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## CHEMISTRY ORGANIC CHEMISTRY II

Level & Board	AQA (A-LEVEL)
TOPIC:	OPTICAL ISOMERISM
PAPER TYPE:	SOLUTION - 3
TOTAL QUESTIONS	10
TOTAL MARKS	/33

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## <u> Optical Isomerism - 3</u>

### Ι.

(a) Name alcohol (CH<sub>3</sub>)<sub>2</sub>C(OH)CH(CH<sub>3</sub>)<sub>2</sub>

2, 3 – dimethylbutan – 2 – ol



Name compound C: Propyl methanoate

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**Reagent:** Sodium bicarbonate (NaHCO<sub>3</sub> )

Observation with C (Propyl methanoate):

No reaction or effervescence will be observed.

Observation with D (Carboxylic acid):

Effervescence (bubbling) will be observed when sodium bicarbonate is added to a carboxylic acid. This is due to the release of carbon dioxide gas  $(CO_2)$  when the carboxylic acid reacts with sodium bicarbonate.

Reagent: acidified K2Cr2O7

Observation with C (Propyl methanoate):

Becomes green

Observation with D (Carboxylic acid):

no change

(3)

The word Chirality / Chiral is used to describe optically active molecules.

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4.

3.

Examples of Chiral Molecules:

*Lactic Acid:* Lactic acid is a chiral molecule. It exists in two enantiomeric forms: L-lactic acid and D-lactic acid.

**2-Amino-I-propanol:** This is a chiral molecule due to the presence of a chiral carbon atom bonded to four different groups.

Lactic Acid:





(2)

#### (c) Type of reaction: Dehydration / elimination

(1)

## (d)

When 2,2-dimethylpropan-I-ol is heated with concentrated sulfuric acid, it does not undergo dehydration because:

**No Hydrogen Atoms:** There are no hydrogen atoms on the carbon atom adjacent to the C-OH group.

**Steric Hindrance:** The carbon atom next to the C–OH group is bonded to three bulky methyl groups, preventing the necessary steps for dehydration.

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#### 6.

Isomer E:



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Isomer F:

(2)

$$CH_3CH_2 - C_{CH_3}^{H} C_{OCH_3}^{H}$$

7.

Structures of Isomers

When 2-butene reacts with HBr, it forms two structural isomers: **2-Bromobutane** 



2-bromobutane

I-Bromobutan

1-bromobutane Mechanism for Formation of the Major Product:

#### Protonation of the Alkene:

- The double bond in 2-butene attacks a proton (H<sup>+</sup> ) from HBr, leading to the formation of a carbocation intermediate.
- $_{\circ}$  The more stable carbocation forms at the more substituted carbon.

 $CH_3 - CH = CH - CH_3 + HBr \rightarrow CH_3 - CH^+ - CH_2 - CH_3 + Br^-$ 

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#### Nucleophilic Attack:

• The bromide ion (Br<sup>-</sup>) then attacks the carbocation, resulting in the formation of 2-bromobutane.

$$CH_3 - CH^+ - CH_2 - CH_3 + Br^-$$

 $\bullet \quad CH_3 - CH(Br) - CH_2 - CH_3 \text{ or}$ 

 $CH_2(Br)-CH_2-CH_2-CH_3$ 

Structures: 2-Bromobutane (Major Product): Structure:

2-bromobutane CH<sub>3</sub>-CH(Br)-CH<sub>2</sub>-CH<sub>3</sub> **I-Bromobutane (Minor Product):** Structure:

1-bromobutane  $CH_2(Br) - CH_2 - CH_2 - CH_3$ 

(6)

#### 8.

The racemic mixture has no effect on plane-polarized light.

This is because in a racemic mixture, there are equal amounts of both enantiomers present.

Each enantiomer rotates plane-polarized light in opposite directions, but to the same extent.

As a result, the rotations caused by the two enantiomers cancel each other out, resulting in no net rotation of plane-polarized light.

Therefore, when a racemic mixture is passed through a polarimeter, no overall rotation is observed.

(2)



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