

**Sample Paper (2023-24)**

**Class 11<sup>th</sup> (Sr. Secondary)**

**Code: CHE-856**

**Roll No.**

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**Chemistry**  
**(English Medium)**  
**Academic**

**[Time allowed: 3 hours]**

**[Maximum Marks: 70]**

**General Instructions:-**

Read the following instructions carefully and strictly follow them.

- (i) This question paper is divided into five sections A, B, C, D and E.
- (ii) This question paper contains 35 questions. All questions are compulsory.
- (iii) In Section A – Question No. 1 to 18 are multiple choice (MCQ) type questions carrying 1 mark each.
- (iv) In Section B – Question No. 19 to 25 are very short answer (VSA) type questions carrying 2 marks each.
- (v) In Section C – Question No. 26 to 30 are short answer (SA) type questions carrying 3 marks each.
- (vi) In Section D – Question No. 31 and 32 are case based questions carrying 4 marks each.
- (vii) In Section E – Question No. 33 to 35 are long answer (LA) type questions carrying 5 marks each.
- (viii) There is no overall choice. However an internal choice has been provided in two questions in Section B, two questions in Section C, two questions in Section D and two questions in Section E.
- (ix) Use of calculators is not allowed.

## SECTION-A

Question No. 1 to 18 are multiple choice (MCQ) type questions, carrying 1 mark each.

(18x1=18)

- How many significant figures are present in 0.0025  
(a) 2                      (b) 4                      (c) 1                      (d) 3
- How many unpaired electrons are present in chromium  
(a) 5 unpaired  $e^-$  s      (b) 6 unpaired  $e^-$  s  
(c) 3 unpaired  $e^-$  s      (d) 1 unpaired  $e^-$
- Among halogens the correct order of amount of energy released during the gain of electron is.  
(a)  $F > Cl > Br > I$                       (b)  $F < Cl < Br < I$   
(c)  $F < Cl > Br > I$                       (d)  $F < Cl < Br > I$
- Which among the following are diamagnetic?  
(a)  $N_2^+$                       (b)  $N_2^{2-}$                       (c)  $O_2$                       (d)  $O_2^{2-}$
- What is the molar mass of  $H_2O$  in gm/mol  
(a) 44                      (b) 18                      (c) 17                      (d) 60
- For the process to occur under adiabatic conduction the correct condition is  
(a)  $\Delta T = 0$       (b)  $\Delta P = 0$                       (c)  $q = 0$                       (d)  $w = 0$
- Find the oxidation number of iron in  $K_4[Fe(CN)_6]$   
(a) +2                      (b) +3                      (c) +4                      (d) +1
- An electrophilic reagent is  
(a) Electron deficient species                      (b) Electron rich species  
(c) Negatively charged species                      (d) Lewis base
- $\Delta U^\circ$  of combustion of methane is  $-X$  KJ/mol. The value of  $\Delta H$  is  
(a)  $= \Delta U^\circ$                       (b)  $> \Delta U^\circ$                       (c)  $< \Delta U^\circ$                       (d) = 0
- Arrange the following in the decreasing order of their boiling point.  
n-butane, n-pentane, 2-methyl butane  
(a) n-pentane > 2-methyl butane > n-butane

- (b) n-butane > 2-methyl butane > n-pentane  
 (c) 2-methyl butane > n-butane > n-pentane  
 (d) 2-methyl butane > n-pentane > n-butane
11. Magnetic Quantum number for the valence electron of Potassium is  
 (a) 0 (b) 1 (c) 2 (d) 7
12. The shape of carbocation is  
 (a) planar (b) linear (c) pyramidal (d) Tetrahedral
13. How many molecules of water are present in 0.01 mole of it.  
 (a)  $6.022 \times 10^{23}$  (b)  $6.022 \times 10^{21}$   
 (c)  $6.022 \times 10^{22}$  (d)  $6.022 \times 10^{24}$
14. How many hydrogen bonded water molecules are associated with  $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$   
 (a) 1 (b) 2 (c) 3 (d) 4

For questions 15 to 18, two statements are given one labelled as Assertion(A) and the other labelled as Reason(R). Select answer to these questions from the codes (a), (b), (c), (d) as given below.

- (a) Both Assertion(A) and Reason(R) are true and Reason(R) is the correct explanation of the Assertion(A)  
 (b) Both Assertion(A) and Reason(R) are true but Reason is not the correct explanation of the Assertion  
 (c) Assertion(A) is true but Reason(R) is false.  
 (d) Assertion(A) is false but Reason(R) is true.
15. Assertion : A substance which gets reduced can act as the reducing agent.  
 Reason : An oxidizing agent itself get reduced.
16. Assertion : Graphite is an element  
 Reason : Element is the pure form of a substance containing same kind of atoms.

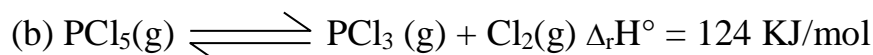
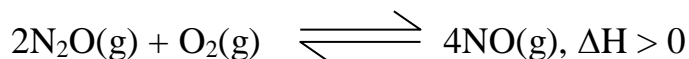
- 17.Assertion : Acetylene is more acidic than ethane.  
Reason : Acetylene has sp character of carbon and therefore more s-character.
- 18.Assertion : Dipole moment of cis Isomer is less than the trans isomer.  
Reason : cis and trans are the geometrical isomers.

### **SECTION-B**

- 19.(a) What is the lowest value of n that allows g orbital to exist?  
(b) How many electrons in an atom may have the following quantum numbers?

$$n = 4, m_s = -\frac{1}{2} \quad (2 \times 1 = 2)$$

- 20.(a) What will be the effect on equilibrium of the following reaction, when volume of vessel increases?



What will be the value of Kc for the reverse reaction, if Kc for the decomposition of phosphorus pentachloride is  $8.3 \times 10^{-3}$  (2 \times 1 = 2)

21. Balance the following equation by ion electron method in acidic medium.



22. Draw the resonance structures for the compound  $\text{CH}_2=\text{CH}-\text{C}=\text{O}$



23. What do you mean by buffer solution. Explain the types of buffer with suitable example. (2)
24. Discuss the mechanism of Friedal Craft Acylation of benzene ring. (2)

OR

Draw Newman projections for the conformations of ethane. Which of these conformations is more stable and why? (2)

25. Arrange the elements N, P, O and S in the order of  
(a) Increasing first ionization enthalpy  
(b) Increasing Non-metallic character (2)

OR

Assign the position of the elements having outer electronic configuration.

(a)  $ns^2 np^4$  for  $n=3$

(b)  $(n-1) d^2 ns^2$  for  $n=4$  (2)

**SECTION-C**

26. In three moles of ethane ( $C_2H_6$ ) calculate the following

(a) Number of moles of carbon.

(b) Number of moles of hydrogen atoms

(c) Number of molecules of ethane (3x1=3)

27. The first  $IE_1$  and second  $IE_2$  ionisation enthalpies (KJ/mol) of three elements

A, B and C are given below

Element	$IE_1$	$IE_2$
A	403	2640
B	549	1060
C	1142	2080

Identify the element which is likely to be

(a) a non metal

(b) An alkali metal

(c) An alkaline earth metal (3x1=3)

28. Enthalpy and entropy changes of a reaction are 40.63 KJ/mol and 108.8 J/K/mol respectively. Predict the feasibility of the reaction at  $27^\circ C$ . (3x1=3)

OR

The standard enthalpies of formation of  $SO_2(g)$  and  $SO_3(g)$  are - 296.6 KJ and -395.6 KJ respectively. Calculate  $\Delta H^\circ$  for the reaction.



29. (a) 0.3780 gm of an organic compound gave 0.5740 gm of silver chloride in carius estimation. Calculate the %age of chlorine present in the compound

(b) What is the principle of paper chromatography? (2+1)

OR

Explain the term Inductive effect and electromeric effect with suitable examples. (3)

30.(a) Calculate the pH value of 0.01 M NaOH

(b) What will be the conjugate base of  $\text{HSO}_4^-$  (2+1)

### **SECTION-D**

The following questions are case based questions. Read the case carefully and answer the questions.

31. In thermodynamics, the energy changes may be measured in the laboratories under two common conditions: One in which the volume of the system is kept constant and other in which the pressure applied on the system is kept constant. The energy change at constant volume is called internal energy change ( $\Delta U$ ) and energy change at constant pressure is called enthalpy change ( $\Delta H$ ).

The two quantities are related to each other as  $\Delta H = \Delta U + P\Delta V$ . The heat changes reported are enthalpy changes because most of the processes are carried out in open vessels i.e. at-constant pressure. The common enthalpy changes are enthalpy of solution, enthalpy of neutralization, enthalpy of hydration etc.

Answer the following questions:-

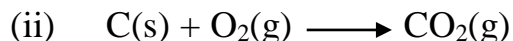
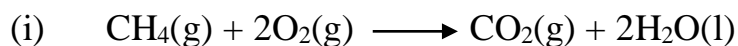
(a) When a reaction is carried out at constant volume, the heat evolved at 298K – 87.425 KJ. Calculate the enthalpy change for this reaction of ammonia formation.



OR

(a) Under what conditions heat exchange at constant volume becomes equal to heat exchange at constant pressure. Explain it by taking suitable example. (2)

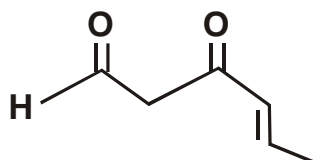
(b) For which of the following reaction  $\Delta H < \Delta U$  and  $\Delta H > \Delta U$



32. Organic compounds are vital for sustaining life on earth. We depend on these compound for our food, clothing medicines etc. For systematic study of organic compounds, we classify these compounds depending upon their structural features and chemical behaviour. We have assigned name to these organic compounds on the basis of certain standard rule as per IUPAC system of naming. The name of organic compound consist of three parts word root, suffix and prefix.

Answer the following questions

(a) Write the IUPAC name of the compound



(b) Draw the bond line formula for the compound pent-3-en-1-yne

(c) Draw the structures and write the IUPAC names of first four homologues members of carboxylic acid series. (1+1+2)

OR

(c) Give IUPAC names of the following compounds



### SECTION-E

33.(a) A golf ball has a mass of 40gm and a speed of 45 m/s. If the speed can be measured with accuracy of 2%. Calculate the uncertainly in position.

(b) Write two difference between orbit and orbital. (3+2)

OR

(a) Two particles A and B are in motion. If the wavelength associated with the particles A is  $5 \times 10^{-8} \text{m}$ , calculate the wavelength of particle B if its momentum is half of A.

(b) Write two differences between emission and absorption spectrum. (3+2)

34.(a) Describe the hybridization in case of  $\text{PCl}_5$ . Why the axial bonds are longer than equatorial bonds in  $\text{PCl}_5$ .

(b) Why  $\text{BeH}_2$  molecule has a zero dipole moment although, the Be-H bonds are polar. (3+2)

OR

(a) Compare relative stability of the following species and indicate their magnetic properties i.e.  $\text{O}_2$ ,  $\text{O}_2^+$ ,  $\text{O}_2^-$  on the basis of Molecular Orbital Theory.

(b) Although  $\text{CO}_2$  and  $\text{H}_2\text{O}$  are triatomic molecules, the shape of  $\text{H}_2\text{O}$  molecule is bent while that of  $\text{CO}_2$  is linear. Explain on the basis of dipole moment.

(3+2)

35.(a) Write a short note on following name reactions

- (i) Sabatier Sanderson's reaction
- (ii) Swartz reaction
- (iii) Wurtz reaction

(b) Write the product and their IUPAC name obtained when hex-1-ene reacts with  $\text{HBr}$  in the presence of peroxide. (3+2)

OR

(a) Convert ethane into butane.

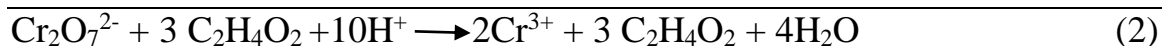
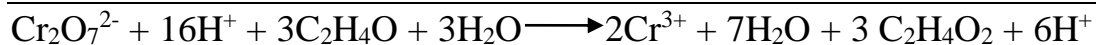
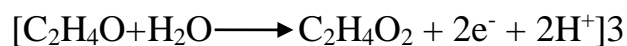
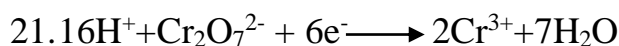
(b) What product is formed when vapours of ethyne are passed over red hot iron tube.

(c) Ozonolysis of an alkene 'x' followed by decomposition with water and a reducing agent gave a mixture of two isomers of the formula  $\text{C}_3\text{H}_6\text{O}$ . Give the structure of the alkene and its IUPAC name. (1+1+3)

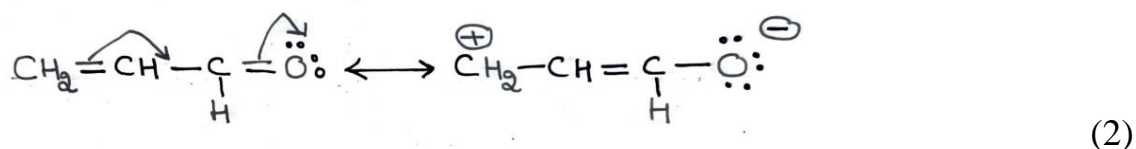
1. (a) 2 (1)
2. (b) 6 unpaired e<sup>-</sup> (1)
3. (c) F<Cl>Br>I (1)
4. (d) O<sub>2</sub><sup>2-</sup> (1)
5. (b) 18 gm/mol (1)
6. (c) q=0 (1)
7. (a) +2 (1)
8. (a) electron deficient species (1)
9. (c) < ΔU° (1)
- 10.(c) (1)
- 11.(a) 0 (1)
- 12.(a) planar (1)
- 13.(b) 6.022 x10<sup>21</sup> molecules (1)
- 14.(a) 1 (1)
- 15.(d) Assertion(A) is false but Reason(R) is true. (1)
- 16.(a) Assertion(A) and Reason(R) are correct and (R) is correct explanation of Assertion(A) (1)
- 17.(a) Assertion(A) and Reason(R) are correct and (R) is the correct explanation of Assertion(A) (1)
- 18.(d) Assertion(A) is wrong statement Reason(R) is correct statement. (1)

**SECTION-B**

- 19.(a) n=5 (1)
- (b) 16e<sup>-</sup> (1)
- 20.(a) as the volume increases, pressure decreases so equilibrium move in forward direction where number of moles increases. (1)
- (b)  $K_C(\text{Reverse}) = \frac{1}{8.3 \times 10^{-3}} = 120.48$  (1)



22.



23. Buffer solution is the solution which resist change in the pH value of solution when small amount of acid or base is added to it.

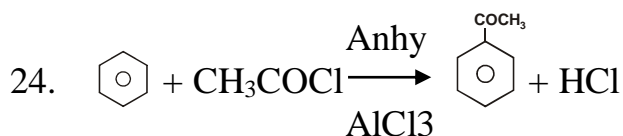
Buffer solutions are of two types

(a) Acidic buffer:- It contains equimolar quantities of weak acid and its salt with strong base.

Example acetic acid and sodium acetate

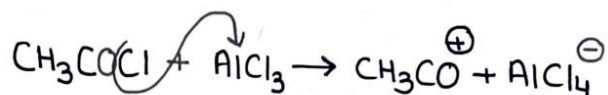
(b) Basic Buffer:- It contains equimolar quantities of weak base and its salt with strong acid

Example ammonium hydroxide and ammonium chloride (2)

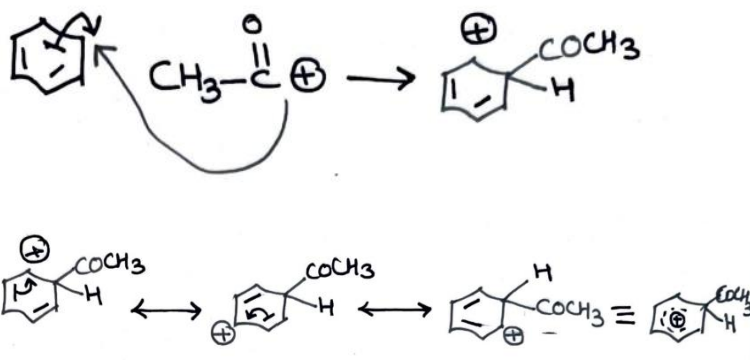


### Mechanism

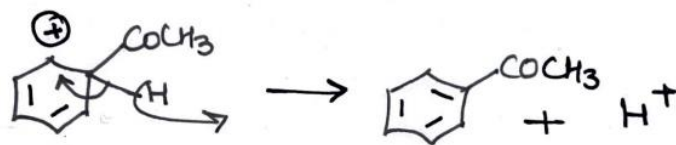
(i) Generation of Electrophile



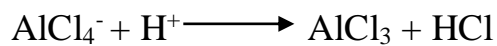
(ii) Attack of electrophile to form intermediate carbocation.



(iii) Loss of proton from carbocation

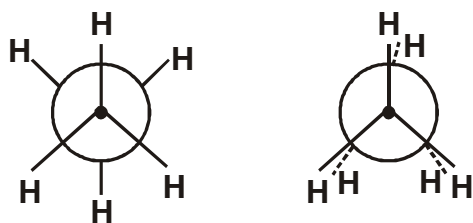


(2)



OR

### Newman's Representation



(1)

Staggered

Eclipsed

Staggered Conformation is more stable due to less repulsions.

(1)

25.(a)  $\text{S} < \text{P} < \text{O} < \text{N}$

(1)

(b)  $\text{P} < \text{S} < \text{N} < \text{O}$

(1)

OR

(a)  $3s^2 3p^4$

Group  $\longrightarrow$  16<sup>th</sup> group

Block  $\longrightarrow$  p-block

Period  $\longrightarrow$  3<sup>rd</sup> period

(1)

(b)  $4s^2 \quad 3d^2$

Group  $\longrightarrow$  4<sup>th</sup> group

Block  $\longrightarrow$  d-block

Period  $\longrightarrow$  4<sup>th</sup> period

(1)

### SECTION-C

26.(a) 1 mole  $\text{C}_2\text{H}_6$  contains 2 moles of carbon, 3 mole  $\text{C}_2\text{H}_6$  contains  $3 \times 2 = 6$  moles carbon.

(1)

(b) 1 mole  $\text{C}_2\text{H}_6$  contains 6 mole hydrogen, 3 mole  $\text{C}_2\text{H}_6$  contains  $6 \times 3 = 18$  mole hydrogen.

(1)

(c) 1 mole  $C_2H_6 = 6.022 \times 10^{23}$  molecules

3 mole  $C_2H_6 = 3 \times 6.022 \times 10^{23}$  molecules

$$= 18.066 \times 10^{23} \text{ molecules} \quad (1)$$

27. (a) C non metal (1)

(a) A alkali metal (1)

(b) B alkaline earth metal (1)

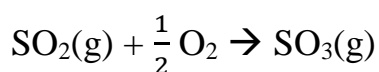
28.  $\Delta G = \Delta H - T \Delta S$  (1)

$$= 40.63 \times 1000 - 300 \times 108.8$$

$$= 7990 \text{ J/mol} \quad (1)$$

$\Delta G = +ve \therefore$  reaction is not feasible (1)

OR



$$\Delta H = \Sigma \Delta_f H^\circ \text{ products} - \Sigma \Delta_f H^\circ \text{ reactants} \quad (1)$$

$$= \Delta_f H^\circ SO_3 - \Delta_f H^\circ SO_2$$

$$= -395.6 - (-296.6) \quad (1)$$

$$= -395.6 + 296.6$$

$$= -99 \text{ KJ} \quad (1)$$

29.(a)  $AgCl \equiv Cl$

$$\begin{array}{r} 108+35.5 \\ 143.5 \end{array} \quad \begin{array}{r} 35.5 \\ 35.5 \end{array}$$

$$\% \text{ of Cl} = \frac{35.5}{143.5} \times \frac{\text{Amt. of AgCl formed}}{\text{Wt. of Organic Compound}} \times 100$$

$$\% \text{ of Cl} = \frac{35.5}{143.5} \times \frac{0.5740}{0.3780} \times 100 \quad (2)$$

$$= 37.57\%$$

(b) Paper chromatography is a type of partition chromatography which is based upon the differences in the tendencies of substance to distribute between two phases. (1)

OR

(c) The process of displacement of  $\sigma$  electrons along the saturated carbon chain due to the presence of a polar covalent bond at one end of the chain is called inductive effect (I effect). It is of two types.

- (i) + I effect  $\rightarrow$  substituent has less  $e^-$  attracting power than H.
- (ii) - I effect  $\rightarrow$  substituent has more  $e^-$  attracting power than H.

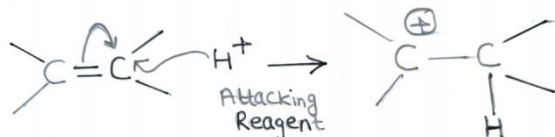
+ I effect      example -  $\text{CH}_3$ , -  $\text{C}_2\text{H}_5$

- I effect      example -  $\text{NO}_2$ , -  $\text{CN}$ , -  $\text{X}$  (1½)

Electromeric effect is the complete transfer of shared pair of  $\pi e^-$  to one of the atom joined by multiple bond. It is of two types

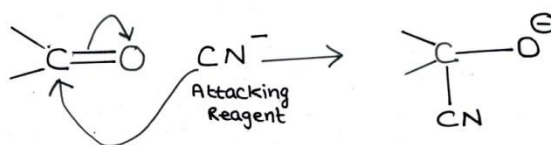
- (i) + E effect
- (ii) - E effect

- (i) + E effect is when  $\pi e^-$ s are transferred to atom to which attacking reagent get attached example

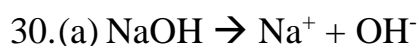


- (ii) - E effect is when  $\pi$  bond  $e^-$  all transferred to atom other than the one to which reagent get attached.

Example



(1½)



0.01 M      0.01M

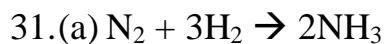
$$K_w = [\text{H}^+] [\text{OH}^-] = 10^{-14}$$

$$[\text{H}^+] = \frac{10^{-14}}{0.01} \times 100 = 10^{-12} \quad (1)$$

$$\text{pH} = -\log [\text{H}^+]$$

$$= -\log 10^{-12}$$

$$= 12 \quad (1)$$



$$\Delta n_g = 2 - 4 = -2$$

$$\Delta H = \Delta U + \Delta n_g RT$$

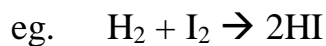
$$= -87.425 \times 1000 + (-2) \times 8.314 \times 298 \quad (2)$$

$$= -87425 - 4955.14$$

$$= -77514.7\text{J}$$

OR

(a)  $\Delta H = \Delta U$  when  $\Delta n_g = 0$



$$\Delta n_g = 2 - 2 = 0$$

$$\therefore \Delta H = \Delta U + 0RT$$

$$\therefore \Delta H = \Delta U$$

$$\Delta H = \Delta U + P\Delta V$$

when  $\Delta V = 0$  then  $\Delta H = \Delta U$  (2)

OR

any other relevant example.

(b) (i)  $\Delta n_g = 1 - 3 = -2$      $\Delta H = \Delta U - 2RT$      $\Delta H < \Delta U$

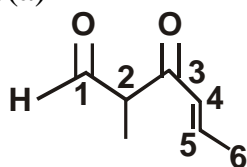
(ii)  $\Delta n_g = 1 - 1 = 0$      $\Delta H = \Delta U + 0$

(iii)  $\Delta n_g = 1 - 0 = 1$      $\Delta H = \Delta U + RT$      $\Delta H > \Delta U$

$\Delta H < \Delta U$  in (i) (1)

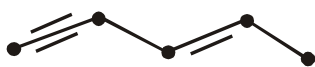
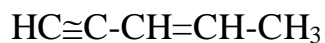
$\Delta H > \Delta U$  in (iii) (1)

32.(a)



3-keto-2-methylhex – 4 – en – 1 – al (1)

(b) pent - 3 – en -1 - yne

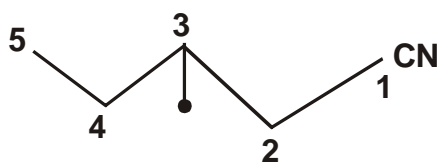


(1)

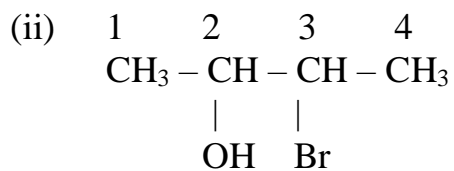


OR

(i)



3-methyl pentane nitrile (1)



3-Bromobutan-2-ol (1)

### SECTION-E

33.(a)  $\Delta V = 45 \times \frac{2}{100} = 0.9$  (1)

$$m = \frac{40}{1000}$$

$$\Delta x = \frac{h}{4\pi m \Delta v} \quad (1)$$

$$= \frac{6.626 \times 10^{-34}}{4 \times 3.14 \times 40 \times 10^{-3} \times 0.9}$$

$$= 1.46 \times 10^{-33} \text{ m} \quad (1)$$

- |     |   |   |
|-----|---|---|
| (b) | Orbit   | Orbital   |
| (a) | It is well defined circular path around which $e^-$ revolve | It is region in space around nucleus where the probability to find $e^-$ is maximum |
| (b) | Represent planar motion                                     | three dimensional motion of electron  |

OR / any other relevant difference. (2)

OR

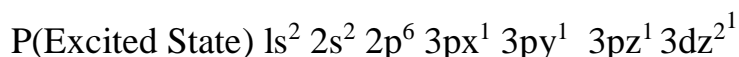
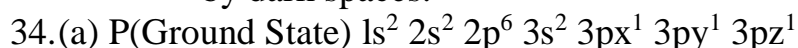
$$(a) \frac{\lambda_A}{\lambda_B} = \frac{h}{P_A} \frac{P_B}{h} = \frac{P_B}{P_A} \quad (1)$$

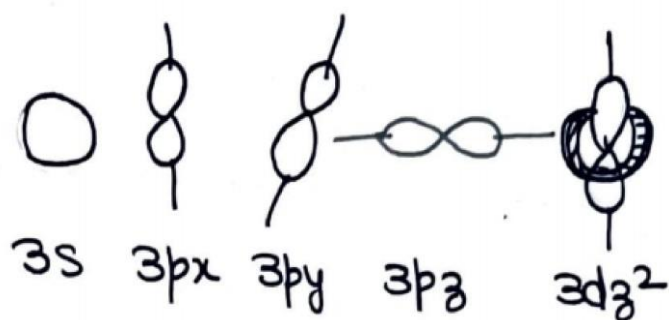
$$P_B = \frac{P_A}{2}$$

$$\frac{\lambda_A}{\lambda_B} = \frac{P_A}{2P_A} = \frac{1}{2} \quad (1)$$

$$\frac{5 \times 10^{-8}}{\lambda_B} = \frac{1}{2} \therefore \lambda_B = 10 \times 10^{-8} \text{ m} = 10^{-7} \text{ m} \quad (1)$$

- |      |   |   |
|------|---|---|
| (b)  | Emission Spectrum   | Absorption Spectrum   |
| (i)  | It is obtained when radiation emitted by the excited substance are analysed with spectroscope | It is obtained when white light is passed through solution and transmitted light is analysed through spectroscope |
| (ii) | Emission spectrum consist of bright coloured lines separated by dark spaces.                  | Consist of dark lines in otherwise continuous spectrum  |
- (2)



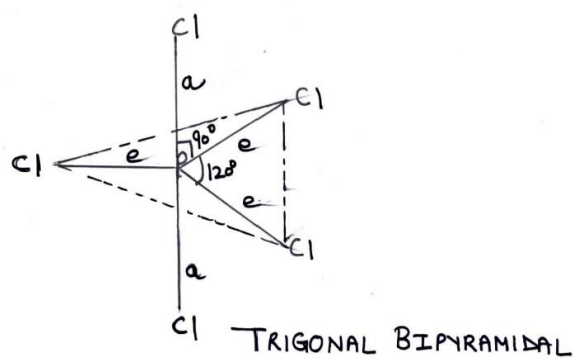
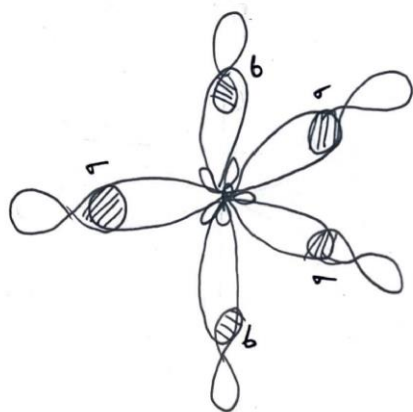


(1)

5 orbital hybridise and form 5 new orbitals of same energy and same shape.



5 sp<sup>3</sup>d hybrid orbitals adopt trigonal bipyramidal arrangement which overlap with 3p<sub>z</sub> orbital to form 5  $\sigma$  bonds



(1)

Axial bonds are longer than equatorial bonds due to greater repulsions from other bonds in axial position. (1)

- (b) BeH<sub>2</sub> is linear molecule. Therefore the resultant dipole moment of two Be-H bonds get cancelled giving zero dipole moment.



OR

$$\begin{aligned} \text{(a) } \text{O}_2 &= \sigma 1s^2 \sigma^* 1s^2 \sigma 2s^2 \sigma^* 2s^2 \sigma 2p_z^2 \pi 2p_x^2 = \pi 2p_y^2 \pi^* 2p_x^1 = \pi^* 2p_y^1 \\ \text{O}_2^+ &= \sigma 1s^2 \sigma^* 1s^2 \sigma 2s^2 \sigma^* 2s^2 \sigma 2p_z^2 \pi 2p_x^2 = \pi 2p_y^2 \pi^* 2p_x^1 = \pi^* 2p_y^0 \\ \text{O}_2^- &= \sigma 1s^2 \sigma^* 1s^2 \sigma 2s^2 \sigma^* 2s^2 \sigma 2p_z^2 \pi 2p_x^2 = \pi 2p_y^2 \pi^* 2p_x^2 = \pi^* 2p_y^1 \end{aligned} \quad (1)$$

$$\text{Bond order } \text{O}_2 = \frac{8-4}{2} = \frac{4}{2} = 2$$

$$\text{O}_2^+ = \frac{8-3}{2} = \frac{5}{2} = 2.5$$

$$\text{O}_2^- = \frac{8-5}{2} = \frac{3}{2} = 1.5 \quad (1)$$

Higher is the bond order, more is the stability.

$$\text{O}_2^+ > \text{O}_2 > \text{O}_2^- \quad (1)$$

- (b) CO<sub>2</sub> has zero dipole moment so CO<sub>2</sub> is linear as two C=O bond moments get cancelled where as H<sub>2</sub>O molecule has resultant dipole moment. Two O-H bonds are arranged in angular shape and the bond moment of two O-H bonds give resultant dipole moment. (2)

35.(a)

