

Solution: CH = CH  $\xrightarrow{\text{H}_2SO_4,\text{H}_2O}$  CH<sub>2</sub> = CHOH  $\rightarrow$  CH<sub>3</sub>CHO

The hydration first gives vinyl alcohol (ethenol) as an intermediate, which tautomerises to acetaldehyde (ethanal)

- 6. The acidic nature of hydrogens in acetylene cannot be explained by the reaction with
  - A) Sodium metal

- B) Ammonical cuprous chloride solution
- C) Ammonical silver nitrate solution D) HCN

### Answer:D

Solution: Acidic H in acetylene reacts with:

Na metal → sodium acetylide

Ammoniacal  $Cu_2Cl_2 \rightarrow red Cu_2C_2$  precipitate

Ammoniacal  $AgNO_3 \rightarrow white Ag_2C_2$  precipitate

HCN → no reaction of acidic H with HCN; HCN is acidic itself, so no acid-base reaction

- 7. Westron is the solvent obtained by the reaction of chlorine with
  - A) Ethylene

B) Ethyne

C) Ethane

D) Methane

## Answer:B

Solution: Westron is a trade name for tetrachloroethane used as a solvent.

It is produced by chlorination of acetylene (ethyne):

$$HC \equiv CH + 2Cl_2 \rightarrow CHCl_2 - CHCl_2$$

- 8. The number of acidic hydrogen atoms in 1- butyne and 2- butyne respectively are
  - A) 1,0
- B) 0,1
- C) 1,1
- D) 1,2

### Answer:A

Solution:An acidic hydrogen is a hydrogen atom in a molecule that can be easily removed (released) as H<sup>+</sup> ion (proton) when it reacts with a base or metal like sodium

1-Butyne:  $CH = C - CH_2 - CH_3$ 

The hydrogen attached to the terminal carbon of the triple bond (sp-hybridized) is acidic.

Only one hydrogen is attached to that terminal sp-carbon.

Acidic Hydrogens =1

2-Butyne:  $CH_3-C \equiv C-CH_3$ 

In 2-butyne, both triple-bonded carbons are attached to alkyl groups (CH<sub>3</sub>).

No hydrogen is directly attached to the sp-hybridized carbons.

Therefore, no terminal hydrogen  $\rightarrow$  no acidic hydrogen.

Acidic Hydrogens = 0

- 9. Acetylene does not show which of the following reactions?
  - A) Condensation

B) Polymerization

C) Addition reactions

D) Combustion reaction

## Answer:A

Solution: Acetylene undergoes addition, polymerization, and combustion reactions — but not condensation.

- 10. The final product formed when ethyne and acetic acid react is
  - A) Vinyl acetate

- B) Ethyl acetate
- C) Acetylene acetic acid
- D) Ethylidene acetate

#### Answer:A

Solution:  $CH = CH + CH_3COOH \xrightarrow{Hg^{2+}} CH_2 = CHOCOCH$ 

Ethyne (acetylene) reacts with acetic acid in the presence of Hg<sup>2+</sup> catalyst to form vinyl acetate

- 11. When alkyl substituted acetylene udergoes addition with hydrogen in presence of Lindlar's cataylst, the alkene formed is
  - A) A mixture of cis and trans
  - B) Trans product only
  - C) cis product only
  - D) In presence of Lindlar's catalyst, addition does not take place

#### Answer:C

Solutions:Lindlar's catalyst is a poisoned palladium catalyst (palladium deposited on calcium carbonate or barium sulfate and treated with lead salts/quinoline).

This catalyst is specifically designed to perform the partial hydrogenation of alkynes to alkenes and stop the reaction at the alkene stage.

The reaction occurs via a syn addition mechanism, where both hydrogen atoms add to the same face (side) of the triple bond.

Syn addition of hydrogen to an alkyne results exclusively in the formation of a cis (or Z) alkene product.

- 12. The compounds used in plastic industry and polymer industry respectively are
  - A) Vinyl acetate & Vinylcyanide
- B) Vinyl cyanide & Vinyl acetate
- C) Acrylonitrile & Ethylidene
- D) Ethylidene cyanide & Vinyl cyanide

### Answer:B

Solution: Vinyl cyanide (also known as acrylonitrile) is a vital monomer used in the plastic industry for the manufacture of important plastics such as polyacrylonitrile (PAN) and ABS plastic.

Vinyl acetate is an important precursor used in the polymer industry for making a wide range of polymers and copolymers, including polyvinyl acetate (PVAc), ethylene-vinyl acetate (EVA) copolymers, and polyvinyl alcohol (PVOH). These polymers are used in adhesives, paints, and other applications within the broader polymer industry.

13. The monosodium salt of acetylene on treating with methyl chloride forms

(FA & SA- 5 Marks/8 Marks)

- A) CH≡C.COOH
- B)  $CH \equiv C CH_3$
- C)  $CH_3C \equiv CCH_3$
- D)  $CH \equiv C.CH_2CH_3$

Answer:B

Solution:  $HC \equiv CH \xrightarrow{Na} HC \equiv C^{-}Na^{+} + CH_{3} - Cl \rightarrow HC \equiv C - CH_{3} + NaCl$ 

- 14. The number of  $\pi$  bonds in westron
  - A) 0

- B) 1
- C) 2

D) 3

Answer:A

Solution: Westron is the common name for 1,1,2,2-tetrachloroethane (chemical formula: CHCl<sub>2</sub>CHCl<sub>2</sub>). The structure of 1,1,2,2-tetrachloroethane consists of a single bond between the two carbon atoms, and each carbon atom is bonded to two chlorine atoms and one hydrogen atom by single bonds. The molecule only contains single (sigma) bonds.

There are no double or triple bonds present in the structure.

Therefore, the total number of pi bonds in westron is 0.

$$Cl \quad Cl$$
 
$$\mid \quad \mid$$
 
$$H-C \equiv C-H \xrightarrow{Cl_2} H-C-C-H$$
 
$$\mid \quad \mid$$
 
$$Cl \quad Cl$$

- 'x' on ozonolysis gives a dial while 'y' reacts with Baeyer's reagent to give a diol. 15. Then 'x' and 'y' respectively are
  - A) C<sub>2</sub> H<sub>2</sub> & C<sub>6</sub> H<sub>6</sub>

C)  $C_2 H_2 & C_2 H_4$ 

B) C<sub>2</sub> H<sub>4</sub> & C<sub>2</sub> H<sub>2</sub> D) C<sub>2</sub> H<sub>4</sub> & C<sub>6</sub> H<sub>6</sub>

Answer:B

Solution:x = C<sub>2</sub>H<sub>2</sub> (acetylene): ozonolysis of acetylene gives glyoxal (OHC-CHO), a dialdehyde.

y =  $C_2H_4$  (ethene): Baeyer's reagent (cold KMnO<sub>4</sub>) converts ethene to ethylene glycol (a diol)

16. In the following sequence of reactions the product (D) is

$$CH \equiv CH \xrightarrow{HBr} A \xrightarrow{HBr} B \xrightarrow{alcKOH} C \xrightarrow{NaNH_2} D. D is$$

- A) Ethanol
- B) Ethyne C) Ethanal

Br

D) Ethene

Answer:B Solution:

$$CH \equiv CH \xrightarrow{HBr} CH_2 = CH - Br \xrightarrow{HBr} CH_3 - C H - Br \xrightarrow{alcKOH} CH_2 = CH - Br \xrightarrow{|HBr|} CH_3 - C H - Br \xrightarrow{alcKOH} CH_2 = CH - Br \xrightarrow{|HBr|} CH_3 - C H - Br \xrightarrow{alcKOH} CH_2 = CH - Br \xrightarrow{|HBr|} CH_3 - C H - Br \xrightarrow{alcKOH} CH_3 - C H$$

$$\xrightarrow{NaNH_2} CH \equiv CH$$

17. 1-Butyne reacts with cold alkaline KMnO<sub>4</sub> to produce

A) CH<sub>3</sub>CH<sub>2</sub>COOH

B) CH<sub>3</sub>CH<sub>2</sub>CH<sub>2</sub>COOH

C)  $CH_3CH_2COOH + CO_2$ 

D)  $CH_3CH_2COOH + HCOOH$ 

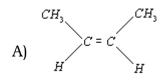
Answer:C

Solution: Cold alkaline KMnO<sub>4</sub> does syn dihydroxylation of double bonds, but 1butyne is an alkyne.

Alkynes with cold, dilute KMnO<sub>4</sub> can give a-diketones, but terminal alkynes get oxidized to carboxylic acids + CO<sub>2</sub>.

1-Butyne:  $CH = C-CH_2-CH_3 \rightarrow terminal alkyne oxidation:$   $CH = C-CH_2-CH_3 \rightarrow CO_2 + CH_3-CH_2-COOH$  (propanoic acid).

Which of the following compound has the lowest dipole moment 18.

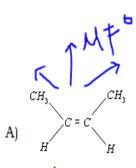


B)  $CH_3C \equiv CCH_3$ 

C)  $CH_3CH_2C \equiv CH$ 

D)  $CH_2 = CH - C \equiv CH$ 

Answer:B



Solution:

19.  $CH = CH \xrightarrow{NaNH_2} A \xrightarrow{CH_3Br} B \cdot B \text{ is}$ 

A)  $CH_2 = CH - CH = CH_2$ 

B)  $HC \equiv C - CH_3$ 

C)  $CH_2 = CH - CH_3$ 

D)  $CH_3 - CH_2 - CH_3$ 

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**Answer:B** 

Solution:

 $CH \equiv CH \xrightarrow{NaNH_2} CH \equiv C^-Na^+ \xrightarrow{CH_3BR} CH \equiv C - CH_3$ 

- Hydration of ethyne to ethanal takes place through the formation of 20.
  - A)  $CH_3CH(OH)_3$
- B)  $CH_2 = CHOH$  C)  $CH_2 = CHO^-$  D)  $CH = C^-$

### Answer:B

Solution: Mechanism:

 $C_2H_2 + H_2O (H^+, Hg^{2+}) \rightarrow \text{vinyl alcohol (CH}_2=CH-OH) (enol) \rightarrow \text{tautomerizes to}$ CH<sub>2</sub>-CHO.

Intermediate = CH<sub>2</sub>=CH-OH.

- 21. A compound on dehydrohalogenation with alcoholic KOH gives alkyne but on dehalogenation with zinc dust gives alkene. The compound is
  - A) C<sub>o</sub>H<sub>e</sub>Br

B) CH<sub>3</sub> CH Br<sub>2</sub>

C) CH<sub>2</sub>Br -CH<sub>2</sub>Br

D) CH Br<sub>2</sub> - CHBr<sub>2</sub>

## Answer:C

- Solution: Alcoholic KOH (strong base) does two eliminations on a vicinal dibromide to give an alkyne (e.g. acetylene from 1,2-dibromoethane). Zinc dust (reductive dehalogenation) converts vicinal dihalides to the alkene (ethene)
- $CaC_2 \xrightarrow{Hydrolysis} A \xrightarrow{HgSO_4 + dil.H_2SO_4} B$ . B is 22.
  - A) Acetylene

B) Acetaldehyde

C) Acetone

D) Acetic acid

Answer:B

$$CaC_2 \xrightarrow{2H_2O} Ca(OH)_2 + C_2H_2(A)$$

Solution:  $C_2 H_2 + H_2 O (HgSO_4, dil H_2 SO_4) \rightarrow CH_3 CHO (B)$ 

 $CaC_2 + H_2O \rightarrow A + B \xrightarrow{1mole Na}$ 23.

$$C \xrightarrow{C_2H_5I} D$$
. D is

A) 1-butene

B) Propene

C) 1-pentene

D) 1 - Butyne

Answer:D

$$CaC_2 + 2H_{20} \longrightarrow (aGH)_2 + C_2H_2$$

$$A = C_2H_2.$$

$$B = Ca(OH)_2.$$

$$A + 1 \text{ mole Na.}$$

$$C_2H_2 + N_C \longrightarrow N_0^+ C_2H^- + \frac{1}{2}H_2$$

$$C = N_0 C = CH \text{ (Monosodium accelylida).}$$

$$C + C_2H_5I \longrightarrow D.$$

$$N_0 C = CH + C_2H_5I \longrightarrow CHEC - C_2H_5 + N_0I$$

$$D = CH = C - C_2H_5$$

24. 
$$CH_2 - CH_2 \xrightarrow{Alc. KOH} A \xrightarrow{One mole \ HCl} B'B' is$$

$$| \qquad | \qquad \qquad Cl \qquad Cl$$

- A) Ethyl chloride
- C) Vinyl chloride

- B) 1,2 dichloro ethene
- D) Ethylidine chloride

### Answer:C

Solutio:

Solution:

25. 
$$CH \equiv CH \xrightarrow{HCl} A \xrightarrow{Polymersiation} B$$
 The polymer 'B' is A) orlon B) PVC C) nylon D) teflon

#### Answer:B

26. 
$$H-C \equiv C-H+NaNH_2 \rightarrow A \xrightarrow{2 \text{ mole} \atop CH_3Cl} B \text{ then `B` is}$$

- A) 1-Butyne
- B) 2-Butyne
- C) 2-Pentyne
- D) Propyne

## Answer:B

Solution:

27. When 2-pentyne is treated with dilute H<sub>2</sub>SO<sub>4</sub> and HgSO<sub>4</sub> the product formed is A) 1-pentanol B) 2-pentanol C) 2-pentanone D) 3-pentanone

Answer:C

Solution:

- 28. The cyclic polymerisation of methyl acetylene produces
  - A) benzene

B) O-xylene

C) 1,3,5 - Trimethyl benzene

D) 1,3,5 - Tri methyl cyclo hexane

Answer:C

Solution:

- 29. The compounds 1-butyne and 2-butyne can be distinguished by using
  - A) Bromine water

B)  $KMnO_4$  solution

C) Tollen's reagent

D) Chlorine gas

### Answer:C

Solution:

- 1-butyne: terminal alkyne (HC≡C-CH<sub>2</sub>-CH<sub>3</sub>)
- 2-butyne: internal alkyne ( $CH_3-C = C-\bar{C}H_3$ )

Terminal alkynes react with Tollen's reagent (ammoniacal AgNO<sub>2</sub>) to form white precipitate of silver acetylide; internal alkynes do not.

Bromine water and KMnO<sub>4</sub> react with both (unsaturation tests). Cl<sub>2</sub> gas also reacts with both.

- 30. Which of the following orders regarding acidic strength is correct

  - A)  $CH_3COOH > CH_3CH_3OH > CH \equiv CH$  B)  $CH_3COOH > CH \equiv CH > CH_3CH_3OH$

  - C)  $CH = CH > CH_3COOH > CH_3CH_2OH$  D)  $CH = CH > CH_3CH_2OH > CH_3COOH$

### Answer:A

Solution: Acetic acid (pKa ~ 4.8) is far stronger than ethanol (pKa ~ 16), which is stronger than acetylene (pKa ~ 25)

An unknown compound 'A' has a molecular formula of  $C_4H_6$  when 'A' is treated 31. with an excess of  $Br_2$  a new substance 'B' with formula  $C_4H_6Br_2$  is formed . A forms a white precipitate with ammonical silver nitrate solution 'A' may be A) Butyne-1 B) Butyne-2 C) Butene-2 D) Butene-1

## Answer:A

Solution: A gives a white precipitate with ammoniacal  $AgNO_3 \rightarrow characteristic$  of a terminal alkyne (forms a silver acetylide). So A is 1-butyne (C<sub>4</sub>H<sub>6</sub>)

- 32. The reduction of 4-octyne with  $H_2$  in the presence of  $Pd/BaSO_4$  quinoline gives
  - A) trans -4 octene
  - B) cis 4 octene
  - C) a mixture of cis and trans-4octene
  - D) A completely reduced product  $C_8H_{18}$

Answer:B

Solution:

- $CH = CH + 2CH_3COOH \rightarrow A \xrightarrow{\Delta} B + C$ . B can be obtained from (X) by hydration and 33. (X)
  - C can be obtained from (Y) by heating with  $P_2O_5$ . Hence A is
  - A)  $CH_2 = CH O COCH_3$  B)  $CH_3 CH(O COCH_3)_2$

C) CH<sub>3</sub>CH<sub>2</sub>COOCH<sub>3</sub>

D)  $CH_3COOCH_2CH_3$ 

#### Answer:A

Solution: 
$$CH = CH + 2CH_3COOH \xrightarrow{\Delta} CH_3 - CH(O - COCH_3)_2$$
  
 $A = CH_3 - CH(O - COCH_3)_2$ 

## JEE ADVANCED LEVEL QUESTIONS

## Multi Correct Answer Type:

- 1. Which of the following statements about alkynes is/are true?
  - A) They contain at least one triple bond between carbon atoms.
  - B) They undergo addition reactions readily.
  - C) They are more reactive than alkenes.
  - D) They can be converted into alkenes by reduction.

## Answer:A,B,D

Solution: A) True — definition of alkynes.

- B) True alkynes undergo electrophilic addition (with Br<sub>2</sub>, HX, etc.) and catalytic hydrogenation.
- C) False alkynes are generally less reactive than alkenes toward electrophilic addition because the sp-hybridized carbons hold p-electrons more tightly.
- D)True using Lindlar's catalyst (cis-alkene) or Na in liquid NH3 (transalkene).
- 2. Which of the following statements about the bond length and strength in alkynes are true?
  - A) The triple bond in alkynes consists of one sigma bond and two pi bonds.

- B) The bond length of the triple bond is shorter than that of the double bond in alkenes.
- C) The triple bond in alkynes is weaker than the double bond in alkenes.
- D) The triple bond in alkynes allows for rotation around its axis.

### Answer:A,B

Solution:A) The triple bond in alkynes consists of one sigma bond and two pi bonds.

→True.

B)True —  $C \equiv C$  bond length ~120 pm, C=C ~134 pm.

- C) False C=C bond is stronger (higher bond energy  $\sim 839 \text{ kJ/mol}$ ) than C=C ( $\sim 611 \text{ kJ/mol}$ ).
- D) False rotation is restricted around triple bond just like double bonds (pi bonds prevent free rotation).

## **Reason And Assertion Type:**

- A) Both (A) and (R) are true and (R) is the correct explanation of(A)
- B) Both (A) and (R) are ture and (R) is not the correct explanation of (A)
- C) (A) is true but (R) is false
- D) (A) is false but (R) is true
- 3. **Assertion**: Acetylene is formed when ethylene chloride or ethylidene
  - chloride is heated with alcoholic KOH
  - **Reason**: Both gem dihalides and vicinal dihalides on dehydrohalogenation

form alkyne

#### Answer:A

Solution: Assertion (A) is true: heating ethylene chloride (vicinal dihalide) or ethylidene chloride (geminal dihalide) with alcoholic KOH leads to successive dehydrohalogenations giving acetylene (terminal alkyne).

Reason (R) is true: both vicinal and geminal dihalides can undergo successive eliminations to form an alkyne.

R correctly explains A

4. **Assertion**: Heavy metal acetylides can be used to purify alkynes

**Reason**: Terminal alkynes form acetylides which are soluble in acids

#### Answer:C

Solution:Assertion (A) is true: heavy-metal acetylides (Ag, Cu salts) are used in detection/purification of terminal alkynes — the terminal alkyne is precipitated as a metal-acetylide and can be reconverted back to the alkyne. Reason (R) is false as stated: terminal alkynes form metal-acetylides which are insoluble salts (not "soluble in acids"). In practice the metal salt is decomposed (protonated) by acid to regenerate the alkyne

5. **Assertion**: Disubstituted acetylene on partial hydrogenation may give trans

isomei

**Reason**: Lindlar's catalyst is used for Partial hydrogenation

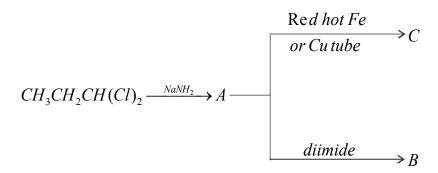
### Answer:B

Solution: Assertion (A) is true: a disubstituted alkyne on partial hydrogenation may give a trans alkene (for example, using Na/NH<sub>3</sub> gives the trans (E) alkene).

Chemistry: Alkynes

Reason (R) is true in itself: Lindlar's catalyst is used for partial hydrogenation — but Lindlar's gives the cis (Z) alkene, not the trans. Therefore R does not correctly explain A

## Comprehension Type:



6. A is

A)Propyne

B) Propene

C)Propanal

D)Propanone

#### Answer:A

Solution: The reactant  $CH_3CH_2CH(Cl)_2$ , 1,1-dichloropropane, undergoes dehydrohalogenation with a strong base like NaNH, to form an alkyne. Two molecules of HCl are removed to form  $CH_3C \equiv CH$ , which is propyne.

7. B is

A) 
$$CH_3CH = CH_3$$
 B)  $CH_3CH_2CH_3$  C)  $CH_3COCH_3$ 

C) 
$$CH_3COCH_3$$

D) 
$$CH_3CH_3$$

#### Answer:B

Solution: Product A (propyne) undergoes catalytic hydrogenation using diimide N<sub>2</sub>H<sub>2</sub>, which is a selective reducing agent that reduces alkynes to alkanes  $(CH_3CH_2CH_3, \text{ propane})$  under these conditions.

8. C is

A)Mesitylene

B)Benzene

C)Cyclooctatetraene

D)Benzaldehyde

#### Answer:A

Solution: When propyne is passed through a red-hot iron or copper tube, it undergoes cyclic polymerization (trimerization) to form 1,3,5trimethylbenzene, commonly known as mesitylene.

## **Integer Type:**

9. Determine the total number of sigma and pi bonds in the molecule of ethyne.

#### Answer:5

Solution:Structure:  $H - C \equiv C - H$ 

Sigma bonds:2 × C-H bonds

1 × C-C (sigma part of triple bond)

Total sigma bonds = 3

Pi bonds:

Triple bond has 2 pi bonds

Total pi bonds = 2

Total sigma + pi bonds = 3 + 2 = 5

10. What is the number of sp hybridized carbon atoms in 2-butyne?

Answer:2

Solution:Structure: CH<sub>3</sub>-C<sub>≡</sub>C-CH<sub>3</sub>

Each triple-bonded carbon is sp hybridized.

There are 2 such carbons (the ones in  $C \equiv C$ ).

The  $CH_3$  carbons are  $sp^3$  hybridized.

So sp hybridized carbon atoms = 2

## **Matrix Matching Type:**

## 11. Column I

A) 
$$R - CH = CH - R \xrightarrow{cold \, dil} \xrightarrow{KMnO_4}$$

cis

B) 
$$R-CH=CH-R \xrightarrow{Br_2} CCl_4$$

trans

C) 
$$RC \equiv CR \xrightarrow{H_2} Ri_2B$$

D) 
$$RC \equiv CR \xrightarrow{Li}_{Liq.NH_3}$$

#### Column II

P) Meso

- R) trans-alkene
- S) controlled reduction
- T) Oxidation

## ${\bf Answer: A-T, B-P, C-Q, D-R}$

Solution:A)  $R - CH = CH - R \xrightarrow{cold dil} KMnO_4$ 

 $\frac{dl}{da}$  T) Oxidation

B)  $R - CH = CH - R \xrightarrow{Br_2} CCL$ 

P) Meso

cis

C)  $RC \equiv CR \xrightarrow{H_2} Ri_{1}B$ 

Q) cis-alkene

D)  $RC \equiv CR \xrightarrow{Li}_{Liq.NH_3}$ 

R) trans-alkene

## LEARNERS TASK

## **CONCEPTUAL UNDERSTANDING QUESTIONS (CUQ'S)**

- 1. The IUPAC name of the compound having the formula CH = C CH = CH,
  - A) Butene -2- yne

B) But-2-yne-3-ene

C) 3- butane 1- ene

D) But-1-ene - 3- yne

#### Answer:D

Solution:IUPAC name of the compound (structure given as a molecule containing both a C=C and a C=C) — best systematic name is But-1-ene-3-yne (double at C-1 and triple at C-3), written as but-1-ene-3-yne.

- 2. Alkynes exhibit.
  - A) Chain isomerism

B)Position isomerism

C)Functional isomerism

D) All the above

#### Answer:D

Solution:Alkynes show chain, position and functional isomerism (they have all those types)

- 3. Alkynes exhibit functional isomerism with
  - A) Alkanes
- B) Alkenes
- C) alkadienes
- D) Alcohols

#### Answer:C

Solution:

Functional isomers: alkynes  $\rightarrow$  alkadienes (same molecular formula  $C_nH_{2n-2}$ )

- 4. The isomer of propyne
  - A) Allene
- B) Propene
- C) Cyclo propane D) Propane

#### Answer:A

Solution: The isomer of propyne (CH<sub>3</sub>-C=CH) is allene (propadiene, CH<sub>2</sub>=C=CH<sub>2</sub>)

- 5. Bond angle between C C in alkyne
  - A) 109° 28′
- B) 120°
- C) 180°
- D) 60°

#### Answer:C

Solution:Bond angle about the C–C axis in an alkyne (sp-hybridised carbon) is 180° (linear)

- 6. The molecule having linear structure is
  - A) Methane
- B) Ethylene
- C) Acetylene
- D) Water

#### Answer:C

Solution: Methane  $\rightarrow$  tetrahedral

Ethylene  $\rightarrow$  trigonal planar

Acetylene  $\rightarrow$  linear

Water →bent

- 7. The C C bond length is shortest in
  - A)  $C_2H_6$
- B)  $C_{2}H_{2}$
- C)  $C_6H_6$
- D)  $C_2H_4$

## Answer:B

Solution:A)  $C_2H_6 \rightarrow \text{Ethane} \rightarrow \text{C-C single bond} \rightarrow ~1.54\text{A}^0$ 

- B)  $C_2H_2 \rightarrow \text{Acetylene} \rightarrow \text{C=C triple bond} \rightarrow \sim 1.20 \text{ Å}$
- C)  $C_6H_6 \rightarrow$  Benzene  $\rightarrow$  C–C bond in aromatic ring  $\rightarrow$  ~1.39 Å (intermediate between single and double)
- D)  $C_2H_4$  Ethene  $\rightarrow$  C=C double bond  $\rightarrow$  ~1.34 Å
- 8. The hydrolysis of  $Mg_2C_3$  produces
  - A) acetylene
- B) propyne
- C) butyne
- D) ethylene

## Answer:B

Solution:  $Mg_2 C_3 + 4H_2 O \rightarrow 2Mg(OH)_2 + CH_3 - C \equiv CH \text{ (propyne)}$ .

- 9. The number of possible alkynes with molecular formula  $C_5H_8$  is
  - A) 3
- B) 4

C) 5

D) 6

## Answer:A

$$CH_{3}-CH_{2}-CH_{2}-C\equiv CH \rightarrow Pent-1-yne$$
 
$$CH_{3}-CH_{2}-C\equiv C-CH_{3} \rightarrow Pent-2-yne$$
 
$$CH_{3}$$

Solution:

$$CH_3 - CH - C \equiv CH \rightarrow 2 - metylBut - 1 - yne$$

- 10. Iodoform on heating with silver powder gives
  - A) CH<sub>4</sub>
- B)  $C_2H_2$
- C)  $C_2H_4$
- D) C<sub>6</sub>H<sub>6</sub>

#### Answer:B

Solution:  $2CHI_3 + 6Ag \rightarrow C_2 H_2(acetylene) + 6AgI$ 

- 11. Acetylene is stored and transported in
  - A) Ethanol solution

B) Methanol solution

C) Ethanal solution

D) Propanone solution

#### Answer:D

Solution: Acetylene is unstable under pressure and can decompose explosively. For safe storage and transportation, it is dissolved in acetone (propanone) in cylinders containing a porous material (like diatomaceous earth) to prevent high pressure buildup.

- 12. PVC is the polymer of the following
  - A) Ethyl chloride

B) Vinyl chloride

C) Allyl chloride

D) Ethynyl chloride

## Answer:B

Solution:PVC (Polyvinyl chloride) is formed by polymerization of vinyl chloride

$\overline{}$	
(9th	Class

Chemistry: Alkynes

Hydrocarbon which gives oxyacetylene flame 13.

- A) ethane
- B) ethene
- C) ethyne

D) ethanal

Answer:C

Solution: Ethyne is the hydrocarbon that produces the oxy-acetylene flame used for welding and cutting metal. This is because the high-temperature flame from burning ethyne (acetylene) in oxygen provides the heat needed to melt metals. Ethane and ethene are not suitable for this purpose

14. The gas obtained when ethylene chloride reacts with alcoholic potash and sodamide is

- A)  $C_{2}H_{4}$
- B)  $C_2H_6$
- C)  $C_2H_2$
- D)  $C_2H_5C1$

Answer:C

Solution: Ethylene chloride with alcoholic KOH / sodamide gives dehydrohalogenation → acetylene

15. Iodoform on heating with silver powder gives

- A) CH<sub>4</sub>
- B)  $C_2H_2$  C)  $C_2H_4$
- D)  $C_6H_6$

Answer:B

Solution:  $2CHI_3 + 6Ag \rightarrow C_2 H_2(acetylene) + 6AgI$ 

## JEE MAINS LEVEL QUESTIONS

1. Pure acetylene has sweet smell, where as impure gives garlic odour due to presence of

- A) NH<sub>2</sub>
- B) PH<sub>2</sub>
- C) SbH<sub>2</sub>
- D) HC1

Answer:B

Solution:Pure acetylene has a sweet smell.

Commercial or impure acetylene (prepared from calcium carbide and water) often contains traces of phosphine (PH<sub>2</sub>) and arsine (AsH<sub>2</sub>).

These impurities are responsible for the garlic-like odor of impure acetylene.

2. Which of the following possess acidic hydrogen

- A)  $C_2H_6$
- B)  $C_2H_4$  C)  $C_2H_2$  D)  $CH_4$

Answer:C

Solution: Acidic hydrogen is present in terminal alkynes (C<sub>2</sub>H<sub>2</sub>) due to sp-hybridized carbon.

3. The reagent used for obtaining trans alkene from alkyl substituted acetylene with hydrogen is (FA & SA- 3 Marks/4 Marks)

A) Na in liq. NH<sub>2</sub>

B) LiAlH<sub>4</sub>

C)  $Z_n + HCl$ 

D) H, in presence of Ni

Answer:A

Solution: Na in liq.  $\mathrm{NH_3}$  — Dissolving-metal hydrogenation (Na / liquid  $\mathrm{NH_3}$  ) of an alkyl-substituted alkyne gives the trans alkene

EdOS - Educational Operating System

The stronger base is

(FA & SA- 5 Marks/8 Marks)

- A)  $CH_3CH_2^-$
- B)  $CH_2 = CH^-$  C)  $CH \equiv C^-$
- D) *Cl*<sup>-</sup>

### Answer:A

Solution: The strength of a base is inversely related to the stability of its conjugate acid. A stronger base will have a weaker, less stable conjugate acid.

The order of stability of the anions (from least stable to most stable) is:

$$CH_3 CH_2^- < CH_2 = CH^- < CH \equiv C^- < Cl^-$$

Since the strongest base corresponds to the least stable negative charge, the strongest base among the options is CH<sub>3</sub> CH<sub>2</sub>.

- Gem dihalides on treatment with alcoholicKOH give 5.
  - A) Alkyne
- B) Alkene
- C) Alkane
- D) Cyclo alkanes

## Answer:A

Solution: Gem-dihalides (both halogen atoms on the same carbon) on treatment with alcoholic KOH undergo double dehydrohalogenation, forming an alkyne

- 6.  $X + 2Zn \xrightarrow{Alcohol} H - C \equiv C - H$  here 'X' is
  - A) 1, 1-Dibromoethane
- B) 1, 2-Dibromoethane

C) Di bromo ethane

D) 1, 1, 2, 2-Tetra - bromoethane

Answer:D

Deha log enation: 1moleZnreactwith2molesBr

$$Br$$
  $Br$   
 $|$   $|$   
 $H_2 C - CH_2 \xrightarrow{Zn} CH_2 = CH_2 + ZnBr_2$ 

2molesZnreactwith4molesBr

Solution:

- 7. 2- Butyne when treated with lithium in presence of liquid ammonia gives
  - A) cis-2-butene

B) trans-2-butene

C) n-butane

D) I-butyne

Answer:B

Solution: The reaction of 2-butyne (a non-terminal alkyne) with lithium in the presence of liquid ammonia is a dissolving metal reduction (also known as a modified Birch reduction).

This reaction is stereoselective and proceeds via an anti-addition mechanism due to the stability of the intermediate radical anion and vinylic anion species, 9th Class

Chemistry: Alkynes

where the bulky groups prefer to be on opposite sides of the forming double bond.

The result of this anti-addition across the triple bond is the formation of a trans-alkene, which in this case is trans-2-butene

8. Which one of the following possess the minimum boiling point (AIEEE 200D)

(FA & SA- 2 Marks)

A) 1-Pentyne

B) 1-Butyne

C) n-Butane

D) Isobutane

Answer:D

Solution: Boiling point increases with molar mass and decreases with branching. n-Butane BP ~0°C, Isobutane BP ~ -12°C (lowest), 1-Butyne BP ~ 8°C, 1-Pentyne BP ~ 40°C.

- 9. 1- pentyne and 2-pentyne can be distinguished by
  - A) Silver mirror test

B) Iodoform test

C) Addition of  $H_2$ 

D) Baevers test

Answer:A

Solution: Terminal alkyne (1-pentyne) gives a precipitate with ammoniacal Ag+ (forms Ag-acetylide), whereas internal alkyne (2-pentyne) does not

- 10. Acetylene on reaction with silvernitrate shows
  - A) Oxidizing property

B) Reducing property

C) Basic nature

D) Acidic nature

Answer:D

Solution: Acetylene (C<sub>2</sub>H<sub>2</sub>) reacts with ammoniacal silver nitrate solution (Tollens' reagent) to form a white precipitate of silver acetylide (C<sub>2</sub>Ag<sub>2</sub>). This reaction occurs because the hydrogen atoms attached to the sp-hybridized carbon atoms in acetylene are weakly acidic and can be replaced by metal ions (in this case, silver). The ability to donate a hydrogen ion (H<sup>+</sup>) is the defining characteristic of an acid, thus demonstrating the acidic nature of acetylene.

11. The compound used in Hawker's lamp and in light houses for illumination purpose is

A) methane

- B) ethane
- C) ethylene
- D) acetylene

Answer:D

Solution: Acetylene (from calcium carbide) was widely used in Hawker's lamps and early lighthouse illumination.

$$CaC_2 + H_2O \rightarrow CH \equiv CH + Ca(OH)_2$$
  
 $CH \equiv CH \xrightarrow{O_2} Light$ 

12. Cold and dil.Alk. *KMnO*<sub>4</sub> will oxidise acytylene to

A) Ethylene glycol B) Ethyl alcohol C) Oxalic acid

D) Acetic acid

Answer:D

Solution: Cold, dilute alkaline KMnO4 oxidises C2H2 to oxalic acid (HOOC-COOH).

- 13.  $X + 2KOH \xrightarrow{Alcohol} H C \equiv C H$  here "X' is
  - A) 1, 1–Dibromoethane
- B) 1, 2-Dibromoethane

C) Both (A) and (B)

D) 1, 1, 2, 2 - Tetrabromoethane

### Answer:C

Solution:From vicinal dihalide (1,2-dibromoethane) or geminal dihalide (1,1-dibromoethane), alcoholic KOH causes double dehydrohalogenation to form acetylene.

So both (A) and (B) can give acetylene.

14. Westrosol is

A) 
$$Cl_{2}C = CHC1$$
 B)  $CHCl_{2} - CHCl_{2}$  C)  $CH_{2}C1 - CH_{2}C1$  D)  $CH_{3} - CCl_{3}$ 

### Answer:A

Solution: Westrosol is trichloroethene (structure: CCl<sub>2</sub>=CHCl)

- 15. Acetylene gives white precipitate with ammonical silver nitrate but ethylene cannot give because
  - A) Acetylene possess sp<sup>2</sup> carbon
  - B) Acetylene posses acidic hydrogen
  - C) Acetylene possess low electronegative carbon
  - D) Acetylene posess  $-C \equiv C$  triple bond

## Answer:B

Solution: Acetylene ( $C_2H_2$ ) has acidic hydrogens attached to sp-hybridized carbons, which can be removed by strong bases or metal ions like  $Ag^+$  (from ammoniacal  $AgNO_3$ ).

This forms a white precipitate of silver acetylide (Ag<sub>2</sub>C<sub>2</sub>).

Ethylene (C<sub>2</sub>H<sub>4</sub>) lacks such acidic hydrogens, so it does not react with ammoniacal AgNO<sub>3</sub>

16. Which of these will not react with acetylene?

(AIEEE 200B)

A) NaOH

B) Na

C) Ammonical AgNO<sub>3</sub>

D) HCl

## Answer:A

Solution: Acetylene is a weak acid and NaOH is a strong base, so the reaction between them is not favorable.

## JEE ADVANCED LEVEL QUESTIONS

## **Multicorrect Answer Type:**

- 1. Which of the following are false
  - A) Acetylene is more reactive than ethylene to an electrophilic attack
  - B) Acetylene is less reactive than ethylene towards electrophilic attack
  - C) Acetylene may show more reactivity or less reactivity towards electrophilic reagent.
  - D) Acetylene and ethylene show identical reactivities towards an electrophilic attack

## Answer:A,C,D

Solution:A)False — acetylene is less reactive than ethylene toward electrophilic

addition because the sp-hybridized carbons hold p-electrons more tightly, and the intermediate vinylic carbocation is less stable.

- B) Acetylene is less reactive than ethylene towards electrophilic attack True so this is not false.
- C) False in general, it's less reactive; "may show more" is not correct in normal electrophilic addition.
- D) False they do not have identical reactivity.
- 2. Which of the following statements regarding the physical properties of alkynes are true?
  - A) Alkynes have higher boiling points than alkanes of similar molecular weight.
  - B) Alkynes have lower boiling points than alkenes of similar molecular weight.
  - C) Alkynes are less dense than water.
  - D) Alkynes are soluble in nonpolar solvents like hexane.

## Answer:A,C,D

Solution:A) True — alkynes are more polarizable (linear, triple bond) and have stronger intermolecular forces than alkanes.

- B) False alkynes generally have slightly higher boiling points than alkenes due to higher linearity and polarizability.
- C)True like most hydrocarbons, they float on water (density < 1 g/mL).
- D) True they are nonpolar or weakly polar, so soluble in organic solvents.

## **Assertion and Reason Type:**

- A) Both (A) and (R) are true and (R) is the correct explanation of(A)
- B) Both (A) and (R) are ture and (R) is not the correct explanation of (A)
- C) (A) is true but (R) is false
- D) (A) is false but (R) is true
- 3. **Assertion** : Alkynes are more reactive than alkene towards catalytic hydrogenation
- Reason : Alkynes are less reactive towards electrophilic reaction than alkenes

#### Answer:B

#### Solution:

Assertion true: Alkynes are easier to hydrogenate (consume H2) under catalytic hydrogenation conditions than alkenes (they are more readily reduced to alkenes/alkanes).

Reason true but irrelevant: Alkynes being less reactive than alkenes toward electrophilic addition is correct, but that fact does not explain their greater reactivity in catalytic (heterogeneous) hydrogenation — catalytic hydrogenation proceeds by surface adsorption and electron transfer, a different mechanism

4. **Assertion**: Addition of HBr to  $HC = CCH_2CH = CH_2$  give  $HC = CCH_2CHBr - CH_3$  and not  $H_2C = CBrCH_2CH = CH_3$ 

Reason : A triple bond is less reactive than a double bond towards electrophilic

#### Answer:A

Solution: Assertion true: In a molecule containing both a C = C and a C = C, electrophilic addition (HBr) attacks the double bond preferentially, giving the product with HBr added to the alkene unit rather than adding to the triple bond.

Reason true and explanatory: The triple bond is generally less reactive than the double bond toward simple electrophilic addition (vinyl cations are less stabilized), so the double bond reacts first — R correctly explains A

5. : Reaction of but-2-yne by Na/liqNH<sub>3</sub> gives trans But-2-ene Assertion

Reason : It is syn addition

### Answer:C

Solution: Assertion true: But-2-yne reduced with Na / liq NH3 (dissolving-metal reduction) gives trans-2-butene.

Reason false: The reaction proceeds by an anti-addition (stepwise electron/ proton transfers producing radical anion intermediates), not by syn addition. Anti-addition is why the trans alkene is formed

## Comprehension Type:

Terminal alkynes have acidic hydrogen/s. Sodium salt of terminal alkynes behave as nucleophile as well as strong base. For primary alkyl halides it behave as nucleophile. Thus primary alkyl halides give SN reaction with its salt. Alkynes undergo electrophilic as well as nucleophilic addition reactions. They also undergo hydroboration, oxidation and ozonolysis.

When 2-pentyne is treated with dil.  $H_{\gamma}SO_{4}$  and  $HgSO_{4}$ , the product formed is 6. A) 1-pentanol B) 2-pentanol C) 2-pentanone D) 3-pentanone. Answer:C

Solution:

7.  $CH_3C \equiv CH + HOX \longrightarrow Y.Y$  is

A)  $CH_3COCH_2X$  B)  $CH_3COCH_3$  C)  $CH_3COCOOH$  D)  $CH_3CHXCHO$ 

#### Answer:A

Solution: HOX adds Markovnikov across the terminal C≡C. The enol/halovinyl intermediate formed is converted by tautomerization to the  $\alpha$ -halo ketone

$$CH_3COCH_2X$$

## **Integer Type:**

Determine the number of pi bonds present in 1-butyne.

### Answer:2

Solution:1-butyne.--->  $CH_3$ - $C \equiv C$ -H

A triple bond contains:1 sigma bond,2 pi bonds Therefore, in 1-butyne: Number of pi bonds = 2

9. Determine the total number of sigma and pi bonds in 1,3-butadiyne.

### Answer:9

Solution: 
$$H - C \equiv C - C \equiv C - H$$

## **Matrix Matching Type:**

#### 10. Compound

- A) Acetylene
- B) Ethylene
- C) Benzene
- D) 2-Butene

## **Ozonolysis products**

- 1) HCHO & CH<sub>3</sub>CHO
- 2) *CH*<sub>3</sub>*CHO*
- 3) One mole of (CHO)<sub>2</sub>
- 4) 3 moles of (CHO)<sub>2</sub>
- 5) CH<sub>2</sub>O

## Answer: A-3, B-5, C-4, D-2

Solution:

A) Acetylene

$$CH \equiv CH + O_3 \rightarrow Ozonide \xrightarrow{Hydrolysis} CHO - CHO$$

3) One mole of (CHO)<sub>2</sub>

B) Ethylene

$$CH_2 = CH_2 + O_3 \rightarrow Ozonide \xrightarrow{Hydrolysis/Reduction} 2HCHO$$

5) CH<sub>2</sub>O

C) Benzene

$$C_6H_6 + 3O_3 \rightarrow Ozonide \xrightarrow{Hydrolysis/Reduction} 3CHO-CHO$$

4) 3 moles of (CHO)<sub>2</sub>

D) 2-Butene

$$CH_3CH = CHCH_3 + O_3 \rightarrow Ozonide \xrightarrow{Hydrolysis/Reduction} 2CH_3CHO$$
 2)  $CH_3CHO$ 

# **KEY**

					TEACHING	TASK				
				JEE MAINS LEVEL QUESTIONS						
	1	2	3	4	5	6	7	8	9	10
D		В	В	В	В	D	В	Α	Α	Α
	11	12	13	14	15	16	17	18	19	20
С		В	В	Α	В	В	С	В	В	В
	21	22	23		25	26				30
С		В	D	С	В	В	С	С	С	Α
	31	32	33							
A		В	Α							
				JEE ADVANCED LEVEL QUESTIONS						
	1	2	3	4	5	6	7	8		10
A,B,D		A,B	Α	С	В	Α	В	Α	5	2
	11									
A-T,B-	P,C-	Q,D-R								
					LEARNERS					
				CONCEPTUAL UNDERSTANDING QUESTIOI						
	1	2			5				9	10
D		D	С	Α	С	С	В	В	Α	В
	11	12				IRA				
D		В	С	С	В					
				JEE MAINS LEVEL QUESTIONS						
	1	2	3	4	5	6		8	9	10
В		С	Α	Α	Α	D	В	D	Α	D
	11									
D		D	С	Α	В	Α				
				JEE ADVANCED LEVEL QUESTIONS						
	1	2	_		5					
A,C,D		A,C,D	В	Α	С	С	Α	2	9	
	10									
A−3, B	-5, (	C-4, D-2								