UNIVERSITY OF SWAZILAND SECOND SEMESTER FINAL EXAMINATION 2014

| TITLE OF PAPER | : | Applied Spectroscopy |
|----------------|---|---------------------------------|
| COURSE NUMBER | : | C603 |
| TIME | : | Three Hours |
| INSTRUCTIONS | : | Answer any FOUR Questions. Each |

This Paper contains five (5) pages.

You must not open this paper until the Chief Invigilator so has granted permission to do.

SECTION A : MASS SPECTROMETRY AND INFRARED SPECTROSCOPY

Question 1

 (a) What fragments might you expect in the mass spectra of the following compounds? Explain how the fragments and their m/z values arise. [18 marks]



(b) Assume that you are in the laboratory carrying out the catalytic hydrogenation of cyclohexene to cyclohexane. Explain how you would use a mass spectrometer to determine when the reaction is finished? [7 marks]

Question 2

(a) Two infrared spectra are shown. One is the spectrum of cyclohexane, and the other is the spectrum of cyclohexene. Identify them and explain your answer. [12 marks]



(b) Assume that you are carrying out the base-induced dehydrobromination of 3-bromo-3methylpentane to yield an alkene. How could you use IR spectroscopy to tell which of the two possible elimination products is formed. [13 marks]

Question 3

(a) 4-Methyl-2-pentanone and 3-Methylpentanal are isomers. Explain how you could tell them apart both by mass spectrometry and by infrared spectroscopy. [18 marks]



(b) Assume that you are carrying out the dehydration of 1-methylcyclohexanol to yield 1methylcyclohexene. Explain clearly how you could use infrared spectroscopy to determine when the reaction is complete. [7 marks]

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SECTION B : NUCLEAR MAGNETIC RESONANCE (¹H AND ¹³C NMR SPECTROSCOPY)

Question 4

Briefly describe the following aspects of nuclear magnetic resonance (NMR) spectroscopy.

| i. | Theory | [3 marks] |
|------|--|-----------|
| ii. | Nature of NMR absorptions | [3 marks] |
| iii. | NMR spectra | [3 marks] |
| iv. | Operation of an NMR Spectrometer. | [3 marks] |
| v. | Spin-Spin Splitting in ¹ HNMR | [3 marks] |

What is the structure of a hydrocarbon that has $M^+ = 120$ in its spectrum and has the following ¹HNMR spectrum: 7.25d (5H,broad singlet); 2,9d (1H,septet, J=7Hz); 1.22d (6H, doublet, J= 7Hz) (10 Marks)

<u>Question 5</u>

Compound A, a hydrocarbon with $M^+ = 96$ in its mass spectrum, has the ¹³C spectral data given below. On reaction with BH₃ followed by treatment with basic H₂O₂, A is converted into B, whose ¹³C spectral data are also given below. Propose structures for A and B.

Compound A

[13 marks]

Broadband – decoupled ¹³C NMR : 26.8, 28.7, 35.7, 106.9, 149.7 &

Dept - 90: No peaks

Dept - 135 : No positive peaks; negative peaks at 26.8, 28.7, 35.7, 106.9 &

Compound B

[12 marks]

Broad band-decoupled 13 C NMR : 26.1, 26.9, 29.9, 40.5, 68.2 δ

Dept - 90 : 40.5 δ

Dept - 135 : positive peak at 40.5 8; negative peaks at 26.1, 26.9, 29.9, 68.2 8

Question 6

To answer the following questions, consider the data and H^1 NMR spectrum below. The mass spectrum of this compound shows a molecular ion at m/z = 113; the IR spectrum has characteristic absorption at 2270 and 1735 cm⁻¹, and the ¹³CNMR spectrum has five (5) signals.



- (a) Based on the mass spectral data and the IR data, what functional groups are present in this compound? [4 marks]
- (b) How many types of non-equivalent protons are there in this molecule? [4 marks]
- (c) Comment or describe the signal at 3.5 delta in terms of integration, splitting pattern and chemical shift. [4 marks]
- (d) Describe the signals at 4.3 delta and 1.3 delta in terms of their integration splitting and chemical shift. [4 marks]
- (e) What is the significance of ¹³C NMR data?

[4 marks]

(f) Analyze all the information deduced from the data provided and then propose a structure for this compound? [5 marks]

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