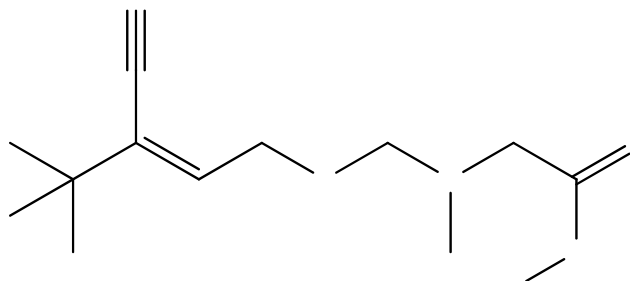


## Lesson 2 For Organic Chemistry

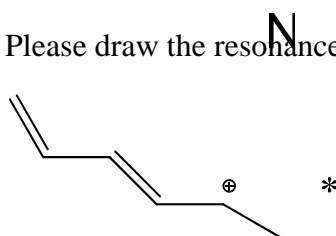
### Revision Exercise and Extra Knowledge --- For Hybridization Theory and Resonance Concept

1. Please state the hybridization state of all the atoms with a \*.



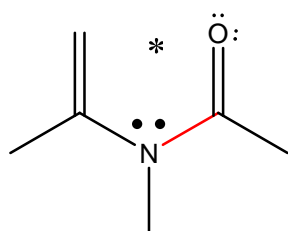
**Hint :** There are \_\_\_ lone pair electrons left on the oxygen atoms and also \_\_\_ on the nitrogen atom.

2. Please draw the resonance structures for the following species and answer the questions.



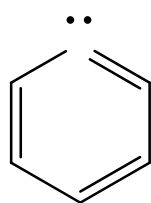
a)

Qu: Which of the carbon atom (s) doesn't carry a positive charge? \_\_\_, \_\_\_ and \_\_\_



b)

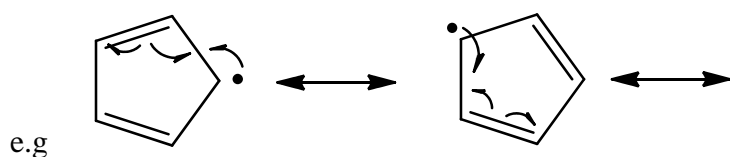
Qu : Can the amide bond in red be rotated easily? \_\_\_. It is because it bears \_\_\_\_\_ bond character.



c)

. It is called **pyridine**, which is a b\_\_\_\_\_.

As you should know, resonance structures can be drawn for those c\_\_\_\_\_ systems in neutral state, positively or negatively charged state. This resonance effect can stabilize the compounds in terms of e\_\_\_\_\_ aspect. Actually, a 'radical' species can also be stabilized by resonance.

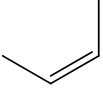



**Isomerism**

Isomers are those compounds with the same molecular formula but different structures in space. Generally, all atoms in different isomers have the \_\_\_\_\_ bonding connectivity. Isomerism is a complicated topic; but we only need to focus on the **geometrical isomerism** and **stereoisomerism** only.

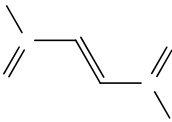
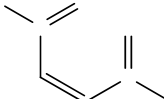
**1. Geometrical Isomerism**

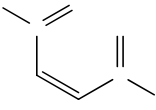
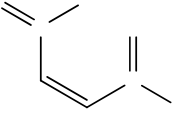
Most importantly, we need to differentiate the two forms of geometrical isomers : \_\_\_\_\_ and \_\_\_\_\_.

For example,  is called \_\_\_\_\_ but-2-ene;  is called \_\_\_\_\_ but-2-ene. Also, you should know that the cis isomer can be made into trans isomer by **breaking and reforming** the \_\_\_\_\_ bonds, or vice versa. (HKAL 2009-2010)

Simply speaking, geometrical isomers have the \_\_\_\_\_ chemical properties. However, when we compare some physical properties such as melting point/ boiling point, they **may not** be the same.

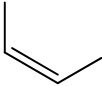
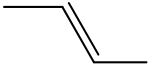
**Illustration 1: Comparison of boiling point**

Explain the fact that  has a **higher** boiling point than  ?

→ Note that  is the same as  . \_\_\_\_\_ molecular hydrogen bonds are formed. In other words, the chance for the cis form to form \_\_\_\_\_ molecular hydrogen bonds is \_\_\_\_\_ . For the trans-form, there are **extensive** formation of \_\_\_\_\_ molecular hydrogen bonds. Hence, the result follows.

→ How about their difference in **acidity**?

**Illustration 2 : Comparison of boiling point**

Explain the fact that  has a **higher** boiling point than  ?

→ The main difference between the cis/trans isomers in this case is their **polarity**. Cis-isomer has a **permanent** dipole moment and hence it is a p\_\_\_\_\_ molecule. However, the trans form is not polar because there is a **cancellation** of dipole moment. Thus, there are \_\_\_\_\_-\_\_\_\_\_ interactions between cis form molecules and more energy is required to separate the stronger interaction to boil it.

→ How about their **melting point**?

HO

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2

C

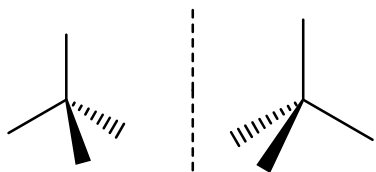
O

O

C

## 2. Enantiometric Isomerism

It is talking about the relationship between a compound (molecule) and its mirror image. What is called **mirror image**? Actually, every compounds has its mirror image. It can be drawn by a simple **reflection** as :



. But, if the compound is enantiomeric, the compound should be

imsuper to its mirror image.

Usually, if a tetrahedral atom ( $sp^3$  hybridized) has \_\_\_ different bonding groups, the molecule formed will likely be enantiomeric. It is because the molecule contains a c\_\_\_\_\_ centre. But you should be careful that when the compound contains **A more than 1** chiral \_\_\_\_\_, it may **not** be a chiral molecule anymore. So, the presence of chiral centre(s) is **not** an absolute indication for the occurrence of enantiomerism. A compound \_\_\_\_\_ be chiral if it contains even **no** chiral centre. However, a simple rule can be stated as :

Enantiomerism occurs when a compound cannot super\_\_\_\_\_ on its mirror image.

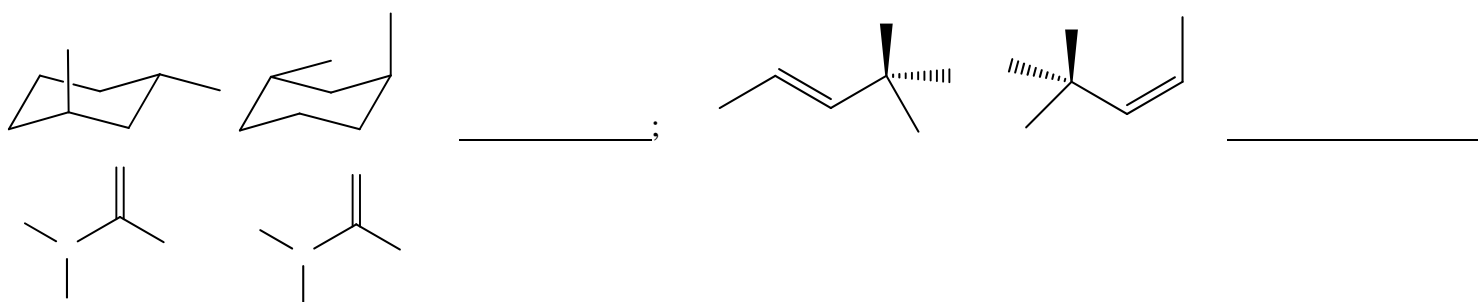
### Exercise For Isomerism

### D

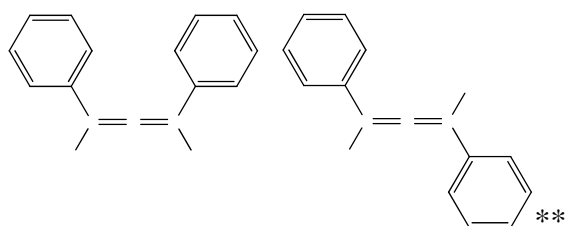
a) Please draw the **isomers** for the following compounds.



b) Please state the relationship (Identical, geometric or enantiomer) between the following pairs of compounds.



They are the \_\_\_\_\_ after a simple r\_\_\_\_\_. They are called **rotamer**.



\_\_\_\_\_ as there is **no internal plane of symmetry**.

## 2. Enantiometric Isomerism

Regarding the pair of enantiomer, they should have the \_\_\_\_\_ **physical** properties except for the direction of rotation of the plane of polarized light. It is because they have the **opposite optical activity**. The ability to interact with the plane polarized light is called \_\_\_\_\_ activity.

→ A **chiral** and **non-racemic** compound can rotate the plane of polarized light since they have optical activity. But for a pair of enantiomer, they will have the o\_\_\_\_\_ optical activity.

→ If we mix a **1:1** ratio of enantiomer, the solution formed will show \_\_\_\_ optical activity. This solution is called **racemic mixture**, which is an important term in the organic chemistry later.

→ The **racemate** (= \_\_\_\_\_ mixture) may have **different** values In terms of boiling point/melting point from those of a **single** enantiomer.

Just to mention, enantiomers also have the \_\_\_\_\_ **chemical** properties, except when interacting with another c\_\_\_\_\_ compounds. (Like some enzymatic reactions happen in our body)

### Important Terms For Organic Chemistry

1. **Acid and Base** : Organic compounds can be an acid (a \_\_\_\_\_ donor) or a base (a proton a\_\_\_\_\_).

→ The reaction between an acid and a base is called A\_\_\_\_\_ B\_\_\_\_\_ reaction.

→ This concept leads to the comparison of **Acidity and Basicity**.

2. **Nucleophiles and Electrophiles** : Nucleophiles are those compounds which are **nucleus loving**. They usually bear lone pair of electrons and \_\_\_\_\_ charged; Electrophiles are those compounds which are **electron loving**. They usually are \_\_\_\_\_-deficient.

→ The reaction between a nucleophile and an electrophile is called I\_\_\_\_\_ reaction.

→ This concept leads to the comparison of **Nucleophilicity and Electrophilicity**.

3. **Intermediate** : A s\_\_\_\_\_ lived and very un\_\_\_\_\_ species that form along the reaction pathway.

Usually, in HKAL, we should be familiar with the c\_\_\_\_\_ intermediate.

→ Note that intermediate is **not** the same as t\_\_\_\_\_ state.

4. **Stability** : It is the term relating to the energy of the species. If the species has a \_\_\_\_\_ energy, it is said to be relatively s\_\_\_\_\_.

5. **Electron-donating and Electron-withdrawing groups** : Electron-donating groups are those group which can \_\_\_\_\_ electrons; Electron-withdrawing groups are those which can \_\_\_\_\_ electrons.

→ This concept leads to the **Inductive effect (through space) and Resonance effect (more powerful)**.

6. **Reaction mechanism** : It is the detailed and step-by-step description of the pathway by which reactants are converted to \_\_\_\_\_. (Frequently asked)