Reg. No. : ..... Name : .....

III Semester M.Sc. Degree (CBSS – Reg./Suppl./Imp.) Examination, October 2021 (2018 Admission Onwards) CHEMISTRY CHE3C.09 : Organic Chemistry - III

one mark.)

Time: 3 Hours

Max. Marks: 60

## (Answer all questions in one word or one sentence. Each question carries

SECTION - A

 Out of cis- and trans- isomer of cinnamic acid which isomer absorbs at higher wavelength?

- In an organic compound, there is no absorption in the region of 1600 cm<sup>-1</sup>, 1580 cm<sup>-1</sup> and 1500 cm<sup>-1</sup>. This gives a sure proof absence of which
- functional group? 3. Write the multiplicity of signals of CH<sub>s</sub> – CH<sub>s</sub> – NH<sub>s</sub> in <sup>1</sup>H NMR spectroscopy.
- How will you account triplet splitting pattern of CDCl<sub>s</sub> at δ 76, 77, 78 ppm in
- 13C NMR ? 5. Write the source of the electrons, which required for the bombardment in El
- 6. What do you mean by Nitrogen rule? 7. By which name reaction oxetanes can be prepared?
- Write the structure of 1,2,4-triazine.

Mass spectrometry .

- SECTION B
- (Answer any eight questions. Answer may be in two or three sentences. Each question carries 2 marks.)

 $(8 \times 1 = 8)$ 

## 9. How will you account the difference in the value of $\lambda_{max}$ in the following two

compounds? Discuss in detail. a) CH<sub>2</sub>CI – 173 nm b) CH<sub>a</sub>I - 258 nm.

P.T.O.

# 11. Calculate the $\lambda_{\text{max}}$ of following compound by applying Woodward Fieser rule.

explanation.

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12. On oxidation 2-propanol will be converted to propanone. How can you monitor

13. P-OMeC<sub>6</sub>H<sub>4</sub>-Me shows in its <sup>1</sup>H NMR spectrum two peaks at δ (ppm) 2.34 and 3.75. Assign these two signals and comment about the signal positions.

the progress of this reaction by IR spectroscopy?

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10. If we add 20 ml of CCl₂ to 4 ml of ethyl alcohol how the position of O-H stretching frequency of ethyl alcohol is varied from the pure state? Give a suitable

- 14. What is the order of chemical shift of protons attached to tertiary, secondary, primary carbons ? Explain why.
- How can you distinguish ethyl benzene from m-xylene by <sup>13</sup>C NMR spectroscopy? 16. The proton decoupled <sup>13</sup>C NMR spectrum of C<sub>6</sub>H<sub>3</sub>Br<sub>3</sub> gives only two signals. Write the suitable structure of the compound and explain why spectrum follows
- this pattern. 17. How will you differentiate 3-methylcyclohexene from 4-methylcyclohexene by mass spectrometry?

18. What is the importance of meta stable peaks? Illustrate with example.

19. Discuss a method for the synthesis of 1,2,4-triazole with reaction. (8×2=16) 20. Explain the method of synthesis of coumarin.

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22. How conjugation affect the stretching frequency of carbonyl compounds?

25. A compound with molecular formula C<sub>7</sub>H<sub>8</sub>O shows peaks at δ (ppm) 9.8, s, 1H

and 7.2, m, 5H. It is on reaction with hydroxylamine followed by P<sub>2</sub>O<sub>5</sub> resulted another compound with molecular formula C,H,N which showed an IR peak 2210 cm<sup>-1</sup> and a <sup>1</sup>H NMR signal at δ (ppm) 7.3, s, 5H. Deduce the structure of

and why? Write examples for important electrophilic substitution reactions in

SECTION - D

SECTION - C

(Short paragraph questions. Answer any four questions. Each question carries 3 marks.) 21. What are batho, hypso, hypo and hyper chromic shift of absorption bands in

UV-V is spectroscopy? Illustrate with examples.

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 $(4 \times 3 = 12)$ 

### Explain the usage of shift reagents in the simplification of NMR spectra. Discuss the working principle of FAB mass spectrometry.

imidazoles.

28. A) Discuss:

iii) CIDNP.

B) Discuss:

ii) DEPT spectroscopy

ii) FT NMR spectroscopy

Explain with different examples.

these two compounds writing the reaction scheme. 26. Which is the preferred position of electrophilic substitution reaction in imidazoles

(Essay type questions. Answer four questions. Each question carries 6 marks.) 27. A) Explain Woodward-Fieser rules in predicting the  $\lambda_{max}$  of dienes,  $\alpha$ , β-unsaturated carbonyl compounds and arenes with examples. OR B) Discuss the concept of group frequencies in IR spectroscopy for structural

elucidation illustrating with different classes of organic compounds.

i) Simplification of complex spectra by use of double resonance method

i) Simplification of complex spectra by using high field NMR

iii) Theory of spin-spin splitting citing suitable example.

OR

29. A) Deduce the structure and stereochemistry of the compound from the following spectral data. Explain the pattern of each and every signals with suitable

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explanation. i) UV - 284 nm, 308 nm ii) IR - 1690 cm-1

(d, J= 16 Hz, 1H), 9.75 (d, J = 8 Hz, 1H)

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iii) <sup>1</sup>H NMR -  $\delta$  (ppm) - 6.7 (dd, J= 16 Hz, 8 Hz, 1H), 7.40 (m, 5H), 7.45

iv) <sup>13</sup>C NMR - δ (ppm) - 128.2, 128.3, 128.8, 131.0, 134.0, 152.0, 193.0.

B) a) Deduce the structure of the compound from the following spectral data. Explain the pattern of each and every signals with suitable explanation. i) Molecular Formula- C, H, O,

iii) <sup>1</sup>H NMR - δ (ppm) - 6.8 -7.3 (m, 5H), 4.3 (t, 2H), 2.93 (t, 2H), 2.0 (s, 3H)

b) Fragmentation pattern in carbonyl compounds. 30. A) Discuss the structure, synthesis and reactions of

ii) IR- 1730, 3000 cm<sup>-1</sup>

v) Mass - m/z - 132, 131, 103

OR

iv) Mass - m/z - 73, 91, 149, 164

OR

- Thietanes ii) Oxadiazoles

ii) Pyrans

iii) Azepins.

iii) Quinolines

B) Discuss the structure, synthesis and reactions of

Selenophenes

 $(4 \times 6 = 24)$