\mathit{Cr}(\mathit{CO})_{6}\right]$ 

 $D. Mn(CO)_6]^+$ 

#### Answer: 1



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### **LEVEL-I**

- 1. Which of the following is not a draw back of Werner's theory?
  - A. does not explain the valency of metal ions in the complex
  - B. does not give any explanation for the colour of complex

compounds

C. does not explain the magnetic behaviour of complex

compounds

D. does not correlate electronic configuratio of the metal with the formation of complex

#### **Answer: 1**



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- **2.** Aqueous solution of  $\left[Co\left(H_2O\right)_5SO_4\right]Cl$  gives precipitate with
  - A.  $BaCl_{2(aq)}$
  - $B.\mathit{AgNO}_{3(\mathit{aq})}$
  - C. both 1 and 2
  - D. neither 1 nor 2

### Answer: 2



**3.** Silver chloride dissolves in excess ammonia due to the formation of a soluble compplex whose formula is

A. 
$$\left[Ag\left(NH_3\right)\right]Cl$$

B. 
$$\left[Ag\left(NH_3\right)_2\right]Cl$$

$$C. \left[ Ag(NH_3)_3 \right] Cl$$

D. 
$$\left[Ag\left(NH_3\right)_4\right]Cl$$

### Answer: 2



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**4.** Which is anionic complex?

A. Ferrous ammonium sulphate

B. Carnallite

- C. Potassium ferrocyaniode
- D. Gypsum



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### **5.** $Zn^{+2}$ dissolves in excess of NaOH due to the formation of

- A. Soluble  $Zn(OH)_2$
- B. Soluble  $Na_2$   $\left[ Zn(OH)_4 \right]$
- C. Soluble  $Na[Zn(OH)_3]$
- D. ZnO

### Answer: 2



<b>6.</b> Example of neutral	complex	compound in	the following is
or = //op. o o o			

- A.  $CoCl_3.6NH_3$
- B.  $CoCl_3.5NH_3$
- C.  $CoCl_3.4NH_3$
- D.  $CoCl_3$ .3 $NH_3$



- **7.** Which of the following releases metal slowly to give uniform coating in electroplating is?
  - A. metal salts
  - B. double salts
  - C. complex salt

D. alums

#### Answer: 3



- **8.** IUPAC name of the complex  $\left[Cu(NH_3)_4\right]SO_4$  is
  - A. cuprammonium sulphate
  - B. copper sulphate tetraammonia
  - C. tetraamminecopper(II) sulphate
  - D. copper ammonium(IV) sulphate

#### **Answer: 3**



9. Which of the following does not exhibit optical isomerism?

A. 
$$\left[ Cr(NH_3)_3 \right] Cl_3$$

$$B. \left[ Cr(en)_3 \right] Cl_3$$

C. 
$$\left[ Cr(en)_2 Cl_2 \right] Cl$$

D. 
$$\left[ Cr(en) \left( NH_3 \right)_2 Cl_2 \right] Cl$$

#### **Answer: 1**



**10.** Co-ordination compounds  $\left[Pt(NH_3)_3(NCS)\right]$  and

$$\left[Pt\left(NH_3\right)_2Cl_2\right]Cl$$
 are example of ...... Isomerism

A. Co-ordination

B. Ionization

- C. Linkage
  D. Optical
- **Answer: 3**



#### 11. ISOMERISM IN COORDINATION COMPOUNDS

- A. Tetrahedral complex
- B. Square planar complex
- C. Linear Complex
- D. Planar triangle complexes

#### Answer: 2



12. Stable complex based on EAN rule

A. 
$$K_4[Fe(CN)_6]$$

$$\mathbf{B.} \left[ Co \left( NH_3 \right)_5 Cl \right] Cl_2$$

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

D. all the above

#### **Answer: 4**



**13.** Deduce the structures of  $\left[NiCl_4\right]^{2-}$  and  $\left[Ni(CN)_4\right]^{2-}$  considering the hybridisation of the metal ion. Calculate the magnetic moment (spin only) of the species.

A.  $sp^3$ 

B.  $dsp^2$ 

 $C. sp^3d$ 

D.  $sp^3d^2$ 

#### Answer: 1



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# **14.** Which one of the following has a square planar geometry?

(Co = 27, Ni = 28, Fe = 26, Pt = 78)

A. 
$$\left[COCl_4\right]^{-2}$$

B.  $\left[PtCl^4\right]^{-2}$ 

C.  $\left[NiCl_4\right]^{-2}$ 

D.  $\lceil FeCl_4 \rceil^{-2}$ 

### Answer: 2



<b>15.</b> Which of the following is paramagnetic?
A. $Ni(CO)_4$
B. $Fe(CO)_5$
C. <i>V(CO)</i> <sub>6</sub>
D. <i>Cr(CO)</i> <sub>6</sub>
Answer: 3
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<b>16.</b> The number of ions formed when cuprammonium sulphate is

**B.** 2

**A.** 1

**C**. 4

D. zero

#### Answer: 2



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**17.** Which of the following is correct arrangement of ligands in terms of field strength

A. 
$$Cl^- < F^- < NCS^- < NH_3 < CN^-$$

$$B. NH_3 < F^- < Cl^- < NCS^- < CN^-$$

$$C. Cl^- < F^- < NCS^- < CN^- < NH_3$$

D. 
$$NH_3 < CN^- < NCS^- < Cl^- < F^-$$

#### **Answer: 1**



**18.** In which of the following octahedral complexes of Co (at. no. 27), will the magnitude of  $\Delta_o$  be the highest?

A. 
$$\left[Co\left(C_2O_4\right)_3\right]^{3}$$

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

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

D. 
$$\left[ Co(CN)_6 \right]^{3}$$

#### Answer: 4



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**19.** If  $\Delta_0 < P$ , the correct electronic configuration for  $d^4$  system will be (P = paiting energy)

A. 
$$t_{2g}^4 e_g^0$$

- $\mathbf{B.}\ t_{2g}^{3}e_{g}^{1}$
- C.  $t_{2g}^{0}e_{g}^{4}$
- D.  $t_{2g}^{2}e_{g}^{2}$



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- 20. Ammonium ions are detected with
  - A. Nessler's reagent
  - B. Borsch reagent
  - C. Tollen's reagent
  - D. Fehling's solution

#### Answer: 1



**21.** 
$$\left[ \left( Ph_3P \right)_3 RhCl \right]$$
 is a familiar catalyst used in

A. hydrogenation of oils

B. hydrogenation of alkenes

C. dehydration of alcohols

D. dehydration of aldehydes

#### Answer: 2



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**22.** Metals those can be extracted commercailly with aqueous solution of sodium cyanide as complexes are

A. Au and Ag

- B. Fe and Ag
- C. Au and Mg
- D. Hg and Fe



**23.** If 
$$Ag^+ + NH_3 \Leftrightarrow \left[ Ag(NH_3) \right]^+$$
,  $K_1 = 3.5 \times 10^{-3}$  and  $\left[ Ag(NH_3) \right]^+ + NH_3 \Leftrightarrow \left[ Ag(NH_3)_2 \right]^+$ ,  $K_2 = 1.74 \times 10^{-3}$ . The

formation constant of 
$$\left[Ag(NH_3)_2\right]^+$$
 is :

- A.  $1.7 \times 10^{-3}$
- B.  $5.92 \times 10^{-6}$
- C.  $1.8 \times 10^3$ 
  - D.  $1.7 \times 10^7$



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#### **LEVEL-II**

- **1.** The oxidation number of cobalt in  $K[Co(CO)_4]$  is
  - A. + 1
  - B. + 3
  - **C**. -1
  - D. -3

#### **Answer: 3**



2. EDTA has coordination number A. Monodentate ligand B. Bidentate ligand C. Quadridentate ligand D. Hexadentate ligand Answer: 4 Watch Video Solution 3. en' is an example of a A. Monodentate ligand B. Bidentate ligand C. Tridentate ligand D. Hexadentate ligand



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- **4.** IUPAC name of  $K_2 \Big[ \mathit{PtCl}_6 \, \Big]$  is
  - A. Potassiumhexachloroplatinum
  - B. Potassiumhexachloroplatinum(IV)
  - C. Potassiumhexachloroplatinum(IV)
  - D. Dipotassium hexa chloro plantinum

#### **Answer: 3**



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**5.** IUPAC name of  $\left[Pt\left(NH_3\right)_3(Br)\left(NO_2\right)CI\right]CI$  is

- A. Trimminechlorobromonitroplatinum (IV) chloride
- B. Trimminebromonitrochloroplatinum (IV) chloride
- C. Trimminebromochloronitroplatinum (IV) chloride
- D. Trimminenitrobromochloroplatinum (IV) chloride



- 6. Tetrammine diaqua copper (II) hydroxide is given by the formula
  - A.  $\left[ Cu \left( NH_3 \right)_4 \right] (OH)_2.2H_2O$
  - B.  $\left[Cu\left(NH_3\right)_4(OH)_2\right].2H_2O$
  - C.  $\left[Cu\left(NH_3\right)_4\left(H_2O\right)_2\right](OH)_2$
  - D.  $\left[ Cu(NH_3)_4 (H_2O)(OH)_2 \right]$



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**7.** The compexes

$$\left[ Co(NH_3)_6 \right] \left[ Cr(C_2O_4)_3 \right] \qquad \text{and} \qquad$$

$$\left[ Cr \left( NH_3 \right)_6 \right] \left[ Co \left( C_2 O_4 \right)_3 \right]$$

- A. Geometrical isomerism
- B. Linkage isomerism
- C. Coordination isomerism
- D. Ionization isomerism

#### **Answer: 3**



**8.** Which of the following complex or the complex ion will show geometrical isomerism?

A. 
$$\left[ Pt \left( NH_3 \right)_2 Cl^2 \right]$$

$$B. \left[ Pt \left( NH_3 \right) Cl_5 \right]^{-1}$$

C. 
$$\left[ Pt \left( NH_3 \right)_5 Cl \right]^{3+}$$

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

#### Answer: 1



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**9.** Which isomerism is exhibited by

$$\left[Co(NH_3)_3(H_2O)_3\right]Cl_3?$$

A. Geometrical isomerism

- B. Linkage isomerism
- C. Coordination isomerism
- D. Ionization isomerism



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## **10.** $\left[Co\left(NH_3\right)_4\left(NO_2\right)_2\right]CI$ exhibits

- A. Linkage isomerism, ionization isomerism and geometrical isomerism
- B. Ionisation isomerism, geometrical isomerism and optical isomerism
- C. Linkage isomerism, geometrical isomerism and optical isomerism

D. Linkage isomerism, ionization isomerism and optical isomerism

#### Answer: 1



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11. Which one of the following is an example of coordination isomerism?

A. 
$$\left[Co(NH_3)_5Br\right]SO_4$$
 and  $\left[Co(NH_3)_5SO_4\right]Br$ 

B. 
$$\left[Co(NH_3)_5NO_2\right]Cl_2$$
 and  $\left[Co(NH_3)_5ONO\right]Cl_2$ 

C. 
$$\left[Cr\left(H_2O\right)_6\right]Cl_3$$
 and  $\left[Cr\left(H_2O\right)_5Cl\right]Cl_2$ .  $H_2O$ 

D. 
$$\left[ Cr \left( NH_3 \right)_6 \right] \left[ Co \left( CN \right)_6 \right]$$
 and  $\left[ Co \left( NH_3 \right)_6 \right] \left[ Cr \left( CN \right)_6 \right]$ 

#### Answer: 4



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**12.**  $\left[\mathit{FeF}_{6}\right]^{3+}$  has Fe atom ...... Hybridized with unpaired .....

Electrons.

A. 
$$d^2sp^3$$
, 4

B. 
$$d^2sp^3$$
,5

C. 
$$sp^3d^2$$
,5

D. 
$$sp^3d^2$$
, 3

#### Answer: 3



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**13.** How many EDTA molecules are required to make an octahedral complex with a  $Ca^{2+}$  ion?

A. two

- B. six
- C. three
- D. one



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**14.** The d electron congfiguration of  $Cr^{2+}$ ,  $Mn^{2+}$ ,  $Fe^{2+}$  and  $Ni^{2+}$  are  $3d^4$ ,  $3d^5$ ,  $3d^6$  and  $3d^8$  respectively. Which one of the following aqua complexes will exhibit the minimum paramagnetic behaviour? (At. No. Cr = 24, Mn = 25, Fe = 26, Ni = 28)

A. 
$$\left[ Fe \left( H_2 O \right)_6 \right]^{+2}$$

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

$$C. \left[ Cr \left( H_2 O \right)_6 \right]^{+2}$$

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



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**15.** The expected spin only magnetic mometum for  $\left[Fe(CN)_6\right]^{4-}$  and

$$[FeF_6]^{3+}$$
 are

- A. 1.73 and 1.73 B.M
- B. 1.73 and 5.93 B.M
- C. 0.0 and 1.73 B.M
- D. 0.0 and 5.92 B.M

#### **Answer: 4**



**16.** The volume (in mL) of  $0.1MAgNO_3$  required for complete precipitation of chloride ions present in 30mL of 0.01M solution of

$$\left[Cr\left(H_2O\right)_5Cl\right]Cl_2$$
, as silver chloride is close to:

- **A.** 3
- B. 4
- **C**. 5
- **D.** 6

#### Answer: 4



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**17.** Among the following ions which one has the highest paramagnetism?

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

$$B. \left[ Fe \left( H_2 O \right)_6 \right]^{2+}$$

$$C. \left[ Cu \left( H_2 O \right)_6 \right]^{2+}$$

D. 
$$\left[ Cu \left( H_2 O \right)_2 \right]^{2+}$$



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### 18. Which of the following complex is an outer orbital complex?

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

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

$$\mathsf{C.}\left[\mathit{Co}\big(\mathit{NH}_3\big)_6\right]^{3+}$$

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

#### Answer: 4

**19.** The EAN of cobalt in the complex ion  $\left[Co(en)_2Cl_2\right]^+$  is

- **A.** 27
- **B.** 36
- **C**. 33
- D. 35

#### Answer: 2



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**20.** For an octahedral complex, which of the following d electron configuration will give maximum crystal-field stabilisation energy?

A. High spin  $d^6$ 

B. Low - spin  $d^4$ 

C. Low spin  $d^5$ 

D. High - spin  $d^7$ 

### Answer: 3



## 21. Chromium compound widely used in tanning of leather is

 $\mathsf{B.}\mathit{Cr}_2O_2\mathit{Cl}_2$ 

A.  $Cr_2O_3$ 

 $C. Cr_2O_3$ 

Answer: 4

( )

D.  $K_2 SO_4 Cr_2 (SO_4)_3 24H_2 O$ 



**22.** Oxidation state of central metal atom and geometry for the Wilkinsons catalyst.  $\left[Rh\left(Ph_3P\right)_3Cl\right]$  used for hydrogenation of alkenes are ......

- A. 1, tetrahedral
- B. 2, tetrahedrak
- C. 1, square planar
- D. 2, square planar

#### **Answer: 3**



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**23.** Coordination compounds have great importance in biological systems. In this context which of the following statements is

incorrect:

A. Chlorophylls are green pigments in plants and contain calcium.

- B. Haemoglobin is the red pigment of blood and contains iron.
- C. Cyanocobalamin is  $B_{12}$  and contains cobalt.
- D. Carboxypeptidase-A is an enzyme and contains zinc.

#### **Answer: 1**



#### **LEVEL-III**

**1.** One mole of complex compound  $Co(NH_3)_5Cl_3$  gives 3 moles of ions on dissolution in water. One mole of same complex reacts with two moles of  $AgNO_3$  to yield two moles of AgCl(s). The complex is:

A. 
$$\left[Co(NH_3)_4Cl\right]Cl_2$$
.  $NH_3$ 

$$\operatorname{B.}\left[\operatorname{Co}\left(\operatorname{NH}_3\right)_5\operatorname{Cl}\right]\operatorname{Cl}_2$$

$$C. \left[ Co(NH_3)_3 Cl_3 \right] 2. NH_3$$

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



- 2. The primary and secondary valencies of chromium in the complex ion, dichlotodioxalatochromoium (III), are respectrively
  - A. 3, 4
  - B. 4, 3
  - C. 3, 6
  - D. 6, 3



**3.** In the complex with formula  $MCl_3.4H_2O$  the co-ordination number of the metal M is six. And there is a no molecule of hydration in it. The volume of 0.1 M  $AgNO_3$  solution needed to precitate the free chloride ions in 200 ml of 0.01 M solution of the complex is

- A. 40 ml
- B. 20 ml
- C. 60 ml
- D. 80 ml

#### Answer: 2



- **4.** The molar ionic conductances of octahedral complexes.
- (I)  $PtCl_4.5NH_3$  (II)  $PtCl_4.4NH_3$
- (III)  $PtCl_4.3NH_3$  (IV)  $PtCl_4.2NH_3$ 
  - A. I < II < III < IV
  - B. IV < III < II < I
  - C.III < IV < II < I
  - D. IV < III < I < II



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**5.** The coordination number of a central metal atom in a complex is determined by:

A. the number of ligands around a metal ion bonded by sigma

bonds

B. the number of ligarnds around a metal ion bonded by  $\pi$ -bonds

C. the number of ligands around a metal ion bounded by sigma and pi bonds both

D. the number of only anionic ligands bonded to the metal ion.

#### Answer: 1



- **6.** Among the following which are ambidentate ligands?
- (a)  $NO_2^-$  (b)  $NO_3^-$  (c)  $EDTA^+$
- (d)  $C_2O_4^{2+}$  (e)  $SCN^-$  (f)  $H_2NCH_2CH_2NH_2$ 
  - A. (a) and (b)

- B. (c) and (d)
- C. (a) and (f)
- D. (a) and (e)



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diethylene triamine and EDTA are respectively

7. The number of donor sites in dimethyl glyoxime, glycinato,

- (a) 2, 2, 3 and 4
- (b) 2, 2, 3 and 6 (c) 2, 2, 2 and 6
- (d)2, 3, 3 and 6.
  - A. 2, 2, 3 and 6
  - B. 2, 2, 3 and 4

- C. 2, 2, 2 and 6
- D. 2, 3, 3 and 6



- **8.** The IUPAC name of the coordination compound  $K_3 \left[ Fe(CN)_6 \right]$  is:
  - A. potassium hexacyanoferrate (II)
  - B. potassium hexacyanoferrate (III)
  - C. potassium hexacyanoiron (II)
  - D. tripotassium hexacyanoion (II)

#### Answer: 2



9. Which one of the following has largest number of isomers?

A. 
$$\left[ Ru \left( NH_3 \right)_4 Cl_2 \right]^+$$

B. 
$$\left[ Co(NH_3)_5 Cl \right]^{2+}$$

C. 
$$\left[ Ir \left( PR_3 P \right)_2 H(CO) \right]^{2+}$$

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

#### Answer: 4



10. Which of the following compounds shows optical isomerism?

A. 
$$\left[ Cr \left( C_2 O_4 \right)_3 \right]^{3-}$$

B. 
$$\left[ Co(CN)_6 \right]^{3}$$

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

D. 
$$\left[ ZnCl_4 \right]^{2-}$$



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**11.** In which of the following pairs both the complexes do not show optical isomerism?

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

B. 
$$\left[Co(en)_3\right]Cl_3$$
,  $Cia - \left[Co(en)_2Cl_2\right]Cl$ 

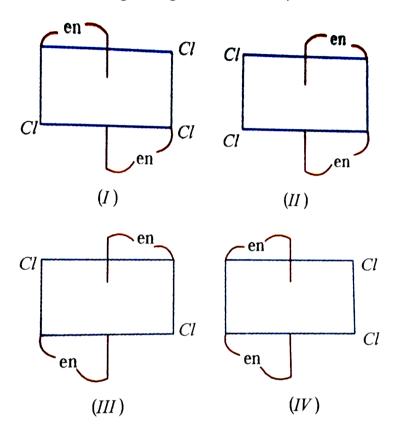
C. 
$$[PtCl_2(en)]$$
,  $[NiCl_2Br_2]^{2-}$ 

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

#### **Answer: 3**



### 12. Of the following configurations, the optical isomers are



A. *I&II* 

B. *I&III* 

C. II&IV

D. II&III



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- **13.** Which kind of isomerism is shown by  $Co(NH_3)_4Br_2Cl$ ?
  - A. Optical and ionisation
  - B. Geometrical and optical
  - C. Geometrical and ionisation
  - D. Only geometrical

### **Answer: 3**



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14. Which of the following does not have optical isomer?

A. 
$$\left[Co(en)_3\right]Cl_3$$

$$B. \left[ Co \left( NH_3 \right)_3 Cl_3 \right]$$

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

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



**15.** The complex  $K_3[Fe(CN)_6]$  should have a spin only magentic of

A. 
$$\sqrt{8}BM$$

B. 
$$2\sqrt{5}BM$$

C. 
$$\sqrt{3}BM$$

D. 
$$\sqrt{6}BM$$



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16. Which one of the following complexes? [Atomic numbers, Mn=25,

A. 
$$[Fe(CN)_6]^{4-}$$

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

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

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

### **Answer: 4**



**17.** The correct order of magnetic moments (spin values in B.M.) among is:

A. 
$$\left[ \mathit{MnCl}_4 \right]^{2-} > \left[ \mathit{CoCl}_4 \right]^{2-} > \left[ \mathit{Fe}(\mathit{CN})_6 \right]^{4-}$$

$$B. \left[ MnCl_4 \right]^{2-} > \left[ Fe(CN)_6 \right]^{4-} > \left[ CoCl_4 \right]^{2-}$$

$$C. \left[ Fe(CN)_6 \right]^{4-} > \left[ MnCl_4 \right]^{2-} > \left[ CoCl_4 \right]^{2-}$$

D. 
$$\left[Fe(CN)_6\right]^{4-} > \left[CoCl_4\right]^{2-} > \left[MnCl_4\right]^{2-}$$

#### Answer: 1



**18.** Which one of the following has lowest value of paramagnetic behaviour?

A. 
$$[Fe(CN)_6]^3$$

B. 
$$\left[ Co(CN)_6 \right]^{3}$$

C. 
$$\left[ Cr(CN)_6 \right]^{3}$$

D. 
$$\left[Mn(CN)_6\right]^{3}$$



- 19. Which of the following statements is not correct?
  - A. The complexes  $\left[NiCl_4\right]^{2-}$  and  $\left[Ni(CN)_4\right]^{2-}$  differ in their magnetic properties.
  - B. The complexes  $\left[NiCl_4\right]^{2^-}$  and  $\left[Ni(CN)_4\right]^{2^-}$  differ in the state of hybridisation of nickel
  - C. The complexes  $\left[NiCl_4\right]^{2-}$  and  $\left[Ni(CN)_4\right]^{2-}$  differ in geometry
  - D. The complexes  $\left[NiCl_4\right]^{2-}$  and  $\left[Ni(CN)_4\right]^{2-}$  differ in primary valancies of nickel.



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20. The compound having a tetrahedral geometry is .

A. 
$$\left[ PdCl_4 \right]^{2-}$$

B. 
$$\left[Ni(CN)_4\right]^{2}$$

C. 
$$\left[ Pd(CN)_4 \right]^{2}$$

D. 
$$\left[NiCL_4\right]^{2}$$

### Answer: 4



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**21.** Geometry, hybridisation and magnetic moment of the ions  $[Ni(CN)_4]^{2-}$ ,  $[MnBr_4]^{2-}$  and  $[FeF_6]^{3-}$  are .

 $dsp^2$ ,  $sp^3$ ,  $sp^3d^2$ : 0, 5.9, 4.9 C. Square planar, tetrahedral, octahedral:  $dsp^2$ ,  $sp^3$ ,  $d^2sp^3$ : 5.9, 4.9, 0 D. Square planar, tetrahedral, octahedral:

square

square

planar,

planar,

octahedral:

octahedral:

# Answer: 4



 $dsp^3$ ,  $sp^3$ ,  $sp^3d^2$ : 0, 5.9, 4.9

A. Tetrahedral,

B. Tetrahedral,

 $sp^3$ ,  $dsp^3$ ,  $sp^3d^2$ : 5.9, 0, 4.9

**22.**  $\left[Cr(H_2O)_6\right]Cl_3$  (at no. of Cr = 24) has a magnetic moment of 3.83*B*. *M*. The correct distribution of 3*d* electrons the chromium of the complex.

A. 
$$3d_{xy}^1$$
,  $3d_{yx}^1$ ,  $3d_{zx}^1$ 

B.  $3d_{xy}^1$ ,  $3d_{yx}^1$ ,  $3d_{z^2}^1$ 

C.  $3d_{(x^2-y^2)^1,3d_{z^2}^1,3d_{xz}^1}$ 

D.  $3d_{xy}^1$ ,  $3d_{(x^2-y^2)^1, 3d_{yz}^1}$ 

### Answer: 1



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will the magnitude of  $\Delta_o$  be the highest?

23. In which of the following octahedral complexes of Co (at. no. 27),

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

$$B. \left[ Co \left( C_2 O_4 \right)_3 \right]^{3-}$$

$$\mathsf{C.}\left[\mathit{Co}\big(H_2\mathsf{O}\big)_6\right]^{3+}$$

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



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**24.** Which of the following is a correct Irving-Williams order? (Tendency of complex formation)

A. 
$$Mn^{2+} < Fe^{2+} < Co^{2+} < Ni^{2+}$$

B. 
$$Ni^{2+} < Co^{2+} < Fe^{2+} < Mn^{2+}$$

$$C. Fe^{2+} < Mn^{2+} < Ni^{2+} < Co^{2+}$$

D. 
$$Co^{2+} < Mn^{2+} < Fe^{2+} < Ni^{2+}$$

### Answer: 1



A. 
$$Cl_1 < F^- < C_2O_4^{2-} < NO_2^- < CN^-$$

B. 
$$CN^- < C_2O_4^{2-} < Cl^- < NO_2^- < F^-$$

$$C. C_2O_4^{2-} < F^- < Cl^- < NO_2^- < CN_1$$

$$D.F^- < Cl^-NO_2^- < C_2O_4^2^-$$



(III)

## **26.** Which of the following ligands is calles $\pi$ - acceptors?

$$CO$$
  $CN^ NO^+$   $(I)$   $(II)$   $(III)$ 

(I)

B. I, II only

C. I, III only

D. III only



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### **27.** In $Fe(CO)_5$ , the Fe - C bond possesses:

- A. π-character only
- B.  $\sigma$  character only
- C. ionic character only
- D. both  $\sigma \& \pi$  character

### **Answer: 4**



1. Which of the following has an optical isomer?

(en=ethylenediamine)?

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

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

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

D. 
$$\left[Co\left(H_2O\right)_4 en\right]^{3+}$$

### **Answer: 3**



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**2.** Which of the following facts about the complex  $\left[Cr(NH_3)_6\right]Cl_3$  is wrong?

- A. The complexes innvolves  $d^2sp^3$  hybridisation and is octahedral
- B. The complex is paramagnetic

shape

- C. The complex is an outer orbital complex
- D. The complex gives white prepipitate which silver nitrate solution

### Answer: 3



**3.** Deduce the structures of  $\left[NiCl_4\right]^{2-}$  and  $\left[Ni(CN)_4\right]^{2-}$  considering the hybridisation of the metal ion. Calculate the magnetic moment (spin only) of the species.

### A. 1.8*BM*

B. 5.86*BM* 

C. 2.82*BM* 

D. 3.82*BM* 

### Answer: 3



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**4.** Which among the following will be named as dibromidobis (ethylene diamine) chromium (III) bromide?

$$A. \left[ CrBr_2(en)_2 \right]^+$$

B. 
$$Cr\left[Br_2(eN)_2^{-}\right]^{-}$$

C. 
$$\left[ CrBr_2(en) \right]^+$$

D. 
$$\left[ CrBr_2(en) \right]^{+2}$$

### Answer: 1

**5.** Which of the following complex species is not expected to exhibit optical isomerism ?

A. 
$$\left[Co(en)_2Cl_2\right]^+$$

$$B. \left[ Co(NH_3)_3 Cl_3 \right]$$

C. 
$$\left[Co(en)\left(NH_3\right)_2Cl_2\right]^+$$

D. 
$$[Co(en)_3]^{3+}$$

### **Answer: 2**



- 1. Which is a low spin complex?
  - A.  $[Fe(CN)_6]^{3}$
  - B.  $\left[ Co(NO_2)_6 \right]^{3-}$
  - $C. \left[ Mn(CN)_6 \right]^{3}$
  - D. All of these



- **2.** Aqueous solution of  $Ni^{2+}$  contains  $\left[Ni\left(H_2O\right)_6\right]^{2+}$  and its magnetic moment is 2.83 B.M. When ammonia is added in it, the predicted change in the magnetic moment of solution is:
  - A. It will remain same

B. It increases from 2.83 B.M.

C. It decreases from 2.83 B.M.

D. It can not be predicted theoretically.

### **Answer: 1**



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3. Which of the following complexes is a paramagnetic complex?

A. 
$$K_2[Ni(CN)_4]$$

$$\mathrm{B.}\,\mathit{Ni}\Big[H_2O\Big)_6\Big]\Big(\mathit{NO}_3\Big)_2$$

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

D. 
$$\left[ Pt \left( NH_3 \right)_4 \right] Cl_2$$

### Answer: 2



Watab Widoo Calation

- 4. Which of the following are paramagnetic?
  - A.  $K_4$   $\left[Ni(CN)_4\right]$
  - $B. K_3 \Big[ Cr(CN)_6 \Big]$
  - $C. K_3 [Co(CN)_6]$
  - D.  $K_2[Ni(CN)_4]$



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**5.** Which of the following pairs of d-electron configuration exhibit both low and high spin tetrahedral complex

A.  $d^1$ ,  $d^2$ 

B.  $d^3$ ,  $d^4$ 

C.  $d^7$ ,  $d^8$ 

D.  $d^9$ ,  $d^{10}$ 

### Answer: 2



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6. Assign the hybridisation, shape and magnetic moment of  $K_2 \left[ Cu(CN)_4 \right]$ 

A.  $sp^3$ , tetrahedral, 1.73 BM

B.  $dsp^2$  square planar, 1.73 BM

C.  $sp^3$ , tetrahedral, 2.8 BM

D.  $dsp^2$ , square planar, 2.8 BM

## Answer: 2

7. Which of the following is most stable?

A. 
$$\left[ Fe \left( H_2 O \right)_6 \right]^{3+}$$

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

C. 
$$\left[ Fe \left( C_2 O_4 \right)_3 \right]^{3-1}$$

D. 
$$[FeCl_6]^{3}$$

### Answer: 3



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**8.** Which of the following complex formed by  $Cu^{2+}$  ions is most stable?

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

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

C. 
$$Cu^{2+} + 2en \Leftrightarrow \left[ Cu(en)_2 \right]^{2+}, \log K = 15.4$$

D. 
$$Cu^{2+} + 4H_2O \Leftrightarrow \left[ Cu(H_2O)_4 \right]^{2+}, \log K = 8.9$$



**9.** The colour of the coordination compounds depends on the crystal field splitting . What will be the correct order of obsorption of wavelength of light in the visible region, for the comolexes,  $\left[Co(NH_3)_6\right]^{3+}$  and  $\left[Co(H_2O)_6^{3+}\right]^{3+}$ ?

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

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

$$C. \left[ Co \left( H_2 O \right)_6 \right]^{3+} > \left[ Co \left( NH_3 \right)_6 \right]^{3+} > \left[ Co (CN)_6 \right]^{3-}$$

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



### 10. Indicate the complex ion which shows geometrical isomerism.

A. 
$$\left[ Cr \left( H_2 O \right)_4 C l_2 \right]^+$$

$$B. \left[ Pt \left( NH_3 \right)_3 Cl \right]^+$$

$$\mathsf{C.}\left[\mathit{Co}\left(\mathit{NH}_{3}\right)_{6}\right]^{3+}$$

D. 
$$\left[ Co(CN)_5(NC) \right]^{3}$$

#### Answer: 1



**11.** When 1 mol  $CrCl_3.6H_2O$  is treated with excess of  $AgNO_3$ , 3 mol of AgCl are obtained. The formula of the coplex is

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

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

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

D. 
$$\left[ Cr \left( H_2 O \right)_6 \right] Cl_3$$

### Answer: 4



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**12.** When 0.1 mol  $CoCl_3(NH_3)_5$  is treated with excess of  $AgNO_3$ , 0.2 mole of AgCl are obtained. The conductivity of solution will correspond to

- A. 1:3 electrolyte
- B. 1:2 electrolyte
- C. 1:1 electrolyte
- D. 3:1 electrolyte



- **13.** The correct IUPAC name of  $\left[Ot(NH_3)_2Cl_2\right]$  is
  - A. diamminedichloridoplatinum (II)
  - B. diamminedichloridoplatinum (IV)
  - C. diamminedichloridoplatinum (O)
  - D. diamminedichloridoplatinum (IV)



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**14.** The stabilization of coordination compound due to chelation is called the chelate effect. Which of the following is the most stable complex species?

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

B. 
$$[Fe(CN)_6]^{3}$$

C. 
$$\left[ Fe \left( C_2 O_4 \right)_3 \right]^{3-1}$$

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

### **Answer: 3**



**15.** The CFSE for octahedral  $\Big[CoCl_6\Big]^{4-}$  is  $18,000cm^{-1}$ . The CFSE for tetrahedral  $\Big[CoCl_4\Big]^{2-}$  will be

A. 18, 
$$000cm^{-1}$$

D. 20, 
$$000cm^{-1}$$

### Answer: 3



**16.** Due to the presence of ambidenate ligands coordination compounds show isomerism. Palladium complexes of the type

 $\left\lfloor Pd \Big(C_6H_5\Big)_2 (SCN)_2 \right\rfloor \& \left\lceil Pd \Big(C_{6H_5} - (2)(NCS)_2 \right\rceil \text{ are }$ 

- B. coordination isomers
- C. ionization isomers
- D. geometrical isomers.



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- 17. The compounds  $\left[Co\left(SO_4\right)\left(NH_3\right)_5\right]Br$  and  $\left[Co\left(SO_4\right)\left(NH_3\right)_5\right]Cl$ 
  - A. Linkage isomerism
  - B. Ionisation isomerism
  - C. coordinatio isomerism
  - D. no isomerism

### Answer: 4



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- 18. Which of the following is not chelating agent
- (a) Thiosulphate
- (b) Oxalato
- (c) Glycinato
- (d) Ethylene diamine.
  - A. Thiosulphate
  - B. Oxalato
  - C. Glycinato
  - D. Ethane-1, 2-diamine

### Answer: 1



**19.** Which of the following species is not expected to be a ligand?

A. NO

 $B.NH_4^+$ 

 $C. NH_2CH_2CH_2NH_2$ 

D. CO

### **Answer: 1**



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**20.** What kind of isomerism exists between  $\left[Cr(H_2O)_6\right]Cl_3$  (violet)

and  $\left[Cr\left(H_2O\right)_5Cl\right]$ .  $H_2O$  (greyish-green)?

A. Linkage isomerism

B. Solvate isomerism

- C. Ionisation isomerism
- D. Coordination isomerism



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- **21.** IUPAC name of  $\left[Pt(NH_3)_2Cl(NO_2)\right]$  is
  - A. platinum diaminechloronitrite
  - B. Chloronitrito-N-ammeniplatinum (II)
  - C. diammenichloridonitrito-N-plantinum(II)
  - D. diamminechloronitrito-N-plantinate(II)

### **Answer: 3**



**22.** On the basis of the following observations made with acqueous solutions. In which of the following complexes the central metal atom exhibits secondary valency is six

Moles of AgC1 precipitated per moles of the compounds with execess of $AgNO_3$
2
2
0
1
0

**A.** *I*, *ii*, *iv* 

B. ii, iii, iv

C. ii, iv, v

D. *I*, *iv*, *v* 



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23. Calculate the overall complex dissociation equilibrium constant

for the  $\left[ {\it Cu} \Big( {\it NH}_3 \Big)_4 \right]^{2+}$  ion, given that  $\beta_4$  for this complex is  $2.1 \times 10^{13}$ 

A. 
$$4.7 \times 10^{-14}$$

B. 
$$2.1 \times 10^{-13}$$

$$C. 2.1 \times 10^{13}$$

D. 
$$4.7 \times 10^{14}$$

### Answer: 1



### LEVEL-I (H.W)

**1.** The complex formed by the combination of calcium ions and ethylene di ammine tetra acetate. (*EDTA*)<sup>-4</sup> Number of moles of calcium ions produced by dissolving of one moles of calcium ions produced by dissolving of one mole of that complex in excess of water is

A. one

B. two

C. four

D. five

### Answer: 2



- 2. Which of the following is a double salt?
  - A. Carnalite
  - B. Potassium ferrocyanide
  - C. Potasium ferricyanide
  - D. Nessler's reagent



- **3.** Bonds present in  $K_4[Fe(CN)_6]$  are
  - A. only ionic
  - B. only covalent
  - C. ionic and covalent

D. ionic, covalent and coordinate covalent

### **Answer: 4**



**4.** Copper sulphate solution forms blue coloured complex with excess of ammonia. Its formula is

A. 
$$\left[ Cu(NH_3)_4 \right]^{+3}$$

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

$$C. \left[ Cu \left( NH_4 \right)_3 \right]^{+2}$$

D. 
$$\left[ Cu \left( NH_4 \right)_3 \right]^{+3}$$

### Answer: 2



A.  $BF_3$ 

 $\mathsf{B.}\mathit{NH}_3$ 

C. *NO* +

D. *CN* -

#### **Answer: 1**



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# 6. Alum in aqueous solution gives positive test for

(A)  $K^+$  (B)  $Al^{3+}$  (C)  $SO_4^{2-}$ 

A. A only

B. B only

- C. A and B
- D. A, B and C



# 7. Chelates are used in

- A. Analytical chemistry
- B. Water softening
- C. Removal of  $Pb^{+2}$  from the blood
- D. All of these

# Answer: 4



8. Hexaaquatitanium (III) chloride is

A. 
$$\left[ Ti \left( H_2 O \right)_6 \right] Cl_3$$

B. 
$$\left[ TiCl_3 \right] 6H_2O$$

$$C. \left[ Ti \left( H_2 O \right)_5 Cl \right] Cl_2$$

$$\mathsf{D}.\left[\mathit{TI}\!\left(H_2O\right)_3\!Cl_3\right]$$

### **Answer: 1**



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**9.** IUPAC name of  $Li[AIH_4]$  is

A. Lithium aluminium hydride

B. Lithium tetrahydrido aluminate (III)

C. Tetrahydride aluminium lithionate

D. Aluminium lithium hydride

**Answer: 2** 



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10. Dipole moment will be zero in the complexes

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

B. 
$$Cis - \left[Pt(NH_3)_2Cl_2\right]$$

C. Trans - 
$$\left[Pt\left(NH_3\right)_2Cl_2\right]$$

D. Both 1 and 3

**Answer: 4** 



11. The number of geometrical isomers of  $\left[Co(NH_3)_3(NO_3)_3\right]$  is

**A.** 0

**B.** 2

**C**. 3

D. 4

# Answer: 2



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**12.** For the given complex  $\left[CoCl_2(en)\left(N_3\right)_2\right]^+$ , the number of geometrical iosmers, the number of optical isomers and total number of isomers of all type possible respectively are

A. 2, 2, &4

B. 2, 2&3

C. 2, 0&2

D. 0, 2&2

# Answer: 2



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# 13. Which does not obey EAN rule?

- A.  $K_4$   $\left[ Fe(CN)_6 \right]$
- B.  $K_3[Fe(CN)_6]$
- $C. \left[ Co(NH_3)_6 \right] Cl_3$
- D.  $\left[Ni(CO)_4\right]$

# Answer: 2



**14.** The following solutions requires three moles of  $AgNO_3$  for the complete precipitation of all the chloride ions present in it

A. One litre of 1M 
$$\left[Co(NH_3)_6\right]Cl_3$$

B. Three litres of 1M 
$$\left[Co(NH_3)_4Cl_2\right]Cl$$

C. One litre of 1.5 M 
$$\left[ Co(NH_3)_5 Cl \right] Cl_2$$

D. all the above

#### Answer: 4



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**15.** AgCl dissolved in excess of  $NH_3$ , KCN and  $Na_2S_2O_3$  solutions the complex produces ions

A. 
$$\left[Ag\left(NH_3\right)_2\right]^+$$
,  $\left[Ag(CN)_2\right]^-$ &  $\left[Ag\left(S_2O_3\right)_2\right]^{3-}$ 

B. 
$$\left[ Ag(NH_3)_2 \right]^{2+}$$
,  $\left[ Ag(CN)_2 \right]^{3-} & \left[ Ag_4(S_2O_3)_5 \right]^{2+}$ 

C. 
$$\left[ Ag(NH_3)_2 \right]^{2+}$$
,  $\left[ Ag(CN)_2 \right]^{+} \& \left[ Ag_2(S_2O_3)_5 \right]^{2-}$ 

D. 
$$\left[ Ag(NH_3)_4 \right]^+$$
,  $\left[ Ag(CN)_4 \right]^{3-} \& \left[ Ag_2(S_2O_3)_2 \right]^{2-}$ 



# **16.** The secondary valency of chromium in $\left[Cr(en)_3\right]Cl_3$ is

- **A.** 6

**B**. 3

- **C**. 2
  - D. 4



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- 17. Stabilisation energy of octahedral complex with  $d^7$  configuration
- (A) -1.8 $\Delta_0$  with one unpaired electron
- (B)  $1.8\Delta_0$  with three unpaired electrons
- (C)  $-0.8\Delta_0$  with one unpaired electron
- (D)  $0.8\Delta_0$  with three unpaired electrons
  - A. A and D
  - B. A and B
  - C. C and D
  - D. B and C

#### **Answer: 1**



**18.** If  $\Delta_0 > P$ , the correct electronic configuration for  $d^4$  system will be (p = pairing energy)

- A.  $t_{2g}^4 e_g^0$
- B.  $t_{2g}^{3}e_{g}^{1}$
- C.  $t_{2g}^{0}e_{g}^{4}$
- D.  $t_{2g}^2 e_g^2$

# **Answer: 1**



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**19.** The ligand that gives dark blue color with curic ion in the laboratory is

A.  $NH_3$ 

 $B.I^-$ 

C. CN

D.  $S_2O_3^{2-}$ 

# Answer: 1



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**20.** The configuration of an elements 'X' is  $3d^{10}4s^1$ . The wrong statement regarding the element 'X' is

A. it forms complexes

B. it exhibits variable velency

C. it forms paramagnetic ions only

D. it can form coloured salts

# Answer: 3



**21.** In the qualitative analysis of group 3 cations blood red colouration is a test for

A. iron using cyanate as ligand

B. chromium using cyanide as ligand

C. iron using thiocyante as ligand

D. chromium using thiocyanate as ligand

**Answer: 3** 



LEVEL-II (H.W)

- 1. An ambidentate ligand is one which
  - A. is linkage to the metal atom at two points
  - B. has two donor atoms but only one of them has the capacity to form a coordinate bond
  - C. has two donor atoms but either of the two can form a coordinate bond
  - D. forms chelate rings



- 2. Which of the following is not chelating agent
- (a) Thiosulphate
- (b) Oxalato

- (c) Glycinato
- (d) Ethylene diamine.
  - A. Thiosulphato
  - B. Oxalato
  - C. Glycinato
  - D. Ethylene diamine



- **3.** IUPAC name of  $Na_3 \left[ CoCl \left( NO_2 \right)_5 \right]$  is
  - A. Sodium chloropentanitrocobaltate (III)
  - B. Sodium cobaltnitrate
  - C. Trisodium chloropentanitro cobalt

D. Pentanitrocobalt (III) trisodium complex

#### **Answer: A**



- **4.** The IUPAC name of Wilkinsons catalyst  $\left[RhCl(PPh_3)_3\right]$  is
  - A. Chlorotris (triphenylphopshine) rhodium (I)
  - B. Chlorotris (triphenylphosphine) rhodium (IV)
  - C. Chlorotris (triphenylphosphine) rhodium (0)
  - D. Chlorotris (triphenylphosphine) rhodium (VI)

## **Answer: 1**



**5.** IUPAC name 
$$\left[Co(NH_3)_5(NO_2)\right]Cl_2$$
 is

- A. Nitrito-N-pentaammine cobalt (III) chloride
- B. Nitrito-N-pentaammine cobalt (II) choride
- C. pentaammine nitrito-N-cobalt (II) chloride
- D. Pentaamine nitroto-N-cobalt (III) chloride



- **6.** Both geometrical and optical isomerism are shown by
- A.  $\left[Co(en)_2Cl_2\right]^+$ 
  - $\mathsf{B.}\left[\mathit{Co}\!\left(\mathit{NH}_{3}\right)_{5}\!\mathit{Cl}\right]^{2+}$
  - $\mathsf{C.}\left[\mathit{Co}\!\left(\mathit{NH}_{3}\right)_{4}\!\mathit{Cl}_{2}\right]^{+}$

D. 
$$[Cr(ox)_3]^{3}$$



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# 7. Which of the following is not optically active?

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

B. 
$$\left[ Cr(ox)_3 \right]^{3}$$

C. 
$$cis - \left[CoCl_2(en)_2\right]^+$$

D. trans - 
$$\left[ CoCl_2(en)_2 \right]^+$$

# Answer: 4



**8.** Cis-trans isomerism is found in square planar complexes of molecular formula: (a and b are monodentate ligands)

- A.  $Ma_4$
- B.  $Ma_3b$
- $C. Ma_2b_2$
- D.  $Mab_3$

#### Answer: 3



- **9.** Which of the following isomeric pairs shows ionization isomerism?
  - A.  $\left[ Co(NH_3)_6 \right] \left[ Cr(CN)_6 \right]$  and  $\left[ Cr(NH_3)_6 \right] \left[ Co(CN)_6 \right]$

B. 
$$\left[Cr\left(H_2O\right)_6Cl_3\text{ and }\left[Cr\left(H_2O\right)_5Cl\right]Cl_2.H_2O\right]$$

C. 
$$\left[Pt\left(NH_3\right)_2Cl_2\right]$$
 and  $\left[Pt\left(NH_3\right)_4\right]\left[PtCl_4\right]$ 

D. 
$$\left[ Co(NH_3)_5 Br \right] SO_4$$
 and  $\left[ Co(NH_3)_5 SO_4 \right] Br$ 



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# **10.** The color of $\left[Ti(H_2O_6)^{3+}\right]^{3+}$ is due to

A. Charge transfer transition

 $B. d \rightarrow d$  transition

C. Ligand to metal charge transfer tranition

D. Metal to ligand charge transfer transition

# Answer: 2

**11.** Coordination number of Cr is six. A complex with ligands  $C_2O_4^{2-}$ , en and superoxide will be in the ratio to make complex

$$\left[ Cr \left( C_2 O_4 \right)_x (en)_y \left( O_2 \right)_z \right]^{\Theta}.$$

- A. 1:1:2
- B. 1:1:1
- C. 1:2:2
- D. 2:1:1

# Answer: 1



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**12.** The geometries of  $Ni(CO)_4$  and  $Ni(PPh_3)_2Cl_2$  are .

A. both square planar B. tetrahedral and square planar C. both tetrahedral D. square planar and tetrahedral Answer: 2 **Watch Video Solution** 13. What are the magnetic moment (in BM) for Ni(II) ion in square planar and octahedral geometry, respectrively? A. 0 and 2.83 B. 2.83 and 2.83 C. 0 and 1.73 D. 2.83 and 0



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**14.** A solution containing 2.675 g of  $CoCl_3.6NH_3$  (molar mass =  $267.5gmol^{-1}$ ) is passed through a cation exchanger. The chloride ions obtained is solution were treated with excess of  $AgNO_3$  to give 4.73 g of AgCl (molar mass =  $143.5gmol^{-1}$ ). The formula of the complex is (At. mass of Ag = 108 u)

A. 
$$\left[CoCl_3(NH_3)_3\right]$$

B. 
$$\left[ CoCl(NH_3)_5 \right] Cl_2$$

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

D. 
$$\left[ CoCl_2 \left( NH_3 \right)_4 \right] Cl$$

#### **Answer: 3**



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**15.** The complex  $\left[Pd(en)_2\right]^{2+}$  has....Structure

- A. Square planner
- B. tetrahedral
- C. pyramidal
- D. pentagonal

#### **Answer: 1**



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**16.** Crystal field stabilization energy for high spin  $d^4$  octahedral complex is

A. -  $0.6\Delta_0$ 

D. -  $1.2\Delta_0$ 

B. -  $1.8\Delta_0$ 

 $C. - 1.6\Delta_0$ 

# **Answer: 1**



17. Which of the following is antidote for lead poisoinig?

B. Cis-platin

A. CoCl<sub>3</sub>

D. DMG

Answer: 3

C. EDTA

18. EDTA is used for the estimation of

- A.  $Na^+$  and  $K^+$  ions
- B.  $Cl^-$  and  $Br^-$  ions
- C.  $Cu^{2+}$  and  $Ag^{+}$  ions
- D.  $Ca^{2+}$  and  $Mg^{2+}$  ions

#### Answer: 4



- **19.** Wilkinson's catalyst is used as a homogeneous hydrogenation catalyst for the conversion of alkenes to alkanes. It is a complex of
  - A. iron
  - B. aluminium

- C. rhodium
- D. cobalt



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# 20. Ziegler-Natta catalyst is

- A. Solution of  $SnCl_4$  + trialkylaluminium
- B. Solution of  $TiCl_4$  + trakylaluminium
- C. Solution of  $TiCl_4$  + trialkylchromium
- D. Solution of  $SnCl_4$  + Tollen's reagent

# Answer: 2



**1.** IUPAC name of complex  $K_3 \left[ Al \left( C_2 O_4 \right)_3 \right]$  is

A. Potassium alumino-oxalate

B. Potassium trioxalatoaluminate(III)

C. Potassium aluminium (III) oxalate

D. Potassium trioxalatoaluminate (IV)

### **Answer: B**



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2. Trioxalato aluminate (III) and tetrafluoro-borate (III) ions are:

A. 
$$\left[Al\left(C_2O_4\right)_3\right], \left[BF_4\right]^{3-1}$$

$$B. \left[ Al \left( C_2 O_4 \right)_3 \right]^{3+}, \left[ BF_4 \right]^{3+}$$

$$C. \left[ Al \left( C_2 O_4 \right)_3 \right]^{3-}, \left[ BF_4 \right]^{-}$$

D. 
$$\left[Al\left(C_2O_4\right)_3\right]^{2-}$$
,  $\left[BF_4\right]^{2-}$ 

# Answer: C



# 3. Which of the ligands can show linkage isomerism:

- A. NCS
  - B.  $NO_2^-$
  - C. *CN* -
  - D. All of these

# Answer: D



D Watal William Calaitian

**4.** In which of the following complexes the nickel metal is in highest oxidation state :

A. 
$$Ni(CO)_4$$

B. 
$$K_2[NiF_6]$$

C. 
$$\left[Ni\left(NH_3\right)_6\right]\left(BF_4\right)_2$$

D. 
$$\left[K_2 \left[Ni(CN)_4\right]\right]$$

## Answer: B



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**5.** A complex of platinum, ammonia and chloride produces four ions per molecule in the solution. The structure consistent with the observation is:

A. 
$$\left[Pt(NH_3)_4\right]Cl_4$$

$$\mathsf{B.}\left[Pt\left(NH_3\right)_2Cl_2\right]$$

C. 
$$\left[Pt\left(NH_3\right)_5Cl\right]Cl_3$$
D.  $\left[Pt\left(NH_3\right)_4Cl_2\right]Cl_2$ 

# Answer: C



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 $\left[Mo_2O_4(C_2H_4)_2(H_2O)_2\right]^{2-}$  is:



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**7.** In which of the following compounds transition metal is in oxidation state zero

A. 
$$\left[ Co(NH_3)_6 \right] Cl_2$$

$$\operatorname{B.}\left[\operatorname{Fe}\!\left(H_2O\right)_{\!6}\!\operatorname{SO}_4\right]$$

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

D. 
$$\left[ Fe \left( H_2 O \right)_3 \right] (OH)_2$$

# **Answer: C**



- 8. Formula of ferrocene is:
  - A.  $[Fe(CN)_6]^{4-}$
  - B.  $[Fe(CN)_6]^{3+}$
  - C.  $\left[ Fe(CO)_5 \right]$
  - D.  $\left[Fe\left(C_5H_5\right)_2\right]$

# **Answer: D**



- **9.** Which of the following is a  $\pi$  complex?
  - A. Trimethyle aluminium
  - B. Ferrocene
  - C. Diethyl zinc

D. Nickel tetra carbonyl

# **Answer: B**



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**10.** Which of the following is most likely structrure of  $CrCI_3.6H_2O$  if 1/3 of total chlorine of the compound is precipitated by adding AgNO<sub>3</sub> to its aqueous solution?

A. 
$$CrCl_3.6H_2O$$

B. 
$$\left[ CrCLl_3 \left( H_2O \right)_3 \right] \cdot \left( H_2O \right)_3$$

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

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

# Answer: C



**11.** The two compounds 
$$\left[Co(SO_4)(NH_3)_5\right]Br$$
 and

$$\left[ Co(SO_4)(NH_3)_5 \right] Cl$$
 represent:

- A. Linkage isomerism
- B. Ionisation isomerism
- C. Co-ordination isomerism
- D. no isomerism

#### **Answer: D**



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12. Diethylene triamine is:

A. Chelating agent

- B. Polydentate ligand
- C. Tridentate ligand
- D. All of these

# Answer: D



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- 13. How many moels of AgCI would be obtained, when 100 mL of 0.1
- M  $CO(NH_3)_5CI_3$  is treated with excess of  $AgNO_3$ ?
  - A. 0.01
  - B.0.02
  - C. 0.03
  - D. none of these

# **Answer: B**

**14.** 0.001 mol of  $Co(NH_3)_5(NO_3)(SO_4)$  was passed through a cation exchanger and the acid coming out of it required 20 ml of 0.1

A. 
$$\left[CoSO_4\left(NH_3\right)_5\right]NO_3$$

M NaOH for neutralisation. Hence, the complex is

B. 
$$\left[CoNO_3(NH_3)_5\right]SO_4$$

$$C. \left[ Co(NH_3)_5 \right] SO_4 NO_3$$

D. none of these

#### **Answer: B**



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15. Which of th efollowing is a bivalent and bidentate ligand?

A. Nitrito B. Oxalato C. Glycinato D. Ethylene diamine **Answer: B Watch Video Solution** 16. Which of the following has six donor (coordinating) sites? A. Triethylene tetramine B. Ethylenediamine tetracetate ion (EDTA) C. Nitrilotriacetic (NTA) D. Diethylene triamine **Answer: B** 

**17.** The IUPAC name of the red coloured complex  $Fe(C_4H_7O_2N_2)_2$ obtained from the reaction of  $Fe^{2+}$  and dimethyl glyoxime

- A. bis (dimethyl glyoxime) ferrate (II)
- B. bis (dimethyl glyoximato) iron (II)
- C. bis (2, 3-butanediol dioximato) iron (II)
- D. bis (2, 3-butanedione dioximato) iron (II)

**Answer: B** 



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**18.** On treatment of 10 ml of 1M solution of the complex  $CrCl_3.6H_2O$ with excess of AgNO<sub>3</sub>, 4.305 g of AgCl was obtained. The complex is

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

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

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

D. 
$$\left[ Cr \left( H_2 O \right)_6 \right] Cl_3$$

#### **Answer: D**



- 19. Which of the following species is not expected to be a ligand
- (a)NO
- (b)  $NH_4^{\oplus}$
- (c)  $NH_2 NH_3^{\oplus}$
- (d) (CO).
  - **A.** *NO* <sup>+</sup>

- $B.NH_4^+$
- $C. NH_2 NH_3^+$
- D. CO

#### Answer: B



- **20.** The number of donor sites in dimethyl glyoxime, glycinato, diethylene triamine and *EDTA* are respectively
- (a) 2, 2, 3 and 4
- (b) 2, 2, 3 and 6
- (c) 2, 2, 2 and 6
- (d)2, 3, 3 and 6.
  - A. 2,2,3 and 4
  - B. 2,2,3 and 6

- C. 2,2,2 and 6
- D. 2,3,3 and 6

#### **Answer: B**



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- **21.** The IUPAC name of  $Ba[BrF_4]_2$  is
  - A. Barium tetrafluorobromate (V)
  - B. Barium tetraflurorbromate(III)
  - C. Barium bis (tetrafluorobromate)(III)
  - D. none of these

#### **Answer: B**



22. The formula of the complex hyderidotrimethoxoborate (III) ion is:

A. 
$$\left[BH\left(OCH_3\right)_3\right]^{2}$$

B. 
$$\left[BH_2(OCH_3)_3\right]^{2-}$$

C. 
$$\left[BH\left(OCH_3\right)_3\right]^{-1}$$

D. 
$$\left[BH\left(OCH_3\right)_3\right]^+$$

#### **Answer: C**



**23.** The complex ion which has no 'd'-electrons in the central metal atom is

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

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

D. 
$$\left[MnO_4\right]^{-}$$
Answer: D

 $C. \left[ Cr \left( H_2 O \right)_6 \right]^{3+}$ 

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 $Na_4[Fe(CN)_5(NOS)]$  is :

number of Fe in violet coloured

complex

- - **A.** 0

24. Oxidation

- **B**. 2
- **C**. 3
- D. 4

**Answer: B** 

**25.** Complexes  $\left[Co\left(SO_4\right)\left(NH_3\right)_5\right]Br$  and  $\left[CoBr\left(NH_3\right)_5\right]SO_4$  can be distinguished by

A. conductance measurement

B. using  $BaCl_2$ 

C. using  $AgNO_3$ 

D. all

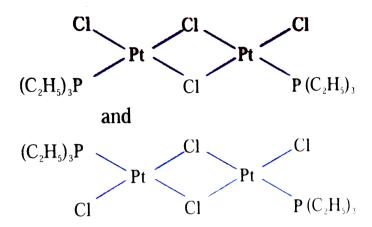
#### **Answer: D**



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**EXERCISE -2** 

1. The Complexes given below show:



- A. Optical isomerism
- B. coordination isomers
- C. Geometrical isomerism
- D. Bridged isomerism

#### **Answer: C**



**2.** For the complex ion dichloro bis (ethylene diamine) cobalt (III), select the correct statement.

A. It has three isomers, two of them are optically active and one is optically inactive

B. It has three isomers, all of them are optical active

C. It has three isomers, all of them are optically inactive

D. It has one optically active isomer and two geometrical isomers

#### **Answer: A**



**3.**  $\left[ Pd \left( NH_3 \right)_2 (SCN)_2 \right]$  and  $\left[ Pd \left( NH_3 \right)_2 (NCS)_2 \right]$  are:

A. Linkage isomers

- B. coordination isomers
- C. Ionisation isomers
- D. geometrical isomers.

#### Answer: A



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- 4. Which one of the following will be able to show geometrical isomerism if complexes are square planar? .
  - A.  $Ma_3b$
  - B.  $M(\forall)_2$
  - C. Mabcd
  - D.  $Ma_{\Lambda}$

#### Answer: C

5. The total number possible isomers for the complex compound

$$\left[ \mathit{Cu^{II}} \left( \mathit{NH}_{3} \right)_{4} \left[ \mathit{Pt^{II}} \mathit{CI}_{4} \right] \right]$$
 are

- **A.** 3
- **B**. 5
- C. 4
- D. 6

#### Answer: C



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**6.** Cis-trans isomerism is found in square planar complexes of molecular formula: (a and b are monodentate ligands)

- A.  $MA_A$
- B.  $MA_3B$
- $C. MA_2B_2$
- D.  $MAB_3$

#### **Answer: C**



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#### 7. Which complex is likely to show optical activity:

- A. Trans- $\left[CoCl_2(NH_3)_4\right]^+$
- $B. \left[ Cr \left( H_2 O \right)_6 \right]^{3+}$
- C. Cis  $\left[Co(NH_3)_2(en)_2\right]^{3+}$
- D. Trans- $\left[Co\left(NH_3\right)_2(en)_2\right]^{3+}$

## Answer: C



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- **8.** The number of bridged carbonyl groups in  $Mn_2(CO)_{10}$  is :
  - **A.** 2
  - **B**. 3
  - **C**. 0
  - D. 1

#### **Answer: C**



**9.** The oxidation number of Co in the complex ion

ion 
$$[(en)_2 Co \stackrel{\text{NH}}{\sim} Co(en)_2]^{3+}$$

- A. + 2
- B. + 3
- C. + 4
- D.+6

#### **Answer: B**



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10. Which of the following statements is correct?

A. Geometrical isomerism is not observed in complexes of C.N. 4

having tetrahedral geometry

- B. Square planar complexes generally do not show geometrical
  - isomerism
- C. The square planar comlex of general formulae  ${\it Ma}_3{\it b}$  or  ${\it Mab}_3$  exhibits cis-trans isomerism
- D. The plantinum glycinato complex,  $\left[Pt(Gly)_2\right]$  does not show geometrical isomerism

#### **Answer: A**



- 11. Geometrical isomerism can be shown by
  - A.  $\left[Ag(CN)\left(NH_3\right)\right]$
  - $\mathrm{B.}\,\mathit{Na}_{2}\Big[\mathit{Cd}\Big(\mathit{N}_{2}\Big)_{4}\Big]$
  - C.  $\left[PtCl_4I_2\right]$

D. 
$$\left[ PtCl \left( NH_3 \right)_3 \right] \left[ Au(CN)_4 \right]$$

#### **Answer: C**



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- **12.**  $\left[Co(en)_3\right]^{3+}$  ion is expected to show
  - A. two optically active isomers: d, I forms
  - B. three optically active isomers: d, l and meso forms
  - C. four optically active isomers: cis, d and l isomers and trans d
    - and I isomers
  - D. none of these

#### **Answer: A**



13. The number of geometrical isomers for octahedral

$$\left[ CoCl_4 \left( NH_3 \right)_2 \right]^{-}$$
, square planar  $\left[ AuBr_2Cl_2 \right]^{-}$  and  $\left[ PtCl_2(en) \right]$  are

- **A.** 2, 2, 2
- B. 2, 2, no isomerism
- C. 3, 2, 2
- D. 2, 3, no isomerism

#### Answer: B



14. Which of the following statements is incorrect?

A. Co-ordination compounds and complexes are synonymous

terms

- B. Complexes must give free metal ions in the solution
- C. Complexes may give ions in the solution or may not give ions the solution
- D. Generally complex ion does not dissociate into its component parts even in the solution

#### **Answer: B**



**15.** Which one of the following is an example of coordination isomerism?

A. 
$$\left[Co\left(NH_3\right)_5 Br\right] SO_4$$
 and  $\left[Co\left(NH_3\right)_5 SO_4\right] Br$ 

- B.  $\left[ Co(NH_3)_5 NO_2 \right] Cl_2$  and  $\left[ Co(NH_3)_5 ONO \right] Cl_2$
- C.  $\left[Cr(H_2O)_6\right]Cl_3$  and  $\left[Cr(H_2O)_5Cl\right]Cl_2$ .  $H_2O$

D. 
$$\left[ Cr(NH_3)_6 \right] \left[ Co(CN)_6 \right]$$
 and  $\left[ Co(NH_3)_6 \right] \left[ Cr(CN)_6 \right]$ 

**Answer: D** 



**16.** The two compounds pentaamminesulphatocobalt (*III*)bromide and pentaamminesulphatocobalt (*III*) chloride represent :

- A. Linkage isomerism
- B. Ionisation isomerism
- C. coordinatio isomerism
- D. no isomerism

**Answer: D** 



**17.** Select the correct code about complex

$$\left[ Cr(NO_2)(NH_3)_5 \right] \left[ ZnCl_4 \right]$$
:

- (I) IUPAC name of compoun is pentaamminenitrito-N- chromium
- (III) tetrachlorozincate (II)
- (II) It shows geometrical isomerism
- (III) It shows linkage isomerism
- (IV) It shows coordination isomerism`

A. III, IV

**B. I, III & IV** 

C. II, III & IV

D. I, II, III & IV

#### **Answer: B**



**18.** Isoomerisms exhibited by  $\left[Cr(NH_3)_2(H_2O)_2Cl_2\right]^+$  are

A. ionisation, optival

B. hydrate, optical

C. geometrical, optical

D. coordinate, geometrical

#### **Answer: C**



**19.** Of the following complex ions which one can form a chelate with ethylenediamine?

A. 
$$cis - \left[Co(en)_2Cl_2\right]^+$$

B. trans-
$$\left[ Co \left( NH_3 \right)_4 Cl_2 \right]^+$$

C. trans-
$$\left[Co\left(NH_3\right)_2Cl_4\right]^{-1}$$

D. trans- $\left[Co(en)_2Cl_2\right]^+$ 

#### Answer: A



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# **20.** $[Pt(NH_3)(NO_2)Ph(NH_2OH)]^+$ , the no. of geometrical isomers

including linkage isomerism is

B. 6

**C**. 4

D. 5

**Answer: B** 



21. Which of the following compound shows optical isomerism?

(en=ethylenediamine)?

(1) 
$$cis - \left[Co(NH_3)_4Cl_2\right]$$

(2) trans - 
$$\left[Co(en)_2Cl_2\right]$$

(3) 
$$cis - \left[Co(en)_2 Cl_2\right]$$

(4) 
$$\left[ Co(en)_3 \right]$$

Select the correct answer using the codes given below:

Codes:

A. I and II

B. II and III

C. III and IV

D. I, II and IV

#### **Answer: C**

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**22.** Oxidation number of Cr in the following complex is

$$\left( (H_2O)_4 Cr < O_2 > Cr (H_2O)_4 \right)$$

- **A.** 3
- B. 6
- **C**. 4
- D. 5

**Answer: A** 



**23.**  $\left[Cr\left(NH_3\right)_5 Br\right] Cl$  and  $\left[Cr\left(NH_3\right)_5 Cl\right] Br$  can be distinguished by/and isomerism shown is:

- A. BaCl<sub>2</sub>, ionisation
- ${\rm B.}\,AgNO_3, {\rm ionisation}$
- C. AgNO<sub>3</sub>, coordinate
- D. BaCl<sub>2</sub>,linkage

#### **Answer: B**



**24.** If excess of  $AgNO_3$  solution is added to 100 mL of a 0.024 M solution of dichlorobis (ethylene diamine) cobalt (III) chloride, how many mol of AgCl be precipitated:

A. 0.0012

B. 0.0016

C. 0.0024

D. 0.0048

#### **Answer: C**



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#### 25. Match the column:

Column I (Co-ordination compound)

- (A)  $Na_2[Pt(SCN)_2(Ox)_2]$
- (B)  $[\operatorname{CrCl}_2(\operatorname{NH}_3)_4]\operatorname{NO}_3$
- (C)  $[Pt(NO_2)(Gly)(NH_3)]$
- (D)  $K_3[Fe(OH)_2(C_2O_4)_2]$

Column II (Type of isomerism shown)

- (P) Ionization isomerism
- (Q) Linkage isomerism
- (R) Geometrical isomerism
- (S) Optical isomerism



**26.** Which of the following is correct IUPAC name of any compound.

A. Tris (acetyl acetonato) iron (III) chloride

B. Hexachloroplantinum (IV) tetraammine dicyano platinate(IV)

C. Ammine bromochloro methylamine platinum (II)

D. Cis dichloro (ethylenediamine) platinum (II)

#### **Answer: C**



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**27.**  $\left[Fe^{II}Cl(CN)_4(O_2)\right]^{4-}$  is named as :

A. chlorotetracyanodioxoferrate (II) ion

B. chlorotetracyanoperoxoferrate(II) ion

 $C.\ chlorotetracyanosuperoxoferrate (II)\ ion$ 

D. none is correct

#### **Answer: C**



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**28.** Which of the following complexes has the least molar conductivity in the solution?

- A.  $CoCl_3.3NH_3$
- B.  $CoCl_3.4NH_3$
- C.  $CoCl_3.5NH_3$
- D.  $CoCl_3.6NH_3$

#### **Answer: A**



**29.** Hybridization and magnetic moment of  $\left[Cu(NH_3)_4\right]^{2+}$  and

 $Mn(CN)_6^{3-}$  ions respectively are

A.  $dsp^2$ , 1.73BM:  $d^2sp^3$ , 2.83BM

B.  $sp^{3}$ , zero,  $sp^{3}d^{2}$ , 4.9BM

C.  $sp^3$ , 1.73BM,  $sp^3d^2$ , 4.9BM

D.  $dsp^2$ , 1.73BM,  $d^2sp^3$ , zero

#### **Answer: A**



30. Lead poisoinig in the body can be removed by:

A. EDTA in the form of calcium dihydrogen salt

B. Cis-platin

C. Zeisse's salt

D. DMG

Answer: A



31.

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 $\left[Pd^{2+}\left(NH_2 - CH\left(CH_3\right) - CO_2^{-}\right)_2\right]^0$ 

How many geometrical isomers are possible

for

**A.** 2

C. 4

**B**. 3

D. 6

**Answer: C** 



**32.** Which kind of isomerism is exhibited by [Co(EDTA)]-

A. Optical & geometrical isomerism

B. Geometrical isomerism

C. Optical isomerism

D. no isomerism

#### **Answer: C**



33. Find the correct pair for Column-I and Column-II:

Column-I

(A)  $[Ma_2bcde]^{n\pm}$  (B)  $[Ma_2b_2c_2]^{n\pm}$  (C)  $[Ma_3bcd]^{n\pm}$  (D)  $[M(AB)c_2d_2]^{n\pm}$ 

Column II

(P) 3 optically inactive isomers

(Q) 4 geometrical isomers

(R) 6 stereo(space) isomers

(S) 2 optically active isomers

(where AB  $\rightarrow$  Unsym. bidentate ligand, a,b,c,d & e

→ monodentate ligands)



**34.** Which of the following complexes exhibit optical isomerism?

A. trans-tetramminebis (thiocyanato)chromium(III)ion

B. cis-diamminedicarbonatocobaltate(III) ion

C. trans-diamminedicarbonatocobaltate(III) ion

D. cis-bis(glycinato)platinum(II)



**35.** Coordination compounds plays many important roles in animals and plants. The are essential in the storage and transport of oxygen as electrons transfer agents as catalysts and in photosynthesis Wide range of application in daily life takes place through formation of complexes Photographic fixing qualitative and quantitative analysis purification of water metallurgical extraction are some specific worth mentioning

The complex  $\left[Fe\left(H_2O\right)_5NO\right]^{2+}$  is formed in the brown ring test for nitrates when freshly prepared  $FeSO_4$  soultion is added to aqueous solution of  $NO_3^{\Theta}$  followed by addition of conc.  $H_2SO_4$  Select correct statement about this complex

- (a) Colour change is due to charge transfer
- (b) It has iron in +1 oxidation state and nitrosyl as  $NO^{\oplus}$

(c ) It has magnetic moment of 3.87BM confirming three unpaired

electrons in Fe

(a) All the above are correct statements .

A. Colour change is due to charge transfer

B. It has iron in +1 oxidation state and nitrosyl as  $NO^{\,+}$ 

C. It has magnetic moment of 3.87 B.M. confirming three

unpaired electrons in Fe

D. All are correct statements

### Answer: D





**1.** Which of the following statements is not true about the complex ion  $\Big[ {\it CrCl}_2(en)_2 \Big]^+$ 

A. It has two geometrical isomers - cis and strans

B. Both the cis and transi isomers display optical activity

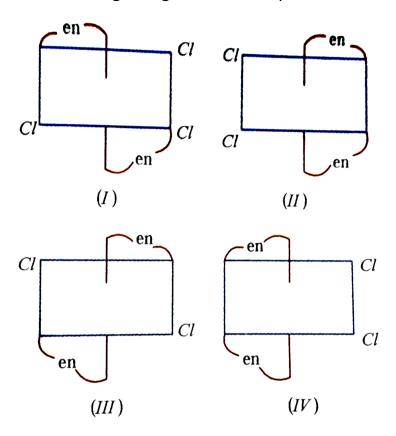
C. only the cis isomers displays optical activity

D. only the cis isomers has non-superimpossible mirror image

#### **Answer: B**



# 2. Of the following configurations, the optical isomers are



A. I and II

B. I and III

C. II and IV

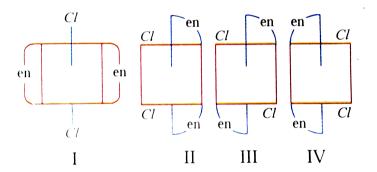
D. II and III

# **Answer: C**



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3. Identify the geometrical isomers of the following:

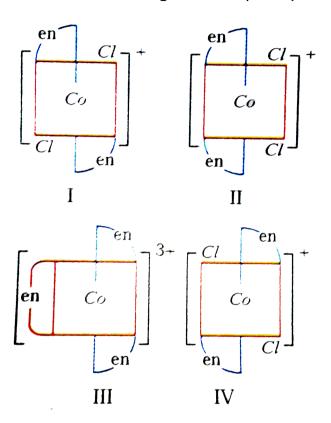


- A. I with III
- B. II with IV
- C. I with II and IV
- D. none of these

# **Answer: C**



**4.** Which of the following ions are optically active?



A. I only

B. II only

C. II and III

D. IV only

# **Answer: C**



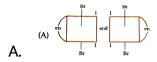
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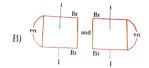
- **5.** How many isomers are possible for the complex ion  $\left[ Cr \left( NH_3 \right) (OH)_2 CI_3 \right]^{2-} ?$ 
  - **A.** 2
  - **B**. 3
  - **C**. 4
  - D. 5

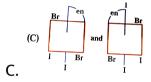
# **Answer: B**



**6.** The complex ion .  $\left[M(en)Br_2I_2\right]^{-1}$ , has two optical isomers. Their correct configurations are:







$$(D) \stackrel{\text{gr}}{\overset{\text{Br}}{=}} \frac{Br}{and} \stackrel{\text{Br}}{\underset{\text{Br}}{=}} \frac{Br}{en}$$

# **Answer: D**



**7.** Match the complexes in column I with their properties listed in column II. Indicate your answer by darkening the appropriate bubbles of the  $4 \times 4$  matrix given in the ORS.

Column I

(A)  $[Co(NH_3)_4(H_2O)_2]Cl_2$ 

(B)  $[Pt(NH_3)_2\dot{C}l_2]$ 

(C) [Co(H<sub>2</sub>O)<sub>5</sub>Cl]Cl (D) [Ni(H<sub>2</sub>O)<sub>6</sub>]Cl<sub>2</sub>

Column II

(P) Geometrical isomers

(Q) Paramagnetic

(R) Diamagnetic

(S) Metal ion with +2 oxidation state



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8. Consider the following complexes:

(I)  $K_2Ptcl_6$  (II)  $PtCl_4.2NH_3$ 

Their electrical conductances in an aqueous solution are:

A. 256, 0, 97, 404

(III)  $PtCl_{A}$ .3 $NH_{3}$  (IV)  $PtCl_{A}$ .5 $NH_{3}$ 

C. 256, 97, 0, 404

D. 404, 97, 256, 0

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Answer: A

9. The total possible co-ordination isomers for the following compounds respectively are

(i) 
$$\left[Co(en)_3\right] \left[Cr\left(C_2O_4\right)_3\right]$$

(ii) 
$$\left[ Cu(NH_3)_4 \right] \left[ CuCl_4 \right]$$

(iii)  $\left[Ni(en)_3\right] \left[Co(NO_2)_6\right]$ 

- A. 4, 4, 4
- **B**. 2, 2, 2
- C. 2, 2, 4
- D. 4, 2, 4

# **Answer: D**



- Geometrical isomerism in coordination compounds is exhibited
  - A. Square planar and tetrahedral complexes
  - B. Square planar and octahedral complexes
  - C. Tetrahedral and octahedral complexes
  - D. Square planar, tetrahedral and octahedral complexes

# **Answer: B**



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11. Which of the following is not optically active?

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

B. 
$$[Cr(ox)_3]^{3-}$$

C. 
$$cis - \left\lceil CoCl_2(en)_2 \right\rceil^+$$

D. trans-
$$\left[CoCl_2(en)_2\right]^+$$

# Answer: D



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**12.** Coordination compounds plays many important roles in animals and plants. The are essential in the storage and transport of oxygen

as electrons transfer agents as catalysts and in photosynthesis Wide range of application in daily life takes place through formation of complexes Photographic fixing qualitative and quantitative analysis purification of water metallurgical extraction are some specific worth mentioning

Arrange of the following in order of decreasing number of unpaired electrons

(I) 
$$\left[Fe\left(H_2O\right)\right)_6]^{2+}$$

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

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

(IV) 
$$\left[ fe \left( H_2 O \right)_6 \right]^{3+}$$

- (a) IV,I,II,III
- (b) *I*, *II*, *III*, *IV*
- (c) III, II, IIV
- (d) II,III,I,IV`.



**13.** Which of the following statements is/are false

A. In  $\left[PtCl_2(NH_3)_4\right]^{2+}$  complex ion, the cis-form is optically active, while trans-form is optically inactive

- B. In  $\left[Fe\left(C_2O_4\right)_3\right]^{3-}$ , geometrical isomerism does not exist, while optical isomerism exists
- C.  $[Mabcd]^{n^{\frac{1}{2}}}$  square planar complexes exhibit both optical as well as geometrical isomerism
- D. In  $[Mabcd]^{n-1 \over 2}$  tetrahedral comlexes, optical isomerism cannot be observed

Answer: A,C,D



14. The number of unpaired electrons expected for the complex ion

$$\left[Cr(NH_3)_6\right]^{2+} is:$$

- **A.** 2
- **B.** 3
- **C**. 4
- **D**. 5

# **Answer: A**



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**15.** Select the correct code regarding total number of space isomers for the following compounds:

(I)  $\left[Ma_3b_2c\right]^{n\pm}$ 

(II)  $\left[M(AB)_3\right]^{n\pm}$ 

(III)  $\left[Ma_2b_2c_2\right]^{n\pm}$ 



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# **EXERCISE -4**

1. 
$$\ln \left[ Cr \left( C_2 O_4 \right)_3 \right]^{3-}$$
, the isorerism shown is

- A. Ligand
- B. Optical
- C. Geometrical
- D. Ionization

# Answer: B



- 2. A similarity between optical and geometrical isomerism is that
  - A. Each gives equal number of iosmers for a given compound
  - B. If in a compound one is present then so is the other
  - C. Both are included in stereoisomerism
  - D. They have similarity

# **Answer: C**



- **3.** A square planar complex is formed by hybridization of which atomic orbitals?
  - A. s, $P_x$ ,  $P_y$ ,  $d_{yz}$
  - B. s, $p_x$ ,  $p_y$ ,  $d_x^2 y^2$

C. s, $p_x$ , p - (y),  $d_z^2$ 

D. s, $p_x$ ,  $p_y$ ,  $d_{xy}$ 

# **Answer: B**



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- **4.** The type of isomerism present in intro pentaamine-chromium(III) chloride is:
  - A. Optical
  - B. Linkage
  - C. Inization
  - D. Polymerization

# Answer: B, C



**5.** In the complex  $\left[Fe\left(H_2O\right)_6\right]^{3+}\left[Fe(CN)_6\right]^{3-}\left[Fe\left(C_2O_4\right)_3\right]^{3-}$  and  $\left[FeCl_6\right]^{3-}$ , that complex that has highest stability is

A. 
$$\left[ Fe \left( H_2 O \right)_6 \right]^3$$

B. 
$$[Fe(CN)_6]^{3-}$$

C. 
$$\left[ Fe \left( C_2 O_4 \right)_4 \right]^{3-1}$$

D. 
$$[FeCl_6]^{3}$$

# Answer: C



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**6.** One mole of complex compound  $Co(NH_3)_5Cl_3$  gives 3 moles of ions on dissolution in water. One mole of same complex reacts with two moles of  $AgNO_3$  to yield two moles of AgCl(s). The complex is:

A. 
$$\left[ Co(NH_3)_4 Cl_2 \right] Cl. NH_3$$

B. 
$$\left[ Co(NH_3)_4 Cl \right] Cl. NH_3$$

$$C. \left[ Co(NH_3)_4 Cl \right] Cl_2$$

$$D. \left[ Co \left( NH_3 \right)_3 Cl_3 \right] 2NH_3$$

# **Answer: C**



**7.** Ammonia forms the complex  $\left[Cu\left(NH_3\right)_4\right]^{2+}$  with copper ions in alkaline solution but not in acid solution. The reasons for it is:

A. In alkaline solution  $Cu(OH)_2$  is precipitated which is soluble in

excess is alkali

B. Copper hydroxide is amphoteric

C. In acidic solution hydration protects  $Cu^{2+}$  ions

D. In acidic solution protons coordinates with ammonia molecule formaing  $NH_4^+$  ions and  $NH_4$  molecules are not available

#### **Answer: D**



- 8. Which one of the following statement is correct?
  - A. Ferric ions give abule colour with ammonium thiocyanate in acidic medium
  - B. On boiling a solution having  $\left[Co\left(H_2O\right)_6\right]^{2+}$  and air is bubbled to get  $CoCO_3$  quantitatively
  - C. A dilute solution of mageanese salt give a pint colour
  - D. Form a mixed precipitate of AgCl and Agl ammonia solution dissolves only AgCl

# **Answer: D**



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- 9. Which has regular tetrahedral geometary?
  - A.  $[Ni(CH)_4]^{2+}$
  - B.  $[Pt(NH (3))_4]^{2+}$
  - C.  $\left[AuCl_4^{-1}\right]$
  - D.  $\left[ Cu \left( NH_3 \right)_4 \right]^{2+}$

# **Answer: C**



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**10.** The coordination number of a central metal atom in a complex is determined by:

- A. The numbe of only anionic ligand bonded to metal ion
- B. The number of ligands aroumnd a metal ion bounded by pi
- C. The number of ligands around a metal a ion bonded by sigma and pi bonds
- D. The number o ligands around a metal ion bonded by sigma bonds

#### **Answer: D**



- 11. Which of the following complex is an outer orbital complex?
  - A.  $\left[Ni\left(NH_3\right)\right]^{2+}$
  - B. [Math Processing Error]

$$\mathsf{C.}\left[\mathit{Co}\left(\mathit{NH}_3\right)_6\right]^{3+}$$

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

# Answer: A



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- 12. Co-ordination compounds have great importance in biological systems. In this context, which statements is incorrect?
  - A. Carboxypeptidase A is an enzyme and contains zinc
  - B. Haemoglobin is the red pigment of blood and contains iron.
  - C. Cyanocobalamin is  $B_{12}$  and contains cobalt.
  - D. Cis platin is used as analgesic.

# Answer: D



13. Which one of the following has largest number of isomers?

A. 
$$\left[Co(en)_2Cl_2\right]^+$$

B. 
$$\left[ Co(NH_3)_5 Cl \right]^{2+}$$

C. 
$$\left[ Ir \left( PR_3 P \right)_2 H(CO) \right]^{2+}$$

D. 
$$\left[ Ru \left( NH_3 \right)_4 Cl_2 \right]^+$$

# Answer: A



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**14.** The correct order of magnetic moments (spin values in B.M.) among is:

A. 
$$\left[ Fe(CN)_6 \right]^{4-} > \left[ CoCl_4 \right]^{2-} > \left[ MnCl_4 \right]^{2-}$$

$$C. \left[ Fe(CN)_6 \right]^{4-} > \left[ MnCl_4 \right]^{2-} > \left[ CoCl_4 \right]^{2-}$$

D. 
$$[MnCl_4]^{2-} > [CoCl_4]^{2-} > Fe(CN)_6]^{4-}$$

B.  $[MnCl_4]^{2-} > Fe(CN)_6]^{4-} > [CoCl_4]^{2-}$ 

# Answer: D



A.  $\pi$ -character only

B. Both  $\sigma$  and  $\pi$  characters

**15.** In  $Fe(CO)_5$ , the Fe - C bond possesses:

C. Ionic characters

D.  $\sigma$ -character only



Answer: B

**16.** Which of the following will show optical isomerism? .

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

- B.  $\left[ZnCl_4\right]^{2}$
- $C. \left[ Cr \left( C_2 O_4 \right)_3 \right]^{3-}$
- D.  $\left[ Co(CN)_6 \right]^{3}$

# Answer: C



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**17.** The value of 'spin only' magnetic moment for one of the following configuration is 2.84*B*. *M*. The correct one is:

A.  $d^4$  (in strong field ligand)

- B.  $d^4$  (in weak as well as strong field ligand)
- C.  $d^3$  (in weak as well as strong field ligand)
- D.  $d^5$  (in strong field ligand)

# **Answer: A**



- **18.** Nickel (Z=28) combines with a uninegative monodentate ligand  $X^-$  to form a paramagnetic complex  $\left[NiX_4\right]^{2-}$ . The number of unpaired electron(s) in the nickel and geometry of this complex ion are, respectively:
  - A. One, tetrahedral
  - B. Two, tetrahedral
  - C. One, square planar
  - D. Two, square planar

# **Answer: B**



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- **19.** IUPAC name  $\left[ Co \left( NH_3 \right)_5 \left( NO_2 \right) \right] Cl_2$  is
  - A. Nitrito-N-pentaammine cobalt (III) chloride
  - B. Nitrito-N-pentaammine cobalt (II) chloride
  - C. pentaamminenitrito-N-cobalt (II) chloride
  - D. pentaamminenitrito-N-cobalt (III) chloride

# Answer: D



# 1. Type of isomerism

Type of isomerism

- A) Ionisation
- B) Linkage
- C)Coordination
- D)Hydrate

Pair of examples

(a) 
$$\left[Cr(H_2O)_6\right]Cl_3$$
 and

$$\left\lceil Cr(H_2O)_5 Cl \right\rceil Cl_2.H_2O$$

(b) 
$$\frac{\left[Cr(NH_3)_6\right]\left[Co(CN)_6\right]}{\&\left[Co(NH_3)_6\right]\left[Cr(CN)_6\right]}$$

$$\& [Co(NH_3)_6][Cr(CN)_6]$$

$$\left[ Co(NH_3)_5(NO_2) \right] Cl_2$$

(c) 
$$\frac{\left[Co(NH_3)_5(NO_2)\right]Cl_2}{\left[Co(NH_3)_5(ONO)\right]Cl_2}$$

(d)

$$\left[ Co(SO_4)(NH_3)_5 \right] Br \& \left[ Co(Br)(NH_3)_5 \right] SO_4$$

The correct match is

$$A$$
.  $a$   $b$   $c$   $d$ 

$$B.b$$
 a  $d$   $c$ 

D.dbca

# Answer: 3



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2. Match the following columns

COLUMN-I COLUMN-II

q) Tetragonal

(Complex) (Geometry)

A)  $[Ni(CN)_4]^{2-}$  p) Tetrahedral

C)  $[Co(en)_3]^{3+}$  r) Sqaure planar

D)  $[C u C l_4]^{3-}$  s) square pyramid t) Octahedral

t p

B. r p t (q, t)

B)  $\left[ZnCl_{4}\right]^{2-}$ 



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3.	Match	the	following	columns
COLUMN-I		COLUMN-II		

(Equivalent conductance) (formula)

- A) 229 p)  $[Pt(NH_3)_5Cl]Cl_3$
- B) 97 q)  $[Pt(NH_3)_3Cl_3]Cl$
- C) 404 r)  $[Pt(NH_3)_4Cl_2]Cl_2$ 
  - D) 523 s)  $[Pt(NH_3)_6]Cl_4$
  - A. s p q r
  - B.r q s p
  - C.rqsp



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**4.** Match the following columns

COLUMN-II

(Complex)  $(\mu \text{ effective})$ 

A)  $K_3[MnF_6]$  p) 1.70

B)  $K_3[VF_6]$  q) 2.8

C)  $K_2[Mn(CN)_6]$  r) 3.8

D)  $K_3[TiF_6]$  s) 4.9

 $A. \frac{A}{s} \frac{B}{q} \frac{C}{r} \frac{D}{p}$ 

B. q s r p

C.s q p r



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5. Match the following columns

Set -1 Set -11

- **A)** Co-ordination number 3 (1) Pentagonal bipyramidal
- **B) Co-ordination** number 2 2) Trigonal bipyramidal
- C) Co-ordination number 5 3) Linear
- **D)** Co-ordination number 7 4) Trigonal planar The correct matching is

A B C D

A. <sub>4</sub> 3 2 1

B. 1 2 3 4

c. 4 3 1 2

COLUMN-I

# **Answer: 1**



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6.	Match	the	following	columns

**COLUMN-II** 

0 0 2	
(Complex Ions)	(No. of unpaired
_	electrons)

A) 
$$[CrF_6]^{4-}$$
 p)One  
B)  $[MnF_6]^{4-}$  q) Two

C) 
$$[Cr(CN)_6]^{4-}$$
 r) Three

D) 
$$[Mn(CN)_6]^{4-}$$
 s) Four t) Five

q p

B. q r s t

D.pqrs

# Answer: 1



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7.	Match	the	following	columns
,	COLUMN-I	CO	OLUMN-II	

(Complex) (O.N. of Co) p) -1  $\mathbf{A)} \left[ Co(NCS)(NH_3)_5 \right] (SO_3)$ 

q) 0 B)  $Na[Co(CO)_4]$ 

C)  $Na_{4}[Co(S_{2}O_{3})_{3}]$  $r)_{+1}$ D)  $Co_2(CO)_8$ s) +2

t) +3

B.tpsq



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**8.** Match the list I and II and pick the correct matching from the codes given below

List - I List - II (complex) (Structure and magnetic moment)

- a)  $\lceil Ag(CN) \rceil$  1) square planar and 1.73 BM
- b)  $\left[ Cu(CN)_{A} \right]^{3-}$  2) Linear and zero
- c)  $\left[ Cu(CN)_{6} \right]^{4-}$  3) Octahedral and zero
- d)  $\left[Cu(NH_3)_4\right]^{2+}$  4) tetrahedral and zero
- e)  $\left[ Fe(CN)_{6} \right]^{4-}$  5) octrahedral and 1.73 BM

A. a-2, b-4, c-5, d-1, e-3

- B. a-5, b-4, c-1, d-3, e-2
- C. a-1, b-3, c-4, d-2, e-5
- D. a-4, b-5, c-2, d-1, e-3



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**9.** Two statements 'A' and 'R' are given below Select your answers to these items using the codes given below:

Assertio (A) : Among the cobalt (III) complexes  $\left[ Co \left( NH_3 \right)_6 \right]^{3+}$  is a diamagnetic but  $\left[ CoF_6 \right]^{3-}$  is paramagnetic.

Reason (R) : Hybridisation of  $\left[Co\Big(NH_3\Big)_6\right]^{3+}$  is  $sp^3d^2$  where as hybridisation of Co in  $\left[CoF_6\right]^{3-}$  is  $d^2sp^3$ .

A. Both A and R are correct and R is the correct explanation of A

B. Both A and R are correct and R is not a correct explanation of

Α

C. A is true but R is false

D. A is false but R is true

# Answer: 3



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**10.** Two statements 'A' and 'R' are given below Select your answers to these items using the codes given below:

Assertion(A) : The complex ion cis -  $\left[Co(en)_2Cl_2\right]^+$  is optically active Reason (R) : It is an octahedral complex

A. Both A and R are correct and R is the correct explanation of A

B. Both A and R are correct and R is not a correct explanation of

Α

- C. A is true but R is false
- D. A is false but R is true

#### Answer: 2



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## **EXERCISE-5**

ligands (in proper number so that coordination number of Pt vecomes six). Which of the following can be its correct IUPAC name?

1. A complex cation is formed by Pt (in some oxidation state) with

- A. Diammineethylenediaminedithiocyanato-S-iplatinum (II)
- B. Diammineethylenediaminedithiocyaanato-S- platinate (IV) ion.
- $\hbox{C. Diammineethylendiamined ithio cyanato-S-platinate (IV) ion.}\\$

D. Dimminebis (enthlenediamine) dithiocyanato-S-platinum (IV)

ion.



2. The correct order of their paramagnetic moment (spin only) is

$$P: [FeF_6]^{3}$$
-,  $Q: [CoF_6]^{3}$ -

$$R: \left[V\left(H_2O\right)_6\right]^{3+}, S: \left[Ti\left(H_2O\right)_6\right]^{3+}$$

A. 
$$P > Q > R > S$$

B. 
$$P < Q < R < S$$

$$C. P = Q = R = S$$

D. 
$$P > R > Q > S$$

- **3.** Which of the following statement is incorrect?
  - A.  $Ti(NO_3)_4$  is a colourless compound.
  - B.  $\left[Cu\left(NCCH_3\right)_4\right]^+BF_4^-$  is a colourless comopound.
  - C.  $\left[Cr(NH_3)_6\right]^{3+3Cl}$  id a coloured compound.
  - D.  $K_3[VH_6]$  is a colourless compound.



- **4.** What will be the 'spin only' magnetic moment of the complex formed when Fe  $(SCN)_3$  reacts with solution containing excess F-?
  - A. 2.83 B.M.
  - B. 3.87 B.M.

C. 5.92 B.M.

D. 1.73 B.M.



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- 5. The compound having a tetrahedral geometry is .
  - A.  $\left[Pd(CN)_4\right]^{2}$
  - B.  $\left[Ni(CN)_4\right]^{2}$
  - C.  $[PbCl_4]^{2-}$
  - D.  $\left[NiCl_4\right]^2$



**6.** It is an experimental fact that  $Cs_2[CuCl_4]$  is orange coloured but  $\left(NH_4\right)_2[CiCl_4]$  is yellow. It id further known that total paramagnetic moment of an unpaired electron is due to spin as well as due to nature of orbitlal, 'd' orbital contributing more than 's' or 'p' Thus the total paramagnetic moment of organe compound is found to be more than that of yellow compound. Then which of the following is correct.

- A. Anion of organe compound is tetrahedral and that of yellow is square planar.
- B. Anion or orange compound is squance and that of yellow is tetrahedral.
- C. Both the anions are terrahedral
- D. Both the anion are square planar.

- A.  $sp^3$
- B.  $dsp^2$
- $C. d^2sp^3$
- D.  $sp^3d^2$



8. Which of the following statements is correct?

A. Potassium ferrocyanide is diamagnetic where as potassium ferricyanide is paramagnetic.

B. Crystal field splititing in ferrocyanide ion is greater than that

C. The geometry of  $Ni(CO)_4$  and  $\left[NiCl_2\Big(PPh_3\Big)_2\right]$  are tetrachedral.

D. (A) & (D)



of ferricyanide ion.

- **9.** Amongst  $\left[Co(ox)_3\right]^{3-}$ ,  $\left[CoF_6\right]^{3-}$  and  $\left[Co\left(NH_3\right)_6\right]^{3+}$ :
  - A.  $\left[Co(ox)_3\right]^{3-}$  and  $\left[CoF_6\right]^{3-}$  are paramagnetic and  $\left[Co\left(NH_3\right)_6\right]^{+3}$  is diamagnetic.
  - B.  $\left[Co(ox)_3\right]^{3-}$  and  $\left[Co\left(NH_3\right)_6\right]^{+3}$  are paramagnetic and

 $[CoF_6]^{3-}$  is paramagnetic.

C.  $\left[Co(ox)_3\right]^{3-}$  and  $\left[Co\left(NH_3\right)_6\right]^{3+}$  are diamagnetic and  $\left[CoF_6\right]^{3-}$  is paramagnetic.

D.  $\left[Co\left(NH_3\right)_6\right]^{3+}$  and  $\left[CoF_6\right]^{3-}$  are paramagnetic and  $\left[Co(ox)_3\right]^{3-}$  is diamagnetic.



10. All the following complexes show decrease in their weights when placed in a magnetic balance then the group of complexes having tetrahedral geometry is:

$$Ni(CO)_4$$

$$K[AgF_4]$$
 $Na_2[Zn(CN)_4]$ 

$$K_2[PtCl_4]$$

$$\left[ RhCl \left( PPh_3 \right)_3 \right]$$

- A. (ii), (iii),(v)
- B. (i), (II), (iii)
- C. (i), (III), (iv)
- D. None of these



- **11.** The geometry of  $\left[NiCl_4\right]^{2-}$  and  $\left[Ni\left(PPh_3\right)_2Cl_2\right]$  are :
  - A. tetrahedral and square planar redpectively.
  - B. both tetrahedral.
  - C. both square planar.
  - D. square planer and tetrahedral respactively.



**12.** 
$$\left[ Co(en)_2 \left( H_2 O \right)_2 \right]^{3+} + en \rightarrow \text{ complex (X) is :}$$

A. it is a low spin complex

B. it is diamagnetic

C. it show geometrical isomerism

D. (A) & (B)



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13. Which one of the following complexes exhibit chirality?

A. 
$$\left[Cr(ox)_3\right]^3$$

$$B.\,cis-\left[PtCl_2(en)\right]$$

C. 
$$cis - \left[RhCl_2(NH_3)_2\right]^{-1}$$

D. mer - 
$$\left[ Co(NO_2)_3(\text{trien}) \right]$$



**14.** On treatment of  $\left[Pt(NH_3)_4\right]^{2+}$  with CI compound I is formed

When  $[PtCl_4]^{2-}$  reacted with aq.  $NH_3$  the complex II is formed.

Point out the correct statement.

A. I cis. II trans, both tetrahedral

B. I cis, II trans, both square planar

C. I trans, II cis, both tetrahedral

D. I trans, II cis, both square planar



**15.** For the empirical formula,  $Pt(NH_3)_2Cl_2$  the number of possiblecoordination isomers would be : (oxidation number of platinum should be +2 in all isomeric forma)

- **A.** 1
- B. 2
- C. 3
- D. 4



**16.** The total number of isomer shown by  $\left[ Co \left( NH_3 \right)_4 \left( NO_2 \right)_2 \right] \left( NO_3 \right)$  complex is :

- A. 10
- B. 6
- C. 4
- D. 12

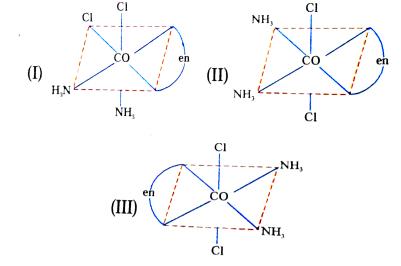


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# **EXERCISE-6**

1. Three arrangements are shown for the complex

$$\left[ {\it Co(en)} \left( {\it NH}_3 \right)_2 {\it Cl}_2 \right]^+$$
 pick up the wrong statement.



- A. I and II are geometrical isomers
- B. II and III are optical isomers
- C. I and III are optical isomers
- D. II and III are geometrical isomers

## Answer: B,C,D



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2. Which of the following isomerism is/are shown by the complex

$$\left[ CoCl_2 \left( H_2O \right)_2 \left( NH_3 \right)_2 \right] Br?$$

- A. Ionization
- B. Linkage
- C. Geometrical
- D. optical

### Answer: A,C,D



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**3.** Co-ordination number of Cr in  $CrCl_3.5H_2O$  is six. The volume of 0.1

N  $AgNO_3$  needed to ppt. the chorine in outer sphere in 200 ml of

0.01 M solution of he possible complex es is/are:

A. 140 ml

- B. 40 ml
- C. 80 ml
- D. 20 ml

### Answer: B,D



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- **4.** Which of the following statement(s) is (are) correct with reference to the ferrous and ferric ions ?
  - A.  $Fe^{3+}$  gives brown colour solution with potassium ferricyanide.
  - B. gives blue precipitate with potassium ferricyanide.
  - C.  $Fe^{3+}$  gives red colour solution with potassium sulphocyanide.
  - D.  $Fe^{2+}$  gives brown colour solution with potassium sulphocyanide.

## Answer: A,B,C



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- **5.** The complex  $\left[Cr(H_2O)_4Br_2\right]$  gives the test for :
  - A. Br
  - B. *Cl* -
  - C.  $Cr^{3+}$
  - $D.H_2O$

#### **Answer: B**



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6. Which of the following ligands can show linkage isomerism?

A. 
$$NO_2$$

B. SCN

C. CNO

D.  $CO_3^2$ 

# Answer: A,B,C



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# **EXERCISE-7**

**1.** Which one of the following has the correct order of  $\Delta_o$ ?

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

$$B. \left[ Co \left( H_2 O \right)_6 \right]^{2+} < \left[ Co \left( H_2 O \right)_6 \right]^{3+}$$

$$C. \left[ Co \left( H_2 O \right)_6 \right]^{3+} > \left[ Rh \left( H_2 O \right)_6 \right]^{3+}$$

D. 
$$\left[ Co(NH_3)_6 \right]^{3+} < \left[ CoF_6 \right]^{3-}$$



**2.** Wilkinson's catalyst react with  $H_2$  to from an octahedral complex in which Rh (Z= 45) has the following electronic configuration in the ligand field  $t_{2g}^{2.2.2,e_g^{0.0}}$ . Then which of the following is (are) correct about this new complex ?

A. It is diamagnetic

B. Its IUPAC name is chlorodihydridotris (triphenylphosphine)

rohidium (III)

C. Hybridisation of Rh (I) is  $d^2sp^3$ 

D. It is a paramagnetic complex

**3.** In the cystal field of the complex  $\left[Fe(Cl)(CN)_4(O_2)\right]^{4-}$  the electronic configuration of metalis found to be  $t_{2g}^6$ ,  $e_g^o$  then which of the following is/are true about this complex ion :

A. It is a paramagnetic complex

B. O-O bond length will be more than found in  $\mathcal{O}_2$  molecule

C. Its IUPAC name will be chloridotetacyanidosuperoxidoferrate

D. It will show geometrical as will as optical isomerism



(II) ion.

4. Select the correct statement (s) for the coordination compound

$$K_2 \Big[ Cr \Big( NO^+ \Big) \Big( NH_3 \Big) (CN)_4 \Big].$$

A. Its IUPAC name is potassium

amminetetracyanidonitrosoniumchromate (I).

- B. Its' spin only' magnetic moment is 8 B.M.
- C. Its hybridisation is  $d^2sp^3$
- D. It shows geometrical isomerism



- **5.** Which of the following complex (s) is/are correctly matched with their geometry?
  - A.  $\left\lceil CoCl_4 \right\rceil^{2-}$  -tetrahedral

- B.  $\left[ Co(PY)_4 \right]^{2+}$  -square planar.
- C.  $\left[ Cu(CN)_4 \right]^{3-}$  -tetrahedral
- D.  $[Fe(CO)_4]^{2-}$  -square planar.

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## **EXERCISE-8**

**1.** Match the complexes listed in column-I with characteristic (s)/type of hybridisation olisted in column-II.

```
Column-I

(A) [Co(OX)<sub>3</sub>]<sup>3-</sup>

(B) [Co(F)<sub>6</sub>]<sup>3-</sup>

(C) [Fe(H<sub>2</sub>O)<sub>5</sub>(NO)]<sup>2+</sup>

(D) [Co(NH<sub>3</sub>)<sub>6</sub>]<sup>3+</sup>

(E) [Mn(CN)<sub>6</sub>]<sup>3-</sup>

Column-II

(p) Chelation

(q) 2.8 BM 'spin only' magnetic moment

(r) d<sup>2</sup>sp<sup>3</sup> and paramagnetic

(s) sp<sup>3</sup>d<sup>2</sup> and paramagnetic

(t) +1 oxidation state of central metal

(u) d<sup>2</sup>sp<sup>3</sup> and diamagnetic
```



2. Match the complexes listed in column-I with characteristic(s)

listed in column-II.

(C) 
$$[Ni(PF_3)_4]$$
 (D)  $[Ni(PPh_3)_2 Br_2]$   
Column – II  
(p) Terahedral  
(q)  $\pi$  back bonding

(B)  $[Fe(CO)_2(NO)_2]$ 

(r) Diamagnetic(s) One of the ligand is three electeron donor.



Column-I

(A)  $[Ni (CO)_4]$ 

hybridisation listed in column-II.

Column –I

(A)  $[Co(H_2O)_6]^{3+}$  (B)  $[CoF_6]^{3-}$ (C)  $[Co(NO)]^{3-}$  (D)  $[CoF_6]^{3-}$ 

3. Match the complexes listed in column-I with characteristic (s)/type

(C)  $[Co(NO_2)_6]^{3-}$  (D)  $[CoF_3(H_2O)_3]$ Column –II (p)  $d^2sp^3$  (q)  $sp^3d^2$ 

(r) Diamagnetic (s) Paramagnetic



**4.** Match the complexes listed in column-I with characteristic (s)/type

of bridisation listed in column -II

Column – I (A)  $[Co (en)_3]^{3+}$  (B)  $[Co (OX)_3]^{3-}$ (C)  $[Co (H_2O)_6]^{2+}$  (D)  $[Co (NO_2)_6]^{4-}$ Column – II

(p) sp<sup>3</sup>d<sup>2</sup> hybridisaion (q) Diamagnetic

(r) d<sup>2</sup>sp<sup>3</sup> hybridisation (s) Paramagnetic

(t) Chelating reagent



5. Match the complexes listed in colmn-I with type hybridisation

listed in column-II.

Column - I

(A)  $[Au F_4]^-$  (B)  $[Cu(CN)_4]^{3-}$  (C)  $[Co(NH_3)_6]^{2+}$  (D)  $[Fe(H_2O)_5 NO]^{2+}$ 

Column - II

(p) dsp² hybridisation (q) sp³ hybridisation (r) sp³d² hybridisation (s) d²sp³ hybridisation

**6.** Select the correct option (s) for the coordination compounds and their respectively isomeric forms.

Column-I

(A) [Co(en)<sub>2</sub>NH<sub>3</sub>Cl]SO<sub>4</sub>

(B)  $[Co(NH_3)_4(NO_2)_2](NO_3)$ 

(C)  $[Co(en)(pn)(NO_2)_2]C1$  (D)  $[Co(gly)_3]$ 

(p) Optical isomer(q) Geometrical isomer(r) Ionization isomer(s) Linkage isomer



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### **EXERCISE-9**

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

STATEMENT-2 : Energy difference between d level ( $i.e.\Delta$ ) for  $H_2O$  complex (paramagnetic) is in the visible region and that for the cyano complex (diamagnetic) is in the UV region.

A. Statement-1 is True, Statement-2 is True, Statement-2 is a correct explanation for statement-1.

B. Statement-1 is True, Statement-2 is True, Statement-2 is NOT a correct explation for statement-1

C. Statement-1 is True, Statement-2 is False.

D. Statement-1 is False, Statement-2 is True.



**2.** STATEMENT-1 The value of  $\Delta_o$  for  $M^{3+}$  complexes are always much higher than value for  $M^{2+}$  complexes

STATEMENT-2 : The crystal field stabilization energy of

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

A. Statement-1 is True, Statement-2 is True, Statement-2 is a correct explanation for statement-1.

B. Statement-1 is True, Statement-2 is True, Statement-2 is NOT a correct explation for statement-1

C. Statement-1 is True, Statement-2 is False.

D. Statement-1 is False, Statement-2 is True.



with strong field as well as with weak field ligands are square planar.

**3.** STATEMENT-1: All the complexes of Pt  $(+ \Box \Box)$  and Au  $(+ \Box \Box)$ 

STATEMENT-2 : The crystal filed spliting energy is larger for secong

and third row transition elements, and for more highly charged species. This large value of crystal field splitting energy energetically favours the pairing of electron for square planar geometry.

A. Statement-1 is True, Statement-2 is True, Statement-2 is a correct explanation for statement-1.

B. Statement-1 is True, Statement-2 is True, Statement-2 is NOT a correct explation for statement-1

C. Statement-1 is True, Statement-2 is False.

D. Statement-1 is False, Statement-2 is True.



**4.** STATEMENT-1 : diamminedichloroplatinum ( $\Box$   $\Box$ ) is more soluble in water than trans-diammine dichloroplatinum ( $\Box$   $\Box$ )

STATEMENT-2 : Diamminedichloroplatinum ( $\square$   $\square$ ) is square planar and diamagnetic.

A. Statement-1 is True, Statement-2 is True, Statement-2 is a correct explanation for statement-1.

B. Statement-1 is True, Statement-2 is True, Statement-2 is NOT a correct explation for statement-1

C. Statement-1 is True, Statement-2 is False.

D. Statement-1 is False, Statement-2 is True.



**5.** STATEMENT-1 :  $\left[Co\left(NH_3\right)_4Cl_2\right]^+$  can exist in cis-and trans-forms but neither can display optical activity.

STATEMENT-2: Each of these possesses at least one plane of symmetry.

A. Statement-1 is True, Statement-2 is True, Statement-2 is a correct explanation for statement-1.

B. Statement-1 is True, Statement-2 is True, Statement-2 is NOT a correct explation for statement-1

C. Statement-1 is True, Statement-2 is False.

D. Statement-1 is False, Statement-2 is True.



**6.** STATEMENT-1: In complex  $\left[Cr\left(NH_3\right)_4BrCl\right]$ CI, the spin only magnetic moment is close to 1.73 B.M.

STATEMENT-2: All known monoculear complexeas of chromim (III)

irrespective of the strength of the ligand field, must hae three unpaired electrons.

Consider the following statement and arrange in the order of trur/flase as given in the codes.

A. Statement-1 is True, Statement-2 is True, Statement-2 is a correct explanation for statement-1.

B. Statement-1 is True, Statement-2 is True, Statement-2 is NOT a correct explation for statement-1

C. Statement-1 is True, Statement-2 is False.

D. Statement-1 is False, Statement-2 is True.



7. S-1:  $\left[Cr(NH_3)_6\right]^{3+}$  is a inner orbital complex with

S-2: The complex formed by joining the CN ligands to  $Fe^{3+}$  ion has theoretical value fo 'spin only' magnetic moment equal to 1.73 B.M.

S-3:  $Na_2S + Na_2\Big[Fe(CN)_5NO\Big] \rightarrow Na_4\Big[Fe(CN)_5NOS\Big]$ , In reactan and product the oxidation states of iron are same

A. FTF

B.TTF

C. T T T

D. F F F



**8.** s-1:  $[MnCl_6]^3$ ,  $[FeF_6]^{3-}$  and  $[CoF_6]^{-3}$  are paramagnetic having four, five and four unpaired electrons respectively.

S-2: Valence bond theory gives a quantitative interpretaion of the thermodynamic stabilities of coordination compounds.

S-3: The crystal field splitting  $\Delta_o$  depends upon the field produced by the ligand and charge onn the metal ion.

A. TTT

B. T F T

C. F T F

D. T F F



**9.** S-1: The  $\left[Co(ox)_3\right]^{3-}$  complex is diamagnetic andgains stability through chelation.

S-2: The  $\left[Co\left(NO_2\right)_6\right]^4$  is inner orbital complex involving  $d^2sp^3$ 

hybridisation and is paramagnetic.

S-3: The complex  $\left[PtCl_4\right]^{2-}$  is square planar and is diamagnetic.

A. TTT

B. F F T

C. TFT

D. T T F



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10. Consider the following satements:

 ${\cal S}_1$ : Generally square planare complexes show geometrical isomerism but do not exhibit optical isomerism because thy do not possess plane of symmetry .

$$S_2: \Delta_t = \frac{4}{9} \Delta_0$$

 $S_3$ : In octahedral complexes each electron entering teh  $t_{2g}$  orbitals

stabilizes the complex ion by 0.4  $\Delta_0$  and each electron entering teh  $e_g$  orbital destabilizes the complex by an amount of 0.6  $\Delta_0$ 

Select the correct statement from the codes given below.

11. The ionisation isomer of  $\left[Cr(H_2O)_4Cl(NO_2)\right]Cl$ 

- A. FTT
- B. F F T
- C. T F T
- D. T T F



- - B. F F T

A. FTT

C. T F T



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

 $S_1$ :Both  $\left[Co(ox)_3\right]^{3-}$  and  $\left[CoF_6\right]^{3-}$  are paramagnetic.

 $S_2$ : $CoCl_3$ 3 $NH_3$  complex is non-conducting.

 $S_3$ :The number of possible isomers for the complex  $\left[Pt\Big(NO_2\Big)(py)(OH)\Big(NH_3\Big)\right] \text{ is six.}$ 

 $S_4$ :The oxidation state of iron in brown ring complex

$$\left[Fe\left(H_2O\right)_5NO^+\right]SO_4$$
 is + || Where NO is  $NO^+$ 

A. FTTF

B. TTTT

C. FTTF



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is the coordination number of Co 13. What in

$$\left[Co(NH_3)_4(H_2O)Br\right](NO_3)_2?$$

- A. FTTF
- B. TTTT
- C. FTTF
- D. TTTF



1. Double salts are addition compounds which lose their identity in aqueous solution whereas complexes which are also addition compounds do not lose their identity in aqueous solution. The coordination compounds show isomerism and find applications in photography, qualitative analysis, metallurgy, water purification and in the treatment of various diseases.

Which of the following statment is incorrect?

- A. Alum is a double salt.
- B. BDTA salt of calcium is used in the treatment of lead poisoning.
- C. Effective atomic number of the metals in complexes  $\left[Ni(CO)_4\right]$  and  $\left[Fe(CN)_6\right]^{4-}$  is same.
- D. Chloridotris (triphenyl phosphine) rhodiuym (I) is effective heterogeneous catalyst for hydrogenation of alkenes.

2. Double salts are addition compounds which lose their identity in aqueous solution whereas complexes which are also addition compounds do not lose their identity in aqueous solution. The coordination compounds show isomerism and find applications in photography, qualitative analysis, metallurgy, water purification and in the treatment of various diseases.

Which of the following statement is true for the complex  $\left[ Co \left( NH_3 \right)_4 Br_2 \right] NO_2 ?$ 

- A. It shows ionisation, linkage and geometrical isomerism.
- B. It does not show optical isomerism because its cis and trans forms each have at least one plane of symmetry.
- C. Its ionisation isomers cannot be differenatiated symmetry

D. (A) and (B) both.



**3.** Double salts are addition compounds which lose their identity in aqueous solution whereas complexes which are also addition compounds do not lose their identity in aqueous solution. The coordination compounds show isomerism and find applications in photography, qualitative analysis, metallurgy, water purification and in the treatment of various diseases .

Choose the correct option if the complex  $\left[PtCl_2(en)_2\right]^{2+}$ .

- A. Platinum is in +2 oxidation state
- B. Racemic mixture is obtained on mixing mirror images of its trans form in 1:1 molar ratio.
- C. It has two five membered chelating rings

D. (B) and (C) both



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### **Comprehension-2**

1. In coordination chemistry there are a veriety of methods applied to find out the structure of complexes. One method involves treating the complex with known reagents and from the nature of reaction, the formula0 of the complex can be predicted. An isomers of the complex  $Co(en)_2(H_2O)Cl_2Br$ , on reaction with concentrated  $H_2SO_4$  (dehydrating agent ) it suffers loss in weight and on reaction with  $AgNO_3$  solution it gives a white precipitate which is soluble in  $NH_3(aq)$ .

The correct formula of the complex is:

A. 
$$\left[ CoCIBr(en)_2 \right] H_2O$$

$$B. \left[ CoCl(en)_2 \left( H_2O \right) \right] BrCl$$

C. 
$$\left[ CoBr(en)_2 \left( H_2O \right) \right] Cl_2$$

D. 
$$\left\lceil CoBrCl(en)_2 \right\rceil Cl. H_2O$$



**2.** In coordination chemistry there are a variety of methods applied to find out the structures of complexes. One method involves treating the complex with known reagents and from the nature of reaction , the formula of the complex can be predicated. An isomer of the complex  $Co(en)_2(H_2O)Cl_2Br$ , on reaction with concentrated  $H_2SO_4$  (dehydrating agent) it suffers loss in weight and on reaction with  $AgNO_3$  solution it gives a white precipitate which is soluble in  $NH_3(aq)$ .

If all the ligands in the coordination sphere of the above complex be replaced by  $SCN^-$ , then the paramagnetic moment of the  $\left[ Co(SCN)_4 \right]^{2-}$  (due to spin only) will be:

- A. 2.8 BM
- B. 5.9 BM
- C. 4.9 BM
- D. 1.73 BM



**3.** In coordination chemistry there are a variety of methods applied to find out the structure of complexes. One method involves treating the complex with known reagents and from the nature of reaction, the formula of the complex can be predicted. An isomer of the complex  $Co(en)_2(H_2O)Cl_2Br$ , on reaction with concentrated  $H_2SO_4$ 

(dehydrating agent) it suffers loss in weight and on reaction with  $AgNO_3$  solution it gives a white precipitate which is soluble in  $NH_3$ 

(aq).

If one mole of original complex is treated with excess  $Pb(NO_3)_2$ solution, then the number of moles of white precipitate (of PbCl<sub>2</sub>) formed will be:

A. 0.5

B. 1.0

C. 0.0

D. 3.0

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**Comprehension-3** 

**1.** 
$$Co^{2+}(aq) + SCN^{-}(aq) \rightarrow complex(X)$$
.

 $NH_{4}OH$ 

 $Ni^{2+}(aq)$  + Dimethylglyoxime  $\rightarrow$  Complex(Y).

The coordination number of cobalt and nickel in complexes X and Y are four.

The IUPAC names of the complexes (X) and (Y) are respectively.

A. tetrathicyanato-S -Cobalt (II) and bis (dimethylglyoximate ) nickel (II).

B. tetrathicyanato-S -Cobaltate (II) and bis (dimethylglyoximato )

nickel (II).

C. tetrathicyanato-S -Cobaltate (II) and bis (dimethylglyoximato )

nickelate (II).

D. tetrathicyanato-S -Cobaltate (II) and bis (dimethylglyoximato) nickel (II).

**2.** 
$$Co^{2+}(aq) + SCN^{-}(aq) \rightarrow complex(X)$$
.

$$NH_4OH$$

$$Ni^{2+}(aq)$$
 + Dimethylglyoxime  $\rightarrow$  Complex( $Y$ ).

The coordination number of cobalt and nickel in complexes X and Y are four.

The geometry of complexes (X) and (Y) are respectively:

- A. tetrahedral and sqaure planar.
- B. both tetrahedral.
- C. square planer and tetrahedral
- D. both square planer.



**3.** 
$$Co^{2+}(aq) + SCN^{-}(aq) \rightarrow complex(X)$$
.

 $NH_{4}OH$ 

 $Ni^{2+}(aq)$  + Dimethylglyoxime  $\rightarrow$  Complex(Y).

The coordination number of cobalt and nickel in complexes X and Y are four.

Select the correct statement for the complexes (X) and (Y).

- A. (X) is paramagnetic with two unpaired electrons
- B. (Y) is diamagnetic and shows intermolecular H-bonding.
- C. (X) is paramagnetic with three unpaired electrons and(Y) is

diamagnetic

D. (X) and (Y) both are diamagnetic .



1. Square planar complexes are formed by  $d^8$  ions with strong field ligands. The crystal field splitting  $\Delta_0$  is larger for the second and theird row transition elements and for more highly charged species. All the complexes having  $4d^8$  and  $5d^8$  configurations are mostly square planar including those with weak field ligands such as halide ions square planar complexes can show geometrical isomerism but they do not show optical isomerism due to the presence of plane of symmetry

Among the following complexes which has a square planar geometry?

(a) 
$$\left[RhCI(CO)\left(PPh_3\right)_2\right]$$

(b) 
$$K_3 \left[ Cu(CN)_4 \right]$$

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

(d) 
$$K_2 \Big[ Zn(CN)_4 \Big]$$
 .

A. 
$$Ni(+\Box\Box)$$
 and Rh  $(+\Box)$  with strong field ligands

B. 
$$Rh(+\Box)$$
 and Ag  $(+\Box\Box)$  with strong field ligand.

C.  $Pd(+ \Box \Box)$  ,Pt  $(\Box \Box)$  and Au  $(+ \Box \Box \Box)$  with strong and weak field ligands

D. None



**2.** Square planar complexes are formed by  $d^8$  ions with strong field ligands. The crystal field splitting  $\Delta_0$  is larger for the second and theird row transition elements and for more highly charged species. All the complexes having  $4d^8$  and  $5d^8$  configurations are mostly square planar including those with weak field ligands such as halide ions square planar complexes can show geometrical isomerism but they do not show optical isomerism due to the presence of plane of symmetry

Among the following complexes which has a square planar geometry?

- (a)  $\left[RhCI(CO)\left(PPh_3\right)_2\right]$ (b)  $K_3\left[Cu(CN)_4\right]$
- (c)  $\left[Ni(CO)_4\right]$ 
  - (d)  $K_2 \left[ Zn(CN)_4 \right]$ .
    - A.  $\left[RhCI(CO)\left(PPH_3\right)_2\right]$ B.  $K_3\left[CU(CN)_4\right]$
    - $C. K_2 \Big[ Zn(CN)_4 \Big]$

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

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## Comprehension-5

1. In metal carbonyls. There is synergic bonding interaction between metal and carbon monoxide. This leads to increase in strength of metal ligand bond and decrease in bond ordeer of CO in carbonyl complex as compared to band order in order in carbon monoxide.

The increases in bond length in CO as compared to carbon monoxide is due to :

A. the donation of lone pair of electrons from a filled d-orbital of the metal atom

B. the donation of pair of electrons from a filled d-orbital of metal into the vacant antibonding  $\pi$  orbital of carbon monoxide.

C. (A) and (B)

D. None

2. In metal carbonyls, there is synergic bonding interaction between metal and carbon monoxide. This leads to increase in strength of metal ligands bond and decrease in bond order of CO in carbonyl complex as compared to bond order in carbon monoxide.

Simple carbonyls are invariably spin-paired complexes except for vanadium metal.

Which one of the following metal carbonyls are inner orbital complexes with diamagnetic property?

$$(P)Ni(CO)_4$$
  $(Q)Fe(CO)_5$ 

$$(R)V(CO)_6$$
  $(S)Cr(CO)_6$ 

Select the correct answer from the codes given below:

### A. I and II only

B. II, III and IV

C. II and IV

D. I,II and IV

### LEVEL - V

- **1.** The hybridization of the metal in  $[CoF_6]^{3-}$  is
  - A.  $sp^3d^2$
  - B.  $d^2sp^3$
  - $C. dsp^3$
  - D.  $sp^3d$

### Answer: a



**2.** The hybridization of Cr in 
$$\left[Cr(en)_3\right]^{3+}$$
 is

A. 
$$d^2sp^3$$

B. 
$$sp^3d$$

$$C. sp^3d^2$$

D. 
$$dsp^3$$

### Answer: a



- **3.** The magnetic moment of the complex  $\left[Ti(H_2O)_6\right]^{3+}$  is
  - A. 3.87 BM
  - B. 1.73 BM
  - C. 2.84 BM

D	5.87	DM
υ.	5.07	DIVI

#### Answer: b



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**4.** Which one of the following coordination numbers can show a square pyramidal geometry?

A. 4

B. 3

C. 7

D. 5

#### Answer: d



**5.** If the formation constant of  $\left[Cu(NH_3)_4\right]^{2+}$  is  $2 \times 10^3$ , then its dissociation constant is

A. 
$$5 \times 10^{-4}$$

B. 
$$2 \times 10^{3}$$

$$C.5 \times 10^{-2}$$

D. 
$$0.2 \times 10^3$$

#### Answer: a



- **6.** The IUPAC of  $\left[Fe\left(H_2O\right)_6\right]Cl_3$  is
  - A. Hexaaquirion(III) chlorine
  - B. Hexaquorion(III) trichloride

- C. Hexaaquairion(III) chloride
- D. Hexa aquairion(III) trichloride

#### Answer: c



# **7.** IUPAC name of $\left[Co(en)_2Cl_2\right]Cl$ is

- A. dichloridobis (ethylene diammine) cobalt (III) Chloride.
- B. dichloridobis (ethylenediammine) cobalt (III) Chloride
- C. dichlorobis (ethlene diamine) cobalt (III) Chloride.
- D. dichloridobis (ethylene diamine) cobalt (III) chloride

#### Answer: d



- 8. Which is not true statement?
  - A.  $\left[Ti(H_2O)_6\right]^{3+}$  is coloured due to d-d transition.
  - B.  $\left[Dy(H_2O)_6\right]^{3+}$  is coloured due tof-f transition.
  - C.  $\left[Sc(H_2O)_6\right]^{3+}$  and  $\left[Ti(H_2O)_6\right]^{4+}$  are coloured complexes
  - D.  $Cu^+$  is colourless ion.

#### Answer: c



**9.** Among the following ions, which one has the highest paramagnetism?

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

B. 
$$\left[ Fe \left( H_2 O \right)_6 \right]^{2+}$$

$$C. \left[ Cu \left( H_2 O \right)_6 \right]^{2+}$$

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

### Answer: b



- **10.** Which of the following statements is correct?
  - A.  $\left[CoF_6\right]^{3-}$  and  $\left[Co\left(NH_3\right)_6\right]^{3+}$  both are paramagnetic
    - complexes
    - B.  $\left[CoF_6\right]^{3-}$  and  $\left[Co\left(NH_3\right)_6\right]^{3+}$  both are high spin complex
    - C.  $\left[ CoF_6 \right]^{3-}$  is octahedral while  $\left[ Co \left( NH_3 \right)_6 \right]^{3+}$  has
      - pentagonal pyramid shape

D.  $[CoF_6]^{3-}$  is outer orbital complex while  $[Co(NH_3)_6]^{3+}$  is inner orbital complex

#### Answer: d



**11.** What is wrong about the compound  $K[Pt(\eta^2 - C_2H_4)Cl_3]$ ?

- A. It is called Zeise's salt
- B. It is only  $\pi$ -bonded complex
- C. Oxidation number of Pt is +2
- D. Four ligands surrounds the Platinum atom.

### Answer: b



**12.** Which complex is likely to show optical activity:

A. Trans-
$$\left[Co\left(NH_3\right)_4Cl_2\right]^+$$

$$B. \left[ Cr \left( H_2 O \right)_6 \right]^{3+}$$

C. Cis - 
$$\left[Co(NH_3)_2(en)_2\right]^{3+}$$

D. Trans-
$$\left[Co\left(NH_3\right)_2(en)_2\right]^{3+}$$

### Answer: c



**13.** Which of the following complex ion has a magnetic moment same as  $\left[Cr\left(H_2O\right)_6\right]^{3+}$ ?

$$A. \left[ Mn \left( H_2O \right)_6 \right]^{4+}$$

$$B. \left[ Mn \left( H_2 O \right)_6 \right]^{3+}$$

C. 
$$\left[Fe\left(H_2O\right)_6\right]^{3+}$$
D.  $\left[Cu\left(NH_3\right)_4\right]^{2+}$ 

### Answer: a



**14.** In which of the following ions, the colour is not due to d-d transition?

A. 
$$CrO_4^2$$

$$CTi(H_{-}O)^{3+}$$

B.  $Cu(NH_3)^2$ 

C. 
$$Ti\left(H_2O\right)_6^{3+}$$

D.  $CoF_6^3$ 

## Answer: a



15. Gold and platnum dissolves in aquaregia to produce respectively

- A.  $H[AuCl_4]$  and  $H[PtCl_3]$
- $\operatorname{B.} H \Big[ \operatorname{AuCl}_4 \Big] \text{ and } H_2 \Big[ \operatorname{PtCl}_6 \Big]$
- $C.H_2$   $AuCl_6$  and H  $PtCl_4$
- $\mathrm{D.}\,H_2\!\left[\mathrm{AuCl}_6\right]$  and  $H\!\left[\mathrm{PtCl}_6\right]$

#### Answer: b



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### **Multiple Answer Questions**

1. Which of the following statements is/are true?

A. In metal carbonyl complexes d increases compared to that in

CO molecule

B. The pair of compounds

$$\left[Cr(H_2O)_6\right]Cl_3$$
 and  $\left[CrCl_3(H_2O)_3\right].3H_2O$  shows hydration isomerism

 ${\rm C.}\,d_{{\rm Z}^2}$  orbital of central metal atom/ion is used in  $dsp^2$  hybridisation

D. Facial and meridional isomers associated with  $\left[Ma_3b_3\right]^{n\pm}$  type complex compound, both are optically inactive

Answer: a,d



**2.** A d-block element forms octahedral complex but its spin magnetic moment remains same either in strong field or in weak field ligand. Which of the following is/are correct?

A. d-block element always forms colourless compound

B. Number of electrons in  $t_{2g}$  orbitals are higher than in  $e_g$  orbitals

C. It can have either  $d^3$  or  $d^8$  configuration

D. It can have either  $d^7$  or  $d^8$  configuration

## Answer: b,c



3. The complexes that have a magnetic moment of 1.73 BM is

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

$$B. \left[ V \left( H_2 O \right)_6 \right]^{4+}$$

$$C. \left[ Mn \left( H_2 O \right)_6 \right]^{2+}$$

$$D. \left[ Mn \left( H_2 O \right)_6 \right]^{3+}$$

## Answer: a,b



# **4.** The square planar complexes are

A. 
$$\left[Pt\left(NH_3\right)_2Cl_2\right]$$

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

$$C. \left[ NiCl_4 \right]^{2-}$$

D. 
$$\left[ PtCl_4 \right]^{2}$$

## Answer: a,b,d

A. 
$$Ni(CO)_4$$

$$\operatorname{B.}\left[\operatorname{FeCl}_4\right]^{\text{-}1}$$

C. 
$$\left[MnCl_4\right]^{2}$$

D. 
$$\left[ CoCl_4 \right]^{2}$$

### Answer: a,b,c,d



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6. The diamagnetic complexes are

A. 
$$\left[ Cr(CO)_6 \right]$$

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

$$\mathsf{C.}\left[\mathit{Fe}(\mathit{CN})_{6}\right]^{4}$$

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

### Answer: a,b,c



### 7. The strong field ligands or moderate ligand are

A.  $NH_3$ 

B. en

C. CO

D. *CN* -

### Answer: a,b,c,d



**8.**  $d \rightarrow d$  transition is possible in

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

$$\mathsf{B.}\mathit{KMnO}_4$$

$$C. K_2 Cr_2 O_7$$

$$\mathrm{D.}\left[\mathit{Co}\!\left(\mathit{NH}_{3}\right)_{6}\right]^{\!3\,+}$$

### Answer: a,d



- 9. Which of the following can act as bridging ligands?
  - A. OH
    - $B.H_2O$
    - C. CO

D.	Cl -

Answer: a,c,d



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10. Which of the following ligands can show linkage isomerism?

A. en

 $B.NO_2$ 

**C**. *CN* <sup>-</sup>

D. SCN

Answer: b,c,d



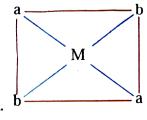
**11.** Which of the following can show geometrical isomers (M=metal,a,b,c-monodentate ligands?

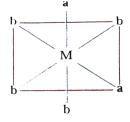
- A.  $Ma_3b$
- B.  $Ma_4b_2$
- C. Ma<sub>2</sub>bc (squareplanar)
- D.  $Ma_2b_2c_2$

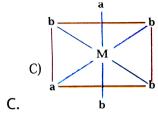
### Answer: b,c,d

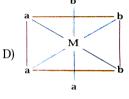


12. Which of the following are cis isomers?









#### Answer: b,c,d

D.

В.



**13.** The following complexs are given?

trans-
$$\left[Co(NH_3)_4 1_2\right]^{\oplus}$$

$$cis - \left[Co(NH_3)_2(en)_2\right]^{3+}$$

$${\rm trans-} \bigg[ {\it Co} \Big( {\it NH}_3 \Big)_2 (en)_2 \bigg]^{3+} \\ \bigg[ {\it NiI}_4 \bigg]^{2-}$$

$$\left[ TiF_6 \right]^2$$

$$[CoF_6]^{3}$$
-
Choose the correct code .

A. (1),(2) are optically active,

B. (2) is optically active, (1), (3) are optically inactive

C. (4),(6) are colourless (5) is coloured

D. (4) is coloured and (5) is colourless

## Answer: b,d



**14.** Among the following which is most stable?

(1) 
$$\left[\operatorname{Fe}(\operatorname{CN})_{6}\right]^{3}$$
 and  $\left[\operatorname{FeF}_{6}\right]^{3}$  (X) (Y)

(2)  $\left[\operatorname{Fe}(\operatorname{CN})_{6}\right]^{3}$  and  $\left[\operatorname{Fe}(\operatorname{CN})_{6}\right]^{4}$  (S) (T)

(3) 
$$[Cr(CN)_6]^{-3}$$
 and  $[Cr(CN)_6]^{-4}$   
(P) (Q)

Choose the correct code

A. 
$$1 - X$$
,  $2 - T$ ,  $3 - Q$ 

#### Answer: c



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15. Which of the following is/are correct?

A. Ligands with filled  $\pi$  orbitals are called  $\pi$  donors

B.  $\pi$  donor ligand forms sigma bond with metal  $t_{2g}$  orbital

C. Ligand to metal charge transfer favoured when the central metal has a high oxidation state

D. Delocalization of  $\pi$  electrons from the ligand to the metal rduces the value of  $\triangle_0$ 

#### Answer: a,d



**16.** The magnitude of crystal field stabilization energy depends on

A. The nature of of ligands

- B. The charge on the metals atom
- C. The charge on the ligand
- D. Position of metal in periodic table

#### Answer: a,b,d



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# 17. Complex ions $\left[NiCl_4\right]^{2}$ , $\left[Ni\left(CN_4\right]^{2}\right]$

Similar in theior given properties:

- A. Oxidation state
- B. Co-Ordination number
- C. Mangnetic moment
- D. Color

## Answer: a,b

18. Select correct statement(s) regarding given complexes:

- A.  $\left[Fe(CO)_5\right]$  the orbitals used for hybridix=zation in Fe atom are s, Px, Py, Pz,  $d_{z^2}$
- B.  $\left[Pt\left(NH_3\right)_2Cl\right]$  the orbital used for hydrization on pt atom are s,Px,Py, $d_{x^2-y^2}$  and it is low spin complex
- C.  $\left[Cr\left(H_2O\right)_6\right]^{3+}$  the orbitals used for hybridiztion in cr atom are s  $P_xP_y$ ,  $P_z$ ,  $D_{z^2}$ ,  $D_{x^2-y^2}$  and it si high spain complex
- D. Ni ( $CO_4$ the orbitals used for hybrization in Ni atom s

 $P_X$ ,  $P_V$ ,  $P_Z$  and it low spin complex

### Answer: a,b,c,d



**19.** Complex compound  $\left[Co(SCN)_2(NH_3)_4\right]Cl$  exhibits:

A. ionzation isomerism

B. Geometrical isomerism

C. optical isomerism

D. Linkage isomerism

#### Answer: a,b,d



**20.** Which of the following compound has/have effective atomic number equal to the atomic number of a noble gas ?

A. 
$$K[Co(CO)_4]$$

$$\mathsf{B.}\, K_2 \Big[ \mathit{Fe}(\mathit{CO})_4 \Big]$$

$$\mathsf{C.}\left[\mathit{Co}\left(\mathit{NH}_{3}\right)_{6}\right]\mathit{Cl}_{2}$$

$$\mathsf{D.}\left[\mathit{CoCl}_{3}\!\!\left(H_{2}O\right)_{\!3}\right]$$

## Answer: a,b,d



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**21.** 
$$K_2 \left[ Ni(CN)_4 \right]^{\text{K in liq.}} \rightarrow NH_3 X'$$

Regarding this reaction correct statement is/are:

A. 
$$'X'$$
 is  $K_4$   $\left[Ni(CN)_4\right]$ 

B. the oxidation state of Ni changed +2 to zero

C. The stucture of 'X'is tertrahedral

D.  $\left[Ni(CN)_4\right]^{2-}$  is a square planar complex

# Answer: a,b,c,d



22. Which one of the following statement(s) is/are false?

A. Weak ligands like  $F^-$  and  $OH^-$  usually form low apin complexes

B. strong liqand like  $CN^-$  and  $NO_2^-$  generally form high spin complexes

- C.  $[FeF_6]^{3-}$  is high spin complex
- D.  $\left[Ni(CO)_4\right]$  is a high spin complex

#### Answer: a,b,d



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23. Which of the following is correct about

Tetraamminedithiocyanato-s cobalt(III) tris(oxalato)cobaltate(III)?

A. Formula of the complex is

$$\Big[ {\it Co(SCN)}_2({\it NH})_3 \Big] \Big[ {\it Co(OX)}_3 \Big]$$

B. It is a chelating complex and show linkage isomerism

C. It shown optical isomerism

D. It shows geometal isomerism

### Answer: b,c,d



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#### 24. Select the correct statement

A. Chelatiojn effect is maximum for five and Six memberd rings

B. Greater the charge on the central metal cation greater ,the

value of  $\Delta(CFSE)$ 

C. In complex ion  $\Delta_{Oct}$  < P(Pairing energy) and it is a low spin

complex

D.  $\left[CoCl_2(NH_3)_2(en)\right]^{\oplus}$  complex ion will have four different isomers

### Answer: a,b



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# **25.** Which of the following obey18e<sup>-</sup>rule?

- A.  $Cr(CO)_6$
- B.  $Ni(CO)_{A}$
- C.  $Cr(C_6H_6)_2$ D.  $Fe(C_5H_5)_2$

# Answer: a,b,c,d

26. Which of the following obey EAN rule?

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

B. 
$$Cr(CO)_6$$

$$C.\left[V(CO)_{6}\right]^{-}$$

D. 
$$[Zn(EDTA)]^{-2}$$

#### Answer: a,b,c,d



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27. Which of the following are four coordinated cpomplexes?

A.  $Be(acac)_2$ 

B.  $[Zn(EDTA)]^2$ 

C.  $[Pt(dien)Cl]^+$  (dien=diethyklene triamine)

D. [Pt(Trien)] +2(triehtylene tetramine)

## Answer: a,c,d



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**28.** In  $Ni^{2+}$  octahedral complex ,which of the following ligands do not pair up the electrous ?

**A.** *NH*<sub>3</sub>

B. *H*<sub>2</sub>*O* 

C. en

D. Dien

Answer: a,b,c,d

#### 29. Explain the following:

- (i) Low spin octahedral complexes of nickel are not known.
- (ii) The  $\pi$  complexes are known for transition elements only.
- (iii) CO is a stroger ligand than  $NH_3$  for many metals.
  - A. Ferrocene
  - B. Zeises salt
  - C. Dibenzenechromimum exhibits optical activity?
  - D. Grignard Reagent

#### Answer: a,b,c



A. 
$$Cis - \left[Co(en)_2Cl_2\right]^+$$

B. Trans - 
$$\left[Co(en)_2Cl_2\right]^+$$

$$C. K_3 \left[ Al \left( C_2 O_4 \right)_3 \right]$$

$$D. \left[ Co \left( NH_3 \right)_4 Cl_2 \right]^+$$

#### Answer: a,c



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# 31. The optically active complex are

A. 
$$\left[As\left(CH_3\right)(Cl)(Br)(I)\right]^+$$

B. 
$$\left[Pt\left(NH_3\right)(Py)(Cl)(Br)\right]$$

C. 
$$\left[ Pt(gly)_2 \right]$$

D. 
$$\left[ Cr(en)_3 \right] Cl_3$$

#### Answer: a,d



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**32.** Which is (are) correct statement (s)?

- A.  $\left[Ag\left(NH_{3}-(2)\right)\right]^{+}$  is linear with sp hybridsed  $AG^{+}$  ions
- B.  $NiCl_4^2$ ,  $CrO_4^2$  and  $MnO_4$  have tetrahedral geometry
- C.  $\left[Cu\left(NH_3\right)_4\right]^{2+}$ ,  $\left[pt\left(NH_3\right)_4\right]^{2+}$  and  $\left[Ni(CN)_4\right]^{2-}$  have  $dsp^2$  hybridisatioi of the metal ion
- ${\rm D.} {\it Fe}({\it CO})_{\rm 5}{\rm has}$  trigonal bipyramidal struure with  $D_{\it Z^2}{\it Sp}^3$  hydridesed ion

#### Answer: a,b,c,d



**33.** which of the following is /are correct relations(s)?

$$A. \Delta_0 = \frac{4}{9} \Delta_t$$

$$\mathsf{B.}\,\Delta_t = \frac{4}{9}\Delta_0$$

$$\mathsf{C.}\ \Delta_s = 1.3\Delta_0$$

D. 
$$\Delta_t = \frac{3}{7}\Delta_0$$

#### Answer: b,c



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**34.** Which of the following is /are correct aboout wikinson's catalst?

A. It is used as homogeneous catalst for selecticve

hydrrogenatuion of organic mole cule at room temparture

and pressure

B. It is tetrahedral complex

C. It does not have unpaired el, ectrons

D. Its formula is 
$$TiCl_4 + Al(C_2H_5)_3$$

Answer: a,c



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## **Comprehension Questions**

**1.** A metal complex having composition  $\left[Cr\left(NH_3\right)_4Br_2I\right]$  was isolated in two forms (X) and (Y). Form (X) reacts with  $AgNO_3$  to give a pale yellow precipitate which is partially soluble in excess of  $NH_4OH$  whereas (Y) gives a greenish yellow precipitate which is insoluble in  $NH_4OH$ .

what is formula of X and Y?

$$\left[Cr\left(NH_3\right)_4BrI\right]Br$$
 and  $\left[Cr\left(NH_3\right)_3Br_2I\right]NH_3$  respectively

B. The formula of (X) and (Y) are 
$$\left[ Cr \left( NH_3 \right)_4 BrI \right] Br \ \& \left[ Cr \left( NH_3 \right)_4 Br_2 \right] I \text{ repectively}$$

C. The formula of (X) and (Y) are both 
$$\left[Cr(NH_3)_4I\right]Br_2$$

D. The formula of (X) and (Y) are 
$$\left[Cr\left(NH_3\right)_2IBr_2\right]\left(NH_3\right)_2$$

#### Answer: b



**2.** A metal complex having composition  $\left[Cr\left(NH_3\right)_4Br_2I\right]$  was isolated in two forms (X) and (Y). Form (X) reacts with  $AgNO_3$  to give a pale yellow precipitate which is partially soluble in excess of  $NH_4OH$  whereas (Y) gives a greenish yellow precipitate which is

insoluble in  $NH_4OH$ .

what is formula of X and Y?

A. linkage isomerism

B. Coordination isomerism

C. Ionization isomerism

D. None of these

#### Answer: c



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**3.** A metal complex having composition  $\left[Cr\left(NH_3\right)_4Br_2I\right]$  was isolated in two forms (X) and (Y). Form (X) reacts with  $AgNO_3$  to give a pale yellow precipitate which is partially soluble in excess of  $NH_4OH$  whereas (Y) gives a greenish yellow precipitate which is

insoluble in  $NH_4OH$ .

Which of the following statement is true?

A. (X)-cis form optically inactive (Y)-cis form optically active

B. (X)-cis form optically inactive (Y)- trans form optically active

C. The cis and trans forms of both X and Y are optically active

D. The cis and trans form of both X and Y are optically inactive.

#### Answer: d



**Matrix Matching Questions** 



the Column - II

following

a) 229

(molar conductance)

 $p) \left[ Pt(NH_3)_5 C1 \right] Cl_3$ 

**b**) 0

q)  $\left[ Pt(NH_3)_2 Cl_4 \right]$ 

c) 404

 $r) \left[ Pt(NH_3)_4 Cl_2 \right] Cl_2$ 

d) 523

s)  $[Pt(NH_3)_6]Cl_4$ 

t)  $\left[ Co(NH_3)_3 Cl_3 \right]$ 



2.

Match Column - I

Column - II

p) sp<sup>3</sup> hybridized

s) digmagnetic

r) outer orbital complex

the

following

following

a) K<sub>2</sub>[NiCl<sub>4</sub>]

b) [Ni(CO),]

q) para magnetic c)  $\left[ \text{Cr}(\text{NH}_3)_6 \right]^{3+}$ 

d)  $\left[ \text{FeF}_6 \right]^{3-}$ 

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3.

Column - I

Match

Column - II

the

a)  $\lceil \text{Fe}(\text{CN})_6 \rceil^{4-}$  p) Paramagnetic,  $\text{sp}^3 \text{d}^2$ 

b)  $\left[ \text{Fe} \left( \text{H}_2 \text{O} \right)_6 \right]^{2+}$  q) Diamagnetic,  $d^2 \text{sp}^3$ 

c)  $\lceil \text{CrCl}_2 \left( \text{NH}_3 \right)_4 \rceil \text{NO}_3 \text{ r} \rangle$  Diamagnetic  $\mathbf{sp}^3 \mathbf{d}^2$ 

d)  $\left[ Zn(H_2O)_6 \right]^{2+}$  s) Ionization isomerism

the 4. Match following

Column - II Column - I

a)  $\left[ \text{Ni}(\text{NH}_3)_6 \right]^{2+}$ p) Diagmagnetic

b)  $\left[ Ni(CO)_4 \right]$ q)  $sp^3 d^2$ 

c)  $\left[ Cr(H_2O)_6 \right]^{3+}$ r)  $d^2sp^3$ 

d)  $\left[ Ni(CN)_4 \right]^{2-}$ s) Inner orbital complex

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d)  $\left[ Cr(H_2O)_6 \right] Cl_3$ **View Text Solution** 

List-I

b)  $\left[ Pt(NH_3)_2 Cl_2 \right]$ 

c)  $\left[ Cr(H_2O)_5 Cl \right] Cl$ 

5.

6.

Column - I

Match List-II (A)  $\left[CoF_6\right]^{3-}$ (p) High spin complex (B)  $\left[ Fe(H_2O)_6 \right]^{3+}$  (q) Low spin complex

Match

a)  $\lceil \text{Co}(\text{NH}_3)_4 (\text{H}_2\text{O})_2 \rceil \text{Cl}_2$ 

s) optical isomers

the

Column - II

isomers

p) Geometrical

q) Diamagnetic

r) Paramagnetic

the

following

following

(s) Inner orbital complex (t) Paramagnetic

(C)  $\lceil Co(en)_3 \rceil^{3+}$  (r) Outer orbital complex

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(D)  $\left[ Cr(CO)_{6} \right]$ 

7. Match following the List-II List-I (p) Dipole Moment (A)  $\left\lceil Pt(NH_3)_2 Cl_2 \right\rceil$ (B)  $K_3 \left[ Cr(OX)_3 \right]$ (q) Polarimeter (C)  $\lceil Co(NH_3) \choose A Cl_2 \rceil Cl$ (r) Precipitation with  $AgNO_{3}$ (D)  $\lceil Co(en)_2 Cl_2 \rceil Cl$ (s) Cation exchange Resin (t) Anion exchange Resin



8. Match the following List-I List-II (A) Wilkinson catalyst (p) +1 O.S.(B) deoxy haemoglobin (q) + 2 O.S.(C) Brown Ring complex (r) Coordination number 4 (D) Vaskas complex (s)Coordination number 6 (t) Neutral complex



9. Match the following Column -I Column-II A) [Co(NH<sub>3</sub>)<sub>4</sub>Cl<sub>2</sub>] p) Optical isomerism B) [Co(en)<sub>3</sub>]Cl<sub>2</sub> q) Ionization isomerism C) [Co(en)<sub>2</sub>(NO<sub>2</sub>)Cl]SCN r) Coordination isomerism D)  $[Co(NH_3)_6][Cr(CN)_6]$ s) Geometrical isomerism



10. Match the following

Column - I Pair of complex

Column - II Property is different among the given pair

p) Magnetic moment

 $K_2[Ni(CN)_4]$ 

A)  $\lceil Ni(CO)_4 \rceil \&$ 

B)  $\left[ \text{Cu}(\text{NH}_3)_4 \right] \text{SO}_4 \&$ 

 $K_3 \lceil Cu(CN)_4 \rceil$ 

C)  $K_2[NiCl_4]$ &

 $K_2[Ni(CN)_4]$ 

D) K<sub>2</sub>[NiCl<sub>4</sub>] & K<sub>2</sub>[PtCl<sub>4</sub>] q) Oxidation no of central

metal

r) Geometry

s) EAN of central atom

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## 11. Match the following

Column - I

Column - H

A)  $MnCl_4^{2-}$ 

p) Square planar

B) NiCl<sub>4</sub>

q) Tetrahedral

C)  $Cs_2(CuCl_4)$ 

r) Diamagnetic

D)  $Ni(CN)_4^{2}$ 

s) Paramagnetic



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#### 12. Match the following

Column - I

Column - I

- a)  $\left[ \text{Fe}(\text{CN})_{6} \right]^{4-}$
- b)  $\left[ \text{Fe}(H_2O)_6 \right]^{2+}$
- c)  $\lceil \text{Co}(\text{NH}_3)_6 \rceil^{3+}$
- d)  $\left[\operatorname{CoF}_{6}\right]^{3-}$

Column - II

- p) Paramagnetic
- q) Diamagnetic
- t) octahedral
- r) Inner orbital complex s) Outer orbital complex

13. Match the following: Complex

Complex

(A) 
$$\left[Pt(ghy)_{2}\right]$$
 (B)  $\left[Co(ghy)_{3}\right]$  (C)  $\left[Pt(en)_{2}\right]^{2+}$  (D)  $\left[Co(en)_{3}\right]^{3+}$ 



### 14. Match the following:

(A)  $Be(acac)_2$ (B)  $\left[ Zn(EDTA) \right]^{-2}$ (q) Square planar

(p) Octahedral

- (C)  $K_3 \left[ Al \left( C_2 O_4 \right)_3 \right]$ (r) Tetrahedral
- (D)  $Ni(DMG)_2$ (s) Diamagnetic (t) Exhibits optical activity



#### 15. Match the following:

List-I

A) deoxyhaemoglobin

(B) Wilkinson catalyst

(C) Sodium nitroprusside

(D) Reineck salt

List-II

- (p) Oxidation state of the metal is +2
- (q) Oxidation state of the metal is +1
- (r) Oxidation state of the metal is +3
- (s) Coordination number: 4
- (t) Coordination number: 6
  - A. deosyhaemoglobin
  - B. Wilkinson catalyst
  - C. Sodium nitroprusside
  - D. Reineck salt

Answer: A-p,t B-q,s C-p,t D-r,t



16. Match the following

I

(1) 
$$\left[ Co(NH_3)_6 \right]^{3+}$$
 (A) 5.87 BM

(2) 
$$\left[Fe(CN)_{6}\right]^{3-}$$
 (B) diamagnetic

(3) 
$$[Mn(H_2O)_6]^{2+}$$
 (C) 1.73 BM

III

1) (P) 
$$sp^3d^2$$
 (U) Outer orbital complex

2) (Q) 
$$d^2sp^3$$
 (V) Inner orbital complex

3) (R) 
$$dsp^3$$
 (W) Having only  $\sigma$  donation from the ligand

A. 1: B,Q,V,W 2:C,Q,V 3:A,P,U,W

B. 1: A,Q,V,W 2:B,P,V,W 3:B,Q,U,W

C. 1: B,Q,V,W 2:A,P,U,W 3:C,R,V

D. 1: A,P,U 2:C,R,U,W 3:A,Q,V

#### 17. Match the following

Complex	I
$(1)  \left[ Pt(CN)_4 \right]^{2-}$	(A) 2.83 BM
(2) $[NiCl_4]^{2-}$	(B) 5.87 BM
$[MnBr_4]^{2-}$	(C) 3.87 BM
4) $\left[CoCl_4\right]^{-2}$	(D) Diamagnetic
II	III
1) (P) $sp^3$	(U) $d^8$ (Low spin)
2) Q) dsp <sup>2</sup>	(V) $d^5$
3) (R) Tetrahedral	(W) $d^7$
(4) (S) Square planar	(X) $d^8$ (high spin)

A. 1: A,P,R,W 2:D,Q,S,U 3:D,Q,S,W 4: A,Q,S,X

B. 1: C,Q,S,W 2:B,P,R,W 3:A,P,R,V 4:B,Q,S,W

C. 1: B,P,R,V 2:C,Q,S,V 3:C,Q,S,X 4:D,P,R,V

D. 1: D,Q,S,U 2:A,P,R,X 3:B,P,R,V 4:C,P,R,W

#### Answer: D



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18. Match the following

List-I

List-II

(A)  $\lceil Co(en)_2 Cl_2 \rceil Cl$  (p) Geometrical

Isomerism

(B)  $\lceil Co(en)_3 \rceil Cl_3$  (q)Optical isomerism

(C)  $\lceil Cr(NH_3)_4 Cl_2 \rceil Cl$  (r) Paramagnetic

(D)  $K_3 \lceil Cr(C_2O_4)_3 \rceil$ 

(s) Diamagnetic

(t) Cationic complex

A. 1: A,P,R,W 2:D,O,S,U 3:D,O,S,W 4: A,O,S,X

B. 1: C,O,S,W 2:B,P,R,W 3:A,P,R,W 4: B,O,S,W

C. 1: B,P,R,V 2:C,Q,S,V 3:C,Q,S,X 4: D,P,R,V

D. 1: D,Q,S,U 2:A,P,R,X 3:B,P,R,V 4: C,P,R,W

Answer: A-p,q,s,t B-q,s,t C-p,r,t D-q,r

19. Match the following

Column - I Column - II

A) 
$$\left[ Cr(gly)_3 \right]$$
 p) Low spin c

A)  $\left[ \text{Cr}(\text{gly})_3 \right]$  p) Low spin complex B)  $\left[ \text{Co}(\text{Br})_2 (\text{CI})_2 (\text{SCN})_2 \right]^{3-}$  q) High spin complex

$$\mu = 4.84 \text{ B.M}$$

C) 
$$\left[ \text{Co}(\text{NH}_3)_6 \right]^{3+}$$
 r) Optical isomerism

D) Na[Pt(Br)(Cl)(NO<sub>2</sub>)(NH<sub>3</sub>)]s) Geometrical isomerism t) Linkage isomerism



# **Statement Type**

**1.** Statement-1:  $Co(CO)_4$  is not stable but  $\left[Co(CO)_4\right]^-$  is stable.

Statement-2:  $\left[Co(CO)_4\right]^{-1}$  obeys EAN rule

A. Statement-1 is True, Statement-2 is True,

Statement-2 is a correct explanation for Statement-1.

B. Statement-1 is True, Statement-2 is True,

Statement-2 is NOTa correct explanation for Statement-1.

C. Statement-1 is True, Statement-2 is False.

D. Statement-1 is False, Statement-2 is True.

#### Answer: a



**2.** Statement-1:  $\left[W(CO)_6\right]$  obeys EAN rule as it attains xe configuration.

Statement-2: It is stable because it attains Rn configuration

A. Statement-1 is True, Statement-2 is True,

Statement-2 is a correct explanation for Statement-1.

B. Statement-1 is True, Statement-2 is True,

Statement-2 is NOTa correct explanation for Statement-1.

C. Statement-1 is True, Statement-2 is False.

D. Statement-1 is False, Statement-2 is True.

## Answer: d



- 3. The questions given below consist of Assertion (A) and Reason (R)
- . Use the following key to select the correct answer.

Assertion : The complex  $\left[ \mathit{Ni(en)}_{3} \right] \mathit{Cl}_{2}$  has lower stability than

$$\left[Ni\left(NH_3\right)_6\right]Cl_2$$
.

Reason : In  $\left\lceil Ni(en)_3 \right\rceil Cl_2$ , the geometry of Ni is triagonal bipyramidal

A. Statement-1 is True, Statement-2 is True,

Statement-2 is a correct explanation for Statement-1.

B. Statement-1 is True, Statement-2 is True,

Statement-2 is NOTa correct explanation for Statement-1.

C. Statement-1 is True, Statement-2 is False.

D. Statement-1 is False, Statement-2 is True.

## Answer: d



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4. 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.

- A. Statement-1 is true, Statement-2 is True, Statement-2 is a correct explanation for Statement-1
- B. Statement- is True, Statement-2 is True, Statement-2 is not a correct explanantion for Statement-1
- C. Statement-1 is True, Statement-2 is False
- D. Statement-1 is False, Statement-2 is True.

#### **Answer: C**



**5.** The magnetic moment of  $\left[Ni(Cl)_4\right]^{2-}$  is

correct explanation for Statement-2

A. Statement-1 is true, Statement-2 is True, Statement-2 is a

- B. Statement- is True, Statement-2 is True, Statement-2 is not a
- C. Statement-1 is True, Statement-2 is False

correct explanantion for Statement-2

D. Statement-1 is False, Statement-2 is True.

## **Answer: D**



- **6.**  $Fe^{3+}$  compounds are more stable than  $Fe^{2+}$  compounds because
  - A. Statement-1 is true, Statement-2 is True, Statement-2 is a correct explanation for Statement-3
    - B. Statement- is True, Statement-2 is True, Statement-2 is not a
      - correct explanantion for Statement-3
    - C. Statement-1 is True, Statement-2 is False

D. Statement-1 is False, Statement-2 is True.

#### **Answer: A**



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- 7. The shape of  $\left[Co(NH_3)_6\right]^{3+}$  is
  - A. Statement-1 is true, Statement-2 is True, Statement-2 is a
  - B. Statement- is True, Statement-2 is True, Statement-2 is not a
    - correct explanantion for Statement-4

correct explanation for Statement-4

- C. Statement-1 is True, Statement-2 is False
- D. Statement-1 is False, Statement-2 is True.

## **Answer: A**



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**8.** The percentage of  $Mg^{2+}$  ions in a solution can be tested by adding a solution of

A. Statement-1 is true, Statement-2 is True, Statement-2 is a correct explanation for Statement-5

B. Statement- is True, Statement-2 is True, Statement-2 is not a correct explanantion for Statement-5

C. Statement-1 is True, Statement-2 is False

D. Statement-1 is False, Statement-2 is True.

## **Answer: C**



**9.** Square planar complexes are formed by  $d^8$  ions with strong field ligands. The crystal field splitting  $\Delta_0$  is larger for the second and theird row transition elements and for more highly charged species. All the complexes having  $4d^8$  and  $5d^8$  configurations are mostly square planar including those with weak field ligands such as halide ions square planar complexes can show geometrical isomerism but they do not show optical isomerism due to the presence of plane of symmetry

Which of these is square planar complex?

(a) All are square planar complexes (b) 
$$\left[Ni(CO)_4\right]$$
 (c)  $\left[Pt\left(NH_3\right)_4\right]^{2+}$  (d)  $\left[Ni\left(NH_3\right)_6\right]^{2+}$ .

A. Statement-1 is true, Statement-2 is True, Statement-2 is a correct explanation for Statement-6

B. Statement- is True, Statement-2 is True, Statement-2 is not a correct explanantion for Statement-6

C. Statement-1 is True, Statement-2 is False

D. Statement-1 is False, Statement-2 is True.

## Answer: A



# **Integer Type Questions**

**1.** The secondary valency in  $[Co(EDTA)]^{2-}$  is

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**2.** How many of the following are diamagnetic?  $\left[ Ag(NH_3)_2 \right]^+, \left[ Cd(NH_3)_4 \right]^{2+},$ 

$$Cr(CO)_6$$
,  $\left[Ni\left(NH_3\right)_6\right]^{2+}$ ,  $\left[Ni(CN)_4\right]^{2-}$ 

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**3.** The number of geometrical isomers and optical isomers of  $\Big[Pt\Big(NH_3\Big)(Br)(Cl)(Py)\Big]$  are respectively.



**4.** The number of oxygen atoms involved in bonding in the coordination sphere of  $[Mq(EDTA)]^{2-}$  is



**5.** The number of unpaired spins in the  $\left[Cr(en)_3\right]^{2+}$  ion is



- 6. How many geometrical isomers are possible for
- $\left[Pt\left(NO_2\right)\left(NH_2OH\right)(Py)\right]^+?$ 
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7. How many steroisomers are possible for

$$\left[Pt(Br)(Cl)\left(NH_3\right)_2\right]?$$



- **8.** Dimethyl glyoxime forms a square planar conplex with  $Ni^{+2}$ . The number of unpaired electrons are
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9. How many number of d-electrons are present in  $\left[ Co \left( NH_3 \right)_5 CO_3 \right] ClO_4 ?$ 



**10.** The number of oxygen atoms coordinated to Cr in Cr (acac)<sub>3</sub> is ?



**11.** Al  $(acac)_X$  the 'X' is

Note: acac is univalent bidentate ligand



**12.** In  $Zn (oxine)_2$  (oxine : 8- quinolinol), the coordination number and the oxidation state of Zn are x and Y respectively, then (x+y) is

**13.**  $\left[ Fe^{3+} (biphy)_X \right]$ , (bipy = 2,2' bipyridine)'X' value is



**14.** Choose the incorrect regarding stability

(a) 
$$\left[ Fe(\text{phen})_3 \right]^{+2} > \left[ Fe(\text{phen})_3 \right]^{+3}$$

(b) 
$$\left[ Fe(CN)_6 \right]^{-4} > \left[ Fe(CN)_6 \right]^{-3}$$

(c) 
$$Fe(\text{dipy})_3^{+3} > Fe(\text{dipy})_3^{+2}$$



**15.** The number of  $t_{2g}$  electrons in  $\left[Fe(H_2O)_6\right]^{3+}$  is



**16.** Which of the following compound statement is correct for

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**17.** How many of the following are  $sp^3$  hybridized?

$$\left[ZnCl_4\right]^{2-}$$
,  $Ni(DMG)_2$ ,  $\left[Ni(CN)_4\right]^{2-}$ ,  $\left[Ni(CO)_4\right]$ 

$$[NiCl_4]^{2-}$$
,  $[PtCl_4]^{2-}$ ,  $[Pd(NH_3)_4]^{2+}$ ,  $[FeCl_4]^{-1}$ 



$$\begin{bmatrix} y_1(y_1) \end{bmatrix}_{2+} \begin{bmatrix} g_1(y_1) \end{bmatrix}_{3+} \begin{bmatrix} g_1(g_1) \end{bmatrix}_{4}$$

**18.** How many of the following have  $d^2sp^3$  hybridization?

$$\left[Ni\left(NH_3\right)_6\right]^{2+}, \left[Co\left(NH_3\right)_6\right]^{3+}, \left[Fe(CN)_6\right]^{4-}$$

$$\left[Cr(NH_3)_6\right]^{3+}, \left[Co(en)_3\right]^{3+}, \left[Co(EDTA)\right]^{2-}$$

$$[CoF_6]^{3-}, [Co(H_2O)_6]^{2+}$$

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<b>19.</b> The hybridization of the d-block element sodium nitropruside is
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**20.** The number of unpaired electrons in the brown ring complex is

**21.** the number of five membered chelate rings in  $[Zn(EDTA)]^{2-}$  is



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**22.** DMG is x-valen and y-dentate ligand. Taking x and y as numbers, find (x+y)



**23.** How many geometrical isomers and stereoisomers are possible for

for 
$$\left[ Pt \left( NO_2 \right) \left( NH_3 \right) \left( NH_2OH \right) (Py) \right]^+ \text{ and } \left[ Pt (Br) (Cl) (I) \left( NO_2 \right) \left( NH_3 \right) (Py) \right]$$



respectively?

**24.** How many molecules of benzene  $\left(C_6H_6\right)$  are there in 1L of benzene? Specific gravity of benzene is 0.88.



25. The oxidation number of the d-block metal in Zeises salt is



**26.** The oxidation stat of the metal in Reinecks salt is +x, x is



**27.** The coordination number of the metal in the Wilkinson catalyst is +xx is



**28.** The oxidation number of the metal in the Wilkinson catalyst is +x

x is,



**29.** The complexes  $\left[ Co \left( NH_3 \right)_6 \right] \left[ Cr(CN)_6 \right]$  and

 $\left[ \mathit{Cr} \left( \mathit{NH}_3 \right)_6 \right] \left[ \mathit{Co} \left( \mathit{CN} \right)_6 \right]$  are the examples of which type of isomerism ?



**30.** In the complex  $\left[Pt(NH_3)_3Cl\right]\left[CuCl_3(NH_3)\right]$  the charges over

$$\left[Pt\left(NH_3\right)_3Cl\right]^{+x}$$
 and  $\left[CuCl_3\left(NH_3\right)^{-y}\right]^{-y}$  then (x+y) is



**31.** The possible number of stereoismers for the formula  $\left[ Ma_2b_2cd \right]^{\pm n}.$ 



**32.** The number of nitrogen atoms coordunated in  $[Zn(EDTA)]^{-2}$  is



**33.** The number of five membered rings in  $[Pt(trien)]^{2+}$  (trien=triethyleneetetra amine) is



**34.** The oxidation number of metal atom is zero in



**35.** The oxidation state amnd coordination number of the metal in Wilkinson catalyst are +x and y respectively. (x+y) is



36. The formula of sodium nitroprusside is



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37. Intermolecular hydrogen bonding strongest in:



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Which of the 38. following isomerism, exhibited by

$$\left[ CrCl_2(OH)_2 \left( NH_3 \right)_2 \right]^{-}?$$



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39. The total possible co-ordination isomers for the following compounds respectively are

- (i)  $\left[Co(en)_3\right]\left[Cr\left(C_2O_4\right)_3\right]$
- (ii)  $\left[ Cu(NH_3)_4 \right] \left[ CuCl_4 \right]$
- (iii)  $\left[Ni(en)_3\right]\left[Co\left(NO_2\right)_6\right]$ 
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**40.** A metal complex of coordination number six having three different types of ligands a, b and c of composition  $Ma_2b_2c_2$  can exist in several geometrical isomeric forms, the total number of such isomers is:



- **41.** The crystel field stabalisation energy of  $Mn(H_2O)_6^{2+}$  is
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**42.** If  $\Delta_0 < P$  then electronic arrangement of metal atom/ion in an octahederal complex with  $d^4$  configuration is  $t_{2g}^x e_g^y$ . What is the value of x



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43. The no. of complexes showing and mer isomerism

(A) 
$$\left[ Co(NH_3)_3 Cl_3 \right]$$
 (B)  $\left[ Co(en)_3 \right]^{+3}$ 

$$\left[Co(gly)_3\right]$$

(D) 
$$\left[ Co\left(NH_3\right)_2 \left(H_2O\right)_2 Cl_2 \right]$$

(E) 
$$\left[ Fe(dipy)_3 \right]^{+2}$$
 (F)  $\left[ Fe(o - phen)_3 \right]^{+2}$ 



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**44.** The no. of comlexes showing hepticity 5

A. ferrocene B. zeise's salt C. ruthenocene D. chromocene Answer: 2 **Watch Video Solution** 45. The no. of chelating ligands are A. Sulphato B. EDTA C. nitrato D. thiosulphato **Answer: 1** 



**46.** In complex  $\left[Cr(NH_3)_6 \left[Co(C_2O_4)_3\right]\right]$  the total No. of ions formed are



**47.** In the complex  $\left[Fe(CO)_2(NO)_2\right]$  the oxidation state of Fe is -x,'x' is



**48.** In the coordination compound,  $K_4\Big[Ni(CN)_4\Big]$  oxidation state of nickel is



49. The no of statements is / are correct?



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**50.** Consider the following complex:

$$\left[Co\left(NH_3\right)_5CO_3\right]ClO_4$$

the coordination number is X and oxidation number is Y, number of d-electrons is Z, and number of unpaired d-electrons of the metal is A are respectively. (X+Y)-(Z+A).



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**51.** The CFSE for  $\left[CoCl_4\right]^{4-}$  is  $18000cm^{-1}$  the value of y ios



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**52.** Amongest  $\left[Co(OX)_3\right]^{3-}$ ,  $\left[CoF_6\right]^{3-}$ ,  $\left[Co(NH_3)_6\right]^{3+}$  and

$$\left[Co(NH_3)_6\right]^{2+}$$
 the no. of coloured complexes



**53.** The theoritical spin-only magnetic moment of cobalt in  $Hg[Co(SCN)_4]$  is 3.87 B.M.



**54.** From  $Ma_3b_3$ , when one b is replaced by c, the total no. of geometrical isomer possible are



- **1.** The number of unpaired electrons in  $\left[Cr(NH_3)_6\right]^{2+}$  is
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- **2.** The hybridization of  $\left[Cr(H_2O)_6\right]^{2+}$  is \_\_\_\_\_\_.
  - Watch Video Solution

- 3. The total number of geometric isomers for Mabcdef are
  - Watch Video Solution

<b>4.</b> $\left[Pt\left(NH_3\right)_5Cl\right]^+$ does not exhibit and				
stereio isomersim .				
Watch Video Solution				
5. The EAN of $Mo(CO)_6$ is  Watch Video Solution				
<b>6.</b> The coordination number of Pt in cisplatin is				
Watch Video Solution				
SINGLE ANSWER QUESTIONS				

1. In Ziesses salt C=C bond length is:

Note: 
$$\left\{ \begin{array}{l} C - C \text{bond length in enthance is 1.54Å} \\ C = C \text{bond length in enthance is 1.34Å} \\ C \equiv C \text{bond length in enthance is 1.20Å} \end{array} \right\}$$

- **A.** 1.37*A* °
- B. 1.19A°
- C. 1.87*A* °
- D. 1.34A °

## Answer: a



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2. Which one of the following can get oxidized easily?

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

B. 
$$\left[Co(NH_3)_6\right]^{2+}$$

$$\mathsf{C}.\mathit{Fe}(\mathit{Co})_5$$

# Answer: b



3.

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magentic momentsn(BM) of

the complexes

 $\left[Ni\left(H_2O\right)_6\right]^{2+}$  and  $\left[Ni\left(NH_3\right)_6\right]^{2+}$  are resectively

The

A. 1.73, 2.84

B. 0, 2.84

C. 2.84, 0

D. 2.84, 2.84

# Answer: d

**4.** The number of electron in  $t_{2g}$  orbitals in  $K_4 \lceil Fe(CN)_6 \rceil$  is

**5.** The crystel field stabalisation energy of  $\left[Mn(H_2O)_6\right]^{2+}$  is

- A. 2
- B. 3
- C. 6
- D. 5

## Answer: c



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A.  $-1.2 \triangle_{\alpha}$ 

B. - 1.9 
$$\triangle_{\theta}$$

C. 0

D. -2.4 △<sub>0</sub>

## Answer: c



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# The theoretical percentage wight loss 6.

when

 $\left[Cr\left(H_2O\right)_4Cl_2\right]Cl.2H_2O$  conc.  $H_2SO_4$  in a closed desicator is

A. 13.5

C. 18.0

B. 6.75

D. 36.0

Answer: a

**7.** 0.2435 g of a complex gave 0.2870 g of AgCl when treated with a excess  $AgNO_3$  solution. The complex is

A. 
$$\left[ Cr(NH_3)_4 Cl_2 \right] Cl$$

$$B. \left[ Cr \left( NH_3 \right)_5 Cl_2 \right] Cl_2$$

$$C. \left[ Cr \left( NH_3 \right)_3 Cl_3 \right]$$

D. 
$$\left[Cr\left(NH_3\right)_6Cl_2\right]Cl_3$$

## Answer: b



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- A. pentacyanocarbonylferrate (II)
- B. carbonylpentacyanidoferrate (II)
- C. pentacyanocarbonyliron (III)
- D. carbonylpentacyanoiron (III)

## Answer: b



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- 9. The IUPAC name of oerrocene is
  - A. bis ( $\mu^5$  cyclopentadienyl) iron (O) )
  - B. bis( $\mu^{10}$ -cyclopentadienyl)iron(O))
  - C. bis( $\mu^5$ -cyclopentadienyl)iron(II))
  - D. bis( $\mu^{10}$ -cyclopentadienyl)iron(II))

## Answer: c

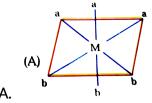
## **10.** Cis di- $\mu$ -chloro bis [chloro(triphenyl phosphine) platinum(II)] is

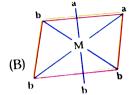
## Answer: b



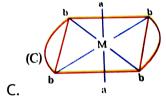
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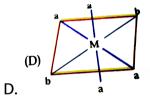
11. Which one of the following represents a Cis isomer?





В.





## Answer: b



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12. The total number of geometric isomers for Mabcdef are

**A.** 4C<sub>2</sub>

2

 $B.5C_1$ 

**c**. 3*C*<sub>2</sub>

D. 6*C*<sub>2</sub>

# Answer: d



**13.** The complex formed by  $Pt^{2+}$  with diethylenetriamine (dien) and chlorido ligand is given by

 $B. \left[ Pt(dien)Cl_2 \right]^{2+}$ 

$$C. [Pt(dien)Cl)]^{1+}$$

D.  $[Pt(dien)Cl]^{2+}$ 

### Answer: c



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14. Which one of the following given an acid whenpassed through cation exchange resin( $RSO_3H$ )?

A. 
$$K_4[Fe(CN)_6]$$

$$C. \left[ CO(NH_3)_6 \right] Cl_3$$

 $B. K_3 \left[ Al \left( C_2 O_4 \right)_3 \right]$ 

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

## Answer: c



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**15.** Which one of the following give a base when passed through anion exchange resin  $\left[R_4N^+\right]OH^-$ ?

A. 
$$K_4$$
 [ $Fe(CN)_6$ ]

$$B. \left[ {\it Co(en)}_3 \right] {\it Cl}_3$$

$$C. \left[ CO(NH_3)_6 \right] Cl_3$$

D. 
$$\left[ Fe \left( H_2 O \right)_6 \right] SO_4$$

#### Answer: a



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**16.** Which one of th efollowing does not form a chelate with oxalate ion by reacting in aqueous solution ?

A. cis-
$$\left[Co\left(NH_3\right)_4Br_2\right]^{+1}$$

B. cis- $\left[Co\left(NH_3\right)_4Cl_2\right]^{+1}$ 

C. cis- $\left[Co(en)_2Cl_2\right]^{+1}$ 

D. trans- $\left[ Co(en)_2 Cl_2 \right]^{+1}$ 

### Answer: d



17. The neutral and positive charged species that can coordinate to a metal atom is

A. Cl

 $B.NH_3$ 

C. Co

D. NO

#### Answer: d



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**18.** Which of the following pair of compounds is expected to exhibit same colour in aqueous solution?

- A. FeCl<sub>2</sub>, CuCl<sub>2</sub>
- B. VOCl<sub>2</sub>, CuCl<sub>2</sub>
- C. VoCl<sub>2</sub>, FeCl<sub>2</sub>
- $D. FeCl_2, MnCl_2$

### Answer: b



19. The oxidation state of Mo in its oxo-complex species

$$\left[Mo_2O_4(C_2H_4)_2(H_2O)_2\right]^{2-}$$
 is:

- A. 2
- B. 3
- C. 4
- D. 5

### Answer: b



- **20.** The donor sites of  $(EDTA)^{4-}$  are ?
  - A. o atoms only
  - B. N atoms only

- C. Two N atoms and four O atoms
- D. Three N atoms and three O atoms

#### Answer: c



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- **21.** In nitroprusside ion the iron and NO exist as Fe (II) and  $NO^+$  rather than the Fe(III) and NO. these forms can be differentiated by
  - A. Estimating the concentration of iron
  - B. Measuring the cancentration of CN-
  - C. Measuring the solid state magnetic moment
  - D. Thermally decomposing the compound

#### Answer: c



**22.** In the Iso-electronic series of metal carbonyls, the C-O bond strength is expected to increase in the order:-

A. 
$$\left[ Mn(CO)_6 \right]^+ < \left[ Cr(CO)_6 \right] < \left[ V(CO)_6 \right]^-$$

$$\mathsf{B.}\left[\mathit{V(CO)}_{6}\right]^{-} < \left[\mathit{Cr(CO)}_{6}\right] < \left[\mathit{Me(CO)}_{6}\right]^{+}$$

$$\mathsf{C.}\left[\mathit{V(CO)}_{6}\right]^{-} < \left[\mathit{Mn(CO)}_{6}\right]^{+} < \left[\mathit{Cr(CO)}_{6}\right]$$

$$D.\left[Cr(CO)_{6}\right] < \left[Mn(CO)_{6}\right]^{+} < \left[V(CO)_{6}\right]^{-}$$

### Answer: b



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23. The magnetic moment of complex given below are in the order:

- (I)  $\left[Ni(CO)_4\right]$
- (II)  $\left[Mn(CN)_6\right]^{4}$

(III) 
$$\left[ Cr \left( NH_3 \right)_6 \right]^{3+}$$

(IV) 
$$\left[CoF_6\right]^{3}$$

- A. I gtll gtlll gtlV
- B. I Itil Itili ItiV
- C. IV gt II gtI gtIII
- D. IV It II ItI ItIII

#### Answer: b



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24. Three arrangements are shown for the complexes

$$\left[ CoBr_2(NH_3)_2(en) \right]^{\oplus}$$
. Which one is wrong statement?

$$\begin{bmatrix} B_{1} & B_{1} & B_{1} & B_{1} & B_{2} & B_{3} & B_{4} & B_{5} & B_$$

- A. I and II are geometrical isomer
- B. II and III are optical isomer
- C. I and III are optical isomers
- D. II and III are geometrical isomers

### Answer: b



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**25.** Among the following, which is not the  $\pi$ -bonded organometallic compound

A. 
$$(CH_3)_4$$
Sn

B. 
$$Cr(n^6 - C_6H_6)_2$$

$$\mathsf{C.} \, Fe \Big( n^5 - C_5 H_5 \Big)_2$$

D. 
$$K[PtCl_3(n^2 - C_2H_4)]$$



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**26.** Complex compounds(s) which is optical active and does ot dpeend upon the orientation of the ligands around metal cation:

(i) 
$$\left[ CoCl_3 \left( NH_3 \right)_3 \right]$$

(ii) 
$$\left[Co(en)_3\right]Cl_3$$

(iii) 
$$\left[Co\left(C_2O_4\right)_2\left(NH_3\right)_2\right]^{-1}$$

(iv) 
$$\left[ CrCl_2 \left( NH_3 \right)_2 (en) \right]^+$$

A. II,IIIand Iv

B. I, liand Iv

C. II and Iv

D. Only II

### Answer: d



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**27.** The total possible co-ordination isomers for the following compounds respectively are

(i) 
$$\left[Co(en)_3\right]\left[Cr\left(C_2O_4\right)_3\right]$$

(ii) 
$$\left[ Cu \left( NH_3 \right)_4 \right] \left[ CuCl_4 \right]$$

(iii) 
$$\left[Ni(en)_3\right]\left[Co\left(NO_2\right)_6\right]$$

A. 4,4,4

B. 2,2,2

C. 2,2,4

D. 4,2,4

28. Which of the following can show geometrical isomerism?

A. 
$$\left[ Pt \left( NH_2 - CH \mid CH_3 - CH \mid CH_3 - NH_2 \right)_2 \right]^{2+}$$

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

C. Both a and B

D. None if these

#### Answer: c



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29. The complex which does not obey EAN rule is

A. 
$$\left[ Fe(CO)_5 \right]^0$$

$$B. \left[ Cr(CO)_3(NO)_2 \right]^0$$

$$\mathsf{C.}\left[\mathit{Fe}(\mathit{CO})_{3}(\mathit{NO})_{2}\right]^{0}$$

D. 
$$\left[ Fe(CO)_2(NO)_2 \right]^0$$

### Answer: c



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# Compreshession type question

1. 
$$(I)K_3[Co(CN)_6] + [Cr(NH_3)_6]Cl_3 \xrightarrow{\text{acetone}} x + 3KCl$$

$$K_3[Cr(CN)_6] + \left[Co(NH_3)_6\right] Cl \rightarrow y + 3KCl$$

the solution was filted x and y are obtained after evoporation of the

solvent from the filtrate.

`X and y are

A. Linkage isomers

B. Geometrical isomerism

C. Coordinaate Isomers

D. Positinal IsoMers.

#### Answer: c



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2. 
$$(I)K_3[Co(CN)_6] + [Cr(NH_3)_6]Cl_3 \xrightarrow{\text{acetone}} x + 3KCl$$

$$K_3[Cr(CN)_6] + [Co(NH_3)_6]Cl \rightarrow y + 3KCl$$

the solution was filted x and y are obtained after evoporation of the solvent from the filtrate.

when x is sent through cation exxhange resin the complex coming out in solution is

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

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

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

D. 
$$\left[ Cr \left( CN_3 \right)_6 \right]^{3}$$

### Answer: c



3. 
$$(I)K_3[Co(CN)_6] + [Cr(NH_3)_6]Cl_3 \xrightarrow{\text{acetone}} x + 3KCl$$

$$K_3[Cr(CN)_6] + [Co(NH_3)_6]Cl \rightarrow y + 3KCl$$

the solution was filted x and y are obtained after evoporation of the solvent from the filtrate.

when y is sent through anion exchange resn , then the complex

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

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

$$C. \left[ Cr \left( NH_3 \right)_6 \right]^{3-1}$$

D. 
$$\left[ Cr(CN)_6 \right]^{3}$$

### Answer: a



**4.** Let  $x_1 < x_2 < x_3 < x_4 < x_5$  and  $y_1 < y_2 < y_3 < y_4 < y_5$  are in AP such that  $\sum_{i=1}^{5} I = 1$   $x_i = \sum_{i=1}^{5} I = 1$   $y_i = 25$  and  $\sum_{i=1}^{5} I = 1$   $y_i = 0$  then  $|y_5 - x_5|$ 

A. 
$$\beta_n$$
 and log  $\beta_n$ 

B.  $K_n$  and  $\log \beta_{n-1}$ 

 $C.\beta_{n-1}$  and  $\log \beta_{n-1}$ 

 $D.K_{n-1}$  and  $log K_{n-1}$ 

### Answer: a



Consider the following complex equilibrium

$$M + L \Leftrightarrow ML, K_1 = \frac{[ML]}{[M][L]}$$
 $ML + L \Leftrightarrow ML_2, K_2 = \frac{[ML_2]}{[ML][L]}$ 
 $ML_2 + L \Leftrightarrow ML_3, K_3 = \frac{[ML_3]}{[ML_2][L]}$ 

5.

$$ML_{n-1} + L \Leftrightarrow ML_n, K_n = \frac{[ML_n]}{[ML_{n-1}][L]}$$

 $K_1, K_2, K_3$  are step wise stability constant

$$M + L \Leftrightarrow ML$$
  $\beta_1 = \frac{\lfloor ML \rfloor}{\lfloor M \rfloor \lfloor L \rfloor}$ 

$$M + 2L \Leftrightarrow ML_2$$
  $\beta_2 = \frac{[ML_2]}{[M][L]^2}$ 

$$M + 3L \Leftrightarrow ML_3$$
  $\beta_3 = \frac{[ML_3]}{[M][L]^3}$ 

$$M + nL \Leftrightarrow ML_n$$
 
$$\beta_n = \frac{[MLn]}{[M][L]^n}$$

 $\beta_P, \beta_2, \beta_3, \beta_n$  are called over all formation constants.

Choose the

correct order

A.  $K_3 > K_2 > K_1$ 

B.  $K_3 = K_2 = K_1$ 

 $C. K_1 > K_2 > K_3$ 

D.  $K_1 = K_2 \neq K_3$ 

### Answer: c



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**6.** (1) 
$$\left[ Cd \left( H_2O \right)_4 \right]^{2+} + 4NH_3 \Leftrightarrow \left[ Cd \left( NH_3 \right)_4 \right]^{2+} + 4H_2O, \log \beta_1$$

$$(2) \left[ Cd \left( H_2O \right)_4 \right]^{2+} + 2en \Leftrightarrow \left[ Cd(en)_2 \right]^{2+} + 4H_2O\log\beta_2$$

 $\log \beta_2 > \log \beta_1$ , is due to

A. Enthalpy change

B. Entropy change

C. Ring strain in the chelate

D. Large change in  $\Delta_0$ 

#### Answer: b



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7. Bonding in metal carbonyls is represented by the end structures.

$$M - C \equiv O \quad \leftrightarrow \quad M = C = O$$

$$I \quad II$$

The contribution of I and II to the real structure depends upon the extent of  $\pi$  - back bonding. Both  $\sigma$  donation and  $\pi$  - back bonding are synergitc. The filled orbitals CO overlap with vacant metal d orbitals which is called  $\sigma$  donation. The filled metal d orbitals overlap with  $\pi$  antibondig moleculat orbital of CO. which is called  $\pi$  - back bonding .

Which one of the following has the highest C-O bond length?

A. 
$$\left[ Cr(CO)_6 \right]$$

- B.  $\left[ Ti(CO)_6 \right]^{2}$
- $C. \left[ Mn(CO)_5 \right]^+$
- D.  $\left[ Fe(CO)_5 \right]$

### Answer: b



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**8.** Bonding in metal carbonyls is represented by the end structures.

$$M - C \equiv O \quad \leftrightarrow \quad M = C = O$$

$$I \quad II$$

The contribution of I and II to the real structure depends upon the extent of  $\pi$  - back bonding. Both  $\sigma$  donation and  $\pi$  - back bonding are synergitc. The filled orbitals CO overlap with vacant metal d orbitals which is called  $\sigma$  donation. The filled metal d orbitals overlap with  $\pi$  antibondig moleculat orbital of CO. which is called  $\pi$  - back bonding .

Which one of the following has the highest C-O bond length?

A. 
$$Ti(C_2H_5)_4$$

B. 
$$Ni(CO)_4$$

$$C. Cr(CO)_6$$

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

#### Answer: a



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**9.** Bonding in metal carbonyls is represented by the end structures.

$$M - C \equiv O \quad \leftrightarrow \quad M = C = O$$

$$I \quad II$$

The contribution of I and II to the real structure depends upon the extent of  $\pi$  - back bonding. Both  $\sigma$  donation and  $\pi$  - back bonding are synergitc. The filled orbitals CO overlap with vacant metal d orbitals which is called  $\sigma$  donation. The filled metal d orbitals overlap with  $\pi$  antibondig moleculat orbital of CO. which is called  $\pi$  - back

bonding.

Which one of the following has the highest C-O bond length?

- A.  $\left[ Ti(CO)_6 \right]^{2-}$
- B.  $\left[V(CO)_6\right]^{-1}$
- C.  $\left[Mn(CO)_5\right]^+$
- D.  $Cr(CO)_6$

#### Answer: c



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10. According to crystal field theory d-orbitals split up in octahedral field into two sets.  $d_{\mathrm{xy}}d_{\mathrm{yx}}d_{\mathrm{zx}}$  have lower energy and  $d_{x^2-y^2}$  and  $d_{z^2}$  have higher energy. The difference in energy of these two sets of d-orbitals is called crystal field splitting energy denoted by  $\Delta_0$ .

In tetrahedral field  $d_{x^2-y^2}$  and  $d_{z^2}$  have lower energy whereas  $d_{xy}d_{yz}d_{zx}$  have higher energy. The difference in energy is denoted  $\Delta_t$ .  $\Delta_0$  can be determined by measuring  $\lambda_{\max}$  for absorption and converting into energy units.  $\Delta_0$  depends upon nature of metal ions as well as nature of ligands. The magnitude of  $\Delta_0$  also decides low energy levels are filled.

which one of the following has the highest magnetic moment

$$A. d_{xy}^2 d_{yz}^2 d_{zx}^1$$

B. 
$$d_{xy}^1 d_{yz}^1 d_{zx}^1 d_{x^2-y^2} d_{z^2}^1$$

C. 
$$d_{xy}^2 d_{yz}^1 d_{zx}^1 d_{x^2-y^2}^1$$

D. 
$$d_{xy}^2 d_{yz}^1 d_{zx}^1 d_{x^2-y^2}^2$$

#### Answer: a



11. According to crystal field theory d-orbitals split up in octahedral field into two sets. $d_{xy}d_{yx}d_{zx}$  have lower energy and  $d_{x^2-y^2}$  and  $d_{z^2}$  have higher energy. The difference in energy of these two sets of d-orbitals is called crystal field splitting energy denoted by  $\Delta_0$ .

In tetrahedral field  $d_{x^2-y^2}$  and  $d_{z^2}$  have lower energy whereas  $d_{xy}d_{yz}d_{zx}$  have higher energy. The difference in energy is denoted  $\Delta_t$ .  $\Delta_0$  can be determined by measuring  $\lambda_{\max}$  for absorption and converting into energy units.  $\Delta_0$  depends upon nature of metal ions as well as nature of ligands. The magnitude of  $\Delta_0$  also decides low energy levels are filled.

Given the following data about absorption maximum of several complex ions, the correct order of  $\Delta_0$  for these ions is

Complexes  $\lambda_{\text{max}}$   $\left[Cr(H_2O)_6\right]^{3+694\text{nm}}$ 

$$\left[Cr\left(NH_3\right)_6\right]^{3+465\text{nm}}$$

$$\left[ CrC1_{6} \right]^{3}$$
 758nm

A. 
$$\Delta_0 \Big[ Cr \Big( NH_3 \Big)_6 \Big]^{3+} > \Delta_0 \Big[ Cr \Big( H_2 O \Big)_6 \Big]^{3+} > \Delta_0 \Big[ CrC1_6 \Big]^{3-}$$

B.  $\Delta_0 \Big[ Cr \Big( NH_3 \Big)_6 \Big]^{3+} = \Delta_0 \Big[ Cr \Big( H_2 O \Big)_6 \Big]^{3+} > \Delta_0 \Big[ CrC1_6 \Big]^{3-}$ 

C.  $\Delta_0 \Big[ Cr \Big( NH_3 \Big)_6 \Big]^{3+} < \Delta_0 \Big[ Cr \Big( H_2 O \Big)_6 \Big]^{3+} < \Delta_0 \Big[ CrC1_6 \Big]^{3-}$ 

D.  $\Delta_0 \Big[ Cr \Big( N_2 O \Big)_6 \Big]^{3+} > \Delta_0 \Big[ CrC1_6 \Big]^{3-} > \Delta_0 \Big[ Cr \Big( NH_3 \Big)_6 \Big]^{3+}$ 

### Answer: a



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12. According to crystal field theory d-orbitals split up in octahedral field into two sets.  $d_{{\bf xy}}d_{{\bf yx}}d_{{\bf zx}}$  have lower energy and  $d_{x^2-y^2}$  and  $d_{z^2}$  have higher energy. The difference in energy of these two sets of d-orbitals is called crystal field splitting energy denoted by  $\Delta_0$ .

In tetrahedral field  $d_{x^2-y^2}$  and  $d_{z^2}$  have lower energy whereas

 $d_{\mathrm{xv}}d_{\mathrm{vz}}d_{\mathrm{zx}}$  have higher energy. The difference in energy is denoted  $\Delta_t$ .

 $\Delta_0$  can be determined by measuring  $\lambda_{\max}$  for absorption and converting into energy units.  $\Delta_0$  depends upon nature of metal ions as well as nature of ligands. The magnitude of  $\Delta_0$  also decides low energy levels are filled.

which one of the following has the highest magnetic moment

A. 
$$[CoF_6]^{3-}$$

$$\mathsf{B.}\left[\mathit{Ni}\Big(H_2O\Big)_6\right]^{2+}$$

$$\mathsf{C.}\left[\mathit{Fe}(\mathit{CN})_{6}\right]^{3}$$

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

#### Answer: a

