

IIT-JAM - CHEMISTRY

MOCK TEST PAPER(According to new pattern)

- Attempt ALL the 60 questions.
- SECTION-A Consists of 30 questions. These questions are Multiple Choice Questions (MCQs), First 20 questions carries one marks for each and remaining 10 questions carries two marks for each.
- Section-B Consists of 10 questions. These questions are Multiple Select Questions (MSQs), each question carries two marks.
- Section-C Consists of 20 Numerical Answer Type (NAT) questions each question carries two marks. For each NAT type question, the answer is a signed real number.
- In Section A, for all 1 mark questions, 1/3 marks will be deducted for each wrong answer and for all 2 marks questions, 2/3 marks will be deducted for each wrong answer. There is no negative marking in Section B and Section C.

• Total marks : 100

• Duration of test : 3 Hours

VPM CLASSES

For IIT-JAM, JNU, GATE, NET, NIMCET and Other Entrance Exams

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SECTION-A (Q. 1-30)**MULTIPLE CHOICE QUESTIONS (MCQs)**

- The correct order of electron affinity among the following is :
 - $F > Cl > Br$
 - $Cl > F > Br$
 - $Br > Cl > F$
 - $F > Br > Cl$
- In which of the following group of species, the hybrid state of carbon is same ?
 - CH_4 , C_2H_2 , charcoal
 - ethene, graphite, CO_2
 - CO_2 , propane, diamond
 - CH_4 , diamond, cyclohexane
- A unit cell consists of a cube in which there are anions (B) at each corner and one at the centre of the unit cell and cations (A) at the centre of each face. What is the simplest formula for this compound ?
 - AB
 - AB_2
 - A_3B_2
 - A_2B_3
- The nature of $B(OH)_3$ boric acid is
 - Triacidic Base
 - Tribasic acid
 - Monoacidic Base

- (D) Mono basic acid
5. XeF_6 on reaction with quartz gives :
- (A) XeO_4
(B) XeO_3
(C) XeF_4
(D) XeOF_4
6. How many diastereomers are shown by the following molecule ?
 $\text{CH}_3\text{CH} = \text{CHCH} = \text{CHC}_2\text{H}_5$
- (A) 1
(B) 2
(C) 3
(D) 4
7. The order of stability of following carbocations is ,
- (i) Tropylium cation
(ii) Allyl carbocation
(iii) Benzyl carbocation
(iv) Triphenylmethyl carbocation
- (A) $i > ii > iii > iv$
(B) $i > iv > iii > ii$
(C) $iv > iii > ii > i$
(D) $ii > iii > iv > i$

8. The order of aromaticity of the following heterocycles

(i) Thiophene,

(ii) Pyrrole,

(iii) Furan

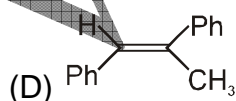
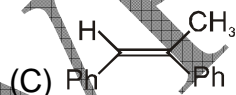
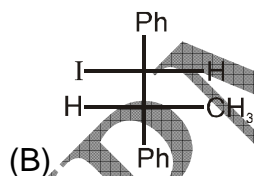
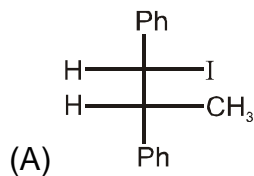
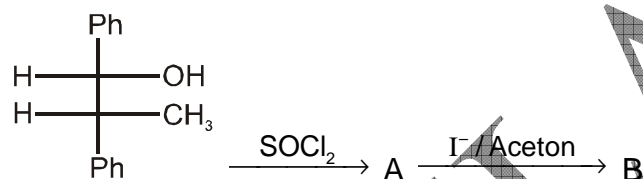
(A) ii > iii > i

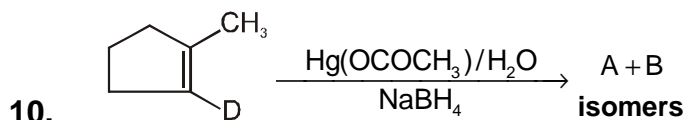
(B) ii > i > iii

(C) i > ii > iii

(D) iii > ii > i

9. The end product formed in the following reaction is





A & B isomers are :

- (A) Identical
(B) Meso
(C) Diastereomers
(D) Pair of Enantiomers
11. What are the component values (in terms of $h/2\pi$) of the orbital angular momentum along the z-direction for a 2p electron ?
- (A) $+\frac{1}{2}, -\frac{1}{2}$
(B) $+\frac{3}{2}, +\frac{1}{2}, -\frac{1}{2}, -\frac{3}{2}$
(C) +2, +1, 0, -1, -2
(D) +1, 0, -1
12. The correct order of first ionization potential is
- (A) He > F > O > N > Mg
(B) N > F > He > O > Mg
(C) He > F > N > O > Mg
(D) F > He > N > Mg > O

13. Which of the following species have the same bond order ?

- (I) CO
- (II) CN^-
- (III) O_2^+
- (IV) NO^+
- (A) I, II, III
- (B) I, II, IV
- (C) I, III, IV
- (D) II, III, IV

14. Match List-I (compound) with List-II (shape) and select the correct answer :

List-I (Compound)

List-II (Shape)

- | | |
|----------------------|--------------------------|
| (a) XeF_5^+ | (1) Tetrahedral |
| (b) SiF_5^- | (2) Square planar |
| (c) AsF_4^+ | (3) Trigonal bipyramidal |
| (d) ICl_4^- | (4) Square pyramidal |
| | (5) Octahedral |

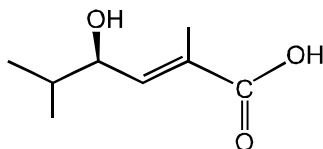
- | | a | b | c | d |
|-----|----------|----------|----------|----------|
| (A) | 1 | 2 | 5 | 3 |
| (B) | 4 | 3 | 1 | 2 |
| (C) | 1 | 3 | 5 | 2 |
| (D) | 4 | 2 | 1 | 3 |

15. HNO_3 on treatment with P_2O_5 products.

- (A) N_2O_5

- (B) NO_2
- (C) HNO_2
- (D) N_2O_4

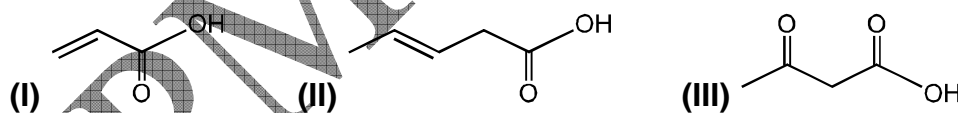
16. For the compound



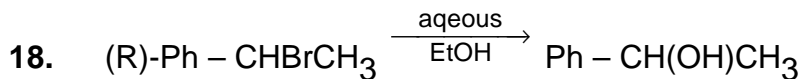
The stereochemical notations are

- (A) 2Z, 4R
- (B) 2Z, 4S
- (C) 2E, 4R
- (D) 2E, 4S

17. The correct order of decarboxylation of the three acids is

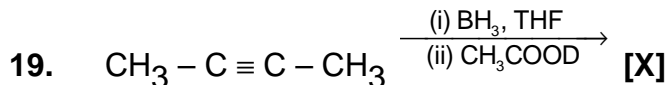


- (A) III > II > I
- (B) III = II > I
- (C) III > II = I
- (D) III = II = I

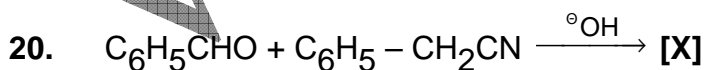
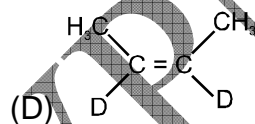
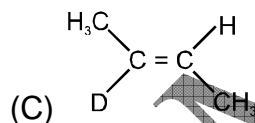
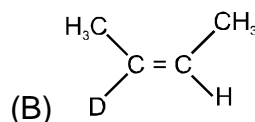
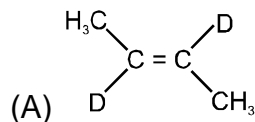


Which statement is the most likely to be correct concerning the given S_N1 reaction.

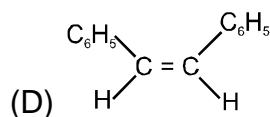
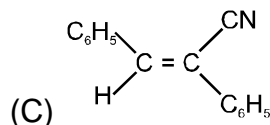
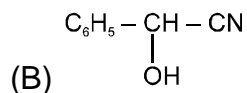
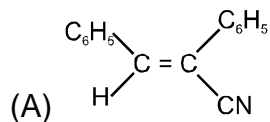
- (A) The reaction proceeds with partial racemization.
- (B) The stereochemistry of the Halide is not inverted.
- (C) The carbonium ion $\text{Ph-CH}_2\text{-CH}_2^+$ is involved.
- (D) The carbonium ion is attacked in each side to same degree.



The major product [X] is



The major product [X] is



21. The wave number of the limiting line in Lyman series of hydrogen is 109678 cm^{-1} .

The wave number of the limiting line in Balmer series of He^+ would be

(in cm^{-1}).

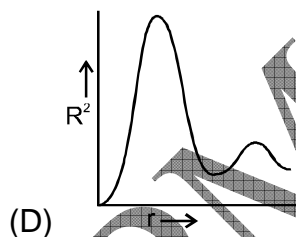
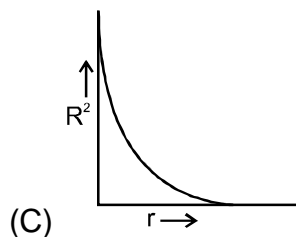
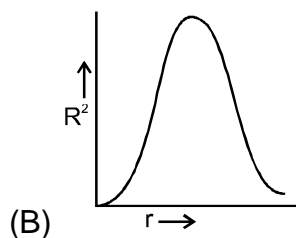
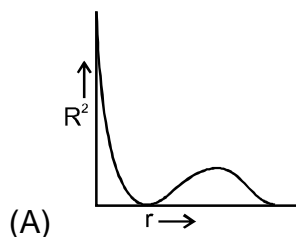
(A) 107879 cm^{-1}

(B) 1096.78 cm^{-1}

(C) 109678 cm^{-1}

(D) 119658 cm^{-1}

22. Which of the following probability distribution curves represents 2s orbital for H-atom.



23. A gas causes heating effect when allowed to expand at room temperature, it shows heating effect because
- (A) A gas has high critical temperature.
 - (B) A gas has inversion temperature much below room temperature.
 - (C) A gas has high Inter molecular forces above room temperature.

- (D) A gas has inversion temperature above room temperature.
24. An ideal gas initially at 710 torr and 30°C occupies 2600 ml. The final temperature if the conditions are changed to a pressure of 1.20 atm and a volume of 3.20 L is :
- (A) 306°C
(B) 206°C
(C) 312°C
(D) 226°C
25. An ideal gas occupying a volume of 2 dm³ at a pressure of 5 bar undergoes isothermal and irreversible expansion against external pressure of 1 bar. The final volume of the system and the work involved in the process is :
- (A) 10 dm³, 1000 J
(B) 10 dm³, -1000 J
(C) 8 dm³, -800 J
(D) 10 dm³, -800 J
26. Which of the following statements are correct about the following complex $[\text{CrCl}_2(\text{OH})_2(\text{NH}_3)_2]^-$
- (A) It shows geometrical isomerism
(B) It does not show optical isomerism
(C) It shows ionization isomerism
(D) It shows hydrate isomerism

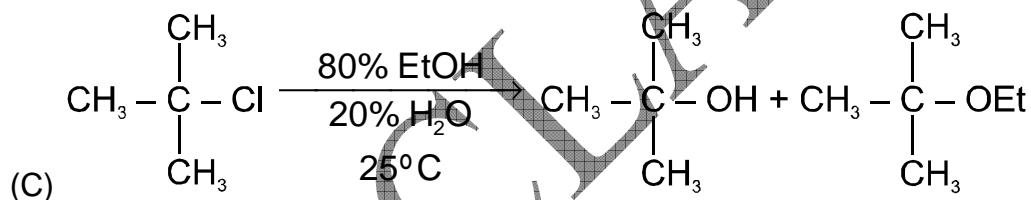
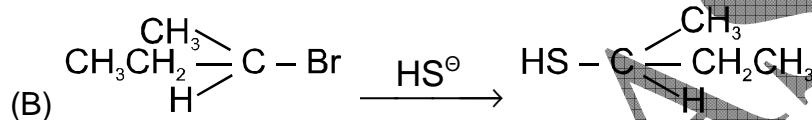
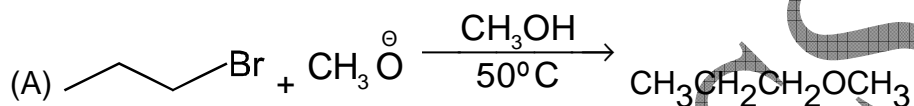
27. Which of the following compound on reaction with NaOH and Na₂O₂ gives yellow colour?
- (A) Cr(OH)₃
(B) Zn(OH)₂
(C) Al(OH)₃
(D) None of these
28. What volume of 5.00 N H₂SO₄ is required to neutralize a solution containing 2.50 g NaOH ?
- (A) 16.4 ml
(B) 10.2 ml
(C) 13.5 ml
(D) 12.5 ml
29. In the trigonal bipyramidal crystal field the d-orbital with the Highest energy is
- (A) dxy
(B) dx² - y²
(C) dyz
(D) dz²
30. What mass of N₂H₄ can be oxidized to N₂ by 24.0 g K₂CrO₄, which is reduced to Cr(OH)₄⁻ ?
- (A) 4.28 g

- (B) 2.15 g
(C) 2.97 g
(D) 3.25 g

SECTION-B (Q. 31-40)

MULTIPLE SELECT QUESTIONS (MSQs)

31. Which of the following reaction occur by S_N^2 mechanism



32. Which of the following metals contain (Al) Aluminium as a metal?

- (A) Bauxite
(B) Cryolite
(C) Corundum
(D) Kaolin

33. In phase equilibria :

- (A) Sulphur system consist of six stable curve and four metastable curve.

- (B) Sulphur system consist of 3 triple point.
(C) At the triple and eutectic point the degree of freedom is zero.
(D) In water system triple point exist at 4.5 mm of Hg pressure.

34. ${}_{90}^{234}\text{Th}$ disintegrates to give ${}_{82}^{206}\text{Pb}$ for this disintegration.

- (A) The mass of α -particle emitted is four.
(B) The mass of β -particle emitted is zero.
(C) The no. of α -particle emitted is 7.
(D) The no. of β -particle emitted is 6.

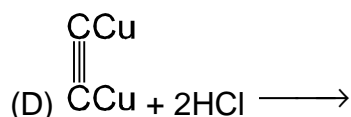
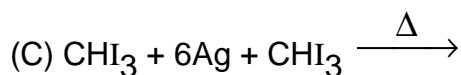
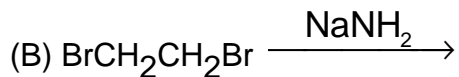
35. An Aliphatic Ketone M.W. 86, when converted into its oxime which is then reduced, gives an amine that can be resolved. What is this Ketone?

- (A) $\text{CH}_3\text{CH}_2\text{COCH}_2\text{CH}_3$
(B) $\text{CH}_3\text{COCH}_2\text{CH}_2\text{CH}_3$
(C) $\text{CH}_3\text{COCH}(\text{CH}_3)_2$
(D) $(\text{CH}_3)_3\text{CC}(\text{O})\text{H}$

36. In NMR spectroscopy :

- (A) In P_4S_3 molecule ${}^{31}\text{P}$ -NMR give quadruplet signal.
(B) In HPF_2 molecule ${}^{19}\text{F}$ -NMR give quadruplet signal.
(C) In HPF_2 molecule ${}^{19}\text{F}$ -NMR give doublet signal.
(D) In P_4S_3 molecule ${}^{32}\text{S}$ -NMR give quintet signal.

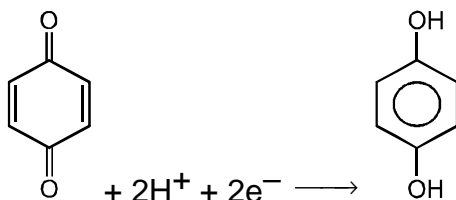
37. Oxyacids of bromine :
- (A) HOBr
 - (B) HBrO₂
 - (C) HBrO₃
 - (D) HBrO₄
38. The vapour pressure of ethanol and methanol are 44.5 mm and 88.7 mmHg respectively. An ideal solution is formed at the same temperature by mixing 60 gm of ethanol with 40 gm of methanol. So :
- (A) The total vapour pressure of solution is 66.13 mm of Hg.
 - (B) mol fraction of ethyl alcohol is .4893.
 - (C) mol fraction of methyl alcohol is .5107.
 - (D) mol fraction of methyl alcohol in vapour is 0.6563.
39. Which of the following pair is buffer :
- (A) Potassium acid phthalate + Di potassium phthalate
 - (B) Borax + NaOH
 - (C) Boric Acid + Borax
 - (D) Na₂HPO₄ + NaH₂PO₄
40. Acetylene is formed by the following reactions
- (A) $6\text{CH}_4 + \text{O}_2 \xrightarrow{\text{high temp.}}$

**PART C (41 - 60)**

41. The amount of ^{14}C isotope in a piece of wood is found to be one-sixth of its amount present in a fresh piece of wood. Calculate the age of wood. (Half life of $^{14}\text{C} = 5577$ years).
42. The heats of hydrogenation of cycloheptene and 1,3,5-cycloheptatriene are 110 kJ/mol (26.3 kcal/mol) and 305 kJ/mol (73.0 kcal/mol), respectively. In both cases cycloheptane is the product. What is the resonance energy of 1,3,5-cycloheptatriene?
43. The EMF of the concentration cell with transference, viz., Pt; H_2 (1 atm), HCl ($a_{\pm} = 0.009048$) : HCl ($a_{\pm} = 0.01751$), H_2 (1 atm); Pt, is 0.02802 V at 25°C. The EMF of the corresponding cell without transference is 0.01696 V. Calculate the liquid junction potential, E_l , and the transference number of the H^+ ion.
44. At 100°C, the specific volumes of water and steam are, respectively, 1 c.c. and 1673 c.c. Calculate the change in vapour pressure of the system by 1°C change in temperature. The molar heat of vaporisation of water in this range may be taken as 9.70 kcal.
45. N_2O_4 is 25% dissociated at 37°C and one atmospheric pressure. Calculate the percentage dissociation at 0.1 atmosphere and 37°C.

46. N_2O_4 is 25% dissociated at 37°C and one atmospheric pressure. What is the minimum pH of a solution 0.10 M in Mg^{2+} from which $\text{Mg}(\text{OH})_2$ will not precipitate?
 $K_{\text{sp}} \text{Mg}(\text{OH})_2 = 1.2 \times 10^{-11} \text{ M}^3$.
47. n-butane is produced by the monobromination of ethane followed by the wurtz reaction. Calculate the volume of ethane at STP required to produce 55gm n-butane if the bromination takes place with 90% yield and the wurtz reaction with 85% yield.
48. Calculate the wave number for the shortest wavelength transition in the Balmer series of atomic hydrogen.
49. Electromagnetic radiation of wave length 242 nm is just sufficient to ionize the sodium atom. Calculate the ionization energy of sodium in KJ mol^{-1} .
50. The composition of a sample of wustite is $\text{Fe}_{0.93}\text{O}_{1.00}$. What % of Iron is present in the form of Fe (III).
51. A metal complex having composition $\text{Cr}(\text{NH}_3)_4\text{Cl}_2\text{Br}$ has been isolated in two forms A and B. The form A react with AgNO_3 to give a white precipitate readily soluble in dilute aqueous ammonia, whereas B gives a pale yellow precipitate soluble in concentrated ammonia. What is their magnetic moment of compound A & B.
52. An acid type indicator, HIn differs in colour from its conjugate base (In^-). The human eye is sensitive to colour differences only when the ratio $[\text{In}^-]/[\text{HIn}]$ is greater than 10 or smaller than 0.1. What should be the minimum change in the pH of the solution to observe a complete colour change ($K_a = 1.0 \times 10^{-5}$)
53. For a first order reaction the energy of activation is 210 kJ mol^{-1} , and the frequency factor is $2 \times 10^{12} \text{ s}^{-1}$. What is the half life of the reaction at 700 k.
54. 36.4 gm of 1,1,2,2- tetrachloropropane was heated with zinc dust and the product was bubbled through ammoniacal AgNO_3 . What is the weight of the ppt.

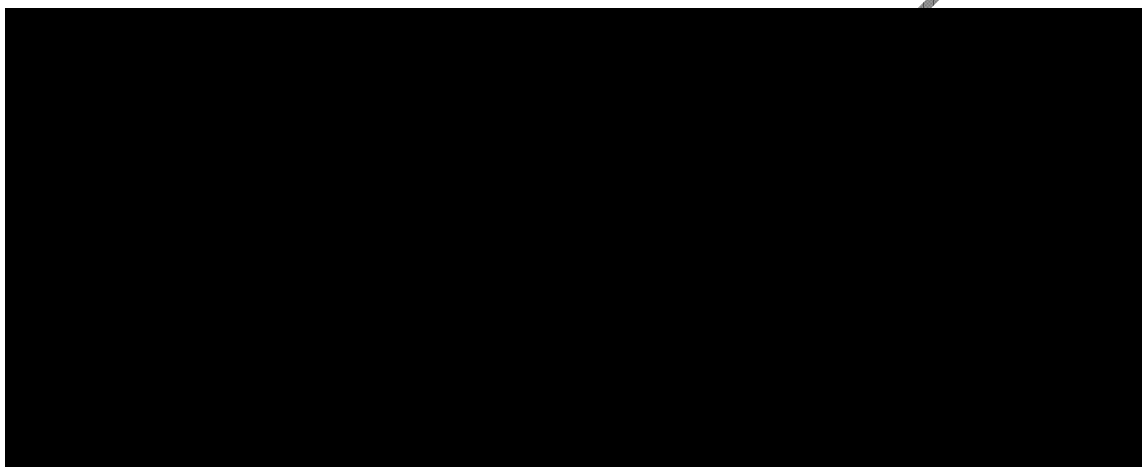
55. Calculate the number of radial nodes in 4F.
56. 0.66 g of H_3PO_2 will require x ml of 0.1 M NaOH for complete neutralization. Find out the value of x.
57. What is the number of dipeptides that can be prepared from alanine and phenylalanine?
58. Calculate the ionic strength of a solution that contains 0.1 mol/kg NaCl and 0.05 mol/kg Na_2SO_4 .
59. Calculate the half cell potential for the quinhydrone electrode



set up at pH = 4 will be (quinhydrone = 1 : 1 molecular compound of quinone (Q) and hydroquinone (QH_2), $E^\circ = 0.699 \text{ V}$).

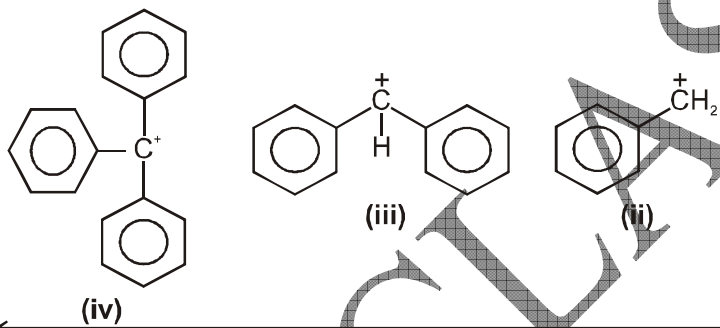
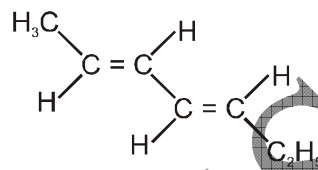
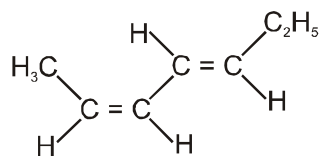
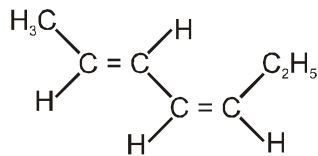
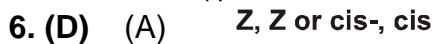
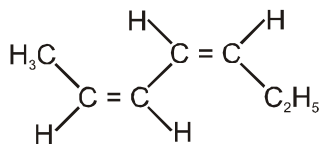
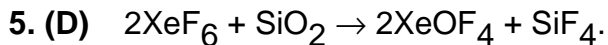
60. The equilibrium pressure of $\text{NH}_4\text{CN}(\text{s})$, $\text{NH}_3(\text{g}) + \text{HCN}(\text{g})$ is 0.298 atm. Calculate K_p . If $\text{NH}_4\text{CN}(\text{s})$ is allowed to decompose in presence of NH_3 at 0.25 atm, calculate partial pressure of HCN at equilibrium.

ANSWER KEY



HINTS & SOLUTIONS

- (B)** ∴ In a group electron affinity is decreased when we go up to down.
 * But electron affinity of F is less than that of Cl due to its extremely small size.
 So order is Cl > F > Br.
- (D)** The hybrid state of C in CH₄, diamond and cyclohexane is sp³.
- (C)** No. of anions (B) present in the unit cell = $\frac{1}{8} \times 8 + 1 \times 1 = 1 + 1 = 2$
 No. of cations (A) present in the unit cell = $6 \times \frac{1}{2} = 3$
 ∴ Formula of the compound = A₃B₂
- (D)** B(OH)₃ is Mono basic acid.



7. (B)

Resonance
Stability of carbocation ↑

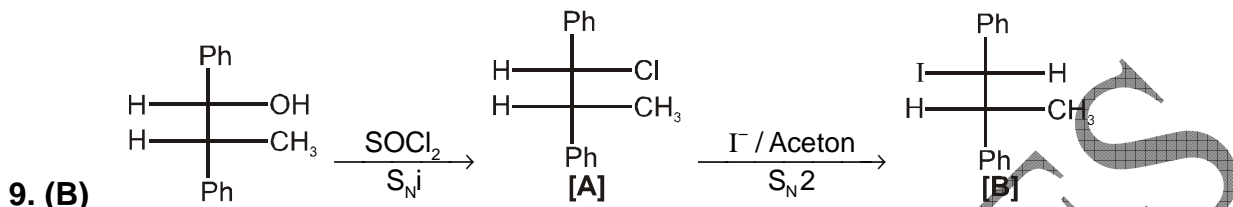
In iv & i, (i) is more stable due to aromaticity.

Aromatic

So order of carbocation stability $i > iv > iii > ii$.

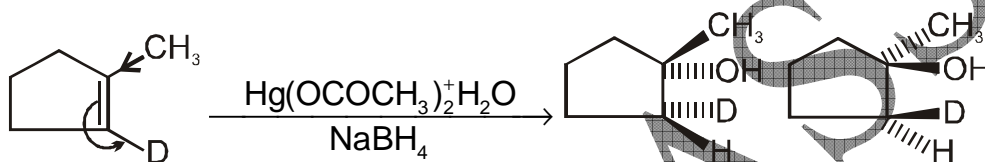
8. (C) The aromatic character in these p-excessive heterocycles depends on the availability of lone pair of electrons on the heteroatom for cyclic delocalization which, in turn, depends on the electronegativity of heteroatom. The more electronegative heteroatom will have stronger attraction for the lone pair of electrons and will make it less available for cyclic delocalization. The aromaticity order in these heterocycles depends on the electronegativity of the heteroatom : O

> N > S and therefore, the aromaticity follows the order as : thiophene > pyrrole > furan.



10.(D) Oximercuration Demercuration : [(i) $\text{Hg}(\text{OCOCH}_3)_2/\text{H}_2\text{O}$; (ii) NaBH_4]

It's give anti addition of H_2O on alkene acc. to marckonicoff rule. So the reaction is



or **Pair of Enantiomer**

11.(D) The component values of orbital angular momentum in z-direction = $m_\ell \times$

m_ℓ for 2p = +1, 0, -1

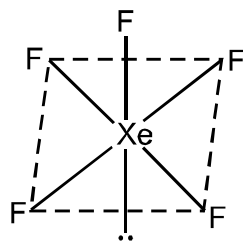
12.(C) He F N O Mg

$1s^2 > 2p^5 > 2p^3 > 2p^4 > 3s^2$

13.(B) CO (10 valence electrons) : B.O. = 3; CN^- (10) : B.O. = 3, O_2^+ (11) : B.O. = 2.5;

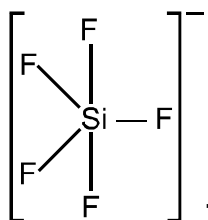
NO^+ (10) : B.O. = 3

14.(B) XeF_5^- — sp^3d^2 Hybridization



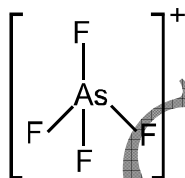
Square Pyramidal

SiF_5^- — sp^3d Hybridization



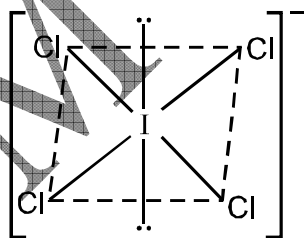
Trigonal bipyramidal

AsF_4^+ — sp^3 Hybridization



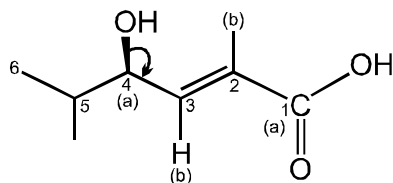
Tetrahedral

ICl_4^- — sp^3d^2 Hybridization



Square planar

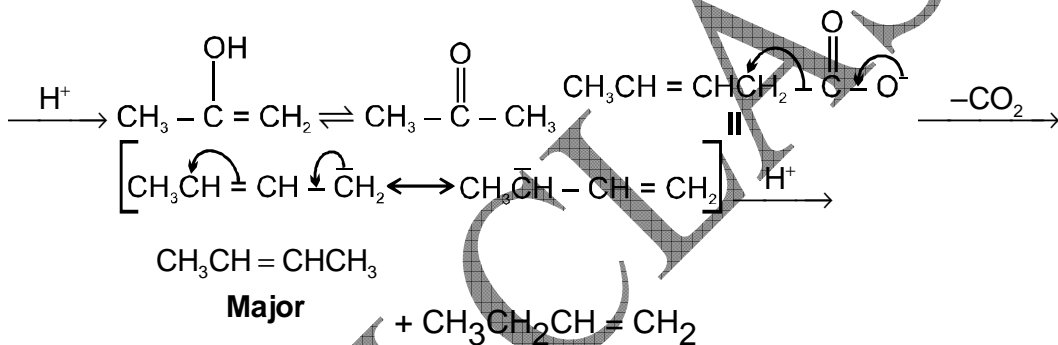
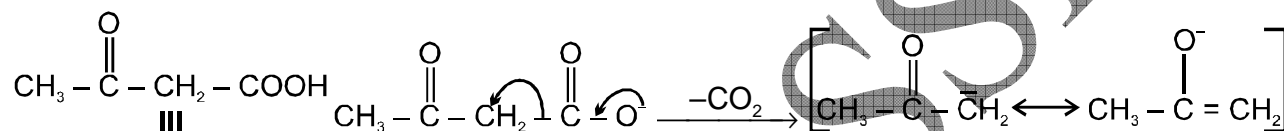
15.(A) $2\text{P}_2\text{O}_5 + 4\text{HNO}_3 \rightarrow 2\text{N}_2\text{O}_5 + 4\text{HPO}_3$



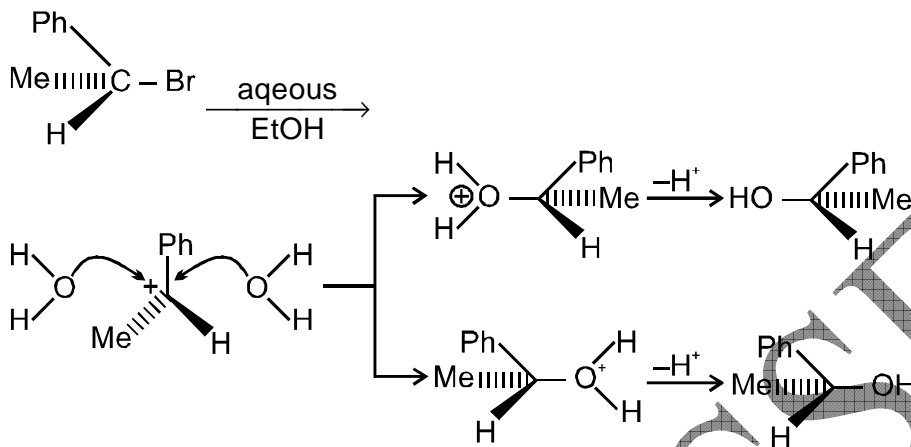
16.(C)

4R, 2E

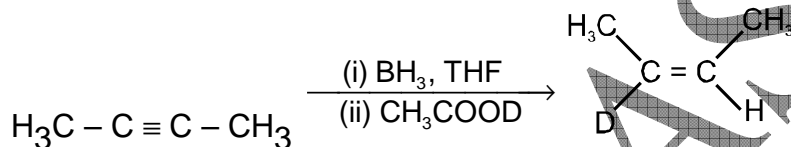
17.(B) β -Keto carboxylic acids and β , γ -unsaturated carboxylic acids undergo decarboxylation easily because the corresponding carbanion is quite stable due to resonance.



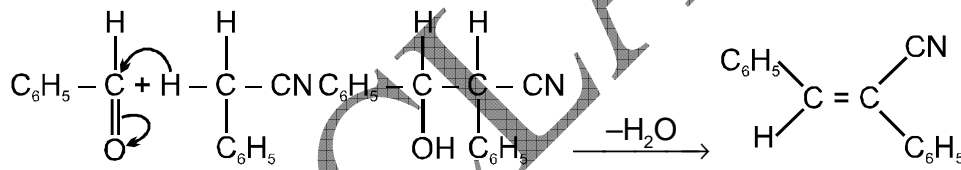
18.(D)



19.(B)



20.(C)

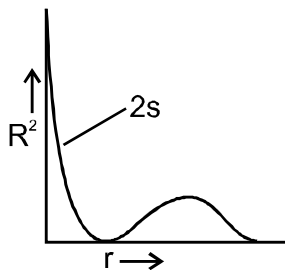


21.(C) $R_H = 109678 \text{ cm}^{-1}$

Wave number of the limiting line in Balmer Series of $\text{He}^+ = R_H \cdot Z^2 =$

$$\left(\frac{1}{n_1^2} - \frac{1}{n_2^2} \right) 109678 \times (2)^2 \left[\frac{1}{(2)^2} - \frac{1}{\infty} \right]$$

$$= 109678 \text{ cm}^{-1}$$



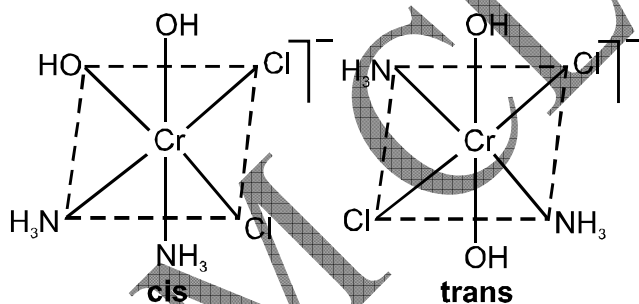
22.(A)

23.(D) If gas is allowed to expand above its inversion temperature it causes heating effect.

$$24.(B) T_2 = \frac{P_2 V_2 T_1}{P_1 V_1} = \frac{1.20 \times 760 \times 3.20 \times 303}{710 \times 2.60} = 479 \text{ K} = 206^\circ\text{C}$$

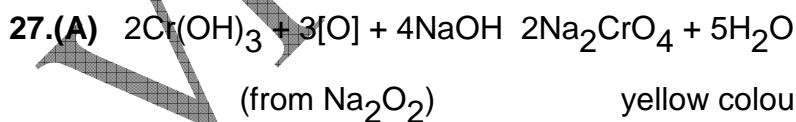
$$25.(D) V_f = \frac{P_i V_i}{P_f} = \frac{(5 \text{ bar}) \times (2 \text{ dm}^3)}{1 \text{ bar}} = 10 \text{ dm}^3$$

$$\text{work } w = -P_{\text{ext}}(V_f - V_i) = -(1 \text{ bar})(10 - 2) \text{ dm}^3 = -1 \times 10^5 \text{ Pa} \times 8 \times 10^{-3} \text{ m}^3 = -800 \text{ J}$$



26.(A)

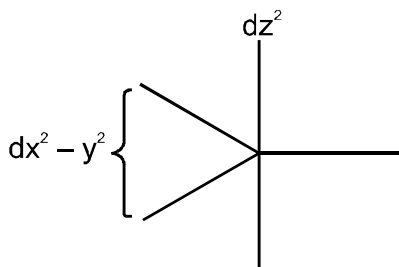
Cis isomer can show optical isomerism also because it has three different types of unidentate ligand.



$$28.(D) \text{Number of equivalents in } 2.50 \text{ g NaOH} = \frac{2.50 \text{ g}}{40.0 \text{ g/eq}} = 0.0625 \text{ eq NaOH}$$

Therefore, 0.0625 eq H₂SO₄ is required.

$$(0.0625 \text{ eq}) \left(\frac{1 \text{ L}}{5.00 \text{ eq}} \right) = 0.0125 \text{ L} = 12.5 \text{ mL of } 5.00 \text{ N solution}$$

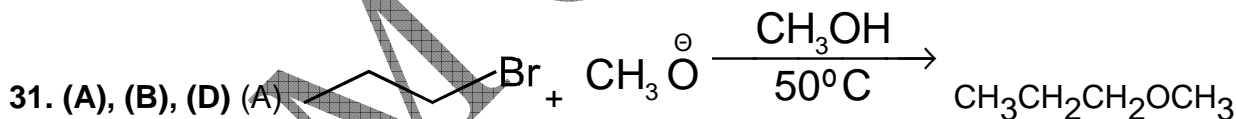


29.(D)

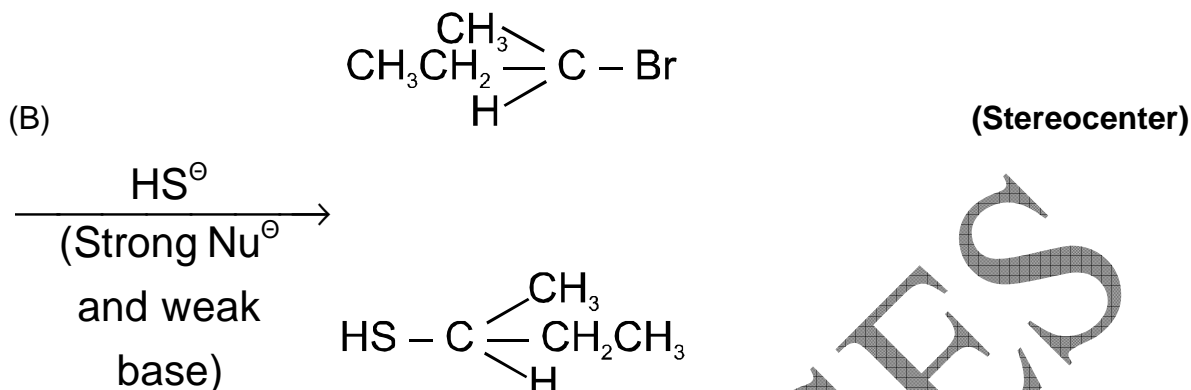
In the trigonal bipyramidal crystal field the d-orbital with the Highest energy is dz².

30.(C) The balanced equation is $4\text{H}_2\text{O} + 3\text{N}_2\text{H}_4 + 4\text{CrO}_4^{2-} \rightarrow 3\text{N}_2 + 4\text{Cr}(\text{OH})_4^- + 4\text{OH}^-$

$$(24.0 \text{ g } \text{K}_2\text{CrO}_4) \left(\frac{1 \text{ mol } \text{K}_2\text{CrO}_4}{194.2 \text{ g } \text{K}_2\text{CrO}_4} \right) \left(\frac{3 \text{ mol } \text{N}_2\text{H}_4}{4 \text{ mol } \text{CrO}_4^{2-}} \right) = (0.0927 \text{ mol } \text{N}_2\text{H}_4) = \left(\frac{32.0 \text{ g}}{\text{mol}} \right) 2.97 \text{ g}$$

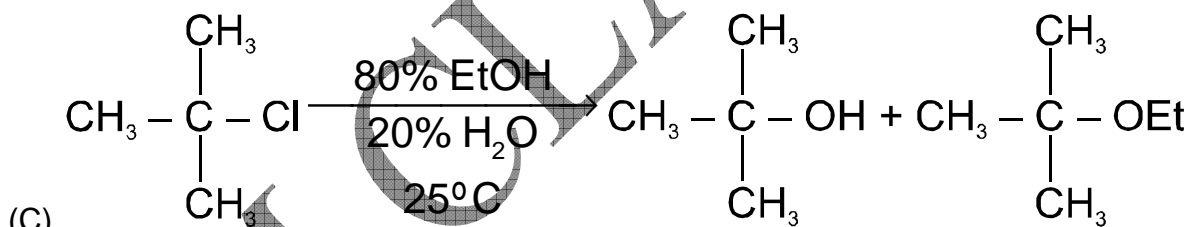


is a 1° halide. The base/Nu⁻ is a CH_3O^- strong base (not a hindered one) and a Nu⁻. Hence it give mainly SN² product.



[(R)Stereoisomer]

SN² with inversion



SN¹ mechanism

32. (A), (B), (C), (D) Bauxite — $\text{Al}_2\text{O}_3 \cdot 2\text{H}_2\text{O}$

Cryolite — Na_3AlF_6

Corundum — Al_2O_3

Kaolin (China Clay) — $\text{Al}_2\text{Si}_2\text{O}_7 \cdot 2\text{H}_2\text{O}$

33. (A), (C), (D) The sulphur system consist 6 stable curve & 4 metastable curve.

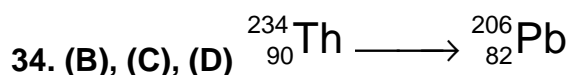
The sulphur system consist four triple point. At the triple point

$$F = C - P + 2 = 1 - 3 + 2 = 0$$

At the eutectic point

$$F = C - P + 1 = 2 - 3 + 1 = 0$$

In water system the triple point exist at .0078°C temperature & 4.58 mm of Hg pressure.



Decrease in mass = $234 - 206 = 28$

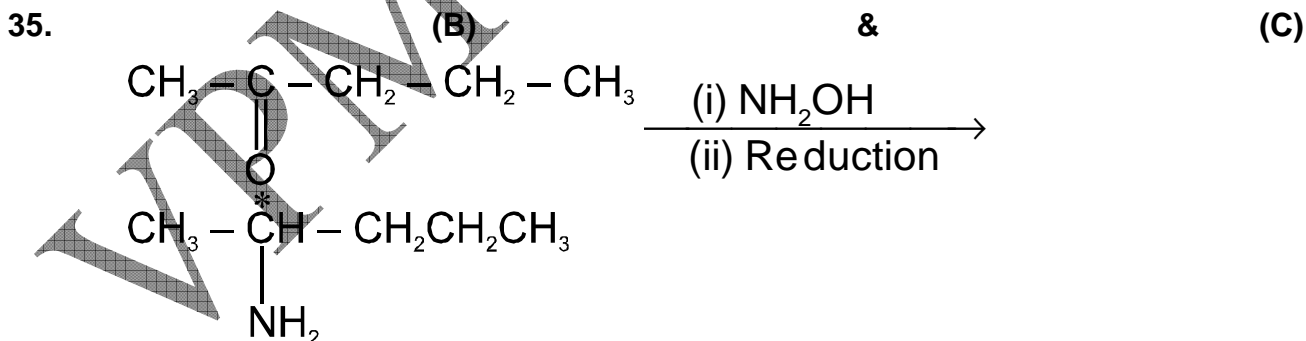
Mass of α -particle = 4

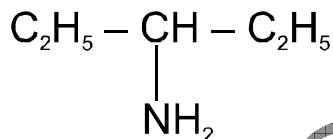
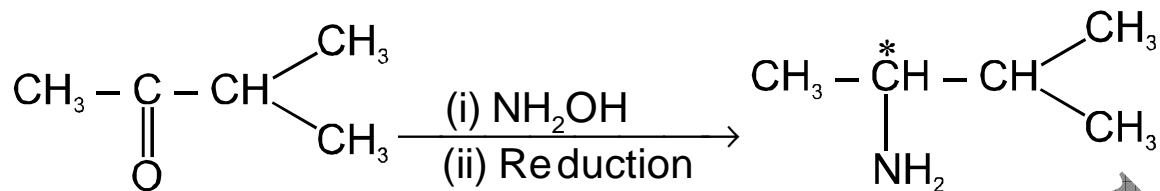
So no. of α -particle = $\frac{28}{4} = 7$

Mass of the α -particle emitted is $7 \times 4 = 28$

No. of β -particle emitted = $(2 \times \text{no. of } \alpha\text{-particle}) - (\text{atomic no. of parent} - \text{at. no. of end product}) = 14 - (8) = 6$

Mass of the emitted β -particle is = zero.

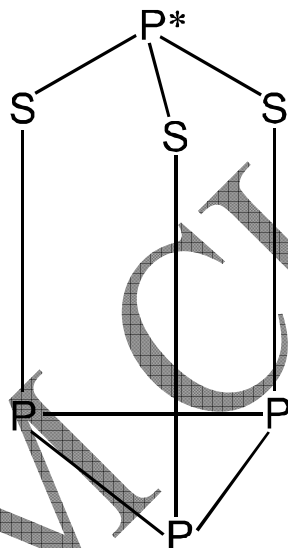




[A] ⇒ Non resolvable compound

[D] ⇒ It is a Aldehyde.

is formed.



36. (A), (B) P_4S_3

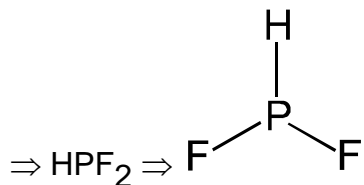
$$\Rightarrow I_p = 2$$

⇒ give 2-signal.

$$\Rightarrow {}^{31}\text{P NMR} = 2 \times 3 \times \frac{1}{2} + 1 = 4 \text{ peak}$$

$$^{31}\text{P NMR} = 2 \times 1 \times \frac{1}{2} + 1 = 2 \text{ peak}$$

So ^{31}P NMR give quadrat doublet signal & for $^{32}\text{S} = I = 0$ so does not give signal in NMR spectroscopy.



$$I_{\text{P}} = \frac{1}{2}, I_{\text{F}} = \frac{1}{2}, I_{\text{H}} = \frac{1}{2}.$$

$$^{31}\text{P NMR} = \left(2 \times 2 \times \frac{1}{2} + 1\right) \times \left(2 \times 1 + \frac{1}{2} + 1\right)$$

$$= 3 \times 2 = 6 \text{ peak}$$

$$^{19}\text{F-NMR} = \left(2 \times 1 \times \frac{1}{2} + 1\right) \times \left(2 \times 1 \times \frac{1}{2} + 1\right) = 4 \text{ peak}$$

$$^1\text{H-NMR give} = \left(2 \times 2 \times \frac{1}{2} + 1\right) \left(2 \times 1 \times \frac{1}{2} + 1\right) = 3 \times 2 = 6 \text{ peak}$$

37. (A, C, D) Oxyacids of bromine HOBr , HBrO_3 and HBrO_4

38. (A, D) Mol. mass of ethyl alcohol = $\text{C}_2\text{H}_5\text{OH} = 46$

$$\text{No. of moles of ethyl alcohol} = \frac{60}{46} = 1.304$$

$$\text{Mol. mass of methyl alcohol} = \text{CH}_3\text{OH} = 32$$

$$\text{No. of moles of methyl alcohol} = \frac{40}{32} = 1.25$$

$$'X_A' \text{ mole fraction of ethyl alcohol} = \frac{1.304}{1.304 + 1.25} = 0.5107$$

$$'X_B' \text{ mol fraction of methyl alcohol} = \frac{1.25}{1.304 + 1.25} = 0.4893$$

$$\text{Partial pressure of ethyl alcohol} = X_A P_A^{\circ} = 0.5107 \times 44.5 = 22.73 \text{ mm Hg}$$

$$\text{Partial pressure of methyl alcohol} = X_B P_B^{\circ} = 0.4893 \times 88.7 = 43.40 \text{ mm Hg}$$

$$\text{Total vapour pressure of solution} = 22.73 + 43.40 = 66.13 \text{ mm Hg}$$

$$\text{Mol. fraction of methyl alcohol in the vapour} = \frac{\text{Partial pressure of CH}_3\text{OH}}{\text{Total vapour pressure}}$$

$$= \frac{43.40}{66.13} = 0.6563$$

39. (A, B, C, D)

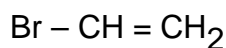
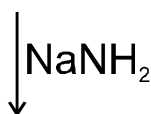
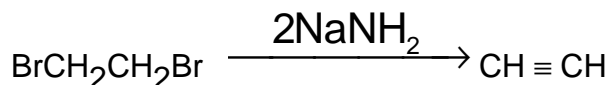
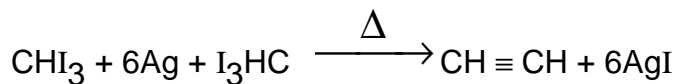
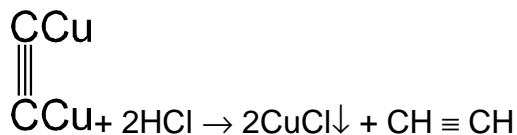
Potassium acid phthalate + Dipotassium phthalate \Rightarrow 4.0 – 6.2

$\text{Na}_2\text{HPO}_4 + \text{NaH}_2\text{PO}_4 \Rightarrow$ 5.9 – 8.0

Boric acid + Borax \Rightarrow 6.8 – 9.2

Borax + NaOH \Rightarrow 9.2 – 11.0

40. (A, C, D) $6\text{CH}_4 + \text{O}_2 \rightarrow 2\text{HC} \equiv \text{CH} + 2\text{CO} + 4\text{H}_2$



41. 14422.9

From the given data, if N_0 denotes original amount of ^{14}C , $t_{0.5} = 5577$ years; $N =$

$$\frac{N_0}{6}$$

From Eqn,

$$t_{0.5} = \frac{0.693}{\lambda} = 5577 \text{ years}$$

∴ The disintegration constant of ^{14}C ,

$$\lambda = \frac{0.693}{5577} \text{ year}^{-1}$$

Substituting the value of

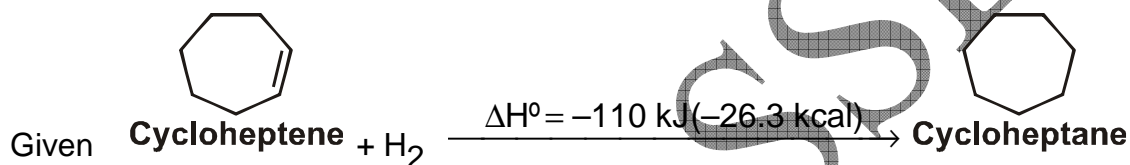
$$\lambda = \frac{0.693}{5577} \text{ and } N = \frac{N_0}{6} \text{ in Eqn, we get}$$

$$\lambda = \frac{2.303}{t} \log \frac{N_0}{N}$$

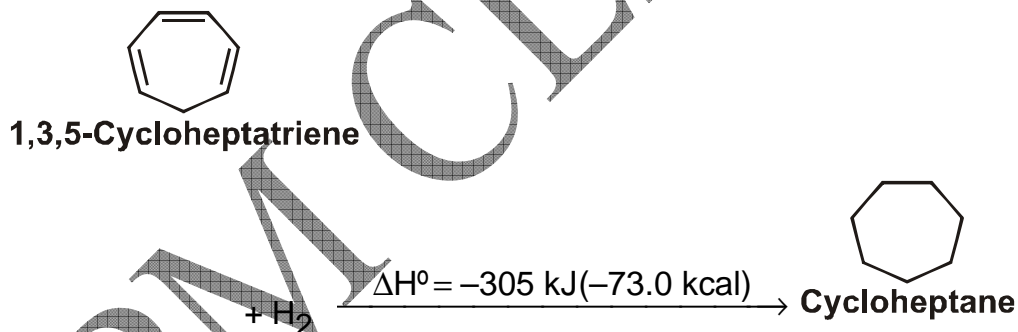
$$\frac{0.693}{5577} = \frac{2.303}{t} \log \frac{N_0}{N_0/6} = \frac{2.303}{t} \log 6 = \frac{2.303 \times 0.7782}{t}$$

$$\text{Thus, } t, \text{ the age of wood} = \frac{2.303 \times 0.7782 \times 5577}{0.693} = 14422.9 \text{ years.}$$

42. 25/5.9



and assuming that there is no resonance stabilization in 1,3,5-cycloheptatriene, we predict that its heat of hydrogenation will be three times that of cycloheptene or 330 kJ/mol (78.9 kcal/mol). The measured heat of hydrogenation is



Therefore

Resonance energy = 330 kJ/mol (predicted for no delocalization) – 305 kJ/mol (observed)

$$= 25 \text{ kJ/mol (5.9 kcal/mol)}$$

The value given in the text for the resonance energy of benzene (152 kJ/mol) is six times larger than this. 1,3,5-Cycloheptatriene is not aromatic.

43. 0.8263

Since the concentration cell with transference includes in it the liquid junction potential while the cell without transference does not, the liquid junction potential would be given by

$$E_l = E_{w.t.} - E_{w.o.t.} = 0.02802 - 0.01696 = 0.01106 \text{ V}$$

As during the operation of the cell, there is transfer of HCl from a solution of higher activity to one of lower activity, therefore we have

$$E_l = (t_- - t_+) \frac{RT}{F} \ln \frac{(a_{\pm})_2}{(a_{\pm})_1} = (t_- - t_+) \times 0.0591 \log \frac{0.009408}{0.01751}$$

$$0.01106 = (t_+ - t_-) \times (-0.016946) \quad \text{or} \quad t_+ - t_- = \frac{0.01106}{0.016946} =$$

0.6527

Since $t_+ + t_- = 1$, hence $t_+ = 0.8263$

Thus, the transference number of H^+ ion = **0.8263**

44. 27.08

Molar volume of liquid water, $V_l = 18 \text{ cm}^3 \text{ mol}^{-1} = 18 \times 10^{-6} \text{ m}^3 \text{ mol}^{-1}$

Molar volume of steam, $V_g = 18 \times 1673 \text{ cm}^3 \text{ mol}^{-1} = 30114 \times 10^{-6} \text{ m}^3 \text{ mol}^{-1}$

Heat of vaporisation, $\Delta H_v = 9700 \text{ cal mol}^{-1} \times 4.184 \text{ J cal}^{-1} = 40584.8 \text{ J mol}^{-1}$

$$\frac{dP}{dT} = \frac{\Delta H_v}{T(V_g - V_l)}$$

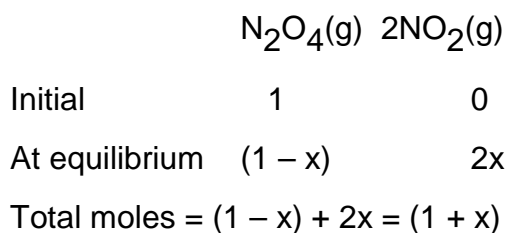
In this case, $dT = 1 \text{ K}$ and $T = 273 + 100 = 373 \text{ K}$

$$\therefore dP = \frac{40584.8 \text{ J mol}^{-1} \times 1 \text{ K}}{373 \text{ K}(30114 - 18) \times 10^{-6} \text{ m}^3 \text{ mol}^{-1}} = 0.00361 \times 10^6 \text{ N m}^{-2} \text{ (J = newton metre)}$$

$$= 0.03561 \text{ atm} = \mathbf{27.08 \text{ mm of Hg}} \quad (1 \text{ atm} = 101325 \text{ N m}^{-2})$$

Thus, the vapour pressure of water increases by about 27 mm of Hg by 1°C rise in temperature, at 100°C.

45. 63.2



$$p_{\text{N}_2\text{O}_4} = \left(\frac{1-x}{1+x}\right)P, p_{\text{NO}_2} = \frac{2x}{(1+x)}P$$

Given, $x = 0.25$ and $P = 1 \text{ atm}$.

$$p_{\text{N}_2\text{O}_4} = \left(\frac{1-0.25}{1+0.25}\right) \times 1 = 0.6 \text{ atm}$$

$$p_{\text{NO}_2} = \left(\frac{2 \times 0.25}{1+0.25}\right) \times 1 = 0.4 \text{ atm}$$

$$K_p = \frac{(p_{\text{NO}_2})^2}{p_{\text{N}_2\text{O}_4}} = \frac{0.4 \times 0.4}{0.6} = 0.267 \text{ atm}$$

Let the degree of dissociation of N_2O_4 at 0.1 atm be ' α ', then

$$p_{\text{N}_2\text{O}_4} = \left(\frac{1-\alpha}{1+\alpha}\right) \times 0.1 \quad \text{and} \quad p_{\text{NO}_2} = \frac{2\alpha}{(1+\alpha)} \times 0.1$$

$$K_p = \frac{0.4\alpha^2}{(1-\alpha^2)} \text{ or } 0.267 = \frac{\left(\frac{2\alpha}{1+\alpha}\right)^2 \times (0.1)^2}{\left(\frac{1-\alpha}{1+\alpha}\right) \times 0.1} = \frac{4\alpha^2 \times 0.1}{(1-\alpha)(1+\alpha)} = \frac{0.4\alpha^2}{(1-\alpha^2)}$$

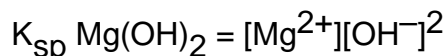
$$\text{or } 0.267 = 667\alpha^2$$

$$\alpha = 0.632$$

Hence dissociation of $N_2O_4 = 63.2\%$.

46. 9.04

Given that



$$1.2 \times 10^{-11} = [0.1][\text{OH}^-]^2$$

$$[\text{OH}^-]^2 = 1.2 \times 10^{-10}$$

$$[\text{OH}^-] = 1.095 \times 10^{-5} \text{ M}$$

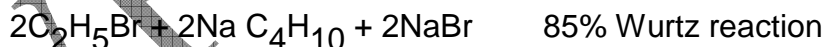
$$\text{pOH} = -\log_{10}(1.095 \times 10^{-5}) = 4.96$$

$$\text{pH} = 14 - 4.96 = 9.04$$

Thus at 9.04 pH, precipitation will not take place.

47. 55.50

The production of n-butane may be represented by the following equation.

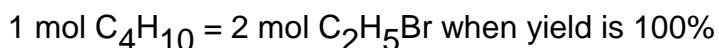


Mass of n-butane to be produced = 55 gm

Molecular mass of $C_4H_{10} = 4 \times 12 + 10 \times 1 = 58 \text{ gm/mol}$.

$$\text{Mole of } C_4H_{10} \text{ in } 55\text{gm} = \frac{55\text{gm}}{58\text{gm/mol}} = 0.948\text{mol.}$$

From wurtz reaction, we find that

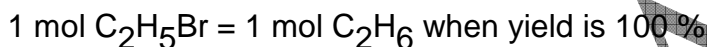


$$0.948 \text{ mol } C_4H_{10} = 2 \times 0.948 \times \frac{100}{85} \text{ mol } C_2H_5Br$$

(when yield is 85%)

$$= 2.23 \text{ mol } C_2H_5Br$$

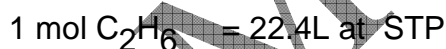
For bromination reaction of 90% yield it is found that



$$= 1 \times \frac{100}{90} \text{ mol } C_2H_6 \text{ when yield is } 90\%$$

$$2.23 \text{ mol } C_2H_5Br = 2.23 \times \frac{100}{90} \text{ mol } C_2H_6$$

$$= 2.478 \text{ mol } C_2H_6$$



$$2.478 \text{ mol } C_2H_6 = 2.478 \times 22.4\text{L} = 55.50 \text{ L}$$

48. 27419.2

$$\text{Wave number} = \frac{1}{\lambda} = \bar{\nu} = R_H \left[\frac{1}{n_1^2} - \frac{1}{n_2^2} \right]$$

$$R_H \Rightarrow \text{Rydberg constant} = 109677 \text{ cm}^{-1}$$

$$n_1 \Rightarrow 2 \text{ for balmer series}$$

$$n_2 \Rightarrow \infty \text{ for shortest wavelength}$$

$$\bar{V} = 109677 \text{ cm}^{-1} \left[\frac{1}{(2)^2} - \frac{1}{(\infty)^2} \right]$$

$$\bar{V} = 27419.2 \text{ cm}^{-1}$$

49. 494.5

$$\text{Energy of electromagnetic radiations } E = hv = \frac{hc}{\lambda}$$

$$h = 6.6 \times 10^{-34} \text{ JS}$$

$$c = 3 \times 10^8 \text{ m/s}$$

$$\lambda = 242 \text{ nm} = 242 \times 10^{-9} \text{ m}$$

$$E = \frac{hc}{\lambda} = \frac{6.6 \times 10^{-34} \text{ Jsec} \times 3 \times 10^8 \text{ m/sec}}{242 \times 10^{-9} \text{ m}}$$

$$= 8.21 \times 10^{-19} \text{ J}$$

Energy required to ionise one mole of Na atom = Energy needed to ionise one Na atom x Avogadro No.

$$= E \times N_A$$

$$= 8.21 \times 10^{-19} \text{ J} \times 6.023 \times 10^{23} \text{ mol}^{-1}$$

$$= 4.945 \times 10^5 \text{ J mol}^{-1}$$

$$\text{I.E. of Na} = 494.5 \text{ KJ mol}^{-1}$$

50. 15.05

The Formula of ore is $\text{Fe}_{0.93}\text{O}_{1.00}$

Number of O atom per 100 unit of wustite = 100

Number of -ive charge due to O atom = 200

Number of Fe atom per 100 unit of wustite = 93

Number of +ive charge due to 93 Fe atom = 200

Total No. of Fe^{+2} ion = X

Total No. of Fe^{+3} ion = y

$$x + y = 93 \text{ \&}$$

$$\& \quad x = 93 - y.$$

$$x \text{Fe}^{+2} + y \text{Fe}^{+3} = 200$$

$$(93 - y) \text{Fe}^{+2} + y \text{Fe}^{+3} = 200$$

$$93 \text{Fe}^{+2} + (y \text{Fe}^{+3} - y \text{Fe}^{+2}) = 200$$

$$93 \times 2 + (3 - 2)y = 200$$

$$186 + y = 200$$

$$y = 200 - 186$$

$$y = 14 = \text{No of } \text{Fe}^{3+} \text{ ion.}$$

$$\% \text{ of } \text{Fe}^{3+} = \frac{14}{93} \times 100$$

$$= 15.05\%$$

51. 4

Identification of complex A

Complex A react with AgNO_3 to give a white precipitate which is readily soluble in dilute aqueous ammonia. This implies that complex A contains ionizable Cl^- group.

Thus

Formula of A is $[\text{Cr}(\text{NH}_3)_4 \text{ClBr}]\text{Cl}$

Identification of complex B.

Complex B react with AgNO_3 to give a pale-yellow precipitate which is soluble in concentrated ammonia. This implies that B contain ionizable Br^- group. Thus

Formula of B is $[\text{Cr}(\text{NH}_3)_4 \text{Cl}_2] \text{Br}$

In Complex A & B

Cr present in +3 oxidation state

So



d^2sp^3 hybridization

& 3 unpaired e^- are present

So magnetic moment $\mu = \sqrt{n(n+2)}$
 $= 3.87 \times 4 \text{ BM}$

52. 2

For the ionisation of indicator $\text{HIn} \rightleftharpoons \text{H}^+ + \text{In}^-$ the ionisation constant (K_a) is given by

$$K_a = \frac{[\text{H}^+][\text{In}^-]}{[\text{HI}]}$$

$$[H^+] = K_a \frac{[HI]}{[In^-]} = \frac{K_a}{[In^-]/[HI]}$$

The colour of In^- is visible when $[In^-]/[HI] > 10$.

Thus

$$[H^+] = \frac{K_a}{[In^-]/[HI]} = \frac{1.0 \times 10^{-5}}{10} = 10^{-6}$$

$$pH = -\log[H^+] = -\log 10^{-6} = 6$$

The colour of HI is visible when $[HI]/[In^-] > 10$ or

$$[In^-]/[HI] < 0.1$$

Thus

$$[H^+] = \frac{K_a}{[In^-]/[HI]} = \frac{1.0 \times 10^{-5}}{0.1} = \frac{10^{-5}}{10^{-1}} = 10^{-4}$$

$$pH = -\log [H^+] = -\log 10^{-4} = 4$$

$$\text{change of pH} = 6 - 4 = 2$$

53. 1611.6

$$\log \left(\frac{k}{A} \right) = \frac{-E_a}{2.303RT}$$

$$= \frac{-210000 \text{ J mol}^{-1}}{2.303 \times 8.314 \text{ J mol}^{-1} \text{ K}^{-1} \times 700 \text{ K}}$$

$$= -15.668$$

$$K/A = \text{Antilog} (-15.668) = 2.1478 \times 10^{-16}$$

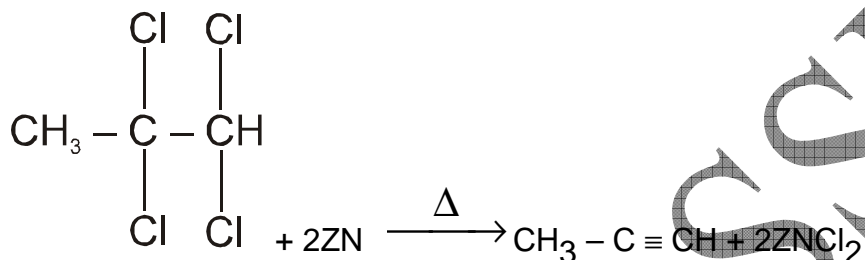
$$k = 1.1478 \times 10^{-16} \text{ A}$$

$$= 2.1478 \times 10^{-16} \times 2 \times 10^{12} \times 5^1 = 4.3 \times 10^{-4} \text{ sec}^{-1}$$

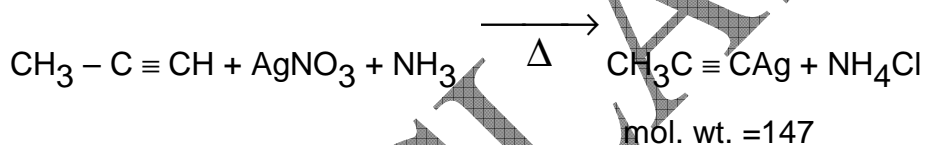
$$L_{\frac{1}{2}} = \frac{0.693}{4.3 \times 10^{-4}} = 1611.6 \text{ sec.}$$

54. 29.4

The dehalogenation of vicinal tetrahaloalkane gives alkyne .



mol. wt. = 182



182 gm of tetrahaloalkanes gives 147 gm ppt.

$$\frac{147 \times 36.4}{182}$$

36.4 gm of tetrahaloalkane gives =

$$182$$

$$= 29.4 \text{ gm ppt}$$

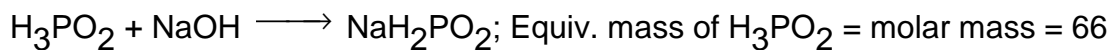
weight of silver propynide = 29.4 gm

55. 0

Angular nodes = $l = 3$

Radial nodes = $n - l - 1 = 4 - 3 - 1 = 0$

56. 100



$$\text{Equivalents of H}_3\text{PO}_2 = \frac{0.66}{66} = 0.01 = x \times 0.1 \Rightarrow x = 0.1 \text{ L} = 100 \text{ ml}$$

57. 4

The number of dipeptides that can be prepared from alanine and phenylalanine is 4 — Ala-phe, Ala-Ala, phe-phe, phe-Ala.

58. 0.25

$$I = \frac{1}{2} [m_1 z_1^2 + m_2 z_2^2 + m_3 z_3^2]$$

$$m_1(\text{Na}^+) = 0.1 + 2 \times 0.05 = 0.2$$

$$m_2(\text{Cl}^-) = 0.1$$

$$m_3(\text{SO}_4^{-2}) = 0.05$$

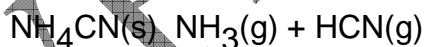
$$z_1 = 1; z_2 = 1; z_3 = 2$$

$$I = \frac{1}{2} [0.1 \times (1)^2 + 0.2 \times (1)^2 + 0.05 \times (2)^2] = \frac{1}{2} [0.5] = 0.25$$

59. 0.463

$$E = E^0 - \frac{0.0592}{2} \log \frac{(\text{QH})_2}{(\text{Q})(\text{H}^+)_2} = E^0 - 0.0592 \text{ pH} = 0.699 - 0.0592 \times 4 = 0.463 \text{ V}$$

60. 0.0689



At equ. $p + 0.25$ p

$$K_p = \frac{p_{(\text{NH}_3)} p_{(\text{HCN})}}{p_{(\text{NH}_4\text{CN})}}$$

$$0.0222 = (p + 0.25)p$$

$$p^2 + 0.25p - 0.0222 = 0$$

$$p = \frac{-0.25 \pm \sqrt{(0.25)^2 + (0.0222) \times 4 \times 1}}{2 \times 1} = 0.0689 \text{ atm}$$

VPM CLASSES