



ALUPE UNIVERSITY
COLLEGE

... Bastion of Knowledge ...

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OFFICE OF THE DEPUTY PRINCIPAL
ACADEMICS, STUDENT AFFAIRS AND RESEARCH

UNIVERSITY EXAMINATIONS

2019/2020 ACADEMIC YEAR

THIRD YEAR SECOND SEMESTER REGULAR

EXAMINATION

FOR THE DEGREE OF BACHELOR OF
EDUCATION SCIENCE

COURSE CODE: CHE 318

COURSE TITLE: COORDINATION CHEMISTRY

DATE: 3RD NOVEMBER, 2020 TIME: 0900 – 1200 HRS

INSTRUCTION TO CANDIDATES

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THIS PAPER CONSISTS OF 4 PRINTED PAGES

PLEASE TURN OVER

REGULAR – MAIN EXAM**CHE 318: COORDINATION CHEMISTRY****STREAM: BED (Science)****DURATION: 3 Hours****INSTRUCTIONS TO CANDIDATES**

- i. Answer **ALL** questions.
- ii. Diagrams may be used whenever they serve to illustrate the answer.

Question One

- a) Define the following terms as applied in coordination chemistry.
- i) Coordination compound (1 Mark)
 - ii) Hard acid (1 Mark)
 - iii) Coordination number (1 Mark)
 - iv) Ligand (1 Mark)
 - v) Nephelauxetic effect (1 Mark)
- b) Draw the shapes of the d orbitals indicating clearly their electron densities distribution with respect to x, y and z axes (2 Marks)
- c) The table below shows some properties of chromium ammonate chlorides. Study it and answer the questions that follow;

Compound	Conductivity
$\text{CoCl}_3 \cdot 6\text{NH}_3$	High
$\text{CoCl}_3 \cdot 5\text{NH}_3$	Medium
$\text{CoCl}_3 \cdot 4\text{NH}_3$	Low
$\text{CoCl}_3 \cdot 3\text{NH}_3$	Low

- i) Determine the primary and secondary valences of chromium in the compounds (1 Mark)
- ii) Draw Werner's representations of the compounds and comment on the trend of conductivity of the compounds (4 Marks)
- iii) What is the effective atomic number of Co in the compounds? (1 Mark)

Question Two

- a) Molecular orbital theory is superior to crystal field theory in accounting for bonding in metal complexes. Give three reasons to defend this statement (3 Marks)
- b) Give two advantages and two disadvantages of the crystal field theory (4 Marks)

- c) Show how the d orbitals are perturbed in a square planar field (1 Mark)
- d) State and explain three factors that determine the magnitude of $\Delta_{\text{splitting}}$ (3 Marks)
- e) The $[\text{Cr}(\text{H}_2\text{O})_6]^{3+}$ is paramagnetic with $\mu_{\text{spin-only}}$ value of 2.83 Bohr Magneton. Is the complex a high-spin (spin free/weak field) or low-spin (spin paired/Strong field)? Briefly explain your answer (3 Marks)

Question Three

- a) Write short notes on the following types of isomerism in co-ordination compounds giving one example in each case:
- Ionization isomerism (2 Marks)
 - Linkage isomerism (2 Marks)
 - Geometric isomerism (2 Marks)
- b) The complex $[\text{RhF}_6]^{3-}$ is octahedral and paramagnetic with a calculated $\mu_{\text{spin-only}}$ value of 4.90 BM. Account for the bonding in this complex using the valence bond theory (3 Marks)
- c) Describe the type of electronic transitions in metal complexes (3 Marks)
- d) Describe the three special features in an electronic spectrum (3 Marks)

Question Four

- a) What is the experimental evidence for the Jahn-Teller distortions in a metal complex? (1 Mark)
- b) Which of the following octahedral complexes will undergo Jahn-Teller distortion? Briefly explain your answer
- Low spin Fe^{2+} (1 Mark)
 - Cu^{2+} (1 Mark)

(Atomic numbers: Fe = 26, Cu = 29)

- c) Work out the ground state term symbols for the following ions
- High spin octahedral Mn^{2+} (1 Mark)
 - Octahedral Cr^{3+} (1 Mark)
- d) Calculate the crystal field stabilization energy (CFSE) of a d^6 complex in,
- Strong field (1 Mark)
 - Weak field (1 Mark)

- e) Briefly defend the following scientific observations
- a) Δ_{oct} is greater than Δ_{tert} (2 Marks)
- b) Square planar Au^{2+} complexes readily oxidize to Au^{3+}
[Hint: Au^{2+} is a d^9] (2 Marks)
- c) Low spin octahedral d^6 complexes are more stable than high spin complexes. (2 Marks)
- d) d-d electronic transition in $(\text{Co}(\text{H}_2\text{O})_6)^{2+}$ is spin allowed while in high-spin $[\text{Mn}(\text{H}_2\text{O})_6]^{2+}$ is spin-forbidden (2 Marks)

Question Five

- a) Draw the structure of the following ligands and indicate whether they are mono or polydentate
- i) Oxalato (1 Mark)
- ii) Pyridine (1 Mark)
- iii) EDTA (1 Mark)
- b) Give the systematic I.U.P.A.C names of the following complexes
- i) $[\text{Co}(\text{NH}_3)_6][\text{Cr}(\text{CN})_6]$ (1 Mark)
- ii) $[\text{Pt}(\text{NH}_3)_4][\text{PtCl}_4]$ (1 Mark)
- c) Explain briefly any two rules of an electronic transition (2 Marks)
- d) State two applications of co-ordination compounds (1 Mark)
- e) Account for the following observations
- i) $[\text{Ni}(\text{Cl})_4]^{2-}$ is paramagnetic but $[\text{Ni}(\text{CN})_4]^{2-}$ is diamagnetic (4 Marks)
- ii) In $[\text{Cr}(\text{H}_2\text{O})_6]^{2+}$, two of Cr-OH₂ bonds are longer than others (1 Mark)
