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(b) Macrocyclic Effect

(c) Werner's theory of Coordination Compounds

(d) Factors affecting stability of the complexes

(5,5,5)

[This question paper contains 8 printed pages.]

Your Roll No.....

Sr. No. of Question Paper: 5526

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Unique Paper Code

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Name of the Paper

: DSC 10: Coordination Chemistry

and Reaction Mechanism

Name of the Course

: B.Sc. (Hons) Chemistry

(NEP-UGCF-2022)

Semester

: IV

Duration: 3 Hours

Maximum Marks: 90

Instructions for Candidates

- 1. Write your Roll No. on the top immediately on receipt of this question paper.
- 2. Attempt six questions in all.
- 3. Question 1 is mandatory.
- 4. All questions carry equal marks.

- 1. Explain the following giving appropriate reasoning:
 - (a) 4d and 5d elements form low spin octahedral complexes.
 - (b) Square planar complexes do not exhibit optical isomerism.
 - (c) [PtF₆] is stable, whereas NiF₆ does not exist.
 - (d) Fe(II) and Fe(III) form complexes with CN- but not with NH₃.
 - (e) The crystal field splitting in tetrahedral complexes is smaller than in octahedral complexes.

(3,3,3,3,3)

- (a) Name the following complexes according to the IUPAC system of nomenclature.
 - (i) $[Pt(NH_3)_4]$ $[PtCl_4]$

- (c) The formation of $[CdBr_4]^{2-}$ from $[Cd(H_2O)_6]^{2+}$ exhibits the successive equilibrium constants K_1 , K_2 , K_3 and K_4 as 1.56, 0.54, 0.06 and 0.37, respectively. Explain why K_4 is larger than K_3 . (5,5,5)
- 7. (a) Cr(II) fluoride and Mn(II) fluoride, both have a central metal ion surrounded by six F-ions. The Mn-F bond lengths are equidistant but four of the Cr-F distances are long and two are short. Explain.
 - (b) What are labile and inert complexes? Explain giving one example each, on the basis of CFT.
 - (c) Differentiate between inner orbital and outer orbital octahedral complexes. (5,5,5)
- 8. Write short notes on (Any three):
 - (a) Crystal Field theory and its limitations

- (b) Describe the factors responsible for strong distortion in the octahedral complexes. Square planar complexes are a special case of octahedral geometry. Justify your answer.
- (c) (i) I and CO have higher trans effects than Cl. Explain.
 - (ii) Predict the products when $[PtCl_4]^{2-}$ is treated with NH₃ followed by C_2H_4 . (5,5,5)
- 6. (a) On the basis of VBT, account for the magnetic properties of $[Ni(NH_3)_6]^{2+}$ and $[Cr(CN)_6]^{3-}$.
 - (b) Determine the CFSE of a d^6 octahedral complex having $10Dq = 25000 \text{ cm}^{-1}$ and $P = 15000 \text{ cm}^{-1}$ (mean pairing energy).

- (ii) $K_3[Fe(CN)_6]$
- (iii) $[Co(NH_3)_3(NO_2)_3]$
- (iv) $[(H_3N)_5Co-NH_2-Co(NH_3)_4(H_2O)]Cl_5$
- (v) $[Co(en)_3]Cl_3$
- (b) What is the chelate effect? Chelate effect is predominantly due to entropy change. Explain.
- (c) Which of the following have higher Δ_0 value and why?
 - (i) $[Fe(H_2O)_6]^{3+}$ or $[Fe(CN)_6]^{3-}$
 - (ii) $[Cr(NH_3)_6]^{2+}$ or $[Cr(NH_3)_4]^{2+}$ (5,5,5)

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- 3. (a) Write the formulae of the following complexes according to the IUPAC system of nomenclature:
 - (i) pentaamminenitrito-O cobalt(III) sulphate
 - (ii) potassium amminedicyanidodioxidoperoxidochromate(VI)
 - (iii) μ-amido μ-hydroxido bis[tetraammine-cobalt(III)]
 - (iv) tetrapyridineplatinum(II) tetrachloridoplatinate(II)
 - (v) Potassium tetrafluridoargentate(I)
 - (b) Define the terms transition state and intermediate state using reaction pathways.
 - (c) What is the effect of π acceptor ligand and π donor ligands on Δ_0 ? Explain on the basis of ligand field theory. (5,5,5)

- 4. (a) Explain why square planar complexes of Pt(II) often undergo associative substitution mechanisms, while octahedral complexes of Cr(III) typically undergo dissociative mechanisms.
 - (b) Draw and explain Crystal Field Splitting diagram for octahedral complexes.
 - (c) How will you distinguish between the following pairs of isomers?
 - (i) cis and trans [Pt(NH₃)Cl₂]
 - (ii) $[Co(NH_3)_5SO_4]$ I and $[Co(NH_3)_5I]$ SO_4 (5,5,5)
- (a) Predict and sketch all the possible isomers of [Cr(gly)₃].