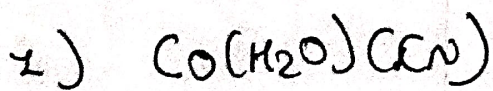


Section - B



$$\text{H}_2\text{O} \Rightarrow 0$$

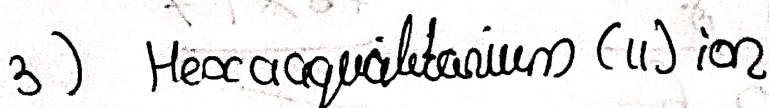
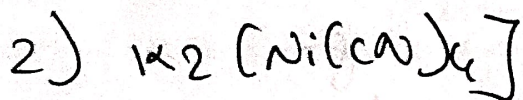
$$\text{CN}^- \Rightarrow -1$$

$$\text{Co} \Rightarrow x$$

$$x + 4(0) + 2(-1) = +1$$

$$x - 2 = +1$$

$$\boxed{x = +3}$$

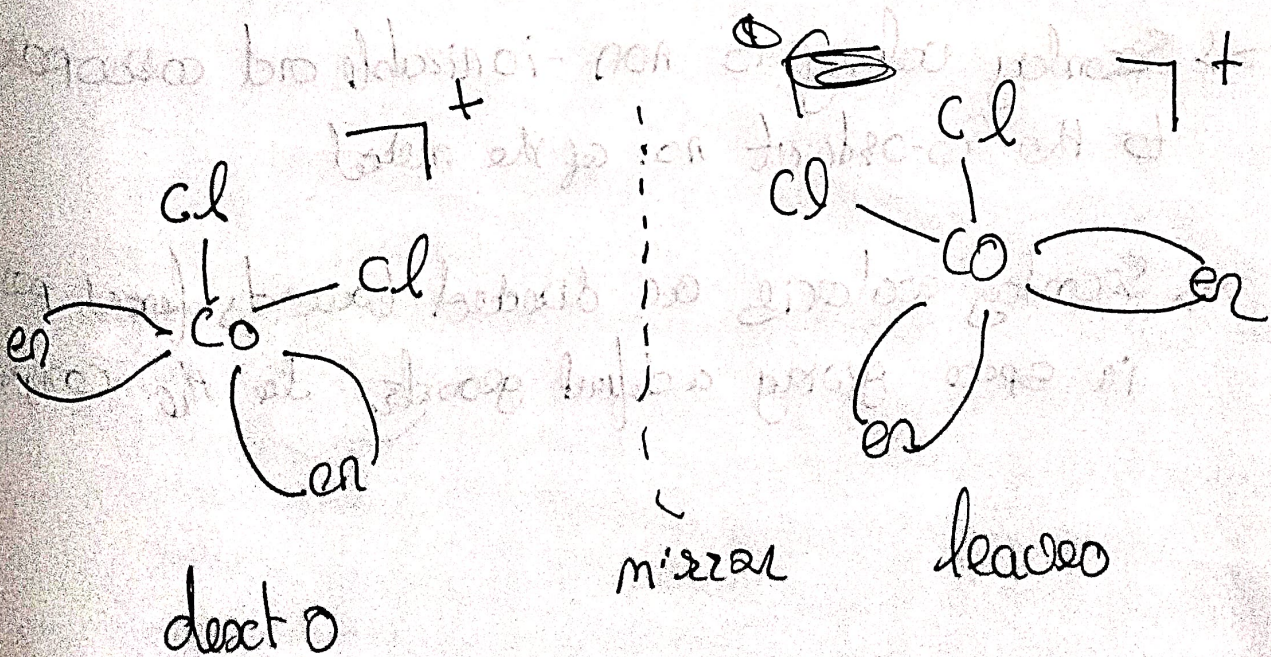
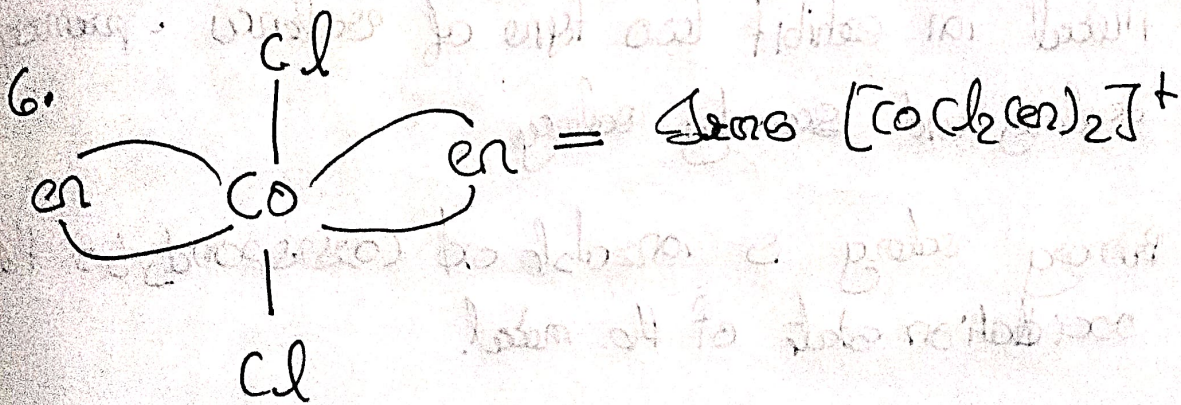


4) Metal carbonyl orbital synergistic bonding involving both σ and π interaction

Section - (

5. Limitations of valence bond theory include its inability to explain the color of complex,

- (ii) the quantitative interpretation of magnetic data
- (iii) the thermodynamic or kinetic stability of Co-ordinate compound.
- (iv) the distribution in shape of certain complex.



7.

i) Potassium tetrakis (oxalato) chromate(III)

ii) dichlorobis (ethylenediamine) cobalt(III) chloride

8) i) $[Cr(Ox)_3]^{3-} \Rightarrow 1$

ii) $[Co(NH_3)_3Cl_3] \Rightarrow 2$

9) Alfred Werner's theory of coordination compound has the following main postulates.

\Rightarrow Metal ion exhibit two type of valency - primary valency and secondary valency.

\Rightarrow Primary valency is ionizable and corresponds to the oxidation state of the metal.

\Rightarrow Secondary valency is non-ionizable and corresponds to the co-ordinate no. of the metal.

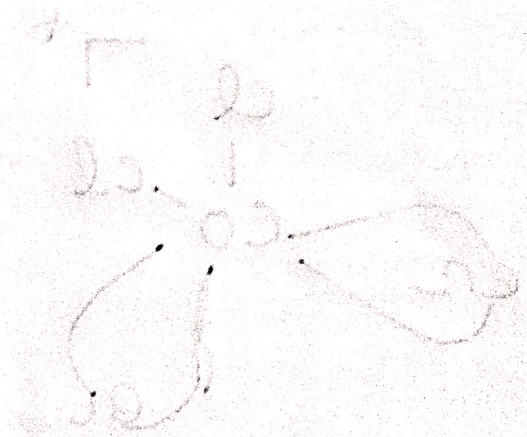
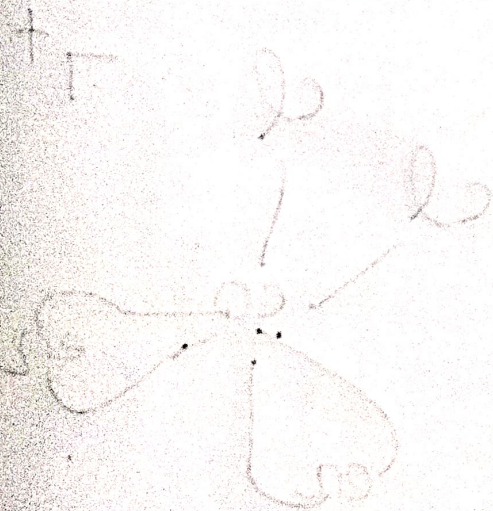
\Rightarrow Secondary valencies are directed towards fixed points in space giving a definite geometry to the complex.

Section - D

20. Primary valence and secondary valence are concepts proposed by Alfred Werner.

⇒ Primary valence: This is the ionizable valence of the metal and is satisfied by negative ions. It is equal to the oxidation state of the central metal atom or ion.

⇒ Secondary valence: This is the non-ionizable valence of the metal and is satisfied by neutral molecules or negative ions. It is equal to the co-ordination number of the central metal atom or ion and determines the geometry of the complex.



tetrahedral

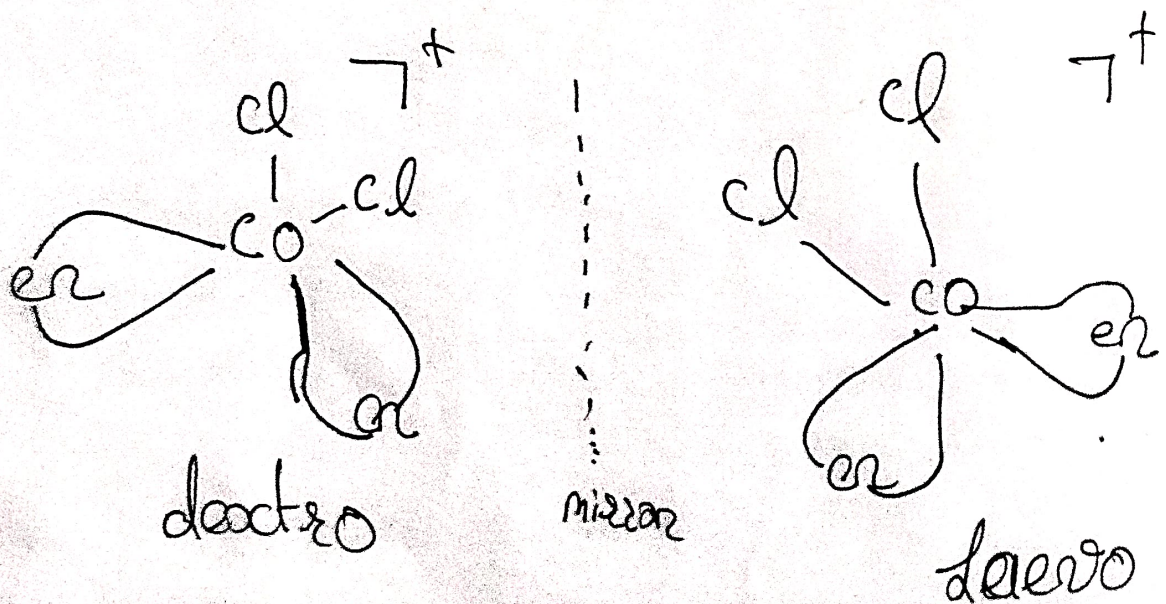
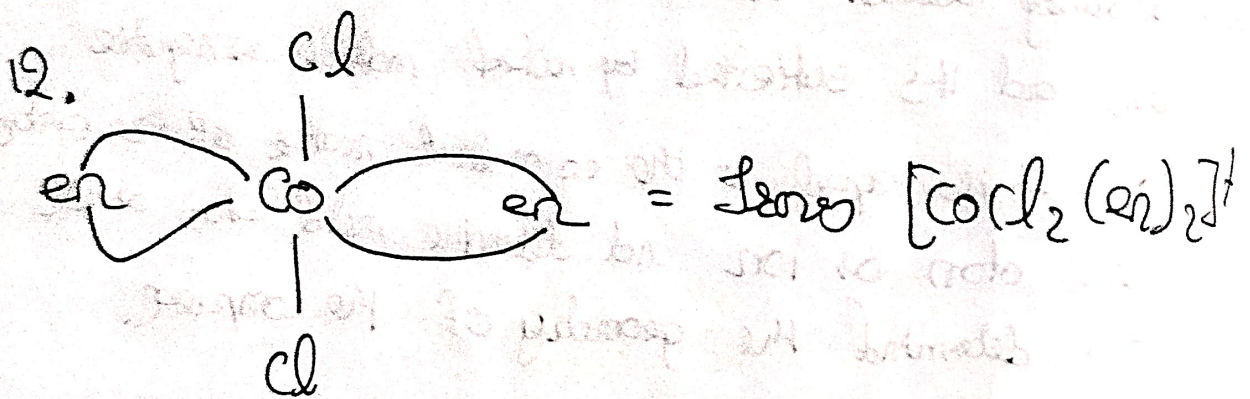
tetrahedral

octahedral

tetrahedral complex has tetrahedral geometry

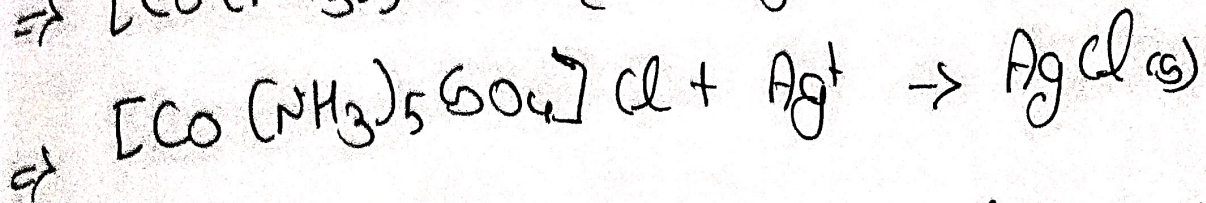
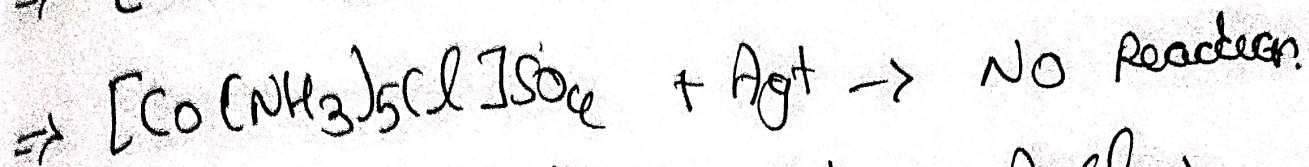
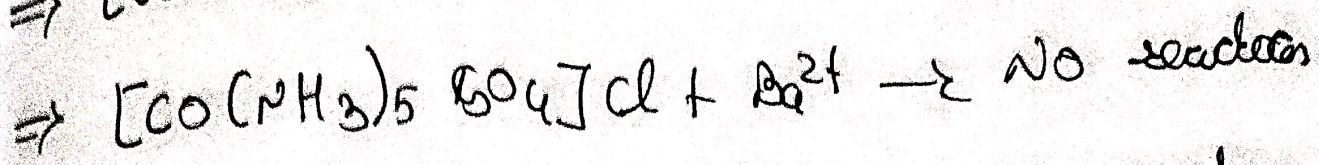
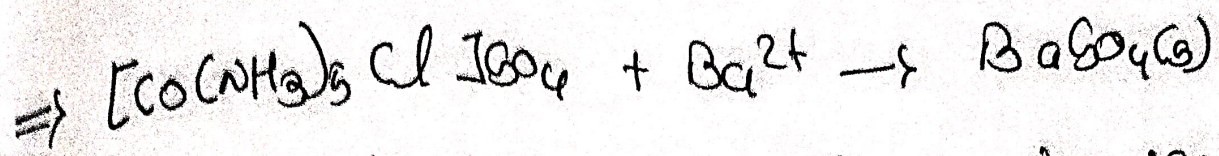
Section - 6

11. The complex $[Ti(H_2O)_6]^{3+}$ is violet because the d^1 ion has a $d-d$ configuration. In the octahedral field, the electron absorbs light in the green yellow region of the spectrum to undergo a $d-d$ transition from the t_{2g} to e_g orbital. The transmitted light is violet the complementary color.



cis $[CoCl_2(en)_2]^+$ optical isomers (d and l)

b) Ionization isomers dissolve in water to give different ions, hence giving different reactions with different reagents.



* Since the given complex give different reactions with reagents, hence they are ionization isomers.