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CHAPTER

CHEMICAL EQUILIBRIUM

The proportion of ingredients is important, but the final result is also a matter of how you put them together. Equilibrium is key.

"ALAIN DUCASSE"

INTRODUCTION

hemical equilibrium are important in numerous biological and environmental processes. For example, equilibrium involving O_2 molecules and the protein hemoglobin play a crucial role in the transport and delivery of O_2 from our lungs to our muscles. Similar equilibrium involving CO moleucles and hemoglobin account for the toxicity of CO. State of chemical equilibrium, these may be classified in three groups.

- (i) The reactions that proceed nearly to completion and only negligible concentrations of the reactants are left. In some cases., it may not be even possible to detect these experimentally.
- (ii) The reactions in which only small amounts of products are formed and most of the reactants remain unchanged at equilibrium state
- (iii) The reactions in which the concentrations of the reactants and products are comparable. When the system is in equilibrium.



Derivation of equilibrium constant : Consider a reversible homogeneous chemical reaction which has attained equilibrium

state at a particular temperature : $A + B \xrightarrow{} C + D$

Let the active masses of A, B, C and D be [A] [B] [C] and [D] respectively at equilibrium.

According to law of mass action :

Rate of forward reaction $(R_f) \propto [A][B]$ Rate of backward reaction $(R_b) \propto [C][D]$

 $R_{c} = k_{c}[A][B]$ and $R_{b} = k_{b}[C][D]$

Where k_f and k_b are forward and backward rate or velocity constants respectively. At equilibrium state :

$$R_{f} = R_{b}$$

$$k_{f}[A][B] = k_{b}[C][D]$$

$$\frac{k_{f}}{k_{b}} = \frac{[C][D]}{[A][B]}$$

$$\mathbf{K}_{\mathrm{C}} = \frac{\begin{bmatrix} \mathbf{C} \end{bmatrix} \begin{bmatrix} \mathbf{D} \end{bmatrix}}{\begin{bmatrix} \mathbf{A} \end{bmatrix} \begin{bmatrix} \mathbf{B} \end{bmatrix}}; \qquad \qquad \ddots \quad \mathbf{K}_{\mathrm{C}} = \frac{\mathbf{k}_{\mathrm{f}}}{\mathbf{k}_{\mathrm{b}}}$$

 K_c is known as equilibrium constant. K_c has a definite value for every chemical reaction at particular temperature.

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PHYSICS FOR NEET & AIIMS



	SOLVED EX	AMPLE	
Ex. 1	When 1.0 mole of N ₂ and 3.0 moles of H ₂ was heated in converted into NH ₃ at equilibrium. Find the value of K ₁ (A) 3.1×10^{-2} atm ⁻² (B) 4.1×10^{-2} atm ⁻²	n a vessel at 873 K and a pressure of 3.55 atr p for the reaction. (C) 5.1×10^{-2} atm ⁻² (D) 6.1×10^{-2} atm	m. 30% of N_2 is
Sol.	$(\mathbb{C}) \qquad N_{2}(g) + 3H_{2}(g) 2NH_{3}(g)$ $1 \text{ mole} 3 \text{ moles} 0$ $1-0.3 \qquad 3.0-0.9 \qquad 0.6 \text{ mole}$ $= 0.7 \text{ moles} = 2.1 \text{ moles}$ $\text{Total no. of moles at equilibrium} = 3.4$ $K_{p} = \frac{\left(\frac{0.6}{3.4} \times 3.55\right)^{2}}{\left(\frac{0.7}{3.4} \times 3.55\right)\left(\frac{2.1}{3.4} \times 3.55\right)^{3}} = 5.1 \times 10^{-2} \text{ atm}^{-2}$	Initial moles les at equilibrium	
Ex. 2 Sol.	$2SO_{2}(g) + O_{2}(g) \rightleftharpoons 2SO_{3}(g)$ If the partial pressure of SO ₂ , O ₂ and SO ₃ are 0.559, 0. pressure of O ₂ gas, to get equal moles of SO ₂ and SO ₃ . (A) 0.188 atm (B) 0.288 atm	.101 and 0.331 atm respectively. What woul (C) 0.388 atm (D) 0.488 atm	d be the partial
	$2SO_{2}(g) + O_{2}(g) \iff 2SO_{3}(g)$ $K_{p} = \frac{[P_{SO_{3}}]^{2}}{[P_{SO_{2}}]^{2}[P_{O_{2}}]} = \frac{(0.331)^{2}}{(0.559)^{2}(0.101)} = 3.47$ If SO ₂ and SO ₃ have same number of moles, their partial $P_{SO_{3}} = P_{SO_{2}} \therefore P_{O_{2}} = \frac{1}{3.47} = 0.288$ atm	al pressure will be equal and	
Ex. 3	K _p for the reaction N ₂ + 3H ₂ \implies 2NH ₃ at 400°C is 3 (A) 0.3 mole ⁻² litre ² (B) 0.4 mole ⁻² litre ²	0.28×10^{-4} . Calculate K _c . (C) 1.0 mole ⁻² litre ² (D) 0.6 mole ⁻² litre	e ²
Sol.	(C) $N_2 + 3H_2 = 2NH_3$ $\Delta n = -2$ and $K_p = K_c (RT)^{\Delta n}$ $3.28 \times 10^{-4} = K_c (0.0821 \times 673)^{-2}$ and $K_a = 1.0$ mole ⁻² litre ² .		
Ex. 4	A mixture of H, and I, in molecular proportion of $2:3$ was heated at 444°C till the reaction		
	H ₂ + I ₂ \longrightarrow 2HI reached equilibrium state. Calculat (K_c at 444°C is 0.02) (A) 3.38 % (B) 4.38%	te the percentage of iodine converted into HI (C) 5.38% (D) 6.38%	
Sol.	(C) H ₂ + I ₂ \longrightarrow 2HI Initial moles 2 3 0 Equi.conc. $\frac{2-x}{v} = \frac{3-x}{v} = \frac{2x}{v}$ $K_c = \frac{4x^2}{(2-x)(3-x)} = 0.02$		
	x = 0.1615 Out of 3 moles, 0.1615 moles I ₂ is converted into HI.		
	$\therefore \text{ Percentage of I}_2 \text{ converted to HI} = \frac{0.1615 \times 100}{3} = 5$	5.38%	
-)			

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	Exercise # 1	SINGLE OB.	JECTI	VE N	NEET LEVEL
1.	According to law of reaction is proportion (A) Concentration of (B) Molar concentrat (C) Concentration of (D) Molar concentrat	mass action rate of a chemical nal to f reactants tion of reactants f products tion of products	8.	4 moles of A are r equilibrium for the moles of C and D constant for the reac (A) $\frac{1}{4}$ (C) 1	nixed with 4 moles of B. At reaction $A + B \rightleftharpoons C + D$, 2 are formed. The equilibrium ction will be (B) $\frac{1}{2}$ (D) 4
2.	In a reaction the rate its active mass, this (A) Law of mass acti (B) Le-chatelier prin (C) Faraday law of e (D) Law of constant	e of reaction is proportional to statement is known as on ciple lectrolysis proportion	9.	On a given condition of HI, H ₂ and I ₂ are The equilibrium H ₂ + I ₂ \rightleftharpoons 2HI wil (A) 64 (C) 8	a, the equilibrium concentration 0.80, 0.10 and 0.10 mole/litre. constant for the reaction 1 be (B) 12 (D) 0.8
3.	The active mass of 6 would be (A) 2 (C) 5	(B) 1 (D) 0.25	10.	In which of the foll towards completion (A) $K = 10^2$ (C) $K = 10$	lowing, the reaction proceeds (B) $K = 10^{-2}$ (D) $K = 1$
4.	Under a given set of increase in the conc rate of a chemical rea (A) Decreases (B) Increases	experimental conditions, with entration of the reactants, the action	11.	A reversible chemica in equilibrium. If the are doubled, then the (A) Also be doubled (C) Become one-fou	al reaction having two reactants e concentrations of the reactants e equilibrium constant will (B) Be halved rth (D) Remain the same
	(C) Remains unalter(D) First decreases a	ed and then increases	12.	The equilibrium con a given temperature (A) Depends on th	stant in a reversible reaction at e initial concentration of the
5.	The law of mass act (A) Guldberg and W (C) Birthelot	ion was enunciated by aage (B) Bodenstein (D) Graham		reactants (B) Depends on the at equilibrium (C) Does not depend	concentration of the products
6.	For the system 3A - equilibrium constan	$+2B \rightleftharpoons C$, the expression for t is [C]	13.	(D) It is not character Pure ammonia is pla where its dissociation	aced in a vessel at temperature on constant (A) is appreciable.
	(A) $\frac{[A]^3 [B]^2}{C}$ (C) $\frac{[A]^3 [B]^2}{[C]}$	(B) $\frac{1}{[3A][2B]}$ (D) $\frac{[C]}{[A]^3[B]^2}$		(A) K_p does not chan (B) α does not chan (C) Concentration α pressure	nge significantly with pressure ge with pressure of NH_3 does not change with
7.	In the reversible reconcentration of eac 0.8 mole/litre, then the be (A) 6.4	eaction $A + B \rightleftharpoons C + D$, the h C and D at equilibrium was ne equilibrium constant K _e will (B) 0.64	14.	(D) Concentration of For the system equilibrium concent (B) 0.12 mole/litre (0 the reaction is (A) 250	A(g) + 2B(g) \rightleftharpoons C(g), the trations are (A) 0.06 mole/litre C) 0.216 mole/litre. The K _{ep} for (B) 416
	(C) 1.6	(D) 16.0	sindia	(C) $K \times 10^{-3}$	(D) 125

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CHEMICAL EQUILIBRIUM

Exercise # 2 SINGLE OBJECTIVE AIIMS LEVEI 5. One litre of 2M acetic acid and one litre of 3M ethyl Equilibrium constant for the reactions, alcohol are mixed to form ester according to the given $2 \text{ NO} + \text{O}_2 \implies 2 \text{ NO}_2$ is K_{C₁}; equation : $CH_{COOH} + C_{H_{c}OH} \longrightarrow CH_{COOC} + H_{O}O.$ $NO_2 + SO_2 \implies SO_3 + NO$ is K_{C_2} and If each solution is diluted by adding equal volume (1 litre) of water by how many times the initial forward $2 \operatorname{SO}_3 \Longrightarrow 2 \operatorname{SO}_2 + \operatorname{O}_2$ is K_{C_3} then correct rate is reduced? reaction is : (A) 4 times (B) 2 times (A) $K_{C_3} = K_{C_4} \times K_{C_5}$ (\mathbb{C}) 0.5 times (D) 0.25 times (B) $K_{C_3} \times K_{C_1} \times K^2_{C_2} = 1$ 6. In the dissociation of N_2O_4 into NO_2 , $(1 + \alpha)$ values with the vapour densities ratio $\left(\frac{D}{d}\right)$ is as given by (C) $K_{C_3} \times K_{C_1} \times K_{C_2} = 1$ (D) $K_{C_3} \times K^2_{C_1} \times K_{C_2} = 1$ $\left[\alpha - \text{degree of dissociation, D-vapour density before}\right]$ At a certain temperature, the following reactions dissociation, d-vapour density after dissociation] have the equilibrium constant as shown below : $S(s) + O_{2}(g) \implies SO_{2}(g); K_{2} = 5 \times 10^{52}$ $(1+\alpha)$ $2S(s) + 3O_{2}(g) \implies 2SO_{3}(g); K_{2} = 10^{29}$ **(A) (B)** What is the equilibrium constant K_a for the reaction at the same temperature? (1+α) $2SO_{2}(s) + O_{2}(g) \rightleftharpoons 2SO_{3}(g)$ (C) (D) (A) 2.5×10^{76} **(B)** 4×10^{23} (C) 4×10^{-77} (D) None of these In the above question, α varies with $\frac{D}{d}$ according Sulfide ion in alkaline solution reacts with solid sulfur 7. to form polysulfide ions having formulae S_2^{2-} , S_3^{2-} , to : S_4^{2-} and so on. The equilibrium constant for the

1.

2.

3.

4.

 S_2^{2-} and S? (A) 11

the K_c for the equilibrium

(C)132

is: (A)4

(C)9





(A) $d_{20} > d_{45} > d_{65} > d_{80}$
B) $d_{80} > d_{65} > d_{45} > d_{20}$
C) $d_{20} = d_{45} = d_{65} = d_{80}$
D) $(d_{20} = d_{45}) > (d_{65} = d_{80})$

 $+ H_2O(\ell)$

formation of S_2^{2-} is $K_1 = 12$ & for the formation of S_3^{2-} is $K_2 = 132$, both from S and S²⁻. What is the

equilibrium constant for the formation of S_3^{2-} from

When alcohol ($C_2H_5OH(\ell)$) and acetic acid ($CH_3COOH(\ell)$) are mixed together in equimolar ratio at 27°C, 33% of each is converted into ester. Then

 $C_{H_{5}}OH(\ell) + CH_{3}COOH(\ell) \Longrightarrow CH_{3}COOC_{H_{5}}(\ell)$

(B) 12

(B) 1/4

(D) 1/9

(D) None of these

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8.

PHYSICS FOR NEET & AIIMS

I	Exercise # 3 PART - 1	MATRIX MATCH COLUMN
1.	Match the following : Column I (Assume only reactant were present initially) (A) For the equilibrium	Column II (n) Forward shift
	(A) For the equilibrium $NH_4I(s) \Longrightarrow NH_3(g) + HI(g),$ if pressure is increased at equilibrium (B) For the equilibrium $N_2(g) + 3H_2(g) \Longrightarrow 2NH_2(g)$	(q) No shift in equilibrium
	volume is increased at equilibrium (C) For the equilibrium $H_2O(g) + CO(g) \Longrightarrow H_2(g) + CO_2(g)$ inert gas is added at constant pressure at equilibrium	(r) Backward shift
	(D) For the equilibrium $PCl_5 \implies PCl_3 + Cl_2$ Cl_2 is removed at equilibrium.	(s) Final pressure is more than initial pressure
2.	Match the following : (Assume only reactants were	e present initially).
	(A) $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g) (t = 300^{\circ}C)$ (B) $PCl_5(g) \rightleftharpoons PCl_3(g) + Cl_2(g) (t = 50^{\circ}C)$ (C) $C(s) + H_2O(g) \rightleftharpoons CO(g) + H_2(g)$	(p) $\Delta n_g > 0$ (q) $K_p < K_c$ (r) K_p not defined
	(D) $CH_3COOH(\ell) + C_2H_3OH(\ell) \Longrightarrow$ $CH_3COOC_2H_5(\ell) + H_2O(\ell)$	(s) $P_{initial} > P_{eq.}$
3.	Column-I (Reactions) (A) Oxidation of nitrogen	Column-II (Favourable conditions) (p) Addition of inert gas at constant pressure
	$N_2(g) + O_2(g) + 180.5 \text{ kJ} \implies 2NO(g)$ (B) Dissociation of $N_2O_4(g)$	(q) Decrease in pressure
	$N_2O_4(g) + 57.2 \text{ kJ} \implies 2NO_2(g)$ (C) Oxidation of NH ₃ (g)	(r) Decrease in temperature
	$4NH_3(g) + 5O_2(g) \implies 4NO(g) + 6H_2O(g) + (D)$ Formation of $NO_2(g)$	- 905.6 kJ (s) Increase in temperature
	$NO(g) + O_3(g) \implies NO_2(g) + O_2(g) + 200 k$	кJ
4.	Column-I (Reaction)	Column-II (If α is negligiable w.r.t. 1)
	(A) $2X(g) \rightleftharpoons Y(g) + Z(g)$	(p) $\alpha = 2 \times \sqrt{K_c}$
	(B) $X(g) \rightleftharpoons Y(g) + Z(g)$	(q) $\alpha = 3 \times \sqrt{K_c}$
	$(\mathbb{C}) 3X(g) \rightleftharpoons Y(g) + Z(g)$	(r) $\alpha = (2K_c)^{1/3}$
	$(\mathbb{D}) 2X(g) \rightleftharpoons Y(g) + 2Z(g)$	(s) $\alpha = \sqrt{K_c}$
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CHEMICAL EQUILIBRIUM

Exercise # 4 PART - 1

- For a reversible reaction, if the concentrations of 6. the reactants are doubled, the equilibrium constant will be [CBSE AIPMT 2000]
 (A) one-fourth (B) halved
 - (C) doubled (D) the same
- 2. For the equilibrium,

 $MgCO_3(s) \xrightarrow{\Delta} MgO(s) + CO_2(g)$ which of the following expressions is correct? [CBSE AIPMT 2000]

(A)
$$K_p = p_{CO_2}$$
 (B) $K_p = \frac{[MgO][CO_2]}{[MgCO_3]}$

(C)
$$K_p = \frac{p_{MgO} \cdot p_{CO_2}}{p_{MgCO_3}}$$
 (D) $K_p = \frac{p_{MgO} + p_{CO_2}}{p_{MgCO_3}}$

- 3. Reaction, $BaO_2(s) \rightleftharpoons BaO(s) + O_2(g), \Delta H = + ve$ In equilibrium condition, pressure of O_2 depends on [CBSE AIPMT 2002]
 - (A) increased mass of BaO_2
 - (B) increased mass of BaO
 - (C) increased temperature of equilibrium
 - (D) increased mass of BaO_2 and BaO both
- 4. The reaction quotient (Q) for the reaction, $N_2(g) + 3H_2(g) \rightleftharpoons 2NH_3(g)$ is given by

 $Q = \frac{[NH_3]^2}{[N_2][H_2]^3}.$

The reaction will proceed towards right side, if [CBSE AIPMT 2003]

(A) $Q > K_c$	$(\mathbf{B})\mathbf{Q}=0$
$(\mathbb{C}) Q = K_{c}$	$(\mathbb{D})Q \leq K_{c}$
where, K is the equil	ibrium constant.

5. In the two gaseous reaction (i) and (ii) at 250°C

(i) NO(g) +
$$\frac{1}{2}$$
O₂(g) \rightleftharpoons NO₂(g), K

(ii) $2NO_2(g) \rightleftharpoons 2NO(g) + O_2(g), K_2$ the equilibrium constants K_1 and K_2 are related as [CBSE AIPMT 2005, 1994]

(A)
$$K_2 = \frac{1}{K_1}$$
 (B) $K_2 =$

(C)
$$K_2 = \frac{1}{K_1^2}$$
 (D) $K_2 = K_1^2$

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7.

8.

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10.

 $CH_4(g) + 2O_2(g) \rightleftharpoons CO_2(g) + 2H_2O(l),$

 $\Delta r H = -170.8 \, kJ \, mol^{-1}$

Which of the following statement is not true?

[CBSE AIPMT 2006]

- (A) At equilibrium, the concentrations of CO_2 (g) and $H_2O(l)$ are not equal
- (B) The equilibrium constant for the reaction is

given by
$$k_p = \frac{[CO_2]}{[CH_4][O_2]}$$

- (C) Addition of CH4(g) or O2(g) at equilibrium will cause a shift to the right
- (D) The reaction is exothermic
- The value of equilibrium constant of the reaction,

$$HI(g) \rightleftharpoons \frac{1}{2}H_2(g) + \frac{1}{2}I_2(g) \text{ is 8.0.}$$

The equilibrium constant of the reaction, $H_2(g) + I_2(g) \rightleftharpoons 2HI(g)$ will be

[CBSE AIPMT 2008]

(A) $\frac{1}{16}$	(B) $\frac{1}{64}$
(C) 16	(D) $\frac{1}{8}$

If the concentration of OH⁻ ions in the reaction, $Fe(OH)_3(s) \rightleftharpoons Fe^{3+}(aq) + 3OH^-(aq)$ is decreased by 1/4 times, then equilibrium concentration of Fe3⁺ will increase by [CBSE AIPMT 2008]

(A) 8 times	(B) 16 times
(C) 64 times	(D) 4 times

The dissociation constants for acetic acid and HCN at 25°C are 1.5×10^{-5} and 4.5×10^{-10} , respectively. The equilibrium constant for the equilibrium, $CN^-+CH_2COOH \rightleftharpoons HCN+CH_3COO^$ would be

	[CBSE AIPMT 2009]
(A) 3.0×10^5	(B) 3.0×10^{-5}
(\mathbb{C}) 3.0 × 10 ⁻⁴	(D) 3.0×10^4

In which of the following equilibrium K_c and K_p are not equal ? [CBSE AIPMT 2010] (A) $2NO(g) \rightleftharpoons N_2(g) + O_2(g)$ (B) $SO_2(g) + NO_2(g) \rightleftharpoons SO_3(g) + NO(g)$ (C) $U_1(g) + U_2(g) \Rightarrow 2UV(g)$

 $(\mathbb{C}) \operatorname{H}_{2}(g) + \operatorname{l}_{2}(g) \rightleftharpoons 2\operatorname{HI}(g)$

$$(D) 2C(s) + O_2(g) \rightleftharpoons 2CO_2(g)$$

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 $K_1^{1/2}$

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MOCK TEST

STRAIGHT OBJECTIVE TYPE

1. Solid ammonium carbamate dissociates to give ammonia and carbon dioxide as follows.

 $NH_2COONH_4(s) \rightleftharpoons 2NH_3(g) + CO_2(g)$ At equilibrium, ammonia is added such that partial pressures of NH_3 now equals the original totoal pressure. Calculate the ratio of the total pressure now to the original total pressure.

(A)
$$\frac{31}{27}$$
 (B) $\frac{60}{40}$ (C) $\frac{31}{9}$ (D) $\frac{62}{27}$

2. In the Haber process for the industrial manufacture of ammonia involving the reaction,

 $N_2 + 3H_2 \rightleftharpoons 2NH_3$ at 200 atm pressure in the presence of a catalyst, a temperature of about 500°C is used. This is considered as optimum temperature for the process because

- (A) yield is maximum at this temperature
- (B) catalyst is active only at this temperature
- (C) energy needed for the reaction is easily obtained at this temperature
- (D) rate of the catalytic reaction is fast enough while the yield is also appreciable for this exothermic reaction at this temperature.

3. For the equilibrium of the reaction, HgO(s) \rightleftharpoons Hg(g) + $\frac{1}{2}$ O₂(g), k_p for the reaction at total pressure of P is :

(A)
$$K_p = \frac{2}{3^{3/2}} p^{3/2}$$
 (B) $K_p = \frac{2}{3^{1/2}} p^{1/2}$ (C) $K_p = \frac{1}{3^{2/3}} p^{3/2}$ (D) $K_p = \frac{1}{3^{2/3}} p^{3/2}$

4. The average person can see the red colour imparted by the complex $[Fe(SCN)]^{2+}$ to an aqueous solution if the concentration of the complex is 6×10^{-6} M or greater. What minimum concentration of KSCN would be required to make it possible to detect 1 ppm (part per million) of Fe(III) in a natural water sample? The instability constant for

$$Fe(SCN)^{2^{+}} \rightleftharpoons Fe^{3^{+}} + SCN^{-} \text{ is } 7.142 \times 10^{-3}.$$
(A) 0.0036 M (B) 0.0037 M (C) 0.0035 M (D) None of these

 $\frac{1}{2} \operatorname{N}_2(g) + \operatorname{O}_2(g) \rightleftharpoons \operatorname{NO}_2(g)$

 $2NO_2(g) \rightleftharpoons N_2O_4(g)$

Given that above reactions have equilibrium constants K_1 and K_2 respectively. What would be the expression for the equilibrium constant K for the following reaction in terms of K_1 and K_2 ?

....K,

...K.

(A)
$$K_1 K_2$$
 (B) $\frac{1}{K_1 (K_2)^2}$ (C) $\frac{1}{K_2 (K_1)^2}$ (D) $\frac{1}{K_1 K_2}$

6. The value of K_{p} for the reaction at 27°C

$$Br_2(\ell) + Cl_2(g) \rightleftharpoons 2BrCl(g)$$

is '1 atm'. At equilibrium in a closed container partial pressure of BrCl gas ia 0.1 atm and at this temperature the vapour pressure of $Br_2(\ell)$ is also 0.1 atm. Then what will be minimum moles of $Br_2(\ell)$ to be added to 1 mole of Cl_2 , to get above equilibrium situation :

(A)
$$\frac{10}{6}$$
 moles (B) $\frac{5}{6}$ moles (C) $\frac{15}{6}$ moles (D) 2 moles

7. The two equilibria, $AB(aq) \rightleftharpoons A^+(aq) + B^-(aq)$ and $AB(aq) + B^-(aq) \rightleftharpoons AB_2^-(aq)$ are simultaneously maintained in a solution with equilibrium constants, K_1 and K_2 respectively. If $[A^+]$ and $[AB_2^-]$ are y and x respectively, under equilibrium produced by adding the substance AB to the solvents, then K_1/K_2 is equal to

(A)
$$\frac{y}{x}(y-x)^2$$
 (B) $\frac{y^2(x+y)}{x}$ (C) $\frac{y^2(x+y)}{x}$ (D) $\frac{y}{x}(x-y)$

[Note : Use the information of the preceding problem]

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11th Class Modules Chapter Details



PHYSICS

CHEMISTRY

Module-1

- 1. Physical World & Measurements
- 2. Basic Maths & Vector
- 3. Kinematics

Module-2

- 1. Law of Motion & Friction
- 2. Work, Energy & Power

Module-3

- **1.** Motion of system of
- particles & Rigid Body
- 2. Gravitation

Module-4

- 1. Mechanical Properties of Matter
- 2. Thermal Properties of Matter

Module-5

- 1. Oscillations
- 2. Waves

Module-1(PC)

- 1. Some Basic Conceps of Chemistry
- 2. Atomic Structure
- 3. Chemical Equilibrium
- **4.** Ionic Equilibrium

Module-2(PC)

- 1. Thermodynamics & Thermochemistry
- 2. Redox Reaction
- **3.** States Of Matter (Gaseous & Liquid)

Module-3(IC)

- 1. Periodic Table
- 2. Chemical Bonding
- 3. Hydrogen & Its Compounds
- 4. S-Block

Module-4(OC)

- 1. Nomenclature of
- Organic Compounds
- 2. Isomerism
- 3. General Organic Chemistry

Module-5(OC)

- 1. Reaction Mechanism
- 2. Hydrocarbon
- **3.** Aromatic Hydrocarbon
- 4. Environmental Chemistry & Analysis Of Organic Compounds

BIOLOGY

Module-1

- 1. Diversity in the Living World
- 2. Plant Kingdom
- 3. Animal Kingdom

Module-2

- 1. Morphology in Flowering Plants
- **2.** Anatomy of Flowering Plants
- **3.** Structural Organization in Animals

Module-3

- 1. Cell: The Unit of Life
- 2. Biomolecules
- 3. Cell Cycle & Cell Division
- 4. Transport in Plants
- 5. Mineral Nutrition

Module-4

- 1. Photosynthesis in Higher Plants
- 2. Respiration in Plants
- 3. Plant Growth and Development
- 4. Digestion & Absorption
- 5. Breathing & Exchange of Gases

Module-5

- Body Fluids & Its Circulation
 Excretory Products & Their Elimination
- **3.** Locomotion & Its Movement
- 4. Neural Control & Coordination
- **5.** Chemical Coordination and Integration

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12th Class Modules Chapter Details



PHYSICS

Module-1

- 1. Electrostatics
- 2. Capacitance

Module-2

- 1. Current Electricity
- 2. Magnetic Effect of Current and Magnetism

Module-3

- 1. Electromagnetic Induction
- 2. Alternating Current

Module-4

- 1. Geometrical Optics
- 2. Wave Optics

Module-5

- 1. Modern Physics
- 2. Nuclear Physics
- 3. Solids & Semiconductor Devices
- 4. Electromagnetic Waves

CHEMISTRY

Module-1(PC)

- 1. Solid State
- 2. Chemical Kinetics
- **3.** Solutions and Colligative Properties

Module-2(PC)

- 1. Electrochemistry
- 2. Surface Chemistry

Module-3(IC)

- 1. P-Block Elements
- 2. Transition Elements (d & f block)
- 3. Co-ordination Compound
- 4. Metallurgy

Module-4(OC)

- 1. HaloAlkanes & HaloArenes
- Alcohol, Phenol & Ether
 Aldehyde, Ketone &
- Carboxylic Acid

Module-5(OC)

- 1. Nitrogen & Its Derivatives
- 2. Biomolecules & Polymers
- 3. Chemistry in Everyday Life

BIOLOGY

Module-1

- 1. Reproduction in Organisms
- 2. Sexual Reproduction in
- Flowering Plants
- 3. Human Reproduction
- 4. Reproductive Health

Module-2

- **1.** Principles of Inheritance and Variation
- 2. Molecular Basis of Inheritance
- **3.** Evolution

Module-3

- 1. Human Health and Disease
- 2. Strategies for Enhancement in
- Food Production
- 3. Microbes in Human Welfare

Module-4

- **1.** Biotechnology: Principles and Processes
- 2. Biotechnology and Its
- Applications
- 3. Organisms and Populations

Module-5

- 1. Ecosystem
- 2. Biodiversity and Conservation
- 3. Environmental Issues

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