

will be the principal functional group and rest as the substituents

$$O = \begin{array}{c} 1 & 2 & 3 \\ C - CH - CH - CH_2 \\ I & I \\ OH & NH_2 & OH \end{array}$$

2-amino-3-hydroxy propanoic acid

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(a)

(a)

**(b)** 

(c)

A transition metal complex absorbs visible light only when it has unpaired electron.  $Ni^{2+}$  in strong field ligand has configuration as



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The directive influence order and tendency to release electron for *o*-and *p*-directing group is,

 $O_2^- > NR_2 > NHR > NH_2 > OH > OCH_3 \approx NHCOCH_3 > CH_3 > X$ 

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(a) In  $MnO_2$ , FeCl<sub>3</sub>oxidation states of Mn and Fe are +4 and +3 respectively.

(b) In  $(MnO_4)^-$ ,  $CrO_2Cl_2$  oxidation states of Mn and Cr are +7 and +6 respectively.

(c) In  $[Fe(CN)_6]^{3-}$ ,  $[Co(CN)_3]$  oxidation states of Fe and Co are +3 and +3 respectively.

(d)  $[NiCl_4]^{2-}$ ,  $[CoCl_4]^{-}$  oxidation states of Ni and Co are +2 and +3 respectively.

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[M(abcd)] complex is square planar so will have three geometrical isomers.

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This is carbylamines reaction.



14 **(a)** 

An experimental fact depending upon the ability of the ligand to cause crystal field splitting (*i.e.*, strength of ligand).

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 $K_3[Fe(CN)_5NO]$ 

Potassium pentacyanonitrosyl ferrate (II).

16 **(c)** 

(d)

The *d*-*d* excitation is responsible for colour of  $Ti(H_2O)_6^{3+}$  which has one unpaired electron.

17 **(c)** 

(b)

The oxidation number of Fe in  $K_4$  [Fe(CN)<sub>6</sub>]is +2.

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Both Ag and Au are extracted by complex formation method. **(b)** 

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 $EAN = 24 - 3 + 2 \times (6) = 33.$ 

ANSWER-KEY										
<b>Q.</b>	1	2	3	4	5	6	7	8	9	10
<b>A.</b>	D	С	D	В	D	Α	В	С	Α	Α
<b>Q</b> .	11	12	13	14	15	16	17	18	19	20
<b>A.</b>	B	С	В	Α	D	С	С	Α	B	В

