1×10

## 2022

# **CHEMISTRY** — HONOURS

#### Paper : CC-7

### (Organic Chemistry)

#### Full Marks : 50

The figures in the margin indicate full marks. Candidates are required to give their answers in their own words as far as practicable.

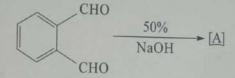
Answer question number 1 (compulsory) and any eight (08) questions from the rest (from question number 2 to question number 12).

1. Answer any ten questions :

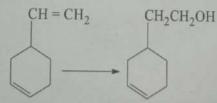
(a) Acetals are stable to bases. Explain.

(b) Explain why the following ylide does not participate in Wittig reaction :

(c) Identify the product [A] in the following reaction :



- (d) What will be the major product when ketene is made to react with HCI?
- (e) In Kolbe's reaction for the synthesis of salicylic acid, NaOH is preferred over KOH. Why?
- (f) Explain why  $\beta$ -hydroxyesters can be prepared by the Reformatsky reaction and not by Grignard reaction.
- (g) How can you accomplish the following conversion?



- (h) Indicate the reagents required for the following conversion :  $CH_3(CH_2)_4C \equiv CH \rightarrow CH_3(CH_2)_5CHO$
- (i) CF<sub>3</sub>CHO gives a stable hydrate. Explain.

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(j) When aniline is subjected to Friedel-Crafts alkylation in the presence of catalytic amount of AlCl<sub>3</sub>, alkylation does not occur; while in presence of large excess of AlCl<sub>3</sub>, a very small amount of *m*-alkylaniline is obtained. Explain precisely.

(k) Ph COOC Me<sub>3</sub> 
$$\xrightarrow{H_2O^{18}}_{H^+}$$
?

Identify the product(s). (Mechanism not needed)

- (1) How can you convert R2CHCOOH to R2CBrCOOH? (No mechanism required)
- (a) cis-2-Butene produces optically inactive racemic 2, 3-butanediol (resolvable) on treatment with mcpba followed by acid catalysed hydrolysis; whereas trans-2-butene produces optically inactive meso-2,3-butanediol (non-resolvable). Justify with plausible mechanism.
  - (b) A six membered cyclic compound gives CH<sub>2</sub>(CHO)<sub>2</sub> as the only product of ozonolysis. Identify the compound. 3+2
- (a) Reaction of acetone with equivalent proportion of bromine yields monobromoacetone under an acid catalysed condition. Explain with proper reaction mechanism.
  - (b) Predict the product of the following reaction with proper reasonings.

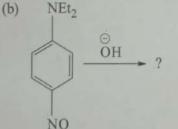
$$CH_2 = C = CH_2 \xrightarrow{HBr} ?$$
 3+2

- 4. (a) Give two Wittig alkene synthesis of 2-methyl-1-hexene. Is one synthesis preferred over the other? Why?
  - (b) Carry out the following conversion :

$$\begin{array}{c} & & O \\ \parallel & & \parallel \\ CH_3 - C - CH_2 - CH_2 - CO_2Et \rightarrow CH_3 - C - CH_2 - CH_2 - CH_2OH \end{array}$$

5. (a) Arrange the following halobenzenes in increasing rate for nitration reaction. Justify your answer.

$$C_6H_5$$
 Br,  $C_6H_5Cl$ ,  $C_6H_5I$ ,  $C_6H_5F$ 



Identify the product(s) with proper justification.

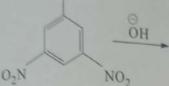
6. (a) Benzene reacts with trimethylacetyl chloride in the presence of anhydrous AlCl<sub>3</sub> at elevated temperatures to give *p*-tertiarybutylbenzene as one of the products; but toluene with the same reagent, at a lower temperature gives, *p*-tertiarybutyl acylated product only. Explain the facts showing proper reactions.

3+2

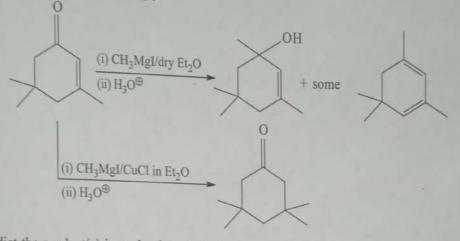
3+2

3+2

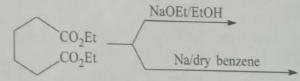
(b) Write down the product of the following reaction showing mechanism : 3+2



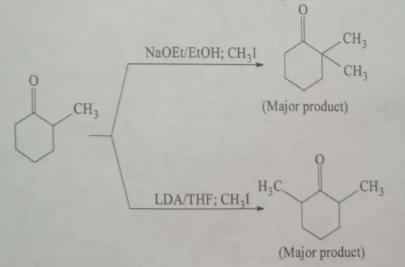
- 7. (a) Justify with proper mechanism, why *m*-chlorotoluene on treatment with  $KNH_2/NH_3(l)$  yields m-toluidine in large excess over o- and p-toluidines.
  - (b) Show mechanistically how o-xylene when heated at 80°C with anhydrous AlCl<sub>3</sub> and HCl converts
- 8. (a) Explain the following reactions :



(b) Predict the product(s) in each of the following reactions giving plausible mechanistic steps : 3+2



9. (a) Explain the formation of following products with plausible mechanism.

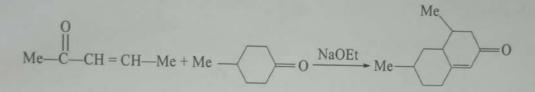


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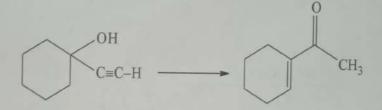
(b) Predict the products expected to be formed from the following alkene on treatment with ozone followed by oxidative work up with H<sub>2</sub>O<sub>2</sub>. 3+2



- 10. (a) Suggest Corey-House synthesis of 3-methyloctane using Gilman reagent. Why can't alkyllithium reagent be used for this synthesis?
  - (b) Suggest mechanistic steps for the following conversion :



11. (a) Carry out the following transformation mentioning proper reagents and show the plausible mechanism.



- (b) Explain why o-nitrobenzaldehyde undergoes benzoin condensation but p-nitrobenzaldehyde does not. 3+2
- 12. (a) When o-hydroxybenzoic acid is treated with excess Br<sub>2</sub>/H<sub>2</sub>O, a heavy white precipitate settles down with the evolution of a colourless gas. Explain mechanistically the products formed along with proper justification of the course of the reaction.
  - (b) Identify the product with proper mechanism.

PhCHO + BrCH<sub>2</sub>CO<sub>2</sub>Me  $K^{\oplus}tBuOH$ 

3+2

3+2