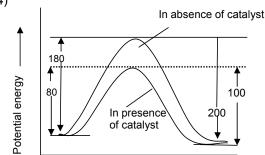
- 81. The energies of activation for forward and reverse reactions for  $A_2 + B_2 \rightleftharpoons 2AB$  are 180 kJ mol<sup>-1</sup> and 200 kJ mol<sup>-1</sup> respectively. The presence of catalyst lowers the activation energy of both (forward and reverse) reactions by 100 kJ mol<sup>-1</sup>. The enthalpy change of the reaction  $(A_2 + B_2 \Longrightarrow 2AB)$  in the presence of catalyst will be (in kJ mol<sup>-1</sup>)
  - (1) 300
  - (3) 280

(2) 120(4) 20

Ans. (4) Sol.



Reaction coordinate\_\_\_\_\_

So, 
$$\Delta H_{Reaction} = E_f - E_b$$
  
= 80 - 100 = -20  
Hence, (4) is correct.

- 82. The cell,  $Zn | Zn^{2+} (1M) | Cu^{2+} (1M) | Cu(E_{cell}^0 = 1.10V)$ , was allowed to be completely discharged at 298
  - K. The relative concentration of  $Zn^{2^+}$  to  $Cu^{2^+}\begin{bmatrix} \boxed{Zn^{2^+}}\\ \boxed{Cu^{2^+}} \end{bmatrix}$  is
  - (1) antilog (24.08)
  - $(3) 10^{37.3}$

- (2) 37.3
- (4) 9.65 × 10<sup>4</sup>

- Ans. (3
- Sol.  $E_{cell} = E_{cell}^o \frac{0.0591}{n} log Q$

Where Q = 
$$\frac{\left[Zn^{2+}\right]}{\left\lceil Cu^{2+}\right\rceil}$$

For complete discharge  $E_{cell} = 0$ 

So 
$$E_{cell}^o = \frac{0.591}{2} log \frac{\left[Zn^{2+}\right]}{\left[Cu^{2+}\right]}$$

$$\Rightarrow \left| \frac{\left[ Z n^{2+} \right]}{\left[ C u^{2+} \right]} \right| = 10^{37.3}$$

Hence, (3) is correct.

- 83. The pKa of a weak acid (HA) is 4.5. The pOH of an aqueous buffered solution of HA in which 50% of the acid is ionized is
  - (1) 4.5

(2) 2.5

(3) 9.5

(4) 7.0

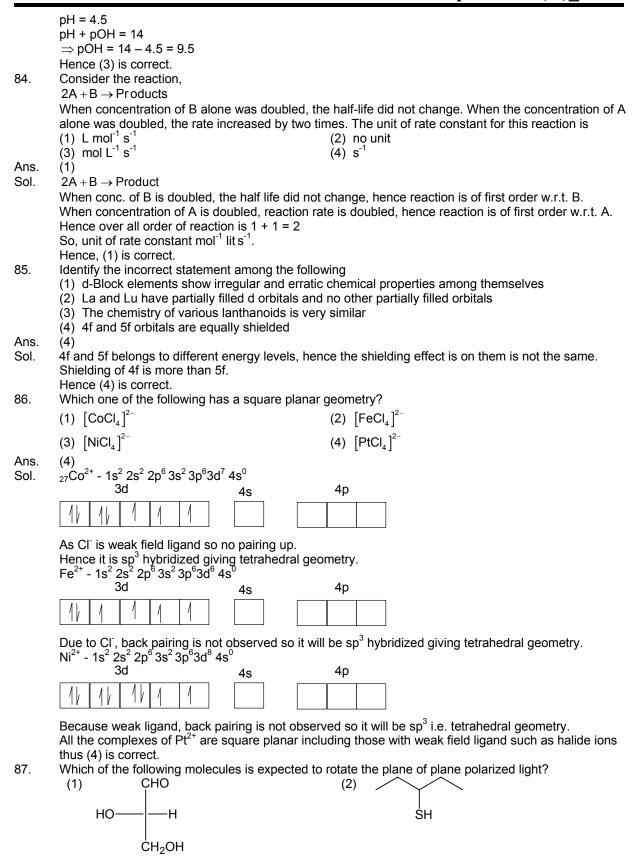
Ans. (3)

Sol. For buffer solution

$$pH = pK_a + log \frac{[Salt]}{[Acid]}$$

= 
$$4.5 + \log \frac{\left[\text{Salt}\right]}{\left[\text{Acid}\right]}$$

as HA is 50% ionized so [Salt] = [Acid]



(3) 
$$H_2N$$
  $NH_2$  (4) COOH  $H_2N$   $H_2N$   $H_2N$   $H_2N$ 

Ans. (1)

Sol. The plane of polarized light is rotated by optically active compound, i.e. it should be chiral. So, (1) has, chiral C-atom. So, it is optically active. In (2), (3) and (4) plane of symmetry is present. Hence, (1) is correct.

88. The secondary structure of a protein refers to

(1) α-helical backbone

- (2) hydrophobic interactions
- (3) sequence of  $\alpha$ -amino acids
- (4) fixed configuration of the polypeptide backbone

Ans. (1)

- Sol. Secondary structure of proteins involves  $\alpha$  helical back bond and  $\beta$  sheet structures. These structures are formed as a result of H-bonding between different peptide groups. Hence, (1) is correct
- 89. Which of the following reactions will yield 2, 2-dibromopropane?
  - (1)  $CH_3 C \equiv CH + 2HBr \longrightarrow$
- (2)  $CH_3CH \equiv CHBr + HBr \longrightarrow$

(3)  $CH \equiv CH + 2HBr \longrightarrow$ 

(4)  $CH_3 - CH = CH_2 + HBr \longrightarrow$ 

Ans. (1)

Sol. 
$$CH_3 - C \equiv CH + HBr \xrightarrow{\text{electrophilic addition of H}^+} CH_3 - C = CH_2 \xrightarrow{\text{HBr}} CH_3 - C = CH_3$$

Hence, (1) is correct.

90. In the chemical reaction,

 $CH_3CH_2NH_2 - CHCl_3 + 3KOH \longrightarrow (A) + (B) + 3H_2O$ , the compound (A) and (B) are respectively

(1) C<sub>2</sub>H<sub>5</sub>CN and 3KCl

(2) CH<sub>3</sub>CH<sub>2</sub>CONH<sub>2</sub> and 3KCl

(3)  $C_2H_5NC$  and  $K_2CO_3$ 

(4) C<sub>2</sub>H<sub>5</sub>NC and 3KCl

Ans. (4)

- Sol. It is example of carbylamine reaction. so, the product will be C<sub>2</sub>H<sub>5</sub>NC and KCl. Hence, (4) is the correct answer.
- 91. The reaction of toluene with Cl<sub>2</sub> in presence of FeCl<sub>3</sub> gives predominantly
  - (1) benzoyl chloride

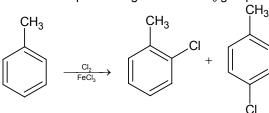
(2) benzyl chloride

(3) o-and p-chlorotoluene

(4) m-chlorotoluene

Ans. (3)

Sol. Due to o- and p- directing nature of CH<sub>3</sub> group.



Hence, (3) is correct answer.

- 92. Presence of a nitro group in a benzene ring
  - (1) activates the ring towards electrophilic substitution
  - (2) renders the ring basic
  - (3) deactivates the ring towards nucleophilic substitution
  - (4) deactivates the ring towards electrophilic substitution

Ans. (4)

Sol. - NO<sub>2</sub> group shows - M effect, so withdraws the electron density from the ring and hence deactivate the ring towards electrophilic aromatic substitution.
 Hence, (4) is correct.

- 93. In which of the following ionization processes, the bond order has increased and the magnetic behaviour has changed?
  - $(1) \quad C_2 \longrightarrow C_2^+$

(2)  $NO \longrightarrow NO^+$ 

 $(3) O_2 \longrightarrow O_2^+$ 

 $(4) N_2 \longrightarrow N_2^+$ 

Ans. (2

Sol. In C<sub>2</sub> – C<sub>2</sub><sup>+</sup> electron is removed from bonding molecular orbital so bond order decreases. In NO

NO<sup>+</sup>, electron is removed from anti bonding molecular orbital so bond order increases and nature changes from paramagnetic to diamagnetic.

Hence, (2) is correct.

- 94. The actinoids exhibits more number of oxidation states in general than the lanthanoids. This is because
  - (1) the 5f orbitals are more buried than the 4f orbitals
  - (2) there is a similarity between 4f and 5f orbitals in their angular part of the wave function
  - (3) the actinoids are more reactive than the lanthanoids
  - (4) the 5f orbitals extend further from the nucleus than the 4f orbitals

Ans. (4)

Sol. The actinoids exhibit more number of oxidation states in general than the lanthanoids. This is because the 5f orbitals extend further from the nucleus than the 4f orbitals.

Hence, (4) is correct.

- 95. Equal masses of methane and oxygen are mixed in an empty container at 25°C. The fraction of the total pressure exerted by oxygen is
  - (1)  $\frac{2}{3}$

(2)  $\frac{1}{3} \times \frac{273}{298}$ 

(3)  $\frac{1}{3}$ 

(4)  $\frac{1}{2}$ 

Ans. (3)

Sol. Let the mass of methane and oxygen is w

mole fraction of oxygen = 
$$\frac{\frac{W}{32}}{\frac{W}{32} + \frac{W}{16}}$$

$$=\frac{\frac{1}{32}}{\frac{1}{32} + \frac{1}{16}} = \frac{\frac{1}{32}}{\frac{3}{32}} = \frac{1}{3}$$

Let the total pressure be P

The pressure exerted by oxygen (partial pressure) =  $X_{O_2} \times P_{total}$ 

$$\Rightarrow P \times \frac{1}{3}$$

Hence, (3) is correct.

- 96. A 5.25 % solution of a substance is isotonic with a 1.5% solution of urea (molar mass = 60 g mol<sup>-1</sup>) in the same solvent. If the densities of both the solutions are assumed to be equal to 1.0 g cm<sup>-3</sup>, molar mass of the substance will be
  - (1) 90.0 g mol<sup>-1</sup>

(2)  $115.0 \text{ g mol}^{-1}$ 

(3)  $105.0 \text{ g mol}^{-1}$ 

(4)  $210.0 \text{ g mol}^{-1}$ 

Ans. (4)

Sol. Solutions with the same osmotic pressure are isotonic

Let the molar mass of the substance be M

$$\pi_1 = C_1RT = C_2RT = \pi_2$$

So, 
$$C_1 = C_2$$

As density of the solutions are same

So 
$$\frac{5.25}{M} = \frac{15}{60}$$

$$M = \frac{5.25 \times 60}{1.5} = 210$$

Hence (4) is correct

- 97. Assuming that water vapour is an ideal gas, the internal energy ( $\Delta U$ ) when 1 mol of water is vapourised at 1 bar pressure and 100°C, (Given: Molar enthalpy of vapourization of water at 1 bar and 373 K = 41 kJ mol<sup>-1</sup> and R = 8.3 J mol<sup>-1</sup>K<sup>-1</sup>) will be
  - (1) 4.100 kJ mol<sup>-1</sup>

(2) 3.7904 kJ mol<sup>-1</sup>

(3) 37.904 kJ mol<sup>-1</sup>

(4) 41. 00 kJ mol<sup>-1</sup>

Ans. (3)

Sol.  $H_2O(\ell) \xrightarrow{\text{vaporisation}} H_2O(g)$ 

$$\Delta n_{\alpha} = 1 - 0 = 1$$

$$\Delta H = \Delta U + \Delta n_{\alpha}RT$$

$$\Delta U = \Delta H - \Delta n_{g}RT$$

$$= 41 - 8.3 \times 10^{-3} \times 373$$

Hence, (3) is correct.

98. In a sautrated solution of the sparingly soluble strong electrolyte AgIO<sub>3</sub> (Molecular mass = 283) the equilibrium which sets in is

$$AgIO_{3(s)} \longrightarrow Ag^{+}_{(aq)} + IO^{-}_{3(aq)}$$

If the solubility product constant  $K_{sp}$  of  $AgIO_3$  at a given temperature is  $1.0 \times 10^{-8}$ , what is the mass of  $AgIO_3$  contained in 100 ml of its saturated solution?

(1) 
$$28.3 \times 10^{-2}$$
 g

(2) 
$$2.83 \times 10^{-3}$$
 g

(3) 
$$1.0 \times 10^{-7}$$
 g

(4) 
$$1.0 \times 10^{-4}$$
 g

Ans. (2)

Sol. 
$$AgIO_3(s) \rightleftharpoons Ag^+(aq) + IO_3^-(aq)$$

Let the solubility of AgIO<sub>3</sub> be s

$$K_{sp} = \lceil Ag^+ \rceil \lceil IO_3^- \rceil$$

$$1.0 \times 10^{-8} = s^2$$

$$s = 10^{-4} \text{ mol/litre}$$

$$= \frac{10^{-4} \times 283}{1000} \times 100$$

$$= 283 \times 10^{-5}$$

$$= 2.83 \times 10^{-3} \text{ g} / 100 \text{ ml}$$

Hence, (2) is correct.

- 99. A radioactive element gets spilled over the floor of a room. Its half-life period is 30 days. If the initial activity is ten times the permissible value, after how many days will it be safe to enter the room?
  - (1) 1000 days

(2) 300 days

(3) 10 days

(4) 100 days

Ans. (4

Sol. Activity 
$$\left(-\frac{dN}{dt}\right) \propto N$$

$$N = N_o \left(\frac{1}{2}\right)^n$$

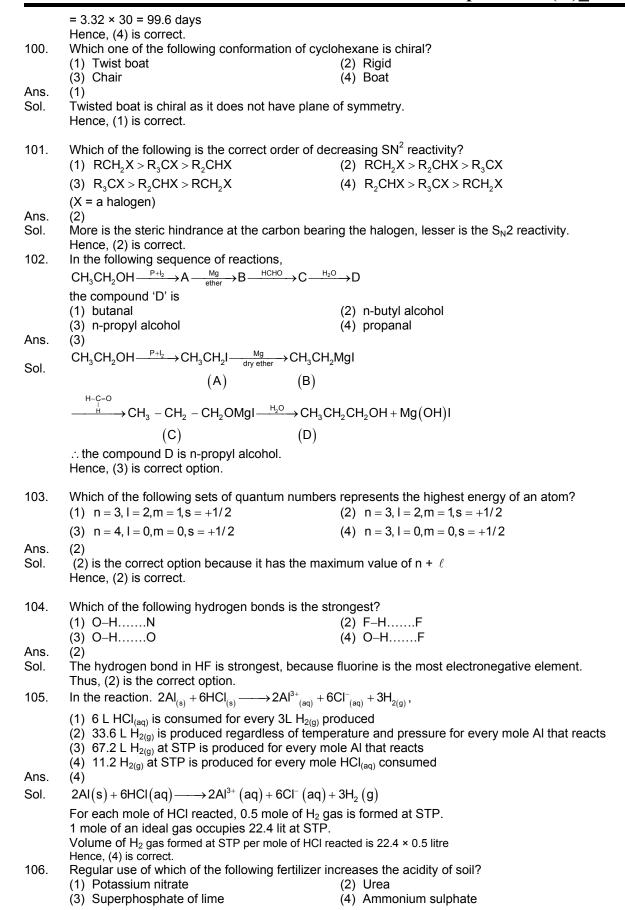
$$\frac{N}{N_0} = \left(\frac{1}{2}\right)^n$$

$$\frac{1}{10} = \left(\frac{1}{2}\right)^n \Rightarrow 10 = 2^n$$

$$log10 = nlog2$$

$$\Rightarrow n = \frac{1}{0.301} = 3.32$$

$$t = n \times t_{112}$$





Sol.  $(NH_4)_2$  SO<sub>4</sub> is a salt of strong acid and weak base, on hydrolysis it ill produce H<sup>+</sup> ion. This will increase the acidity of soil.

$$(NH_4)_2 SO_4 \longrightarrow 2NH_4^+ + SO_4^{2-}$$

$$NH_4^+ + H_2O \Longrightarrow NH_4OH + H^+$$

Hence, (4) is correct answer.

- 107. Identify the correct statement regarding a spontaneous process
  - (1) For a spontaneous process in an isolated system, the change in entropy is positive
  - (2) Endothermic processes are never spontaneous
  - (3) Exothermic processes are always spontaneous
  - (4) Lowering of energy in the reaction process is the only criterion for spontaneity

Ans. (1)

Sol. For a spontaneous process in an isolated system, the change in entropy is positive.

Hence, (1) is correct.

- 108. Which of the following nuclear reactions will generate an isotope?
  - (1) neutron particle emission
- (2) positron emission

(3)  $\alpha$ -particle emission

(4) β-particle emission

Ans. (1)

Sol. 
$${}^{A}_{Z}X \longrightarrow {}^{A-1}_{Z}X + {}^{1}_{0}n$$

Hence, (1) is correct.

109. The equivalent conductances of two strong electrolytes at infinite dilution in H<sub>2</sub>O (where ions move freely through a solution) at 25°C are given below:

$$^{\circ}_{CH_3COONa} = 91.0 \text{ S cm}^2 / \text{equiv}$$

$$^{\circ}_{HCI} = 426.2 \text{ S cm}^2 / \text{equiv}$$

What additional information/quantity one needs to calculate \( \sigma^\circ\) of an aqueous solution of acetic acid?

- (1) ∧° of NaCl
- (2) ∧° of CH<sub>3</sub>COOK
- (3) The limiting equivalent conductance of  $H^+(\wedge^{\circ}_{H^+})$
- (4) ∧° of chloroacetic acid (C/CH<sub>2</sub>COOH)

Ans. (1)

Sol. From Kohlrausch's law

$$\Lambda_{\text{CH}_3\text{COOH}}^{\circ} = \Lambda_{\text{CH}_3\text{COONa}}^{\circ} + \Lambda_{\text{HCI}}^{\circ} - \Lambda_{\text{NaCI}}^{\circ}$$

Hence, (1) is the correct answer.

- 110. Which one of the following is the strongest base in aqueous solution?
  - (1) Trimethylamine

(2) Aniline

(3) Dimethylamine

(4) Methylamine

Ans. (3

Sol. In aqueous solution basicity order of  $1^{\circ}$ ,  $2^{\circ}$  and  $3^{\circ}$  amine with methyl group is  $2^{\circ} > 1^{\circ} > 3^{\circ}$ 

In case of aniline lone pair of nitrogen is involved in resonance, so it is weaker base than aliphatic amines.

Hence, (3) is correct.

- 111. The compound formed as a result of oxidation of ethyl benzene by KMnO₄ is
  - (1) benzophenone

(2) acetophenone

(3) benzoic acid

(4) benzyl alcohol

Ans. (3)

Sol. Any aliphatic carbon with hydrogen attached to it, in combination with benzene ring, will be oxidized to benzoic acid by KMnO<sub>4</sub>/H<sup>+</sup>.

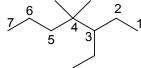
Hence, (3) is correct.

#### 112. The IUPAC name of

- (1) 1, 1-diethyl-2,2-dimethylpentane
- (3) 5, 5-diethyl-4, 4-diemthylpentane
- (2) 4, 4-dimethyl-5, 5-diethylpentane(4) 3-ethyl-4, 4-dimethylheptane

#### Ans. (4)

Sol.



The correct answer is 3-ethyl-4, 4-dimethylheptane.

Hence, (4) is correct.

- 113. Which of the following species exhibits the diamagnetic behaviour?
  - (1)  $O_2^{2-}$

(2)  $O_2^+$ 

(3)  $O_2$ 

(4) NO

is

Ans. (1)

Sol. The correct option is  $O_2^{2-}$ 

This species has 18 e<sup>-</sup>, which are filled in such a way that all molecular orbitals are fully filled, so diamagnetic.

$$\sigma 1s^2\sigma^* 1s^2, \sigma 2s^2\sigma^* 2s^2, \sigma 2p_z^2, \ \pi 2p_x^2 = \pi 2p_y^2, \ \pi^* 2p_x^2 = \pi^* 2p_y^2$$

Hence, (1) is correct.

- 114. The stability of dihalides of Si, Ge, Sn and Pb increases steadily in the sequence
  - (1)  $GeX_2 \ll SiX_2 \ll SnX_2 \ll PbX_2$
- (2)  $SiX_2 \ll GeX_2 \ll PbX_2 \ll SnX_2$
- (3)  $SiX_2 \ll GeX_2 \ll SnX_2 \ll PbX_2$
- (4)  $PbX_2 \ll SnX_2 \ll GeX_2 \ll SiX_2$

Ans. (3)

Sol. Due to inert pair effect, the stability of +2 oxidation state increases as we move down this group.

$$\therefore \mathsf{SiX}_2 \ll \mathsf{GeX}_2 \ll \mathsf{SnX}_2 \ll \mathsf{PbX}_2$$

Hence, (3) is correct.

- 115. Identify the incorrect statement among the following
  - (1) Ozone reacts with SO<sub>2</sub> to give SO<sub>3</sub>
  - (2) Silicon reacts with NaOH<sub>(aq)</sub> in the presence of air to give Na<sub>2</sub>SiO<sub>3</sub> and H<sub>2</sub>O
  - (3) Cl<sub>2</sub> reacts with excess of NH<sub>3</sub> to give N<sub>2</sub> and HCl
  - (4) Br<sub>2</sub> reacts with hot and strong NaOH solution to give NaBr, NaBrO<sub>4</sub> and H<sub>2</sub>O

Ans. (4)

Sol. Br<sub>2</sub> reacts with hot and strong NaOH to give NaBr, NaBrO<sub>3</sub> and H<sub>2</sub>O.

Hence, (4) is incorrect statement.

- 116. The charge/size ratio of a cation determines its polarizing power. Which one of the following sequences represents the increasing order of the polarizing order of the polarizing power of the cationic species, K<sup>+</sup>, Ca<sup>2+</sup>, Mg<sup>2+</sup>, Be<sup>2+</sup>?
  - (1) Mg<sup>2+</sup>,Be<sup>2+</sup>,K<sup>+</sup>,Ca<sup>2+</sup>

(2) Be<sup>2+</sup>,K<sup>+</sup>,Ca<sup>2+</sup>,Mg<sup>2+</sup>

(3)  $K^+, Ca^{2+}, Mg^{2+}, Be^{2+}$ 

(4)  $Ca^{2+}$ ,  $Mg^{2+}$ ,  $Be^{2+}$ ,  $K^{+}$ 

Ans. (3)

Sol. Higher the charge/size ratio, more is the polarizing power.

$$\therefore K^{+} < Ca^{2+} < Mq^{2+} < Be^{2+}$$

Hence, (3) is correct.

- 117. The density (in g mL<sup>-1</sup>) of a 3.60 M sulphuric acid solution that is 29%  $H_2SO_4$  (Molar mass = 98 g mol<sup>-1</sup>) by mass will be
  - (1) 1.64

(2) 1.88

(3) 1.22

(4) 1.45

Ans. (3)

Sol. Let the density of solution be 'd'

Molarity of solution given = 3.6

- i.e. 1 litre of solution contains 3.6 moles of H<sub>2</sub>SO<sub>4</sub>
- or 1 litre of solution contains 3.6 × 98 gms of H<sub>2</sub>SO<sub>4</sub>

Since, the solution is 29% by mass.

100 gm solution contains 29 gm H<sub>2</sub>SO<sub>4</sub>

 $\frac{100}{d}$  ml solution contains 29 gm of H<sub>2</sub>SO<sub>4</sub>

1000 ml solution contains 3.6 × 98 gm of H<sub>2</sub>SO<sub>4</sub>

$$\therefore 3.6 \times 98 = \frac{29 \times d}{100} \times 1000$$

d = 1.22

Hence, (3) is correct.

The first and second dissociation constants of an acid  $H_2A$  are  $1.0 \times 10^{-5}$  and  $5.0 \times 10^{-10}$  respectively. 118. The overall dissociation constant of the acid will be

(1) 
$$5.0 \times 10^{-5}$$

(2)  $5.0 \times 10^{15}$ 

$$(3)$$
 5.0 × 10<sup>-15</sup>

 $(4) 0.0 \times 10^5$ 

Ans. (3)

Sol. 
$$H_2A \Longrightarrow HA^- + H^-$$

$$H_2A \rightleftharpoons HA^- + H^+ \qquad K_1 = \frac{\left[HA^-\right]\left[H^+\right]}{\left[H_2A\right]}$$

$$HA^- \rightleftharpoons H^+ + A^{2-}$$

$$\mathsf{H}\mathsf{A}^- \Longrightarrow \mathsf{H}^+ + \mathsf{A}^{2-} \qquad \mathsf{K}_2 = \frac{\left[\mathsf{H}^+\right]\left[\mathsf{A}^{2-}\right]}{\left[\mathsf{H}\mathsf{A}^-\right]}$$

For the reaction

$$H_2A \Longrightarrow 2H^+ + A^{2-}$$

$$K = \frac{\left[H^{+}\right]^{2} \left[A^{2-}\right]}{\left[H_{2}A\right]} = K_{1} \times K_{2}$$

$$= 1 \times 10^{-5} \times 5 \times 10^{-10}$$
$$= 5 \times 10^{-15}$$

Hence, (3) is correct.

119. A mixture of ethyl alcohol and propyl alcohol has a vapour pressure of 290 mm at 300 K. The vapour pressure of propyl alcohol is 200 mm. If the mole fraction of ethyl alcohol is 0.6, its vapour pressure (in mm) at the same temperature will be

(1) 350

(2) 300

(3) 700

(4) 360

(1) Ans.

Let the vapour pressure of pure ethyl alcohol be P, Sol.

According to Raoult's law

$$290 = 200 \times 0.4 + P \times 0.6$$

$$P = \frac{290 - 80}{0.6} = 350 \text{ mm Hg}$$

Hence, (1) is correct.

120. In conversion of lime-stone to lime,

$$CaCO_3(s) \longrightarrow CaO(s) + CO_2(g)$$

the vales of  $\Delta H^{\circ}$  and  $\Delta S^{\circ}$  are +179.1 kJ mol<sup>-1</sup> and 160.2 J/K respectively at 298 K and 1 bar. Assuming that  $\Delta H^{\circ}$  do not change with temperature, temperature above which conversion of limestone to lime will be spontaneous is

(1) 1008 K

(2) 1200

(3) 845 K

(4) 1118 K

Ans. (4)

We know,  $\Delta G = \Delta H - T\Delta S$ Sol.

So, lets find the equilibrium temperature, i.e. at which

$$\Delta G = 0$$

$$\Delta H = T\Delta S$$

$$T = \frac{179.1 \times 1000}{160.2}$$

So, at temperature above this, the reaction will become spontaneous.

Hence, (4) is correct answer.