



**JARAMOGI OGINGA ODINGA UNIVERSITY OF SCIENCE &
TECHNOLOGY UNIVERSITY EXAMINATIONS 2013
4TH YEAR 1ST SEMESTER EXAMINATION OF BACHELOR OF
EDUCATION (SCIENCE)
REGULAR**

COURSE CODE: SCH 405

COURSE TITLE: SYNTHETIC ORGANIC CHEMISTRY

DATE: 12/8/13

TIME: 9.00 -11.00 AM

DURATION: 2 HOURS

INSTRUCTIONS

- 1. This paper contains five (5) questions.**
- 2. Answer question 1 (compulsory) and ANY other TWO questions.**
- 3. Write all answer in the booklet provided.**

SECTION A: ANSWER ALL QUESTIONS

Question 1

- (a) Define the following terms: (2 marks)
- (i) Organic synthesis
 - (ii) Retrosynthetic analysis
- (b) What is the significance of “organic synthesis.” Discuss any **TWO** counts. (2 marks)
- (c) Name the four interrelated aspects which need to be considered in planning and designing synthesis. (2 marks)
- (d) Distinguish between the following terms; (4 marks)
- (i) Convergent and divergent synthesis
 - (ii) Carbocation and carbene
- (e) What are the limitations of organic synthesis? (4 marks)
- (f) Starting with 1-bromopropane and 1,2-dibromoethane outline the synthesis of 2-pentanone. (6 marks)
- (g) Isopropyl methyl ether $[(\text{CH}_3\text{OCH}(\text{CH}_3)_2]$ may be prepared by any one of the following reactions. Which reaction would give the better yield? Explain your choice. (5 marks)
- (I) $\text{CH}_3\text{ONa} + (\text{CH}_3)_2\text{CHI} \rightarrow \text{CH}_3\text{OCH}(\text{CH}_3)_2$
 - (II) $(\text{CH}_3)_2\text{CHONa} + \text{CH}_3\text{I} \rightarrow \text{CH}_3\text{OCH}(\text{CH}_3)_2$
- (h) Synthesis of symmetrical ethers is achieved by dehydration of primary alcohols. An exception to this restriction is the synthesis of unsymmetrical ether from tert-butyl alcohol with a primary alcohol in sulphuric acid at room temperature. Give a likely mechanism for this reaction and explain why it is successful. (5 marks)

SECTION B (40 MARKS): ANSWER ANY TWO QUESTIONS FROM THIS SECTION
EACH QUESTION CARRIES 20 MARKS

Question 2

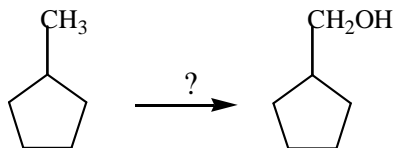
- (a) Synthesis of n-propylalcohol from n-propylbromide follows SN^2 mechanism while synthesis of tert-butyl alcohol from tert-butyl bromide follows SN^1 .
- (i) Outline **THREE** differences between SN^1 and SN^2 (3 marks)
- (ii) Give the mechanisms for the SN^1 hydrolysis with hot silver hydroxide and the SN^2 hydrolysis with hot potassium hydroxide. (7 marks)
- (b) Which of the alkyl halides given below would you expect to react more rapidly by an SN^2 mechanism.
Explain your answer. (4 marks)
- (I) $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{Br}$ vs $\text{CH}_3\text{CH}_2\underset{\text{Br}}{\text{CH}}\text{CH}_3$
- (II) $\text{CH}_3\text{CH}_2\text{CH}_2\text{Cl}$ vs $\text{CH}_3\text{CH}_2\text{CH}_2\text{Br}$
- (c) With the aid of energy-reaction path diagrams, explain why the energy profile of SN^2 has only one transition state while SN^1 has two transition states. (6 marks)

Question 3

- (a) Explain each of the following observations:
- (i) Regiospecificity of hydrogenation of alkenes fails without a peroxide in the reaction matrix. (3 marks)
- (ii) P-2 catalyst is preferred in place of Nickel metal. (2 marks)
- (iii) Acetylene is synthesizable from calcium carbonate. (3 marks)
- (iv) Proton of acetylene is easily replaced by a metal. (2 marks)
- (b) Outline the synthesis of 1-bromobutane from 1,2-dibromoethane and ethylbromide. Show all the necessary steps and the reaction mechanisms involved. (10 marks)

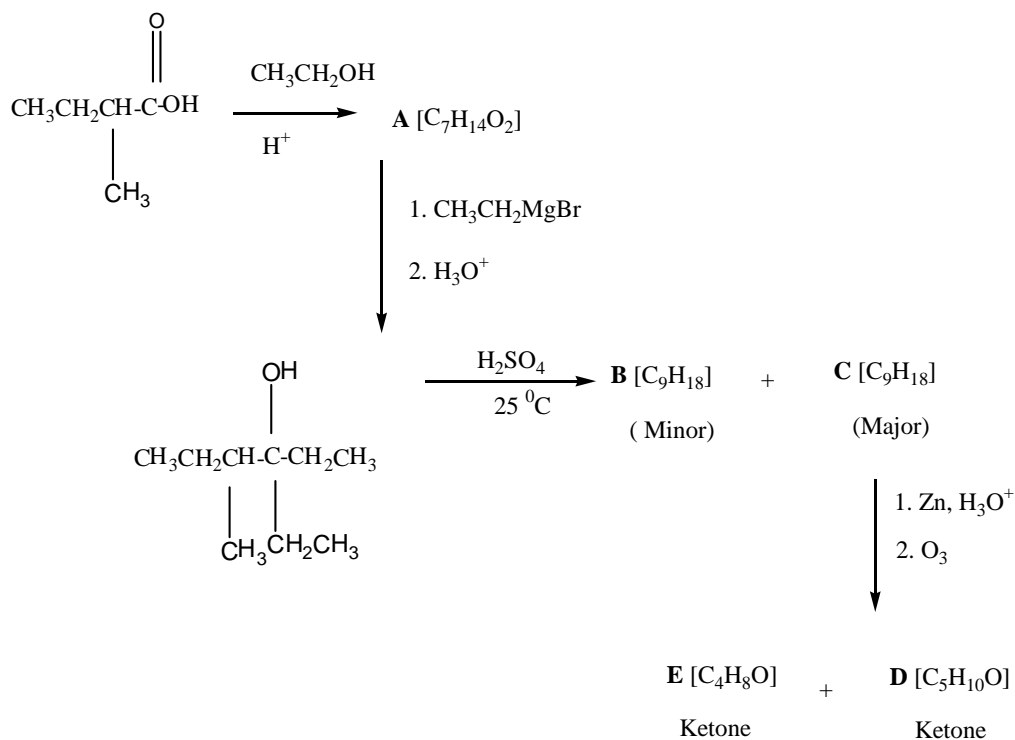
Question 4

- (a) Outline the multistep synthesis of methylenehydroxycyclopentane from Methylcyclopentane: (8 marks)



- (b) Why is chlorination not a preferred route. (2 marks)

- (c) Give the structures and names for compounds A-E. (10 marks)



Question 5

- a) Briefly discuss the synthetic pathway of any **ONE** of the following compounds. (15 marks)
- (i) Quinines
 - (ii) Nicotine
 - (iii) Chloroquine
 - (iv) Pyrethrins
- b) Discuss how organic chemical reactions can be classified with respect to structural change. (5 marks)

E *N* *D*