Ex/SC/CHEM/UG/CORE/TH/12/2024

B. Sc. CHEMISTRY EXAMINATION, 2024

(5th Semester)

CHEMISTRY (CORE)

PAPER: CORE 12

Time: Two Hours Full Marks: 40

(20 marks for each Unit)

Use a separate answer scripts for each Unit.

The figures in the margin indicate full marks.

UNIT-5121 - O

- 1. (a) Synthesize anthracene by Diels–Alder reaction involving 1,4-naphthaquinone and 1,3-butadiene. 1½
 - (b) Predict the product(s) with plausible mechanism of the following reactions (any two): 1½×2

(iii)
$$\frac{(1) \operatorname{Br}_2/\operatorname{CCl}_4}{(2) \text{ Heated with alkali}}$$

[Turn Over]

CHEM-**12**

- (a) Answer **any one** of the following questions: $1\frac{1}{2}$
 - (i) Pyrrole undergoes electrophilic attack at C-2 position whereas indole undergoes electrophilic attack at C-3 position justify the statement.
 - (ii) Furan undergoes Diels-Alder reaction with maleic anhydride whereas pyrrole does not. Explain.
- (b) Predict the product(s) of the following reactions with plausible mechanism (any two): 2×2

(i) OH
$$\frac{1. \text{ Br}_2/\text{MeOH}}{2. 1-2\% \text{ H}_2\text{SO}_4}$$

3. 50 °C

(ii)
$$N$$
 CHCl₃ NaOH

(iii)
$$\xrightarrow[\text{H}_2\text{O}, \text{ heat}]{\text{CH}_2\text{O}, \text{ Me}_2\text{NH}} \times [A] \xrightarrow{\text{1. MeI}} \times [E]$$
H

$$3. \text{ H}_3\text{O}^+$$

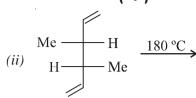
(iv)
$$\frac{\text{NaNH}_2, \text{PhNH}_2}{100 \text{ °C}}$$

(5)

- (iii) Although 'Dewar benzene' is less stable by 60 kcal than its isomer benzene, its conversion into benzene is surprisingly slow (E_{act} is 37 kcal). Explain this observation.
- **5.** Answer *any five* of the following questions: 2×5
 - (a) Explain the stepping-up or stepping-down methods of aldoses.
 - (b) Explain Lobry de Bruyn-van Ekenstein rearrangement.
 - (c) What is anomeric effect? Explain.
 - (d) What is isoelectric point? How is it measured for amino acids with neutral, acidic and basic side chain residue?
 - (e) Explain the Strecker and Azlactone method of synthesis of α -amino acid.
 - (f) Explain Fischer glycosidation procedure.
 - (g) Write down the structure of four different nucleobases found in DNA and the H-bonding pattern found in Watson-Crick model of DNA.







COOMe
$$(iii) \bigcirc O + | | | (a) \text{ heat}$$

$$COOMe \bigcirc (b) \text{ H}_2/\text{Pd-C}$$

$$COOMe \bigcirc (c) \text{ heat}$$

$$(iv) \qquad \stackrel{\text{Me}}{\longleftarrow} \qquad \text{hv} \rightarrow \qquad \qquad \text{Me}$$

(b) Answer *any two* of the following questions: 2×2

(i) Identify [C] & [D] in the following reaction sequence. Explain with mechanism.

$$R \xrightarrow{O} \underbrace{MgBr, THF}_{work up} [C] \xrightarrow{KH in THF}_{\Delta} [D]$$

(ii) Heating of indene derivative [E] causes the scrambling of the labeled deuterium atom to all the three non-aromatic positions. Explain mechanistically the above observation.

[Continued]

(3)

- **3.** (a) Describe schematically resolution of *racemic*-2-aminobutane using (S)-2-hydroxy-2-phenylethanoic acid as resolving agent. Give also the chemical reaction(s) involved.
 - (b) Draw the most stable conformation of cis- and trans 1,2-dimethylcyclohexane in Newman projection.
 Compare their stability with justification.
 - (c) What is/are the product(s) obtained on acetolysis of optically pure *trans*-2-acetoxycyclohexyl tosylate? Give suitable mechanistic and stereochemical explanation.

 $2\frac{1}{2}$

(d) How many pairs of enantiomers are possible for 2-isopropyl-5-methylcyclohexanol? Draw its most stable enantiomeric pair in chair form. What happens when any one of these two enantiomers is treated with sodium hydride/carbon disulfide followed by methyl iodide and then the product(s) obtained is/are strongly heated? Explain this phenomenon with appropriate mechanistic and stereochemical interpretation.

UNIT-5122 - O

4. (a) Predict the product(s) with proper stereochemistry of the following reactions and explain with FMO consideration (Answer *any three*): 2×3

(i)
$$H_3PO_4 \rightarrow Me$$
 Me Me

CHEM-**12**

[Turn Over]