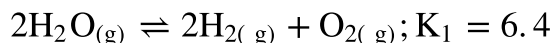
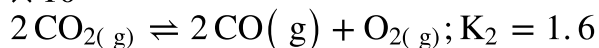


Q1 Equilibrium constants for the following reactions at 1200 K are given :



$$\times 10^{-8}$$



$$\times 10^{-6}$$

The equilibrium constant for the reaction $\text{H}_2(\text{g}) + \text{CO}_2(\text{g}) \rightleftharpoons \text{CO}(\text{g}) + \text{H}_2\text{O}(\text{g})$ at 1200 K will be

- (A) 0.05 (B) 20
(C) 0.2 (D) 5.0

Q2 A chemical reaction $\text{A} \rightleftharpoons \text{B}$ is said to be in equilibrium when

- (A) complete conversion of A to B has taken place
(B) conversion of A to B is only 50% complete
(C) only 10% conversion of A to B has taken place
(D) the rate of transformation of A to B is just equal to rate of transformation of B to A in the system.

Q3 In liquid-gas equilibrium, the pressure of vapours above the liquid is constant at:

- (A) Increase temperature
(B) low temperature
(C) constant temperature
(D) high temperature

Q4 In the reaction, $2\text{SO}_2(\text{g}) + \text{O}_2(\text{g}) \rightleftharpoons 2\text{SO}_3(\text{g}) + X \text{ cal}$, most favourable condition of temperature and pressure for greater yield of SO_3 are

- (A) Low temperature and low pressure
(B) High temperature and low pressure
(C) High temperature and high pressure
(D) Low temperature and high pressure

Q5 A buffer solution contains 0.1 mole of sodium acetate in 1000 cm^3 of 0.1 M acetic acid. To the above buffer solution, 0.1 mole of sodium acetate is further added and dissolved. The pH of the resulting buffer is equal to

- (A) $\text{pK}_a - \log 2$
(B) pK_a
(C) $\text{pK}_a + 2$
(D) $\text{pK}_a + \log 2$

Q6 Hydroxyl ion concentration of 10^{-2} M HCl is

- (A) $1 \times 10^1 \text{ mol dm}^{-3}$
(B) $1 \times 10^{-12} \text{ mol dm}^{-3}$
(C) $1 \times 10^{-1} \text{ mol dm}^{-3}$
(D) $1 \times 10^{-14} \text{ mol dm}^{-3}$

Q7 The K_{sp} of Ag_2CrO_4 , AgCl , AgBr and AgI are respectively, 1.1×10^{-12} , 1.8×10^{-10} , 5.0×10^{-13} , 8.3×10^{-17} . Which one of the following salts will precipitate last if AgNO_3 solution is added to the solution containing equal moles of NaCl , NaBr , NaI and Na_2CrO_4 ?

- (A) Ag_2CrO_4 (B) AgI
(C) AgCl (D) AgBr

Q8 5 moles of PCl_5 are heated in a closed vessel of 5 litre capacity. At equilibrium 40% of PCl_5 is found to be dissociated. What is the value of K_c ?

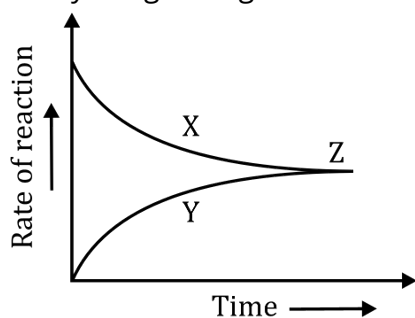
- (A) 0.266 M (B) 0.133 M
(C) 2.5 M (D) 0.20 M



- Q9** The K_p/K_c ratio will be highest in case of
 (A) $\text{CO(g)} + \frac{1}{2} \text{O}_2\text{(g)} \rightleftharpoons \text{CO}_2\text{(g)}$
 (B) $\text{H}_2\text{(g)} + \text{I}_2\text{(g)} \rightleftharpoons 2\text{HI(g)}$
 (C) $\text{PCl}_5\text{(g)} \rightleftharpoons \text{PCl}_3\text{(g)} + \text{Cl}_2\text{(g)}$
 (D) $7\text{H}_2\text{(g)} + 2\text{NO}_2\text{(g)} \rightleftharpoons 2\text{NH}_3\text{(g)} + 4\text{H}_2\text{O(g)}$
- Q10** A 20 litre container at 400 K contains $\text{CO}_2\text{(g)}$ at pressure 0.4 atm and an excess of SrO (neglect the volume of solid SrO). The volume of the container is now decreased by moving the movable piston fitted in the container. The maximum volume of the container, when pressure of CO_2 attains its maximum value, will be (Given that: $\text{SrCO}_3\text{(s)} \rightleftharpoons \text{SrO(s)} + \text{CO}_2\text{(g)}$, $K_p = 1.6 \text{ atm}$)
 (A) 10 litre (B) 4 litre
 (C) 2 litre (D) 5 litre
- Q11** In HS^- , I^- , RNH_2 and NH_3 , order of proton accepting tendency will be
 (A) $\text{I}^- > \text{NH}_3 > \text{RNH}_2 > \text{HS}^-$
 (B) $\text{HS}^- > \text{RNH}_2 > \text{NH}_3 > \text{I}^-$
 (C) $\text{RNH}_2 > \text{NH}_3 > \text{HS}^- > \text{I}^-$
 (D) $\text{NH}_3 > \text{RNH}_2 > \text{HS}^- > \text{I}^-$
- Q12** $\text{H}_2\text{O} + \text{H}_3\text{PO}_4 \rightleftharpoons \text{H}_3\text{O}^+ + \text{H}_2\text{PO}_4^-$, $\text{p}K_1 = 2.15$
 $\text{H}_2\text{O} + \text{H}_2\text{PO}_4^- \rightleftharpoons \text{H}_3\text{O}^+ + \text{HPO}_4^{2-}$, $\text{p}K_2 = 7.20$
 hence pH of 0.01 M NaH_2PO_4 is
 (A) 4.675 (B) 7.350
 (C) 9.35 (D) 2.675
- Q13** The reaction quotient (Q) for the reaction $\text{N}_2\text{(s)} + 3\text{H}_2\text{(g)} \rightleftharpoons 2\text{NH}_3\text{(g)}$ is given by $Q = \frac{[\text{NH}_3]^2}{[\text{N}_2][\text{H}_2]^3}$
 The reaction will proceed from right to left if
 (A) $Q = 0$ (B) $Q = K_c$
 (C) $Q < K_c$ (D) $Q > K_c$
- Q14** Find out the solubility of Ni(OH)_2 in 0.1M NaOH. Given that the ionic product of Ni(OH)_2 is 2×10^{-15} .
 (A) $2 \times 10^{-8} \text{ M}$
 (B) $1 \times 10^{-13} \text{ M}$
 (C) $1 \times 10^8 \text{ M}$
 (D) $2 \times 10^{-13} \text{ M}$
- Q15** The pOH of a buffer solution made by mixing 25 mL of 0.02 M NH_4OH and 25 mL of 0.2 M NH_4Cl at 25°C is _____.
 [$\text{p}K_b$ of $\text{NH}_4\text{OH} = 4.8$]
 (A) 5.8
 (B) 2.8
 (C) 3.8
 (D) 4.8
- Q16** An acid/base dissociation equilibrium is dynamic involving a transfer of proton in forward and reverse directions. Now, with passage of time in which direction equilibrium is favoured?
 (A) in the direction of stronger base and stronger acid
 (B) in the direction of formation of stronger base and weaker acid
 (C) in the direction of formation of weaker base and weaker acid
 (D) in the direction of formation of weaker base and stronger acid
- Q17** In the reaction, $\text{Fe(OH)}_3\text{(s)} \rightleftharpoons \text{Fe}^{3+}\text{(aq)} + 3\text{OH}^-\text{(aq)}$, if the concentration of OH^- ions is decreased by 1/4 times, then the equilibrium concentration of Fe^{3+} will increase by
 (A) 8 times (B) 16 times
 (C) 64 times (D) 4 times



Q18 Study the given figure and label X, Y and Z.



- (A) X Y Z Backward reaction Forward reaction
Products
Equilibrium
- (B) Forward reaction Backward reaction
Equilibrium
- (C) Reversible reaction Irreversible reaction
Equilibrium
- (D) Forward reaction Forward reaction
Backward reaction

Q19 For the reaction $C(s) + CO_2(g) \rightleftharpoons 2CO(g)$, the partial pressures of CO_2 and CO are 2.0 and 4.0 atm respectively at equilibrium. The K_p for the reaction is.

- (A) 0.5 (B) 4.0
(C) 8.0 (D) 32.0

Q20 The following reaction is at equilibrium,
 $Fe_{(aq)}^{3+} + SCN_{(aq)}^- \rightleftharpoons [Fe(SCN)]_{(aq)}^{2+}; K_c$
 Yellow Colourless Deepred

$$K_c = \frac{[Fe(SCN)]^{2+}}{[Fe^{3+}][SCN^-]}$$

In the above reaction, colour intensity of red colour can be increased by

- (A) addition of KSCN
 (B) addition of oxalic acid which reacts with Fe^{3+} ions
 (C) addition of Hg^{2+} ions which react with SCN^- ions
 (D) red colour intensity cannot be changed.

Q21 For the equilibrium, $PCl_5(g) \rightleftharpoons PCl_3(g) + Cl_2(g)$, the total pressure at equilibrium is P and degree of dissociation of PCl_5 is x .

Which of the following is the partial pressure of PCl_3 ?

- (A) $\frac{x}{x-1} \times P$
 (B) $\frac{x}{1-x} \times P$
 (C) $\frac{x}{x+1} \times P$
 (D) $\frac{x}{x-1} \times P$

Q22 A buffer solution of $pH = 9$ can be prepared by mixing

- (A) CH_3COONa and CH_3COOH
 (B) $NaCl$ and $NaOH$
 (C) NH_4Cl and NH_4OH
 (D) KH_2PO_4 and K_2HPO_4 .

Q23 The equilibrium constant for the reversible reaction $2A(g) \rightleftharpoons 2B(g) + C(g)$ is K_1 and for the reaction $\frac{3}{2}A(g) \rightleftharpoons \frac{3}{2}B(g) + \frac{3}{4}C(g)$ is K_2 . K_1 and K_2 are related as:

- (A) $K_2 = K_1^{\frac{3}{4}}$ (B) $K_1 = \sqrt{K_2}$
 (C) $K_2 = \sqrt{K_1}$ (D) $K_1 = K_2^{\frac{3}{4}}$

Q24 The conjugate base of NH_3 is

- (A) NH_4OH (B) NH_2OH
 (C) NH_2^- (D) NH_4^+

Q25 The dissociation constants for acetic acid and HCN at $25^\circ C$ are 1.5×10^{-5} and 4.5×10^{-10} respectively. The equilibrium constant for the equilibrium,
 $CN^- + CH_3COOH \rightleftharpoons HCN + CH_3COO^-$
 would be

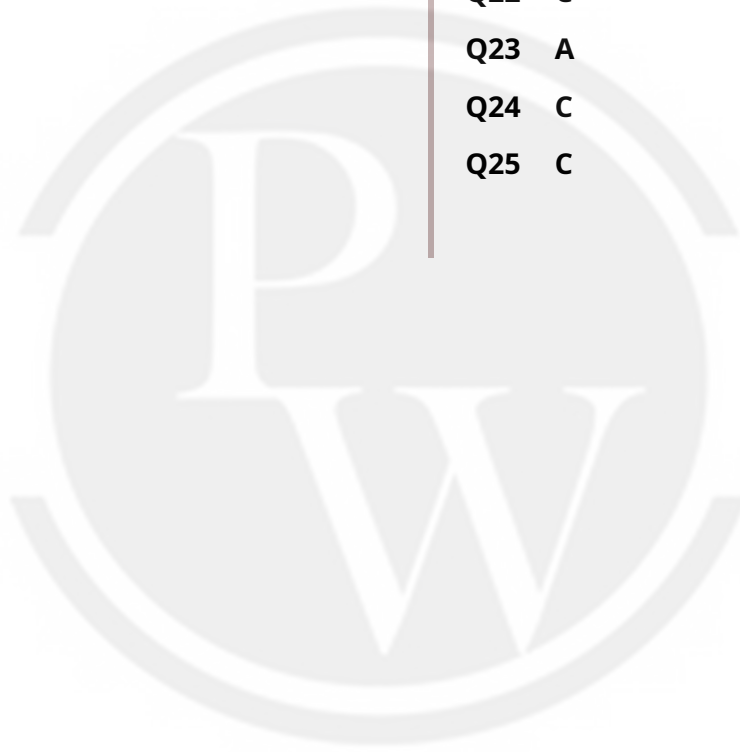
- (A) 3×10^{-5}
 (B) 3×10^{-5}
 (C) 3×10^4
 (D) 3×10^5



Answer Key

Q1 D
Q2 D
Q3 C
Q4 D
Q5 D
Q6 B
Q7 A
Q8 A
Q9 C
Q10 D
Q11 C
Q12 A
Q13 D

Q14 D
Q15 A
Q16 C
Q17 C
Q18 B
Q19 C
Q20 A
Q21 C
Q22 C
Q23 A
Q24 C
Q25 C

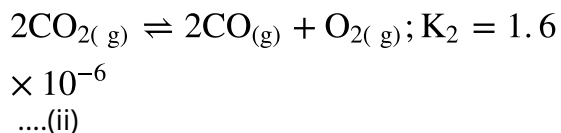
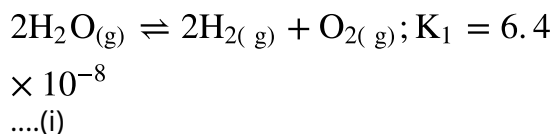


Hints & Solutions

Note: scan the QR code to watch video solution

Q1 Text Solution:

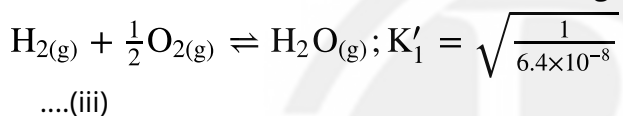
Given:



Required equation is :



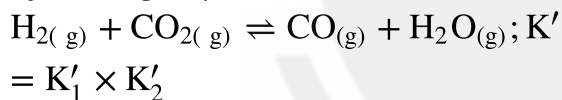
By reversing equation (i) and by multiplying it with $\frac{1}{2}$, we get



And by multiplying equation (ii) with $\frac{1}{2}$, we get $\text{CO}_{2(g)} \rightleftharpoons \text{CO}_{(g)} + \frac{1}{2}\text{O}_{2(g)}; K'_2$ (iv)

$$= \sqrt{1.6 \times 10^{-6}}$$

By adding equations (iii) and (iv), we get



$$K' = \sqrt{\frac{1.6 \times 10^{-6}}{6.4 \times 10^{-8}}} = \sqrt{25} = 5$$

Video Solution:



Q2 Text Solution:

At equilibrium, the rate of forward reaction is equal to rate of backward reaction thus rate of transformation of A to B is equal to rate of transformation of B to A.

Video Solution:



Q3 Text Solution:

For liquid \rightleftharpoons gas

$$K_p = P_{\text{gas}}$$

Vapour pressure of a liquid is constant at constant temperature.

Video Solution:



Q4 Text Solution:

For $2 \text{SO}_2(\text{g}) + \text{O}_2(\text{g}) \rightleftharpoons 2 \text{SO}_3(\text{g})$ (+ X cal, exothermic)

- **Low temperature** favours the exothermic forward reaction.
- **High pressure** favours the side with fewer gas moles (3 → 2).

Hence the greatest SO_3 yield is obtained at **low temperature and high pressure**.

Video Solution:**Q5 Text Solution:**

$$\text{pH} = \text{p}K_a + \log \frac{[\text{Salt}]}{[\text{Acid}]}$$

$$[\text{salt}] = \frac{0.1+0.1}{1000\text{mL}} = \frac{0.2}{1} = 0.2$$

$$[\text{Acid}] = \frac{0.1}{1000\text{mL}} = \frac{0.1}{1} = 0.1$$

$$\text{pH} = \text{p}K_a + \log 2$$

Video Solution:**Q6 Text Solution:**

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

$$[10^{-2}][\text{OH}^-] = 10^{-14}$$

$$[\text{OH}^-] = \frac{10^{-14}}{10^{-2}} = 10^{-12} \text{ mol dm}^{-3}$$

So, the hydroxyl ion concentration of HCl is $1 \times 10^{-12} \text{ mol dm}^{-3}$

Video Solution:**Q7 Text Solution:**

Lower the higher will be precipitation of the salts:

For,

$$\text{Ag}_2\text{CrO}_4 : K_{\text{SP}} = 4S^3 = 1.1 \times 10^{-12} \Rightarrow$$

$$S = 0.65 \times 10^{-4}.$$

$$\text{AgCl} : K_{\text{SP}} = S^2 = 1.8 \times 10^{-10} \Rightarrow S = 1$$

$$.34 \times 10^{-5}.$$

$$\text{AgBr} : K_{\text{SP}} = S^2 = 5 \times 10^{-13} \Rightarrow S = 0.71$$

$$\times 10^{-6}.$$

$$\text{AgI} : K_{\text{SP}} = S^2 = 8.3 \times 10^{-17} \Rightarrow S = 0.9$$

$$\times 10^{-8}.$$

Since, Ag_2CrO_4 has high solubility, it precipitates last.

Video Solution:

Q8 Text Solution:

$$\text{Initial conc.} \quad \frac{5}{5} = 1 \quad 0 \quad 0$$

$$\text{At equilibrium} \quad 1-0.4 \quad 0.4 \quad 0.4$$

$$K_c = \frac{[\text{PCl}_3][\text{Cl}_2]}{[\text{PCl}_5]} = \frac{0.4\text{M} \times 0.4\text{M}}{0.6\text{M}} = 0.266\text{M}$$

Video Solution:**Q9 Text Solution:**

Using the relation $K_p = K_c \cdot (RT)^{\Delta n}$, we get

$$\frac{K_p}{K_c} = (RT)^{\Delta n}$$

Thus $\frac{K_p}{K_c}$ will be highest for the reaction having highest value of Δn .

The Δn values for various reactions are

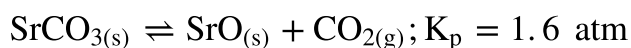
$$(a) \Delta n = 1 - \left(1 + \frac{1}{2}\right) = -1/2$$

$$(b) \Delta n = 2 - (1 + 1) = 0$$

$$(c) \Delta n = (1 + 1) - 1 = 1$$

$$(d) \Delta n = (2 + 4) - (7 + 2) = -3$$

Thus maximum value of $\Delta n = 1$

Video Solution:**Q10 Text Solution:**

$$K_p = \frac{P_{\text{CO}_2} \times P_{\text{SrO}}}{P_{\text{SrCO}_3}} \Rightarrow 1.6$$

$$= P_{\text{CO}_2} (\because P_{\text{SrO}} = P_{\text{SrCO}_3} = 1)$$

\therefore Maximum pressure of $\text{CO}_2 = 1.6 \text{ atm}$

Let the maximum volume of the container when pressure of CO_2 is 1.6 atm be VL

During the process, $PV = \text{constant}$

$$\therefore 0.4 \times 20 = 1.6 \times V$$

$$\Rightarrow V = \frac{0.4 \times 20}{1.6} = 5 \text{ L.}$$

Video Solution:**Q11 Text Solution:**

Strong base has higher tendency to accept the proton. Increasing order of base and hence the order of accepting tendency of proton is $\text{I}^- < \text{HS}^- < \text{NH}_3 < \text{RNH}_2$

Video Solution:**Q12 Text Solution:**

H_2PO_4^- is amphoteric.



$$\text{pH} = \frac{\text{p}K_1 + \text{p}K_2}{2} = \frac{2.15 + 7.20}{2} = 4.675$$

Video Solution:

Q13 Text Solution:

If $Q > K_c$ it means concentration of product is more than that reactants at equilibrium, so reaction will proceed from products to reactants i.e., right to left.

Video Solution:**Q14 Text Solution:**

If 'S' is the solubility of $\text{Ni}(\text{OH})_2$ then for the equilibrium, $\text{Ni}(\text{OH})_2 \rightleftharpoons \text{Ni}^{+2} + 2\text{OH}^-$, solubilities will be: S S 2S.

NaOH is a strong electrolyte and dissociates completely as: $\text{Ni}(\text{OH})_2 \rightleftharpoons \text{Ni}^{+2} + 2\text{OH}^-$, the concentrations

will be 0.1 0.1 0.1

\therefore Total $[\text{OH}^-] = 0.1 + 2s \approx 0.1$ Ionic

$\therefore 2s \ll 0.1$

product of,

$$\text{Ni}(\text{OH})_2 = 2 \times 10^{-15} = [\text{Ni}^{+2}] [\text{OH}^-]^2$$

$$2 \times 10^{-15} = s[0.1]^2$$

$$\therefore S = \frac{2 \times 10^{-15}}{(0.1)^2} = 2 \times 10^{-13} \text{ M}$$

Video Solution:**Q15 Text Solution:**

$$pOH = pK_b + \log \frac{[\text{Salt}]}{[\text{base}]}$$

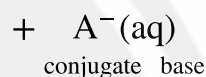
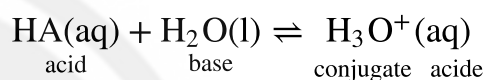
$$pOH = 4.8 + \log \frac{0.2}{0.02}$$

$$pOH = 4.8 + \log 10$$

$$= 5.8$$

Video Solution:**Q16 Text Solution:**

Dissociation of a weak acid in water :



If HA is a stronger acid than H_3O^+ then HA will donate protons and not H_3O^+ and the solution will mainly contain A^- and H_3O^+ ions. Hence, the equilibrium moves in the direction of formation of weaker acid and weaker base. Similarly, this is true for dissociation of a weak base also.

Video Solution:**Q17 Text Solution:**

Given relation,



$$K_{\text{eq}} = \frac{[\text{Fe}^{3+}][\text{OH}^-]^3}{[\text{Fe}(\text{OH})_3]} = \frac{[\text{Fe}^{3+}][\text{OH}^-]^3}{1}$$

When $[\text{OH}^-]$ is decreased by 1/4 times, K becomes

$$K_{\text{eq}} = [\text{Fe}^{3+}] \left(\frac{1}{4} \cdot [\text{OH}^-] \right)^3$$



$$= [\text{Fe}^{3+}] \frac{1}{64} [\text{OH}^-]^3$$

To keep the value of K_{eq} constant, the $[\text{Fe}^{3+}]$ must be increased by 64 times so that K_{eq} remains as such

$$K_{\text{eq}} = 64 \times \frac{1}{64} [\text{Fe}^{3+}] [\text{OH}^-]^3$$

$$K_{\text{eq}} = [\text{Fe}^{3+}] [\text{OH}^-]^3$$

Video Solution:



Q18 Text Solution:

The concentration of reactants decreases and that of products increases with time. Rate of reaction increases with time. At equilibrium, $R_f = R_b$

Video Solution:



Q19 Text Solution:

$$K_p = \frac{(P_{\text{CO}})^2}{(P_{\text{CO}_2})}; K_p = \frac{4 \times 4}{2} = 8; C(s) = 1;$$

The concentration of solids and liquids are taken as unity.

Video Solution:



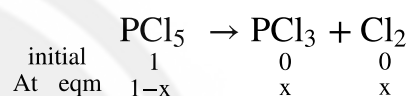
Q20 Text Solution:

Addition of KSCN increases the colour intensity of the solution as it shifts the equilibrium to right. Addition of reagents like oxalic acid or Hg^{2+} ions which remove Fe^{3+} or SCN^- ions shift the equilibrium to the left and colour intensity decreases.

Video Solution:



Q21 Text Solution:



total number of moles at eqm = $1-x+x+x = 1+x$

number of moles of $\text{PCl}_3 = x$

$$\text{molefraction of } \text{PCl}_3 = \chi_{\text{PCl}_3} = \frac{x}{1+x}$$

Partial pressure of

$$\text{PCl}_3 = P_{\text{PCl}_3} = \chi_{\text{PCl}_3} \times P = \frac{x}{1+x} \cdot P$$

Video Solution:

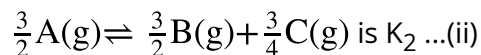
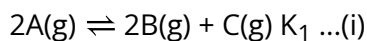


Q22 Text Solution:

$\text{NH}_4\text{OH} + \text{NH}_4\text{Cl}$ is a basic buffer solution.

Video Solution:

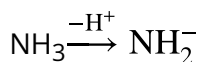
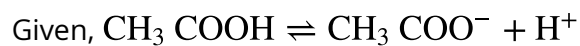


Q23 Text Solution:

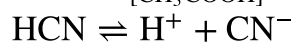
eq. (ii) is $\frac{3}{2}$ times of eq. (i), hence, $K_2 = (K_1)^{\frac{3}{4}}$

Video Solution:**Q24 Text Solution:**

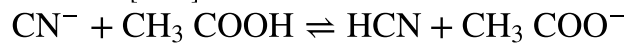
Conjugate base is formed by removal of a proton. Hence conjugate base of NH_3 is NH_2^- .

**Video Solution:****Q25 Text Solution:**

$$K_1 = \frac{[\text{CH}_3\text{COO}^-][\text{H}^+]}{[\text{CH}_3\text{COOH}]} = 1.5 \times 10^{-5}$$



$$K_2 = \frac{[\text{CN}^-][\text{H}^+]}{[\text{HCN}]} = 4.5 \times 10^{-10}$$



$$K = \frac{[\text{HCN}][\text{CH}_3\text{COO}^-]}{[\text{CN}^-][\text{CH}_3\text{COOH}]}$$

$$K = \frac{K_1}{K_2} = \frac{1.5 \times 10^{-5}}{4.5 \times 10^{-10}} \approx 0.3 \times 10^5 \text{ or } K = 3$$

$$\times 10^4$$

Video Solution:
[Android App](#)
[iOS App](#)
[PW Website](#)