

Q.18 The rate at which a substance reacts depends on its :

[1] Atomic weight [2] Molecular weight [3] Equivalent weight [4] Active mass

Q.19 The equilibrium which remains unaffected by pressure change is :

 $[1] N_2(g) + O_2(g) \longrightarrow 2NO(g)$ [2] $2SO_2$ $(g) + O_2(g) \longrightarrow 2SO_3(g)$ $[3] 2O_3(9) \rightleftharpoons 3O_2$ (g) $[4] 2NO₂(g) \longrightarrow N₂O₄(g)$

Q.20 Le Chatelier's principle is applicable only to :

[1] Reaction under equilibrium [2] Reaction without equilibrium

[3] Ionization of electrolytes [4] None of these

298 K, the equilibrium between N,O_s and NO_s may be represented by the followin

(g) \Rightarrow 2NO₂(g). If the total pressure of the equilibrium mixture is P and the degree of dis

(g) at 298 K is x, which one of the foll **Q.21** At 298 K, the equilibrium between N_2O_4 and NO_2 may be represented by the following equation $\mathsf{N}_2\mathsf{O}_4(\mathsf{g})$ \Longrightarrow 2NO₂(g). If the total pressure of the equilibrium mixture is P and the degree of dissociation of $\mathsf{N}_2\mathsf{O}_4(\mathsf{g})$ at 298 K is x, which one of the following is the pressure of $\mathsf{N}_2\mathsf{O}_4(\mathsf{g})$ under this condition :

Q.22 For the reaction : $\text{PCl}_5(g) \longrightarrow \text{PCl}_3(g) + \text{Cl}_2(g)$

the forward reaction at a constant temperature is favoured by

- [1] Introducing an inert gas at constant volume
- [2] Introducing chlorine gas at constant volume
- [3] Introducing an inert gas at constant pressure
- [4] None of these
- **Q.23** One mole of SO₃ was placed in one litre vessel at a certain temperature. The following equilibrium was established $2\mathrm{SO}_3$ \Longrightarrow $2\mathrm{SO}_2$ + O_2 . At equilibrium 0.6 moles of SO_2 were formed. The equilibrium constant of the reaction will be :

[1] 0.36 [2] 0.45 [3] 0.54 [4] 0.675

Q.24 Raising the temperature of a reversible chemical reaction :

[1] Favours the forward rate only [2] Favours the backward rate only

[3] Favours both the forward and backward rate [4] Favours neither the forward nor the backward rates

- **Q.25** The equilibrium constant for the reaction H₂ (g) + I₂(g) \Longrightarrow 2HI(g) is 49. If the equation of the reaction is written as $\frac{1}{2}$ H₂+ $\frac{1}{2}$ I₂(g) \Longrightarrow HI (g) then :
	- [1] The value of equilibrium constant would change
	- [2] The value of equilibrium constant would remain the same
	- [3] Mole fraction of hydrogen reacted would change
	- [4] Mole fraction of HI formed will change

Q.26 ''If a system at equilibrium is subjected to a change in any one of the factors such as temperature, pressure or concentration, the system adjusts itself in such a way as to annual the effect of that change''. This is called :

[1] Avogadro's hypothesis [2] Gay Lussac's law

- [3] Boyle's law [4] Le Chatelier's principle
-
- **Q.27** For the reaction : 2SO_3 \Longrightarrow $2\text{SO}_2 + \text{O}_2$ the equilibrium constant expression is :

$$
\begin{bmatrix} 1 \end{bmatrix} \begin{bmatrix} K_c = \frac{2[SO_2] \times [O_2]}{2[SO_3]} & [2] \end{bmatrix} \begin{bmatrix} K_c = \frac{[SO_2] \times [O_2]}{[SO_3]} & [3] \end{bmatrix} \begin{bmatrix} K_c = \frac{[SO_3]^2}{[SO_2]^2 \times [O_2]} & [4] \end{bmatrix} \begin{bmatrix} K_c = \frac{[SO_2]^2 \times [O_2]}{[SO_3]^2} \end{bmatrix}
$$

- **Q.28** Which of the following reactions will be favoured at low pressure :
	- $[1] H_2 + I_2$ 2HI $[2] N_2 + 3H_2 \longrightarrow 2NH_3$ [3] $\mathsf{PCl}_{_5} \Longleftarrow \mathsf{PCl}_{_3} + \mathsf{Cl}_{_2}$ $[4]$ N₂ + O₂ \Longrightarrow 2NO
- **Q.29** CH₃COOH + C₂H₅OH \Longrightarrow CH₃COOC₂H₅ + H₂O (K_C = 4)

In the above reaction one mole each of acetic acid and alcohol are heated in the presence of a little conc. ${\sf H_2SO}_4$. On equilibrium being attained :

- [1] One mole of ethyl acetate is formed
- [2] 2 moles of ethyl acetate are formed
- [3] 1/3 moles of ethyl acetate is formed
- [4] 2/3 moles of ethyl acetate is formed
- **Q.30** When a catalyst is added to a system in equilibrium :
	- [1] The equilibrium concentrations are increased
	- [2] There is no effect on equilibrium concentrations
	- [3] The equilibrium concentrations are decreased
	- [4] The rate of forward reaction increase and that of backward reaction decreases
- $H_2 + I_2 \Longleftrightarrow PCl_3 + Cl_2$
 $PCl_3 \Longleftrightarrow PCl_3 + Cl_2$
 $PCl_3 + Cl_2 \Longleftrightarrow CH_3COOC_2H_3 + H_2O (K_0 = 4)$
 $QCOOH + C_2H_3OH \Longleftrightarrow CH_3COOC_2H_3 + H_2O (K_0 = 4)$
 $QCOOH + C_2H_3OH \Longleftrightarrow CH_3COOC_2H_3 + H_2O (K_0 = 4)$
 $QCOOH + C_2H_3OH \Longleftrightarrow CH_3OOC_2H_3 + H_2O (K_0 = 4)$
 $QCOOH + C_2H_3OH \Longleftrightarrow CH_3O$ **Q.31** Suppose the reaction : $PCI_{5}(s) \implies PCI_{3}(g) + CI_{2}(g)$ is at equilibrium in a closed vessel. At a constant temperature on addition of PCI₅, what will be effect on equilibrium concentration of CI₂ (g) :
	- [1] Will be decreased
	- [2] Will be increased
	- [3] Will remain unaffected
	- [4] Can not be predicated without the value of K_p
- **Q.32** Equilibrium concentrations of HI, I₂ and H₂ is 0.7, 0.1 and 0.1 moles/litre. Calculate equilibrium constant for reaction $I_2 + H_2 \rightleftharpoons 2HI$ is

[1] 0.36 [2] 36 [3] 49 [4] 0.49

Q.33 Which of the following will shift the reaction $\text{PCl}_3 + \text{Cl}_2 \rightleftharpoons \text{PCl}_5$ to the left side :

 $x = 10^{-8}$
 $(21 + x 10^{-4})$
 $(315 x 10^{-4})$
 $(412 x 10^{-8})$
 $(51 x 10^{-4})$
 $(61 x 10^{-4})$
 $(71 x 10^{-4})$
 $(81 x 10^{-4})$
 $(141 x 10^{-4})$
 $(1$ **Q.34** In which of the following equilibria the value of K_{ρ} is less than K_{ε} : $[1] H_2 + I_2$ 2HI $[2] N_2 + 3H_2 \longrightarrow 2NH_3$ $[3] N_2 + O_2$ 2NO $[4] CO + H_2O \rightleftharpoons CO_2 + H_2$ **Q.35** An equilibrium mixture for the reaction $:$ 2H₂S (g) \Longrightarrow 2H₂ (g) + S₂(g) has 0.5 mole H₂S, 0.1 mole of H₂ and 0.4 mole S_2 in a one litre vessel. The equilibrium constant of this reaction is given by : $[1] 0.004$ mole litre⁻¹ $[2] 0.08$ mole litre⁻¹ $[3] 0.016$ mole litre⁻¹ $[4] 0.160$ mole litre⁻¹ **Q.36** What will be the rate of decomposition of a gas at a particular temperature, if concentration of the gas is 0.05 mole/litre ? Rate constant of decomposition of gas at this temperature is 10⁻⁴ min⁻¹. $[1] 5 \times 10^{-6}$ $[2] 1 \times 10^{-4}$ $[3] 5 \times 10^{-4}$ $[4] 2 \times 10^{-6}$ **Q.37** If the concentration of reactants and products is doubled, then the equilibrium constant (K_c) will be : $[1]$ 2K. $[2] K/2$ $[3] K/4$ [4] unchanged **Q.38** $\,$ 0.96 gram hydrogen iodide is heated at 400°C till equilibrium is established. 14.0 ml of N/10 Na $_{2}$ S $_{2}$ O $_{3}$ solution is needed to neutralize iodine obtained from this reaction, then calculate the percent amount of dissociation of HI : [1] 28% [2] 18.6% [3] 20% [4] 62.3%

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Qus.								8	9	10	11	12	13 [°]	14	15	16	17	18		20
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Ans.		⌒		ົ				⌒		⌒	⌒				ົ					

Aewer Key

Exercise # 2

stant K_y, by the expression
 $\frac{1+2\sqrt{K_p}}{2}$ [2] $\sqrt{\frac{1+2K_p}{2}}$ [3] $\sqrt{\frac{2K_p}{1+2K_p}}$
 A B heat

(i) Equilibrium constant

r./r,

r./r,
 $\frac{1+2\sqrt{K_p}}{2}$ [13] $\frac{1}{\sqrt{1+2K_p}}$ [14] $\frac{2\sqrt{K_p}}{1+2\sqrt{K_p}}$

(iii) Ad **Q.9** Some gaseous equilibrium have following relation in equilibrium constants $CO + H_2O$ K $CO_2 + H_2$; $CO + O_2$ K₁ CO_2 ; $H_2 + O_2$ K₂ H_2O [1] $K = K_1K_2$ [2] $K = (K_1K_2)^2$ [3] $K = (K_1K_2)$ $^{-1/2}$ [4] K = K₁/ K₂ **Q.10** Effect of temperature on equilibrium constant is given by log K₂ –log K₁ = $\frac{1}{2.303R}$ $\boxed{\frac{1}{T_2} - \frac{1}{T_1}}$ H | 1 1 2.303R \mid T, T. $\frac{-\Delta H}{\Delta 303R} \left[\frac{1}{T_2} - \frac{1}{T_1} \right]$. Then for a endothermic reaction the false statement is $\left[1\right]$ $\left[\overline{T_2}^{-} \right]$ $\overline{T_1}$ 1 1 T_2 T. $\begin{bmatrix} 1 & 1 \end{bmatrix}$ $\left[\frac{1}{T_2} - \frac{1}{T_1}\right]$ = positive [2] log K₂ > log K₁ [3] ∆H = positive [4] K₂ [4] $K_2 > K_1$ **Q.11** For the reaction : $2HI(q)$ (g) + I_2 (g), the degree of dissociation (α) of HI(g) is related to equilibrium constant K_p by the expression [1] $\frac{1+2\sqrt{K_{\rm P}}}{2}$ 2 + $\frac{1+2K_{\rm P}}{2}$ 2 + [3] $\sqrt{\frac{2R_{p}}{1+2K_{p}}}$ 2K $1 + 2K$ [4] P P 2 $\sqrt{\mathsf{K}}$ 1 + 2 $\sqrt{\mathsf{K}}$ **Q.12 List X List Y** (A) A B heat (i) Equilibrium constant (B) r_{b}/r_{f} (ii) Adaptation of low temp. (C) $r/r_{\rm b}$ (iii) (Equilibrium constant)–1 (D) 2A(g) + B(g) C(g) (iv) A(g) + B(g) C(g) + D(g) (E) No effect of pressure (v) $\Delta n < 0$ Correct match list X and Y $[1]$ A–(ii), B–(iii), C–(i), D–(v), E–(iv) $[2]$ A–(iii), B–(ii), C–(i), D–(v), E–(iv) $[3]$ A–(iv), B–(iii), C–(i), D–(v), E–(ii) [4] None of these **Q.13** aA + bB cC + dD, ∆H = QKJ. If the higher yield of product is obtained by the increase in pressure and decrease in temp. then $[1] (c+d) > (a+b)$ and Q positive [2] (c+d) $> (a+b)$ and Q negative $[3] (c+d) < (a+b)$ and Q negative [4] $(c+d) < (a+b)$ and Q positive ${\sf Q.14}$ $\;\;\;$ When heating PCI₅ then it decompose PCI $_3$ and CI $_2$ in form of gas, The density of gas mixture is 70.2 and 57.9 at 200°C and 250°C. The degree of dissociation of PCI_s at 200°C and 250°C if [1] 48.50% & 80% [2] 60% & 70% [3] 70% & 80% [4] 80% & 90% **Q.15** If log kp₂ – log kp₁ = $\frac{x}{T_1} - \frac{y}{T_2}$ $x\left(\frac{1}{T_1}-\frac{1}{T_2}\right)$ $\begin{bmatrix} 1 & 1 \end{bmatrix}$ $\left(\frac{1}{\mathsf{T}_{1}} - \frac{1}{\mathsf{T}_{2}}\right)$ then value of x is $[1]$ Δ H°/2.303R [2] Δ H° $[3] \frac{1}{2.3021}$ [4] None of these **Q.16** Series of equal reaction Represent by general equation E + F = G +H. In this equation each reaction of series reached at equilibrium and their equilibrium constant are follows which of the following reaction equilibrium established too late [1] K_c= 2 \times 10⁵ [2] $K_c = 85$ $= 85$ [3]K_c = 0.010 [4] K_c [4] $K_s = 7 \times 10^8$ **Q.17** Equilibrium constant in condition of there reversible equation ; $Cu^{+2} + 4NH_{3}$ 1 K K_{-1} Cu(NH₃)₄⁺² $[1]$ k₁k₋₁ $[2]$ k₁/k₋₁ [3] k_{-1} / k_{1} $[4]$ $(k_{-1})^2$

ction is -2SO₂ + O₂ Li^{ut}uti 2SO₃

the formation of one ton of SO₃, what, would be the quantity of O₂ required

12) 2.0 Ton

13) 2.0 Ton

131 2.0 Ton

141 0.002 Tor

141 0.002 Tor

141 0.002 Tor

141 0.002 Tor **Q.18** In a gaseous mixture moles of A, B and C are 'a' 'b' and 'c' if total volume is V litre and mole fraction of A, B and C are $\mathsf{X}_{\sf A}$, $\mathsf{X}_{\sf B}$ and $\mathsf{X}_{\sf C}$ then which of the following relation is incorrect. [1] $X_{A} = \frac{a}{a + b}$ $a + b + c$ $[2] X_{B} = \frac{b}{a+b}$ $a + b + c$ [3] $X_c = \frac{c}{a + b}$ $a + b + c$ $[4] X_{\rm A} + X_{\rm B} = 1 + X_{\rm c}$ $Q.19$ + Y₂ \Box \Box \Box 2XY reaction was studied at a certain temperature. In the beginning 1 mole of X₂ was taken in a one litre flask and 2 moles of Y_2 was taken in another 2 litre flask. What is the equilibrium concentration of X_2 and Y_{2}^{\prime} (Given equilibrium concentration of [XY] = 0.6 moles/lit. $[1]$ $\left(\frac{1}{3} - 0.3\right)$ $\left(\frac{2}{3} - 0.3\right)$ $\left(\frac{1}{3}-0.3\right).\left(\frac{2}{3}-0.3\right)$ $\qquad [2]\left(\frac{1}{3}-0.6\right).\left(\frac{2}{3}-0.6\right)$ $\left(\frac{1}{3}\text{-}0.6\right) \cdot \left(\frac{2}{3}\text{-}0.6\right)$ [3] (1–0.3), (2–0.3) [4] (1–0.6), (2–0.6) $\overline{\bf Q.20}$ Reaction is $-$ 2SO₂ + O₂ $\overline{\boxplus}$ $\overline{\boxplus}$ $\overline{\boxplus}$ 2SO₃ For the formation of one ton of SO₃, what, would be the quantity of O₂ required [1] 0.20 Ton [2] 0.02 ton [3] 2.0 Ton [4] 0.002 Ton **Q.21** For reaction aA \Box \Box \Box \Box + mM. In condition of suddenly volume increase degree of dissociation a is decrease it represent that. $[1]$ a < (ℓ + m) $[2]$ a = (ℓ + m) $[3]$ a = (ℓ - m) $[4]$ a > (ℓ + m) $Q.22$ dissociates as $XY_2(g) \Box \Box \Box$ $XY(g) + Y(g)$ Initial pressure of $\rm~XY_{2}$ is 600 mm Hg. At equilibrium the total pressure is 800 mm Hg. Calculate the value of K_p. It is assumed that the volume of the system remains unchanged $[2] 400$ $[3] 200$ $[4] 50$ **Q.23** In a .25 litre tube occur dissociation of NO. Its initial mole is 4. If its degree of dissociation is 10%.The K_p for Rxn 2 NO $\overline{\oplus \oplus \oplus}$ N₂ + O₂ $[1]$ $\overline{(18)}^2$ 1 $\overline{(2)}$ $\overline{(8)^2}$ 1 8 $[3] \frac{1}{16}$ $\frac{1}{16}$ [4] 1 32 ${\bf Q.24}$ The equilibrium constant for the reaction $\mathsf{N}_2+\mathsf{O}_2$ $\overline{\boxplus}\,\overline{\boxplus}\,\overline{\boxplus}\,\,$ 2NO is 0.0842 at 3500K. The fraction of equilibrium mixture of N_{2} and O_{2} converted into NO is [1] 12.66% [2] 17.2% [3] 15.9% [4] 16.0% **Q.25** The vapour density of PCI₅ is 104 but when heated to 230°C, its V.D. is reduced to 52. The degree of dissociation of PCI₅ at this temperature will be [1] 6.8% [2] 100% [3] 46% [4] 64% ${\bf Q.26}$ $\;\;\;\;$ The value K for H₂ (g) + CO₂ (g) $\bar{\boxplus}\,\bar{\boxplus}\,\bar{\boxplus}\,\,$ H₂O (g) + CO (g) is 1.80 at 1000°C. If 1.0 mole of each H₂ and CO₂ are placed in 1 litre flask, the final equilibrium concentration of CO at 1000°C will be [1] 0.295 M [2] 0.385 M [3] 0.572M [4] 0.473 M **Q.27** The decomposition of N₂O₄ to NO₂ is carried out at 280°C. When equilibrium is reached, 0.2 mol of N₂O₄ and 2 × 10⁻³ mol of NO₂ are present in 2 litres solution. The K_c for the reaction N₂O₄ 日回画 2NO₂ is $[1] 1 \times 10^{-2}$ $[2] 1 \times 10^{-5}$ $[3] 2 \times 10^{-3}$ $[4] 2 \times 10^{-5}$

Q.28 If K_p for a reaction A (g) + 2 B (g) 日回画 3C (g) + D (g) is 0.05 atm at 1000 K. Its K_c in terms of R will be

$$
[1] 20000 \text{ R} \qquad [2] 0.02 \text{ R} \qquad [3] 5 \times 10^{-5} \text{ R} \qquad [4] \frac{5 \times 10^{-5}}{\text{ R}}
$$

 ${\bf Q.29}$ $\;\;$ An equilibrium mixture for the reaction 2 H₂S (g) $\pm\,\pm\,\pm\,$ and $\pm\,$ As (g) had one mole of hydrogen sulphide 0.2 mole of ${\sf H}_{_2}$ and 0.8 mole of ${\sf S}_{_2}$ in a 2 litre vessel. The value of ${\sf K}_{_{\rm c}}$ in mole litre $^{\scriptscriptstyle -1}$ is [1] 0.004 [2] 0.016 [3] 0.080 [4] 0.160

 $Q.30$ and $K₂$ are equilibrium constant for reactions [1] and [2]

N2 (g) + O² (g) 2NO (g) [1]

NO (g)
$$
\Box \Box \Box
$$
 \Box $N_2(g) + \frac{1}{2} O_2(g)$

Then

$$
[1] K_1 = \left(\frac{1}{K_2}\right)^2
$$
 [2] K_1 = K_2

- K_2^2
- (g) $\overline{a} \equiv \frac{1}{m} \int_{2}^{m} N_{2}(g) + \frac{1}{2} O_{2}(g)$
 $S_{1} = \frac{1}{K_{2}}$
 $\left(2 | K_{1} = K_{2}^{2} \right)$
 $\left(2 | K_{1} = K_{2}^{2} \right)$
 $\left(3 | K_{1} = \frac{1}{K_{2}} \right)$
 $\left(4 | K_{1} = (K_{2})^{2} \right)$
 $\left(5 | K_{1} = (K_{2})^{2} \right)$
 $\left(6 | K_{2} = (K_{1})^{2} \right)$
 \left (g) [2] $[3] K_1 =$ 2 1 K [4] $K_1 = (K_2)^0$
- **Q.31** The equilibrium constant for the reaction

Zn (s) + Cu $^{2+}$ (aq) $\bar{\boxplus} \bar{\boxplus}$ $\bar{\boxplus}$ Zn $^{2+}$ (aq) + Cu (s) + and Cu (s) 2 Ag⁺ (aq) $\bar{\boxplus} \bar{\boxplus}$ Cu $^{2+}$ (aq) + 2Ag (s) $\bar{\boxplus}$ and K $_2$ respectively. Then the equilibrium constant for the reaction

Zn (s) + 2 Ag⁺ (aq) $\overline{\boxplus} \overline{\boxplus} \overline{\boxplus}$ Zn²⁺ (aq) + 2 Ag (s) will be

$$
[1] K_1 + K_2 \qquad [2] K_1 \times K_2 \qquad [3] K_1 / K_2 \qquad [4] K_1 - K_2
$$

Q.32 For an equilibrium change involving gaseous phase, the forward reaction is firs order while the reverse reaction is second order. The unit of $\mathsf{K}_\mathrm{_p}$ for the forward equilibrium is

$$
[1] Atm
$$
 [2] Atm² [3] Atm⁻¹ [4] Atm⁻²

Q.33 Ammonia forms complexes with Ag⁺ according to the following reactions

$$
(\text{i}) \text{ Ag[H}_2\text{O]}_2 + \text{NH}_3(\text{aq}) \overset{\text{f.}}{\oplus} \overset{\text{f.}}{\oplus} \text{ [Ag[NH}_3] [H_2\text{O (aq)}]^+ + H_2\text{O}(\ell)
$$

(ii)
$$
[Ag(NH_3)(H_2O)(aq)]^* + NH_3(aq) \Box \Box \Box
$$
 Ag $(NH_3)_2^*(aq) + H_2O(\ell)$

 $[2]$ 2.0 \times 10³

The equilibrium constants of equilibrium (i) and (ii) are 2.0 \times 10³ and 8.3 \times 10³ respectively. Equilibrium constant of the following reaction $\mathtt{[Ag(H_2O)_2(aq)^+ + 2NH_3(aq) \; \exists \; \exists \; \mathbb{\boxplus} \; \; \text{Ag(NH}_3)_2^{\texttt{+}} + 2\text{H}_2\text{O} \left(\ell \right) }$ will be -

$$
[1] 4.15
$$

[3] 8.3×10^3

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[4] 16.6 \times 10^6
```
 $[4] \frac{\sqrt{2}}{2}$ $1 + \sqrt{2}$

 ${\bf Q.34}$ Two is the equilibrium constant for the reaction ${\sf A_2+B_2}$ $\overline{\uplus \uplus \emptyset}$ 2AB at a given temperature. What is the degree of dissociation for $\mathsf{A}_2^{\vphantom{\prime}}$ or $\mathsf{B}_2^{\vphantom{\prime}}$

 $[1] 0.2$ $[2] 0.5$

$$
[3] \frac{1}{1+\sqrt{2}}
$$

 ${\bf Q.35}$ At temperature, T, a compound AB₂(g) dissociates according to the reaction 2AB₂(g) $\overline{\boxplus} \overline{\boxplus}$ 2AB(g) + B₂(g) with a degree of dissociation x, which is small compared with unity, the expression for kp, in terms of x and the total pressure, P is -

$$
[1] \frac{Px^3}{2} \qquad [2] \frac{Px^2}{2} \qquad [3] \frac{Px^3}{3} \qquad [4] \frac{Px^2}{2}
$$

represent the equilibrium constant for reaction $H_2 + I_2 \overline{H_2H_1R_2}$ 2H & K₂ for % $I_2 \overline{H_2H_1R_2}$ HI,
wen K, & K₂ would be
 $\langle x K_2 = 1$ [2] $\sqrt{K_1} \times K_2 = 1$ [3] $\sqrt{K_1} = K_2$
Bodenstein experiment, one mol H₂ ${\bf Q.36}$ $\rm\,$ For the equilibrium CaCO $_3$ (s) $\overline{10}$ $\overline{10}$ $\rm\,$ CaO(s) + CO $_2$ (g) which of the following expressions is correct [1] K_p = [CaO] [CO₂] / [CaCO₃] $[2] K_p = [P_{CaO} \times P_{CO2}] / P_{CaCO3}$ [3] $K_{p} = P_{CO_{2}}$ [4] All of these **Q.37** One mole of the N₂ is mixed with 3 moles of H₂ in a litre container. If 50% of H₂ is converted into ammonia by the reaction. N $_2$ (g) + 3H $_2$ (g) $\overline{\boxplus} \overline{\boxplus}$ 2NH $_3$ (g), then the total number of moles of gas a the equilibrium are $[1] 1.5$ $[2] 4.5$ $[3] 3.0$ $[4] 6.0$ **Q.38** For the system A(g) + 2B (g) $\overline{\oplus \oplus \oplus}$ C (g), the equilibrium concentration are A = 0.06 mol L⁻¹; B = 0.12 mol L⁻¹; C = 0.216 mol L⁻¹. The Keq of the reaction is $[1] 250$ $[2] 416$ $[3] 4 \times 10^{-3}$ $[4] 125$ **Q.39** If K₁ represent the equilibrium constant for reaction H₂ + I₂ $\overline{\boxplus}$ $\overline{\boxplus}$ \oplus \mathbb{R} \times \mathbb{R} $\overline{\boxplus}$ $\overline{\boxplus}$ \oplus $\overline{\boxplus}$ $\overline{\boxplus}$ $\overline{\boxplus}$ $\overline{\boxplus}$ $\overline{\boxplus}$ $\overline{\boxplus}$ $\overline{\boxplus}$ between K $_{_{1}}$ & K $_{_{2}}$ would be [1] $K_{1} \times K_{2}$ $= 1$ [2] $\sqrt{K_1} \times K_2$ $= 1$ [3] $\sqrt{K_1} = K_2$ [4] \sqrt{K} ₁ = \sqrt{K} ₂ **Q.40** In a Bodenstein experiment, one mol $\mathsf{H}_{_2}$ and two mol I $_2$ are taken initially in a one litre flask. If at equilibrium 0.5 mol H₂ are present, the equilibrium concentrations of I₂ and HI $\,$ in mol $\,$ / $^{\rm t}$ are $[1] 0.5, 1.0$ (20 1.0, 0.5 $[3] 1.5, 1.0$ [4] 1.5 , 0.5 ${\bf Q.41}$ The reaction : CH $_{3}$ COOH + C $_{2}$ H $_{5}$ OH $\overline{\boxplus}$ $\overline{\boxplus}$ CH $_{3}$ COOC $_{2}$ H $_{5}$ + H $_{2}$ O is studied in a 2 ℓ vessel by taking initially a mol acetic acid and b mole ethanol. At equilibria, x mol each of ester and water are formed. The value of K for this reaction is equal to $[1]$ $\overline{(a-x)(b-x)}$ $4x^2$ $(a-x)(b-x)$ [2] $\overline{(a-x)(b-x)}$ x^2 $(a-x)(b-x)$ [3] $(a-x)(b-x)$ $2x^2$ $(a-x)(b-x)$ [4] $\overline{2(a-x)(b-x)}$ x^2 $2(a - x)(b - x)$ **Q.42** At a total equilibrium pressure of 1.0 atmosphere, the degree of dissociation of phosgene is 0.2 $COCI₂(g)$ \Box \Box \Box $CO (g) + Cl₂(g)$ Now the same equilibrium is established at the same temperature in presence of N₂ gas at a partial pressure of 0.4 atm. in a total pressure of 1.0 atm. The new degree of dissociation, α , is –

$$
[1] \alpha < 0.2
$$
 $[2] \alpha > 0.2$ $[3] \alpha = 0.2$ $[4] \alpha = 0$

Q.43 In the study of the reaction

$$
\text{Cl}_2 + \text{PCI}_3 \stackrel{\frown}{\Box} \stackrel{\frown}{\Box} \stackrel{\frown}{\Box} \text{PCI}_5
$$

partial pressures of Cl₂, PCl₃ and PCl₅ at equilibrium are 0.1, 0.1 and 0.2 atm respectively at 250°C. At the same temperature, in another experiment on the same reaction, at equilibrium the partial pressures of PCI $_{_3}$ and CI $_{_2}$ are half those in the first experiment. The partial pressure of the PCI₅ at equilibrium in the second experiment is

```
[1] One–fourth of the first [2] Half of the first [3] One–eight of the first [4] One–third of the first
```
-
- **Q.44** At a given temperature the following reaction is allowed to reach equilibrium in a vessel of volume V₁ litre. The degree of dissociation is $\alpha_{_1}$. If by keeping the temperature fixed the volume of the reaction vessel is doubled (assuming the degrees of dissociation to be small) the new degree of dissociation shall be

$$
\mathsf{PCI}_5^\perp \boxplus \boxplus \mathsf{PCI}_3 + \mathsf{CI}_2
$$

$$
[1] 2\alpha_{1}
$$

 $[2] \sqrt{\frac{u_1}{2}}$ α

[3] $\sqrt{2\alpha_1}$

 $\sqrt{2}$ α .

Q.45 For the general reaction

^A nB

the degree of dissociation is measured from vapour density measurements. If the degree of dissociation, the observed and theoretical vapour densities be α , $\mathsf{d}_{_{\mathrm{O}}}$ and $\mathsf{d}_{_{\mathrm{f}}}$ respectively, then

$$
[1] \alpha = \frac{d_0 - d_t}{d_t (n+1)}
$$
\n
$$
[2] \alpha = \frac{d_0 - d_t}{d_t}
$$
\n
$$
[3] \alpha = \frac{(d_0 - d_t)(n-1)}{d_t}
$$
\n
$$
[4] \alpha = \frac{d_0 - d_t}{d_0 (n-1)}
$$

Q.46 The correct expression for equilibrium constant K_c, for the reaction

H₂(g) +
$$
\frac{1}{2}
$$
O₂(g) H₂O (g) is
\n[1] [H₂O] / [H₂] [L₂] [2] P<sub>H₂O / P₀^{1/2} P₀² P<sub>H₂³]
\nQ.47 The reversible reaction
\nCu(NH₃)₄² + SO₃² Cu (NH₃)₃ SO₃ + NH₃
\nis at equilibrium. What would not happen if ammonia is added –
\n[1] [SO₃²] would increase
\n[3] The value of equilibrium constant would not change [4] [Cu (NH₃)₃⁵ [would increase]
\n[3] The number of moles in the time of the calculation 1₂ + 2S₂O₃² — 21 + S₄O₆² the correct statement is
\n[1] Addition of potassium iodide would lead to depending of the violet colour
\n[2] Addition of EACI₂ will make the reaction faster due to removal of I as Bal₂
\n[3] The number of moles of tertanhisculphate formed at the end of the reaction would be equal to half the moles of thisculphate for the case of the reaction would be equal to half the moles of thisculphate in the case of the reaction (2) and 2.49
\nAt a certain temperature, K_c for
\nSO₂(g) + NO₂(g) SO₃(g) + NO(g) is 16
\nIf we take one mole each of all the four gases, what will be equilibrium concentration of NO₂ and NO respectively
\n[1] 0.4, 1.6 (2) 0.6, 0.6 (3) 0.6, 4 (4) 0.4, 0.4
\nQ.50 The reaction between thiocyanate and ferric ion is represented by the reaction
\nSCN⁻(aq) + Fe⁻³ (aq)
\nEF (SCN)⁻²
\nColourless yellow deep red
\nAddition of triocyanate ion to the equilibrium mixture will make
\n[1] The solution more deep red
\n[2</sub></sub>

[3] The solution yellow [4] The concentration of the complex ion to decrease

Answer Key

CHEMICAL EQUILIBRIUM

Exercise # 3

[1] Temperature and pressure [2] Temperature only **[IIT-1999]**

[3] Pressure only **[4] Temperature, pressure and catalyst**

