ludsid.µK CHEMISTRY Aliphatic Hydrocarbons



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CHAPTER 8 ALIPHATIC HYDROCARBONS

The Compounds which contain Hydrogen and Carbon only are Called hydrocarbons.

e-g methane (CH4), Ethane (C2H6).

There are two classes of hydrocarbons

- (1) open Chain hydrocarbons
- (2) Closed Chain hydrocarbons
- 1 open Chain RydroCarbons:-

HydroCarbons Which contain open Chain of Carbon atoms are Called open Chain hydro-Carbons or aliPhatic hydroCarbons The open Chain hydrocarbons have hoo types.

- (a) Saturated hydrocarbons (Alkanes)
- (b) Unsaturated Rydro Carbons (Alkenes and alkynes)
- (a) Saturated Rydro Carbons :-

The hydrocarbons in which all Carbon atoms are bonded by Single bonds are Called Saturated hydro Carbons or alkanes or farassins.

e-g methane, ethane, Peopune etc.

In Saturated hydrocarbons all four Valencies of Carbon are fully satisfied. Hence it can not take up any more hydrogien

(b) Un-Saturated Rydrocartuns:- The hydrocarbons Which Contain either a doubt or a tille bond

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are Called unsaturated RydroCarbons. e.g. $CH_2 = CH_2$ (e.Rene), $CH_2 = CH - CH_3$ (Propene), $CH = C - CH_3$ (Propyne) etc. The unsaturated RydroCarbons have two types.

(i) Alkenes: - The Unsaturated hydrocarbons Which Contain a double bond are called alkenes or olefins. e.g. CH2 = CH2, CH2 = CH-CH3 (ii) Alkynes: - The Unsaturated hydrocarbons Which Contain a triple bond are Called alkynes or acetylenes. e.g. HC = CH (EHYNE) HC = C-CH3 (Propyre)

(2) Closed Chain Rydrocarbons:-

The hydrocarbons which Contain ring of Carbon atoms are Called closed Chain hydrocarbons or Cyclic hydrocarbons. There are two types of cyclic hydrocarbons.

(a) Alicyclic hydrocarbons:- The hydrocarbons
Which Contain ring of Carbon atoms but
resemble open chain hydrocarbons are
Called alicyclic hydrocarbons. e.g. cycloPropane, cyclobutane, cyclo hexane
(b) Aromatic hydrocarbons:- The hydrocarbons
Which Contair at least one benzene ring
are called aromatic hydrocarbons. e.g.
benzene, Toluene, Maphthalene etc.

John i امولی فرست Nomenclature Common or Trivial system: In the Early days Organic compounds were named according to their history or method of Preparation. These names were called Common or trivial names. e.g is Methane as march gas because it is found in marshy Places Elected (ii) Methyl alcohol is called as wood spirit (iii) Acetic acid (Latin, acetum means vine gar /) (iv) Barbituric acid taken from the name of Chemist Barbara he common names are applicable to all the isomers. e.g CH3-CH2-CH2-CH3, CH3-CH-CH3 (n-butane)Three isomers of Pentane are (iso-butane) (n-pentane) CH-C-CH3 (iso-pentane)

(neo-Pentane) CH3

(neo-Pentane) CH3 The Common names of alkenes end at "ylene" $CH_2 = CH_2$, $CH_3 - CH = CH_2$, $CH_3 - CH = CH_2$ eBylene psopylene psopylene psopylene iso-butylene Defects of Common System:-(i) Common system is not applied for all Compounds (ii) Common system is not valid for Complex Compounds (iii) Common system does not explain molecular structure

JUPAC System: - The systematic method of nomenclature was formulated (given) by the International union of Pure and Applied Chemistry It is called IUPAC system. In this field first try was made by International Chemical Congress in 1889 - This Congress gave an incomplete report in 1892. In 1930, International union of Chemistry (IUC) gave Leige Rules For nomenclature. In 1947, Liege rules were modified by IUPAC. Since that date union has been issuing rules for systematic nomenclature of organic Compounds. The most recent form of these rules was Published in 1979. IUPAC system is based on the following Principle "Each different Compound Should have a different name By IUPAC System more than Seven million Organic Compounds have been named. Alkyl Groups: - When we remove one hydrogen atom from an alkane, we get an alkyl group. Its general symbol is R-. The name of an alkyl group ends in - yl. For example

(methane)

CH₃-CH₃
$$\xrightarrow{-H}$$
 $\xrightarrow{-H}$ $\xrightarrow{-H}$

Nomenclature of alkanes:- (

1:- Select the longest Continuous Chain of Carbon atoms. It gives the Parent name of alkane.

2:- Numbering begins from that End Which is neaver to the substituents (Side groups)

3:- Indicate the Position of all Substituents by the numbers.

4:- The substituents are named in an (alphabetical order) (التساوية)

5:- When two substituents are present on the same Carbon, use that number twice:
6:- When two or more identical substituents are present, then we use Prefix di, tri, tetra etc.

7: In Case of two equal Chains, the Chain with greater number of Substituents is Choosen. 8: When branching occurs at an equal distance from either end, Choose the Chain which gives lower number at the first point of difference 9: The names of first ten alkanes are methane ethane, Propane, butane, Pentane, hexane heptane, octane, nonane and decane.

i) H3C-CH2-CH2-CH2-CH3

CH3 2 - Methyl hexane

(ii) CH3-CH2-CH2-CH2-CH-CH3
2 CH2
1 CH3

3-Methyl Reptane
(iii) CH3-CH-CH2-CH3

CH3

CH3

CH3

4-EKyl-2 mcKyl hexanc



2,3,4-Trimethyl Pentane

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(V)
$$CH_3-CH_2-C-CH_2-CH_2-CH_3$$

 CH_3 3-EHyl-3 methyl hexane

2,3,5 trimethyl - 4-propyl heptane

2,3,5-Trimethylhexane (Not 2,4,5-trimethylhexane)

Nomenclature of Alkenes:-

1:- Select he longest continuous Chain of Carbon atoms Containing double bond (C=C

2:- Numbering begins from that end of Chair. which is nearer to the double bond.

3:- Indicate (1:/215) the Position (15) of the double bond by a number.

4: - Indicate the Position of each substituent group by a number.

5: - The name of an alkene ends at "ene"

6:- More han one double bonds are indicated by diene, triene, tetraene and so on.

Examples

(i)
$$CH_2 = CH - CH_2 - CH_3$$
 1 - Butene

(iv)
$$CH_3-CH=C-CH_3$$

 CH_3 $2-MelRyl-2-butene$
 CH_3 $CH_3-C-CH-CH=C-CH_3$

2.5.5-Trimethyl-2 hexene

(VI)
$$CH_3 - CH_2 - CH_2 - C = CH - CH_3$$

 $CH_2 - CH_2 - CH_3$

3-PROPYL -2 hexene



(Vii) $CH_2 = CH - CH = CH_2$ * 1, 3-Buta diene or Buta - 1, 3 - diene

(VIII) $CH_2 = CH - CH = CH - CH = CH_2$ 1, 3,5 - Hexatriene

Nomenclature of Alkynes

1:- Select the longest Continuous Chain of Carbon atoms Containing teiple bond (C≡C)

2:- Numbering begins from that end of Chain Which is nearer to the triple bond

3:- Indicate the position of Kiple bond by a number

4:- Indicate the Position of each substituent group by a number

5:- The name of an alkyne ends at "yne".

6:- A hydrocarbon Containing more than one triple bonds is named as digne, trigne tetragne and so on

7:- A hydrocarbon containing both double and triple bonds is Called enyne. It is named as followed.

(a) If double and triple bonds are present at identical fasitions, the double bond is



given the lower number.

(b) If double and triple bonds are not Present at identical Positions, the lowest Possible number can be given to a double or a triple bond.

Examples

(i)
$$CH \equiv CH$$
, $HC \equiv C - CH_3$
Ethyne Propyne

(ii)
$$CH \equiv C - CH_2 - CH_3 \quad 1 - Butyne$$

(iii)
$$CH_3-CH-CH_2-CH_2-C=CH$$

 CH_3 $5-Methyl-1-hexyne$

$$(iv) \quad CH = C - CH_2 - CH_2 - C = CH$$

1,5 Hexadiyne

$$(V)$$
 $CH = C - CH = CH - CH_3$

3 - Pentene - 1-yne

(Vi)
$$CH_2 = CH - C \equiv C - CH_3$$

1- Pentene - 3-Yne

(Vii)
$$CH = C - CH - CH = CH_2$$

 CH_3

3-Methyl-1-Penten-4-yne



Atto://www.guldasta.pk/f.sc Atkanes OR Paraffins

HydroCarbons in which all Carbon atoms are bonded by Single bonds are Called Saturated hydrocarbons or alkanes or Paraffins.

e.g. Methane (CH4), Ethane (C2H6), Propane and butane etc. Their general formula is CnH2n+2. In these Compounds all four valencies of Carbon are fully Satisfied. Hence they can not take up any more hydrogen

Preparation of Alkanes (1) By hydrogenation of Alkenes or alkynes (Sabatier and Sendern's Reaction)

Hydrogenation of alkene or alkyne in Presence of Nickel at 200–300°C gives alkane. It is called Sabatier and Sendern's reaction

 $\begin{array}{c} e \cdot g \\ R - CH = CH_2 + H_2 \xrightarrow{Ni} R - CH_2 - CH_3 \\ (Alkene) \end{array}$ (Alkene)

 $CH_2 = CH_2 + H_2 \xrightarrow{N_1} CH_3 - CH_3$ Ethene ethane

In hydrogenation Platinum or Palladium Can be also used as Catalyst . But they are very expensive (Sir) than Nickel.



(2) From alkyl Ralide :-

When alxyl Balide reacts with Zinc in Presence of an aqueous acid, then alkane is Produced . Examples are given below. $R = x + Z_n + H + \bar{x} \longrightarrow R - H + Z_n x_2$ (alxyl Ralide) $CH_3-I+Zn+H+I \longrightarrow CH_4+ZnI_2$ (methyliodide) CH3-CM-CH-CH3+ Zn+H+By -> CH3-CM-CH3+ InBr Hydrogenolysis: - Addition of Rydrogen accompanied by bond cleavage is called hydrogenolysis. Reaction of alight halide with Hz in Presence of Palladium-Charcoal gives alkane. $R-X+H_2$ $\frac{Pd/Charcool}{\Lambda} > R-H+H-X$

(3) By decarboxylation of mono-Carboxylic Acid Removal of Carbon dioxide from a Compound is called decarboxylation. The reaction of Sodium salt of Carboxylic acid with Soda Lime gives an alkane.

 $R-Co\bar{o}$ Na + NAOH <u>Cao</u> R-H + Na₂Co₃ CH3-CH2-COONA + NAOH CAO CH3-CH3+NAZCO3 Sodium Propionate



(4) Kolbe's Method: - It is an electrolytic method for Preparation of an alkane The electrolysis of Solution of Sodium Salt of mono-Carboxylic acid gives alkane. This method is used only for Preparation of symmetrical alkane (R-R) 2 RCOONA + 2 H20 Electrolysis - R-R+2CO2 + H2 + 2 NaOH 2 CH3COOK + 2H2O Electroly sis > CH3-CH3 + 2CO2 +2KOH + H2 The mechanism of reaction is given below. 2 CH3COOK water 2 CH3COO + 2K Pot acetate 2 CH3 COO ---> 2 CH3 COO + 2€] 2 CH3 COO ---> 2 CH3 + 2 CO2 | At anode CH3 + CH3 ---> CH3-CH3. 2H2O + 2E -> 2OH + H2] At Calhode $2K^{+} + 2OH \longrightarrow 2KOH$ Kolbe's method can not be used for the Preparation of methane



(5) From Grignard Reagents: Reachon of Grignard reagent with water or aqueous acid gives alkane.

CH3-M9-Br + H-OH EHREY CH4 + M9 OH

CH3-CH2-M9-Br + H-OH EHRY CH3-CH3 + M9-OH

Ethyl Magnesium Bromide

(ethane)

(6) From Carbonyl Compounds (Aldehydes or Ketones)

Aldehydes and Ketones are Called Carbonyl Compounds
Their reduction gives alkanes

in Clemmenson Reduction:

Reduction of Ketone in Presence of Zinc amalgam and hydrochloric acid gives alkane. In this case Carbonyl group of Ketone is reduced to methylene group (-CH2-)

 $\begin{array}{ll} CH_3 - C - CH_3 + 4[H] & \frac{Zn - Hg}{HCL} > CH_3 - CH_2 - CH_3 + H_2O \\ & \text{acetone} \end{array}$

(ii) Wolf-Kishner's Reduction: -

The reduction of an aldelyde in Presence of Rydrazine (N2H4) and KOH gives alkane. In this case carbonyl group of an aldelyde is reduced to methyl group.

 $\frac{CH_3 - C' - H + 4[H]}{200^{\circ}C} \xrightarrow{N_2H_3/KOH} CH_3 - CH_3 + H_2O$ acetaldehyde
(Ellane)



Physical Properties of alkanes

in Physical State: First four alkanes are

Colourless and odourless gases. The next 13

members (C5 — C17) are Colourless, odourless

liquids and higher members are waxylige solic

ii) Solubility: Alkanes are Soluble in non—

Polar Solvents like benzene, ether, CCly etc.

It is due to their non-Polar nature. Moreover

their Solubility decreases with increase in

molecular mass.

(iii) Melting points: The melting points of al-Kanes increase with the increase of the molecular mass but this increase is not very regular.

(iv) Boiling Points: — The boiling points of alkanes increase with the number of Carban atoms in a regular way. e-g boiling point increases by 20-30°C for addition of cach—CH2-group in alkanes. Moreover the normal or straight Chain isomer has higher boiling to point than its brunched Chain isomer. For example boiling point of n-butane is higher (55°C) than iso butane (-102°C)



Reactivity of alkanes: - Alkanes or Paraffins are less reactive compounds (Letin: Parum = titlle, Affins = affinity) The un-reachvity of alxanes is due to their non-polar bonds and inertness of s-bonds. The electronegativity of Carbon (2.5) and hydrogen (2.1) are nearly same. So bonding electrons show equal sharing between C-C and C-4 bonds Thus bonds in alkanes are non-folar Inertness of o-bond: In a or-bond the electrons are very typhtly hound between the nuclei. So a o-bond is very stable and a lot of energy is required to break it. Thus electrons of a o-bond can neither attack on an electrophile nor a nucleophile can attack on them. It is called incitness of o-bond However alkanes can show two types of reactions in Substitution reactions (ii) Thermal and Catalytic reactions (1) Combustion (Lie): - The burning of a substance in Presence of Oxygen is called Combustion. The complete combustion of alkane gives CO2, H20 and heat. The amount of heat evolved when one well of hydrocartor is burnt to CC2 and H20 is Called Reat of Combustion. E.g. CH4 + 2 02 Flant > CO2 + 2420 + 891 Kj mol (2) Oxidation: - Incomplete oxidation of CHE gives (O and Carbon It occurs in limited supply of oxygen 3CH4+402 Flame > 2CO+6H2O+ C

Catalytic Oxidation: The Catalytic Oxidation of methods occurs at high temperature and Pressure. It gives many useful Products

CH4 + [0] <u>Cu, 400°C/200 atm</u>, CH3OH (methyl alcohol.)

CH3OH + [0] <u>Cu, 400°C/200 atm</u>, HCHO + H2O

HCHO + [0] <u>Cu, 400°C/200 atm</u>, HCDOH (forme acid)

(Formaldelyde)

HCOOH + [0] <u>Cu, 400°C/200 atm</u>, CO2 + H2O

(3) Nitration: The Substitution reaction in which kydrigen atom of an alkane is replaced by nitro group $(-NC_2)$ is called Nitration $e \cdot g$

CH4 + HNO3 (conc) $\frac{450^{\circ}C}{}$ CH3NO2 + 420 (Nitromethane)

The nitroalkanes are used as fuels, Solvents and in organic synthesis.

or more hydrogen atoms of an alkane are replaced by halogen atoms is Called halogenation. It takes place in Presence of Sunlight or UV light. The yeachon of Iodine with alkane is highly violent and gives a mixture of Carbon, HF and fluorinated alkane.

The order of reactivity of hatogens is

$$F_2 > (\ell_2 > \beta_{r_2}) I_2$$

e.g CH4+Cl2 BD > CH3Cl +HCl

The reaction Proceeds through free radical mechanism and involves following three steps



Step II:- $Cl-Cl \xrightarrow{RD} \dot{Cl} + \dot{Cl}$ (Initiation) Step II:- $CH_3-H+\dot{Cl} \xrightarrow{RD} \dot{CH}_3 + HCl$ $\dot{CH}_3+cl-cl \xrightarrow{RD} \dot{CH}_3-cl+\dot{cl}$ (Propagation)

When Step II repeat again and again, a mixture of Products is obtained

 $CH_2Cl_2 + il \longrightarrow CHCl_2 + HCl$ $CHCl_2 + Cl - Cl \longrightarrow CHCl_3 + il$ (Chloroform or trickloromethane)

 $\begin{array}{ccc} \text{CHCl}_3 + \text{il} & \longrightarrow & \text{icl}_3 + \text{Hcl} \\ \text{icl}_3 + & \text{cl} - \text{cl} & \longrightarrow & \text{ccl}_4 + \text{il} \end{array}$

Carbon tetrachloride or Tetra Chloro me thane

Thus we may say that reaction of methane with Chlorine in Presence of sunlight gives a mixture of froducts. These Products are CH3Cl, CH2Cl2, CH2Cl3 and CCl4. These Products are important but this reaction is not synthetically important



Uses of Methane :-

(i) Methane is used as a fuel (varie)

(ii) Methane is used as an illuminating gas

(iii) Methane is used for Preparation of Carbon

black which is used in Paints, inks and tyres

(iv) Methane is used in manufacturing of Urea

(v) Methane is used for industrial Preparation

of methyl alcohol, formaldehyde, H2 and HCN.

(vi) Methane is used for Preparation of CH3CL,

CH2Cl2, CHCl3 (Chloroform) and CCl4

Alkenes

The unsaturated Rydrocarbons which Contain a double bond are Called alkenes or olefins.

e.g. CH2=CH2 (ekene), CH3-CH=CH2 (Propene)

Olefin is a Lahin word which means an oil forming compound. It is due to the reason that lower alkenes give oily Products with Cl2 or Br2.

Alkenes Containing one double bond are Called mono-enes. Their general formula is Cn H2n

Alkenes containing two double bonds are Called dienes and so on.

The simplest alkene (olefin) is ekene, C2 H4.



Preparation of alkenes

(1) By dekydrokalogenation of alkyl kalides

Removal of hydrogen builde (Hx) from hoo authorent carbon aloms of a compound is called dehydrohalogenation, e.g

 $CH_2 - CH_2Er + KCH(alc) - > CH_2 = CH_2 + H_2O + KBr$ Ehilbromide . (ethene)

 $CH_3-CH_1-BY+KOH(alc)->CH_3-CH_2-CH_2+110+KBY$ (n-Propyl bramide) (Propene)

(2) By delydration of alcohols:-

Removal of Water molecule from a substance is called dehydration. In this reaction Al_2O_3 , P_4O_{10} . Cons H_2SO_4 or H_3PO_4 is used as dehydrating agent. e.g.

CH3- CH2ON CORE H2504 -> CH2 = CH2 + H2O The case I terydrahon of alcohols is in the Yollowy I (1) by

Ter acetric > see alcohol > Pri-alcohol

CH3-CH2-CH2-CH- - 55 HASO CH3-CH3-CH3-CH3+H3O

CH3-C+-(H-0) (1/280) CH3-CH6-CH + H20

secondusy alcohol

$$CH_{3}$$
 CH_{3} C

(3) Dehalogenation of Vic-Dihalides

The dihalides having two hitogen atoms on two adjacential at Carbon atoms are Called Vicinal dihalides. The dehalogenation (removal of halogens) of Vic-dihalide gives alkene

2 F/12086 + Zn88

(4) By Partial Rydrogenation of alkynes:-

The Parhal or Controlled Endregenation of ARRINGS five alkenes. It is done in Presence of Suitable Catalist For example, in Presence of Lindton's Catalist an alkyne gives his alkene. (The finely divided Palladium Sufforted on Basou and Poisoned by quinoline is Catalist)

Called Lindlar's Catalyst)
$$R-C \equiv C-R + H_2 - \frac{Pd(Bascu)}{quinoline} = C = C - (Cis-alkene)$$

$$R-C \equiv C-R + H_2 - \frac{Na}{-33}C + R = C - R$$



(5) By electrolysis of Salt of dicarboxylic Acid (Kolbe's electrolytic method)

The electrolysis of aqueous solution of sodium succinate gives ethene. It is called Kolbe's method.

$$CH_2$$
—COONA $+2H_2O$ —Electrolysis $>$ $CH_2=CH_2+2CO_2+H_2$ CH_2 —COONA $+2H_2O$ — $+2NAOH$

The mechanism of reaction is given below

$$2H_2O + 2\overline{\epsilon} \longrightarrow 2OH + H_2$$

$$2Na + 2OH \longrightarrow 2NaOH$$
At Cathode

Physical Properties: - in First three ulkenes are gases, C5 to C15 are liquids and higher are solids (ii) They are water insoluble but alcohol soluble.

(iii) They have Larticular Smell.

(iv) They are Weakly Polis moustules.

Reactivity of a 11-bond:-

A TI-bond is formed by Parallel overlap of P-orbitals. In this case overlap of the P-orbitals is not as food as in a o'-bond. So TI-electrons are less firmly held by the atoms. Thus TI-bond is a weak bond as Compared to a o-bond. Therefore, it breaks easily due to attack of electrophilic reagent. Hence alkenes Show electrophilic reactions.

Reactions of Alkenes

(1) Addition of Hydrogen (Hydrogenation)

The reaction in Which Rydrogen is added to an alkene in Pressure of a Catalyst at a Pressure of 1—5 atm is called hydrogenation e-9

$$CH_{2} = CH_{2} + H_{2} \xrightarrow{Ni} CH_{3} - CH_{3}$$

$$FH_{3}$$

$$CH_{3} - CH - CH = CH_{2} + H_{2} \xrightarrow{Ni} CH_{3} - CH - CH_{2} - CH_{3}$$

$$3 - me flyl - 1 - butene$$

Raney Nickel:- When Ni-Al alloy is treated with NaOH, then Nickel is obtained in the form of fine suspension. It is called Raney Nickel. It is used as catalyst in hydrogenation $Ni-Al+NaOH+H_2O \longrightarrow Ni+NaAlO_2+\frac{3}{2}H_2$ eq $O+3H_2-\frac{Ni}{2}$ Cyclohexane

Markownikov's Rule: The rule states that in the addition of an unsymmetrical reagent to an unsymmetrical alkene, the negative Part of adding reagent goes to that Carten which has the least number of hydrogen atoms.

e-
$$g_{ii}$$
, CH_3 - CH = CH_2 + HBY -> CH_3 - CH - CH_3
 CH_3

(3) Addition of H2SO4:- When an alxene reacts with cold conc H2SO4; then alkyl hydrogen sulphate is formed. Then ethyl H-sulphate reacts with boiling water to give alcohol the overall reaction is the addition of water to an alxene So at is called hydration reaction. e.g.

 $CH_2 = CH_2 + H-O-SO_3H \longrightarrow CH_3-CH_2-O-SO_3H$ Ethyl kydrogen sulphate

CH₃-CH₂-0-SO₃,4 + H₂O $\xrightarrow{100^{\circ}C}$ CH₃-CH₂-0H + H₂SO₄

(4) Addition of Halogens:- Reaction of an alkene with halogen gives a vicinal distalide (1,2 distale alkane) $CH_2 = CH_2 + X_2 - \frac{CCL_4}{CH_2} \qquad CH_2 - CH_2$

 $CH_2 = CH_2 + 8Y_2 \longrightarrow CH_2 - CH_2 \qquad \text{vicinal di halide}$ $MeChanism: \qquad \qquad BY \qquad BY$ $C = CH_2 + 8Y_2 \longrightarrow CH_2 - CH_2$ $H = CH_2 - CH_2$ $H = CH_2 + 8Y_2 \longrightarrow CH_2 - CH_2$ $H = CH_2 - CH_2$ $H = CH_2 + 8Y_2 \longrightarrow CH_2 - CH_2$ $H = CH_2 - CH_2$ $H = CH_2 + 8Y_2 \longrightarrow CH_2 - CH_2$ $H = CH_2 - CH_2$ $H = CH_2 + 8Y_2 \longrightarrow CH_2 - CH_2$ $H = CH_2 - CH_2$ $H = CH_2 + 8Y_2 \longrightarrow CH_2 - CH_2$ $H = CH_2 - CH_2$ $H = CH_2 + 8Y_2 \longrightarrow CH_2 - CH_2$ $H = CH_2 - CH_2$ $H = CH_2 + 8Y_2 \longrightarrow CH_2 - CH_2$ $H = CH_2 + 8Y_2 \longrightarrow CH_2 - CH_2$ $H = CH_2 + 8Y_2 \longrightarrow CH_2 - CH_2$ $H = CH_2 + 8Y_2 \longrightarrow CH_2 \longrightarrow CH_2$ $H = CH_2 + 8Y_2 \longrightarrow CH_2 \longrightarrow CH_2$ $H = CH_2 + 8Y_2 \longrightarrow CH_2 \longrightarrow CH_2$ $H = CH_2 + 8Y_2 \longrightarrow CH_2$ $H = CH_2 \longrightarrow CH_2$ $H = CH_2$

 $CH_2 = CH_2 \xrightarrow{\mu} Br$ H = Br H = Br $H = CH_2 \xrightarrow{\mu} Br$ H = Br $H = CH_2 \xrightarrow{\mu} Br$ H = Br H =

In this reaction brown colour of bromine is discharged.

This test is used for the detection of a double bond.

(5) Addition of Hypo Ralous Acid (HOX): - Aqueous Solution of halegen reacts with an alxene and gives haleafooked or Ralohydrin. In this reaction Solvent molecule (120) also react.

 $X_2 + H_2O \longrightarrow HOX + HX$ $CH_2 = CH_2 + HOX \longrightarrow CH_2 - CH_2 \atop X OH (Ralo Kydrin)$

56 Oxidation Reactions of Alkenes

(1) Addition of Oxygen:- Alkenes react with oxygen in Presence of Silver as a catalyst and give epoxides. e.g

(i) $CH_2 = CH_2 + \frac{1}{2}O_2 \xrightarrow{AB_2O} CH_2 - CH_2$

Ethylene Oxide (Epoxide)

 $CH_3-CH=CH_2+\frac{1}{2}O_2\xrightarrow{A8_2O}CH_3-CH-CH_2$

The epoxides are starting materials for Preparation of glycols (Propylene Oxide)

(2) Combustion: Alkenes burn in air to Produce CO2 and 420.

 $C_2H_4 + 3O_2 \longrightarrow 2CO_2 + 2H_2O + heat$ Ethene

(3) Hydroxylation: - When an alkene is treated with 1% alkaline solution of KMnO7, (Baeyer Rengent) then vicinal glycol is formed. It is called hydroxylation. In this reaction Pink Colour of KMn04 discharges. It is called Baeyer's test. It is used for detection by of a double bond in

An organic molecure. C_q $3CH_2 = CH_2 + 2KMnO_4 + 4H_2O \xrightarrow{Cold} 3CH_2 - CH_2 + 2KOH$ $OH OH OH +2MnO_2$ Ethylene flycol

(4) Ozonolysis:- Ozone (03) reacts with an alkene to form an unstable molozonide. It further rearranges to form an Ozonide

$$CH_2 = CH_2 + O_3 \longrightarrow \begin{array}{c} O \\ O \\ CH_2 = CH_2 + O_3 \longrightarrow \\ CH_2 \longrightarrow CH_2 \end{array} \longrightarrow \begin{array}{c} CH_2 \\ CH_2 \longrightarrow CH_2 \end{array}$$

$$\begin{array}{c} CH_2 \\ CH_2 \longrightarrow CH_2 \end{array}$$

$$\begin{array}{c} CH_2 \\ O \longrightarrow O \\ O \longrightarrow O \end{array}$$

$$\begin{array}{c} O \\ O \longrightarrow O \\ O \longrightarrow O \end{array}$$

$$\begin{array}{c} O \\ O \longrightarrow O \\ O \longrightarrow O \end{array}$$

$$\begin{array}{c} O \\ O \longrightarrow O \\ O \longrightarrow O \end{array}$$

$$\begin{array}{c} O \\ O \longrightarrow O \\ O \longrightarrow O \end{array}$$

$$\begin{array}{c} O \\ O \longrightarrow O \\ O \longrightarrow O \end{array}$$

$$\begin{array}{c} O \\ O \longrightarrow O \\ O \longrightarrow O \end{array}$$

$$\begin{array}{c} O \\ O \longrightarrow O \\ O \longrightarrow O \end{array}$$

$$\begin{array}{c} O \\ O \longrightarrow O \\ O \longrightarrow O \end{array}$$

The ozonide reacts with Zinc and H2O to form aldehyde or Ketone.

$$CH_2 CH_2 + H_2O \longrightarrow 2 MCHO + H_2O_2$$

$$O \longrightarrow O \qquad \qquad (formaldeRyde)$$

$$H_2O_2 + Zn \longrightarrow ZnO + H_2O$$

This whole reaction is called 020nolysis

It is used to locate the Position of a double

bond in an alkene

Uses of Ethene:-

- is Ethene is used for artificial physics ripening of the fruit
- (II) Ethene is used as anaesthetic (is is)

 (III) Ethene is used for manufacture of Polythene

 which is used for making toys, bags, Cables

 (IV) Ethene is used as a starting material for

 many Chemicals c-J jlycol, ethyl alcohol etc.

The mustard gas is used as Chemical weapon. It was used in World War I. It is a fowerful vesicant and Causes Historia 121. Its Smell is mustard like the series in air like a mist in boiling liquid. It spreads in air like a mist in

2 CH2 = CH2 + S2Cl2 - S + S

SUEPhur monochloride CH2-CH3-Cl

Nustured gas

(b.f.-dichlero ethve suithinde)

(Vi) Oxy-Ethylise flame is used for wilding and cutting metals.

Alkynes

The unsaturated kidrocarbons which contain a triple bond are Called alkynes. Their general molecular formula is C_nH_{2n-2} . The first member of alkynes series is ethyne (Acetylene) Its formula is C_2H_2

Preparation of alkynes

(1) By dekydri Rologenation of dihalides

Removal of two molecules of hydrogen halide
from two adjacent Carbon gives an alkyne.

$$R-CH-CH-R_{+2KOH} Alcchol} > R-C=C-R+2KX+2H_2O$$

$$E \cdot g$$

$$CH_2-CH_2 + KOH Alcohol} > CH_2=CH+KSY+H_2O$$

$$CH_2=CH+KOH Alcohol} > CH=CH+KSY+H_2O$$

$$(EBYNC)$$

(2) By dehalogenation of tetrahalide

Removal of two molecules of halogens from a vicinal tetrahalide gives an alkyne

$$BY$$
 BY
 $CH-CH+Zn-CH=CH+3nBY_2$
 BY BY

$$CH = CH$$

$$| | + 2n \longrightarrow CH = CH + 2nB\gamma_2$$

$$BY$$

(3) By Kolbe's Electrolytic method:

The electrolysis of aqueous solution of Potassium maleate gives ethyne

CH-COOK
$$+2H_{2}O$$
 Electricity, CH $+2CO_{2}+2KOH+H_{2}$

CH-COOK CH

Potassium maleate $(Efkyne)$

The mechanism of the reaction is given as

 $CH-COOK$
 $CH-COOK$
 $CH-COO$
 $CH-COO$

Industrial Preparation of Ethyne: - Reaction of Calcium

CaC2 + 2H20 ----- C2H2 + Ca(OH)2

Physical Properties:-

in Alkynes are Colourless and odourless but ethyne has a garlic like odour.

(ii) First three alkynes are gases, next eight members are liquids and higher members are solids (iii) The melting Points, boiling points and densities of alkynes increase gradually with molecular mass (IV) Alkynes are Soluble in non-Polar Solvent like ether, benzene and CCl4

Reactivity of alkynes

Alkynes have Carbon to Carbon triple bond. In a triple bond, one bond is and how are 11-honds. The electrons are tightly held by two atoms. Thus these electrons are not easily available to he elechophilic reagents

Addition Reactions

(1) Addition of Hydrogen:

Hydrogenation of alkynes gives an alxane

$$CH \equiv CH + H_2 - \frac{Ni}{Heat} > CH_2 = CH$$

(2) Addition of Halogens :-

(2) Addition of Halojens:

$$HC = CH + Cl_2 - \frac{CCl_4}{CCl_4} \rightarrow \frac{CH = CH}{Cl_5} \stackrel{CH = CH}{Cl_5} \stackrel{CH}{Cl_5} \stackrel{CH}{C$$

(3) Addition of Halogen acids:

Ha Alkynes react with halogen acids (H-X) to form dihaloulkanes

(4) Addition of Water:-

Alkynes react with water to form an enol which rearranges to give aldehyde or ketone.

e-g

(i)
$$CH = CH + H - OH + H9SO_4, H_2SO_4 > CH_2 = CHOH$$

Vinyl alcohol

 $CH_2 = CHOH + Rearrangement$

(enol)

CH2 = CHOH Rearrangement CH3-CHO

$$(H_2 = CHOH) \xrightarrow{KCarrangement} CH_3 - CHO$$

$$(acetaldehyde)$$

$$(ii) CH_3 - C = CH + H_2O \xrightarrow{HgSO_4, H_2SO_4} CH_3 - C = CH_2$$

$$CH_3 - \dot{C} = CH_2 \xrightarrow{Rearrangement} O \xrightarrow{Rearrangement} O \xrightarrow{Rearrangement} CH_3 - C - CH_3$$

$$(enol) \qquad Acetone$$

(5) Addition of NH3 and Hydrogen Cyanide:-

$$CH = CH + NH_3 - \frac{Al_2O_3}{300C} -> CH_3 - C = N + H_2$$

$$CH = CH + HCN - \frac{Cu_2Cl_2 + NH_4Cl}{Heat} - CH_2 = CHCN$$

$$Vinyl Cyanide$$

$$(ACSYlonitrile)$$

6) Oxidation Reactions of alkanes:-

oxidation of ethyne gives alyoxal and further oxidation of glyoxal gives oxalic acid. It takes place in Presence of Strong alkaline KMn04

 $CH = CH + 2H_2O + 2(O) \xrightarrow{KM_{h}O_{4}} HC - CH$ OH OH OH OO OH OH $HC - CH \xrightarrow{-2H_2O} H-C-C-H(glyoxal)$ OH OH OO O OH $H-C-C-H + 2(O) \xrightarrow{KM_{h}O_{4}} COOH (oxabic acid)$

(ii) Combustion of alkyne: An alkyne burns in air or oxygen and gives CO2, H2O and Reat. e.g Combustion of acetylene is highly exothermic and oxyacetylene flame is used for Welding and Cutting metals

2 HC = CH + 502 ----> 4CO2 + 2H20 + Reat (acetylene)

(7) Polymerization: Alxynes show Polymerization and give linear and Cyclic Compounds. It depends upon temperature and Catalyst. It gives low molecular weight Polymers

in Linear Polymerization: Linear Polymensiden of acetylene gives vinyl acityline and di. 186 acetylene.

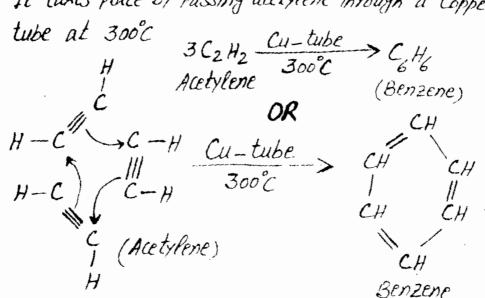
 $HC = CH + HC = CH - \frac{Cu_2 cl_2 + NH_3 cl_4}{Several hours} HC = CH - C_3 = CH$ Acelylene

(B.1. - 1.0. - 1.0.

 $H_2C = CH - C = CH + HC = CH - \frac{Cu_2Cl_2}{NH_4Cl} > CH_2 = CH - C = C - CH = CH_2$ (divinyl acetylene) Hexa-1,5 diene -3-yne

(ii) Cyclic Polymerizanan:

The cyclic Polymerization of acetylene gives benzene Il taxes Place by Passing acetylene through a Copper



Acidic Nature of Alkynes:-

Ethyne and any other terminal alkyne shows acidic, Properties. There is sp-hybridization in ethyne. An sp-hybrid orbital has so S - Character and approaches spherical shape. So elections are close and strongly held by Carbon atom. Therefore H-atom becomes slightly acidic e 9 +8 -8 -8 +8 H-C=C-H

```
For example
                           > Na C = CNa + H
   HC = CH +2Na
                            Sodium Acetylide
   Acetylene
                          > R-C=CNa+NH3
R-CECH + NaNH2-
          Sodamide
                               Sodium Alkynide
 Copper acetylide is Prepared by reaction of
 acetylene with ammonical cuprous Chloride
HC=CH+CU2Cl2+2NHyOH->CUC=CCU+2NHyCl+
                          Copper acetylide
Silver acetylide is formed as follows. (red PPt)
HC=CH+2AgNO3+2NHyOH->AgC=CAg+2H2O+2NHNO3
                        Disilver acetylide
                          (White ppt)
Acetylides react with dilute acids to regenerate
```

acetylene.

Ag C = CAg + H2SO4 (dil) -> HC=CH + Ag2 SO1, AgC = CAg + 2HNO3(dil) -> HC = CH + 2AgNO3

Uses : - Acetylides (Alkynides) are used for the Preparation, Purification and Separation of alkynes (ii) Acetylides are used to identify the terminal and non-terminal alkynes. Terminal alkynes give acetylides e g CH3-C=CH(Propyne) Non-terminal alkynes do not give acelylides (2-Butyne)

Uses of Ethyne:-

in Ethyne is used for artificial ripening of fruit.

(ii) Oxy-acetylene flame is used for cutting and welding metals

illi) Acetylene is used for Preparation of alcohol acetic acid and acetaldehyde

iv: Acetylene is used for manufacture of PVC, Polyvinyl acetate, Orlon and neoprene rubber.

(V) Acetylene is used to Prepare acetylene tetrackloride which is used as Solvent for rubber, Varnish, resins.

Chloroprene and Neoprene

Vinyl acetylene reacts with Conc HCl to give a froduct Called Chloroprene cl

 $CH_2 = CH - C = CH + HCl - CH_2 = CH - C = CH_2$ Vinylacetylene (Chloroprene)

2-Chloro-1, 3-butadiene

The Chlorofrene on folymerization gives neofsene which is used as synthetic rubber

Chlorofrene Polymerization > neofrene (Synthetic rubber)

Soda Lime :-

When lime (CaO) is Soaked with Solution of NaOII. Then the dried Psoduct is Called Soda - Lime -

Comparison of Reactivities of alkanes, alkenes and alkynes.

The yeachivity of alkanes, alkenes and alkynes decrease in the following order.

Alkenes > Alkynes > Alkunes

Alkanes have all sonds. A solond is a strong bond and Cannot be easily broken. Alkenes have double bond in which one is solond and other is TI-bond. The TI-bond is a weak bond and breaks easily. Thus an electrophile Can attack TI-electrons of alkenes Alkynes have a triple bond. In a triple bond, one is solond and other two are TI-bonds. The bond length of two triply bonded Carbons is very short. So electrons are tightly held by two atoms. Thus TI-electrons in alkynes are not available to the electrophilic reagents. Hence Order of reactivity is.

Alkenes > Alkynes > Alkanes

Hydrogenation of vegetable oils

The vegetable oils are unsaturated compounds
When hydrogen is added to vegetable oil in
Isesence of Nickel, then vegetable ghee is Producea
It is called hydrogenation or hardening of oils.

EXERCISE

Q1.	Fill in the Blanks.								
(i)	Ozone macts with othere to form of alkynes.								
(ii)	Lindlar's catalyst is used for of alkynes.	of aikynes.							
1111	Divinylacetylene is a acetylene.								
(iv)	Vicinal dinaides have two halogens oncarbon atoms.								
(V)	Ethyne is acidic in character because of hybridization.								
(vi)	Ethyne is acidic in character because of hybridization. Rainhydrins are formed due to addition of in ethene.								
{vii)	Ethylene glycol is produced whenreacts with cold alkaline KMnO ₄ solution.								
(viii)	Mustard gas is a highly bolling 65								
(x!)	Ethyne haslike odour.								
(x)	Ethyne is obtained by the reaction ofwith calcium carbide) .							
Ans	wer:-(i) ozonide (ii) partial hydrogenation (iii) polymer of (iv) adjacent (v) sp- (vi) hypohalous acid (vi) ethene (viii) liquid (ix) garlic (x) water								
	(Att) attients (Att) tident (IX) Aprile (X) aprile								
Q2.	Indicate True or False.								
(1)	Addition of HX to unsymmetriacal alkenes takes place according to Markowinkov's rule.								
(iii)									
	Mustard gas is a blistering agent.								
	Resident gas is a bustering agent. Baeyer's reagent is used to locate a double bond in an alkene.								
	Ethyne is a saturated compound.								
(vi)	Baeyer's reagent is used to locate a double bond in an alkene.								
(VIII)	Alkanes usually undergo substitution reactions.								
	Benzene is a polymer of ethene.								
	Acrylenitrile can be obtained from ethyne.								
	Ethyne is more reactive towards electrophilic reagents than ether	16							
Ans	swer:-(i) true (ii) false(iii) true (iv) true (v) false								
	(vi) true (vii) true (viii) false(ix) true (x) false								
Q3.	Multiple Choice questions. Encircle the correct answer.								
(1)	Preparation of vegetable ghee involves.								
	(a) Halogenation (b) Hydrogenation								
	(a) Hatogenation (b) Hydrogenation (c) Hydroxylation (d) Dehydrogenation								
$\{i^j\}$	Formula of Chloroform is.								
` '	(a) CH ₃ Ci (b) CCl ₄ (c) CH ₂ Ci ₂ (d) CHCi ₃								
(111)									
1124	(a) Saturation (b) Unsaturation (c) Substitution (d) None								
livi	Vinviacetylane combines with HCl to form								
·1	(a) Poly acetylene (b) Benzene								
	(a) Chloropene (d) Divinul acetylene								

(v)	The addition	n of unsyn	nmetrical :	eagent t	o an uns	ymmetr	ical
	alkene is in	accordan	ce with the	rule		•	
(a)) Hund's re	ulo		(b) Ma	rkowink	liov's ru	le
(c) Pauli's E	xclusion P	rinciple				
Ivil Su	nthetic rubl	ber is mad	e by Polyn	nerizatio	n of		
	Chlorofo					00 (d) 5	lutono
		, ,	•		-	, .	outene
	3' – dichloro						Dia
(a)	Mustard ga	is (b) Laii	ghing gas	(c) Ph	osgene	gas (a)	Bio-gas
(viii) W	hen methand	e reacts w	ith Cl ₂ in tl	ne prese	nce of di	ffused !	ight the
pr	oducts obta	ined are.		-			_
(a)	Chlorofo	rm only	(b)	Carbon	Tetrach	toride or	niv
	Chlorome						
	nich one of th						
	Ethene						
/~/		(2) 2:::	(0)		(4)	, .ора	•
Answe	r:-(i)b	(ii) d	(iii) b	(iv) c	(v) t)	•
		(vii) a			` '		
			,				
	rite the Struc					ig comp	ounds.
(i)	2-Methylp	propane	(ii)	Neopen	tane.		
(111)) 3-Ethype	ntane	(iv)	4-Ethyl-	3.4-dime	thylhep	tane.
	2,2,3,4-Te						
fvi	i) 22 Dime	thulbutan		1988 2 2	Dimath	vincons	20
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Answe	r:-			111			
				<i>Υ</i> ′3			. /
1. 2	MOKUP (enhama.	211		. ::	Neo	bentane
(1) 2-	ין אינוארן	ruparie	CH3-6	\mathcal{H} \mathcal{L} \mathcal{H}	2 (11)	,	/11
		•	_	`			pentane CH3 - C-CH3 CH3
.iii. 2	0.90	Dontan	0			CH	-6 011
(111)	-CDYC-1	resilusi	C			~73	-C-CH3
`				, ,			,,,
	CH3-CH2	-CH-	CH3-C1	43			CH2
· ·	-13 -12	• /	9	~			J
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			52	<i>4</i> ~			
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	<i></i>		1	-112	~2		•
		באי.	2 /11				
		-10	5 673				
	4-EMYC CH3-C		_		^ /		
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	CH2-1	J-CH	C (Ha			
	J. 1	5. /	1	3			
		H3 CH2	CHZ				
		5 -7.5	T'/_3				

Q5. Write down names of the following compounds according to IUPAC system.

(v) CH₃CH₂C(CH₃) ₂CH(CH₂CH₃)CH₃ (vi) (CH₃CH₂)₃CH (vii) CH₃C(CH₃)₂(CH₂)₂CH₃ (viii) (C₆H₅) ₃CH

Answer:- (i)
$$CH_3-CH_2-CH-CH_2-CH_3$$

$$CH_2-CH_3 \quad 3-Ethyl Pentane$$

$$CH_3 \qquad CH_3$$

$$(ii) \quad CH_3-CH-CH_2-C-CH_3$$

$$CH_3 \qquad CH_3$$

$$CH_3 \qquad CH_3$$



2,2,4:trimethyl Pentane



Q6. What are the rules for naming alkanes? Explain with suitable examples.

Answer:- see page No. 35

 (a) Write down the structural formulas for all the isomeric hexanes and name them according to IUPAC system.

The following names are incorrect. Give the correct IUPAC names.

4-Methylpentane

(ii) 3,5,5,-Trimethylhexane

(iii) 2-Methyl-3ethylbutane

Answer: (a) ISOMEYS OF Rexanes:-

4-Methyl-Pentane (incorrect)

2,2,4 teimethyl hexane (correct)
(iii) CH3-CH-CH-CH3 2-methyl 3-ethyl butan.
CH3 CH2-CH3 (incorrect)

2,3-Dimethyl-Pentane (Correct name)

- Q8. (a) Explain why alkanes are less reactive than alkenes?
 What is the effect of branching on the melting point of alkanes?
 - (b) Three different alkenes yield 2-methylbutane when they are hydrogenated in the presence of a metal catalyst. Give their structures and write equations for the reactions involved.

Answer:- (a) see page No. 46

(b)
$$CH_2 = C - CH_2 - CH_3 + H_2 \xrightarrow{Ni} > CH_3 - CH - CH_2 - CH_3$$

 CH_3
 $CH_3 - C = CH - CH_3 + H_2 \xrightarrow{Ni} > CH_3 - CH - CH_2 - CH_3$
 CH_3
 $CH_3 - CH - CH = CH_2 + H_2 \xrightarrow{Ni} > CH_3 - CH - CH_2 - CH_3$
 CH_3
 $CH_$

- Q9. (a) Out line the methods available for the proparation of alkanes.
 - (b) How will you bring about the following conversion?
 - (i) Methane to ethane (iii) Ethane to methane
 - (iii) Acetic acid to ethane (v) Methane to nitromethane

Answer:- (a) see page No. 41,42,43

CH3-CHO + [O]
$$\frac{K_2C_2O_7}{H_2SO_4}$$
 > CH3 COOH

CH3 COOH + NAOH — > CH3 COONA + H2O

CH3 COONA + NAOH $\frac{CaO}{O}$ > CH4 + NA2 CO3

(iii) Acetic acid to ethane:

CH3 COOH + NAOH — > CH3 COONA + H2O

2 CH3 COONA + 2H2O Electric > CH3 - CH3 + 2CO2 + 2NAOH + H2

(iv) Methane to nitromethane:-

CH4 + HNO3 $\frac{450^{\circ}C}{O}$ > CH3 NO2 + H2O

(CONC) Nitromethane

Q10. (a) What is meant by octane number? Why does a high octane fuel has a less tendency to knock in an automobile engine?

(b) Explain free radical mechanism for the reaction of chlorine with methane in the presence of sunlight.

- Q11. (a) Write structural formulas for each of the following compounds.
 - (i) iso-Butlyene (ii) 2,3,4,4-Tetramethyl-2-pentene

(iii) 2,5-Heptadiene

(iv) 4,5-Dimethyl-2-hex,ene

(v) Vinylacetylene

(vi) 1,3-Pentadliene

(vii) 1-Butyne

(viii) 3-n-Propy_i 1,4-pentadiene

(ix) Vinylbromide

(x) But-3-en-1-yne

(xi) 4 Methyl-2 pentyne (xii) iso-Pentane

(b) Name the foll-owing compounds by IUPAC system.

(i) $H_3C--CH = CH(CH_2)_2CH_3$ (ii) $(CH_3)_2C = CH_2$

(iii)
$$CH_3 - CH_2 - CH_2 - C = CH_2$$
 (iv) $CH_2 = CH - CH = CH_2$

$$CH_1 - CH_2 - CH_2 - CH_2$$

$$CH_2 - CH_3 - CH_2 - CH_2 - CH_3$$

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(v)
$$CH_2=C-CH_2CH_2CH_3$$
 (vi) $CH=C-CH_3$
 C_2H_5

(vii) $CH_3=C=C-CH_5$ (viii) $CH_2=CH-C=C-CH=CH_2$

(ix) $CH=C-CH=CH-C=CH$ (x) $CH_2=CH-C=CH$

Answer:-
(i) $CH_3-C=CH_2$

CH3 (iSObutylene)

(ii) 2,3,4,4 tetramethyl 2-pentene

CH3 CH3 CH3

CH3

CH3-C=C-C-CH3

CH3

CH3-CH=CH-CH2-CH=CH-CH3

(iV) 4,5-dimethyl-2-kexene

CH3-CH=CH-CH-CH-CH-CH3

(V) Vinyl acetylene (Vi) 1,3 pentadiene

 $CH_2=CH-C=CH$
 $CH_2=CH-CH=CH-CH_2$

(Vii) 3-n-peopyl 1,4 pentadiene

 $CH_2=CH-CH-CH-CH_2$
 $CH_2=CH-CH-CH-CH_2$
 $CH_2=CH-CH-CH-CH_2$
 $CH_2=CH-CH-CH_2$
 $CH_2=CH-CH-CH_2$
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 $CH_2=CH-CH_2$

(*) But -3-ene-1-yne,
$$CH = C - CH = Cn_2$$
.

(**) $4 - Methyl, 2 + Pentyne, CH_3 - C = C - CH - Ch_2$.

(**) iso Pentane

 $CH_3 - CH - CH_2 - CH_3$
 $CH_3 - CH_2 - CH_2 - CH_3$
 $CH_3 - CH_2 - CH_2 - CH_3$

(ii) $(CH_3)_2 C = CH_2$ or $CH_3 - C = CH_2$
 $CH_3 - CH_2 - CH_2 - C = CH_2$

(iii) $CH_3 - CH_2 - CH_2 - C = CH_2$
 $CH_3 - CH_2 - CH_2 - CH_2 - CH_3$

(iv) $CH_2 = CH - CH - CH_2$, 1, 3 but adiene

(v) $CH_2 = CH - CH_2 - CH_2 - CH_3$
 $CH_3 - CH_3 - CH_3 - CH_3 - CH_3 - CH_3 - CH_3$

(viii) $CH_2 = CH - C = CH_3 - CH_3 - CH_3 - CH_3$

(viii) $CH_2 = CH - C = CH_3 - CH_3 - CH_3$

(viii) $CH_2 = CH - C = CH_3 - CH_3 - CH_3$
 $CH_3 - CH_3 - CH_3 - CH_3 - CH_3 - CH_3$

(viii) $CH_3 - CH_3 - CH_3 - CH_3 - CH_3 - CH_3 - CH_3$

(viii) $CH_3 - CH_3 -$

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Q12. (a) Describe different methods for the preparation of alkenes. How would you establish that ethylene contains a double bond? (b) Give structural formulas of the alkenes expected to form by the dehydrogenation of the following compounds by a strong base. 1-Chloropentane (ii) 3-Chloro-2-methyl butane 1-Chloro-2,2-dimethylpropane. (iii) Answer:- (a)see page No. 50.51,52 Butane CH2-C-CH3+KOH ale > NO Reaction In this case no reaction takes Place because B-hydrogen is not available Q13. (a) Write down chemical equation for the preparation of propens from the following compounds. CTT CH (ii) CH₃ CH3-CH2-CH2 OH (iii) iso-Propyl chloride (b) Write skeleton formula showing only the arrangement of carbon atoms for all the possible alkenes of the molecular formula C5H10. Answer:- (a) (i) $CH_3-CH_2-CH_2OH\frac{Conc\ H_2SO_4}{170°C}$ $CH_3-CH=CH_2+H_2O$ I-Propanol Propenc(ii) $CH_3-C=CH+H_2\frac{Pd}{BaSO_4}$ $CH_3-CH=CH_2$ Propene

(iii)
$$CH_3-CH-CH_3+KOH\frac{alc}{-}CH_3-CH=CH_3+KCl+H_2CCl}(isofropyle Chloride)$$

Professible

alkenes of molecular formula C_5H_{10}

(i) $CH_2=CH-CH_2-CH_3$ 1-Pentene

(ii) $CH_3-CH=CH_2-CH_3$ 2-Pentene

(iii) $CH_3-CH=CH_2-CH_3$ 2-Pentene

(iii) $CH_2=C-CH_2-CH_3$ 2-Pentene

(iv) $CH_2=C-CH_2-CH_3$ 2-Pentene

(iv) $CH_2=C-CH_2-CH_3$ 2-Pentene

(iv) $CH_3-CH_3-CH_3$ 2-Pentene

(V)
$$CH_3-C_1=CH-CH_3$$
 2 methyl-2-Bulene CH_3

Q14. (a) How may ethene be converted into ethyl alcohol?

(b) Starting from ethene buttine the reactions for the preparation of following corepounds.

(i) Ethylene dibromide (ii) Ethyne (iii) Ethane (iv) Ethylene glycol

(c) How will you bring about the following conversions.

(i) 1-Butene to 1-Butyne

(ii) 1-Propanol to CH₃ = CH CH₂Ct

Answer:-(a)
$$CH_2 = CH_2 + H_2O \frac{10\% H_2SO_4}{H_9SO_4} > CH_3 - CH_2OH$$

(b) (II) $CH_2 = CH_2 + B\gamma_2 - > CH_2 - CH_2$
 $S\gamma$ $S\gamma$
 $CH_2 - CH_2 + 2KOH alc$
 $S\gamma$ $S\gamma$
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For facts (i), (iii) and (iv) see reactions of alkenes

(C):-
$$CH_2 = CH - CM_2 - CM_3 + Br_2 \longrightarrow CH_2 - CM - CM_3 - CM_3$$

(i) $1-Butene$
 Ry
 By
 By

(ii)
$$CH_2 = CH - CH_2 - CH_3 + BY_2 - CH_2 - CH - CH_2 - CH_3$$

(iii) $BY BY$
 $CH_2 = CH - CH_2CH_3 + 2KM_DO_4 + 4H_2O - OH_3 - CH_2CH_2CH_3$
 $CH_2 = CH - CH_2CH_3 + 2KM_DO_4 + 4H_2O - OH_3 - CH_2CH_2CH_3$
 $CH_2 = CH - CH_2CH_3 + HBY - CH_3 - CH_3 - CH_2CH_3$
 $CH_3 - CH_3 - CH_2CH_3$
 $CH_3 - CH_3 - CH_3 - CH_3CH_3$
 $CH_3 - CH_3 - CH_3CH_3$
 $CH_3 - CH_3 - CH_3$
 $CH_3 - CH_3$
 CH

(ii) Propene
$$\xrightarrow{Br_2}$$
 $\xrightarrow{KOH(alc)}$ \xrightarrow{HCN} \xrightarrow{F}

Answer:-(i) CH_3-CH_2OH \xrightarrow{ConC} $\xrightarrow{H_2SO4}$ \xrightarrow{C} $\xrightarrow{H_2}$ \xrightarrow{C} $\xrightarrow{H_2}$ \xrightarrow{C} $\xrightarrow{H_2}$ \xrightarrow{C} $\xrightarrow{H_2}$ \xrightarrow{C} $\xrightarrow{H_2}$ \xrightarrow{C} $\xrightarrow{H_2}$ \xrightarrow{Br} \xrightarrow{Br} \xrightarrow{Br}

$$CH_2-CH_2 + 2KOH \xrightarrow{alc} > CH = CH + 2KBY + 2H_2O$$

$$BY \quad BY$$

$$(II) \quad CH_2 = CH - CH_3 + BY_2 \longrightarrow CH_2 - CH - CH_3$$

$$PYOPENE \quad BY \quad BY$$

$$CH_2-CH-CH_3 + 2KOH \xrightarrow{alc} > HC = C-CH_3 + 2KBY + 2H_2O$$

$$BY \quad BY$$

$$HC = C-CH_3 + HCN \longrightarrow CH_2 = C-CH_3$$

$$PYOPYNE \quad CN 2-CYANO PYOPENE$$

Q18. After an ozonolysis experiment, the only product obtained was acetaldehyde, CH₃CHO. The chemist who did the experiment correctly claimed that two possible starting materials can give this product. Can you guess the structural formulas of these compounds.

Q.19.(a) The addition of Sulphuric acid to an alkene obeys
Markownikov's rule. Predict the structures of the alcohols
obtained by the addition of the acid to the following compounds.

(i) Propene (ii) 1-Butene (iii) 2-Butene

(b) Predict the most likely product of the addition of hydrogen chloride to 2-methyl-2-butene. Explain the formation of this product.

Answer:- (a) (i)
$$CH_2 = CH - CH_3 + H_2SO_0 \longrightarrow CH_3 - CH - CH_3$$

 $CH_3 - CH - CH_3 + H_2O \longrightarrow CH_3 - CH - CH_3$
 $OSO_3 + H_2SO_0$
 $OSO_3 + H_2SO_0$
 $OSO_3 + H_2SO_0$

(ii)
$$CH_2 = CH - CH_2 - CH_3 + H_2SO_4 \longrightarrow CH_3 - CH - CH_2 - CH_3$$

$$CH_3 - CH - CH_2 - CH_3 + H_2O \longrightarrow CH_3 - CH - CH_2 - CH_3$$

$$OSO_3H + H_2SO_4 \longrightarrow CH_3 - CH_2 - CH_3 + H_2SO_4$$

$$CH_3 - CH = CH - CH_3 + H_2SO_4 \longrightarrow CH_3 - CH_2 - CH_3 - CH_3 - CH_2 - CH_3 - CH_3 - CH_2 - CH_3 -$$

Q.20.Why are some hydrocarbons called structured and others unsaturated? What type of reactions are characteristics of them?

Answer:- see page No. 31, 32

- Q21. (a) Describe methods for the preparation of Ethyne.
 - (b) How does ethyne react with:
 - (i) Hydrogen (ii) Halogen acid (iii) Alkaline KMnO₄
- (iv) 10% H2SO4 in the presence of HgSO4. (v) Ammonical cuprous chloride
 - (c) Mention some important uses of methane, ethene and ethyne.

Answer: - see page No. 59,60,6/

Q22. Describe how you could distinguish ethane, ethene and ethyne from one another by means of chemical reactions.

Answer: Distinction between Ethane, Ethene and Ethyne

(2) Reachon with KMn04:- Ethane does not react with alkaline KMn04 but ethene and ethyne react

with KMnOy and decolourise its fink colour.

CH3-CH3 + KMnO4 - OH > NO reaction

3CH2=CH2+2KMnC1+4H20-01->3CH-CH2+2KON42KONO. ethene OH OH + 2KON + 8KON +

exalic acid

(22) Reaction with Brz Water: - Ethane does not decolourise By, water but elbere and offyne decolourise bromine water.

CH3-CH3 + BY2 ----> NO YEachon $CH_2 = CH_2 + Br_2 \longrightarrow CH_3 - CH_3$ $CH = CH + 2Br_3 \longrightarrow F$ CH = CH + 28%, ____

(222) Reaction with Ammonical 67 67 Silvey Nitrate

Ehane and essence do not react with ammonical Ag NO3 but ethyne forms white 17th of Silver acetylide.

CH=CH+2A9NO3+2NH,OH-> A9C= CA9+2NH,NO +211,0 (Silver acetylide)

(iv) Ehane and ehene do not react wif ammenical Cuprous Chloride but acetylene forms red Ist with ammonical cuprous Cheoride.

HCECH+ Cu2Cl2+2NHGCH--> CUCECCU+INNGC+2HO nd Mi of corrows acetylide Q23. (a) How will you synthesize the following compounds starting from ethyne.

(i) Acetaldehyde (ii) Benzene (iii) Chloroprene

(vi) Glyoxal (v) Oxalic acid (vi) Acrylonitrile

(viii) Ethane (viii) Methyl nitrile

(b) Write a note on the acidity of ethyne.

Answer:- (a) see page No. 61,62,63

(b) see page No. 64

Q24. (a) Compare the reactivity of ethane, ethene and ethyne.

(b) Compare the physical properties of alkanes, alkenes and alkynes.

Answer: - see page No. 67

Q25. How does propyne react with the following reagent.

(a) AgNO3/NH4OH (b) Cu2Cl2/NH4OH (c) H2O/H2SO4, HgSO4

Answer:-

(b)2(
$$H_3$$
- $C \equiv CH + Cu_2(l_2 + 2NH_4OH \rightarrow 2CH_3-C \equiv CCu + 2NH_4Cl$
Propyre (Copper Propylide) +2H2O

(C)
$$CH_3-C=CH+H_2O \xrightarrow{10\%} H_2SO_4 > CH_3-C=CH_2$$

$$CH_3-C=CH_2 \xrightarrow{\text{rearrangement}} CH_3-C=CH_3 \text{ (acetone)}$$

Q26. A compound has a molecular formula C₄H₆, when it is treated with excess hydrogen in the presence of Ni-catalyst, a new compound C₄H₁₀ is formed. When C₄H₆ is treated with Ammonical silvernitrate a white precipitate is formed. What is the structure formula of the given compound.

Answer:- Reaction of Compound (C4Hz) with H2 indicates that Compound Contains two double bonds or one triple bond in it. But reaction of the Compound with ammonical Siever niteate indicates the



Presence of acidic hydrogen or terminal triple bond. Thus the Compound (C4H6) is 1-butyne. The reactions of 1-Butyne with H2 and ammonical AgNO3 are given below.

 $HC = C - CH_2 - CH_3 + 2H_2 \xrightarrow{N_1} CH_3 - CH_2 - CH_2 - CH_3$ 1 - butyne n - butane

HC=C-CH2-CH3+A9NO3+NH40H-->A9C=C-CH2-CH3+H20H--->

Q27. (a) Identify A and B.

Conc H₂SO₄ Br₂
CH₃CH₂CH₂OH ———— A . - —— B

(b) Give the general mechanism of electrophilic addition reactions of alkenes.

Answer:-

(a)
$$(H_3-CH_2-CH_2OH) \frac{Conc H_2SO_4}{>} CH_3-CH=CH_2+H_2O$$

Propene

 $CH_3-CH=CH_2+Br_2 \longrightarrow CH_3-CH-CH_2$
 Br
 Br
 Br
 $1,2$ dibromo-Propane



These Notes Have been Prepared and Developed By

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السلام عليكم ورحمته الله وبركاته

مخقب تعبادني

کافی عرصہ سے خواہش تھی کہ ایک ایسی ویب سائٹ بناؤں جس پر طالب العلموں کیلئے تعلیمی مواد جمع کر سکوں۔ اللہ تعالی نے توفیق دی اور میں نے ایک سال کی محت کے بعد ایک سائٹ "گلدستہ ڈاٹ پی کے "کے نام سے بنائی جو کہ قرآن و حدیث، اصلاحی، دلچسپ، تاریخی قصے واقعات، اُردو انگاش تحریریں، شاعری و اقوال زریں، F.Sc اور B.Sc کے مضامین کے آن لائن نوٹس، اسلاک، تفریحی، معلوماتی وال پیپرز، حمد و نعت، فرقہ واریت سے پاک اسلامی بیانات، پنجابی تظمیس و ترانے اور کمپیوٹر و انٹرنیٹ کی و نیا کے بارے میں ٹمپس، آن لائن کمائی کرنے کے مستند طریقہ کار۔ کے ساتھ ساتھ اور بھی بہت سی چیزوں پر مشمل ہے۔ اور انشاء اللہ میں مزید وقت کے ساتھ ساتھ اور بھی بہت سی چیزوں پر مشمل ہے۔ اور انشاء اللہ میں مزید وقت کے ساتھ ساتھ اضافہ کرتا جاؤں گا۔ آپ کی قیمتی رائے کی ضرورت ہے۔ عرفان شفیق ساتھ ساتھ اضافہ کرتا جاؤں گا۔ آپ کی قیمتی رائے کی ضرورت ہے۔ عرفان شفیق

انهم نوط

ذیل میں جو نوٹس مہیا کیے گئے ہیں وہ کئی گھنٹوں کی لگاتار محنت کے مرتب ہوئے ہیں۔ اور آپ کو بالکل مفت مہیا کر رہے کیے جارہے ہیں۔ ان کی قیمت صرف اتن سی متوقع ہے کہ ایک بار ہیں۔ آپ سے ان کی قیمت صرف اتن سی متوقع ہے کہ ایک بار ورود ابراھیمی اپنی زبان سے ادا کر دیں۔

يئمني لأكمال يتحمل لتصحيف <u> اللهُ يَّصَلِّعُ إِلَّهُ مُحَمَّلًا مُحَمَّلًا اللهُ يَحَمَّلُهُ اللهُ عَلَيْهُ اللهُ مُحَمَّلًا ل</u> وَتَكُولُونَا لَنُ مُحَمِّدُ لَكُمَا صَلَّتُكَ عَلِي إِبْرَاهِمْ وَعَهِلِ اللهِ إِبْرَاهِمُ مَ انَّكَ *جَمَّنْ*كُ هُجَنْكُ هُ ٲڵڵڮؙڂؾؠٙڽٳۯػ^ۼڸٳؽ۫ۼؙڲؠۜڒٷۜۼڵؚؖؽ النجائك بالأثاكات عالى ابراهمي وعكاني ال إبراهمي اِنَّاكَ حَمَٰكُ أَجْجَيُكُهُ