(8 pages) **Reg. No.:** .....

Code No.: 5884 Sub. Code: PCHM41

M.Sc. (CBCS) DEGREE EXAMINATION, APRIL 2020.

Fourth Semester

Chemistry — Core

## ORGANIC CHEMISTRY — IV

(For those who joined in July 2017 onwards)

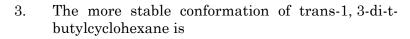
Time: Three hours Maximum: 75 marks

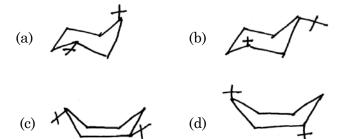
PART A —  $(10 \times 1 = 10 \text{ marks})$ 

Answer ALL questions.

Choose the correct answer:

- 1. In Reimer-Tiemann reaction product formation takes place by which reaction/s:
  - (a) A<sub>r</sub> SE
- (b) SN reaction
- (c) Elimination
- (d) All of these
- 2. Wittig reaction is given by
  - (a) Only 1°-halide
- (b) Only 2°-halide
- (c) Only 3°-halide
- (d) 1° and 2° halides





4. The number of butane gauche interaction Cis-decalin is

(a) 2

(b) 3

(c) 4

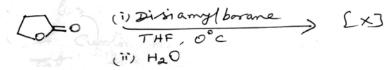
(d) 5

5. The synthetic equivalent for  $R^{\Theta}$  is

- (a) Rmg Br (b) R CH
- (c) R IN OFE (d) R

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- 6. Which one of the following compounds act as blocking group for amines?
  - (a) Benzyloxy carbonyl chloride
  - (b) 9-Fluorenyl methyloxy carbonyl chloride
  - (c) Benzoyl chloride
  - (d) All of these
- 7. In the reaction sequence



[X] will be

- $\begin{array}{cccc} \text{(a)} & CH_2-CH_2 \\ & | & | \\ & CHO & CHO \end{array}$
- $\begin{array}{cccc} \text{(b)} & CH_2 & & CH_2 \\ & | & | \\ & CH_2OH & CH_2OH \end{array}$
- $\begin{array}{cccc} \text{(c)} & CH_2 & & CH_2 \\ & | & | \\ & CH_2OH & CHO \end{array}$
- (d) HC = CH / $H_2COH$  CHO

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## 8. In the reaction sequence

Product will be

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9. The functional groups present in oestrone are

10. In the given reaction

[X] will be

- (a) 7-Dehydrocholesterol
- (b) Ergocalciferol
- (c) Vitamin D<sub>5</sub>
- (d) Vitamin D<sub>3</sub>

PART B — 
$$(5 \times 5 = 25 \text{ marks})$$

Answer ALL questions, choosing either (a) or (b).

Each answer should not exceed 250 words.

11. (a) Discuss the mechanism of

- (i) Julia olefination
- (ii) Mc Murry coupling

Or

(b) Illustrate Bamford-Stevens reaction with suitable examples.

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- 12. (a) (i) State Curtin-Hammett principle. Give its limitations.
  - (ii) What is conformational free energy of difference?

Or

- (b) Discuss the stability of Cis and Trans decalins.
- 13. (a) Discuss the role of activating groups in organic synthesis.

Or

- (b) Give an account of functional group interconversion.
- 14. (a) Reaction with OsO<sub>4</sub> is the best method for Cis-perhydroxylation of alkenes. Explain with mechanism.

Or

- (b) Narrate the mechanism of Suzuki coupling.
- 15. (a) Give an account on prostoglandins.

Or

(b) Bring out the relationship between oestrone, oestradiol and oestriol and also state how they are interconvertible.

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## PART C — $(5 \times 8 = 40 \text{ marks})$

Answer ALL questions, choosing either (a) or (b).

Each answer should not exceed 600 words.

- 16. (a) Write short notes on:
  - (i) Acyloin condensation
  - (ii) Baeyer-Villiger oxidation
  - (iii) Pschorr reaction

Or

- (b) Describe the reaction mechanism for halolactorisaton and Gomberg-Bechmann reactions.
- 17. (a) Explain how conformational features affect the reactivity of cyclohexyl systems in the following reactions:
  - (i) Oxidation of alcohols
  - (ii) Elimination of reactions

Or

- (b) (i) Give the preferred conformation of trans-syn-trans, cis-anti-cis and trans-anti-trans perhydrophenanthrenes.
  - (ii) Trans 4-t-butylcyclohexane carboxylic esters hydrolyse faster than the corresponding cis-esters. Explain.

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- 18. (a) (i) Analyse the synthetic plan for preparing 2, 4-dimethyl-2-hydroxypentanoic acid.
  - (ii) Bring out the importance of protecting groups in organic synthesis.

Or

- (b) (i) Give an account on Robinson annulations reaction.
  - (ii) Write about the Retro-synthetic scheme for Cis-Jasmone.
- 19. (a) Indicate the applications of the following reagents in organic synthesis:
  - (i) DDQ,
  - (ii) 9-BBN

Or

- (b) Describe any four synthetic uses of R<sub>u</sub>O<sub>2</sub>.
- 20. (a) How will you confirm the following in the structure of cholesterol?
  - (i) Presence of OH group.
  - (ii) Nature and position of side chain.

Or

- (b) (i) Formulate the conversion of cholesterol into progesterone.
  - (ii) Discuss the conformational structure of coprostane.

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