

CHEMISTRY

BOOKS - ARIHANT PUBLICATION

COORDINATION COMPOUNDS

Questions For Practice Part I Multiple Choice Questions 1 Mark

1. The coordination number of cobalt in $\lceil Co(NH_3)_3Cl_3 \rceil$ is

A. 3

B. 2

C. 4

D. 6

Answer: D



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- **2.** Which of the following is the IUPAC name of $K_4 \lceil Fe(CN)_6 \rceil$?
 - A. Potassium ferricyanide
 - B. Potassium ferrocyanide
 - C. Potassium hexacyanoferreate (III)
 - D. Potassium hexacyanoferreate (II)

Answer: D



- 3. To which isomers the following compounds belong?
- $\left[Co(NO_2)(NH_3)_5\right]Cl_2$ and $\left[Co(ONO)(NH_3)_5\right]Cl_2$
 - A. Geometrical isomerism
 - B. Linkage isomerism

C. Ionisation isomerism
D. Ligand isomerism
Answer: B
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4. Which kinds of isomerism are exhibited octahed
?

ral $igl[Co(NH_3)_4 Br_2 igr] Cl$

A. Geometrical and ionisation

B. Geometrical and optical

C. Optical and Ionisation

D. Geometrical only

Answer: A



Questions For Practice Part I Very Short Answer Type Questions 1 Mark

- 1. Potash alum is an example of
- (simple, salt, double salt, complex salt)
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- **2.** How many ions are produced from the complex, $\left[Co(NH_3)_6\right]Cl_2$ in solution ?
 - Watch Video Solution

- **3.** The oxidation number of ${}'Co'$ in the complex K[Co(CO)4] is _____
 - Watch Video Solution

4. Write the IUPAC name of $K_2igl[CrCO(CN)_5igr]$

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5. Write the IUPACname of $Naig[Al(OH)_4ig].$
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6. Write the IUPAC name of $\left[Cr(NH_3)_6 ight]\left[Co(CN)_6 ight].$
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7. Write the IUPAC name of $K_3igl[Fe(CN)_5NOigr].$
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8. Write the IUPAC name of $K_2[PtCl_6]$.
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9. Write the IUPAC name of $\left[Ni(\mathrm{en})_3
ight]Cl_2.$ Watch Video Solution



10. Write the IUPAC name of $[Pt(NH_3)_4][PtCl_4]$.



11. Which of the following is more stable complex and why? $\left[Co(NH_3)_6\right]^{3+}$ and $\left[Co(\text{en})_3\right]^{3+}$







13. Why is optical isomerism not shown by square planar complexes?

12. Why is geometrical isomersim not possible for tetrahedral complexes?

14. What type of isomerism is exhibited by the complex $\left[Co(\text{en})_3\right]^{3+}$?



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

$$\left[Cr(C_2O_4)_3
ight]^{3}$$



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

 $igl[{Co(NH_3)}_3 Cl_3 igr]$



17. Which type of ligand is EDTA.



Questions For Practice Part I Short Answer Type I Questions 2 Marks

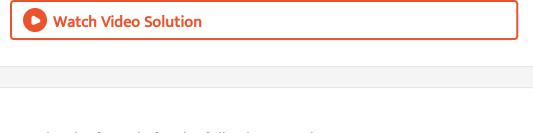
1. What is meant by the chelate effect? Give an example.



- 2. Amongst the following, the most stable complex is
- (i) $Fe(H_2O)_6ig]^{3\,+}$ (ii) $ig[Fe(NH_3)_6ig]^{3\,+}$
- (iii) $\left[Fe(C_2O_4)_3\right]^{3-}$ (iv) $\left[FeCl_6\right]^{3-}$
 - Watch Video Solution

3. Write t he IUPAC name of the following complex.

 $\left[Cr(NH_3)_2Cl_2(ext{en})\right]Cl$



4. Write the formula for the following complex.

Pentaamminenitrito-O--cobalt (III)



5. Name the type of isomerism when ambidentate ligands are attached to central metal ion. Give two examples of ambidentate ligands.



6. Write the IUPAC name of the complex $\left[Cr(NH_3)_4Cl_2\right]^+$. What type of isomerism does it exhibit?



7. Give the evidence that $\left[Co(NH_3)_5Cl\right]SO_4$ and $\left[Co(NH_3)_5SO_4\right]Cl$ are ionisation isomers.



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



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



10. $FeSO_4$, solution mixed with $(NH_4)_2SO_4$, solution in 1:1 molar ratio gives the test for Fe^{2+} ion but $CuSO_4$, solution mixed with aqueous ammonia in1:4 molar ratio does not give the test for Cu^{2+} ion. Explain.



11. A coordiantion compound $CrCl_3\cdot 4H_2O$ percipitates silver chloride when treated with silver nitrate. The molar conductance of its solution corresponds to a total of two ions. Write the structural formula of the compound and name it.



Questions For Practice Part I Short Answer Type Ii Questions 3 Marks

1. Explain with two examples each of the following. Coordination entity, ligand, coordination number, coordination polyhedron.



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2. What is meant by unidentate, didentate and ambidentate ligand? Give two examples for each.



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3. Specify the oxidation number of metal in the following coordination entities.

$$\big[Co(H_2O)(CN)(\mathrm{en})_2\big]^{2\,+}$$



4. Specify the oxidation number of metal in the following coordination entities.

$$\left[CoBr_2(\mathrm{en})_2
ight]^+$$



5. Specify the oxidation number of metal in the following coordination entities.

 $\left[PtCl_4\right]^{2\,-}$



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



7. Using IUPAC norms, write the systematic names of the following.

 $igl[{\it Co(NH_3)}_6 igr] {\it Cl}_3$



8. Using IUPAC norms, write the systematic names of the following.

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



9. Using IUPAC norms, write the systematic names of the following.

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



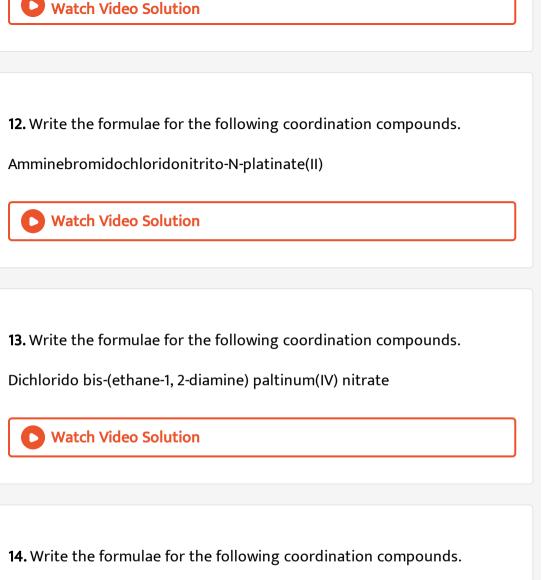
10. Write the formulae for the following coordination compounds.

Tetraamminediaquacobalt(III) chloride



11. Write the formulae for the following coordination compounds.

Potassium tetracyanonickelate(II)



Iron(III) hexacyanoferrate(II)



15. Write the formulae for the following coordination compounds.

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16. Write the types of isomerism exhibited by the following complexes.

 $\left[Co(NH_3)_5Cl\right]SO_4$



17. Write the types of isomerism exhibited by the following complexes.

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



18. Write the types of isomerism exhibited by the following complexes.

 $igl[Co(NH_3)_6 igr] igl[Cr(CN)_6 igr]$



19. What type of isomerism is exhibited by $\left[Co(NH_3)_4Cl_2\right]^+Br^-$? Write the structures of the possible isomers and the state of hybridisation of the central metal atom.



20. Draw all the isomers (geometrical and optical) of

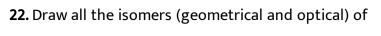
$$\left[CoCl_2(\mathrm{en})_2\right]^+$$



21. Draw all the isomers (geometrical and optical) of

$$\left[Co(NH_3)_2Cl_2(\mathrm{en})\right]^+$$





 $\left[Co(NH_3)_2Cl_2(ext{en})
ight]^+$



Questions For Practice Part I Long Answer Type Questions 7 Marks

1. Write short notes on: Werner's theory of co-ordination compounds.



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



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

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



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

 $[Co(en)_3]Cl_3$



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

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



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

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



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Questions For Assessment Part I Multiple Choice Type Questions 1 Mark

1. A complex compound in which the oxidation number of a metal is zero,

is

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

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

C. $[Ni(CO)_{A}]$

D. $\lceil Pt(NH_3)_4 \rceil Cl_2$

Answer: C



2. The primary and secondary valencies of chromium in the complex ion, dichlorodioxalatochromium (III), respectivelr are

A. 3, 4

B. 4, 3

C. 3, 6

D. 4, 3

Answer: A



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Questions For Assessment Part I Very Short Answer Type Questions 1 Mark

1. Write t he oxidation number of Pt in $\left[PtCl_2(NH_3)_2\right]$.





3. How many ions are produced from the complex, $\left[Cr(H_2O)_6\right]Cl_3$ in solution ?



4. Give the examples of two coordination complexes where central metal atom has coordination number

(i) 5 (ii) 6



5. Which of the following complex is more stable and why?

$$\left[Cr(ox)_3
ight]^{3-}$$
 or $\left[PtCl_2(ext{en})_2
ight]^{2+}$

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6. Which type of isomerism is exhibited by

 $\left[Co(NH_3)_3(NO_2)_3\right]?$



Questions For Assessment Part I Short Answer Type I Questions 2 Marks

1. What are ambidentate ligand? How these ligands affects the coordination complex?



2. What is the primary and secondary valency of chromium in the complex ion, dichlorodioxalatochromium(III) ?



3. How many isomers can be formed for the complex

$$\lceil Co(C_2O_4)_2(NH_3)_2 \rceil$$
 ?



4. The complexes

 $\left[Co(NH_3)_6 \right] \left[Cr(CN)_6 \right] \ ext{and} \ \left[Cr(NH_3)_6 \right] \left[Co(CN)_6 \right]$ are the examples of which type of isomerism? State its defination.



Questions For Assessment Part I Short Answer Type Ii Questions 3 Marks

1. Write the IUPAC name of the following coordiantion compounds.

$$\big[Co(H_2O)_4(\mathrm{en})\big]^{3\,+}$$



2. Write the IUPAC name of the following coordination compounds.

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



3. Write the IUPAC name of the following coordination compounds.

4. Using IUPAC norms, write the formulae for the following?

5. Using IUPAC norms, write the formulae for the following?

$$[FeF_6]^{3-}$$



Sodium tetrafluoridosilicate (IV)



Tris-(ethane- 1,2-diamine) chromium (III) chloride



6. Using IUPAC norms, write the formulae for the following? Hexa aquachromium (III) chloride **Watch Video Solution** 7. Draw all the possible isomers (structural and stereoisomeric) having the composition $CrBr_2Cl(NH_3)_A$. **Watch Video Solution** Questions For Assessment Part I Long Answer Type Questions 7 Marks 1. Indicate the types of isomerism exhibited by the following complexes and draw the structures for these isomers. $[Co(NH_3)_5NO_3]SO_4$ **Watch Video Solution**

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

 $\lceil CoCl(NH_3)_4(H_2O) \rceil Br_2$



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

 $\big[Cr(H_2O)_5(SCN)\big]^{2\,+}$



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

 $\left[\operatorname{Co(en)}_3\right]\left[\operatorname{Cr(CN)}_6\right]$



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

 $\left[PtNH_{3}Cl(\mathrm{Py})_{2}
ight]$



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6. $PtCl_4$ and NH_3 may form five complexes, $A(PtCl_4 \cdot 6NH_3), B(PtCl_4 \cdot 5NH_3), C(PtCl_4 \cdot 4NH_3), D(PtCl_4 \cdot 3NH_3)$

. One mole of each A, B, C, D and E reacts with excess of $AgNO_3$ to yield 4,

3, 2 and 1 mole(s) of AgCl respectively, while E gives, no AgCl. The

conductance of their solutions are in the order A > B > C > D > E.

On the basis of Werner's theory, write their strucutre and give the total number of ions given by one complex.



Questions For Practice Part Ii Multiple Choice Type Questions 1 Mark

1. Which of the following complex species does not involve in the inner orbital hybridisation?

A.
$$\left[CoF_{6}
ight]^{3}$$
 –

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

C.
$$\left \lceil Fe(CN)_6 \right \rceil^{3}$$

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

Answer: A



- **2.** In which of the following coordination entities the magnitude of Δ_0 (CFSE in octahedral field) wiil be maximum? (At. no. of Co = 27)
 - A. $\left[CoH_2O
 ight)_6
 ight]^{3\,+}$
 - B. $\left[Co(NH_3)_6 \right]^{3+}$
 - C. $\left[Co(CN)_6 \right]^{3}$

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

Answer: C



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- **3.** In spectrochemical series, chlorine is above than water, i.e. $Cl>H_2O$, this is due to
 - A. good π -acceptor properties of Cl
 - B. strong $\sigma\text{-donor}$ and good $\pi\text{-acceptor}$ properties of Cl
 - C. good π -donor properties of Cl
 - D. larger size of CI than H_2O

Answer: B



1. In which case splitting will belarger, 3d-orbital or 4d-orbital?



2. Why are low spin tetrahedral complexes not formed?



3. Arrange the following complex ions in the increasing order of srystal field splitting energy

```
(\Delta_0), \left[Cr(CN)_6
ight]^{3-}, \left[Cr(CN)_6
ight]^{3-}, \left[Cr(NH_3)_6
ight]^{3+}.
```



4. Why is CO a stronger ligand than Cl^- ?



5. Magnetic moment of $\left[MnCl_4
ight]^{2-}$ is 5.92 BM. Give reason.



6. Anhydrous $CuSO_4$ is white while hydrated $CuSO_4$ is blue. Why?



Questions For Practice Part Ii Short Answer Type I Questions 2 Marks

IUPAC names and calculate spin only magnetic moment of B.

1. Identify A and B in the given sequence of reaction. Also, write their

$$Fe^{3+}(ext{aq}) \xrightarrow{ ext{Excess of }SCN^{ heta}} ext{Blood red colour (A)}$$

 $\xrightarrow{\operatorname{Excess\ of} F} \operatorname{Colourless\ }(\operatorname{B})$



2. Why do compounds having similar geometry have different magnetic moment?



3. 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.



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

$$K_3igl[Co(C_2O_4)_3igr]$$



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

 $ig[Mn(H_2O)_6ig]SO_4$



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

 $[Mn(H_2O)_6]SO_4$



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

 $[Mn(H_2O)_6]SO_4$



8. Why are different colours observed in octahedral and tetrahedral complex for the same metal and same ligands?



9. What will be the correct order for the wavelength of absorption in the visible region for the following? $\left[Ni(NO_2)_6\right]^{4-}, \left[Ni(NH_3)_6\right]^{2+}, \left[Ni(H_2O)_6\right]^{2+}.$

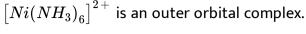


Questions For Practice Part Ii Short Answer Type Ii Questions 3 Marks

1. Expalin the basis of valence bond theory that $\left[Ni(CN)_4\right]^2$ ion with square planar structure is diamagnetic and the $\left[NiCl_4\right]^2$ ion with tetrahedral geometry is paramagnetic.



2. Explain $\left[Co(NH_3)_6
ight]^{3+}$ is an inner orbital complex whereas





3. Predict the number of unpaired electrons in the square planar $\left\lceil Pt(CN)_4 \right\rceil^{2-}$ ion.



4. Write the assumptions of crystal field theory. Discuss the patterm of splitting of d-orbitals under the effect of an octahedral crystal field.



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

6. What is the crystal field stabilisation energy? How does the magnitude of Δ_0 decide the actual configuration of d-orbital in a coordination entiity?



7. What is meant by crystal field splitting energy? On the basis of crystal field theory, write the eelctrone configuration of d^4 in terms of t_{2g} and e_g in an octahedral field when

(i)
$$\Delta_0 > P$$
 (ii) $\Delta_0 < P$



8. The haexaaquamanganese (II) ion contains five unpaired electrons, while the hexacyano ion contains only one unpaired electron. Expalin using crystal field theory.

9. Give the electronic configuration of the following complexes on the basis of crystal field splitting theory.



 $[CoF_6]^{3-}, [Fe(CN)_6]^{4-} \text{ and } [Cu(NH_3)_6]^{2+}.$

10. Amongst the following ions which one has the highest magnetic moment value?

(i)
$$\left[Cr(H_2O)_6
ight]^{3+}$$
 (ii) $\left[Fe(H_2O)_6
ight]^{2+}$ (iii) $\left[Zn(H_2O)_6
ight]^{2+}$



11. $\left[Fe(CN)_6\right]^{4-}$ and $\left[Fe(H_2O)_6\right]^{2+}$, are of different colours in dilate solution, although in both the cases iron is in +2 oxidation state explain.

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12. A solution $igl[Ni(H_2O)_6igr]^{2+}$ is green while a solution of $igl[Ni(CN)_4igr]^{2-}$ is colourless. Explain.



Questions For Practice Part Ii Long Answer Type Questions 7 Marks

1. Discuss the nature of bonding in the following coordination entities on the basis of valence bond theory.

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



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2. Discuss the nature of bonding in the following coordination entities on the basis of valence bond theory. $[FeF_6]^{3}$

3. Discuss the nature of bonding in the following coordination entities on the basis of valence bond theory.

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



4. Discuss the nature of bonding in the following coordination entities on the basis of valence bond theory. $[CoF_6]^{3-}$



 $K[Cr(H_2O)_2(C_2O_4)_2] \cdot 2H_2O$

5. Write down t he IUPAC name for each of the following complexes and indicate the oxidation state, electronic configuration and coordination number. Also give magnetic moment of the complex.



6. Write down t he IUPAC name for each of the following complexes and indicate the oxidation state, electronic configuration and coordination number. Also give magnetic moment of the complex.

 $\big[Co(NH_3)_5Cl\big]Cl_2$



7. Write down t he IUPAC name for each of the following complexes and indicate the oxidation state, electronic configuration and coordination number. Also give magnetic moment of the complex.

 $\left[CrCl_{3}(\mathrm{py})_{3}\right]$



8. Write down t he IUPAC name for each of the following complexes and indicate the oxidation state, electronic configuration and coordination

number. Also give magnetic moment of the complex.

 $Cs[FeCl_4]$



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9. Write down t he IUPAC name for each of the following complexes and indicate the oxidation state, electronic configuration and coordiantion number. Also give magnetic moment of the complex.

$$K_4ig[Mn(CN)_6ig]$$



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Questions For Assessment Part Ii Multiple Choice Type Questions 1 Mark

1. Which complex compound possesses sp^3d^2 hybridisation ?

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

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

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

D.
$$\left[Fe(Cl)_6
ight]^{3}$$
 –

Answer: D



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2. Crystal field stabilisation energy for high spin d^4 octahedral complex is

A.
$$-0.6\Delta_0$$

$$B.-1.8\Delta_0$$

C.
$$-1.6\Delta_0+P$$

D.
$$-1.2\Delta_0$$

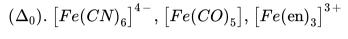
Answer: A



1. Give one example for. Neutral ligand of carbon **Watch Video Solution** 2. Give one example for . Positive ligand of nitrogen **Watch Video Solution** 3. What are the factors on which internal energy depends? Watch Video Solution 4. An elements 'X' has one unpaired eelctron. Calculate its magnetic moment.

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5. Arrange thefollowing complex ions is the increasing order of crystal field splitting energy





Questions For Assessment Part Ii Short Answer Type I Questions 2 Marks

1. Using the valence bond theory, predict the geometry and magnetic behaviour of $\left[CoF_6\right]^{3-}$.

(Atomic number of Co = 27)



2. deduce the structure 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.

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3. Give the name, stereochemistry and the magnetic behaviour of the following complexes.

(i) $\left[Co(NH_3)_5Cl\right]Cl_2$ (ii) $K_2\left[Ni(CN)_4\right]$



4. Give the formula of each of the following coordination entities.

 $Co^{3\,+}$ ion is bound to one Cl^- , one NH_3 molecule and two bidentate ethylenediammine (en) molecules.



5. Give the formula of each of the following coordination entities.

 $Ni^{2\,+}$ ion is bound to two water molecules and two oxalate ions. Write

the name and magnetic behaviour of each of the above coordiantion entities. (Atomic number of, Ni = 28)



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Questions For Assessment Part Ii Short Answer Type Ii Questions 3 Marks

1. Compare the following complexes with respect to structural shapes of untis, magnetic behaviour and hybrid orbitals involved in units.

$$\big[Ni(CN)_4\big]^{2\,-}$$

(Atomic number of Ni = 28, Co = 27).



 $[NiCl_4]^{2-}$

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2. deduce the following complexe with respect to structural shapes of untis, magnetic behaviour and hybrid orbitals involved in units.

(Atomic number of Ni = 28).

3. Compare the following complexes with respect to structural shapes of untis, magnetic behaviour and hybrid orbitals involved in units.

 $[CoF_{6}]^{3}$

(Atomic number of Ni = 28, Co = 27).



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4. Explain the following.

Low spin octahedral complexes of nickel are not known.



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5. Explain the following.

CO is a stronger ligand than NH_3 for many metals.



6. $\left \lceil Ti(H_2O)_6 \right
ceil^{3+}$ absorbs light of wavelength 500 nm. Name one ligand which would from Ti(III) complex absorbing light of lower wavelength than 500 nm and one ligand which would form a complex absorbing light of wavelength higher than 500 nm.



7. $\left[Co(H_2O)_4
ight]^{2+}, \left[CoCl_4
ight]^{2-} ext{ and } \left[Co(ext{dmg})_2
ight]$ are complexes or Co(II) but magnetic moments of $igl[Co(H_2O)_4 igr]^{2+}$ and $igl[CoCl_4 igr]^{2-}$ is higher ($\mu=3.87$ Bm for each) than $\left[Co(\mathrm{dmg})_2\right][\mu=1.73\mathrm{BM}]$, explain.



Questions For Assessment Part Ii Long Answer Type Questions 7 Marks

1. What is spectrochemical series?



2. Draw the structures of geometrical isomers of the following coordination complexes:

$$\left[Co(NH_3)_3 Cl_3
ight] ext{ and } \left[CoCl_2(ext{en})_2
ight]^+$$

(en = ethylene diamine and atomic number of Co is 27).



3. Describe the limitations of valence bond theory.



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Questions For Practice Part Iii Multiple Choice Type Questions 1 Mark

1. The most stable complex among the following is

A.
$$\left[Pd(Cl)_4\right]^{4-}$$

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

C.
$$\left[Ni(Cl)_4
ight]^{4-}$$
D. $\left[Ni(CN)_4
ight]^{3-}$

Answer: D



- 2. In metal carbonyl (organometallic) complexes, the M-C bond is
 - A. ionic
 - B. covalent with ionic character
 - C. covalent
 - D. coordinate covalent

Answer: D



3. Give an example of a metal carbonyl having metal-metal bond. Watch Video Solution
Questions For Practice Part Iii Very Short Answer Type Questions 1 Mark
1. CO is a stronger complexing agent than NH_3 , why? Watch Video Solution
2. Which of the following complex is used for treatment of cancer? Watch Video Solution
3. Name the complex used as oxygen carrier in the blood. Watch Video Solution

4. What is Wilkinson catalyst?
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5. Name the complex which is responsible for photosynthesis.
Watch Video Solution
6. $Ni(CO)_4$ is an example of and bonded organo metallics.
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7. Use EAN rule and predict the molecular formula of nickel carbonyl?
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Ouestions For Practice Part Iii Short Answer Type I Questions 2 Marks

1. Discuss the nature of bonding in metal carbonyls.



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



3. What do you understand by stepwise stability constant and overall stability constant of a coordination compound? How are these two constants related?



4. Discuss briefly giving an example in each case, the role of coordiantion compounds in

(i) biologycal systems (ii) medical chemistry (iii) analytical chemistry (iv) extraction / metallurgy of metals **Watch Video Solution** 5. What are organometallic compounds? Give two examples. **Watch Video Solution** Questions For Practice Part Iii Short Answer Type Ii Questions 3 Marks 1. Calculate The ratio of $\left[Ag(NH_3)_2
ight]^+ \; ext{and} \; \left[Ag^+
ight]$ in 0.1 M NH_3 solution. **Watch Video Solution**

2. Calculate

The ratio of $\left[Ag(S_2O_3)_2
ight]^{3-}$ and $\left[Ag^+
ight]$ in 0.1 M $S_2O_3^{2-}$ solution. Given the stability / formation constants (K_f) for that $\left[Ag(NH_3)_2
ight]^{2+}$ and $\left[Ag(S_2O_3)_2
ight]^{3-}$ are $1.7 imes10^7$ and $1.0 imes10^{13}$



respectively.

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Questions For Practice Part Iii Long Answer Type Questions 7 Marks

1. Draw the structure of following homoleptic metal carbonyl.

 $[Cr(CO)_6]$



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2. Draw the structure of following homoleptic metal carbonyl.

 $\lceil Fe(CO)_5 \rceil$

3. Draw the structure of following homoleptic metal carbonyl.

 $Nl(CO)_4$



4. Discuss important of coordination compounds in boiligical system.



Questions For Assessment Part Iii Multiple Choice Type Questions 1 Mark

1. Which of the following organometallic compound has σ and π -bonds?

A.
$$\left[Feig(\eta^5-C_5H_5ig)_2
ight]$$

B.
$$Kigl[PtCl_3igl(\eta^2-C_2H_4igr)igr]$$

C. Both (a) and (b)

D. None of the above

Answer: B



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Column 1

2. Match the Column and choose the correct option from the codes given below.

	(Coordination compounds)		
A.	Chlorophyll	1.	Cobalt
в.	Vitamin B ₁₂	2.	Iron
C.	Haemoglobin	3.	Magnesium
40.	Haemogioon	3.	Magnesium

Codes

	\boldsymbol{A}	B	C
A.	1	$B \ 2$	3
ъ	\boldsymbol{A}	B	C
В.	3	B 1	2
C	\boldsymbol{A}	$B \ 2$	C
С.	3	2	1
	Α	B	C

Answer: B



Questions For Assessment Part Iii Very Short Answer Type Questions 1 Mark

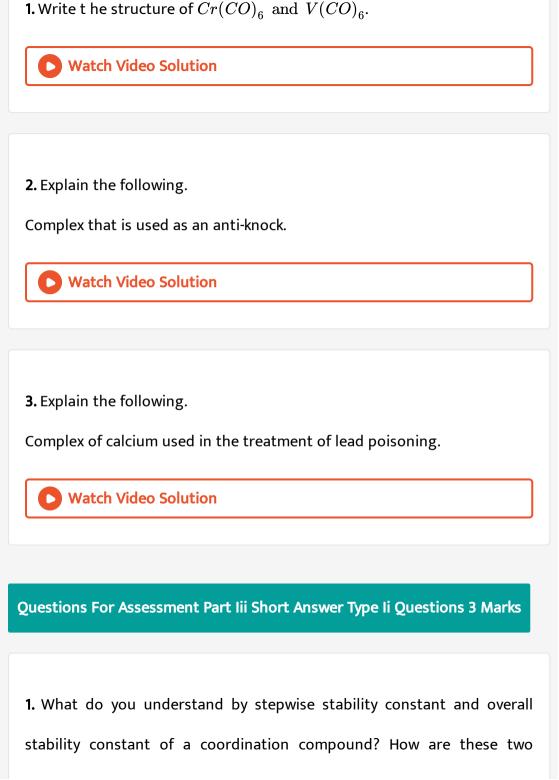
1. If an element have high positive charge then what will be its effect on the stability of complex?



2. What is the reason for wide application and significance of coordiantion complex?



Questions For Assessment Part Iii Short Answer Type I Questions 2 Marks



constants related? **Watch Video Solution** Questions For Assessment Part Iii Long Answer Type Questions 7 Marks 1. Name the following: Ring that is attached to Fe^{2+} in haemoglobin. **Watch Video Solution** 2. Name the following: Vitamin B_{12} is a coordination compound of which element. **Watch Video Solution** 3. Name the following:

Comples that is used in homogeneous and heterogeneous catalysis.



4. Give an example for each when coordination compounds play an important role in



photosynthesis by plant.

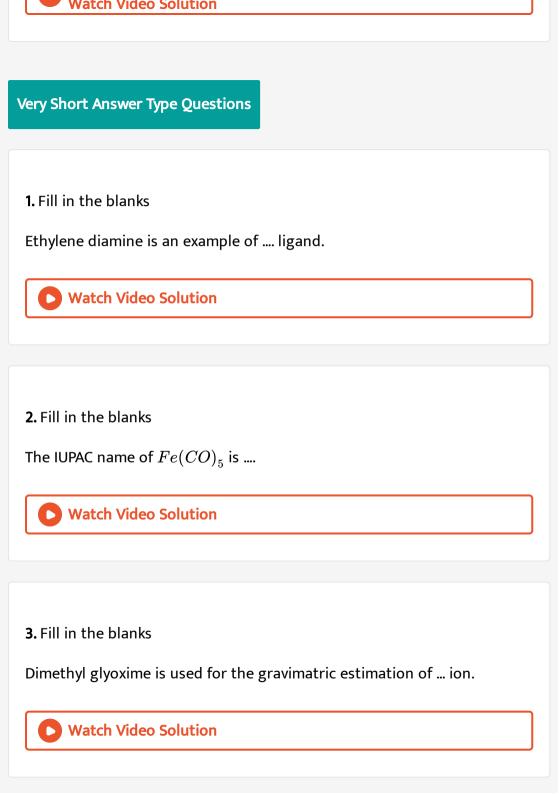
5. Give an example for each when coordination compounds play an important role in



removal of excess of iron present in our body.

6. Give an example for each when coordination compounds play an important role in

development of photographic film.



4. Fill in the blanks

EDTA is a ligand.



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5. Fill in the blanks

 $K_4[Fe(CN)_6]$ is a complex compound, but potash alum is a



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6. Fill in the blanks

One molecule of $\lceil Pt(NH_3)_6 \rceil Cl_4$ gives Ions is solution and requires

of $AgNO_3$ for complete precipitation of chloride ions.



7. Fill in the blanks

Geometrical isomerism is not observed in complex of coordination number 4 of geometry.



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8. Fill in the blanks

The type of magnetism exhibited by $\left[Mn(H_2O)_6\right]^{2+}$ ion is



- 9. Match the following
- I. Double salt (a) $[Co(NH_3)_3Cl_3]$
 - II. Neutral molecule (b) Hexadentate
 - III. EDTA (c) Bidentate
 - IV. Ni(CO)₄ (d) Paramagnetic V. [Cr(NH₃)₆]³* (e) FeSO₄(NH₄)₇SO₄ 6H₇O
 - VI. Low spin complex (f) K₄[Fe(CN)₆]
 - VII. Glycine (g) Diamagnetic
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10. What type of isomers are the following

 $[Co)_5 \text{MnSCN}$ and $[(Co)_5 \text{MnNCS}]$



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11. What type of isomers are the following

 $\left[{Co(\text{en})_3} \right] Cr(CN)_6 \big] \ \ \text{and} \ \ \left[{Cr(\text{en})_3} \right] \left[{Co(CN)_6} \right]$



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12. What type of isomers are the following

 $\left[Co(NH_3)_5NO_3\right]SO_4$ and $\left[Co(NH_3)_5SO_4\right]NO_3$



13. What type of isomers are the following

 $[Co(py)_2(H_2O)_2Cl_2]$ and $[Co(py)_2(H_2O)Cl_2]H_2O$



Short Answer Type I Questions

1. What is the difference between molecular compound and complex compound?



2. Define the terms : central ion, complex ion, ligand, coordination number.



3. Give IUPAC names of the following compounds

 $\left[Ag(NH_3)_2\right]^+, \left[Cu(NH_3)_4\right]^+, \left[Cr(H_2O)_4Cl_2\right],$

 $K_2[PtCl_6], [Fe(CN)_6]^{4-}, [Co(en)_2Cl_2]SO_4,$

 $[Cr(NH_3)_3Cl_3](H_2O)_2, Na_3[Co(NO_2)_6],$

 $[Co(en)_2Cl(NO_2)]^+, [Pt(NH_3)Cl_5]^-$



4. Write the formula of the following complexes:

Hexaamminaenickel (II) chloride,

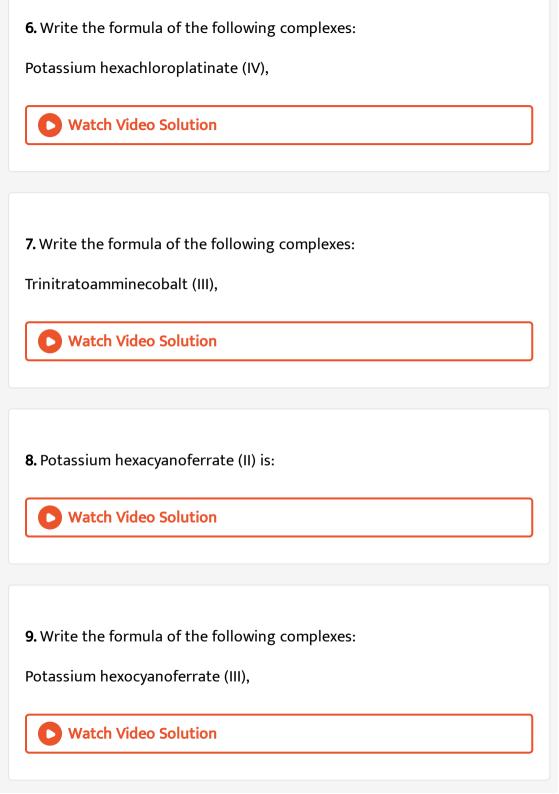


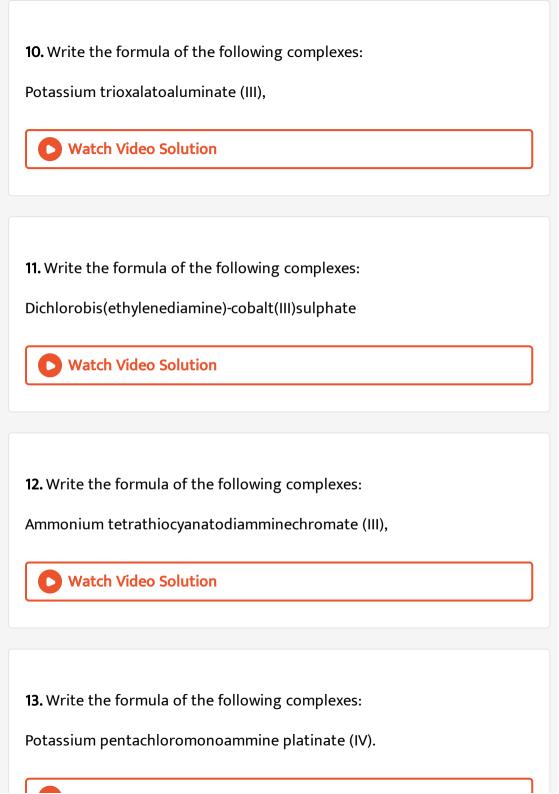
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5. Write the formula of the following complexes:

Hexaammine chromium (III) nitrate,







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14. Write a note on Werner's theory of coordination compounds.				
·				
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15. Give some examples of molecular (non-ionic) complexes.				
Watch Video Solution				
16. Give some examples of ionic complexes.				
Watch Video Solution				
17. What is the symbol used before the bridging ligand while naming a				
17. What is the symbol used before the bridging ligand while halling a				
complex.				
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18. Give the order of naming the ligands.



19. What type of numeral is used to indicate oxidation state of central ion? Write t he structure of $Fe(CO)_5$.



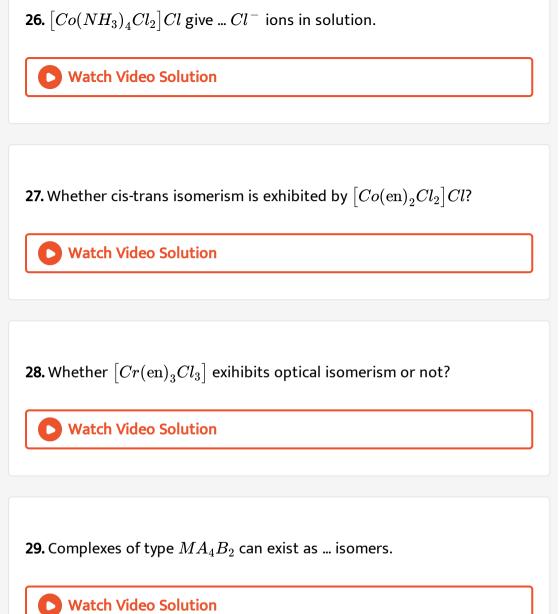
20. Give examples od cic-trans isomerism exhibited by complexes of coordination number 4 and 6.



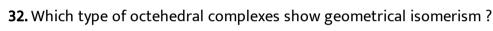
21. Give some examples of optical isomerism exhibited by complexes of coordination number 6.



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22. What are primary and secondary valencies?
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23. How optical isomers are designated?
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24. $\left[Co(NH_3)_5Br\right]SO_4$ and $\left[Co(NH_3)_5SO_4\right]Br$ show isomerism.
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25. $KCl \cdot MgCl_2 \cdot 6H_2O$ is a , but $K_4ig[Fe(CN)_6ig]$ is a
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31. How many isomers are there for the complex $igl[Co(NH_3)_3Cl_3igr]$?
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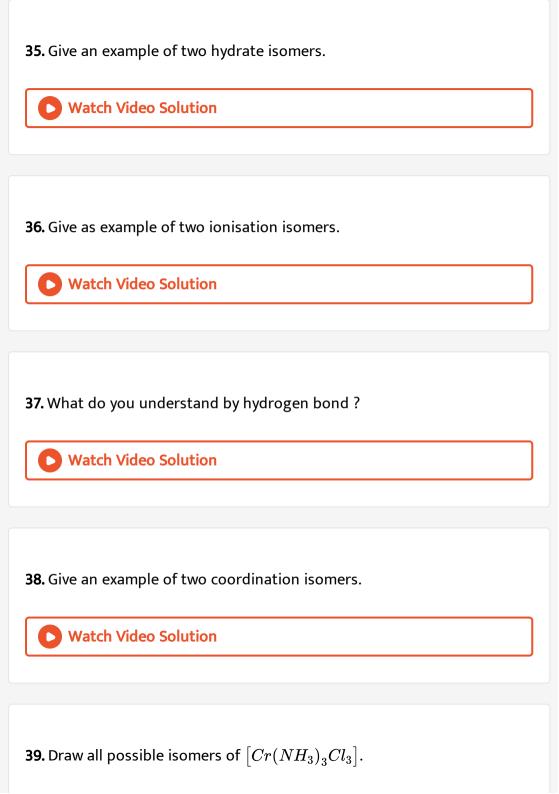


33. Which type of octehedral complexes show geometrical isomerism?



34. Does $\left[Co(NH_3)_6
ight]Cl_3$ show optical isomerism?







40. Give one example of both ionisation and hydrtae isomerism in single compound.



41. Draw the structure of triamminetrichloroplatinum (IV) chloride.



42. $[Rh(en)_3][IrCl_6]$ and $(Rh(en)_2Cl_2]Ir(en)Cl_4]$ are which type isomers?



43. How many chloride ions can be precipitated from an aqueous solution of

 $[Cr(py)_2(H_2O)_2Cl_2]Cl$ with $AgNO_3$?



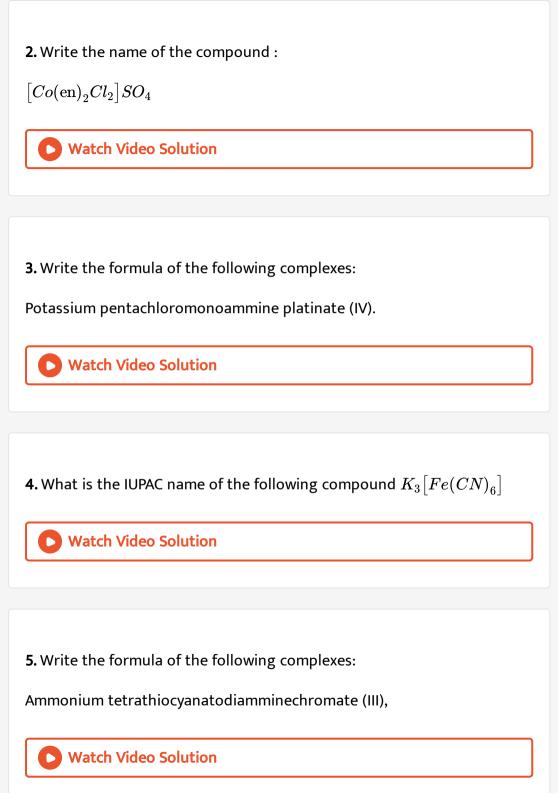
44. How a double salt can be distinguished from a complex salt?



Short Answer Type Ii Questions 3 Marks

1. What is the coordination number of each atom in ccp structure?





6. Write the formula of the following complex:

Tris (ethylene diamine) cobalt (III) sulphate.



7. Write short notes on: Werner's theory of co-ordination compounds.



8. Write the name of the compound : $\left[(C_2H_5)_5N\right]_2[ZnCl_4]$



Long Answer Type Questions 7 Marks

1. Write various postulates and applications of valence bond theory in coordination compounds.



2. Distinguish between inner and outer orbital complex with suitable examples.



3. Account of the following on the basis of valence bond theory

 $\left[Ni(CN)_4\right]^2$ is diamagnetic and square planar.



4. Expalin the basis of valence bond theory that $\left[Ni(CN)_4\right]^{2-}$ ion with square planar structure is diamagnetic and the $\left[NiCl_4\right]^{2-}$ ion with

tetrahedral geometry is paramagnetic.

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5. Dicuss in brief the crystal field theory. How does it differ from valence bond theory?

bond theory? ${\it Explain the fact the } \left[{\it CoF}_6 \right]^{3-} {\it is paramagnetic whereas } \left[{\it Co(NH_3)}_6 \right]^{3+}$



is daimagnetic although both are octahedral.

6. Between $\left[Fe(H_2O)_6\right]^{3+}$ and $\left[Fe(CN)_6\right]^{3-}$ which is a low spin complex and why ?



7. Tetrahedral complexes do not show low spin configuration. Explain.



8. Discuss briefly how crystal field theory explains the magnetic properties of complexes of transition metals.



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- **9.** Discuss the structure of following complexes on the basis of CFT.
- (a) $\left[Cu(NH_3)_6
 ight]^{3+}$ (b) $\left[Fe(CN)_6
 ight]^{4-}$



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Chapter Practice Multiple Choice Type Questions 1 Mark

- 1. Which of the following is diamagnetic?
 - A. $\left[Fe(CN)_6
 ight]^{3}$
 - B. $\left[NiCl_4^{2\,-}
 ight]$

C.	$ig[Ni(CO)_4ig]$
----	------------------

D.
$$\left(MnCl_4\right]^{2}$$

Answer: B



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- **2.** The complexes $\left[Co(NH_3)_6\right]\left[Cr(CN)_6\right]$ and $\left[Cr(NH_3)_6\right]\left[Co(CN)_6\right]$ are the examples of which type of isomerism? State its defination.
 - A. Ionisation isomerism
 - B. Linkage isomerism
 - C. Coordination isomerism
 - D. Solvate isomerism

Answer: C



Chapter Practice Very Short Answer Type Questions 1 Mark

1. The coordination number of Fe in $\left[Fe(CN)_6\right]^{4-}$ is



2. Write the IUPAC name of $\left[Co(NH_3)_5ONO\right]^{2+}$.



3. Write the ionisation isomer of $[Pt(NH_3)NO_2]Cl$.



4. What is the role of synergic bonding in metal-carbonyl complexes?



Chapter Practice Short Answer Type I Questions 2 Marks

1. Discuss the role of complex in analytical chemistry



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2. Discuss the role of complex in medicines



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3. What is crystal field splitting among octahedral and tetrahedral crystal fields? In which case, the magnitude of crystal field splitting is larger?



4. Draw all the isomers (geometrical and optical) of

 $\left[Co(NH_3)_2Cl_2(ext{en})
ight]^+$



5. Write the structures and names of all stereoisomers of the following compounds:

 $[Co(en)_3]Cl_3$



6. Write the structures and names of all stereoisomers of the following compounds:

 $\lceil Pt(NH_3)_2 Cl_2 \rceil$



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



Chapter Practice Short Answer Type Ii Questions 3 Marks

1. Differentiate between weak field and strong field coordination entity.



2. Write the assumptions of crystal field theory. Discuss the patterm of splitting of d-orbitals under the effect of an octahedral crystal field.



3. The most stable oxidation state of iron is



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4. Using crystal field theory draw energy level daigram, write electronic configguration of the central metal atom or ion and determine the magnetic moment value in the following:

 $igl[Fe(H_2O)_6igr]^{2\,+}$



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5. Using crystal field theory draw energy level daigram, write electronic configguration of the central metal atom or ion and determine the magnetic moment value in the following:

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



6. Using valence bond theory, explain the following in relation to the complexes, given below: $\left[Mn(CN)_6\right]^{3-}$, $\left[Cr(H_2O)_6\right]^{3+}$.

Type of hybridisation



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7. Using valence bond theory, explain the following in relation to the complexes, given below: $\left[Mn(CN)_6\right]^{3-}, \left[Cr(H_2O)_6\right]^{3+}.$

Magnetic behaviour



8. Using valence bond theory, explain the following in relation to the complexes, given below: $\left[Mn(CN)_6\right]^{3-}, \left[Cr(H_2O)_6\right]^{3+}.$

Magnetic behaviour



Chapter Practice Long Answer Type Questions 7 Marks

1. Give detailed description of CFT for different types of ligand and explain how CFT explain colour of coordination compound? Use suitable examples to explain.



2. Explain the hybridisation, geometry, magnetic properties, IUPAC nomenclature and comparative stability of $\left[Fe(CN)_6\right]^{4-}$ and $\left[Fe(CN)_6\right]^{3-}$.



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

Explain why?



- **4.** Giving a stuitable exmaple for each, explain the following.
- (i) Linkage isomerism (ii) Chelating ligand



5. $\left[Co(NH_3)_6\right]^{3+}$ and $\left[CoF_6\right]^{3-}$ is contain cobalt in +3 oxidation state, but $\left[Co(NH_3)_6\right]^{3+}$ is diamagnetic while $\left[CoF_6\right]^{3-}$ is paramagnetic with magnetic moment 4.90 BM. Explain.



6. $\left[Co(NH_3)_6\right]^{3+}$ and $\left[CoF_6\right]^{3-}$ is contain cobalt in +3 oxidation state, but $\left[Co(NH_3)_6\right]^{3+}$ is diamagnetic while $\left[CoF_6\right]^{3-}$ is paramagnetic with magnetic moment 4.90 BM. Explain.

