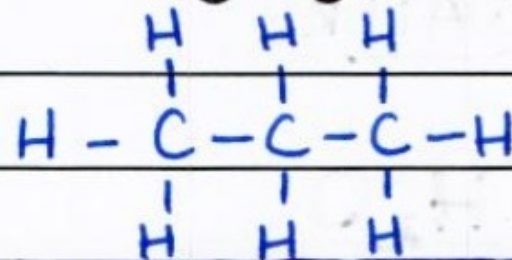


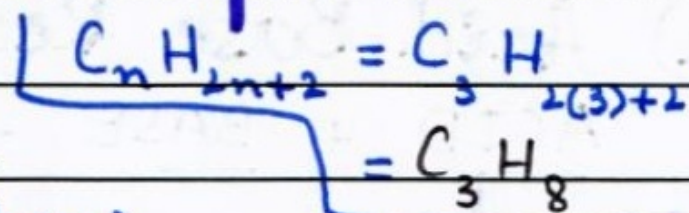
Q. No. 2 Part (i) Names of Organic Compounds:

a) $C_3H_8 \rightarrow$ **PROPANE**

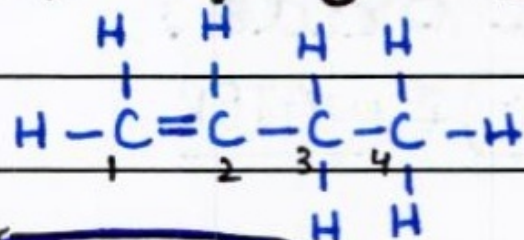


• Stem: Prop -
(3 carbons)

• Class of Compound: Alkane

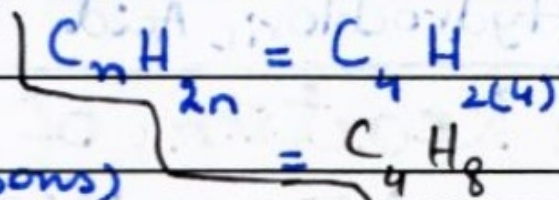


b) $C_4H_8 \rightarrow$ **BUTENE**



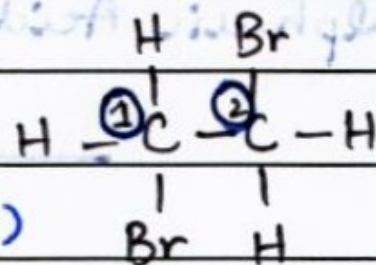
• Stem: But -
(4 carbons)

• Class of Compound: Alkene



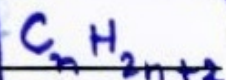
1-Butene

c) $C_2H_4Br_2 \rightarrow$ **1,2-Dibromo Ethane**



• Stem: Eth - (2 carbons)

• Class of Compound: Alkane



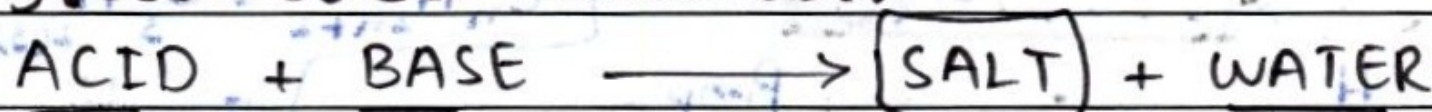
Q. No. 2 Part (ii) Raw Materials for Solvay Process:

Following RAW MATERIALS are used in Solvay process for the manufacture of sodium carbonate (Na_2CO_3):

- 1) Ammonia (NH_3)
- 2) Concentrated solution of NaCl (BRINE).
- 3) Calcium carbonate or Limestone (CaCO_3) as source of carbondioxide (CO_2) and slaked lime Ca(OH)_2 .

Q. No. 2 Part (iii) Formation of Salts:

a) Acid-Base Reaction

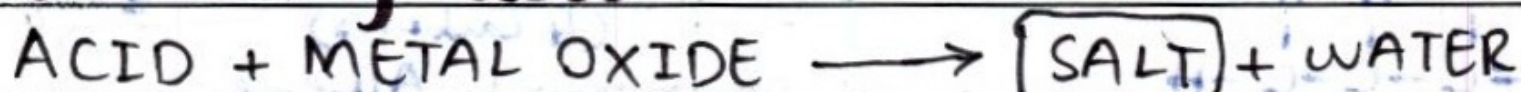


Example:

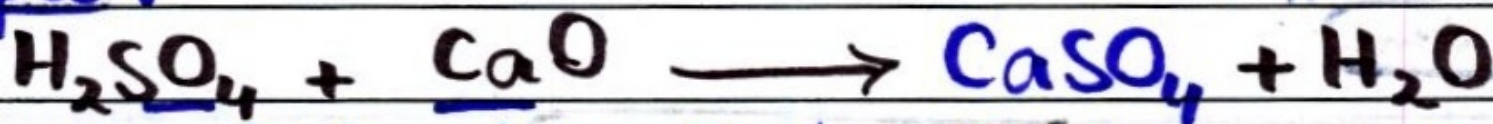


Hydrochloric Acid Sodium Hydroxide Sodium chloride water

b) Reaction of acid with metal oxide:



Example:



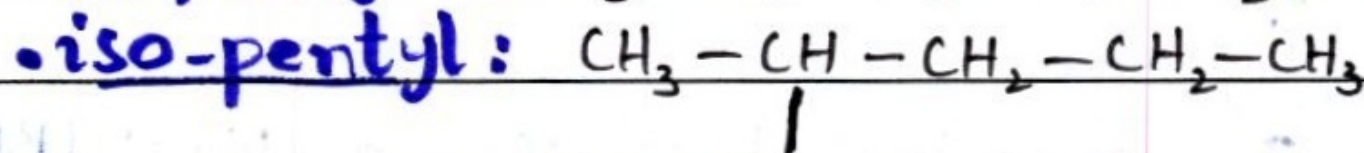
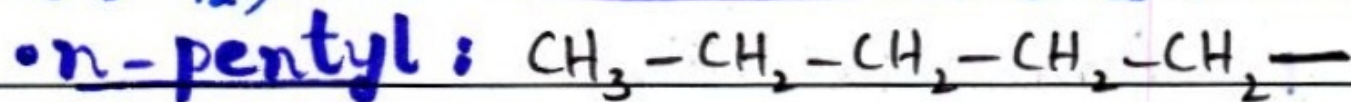
Sulphuric Acid Calcium Oxide Calcium Sulphate water

Q. No. 2 Part (iv) Alkyl Radicals:

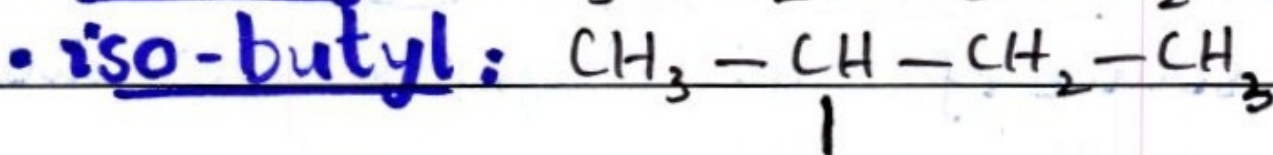
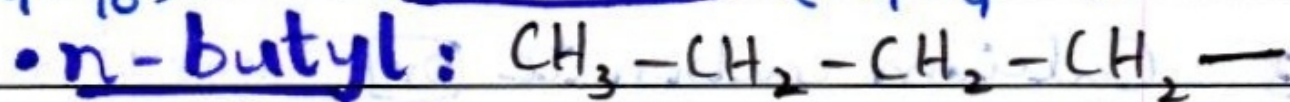
a) Ethane (C_2H_6) \longrightarrow **ETHYL** (C_2H_5-)



b) Pentane (C_5H_{12}) \longrightarrow **PENTYL** ($C_5H_{11}-$)

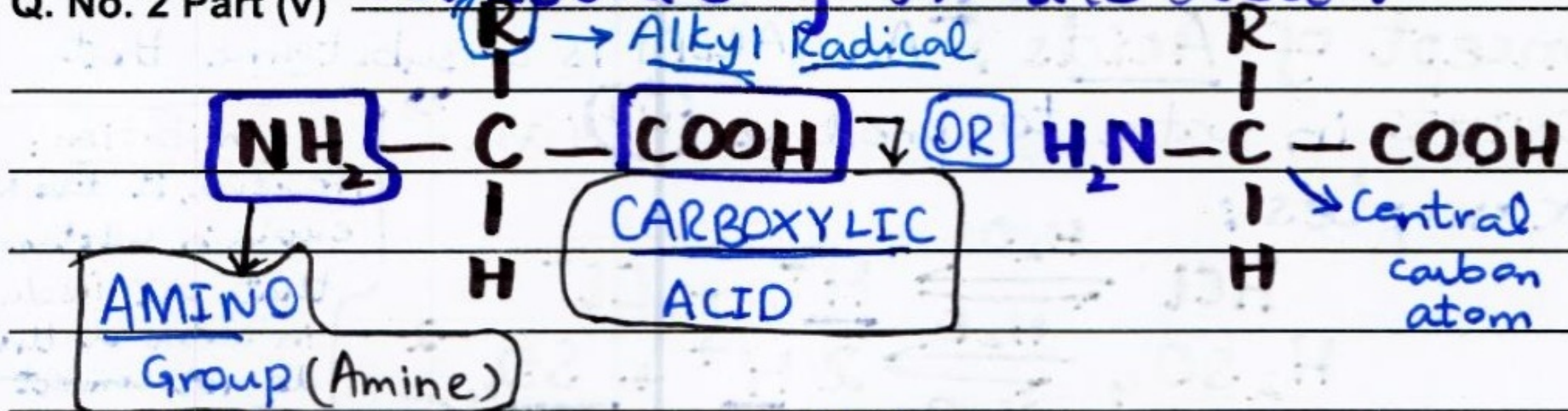


c) Butane (C_4H_{10}) \longrightarrow **BUTYL** (C_4H_9-)



Q. No. 2 Part (v)

Structure of Amino Acid:



Functional Groups : ① Amino Group / AMINE ($\text{H}_2\text{N}-$)
② Carboxylic Acid ($-\text{COOH}$)

Structure :

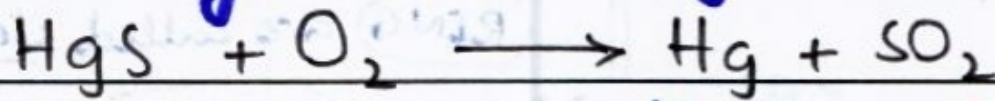
In structure of AMINO ACID, the aforementioned TWO functional groups are ATTACHED to the (central) carbon atom. An HYDROGEN atom is also attached to carbon along with an alkyl radical (R).

Q. No. 2 Part (vi) **Roasting:**

DEFINITION: "Some METALS are converted to oxides by heating in the AIR at temperature below their melting point. This process is called ROASTING."

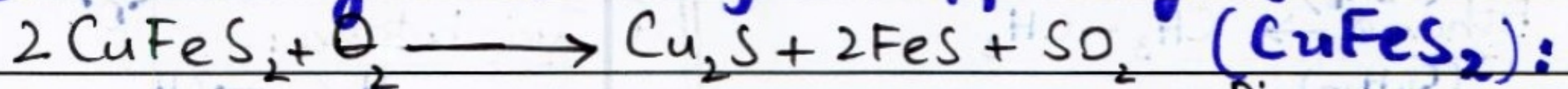
EXAMPLES:

① **Roasting Reaction for Cinnabar (HgS):**



Cinnabar Oxygen Mercury Sulphur dioxide

② **Roasting Reaction for Copper Pyrite Ore**



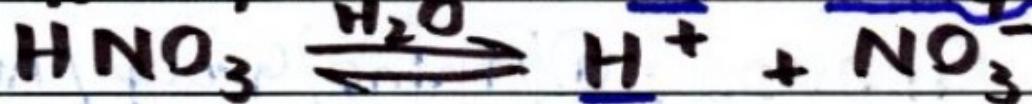
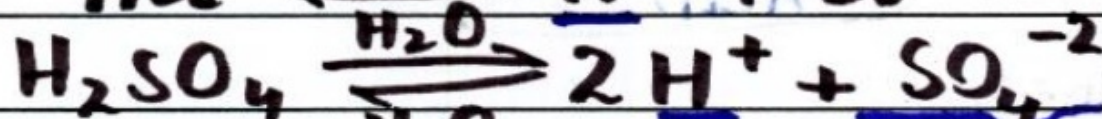
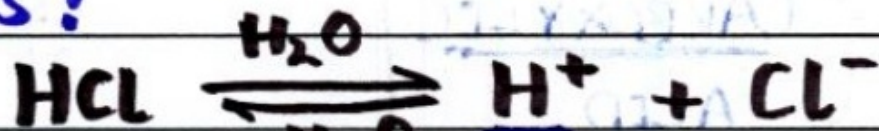
Copper Pyrite Ore Cuprous Sulphide Iron Sulphide Sulphur Dioxide

⇒ In roasting, addition of OXYGEN (oxidation) occurs and METAL OXIDES are formed at LOW TEMPERATURE.

Q. No. 2 Part (vii) Arrhenius Acids and Bases:

Concept of Acids: "An **ACID** is a substance that IONIZES in water to produce **H⁺** ions."

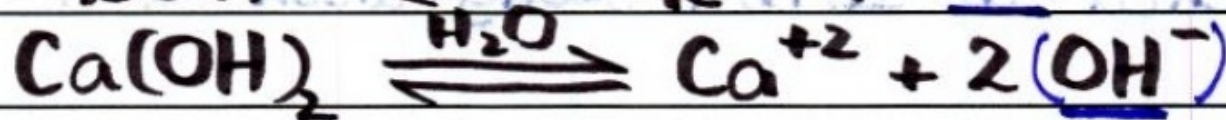
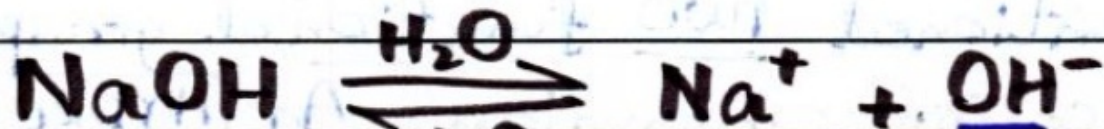
Examples:



Limitation:
However, it does NOT explain substances that are insoluble in water or that do not convert to contain H⁺ or OH⁻ ions.

Concept of Bases: "A **BASE** is a substance that IONIZES in water to produce **OH⁻** ions."

Examples:



Q. No. 2 Part (viii) Open Chain

Closed Chain


Definition

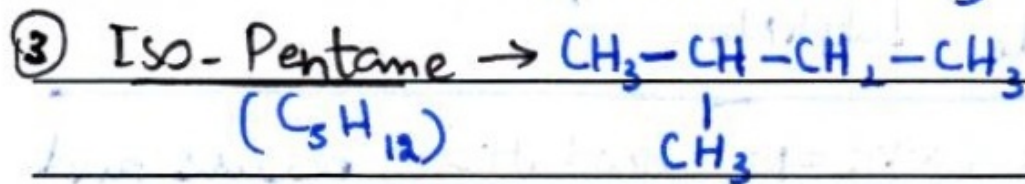
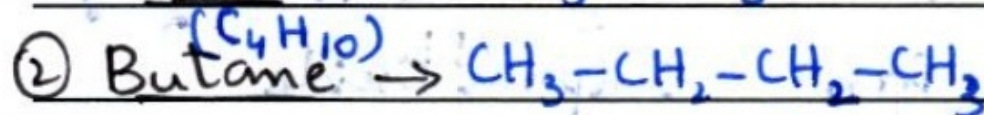
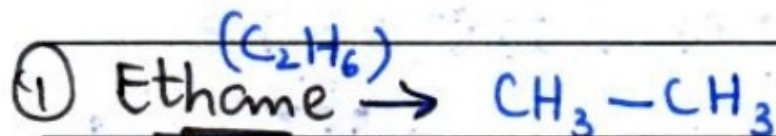
“Open chain compounds CONTAIN an open-ring of carbon atoms.”

“Those compounds which contain ANY number of carbon atoms joined with each other in a CHAIN or RING are called closed-chain compounds.”

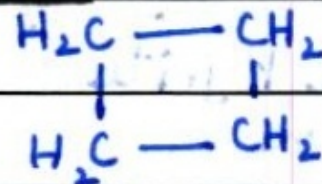
A cyclic compounds

Other Name Cyclic compounds.

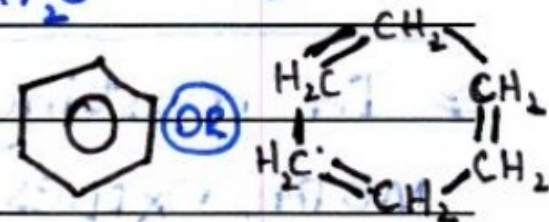
Examples ③ Furan: 



① Cyclo butane:



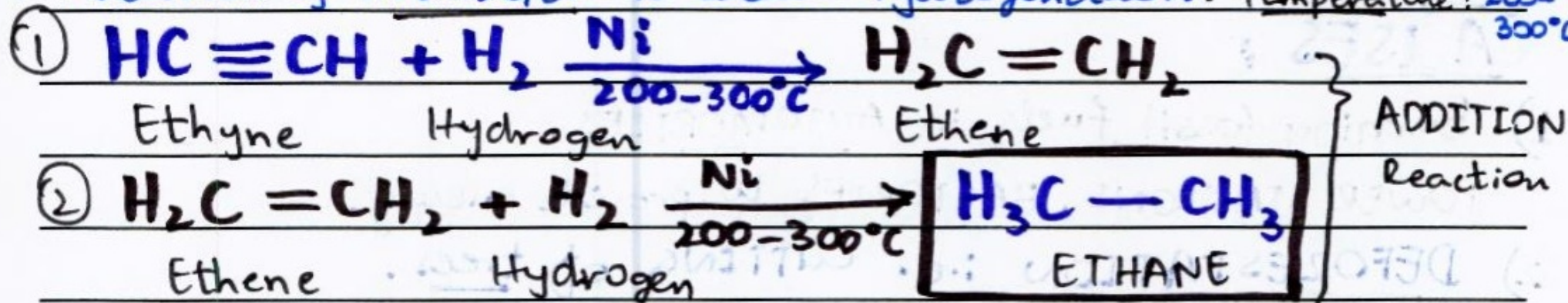
② Benzene:



Q. No. 2 Part (ix) Preparation of Ethane:

1) By Hydrogenation of Alkenes and Alkynes:

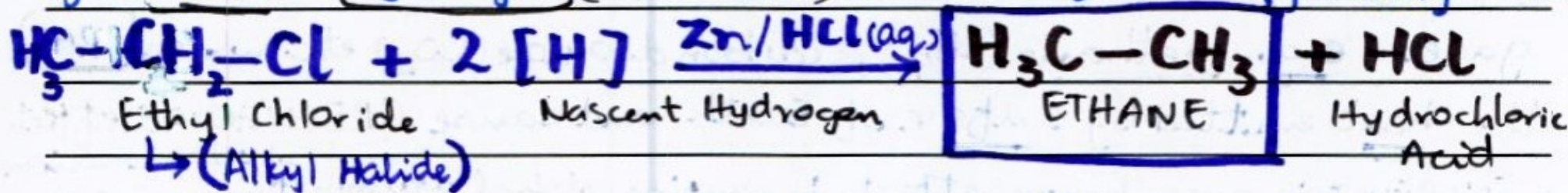
Addition of HYDROGEN is called hydrogenation. Catalyst: Ni or Pt
Temperature: 200-300°C



2) By the Reduction of Alkyl Halides: Zn reacts

with HCl or CH₃COOH (AQUEOUS ACID) to give NASCENT hydrogen. Addition

of NASCENT hydrogen (reduction) reduces alkyl halide, forming ALKANE.



Q. No. 2 Part (x) Ozone Layer Formation: Ozone is the protective layer found in STRATOSPHERE which blocks HARMFUL ultraviolet radiation coming from the Sun.

Natural Balance: Naturally, OZONE formation and its DEPLETION naturally, occur at the SAME rate and natural balance is maintained.

Ozone Depletion: $O_3 + O \longrightarrow 2O_2$

Ozone Reformation: $O_2 + O \longrightarrow O_3$

$3O_2 \xrightleftharpoons[\text{Discharge}]{\text{Electrical}} 2O_3$

Presence of OZONE felt near photocopier

Molecular Oxygen

Atomic Oxygen

Ozone

⇒ However, due to human activities like burning of FOSSIL FUELS, this NATURAL balance is DISTURBED.

⇒ Chlorofluorocarbons (CCl_3F) react with ozone to form CHLORINE FREE RADICALS ($Cl\cdot$) which cause further thinning of ozone layer (OZONE HOLE).

Q. No. 2 Part (xi)

Global Warming:

Definition: The **RISE** in the natural **TEMPERATURE** of Earth due to human activities is called **GLOBAL WARMING**.

CAUSES:

- 1) Burning **fossil fuels** in **AUTOMOBILES**, **POWER STATIONS**, **FACTORIES** to provide energy.
- 2) **DEFORESTATION** i.e. **CUTTING** of **trees**.
- 3) **Exhaust gases / fumes** of **HEAVY industrial units** and **MACHINERY**.
- 4) All the above and various other **NATURAL** processes like **forest fires**, **volcanoes** emit **LARGE AMOUNTS** of **GREENHOUSE gases** e.g. **methane (CH₄)**, **carbon dioxide (CO₂)** etc. which **TRAP** the **heat** emitted by **surface** of **Earth** and cause **GREENHOUSE effect** which has serious consequences.
⇒ **INCREASING** greenhouse effect is causing **global warming**.

Use for rough work

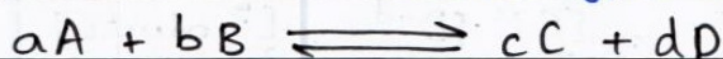
Q. No. 3 (Page 1) LAW OF MASS ACTION

Discovery: In 1864, two chemists C.M. Guldberg and P. Waage proposed Law of Mass Action to describe EQUILIBRIUM state.

Statement: Law of Mass Action states that,
"The rate at which a substance reacts is DIRECTLY PROPORTIONAL to its ACTIVE MASS. The rate at which the reaction proceeds is DIRECTLY PROPORTIONAL to the product of active masses of the reactants."

Active Mass: Active mass describes the CONCENTRATION of reactants and products in mol dm^{-3} for a dilute solution and is represented in terms of (SQUARE) brackets [].

DERIVATION: Consider a hypothetical equation; -



⇒ According to the Law of Mass Action;

Rate of forward reaction $\propto [A]^a \cdot [B]^b$

$$\text{Rate of forward reaction } (R_f) = k_f [A]^a \cdot [B]^b$$

Also,

Rate of reverse reaction $\propto [C]^c \cdot [D]^d$

$$\text{Rate of reverse reaction } (R_r) = k_r [C]^c \cdot [D]^d$$

At EQUILIBRIUM,

Rate of forward reaction = Rate of reverse reaction

$$k_f [A]^a \cdot [B]^b = k_r [C]^c \cdot [D]^d$$

$$\frac{k_f}{k_r} = \frac{[C]^c \cdot [D]^d}{[A]^a \cdot [B]^b}$$

$$\therefore \frac{k_f}{k_r} = K_c$$

$$K_c = \frac{[C]^c \cdot [D]^d}{[A]^a \cdot [B]^b} = \frac{[\text{Products}]}{[\text{Reactants}]}$$

Q. No. 3 (Page 2)

Here " K_c " is the equilibrium constant.

EQUILIBRIUM CONSTANT:

Definition: "Equilibrium constant is defined as the ratio of ^{the} concentration of PRODUCTS to the ratio of concentration of REACTANTS each raised to the **POWER** equal to the coefficient in its balanced chemical equation."

Dependence:

⇒ Equilibrium constant does **NOT** depend on the initial concentration of REACTANTS.

⇒ However, it **DEPENDS** on TEMPERATURE.

EQUILIBRIUM CONSTANT EXPRESSION:

Eq. (A) i.e.

$$K_c = \frac{[C]^c \cdot [D]^d}{[A]^a \cdot [B]^b} = \frac{[\text{Products}]}{[\text{Reactants}]}$$

is the **REQUIRED** equilibrium constant expression with C and D as PRODUCTS and A and B as REACTANTS.

Soft Water:

Definition: "The water that gives LATHER with SOAP and does NOT form scum is called soft water."

For Example: DISTILLED water is soft water.

Methods to Remove Permanent Hardness:

Permanent Hardness: Permanent hardness is called so because it CAN NOT be removed by BOILING.

Caused By: Permanent hardness of water is CAUSED by:

- ① Chlorides of CALCIUM and MAGNESIUM i.e. $\text{CaCl}_2, \text{MgCl}_2$
- ② Sulphates of CALCIUM and MAGNESIUM i.e. $\text{CaSO}_4, \text{MgSO}_4$

Methods:

Following Two methods are used to remove permanent hardness of water;

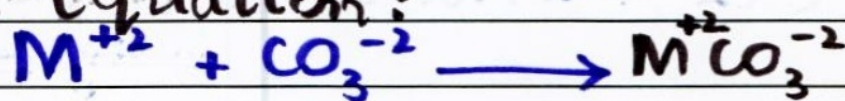
1) By adding Washing Soda ($\text{Na}_2\text{CO}_3 \cdot 10\text{H}_2\text{O}$):

⇒ On the LARGE scale, permanent hardness in water is removed by adding appropriate/estimated amount of WASHING SODA ($\text{Na}_2\text{CO}_3 \cdot 10\text{H}_2\text{O}$).

⇒ Ca^{+2} and Mg^{+2} ions causing the HARDNESS are removed as INSOLUBLE carbonates i.e. $\text{CaCO}_3, \text{MgCO}_3$

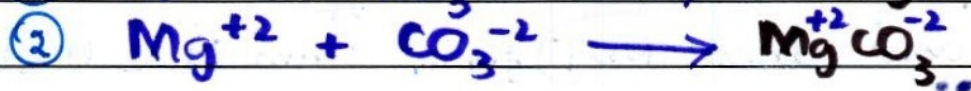
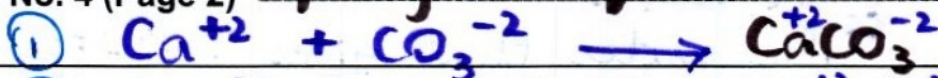
⇒ Thus, water becomes soft.

General Equation:



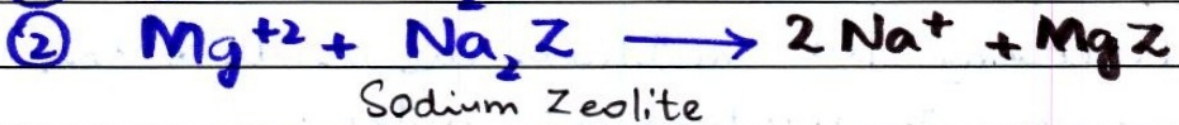
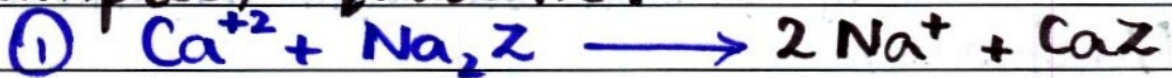
where $\text{M} = \text{Ca}^{+2}, \text{Mg}^{+2}$

Q. No. 4 (Page 2) Specific Equations

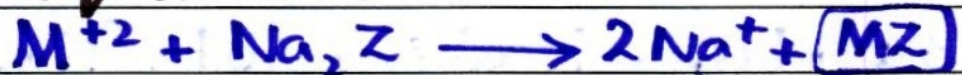


2) By Ion Exchange Resins: Permanent hard water is stored in a tank of suitable resin containing SODIUM ions. ZEOLITE is used as natural ion exchanger. SODIUM ZEOLITE is usually written as Na_2Z . Chemically, it is sodium aluminium silicate. It reacts with Ca^{+2} or Mg^{+2} ions causing the HARDNESS to form calcium zeolite or magnesium zeolite respectively and SODIUM IONS, thus REMOVING hardness.

Examples/Equations:



General Equation:



where $M = Ca^{+2}, Mg^{+2}$

⇒ Sodium zeolite can be Regenerated by reacting the formed zeolites with concentrated $NaCl$.

⇒ Thus, making the process ECONOMICAL.

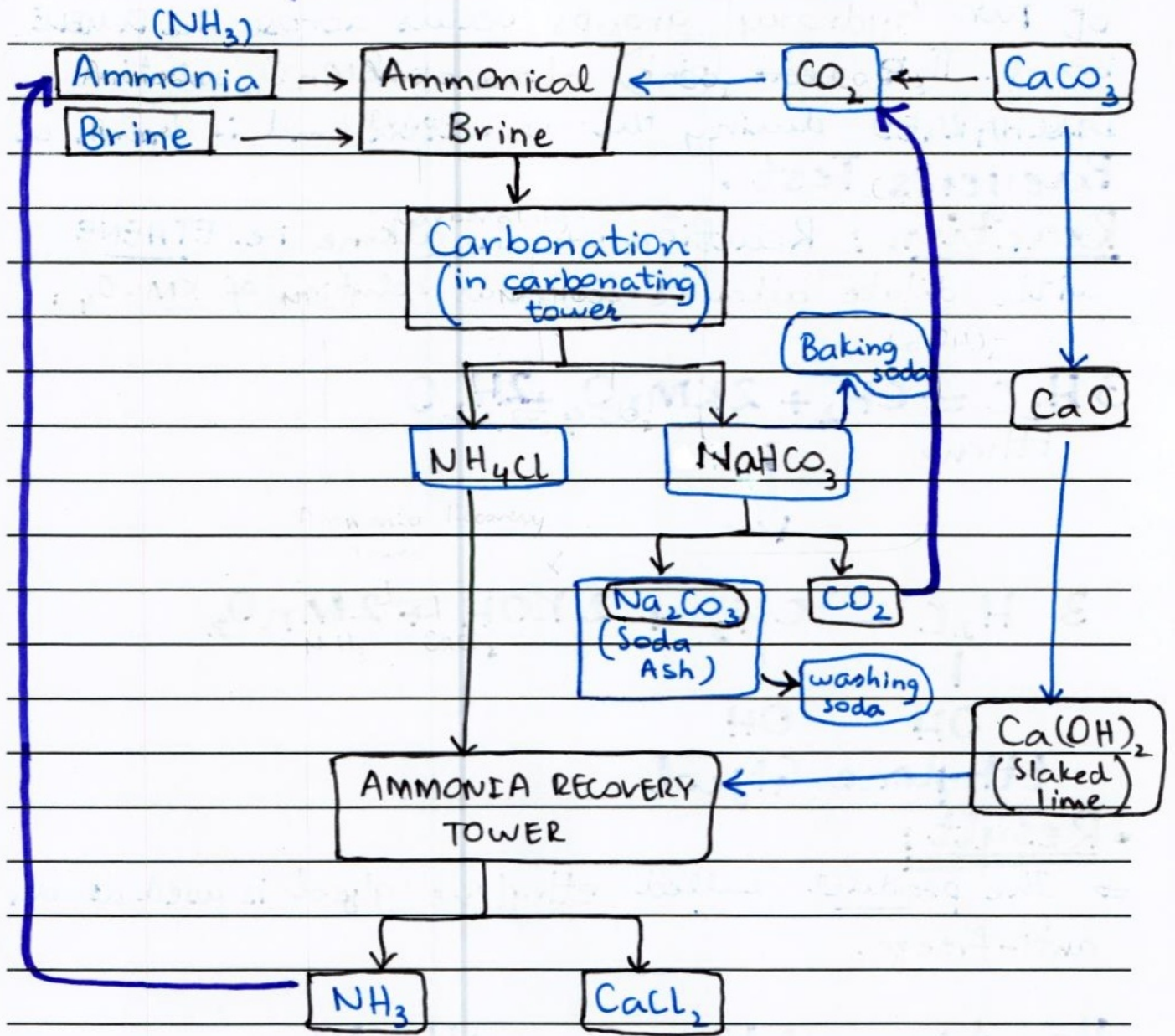
For Example:



Sodium Zeolite

↓
REGENERATED

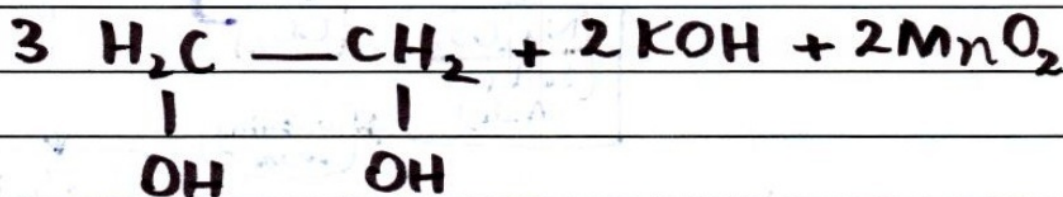
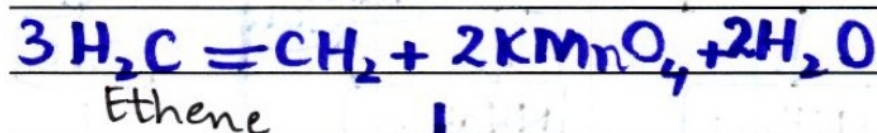
Q. No. 5 (Page 1) **Flow Sheet Diagram of SOLVAY PROCESS,**



Q. No. 5 (Page 2) Reaction of KMnO_4 with Alkenes:

Process: When an alkene is treated with dilute alkaline aqueous solution of KMnO_4 (1%), addition of two hydroxyl groups occurs across the double bond. The ~~rose~~ (pink) colour of KMnO_4 solution is DISCHARGED during the reaction and is known as Baeyer's Test.

Reaction: Reaction of an alkene i.e. ETHENE with dilute alkaline aqueous solution of KMnO_4 :



Ethylene Glycol

• Result:

⇒ The product called ethylene glycol is used as an anti-freeze.

Reaction of KMnO_4 with Alkynes:

Process: Alkynes do NOT react with dilute alkaline solution of KMnO_4 . However, they are oxidized by strong solution of KMnO_4 to give OXALIC ACID.

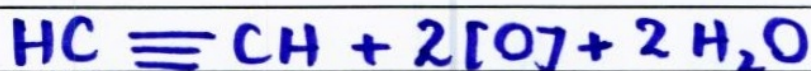
Reactions:

Q. No. 6 (Page 1) First Step:

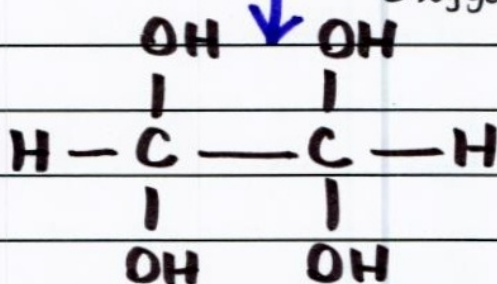


Second Step:

Secondly, four hydroxyl groups are added across the TRIPLE bond, to form tetrahydroxy ethane.

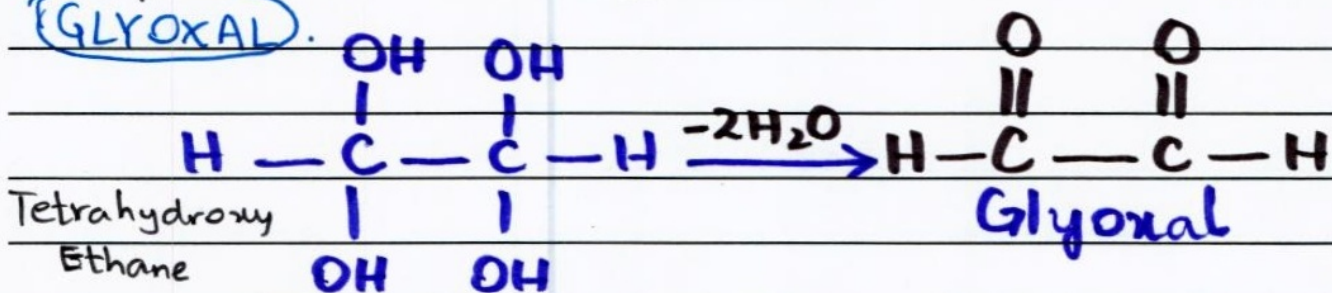


Ethyne Nascent Oxygen water

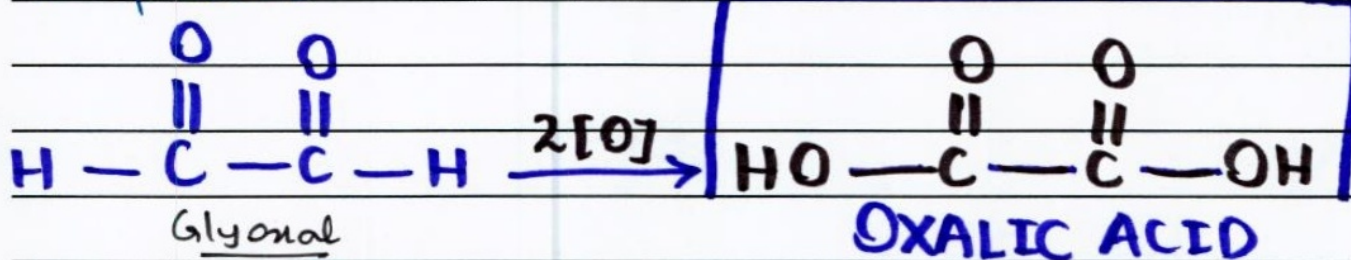


Tetrahydroxy Ethane

Third Step: Tetrahydroxy ethane is an UNSTABLE compound. It loses TWO water molecules to form GLYOXAL.



Fourth Step: Finally, Glyoxal is oxidized to produce OXALIC ACID.



• Result: Thus, OXALIC ACID is produced in this way by the REACTION of alkynes with KMnO₄.

