(b) Identify the reagents and intermediate in the following reaction. Propose a mechanism for the second step.

1+1+1+2=5

(c) What product is expected to be formed in the following reaction?

$$\bigcirc + \stackrel{Cl}{\swarrow} \xrightarrow{AlCl_3} ?$$

Write the name of the above reaction and propose a mechanism, clearly explaining the steps involved.

1+1+3=5

(d) Write the steps involved in a E2 mechanism. Provide one evidence in favour of E2 mechanism. Under what condition E2 is favoured over E1 mechanism? 2+1+2=5

4

- (e) (i) State one method by which carbocations can be generated.
  - (ii) Account for the stability of a benzyl cation.
    - (iii) Why is it difficult to form carbocations at bridgehead positions? 1+2+2=5
- 4. Answer any three questions: 10×3=30
  - (a) (i) Toluene undergoes benzylic bromination when heated with NBS. Write the product obtained in the reaction Propose a mechanism for the reaction.

1+4=5

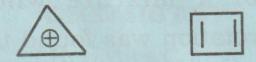
- (ii) Why are terminal alkynes acidic?
  Write the reaction involved in the conversion of propane to pent-2-yne.

  1+2=3
- (iii) How can you convert propyne to propan-2-one?

- (b) Give the 1,2- and 1,4- products of the addition of one equivalent of HBr to 2,4-hexadiene. Draw the transition states involved and predict which of them would be the major product and which will be the minor product. What are the 1,2- and 1,4- addition products of HBr to 2-methyl-1, 3-cyclohexadiene? What is about the products of 1,2- and 1,4-addition of HX to an unsubstituted cyclic-1, 3 diene?
- (c) (i) What do you mean by a racemic mixture? Why is resolution of a racemic mixture a difficult process? How can you resolve a recemic mixture? Suggest one method.

  1+1+3=5
  - (ii) The addition of HBr to propene is regioselective. Write the reaction involved. Propose a mechanism to justify the regioselectivity. 1+4=5

(d) (i) State whether the following compounds are aromatic, non-aromatic or antiaromatic. Give reasons.



- (ii) Define hyperconjugation. How many hyperconjugation structures are possible for an isopropyl radical?
- (iii) What do you mean by partial bond fixation? Which position of anthracene is attacked by electrophiles and why?

  1+(1+2)=4
- (e) (i) Explain Baeyer strain theory. 2
  - (ii) Draw the energy profile diagram of cyclohexane.
  - (iii) Why is the chain form of cyclohexane the most stable? 1
  - (iv) Convert meso-tartaric acid from Fischer to Newman projection and Sawhorse projection. 2

(v) Let (S)-2-bromobutane have a specific rotation of +23.1° and (R)-2-bromobutane have a specific rotation of -23.1°. What is the percentage purity and % composition of a mixture whose specific rotation was found to be +18.4°?

2

(f) (i) What are the factors which determine whether an aliphatic nucleophilic substitution reaction proceeds by SN1 or SN2 reaction?

Discuss any two factors in brief.

1+4=5

(ii) Using appropriate example write briefly about Saytzeff and Hofmann elimination.

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3 (Sem-2/CBCS) CHE HC 1

## 2023

## CHEMISTRY

(Honours Core)

Paper: CHE-HC-2016

(Organic Chemistry-I)

Full Marks: 60

Time: Three hours

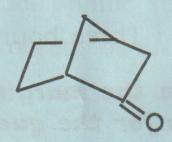
The figures in the margin indicate full marks for the questions.

1. Answer all the questions:

 $1 \times 7 = 7$ 

- (a) Draw the orbital diagrams of singlet and a triplet carbon.
- (b) Write the structure of (R,R)-tartaric acid.
- (c) Draw the Newman projection formula of the lowest energy conformer of butane.

- (d) Between ammonia and trimethylamine which one is more likely to favour elimination over substitution and why?
- (e) Define a meso compound.
- (f) Is cyclopentadiene acidic? Give reasons.
- (g) Write the IUPAC name of the following compound:



- 2. Answer all the questions: 2×4=8
  - (a) What product is obtained when cyclohexane is subjected to ozonolysis?

    Write the reaction involved.
  - (b) Suggest two ways by which you can convert OH group into good leaving group.

    1+1=2
  - (c) Invoking hybridisation, explain the structure of methyl free radical.

(d) Label the following as E- or Z-isomer:

(i) 
$$H \sim CH_2CH_2OH$$

$$H_3CH_2CH_2C \qquad Cl$$

(ii) 
$$NH_2$$
  $CO_2H$   $CH_2OH$ 

- 3. Answer any three questions: 5×3=15
  - (a) When 1-iodo-1-methylcyclohexane is treated with  $NaOCH_2CH_3$  as the base, the more highly substituted alkene product predominates. When  $KOC(CH_3)_3$  is used as the base, the less highly substituted alkene predominates. Write the reactions, giving the structure of the two products and offer an explanation. 3+2=5