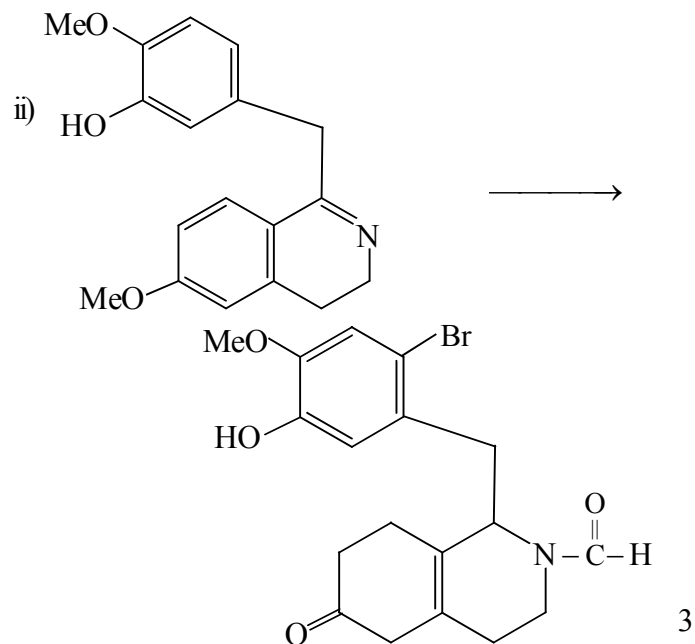
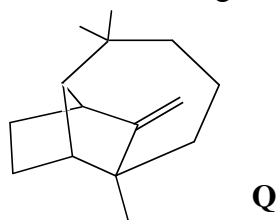


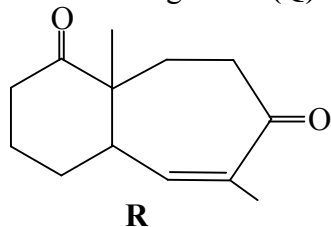
[6]



- f) i) Predict the structure of longifolene hydrochloride.
Explain its formation from longifolene (**Q**). 2



- ii) Discuss the advantages of the intermediate **R** in Prof. Corey's synthesis of longifolene (**Q**). 2



Ex/P/XIV-O/2019

M. SC. CHEMISTRY EXAMINATION, 2019

(4th Semester)

ORGANIC CHEMISTRY SPECIAL

PAPER - XIV-O

Time : Two hours

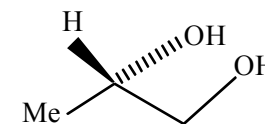
Full Marks : 50

(25 marks for each unit)

Use a separate answerscript for each unit.

UNIT - O - 4141

1. a) Design the synthesis of the compound **A** starting from a suitable easily available enantiopure chiral compound applying Chiron approach. 4



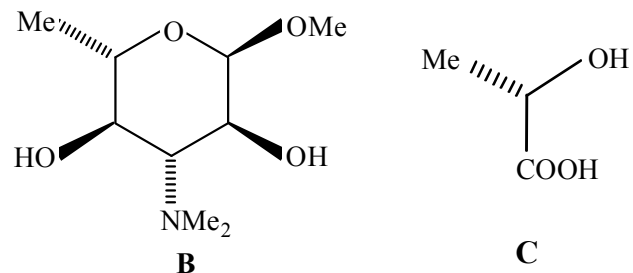
A

- b) Describe the asymmetric synthesis of *l*-menthol which was developed by Prof. Ryoji Noyori. Highlight the stereochemical features of the cyclization step. 3

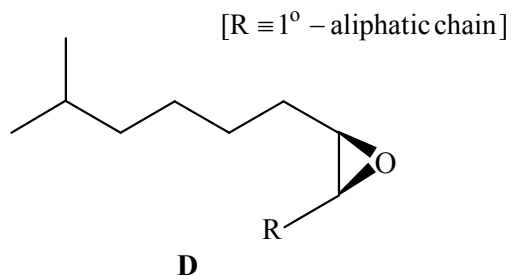
[Turn over

[2]

- c) Discuss the synthesis of the compound **B** starting from the compound **C** and account for the stereochemical aspects of the steps involved (wherever applicable). 4

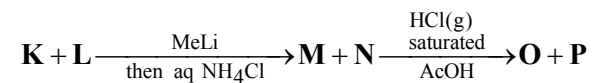


- d) Suggest the steps for the asymmetric synthesis of the compound **D** using Sharpless asymmetric epoxidation in one of the steps [only mention the steps with reagents, no mechanism is needed]. 3

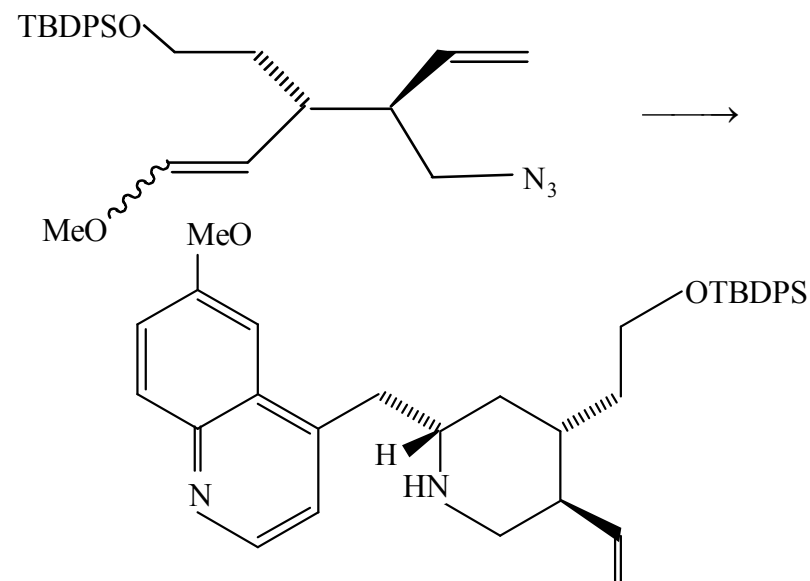


- e) Predict the product **E** when S-BINOL, LiAlH₄ and EtOH are mixed together in equimolecular proportion. When **E** reacts with acetophenone, suggest the absolute configuration of the major product obtained from acetophenone with proper justification. 3

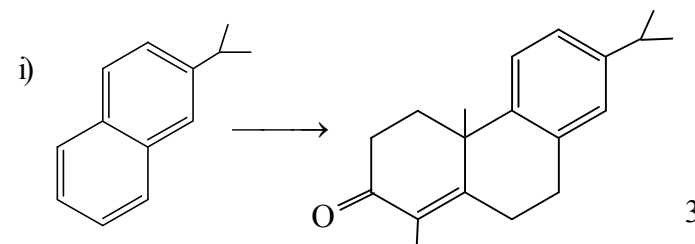
[5]



- d) Illustrate the steps involved in the following conversion highlighting the stereochemical control involved in the last step. 3



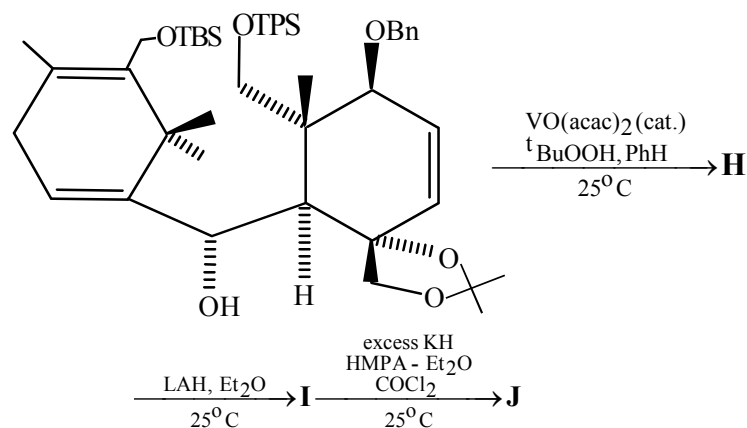
- e) Carry out the following transformations. Show all the intermediate products (mechanism is not required).



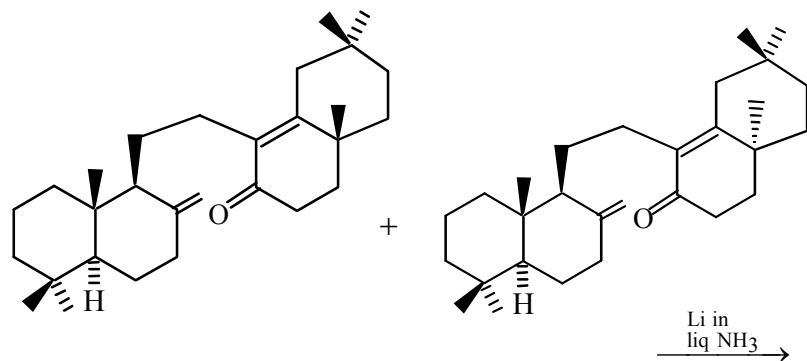
[Turn over

[4]

- b) Predict the products **H**, **I** and **J** of the following reactions (mechanism is not required). Mention at least three reasons for which COCl_2 was chosen as the appropriate reagent for the last step of the following reaction sequence. 3



- c) Identify the products (**K** to **P**) of the the following reactions. Comment on the stereochemistry of any newly generated chiral center(s) of all the products formed. 4

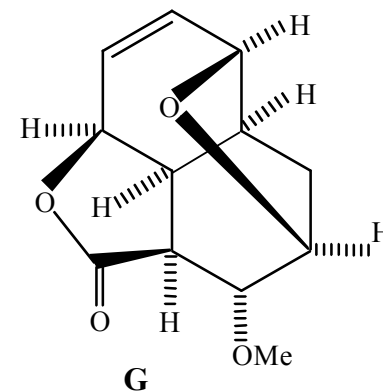


[3]

- f) Suggest the structure of the product **F** obtained in the reaction of D-fructose with dry acetone in the presence of anhydrous ZnCl_2 followed by PCC oxidation. If *E*-stilbene is treated with H_2O_2 in CH_3CN in the presence of catalytic amount of **F**, predict the outcome with absolute configuration of the stereocentres in the predominant product. 1+3
- g) Define the term “Sustainability” 1
- h) Logically suggest the product when *n*-propylbenzene reacts with potassium permanganate adsorbed on neutral alumina. 3

UNIT - O - 4142

2. a) Discuss synthesis of the following compound **G** (racemic variety) from two achiral starting materials. Suggest appropriate mechanistic and stereochemical interpretations as necessary. 5



[Turn over