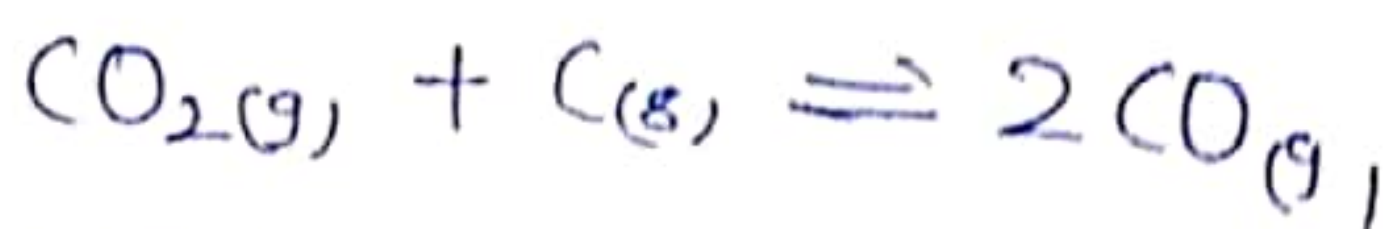


JEE-Main s

Solution

1



$$t=0 \quad 0.5 \text{ atm}$$

$$t=t_{\text{eq}} \quad \begin{array}{l} 0.5-p \\ 0.2 \text{ atm} \end{array}$$

$$\begin{array}{l} 2p \\ 0.6 \text{ atm} \end{array}$$

$$P_T = 0.5 - p + 2p = 0.5 + p = 0.8$$

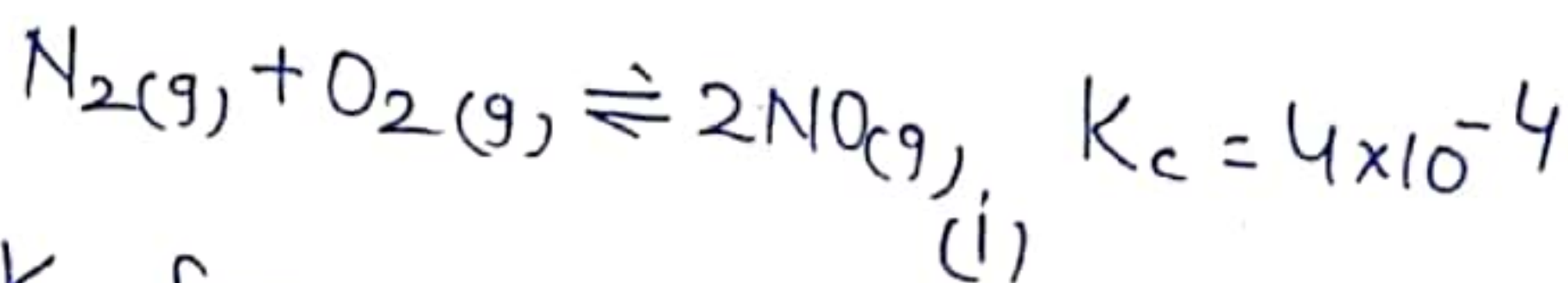
$$p = 0.3 \text{ atm}$$

$$K_p = \frac{P_{\text{CO}}^2}{P_{\text{CO}_2}} = \frac{(0.6)^2}{0.2}$$

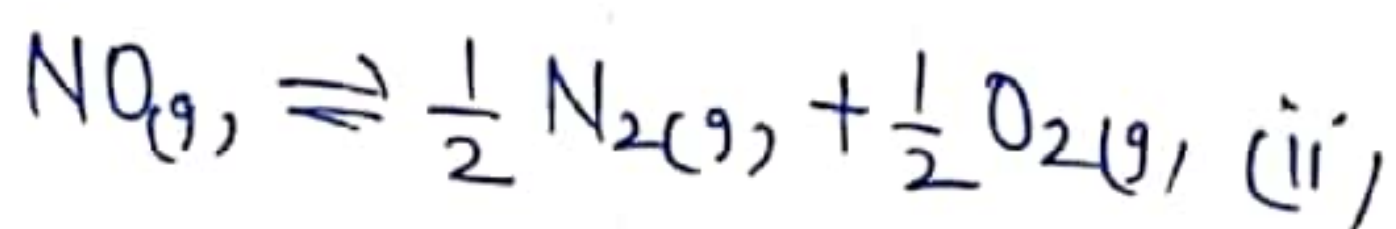
$$= \frac{0.6 \times 0.6}{0.2} = 1.8 \text{ atm}$$

2

For Reaction



then K_c for



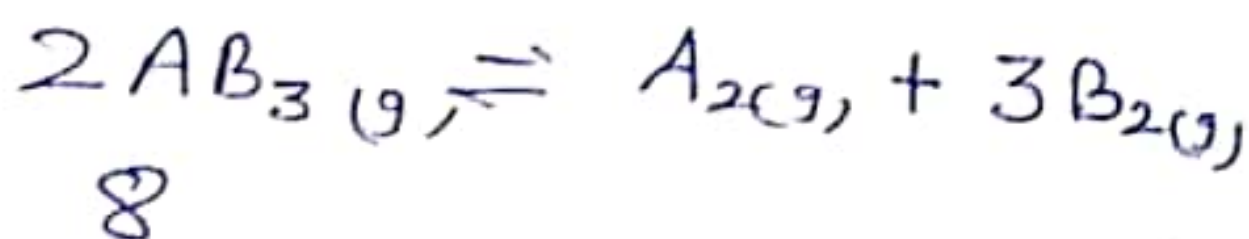
$$\text{reaction (ii)} \quad K_{c2} = \frac{1}{(K_c)^{1/2}}$$

$$K_c = \frac{1}{(4 \times 10^{-4})^{1/2}} = \frac{1}{2 \times 10^{-2}}$$

$$= 50$$

Ans: 3

$t=0$
Mole



$t=t_{eq}$
Mole

$8-2x$
4

x
2

$3x$
6

Mole of $A_2 = x = 2$

$V_L = 12$

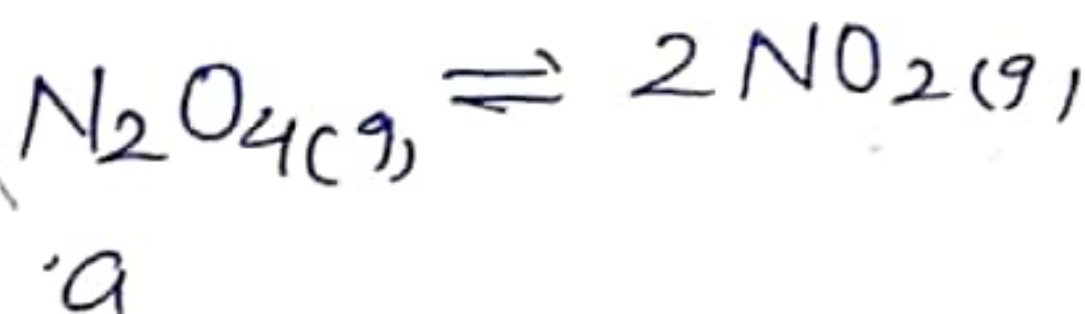
$$K_c = \frac{[A_2][B_2]^3}{[AB_3]^2} = \frac{2(6)^3}{(4)^2}$$

$$K_c = \frac{2 \times 6 \times 6 \times 6 \times 6}{4 \times 4 \times 4}$$

$K_c = 27$

: 4

$t=0$
mole



$t=t_{eq}$

$a-a\alpha$

$2a\alpha$

$\alpha = \text{Degree of dissociation}$

$n_T = a + a\alpha$

$P_T = P$

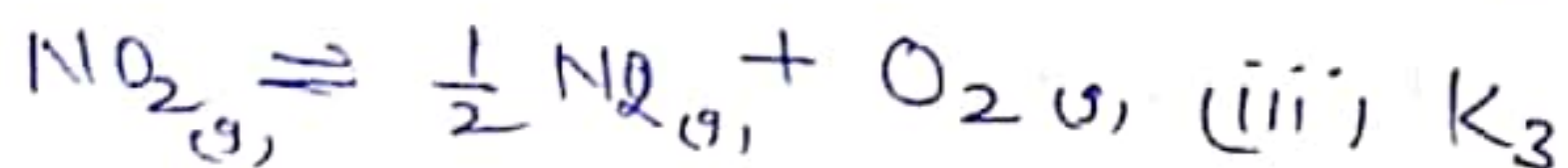
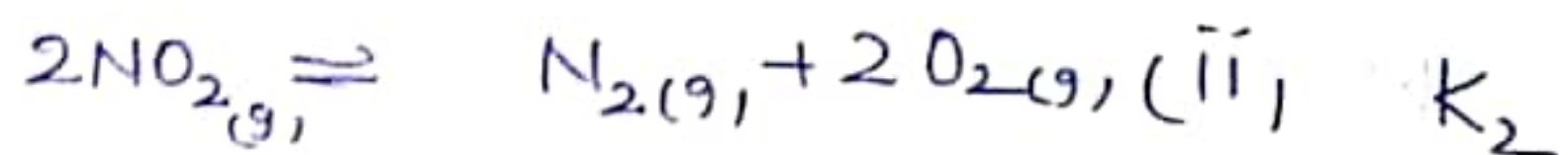
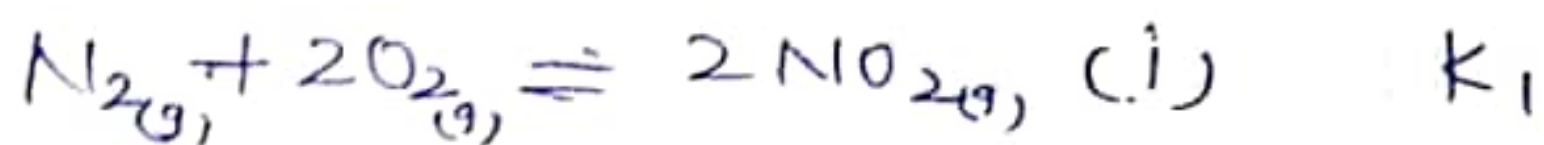
$$K_p = \frac{P_{NO_2}^2}{P_{N_2O_4}} = \frac{\left(\frac{2a\alpha}{a(1+\alpha)}P\right)^2}{\left[\frac{a(1-\alpha)}{a(1+\alpha)}P\right]}$$

$$K_p = \frac{4\alpha^2 P}{1-\alpha^2} = 2 = \frac{4\alpha^2 \times 0.5}{1-\alpha^2}$$

$$\frac{\alpha^2}{1-\alpha^2} = 1 \quad \alpha = \frac{1}{\sqrt{2}} = 0.707$$

$\alpha\% \approx 71\%$

Ans: -5



Reaction (ii) is reverse of (i)

$$K_2 = \frac{1}{K_1}$$

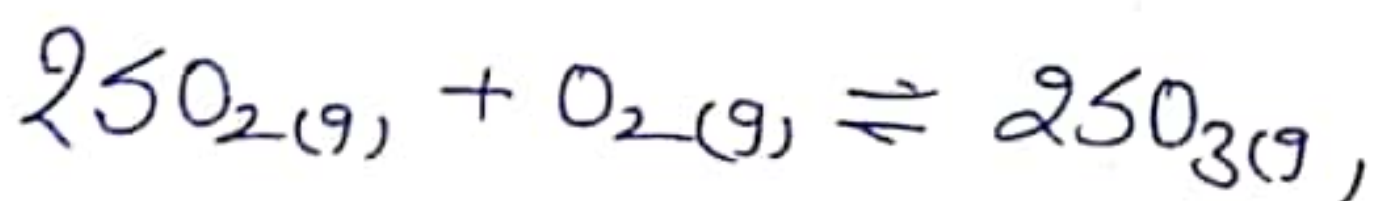
Reaction (iii) is obtained by multiply by $\frac{1}{2}$ and reversed.

$$K_3 = \frac{1}{\sqrt{K_1}}$$

Ans 4

$$K_1 = \frac{1}{K_2} = \frac{1}{K_3^2}$$

Ans: 6



t=0
Mole

2

1

t=t_{eq}
Mole

2-2x
0.4

1-x
0.2

2x
1.6

$$\text{Mole of SO}_3 = 2x = 1.6$$

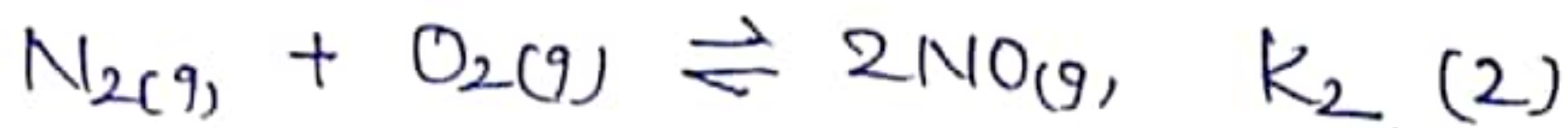
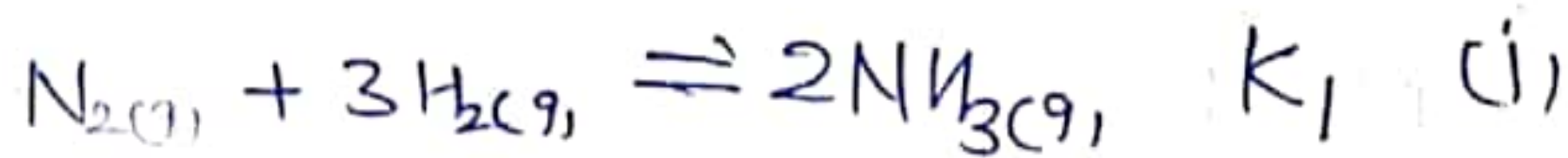
Volume = 1L

$$x = 0.8$$

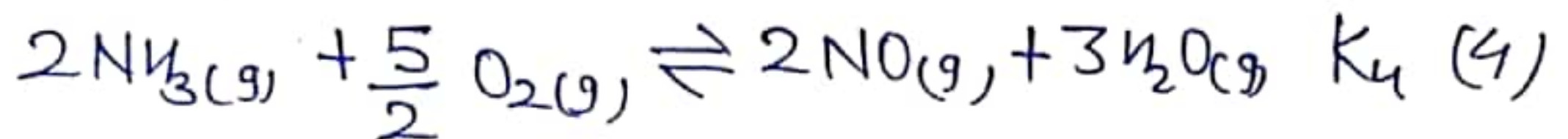
$$K_c = \frac{[\text{SO}_3]^2}{[\text{SO}_2]^2[\text{O}_2]} = \frac{(1.6)^2}{(0.4)^2 \times (0.2)}$$

$$K_c = 80$$

Ans: 7



The equilibrium constant for



Reaction 4 is obtained by

$$(4) = 2 + 3 \times (3) - 1$$

$$K_4 = \frac{K_2 \times K_3^3}{K_1}$$

Ans: 8

For Reaction



$t=0$
Conc.

$$a \quad 1.5a$$

$t=t_{\text{eq}}$
Conc.

$$a-x \quad 1.5a-2x \quad 2x \quad x$$
$$a/2 \quad a/2 \quad a \quad a/2$$

at Eq:-

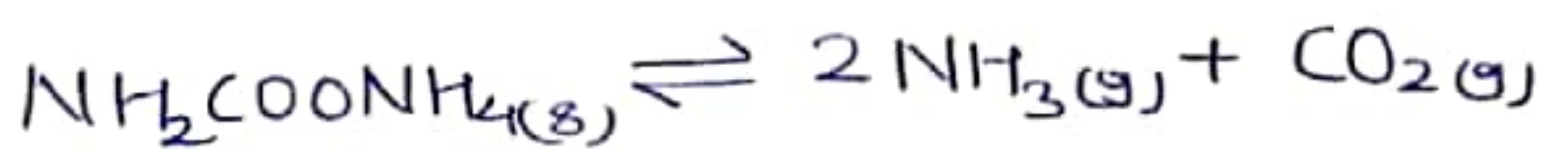
$$[\text{A}] = [\text{B}]$$

$$a-x = 1.5a-2x$$

$$x = a/2$$

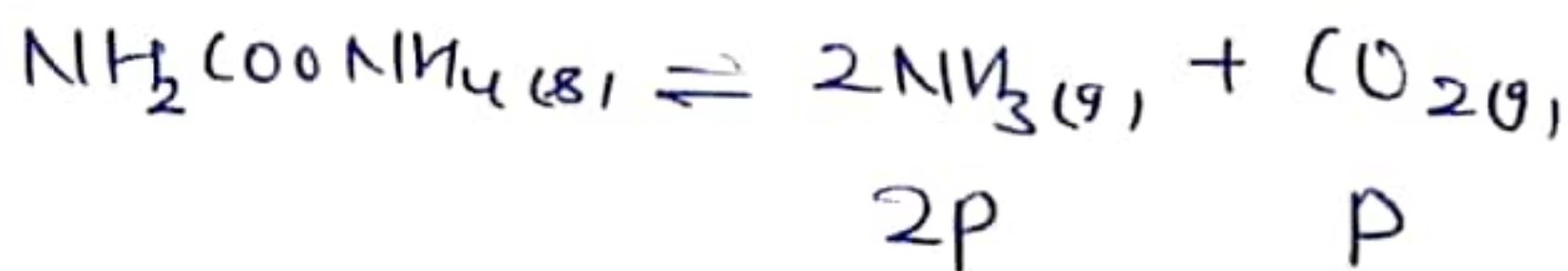
$$K_c = \frac{[\text{C}]^2 [\text{D}]}{[\text{A}] [\text{B}]^2} = \frac{[a]^2 [a/2]}{[a/2] [a/2]^2} = 4$$

Sol. 9



at $t = t_{eq}$ mole Ratio of NH_3 & CO_2 is 2 : 1

So pressure Ratio is 2 : 1



$$K_p = P_{\text{NH}_3}^2 \cdot P_{\text{CO}_2}$$

$$= (2p)^2 p$$

$$= 4p^3 = 2.9 \times 10^{-5} = 29 \times 10^{-6}$$

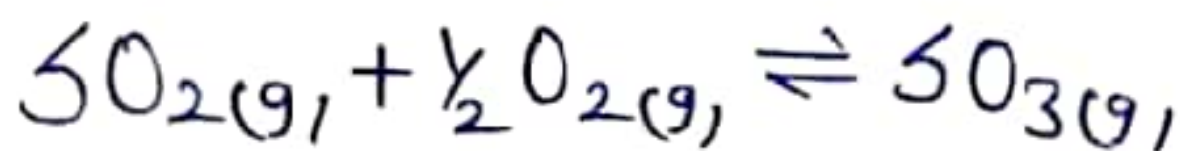
$$p^3 = 7.25 \times 10^{-6}$$

$$p \approx 1.94 \times 10^{-2}$$

$$P_T = 3p$$

$$= 5.82 \times 10^{-2}$$

Solⁿ: 10



Relation b/w K_p & K_c

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

$$\Delta n_g = \sum n_{\text{gas}}(\text{prod}) - \sum n_{\text{gas}}(\text{React})$$

$$\text{For the } \rightarrow \Delta n_g = 1 - [1 + \frac{1}{2}] = -\frac{1}{2}$$

$$K_p = K_c [RT]^{-1/2}$$

Solution-11



$$t=0 \quad 1M \quad 1M \quad 1M \quad 1M \quad ; Q_{\text{initial}} = \frac{[C][D]}{[A][B]} = \frac{1 \times 1}{1 \times 1} = 1$$

(Initially)

$$(Q=1) < (K_c=100)$$

$$t=t_{\text{eq.}} \quad 1-x \quad 1-x \quad 1+x \quad 1+x$$

(Equilibrium)

So reaction proceed
to forward direction

$$K_c = \frac{[C][D]}{[A][B]}$$
$$= \frac{(1+x)(1+x)}{(1-x)(1-x)}$$

$$\Rightarrow 100 = \left(\frac{1+x}{1-x} \right)^2$$

$$\Rightarrow 10 = \frac{1+x}{1-x}$$

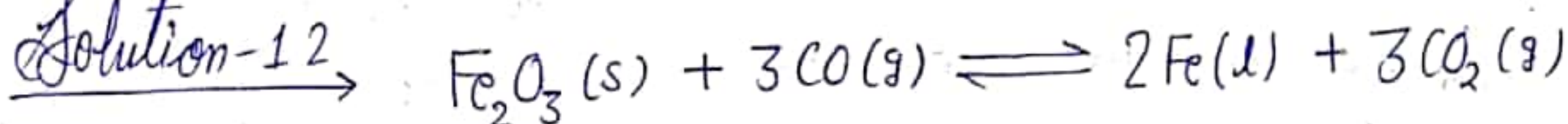
$$\Rightarrow 10 - 10x = 1 + x$$

$$\Rightarrow 11x = 9 \quad \Rightarrow x = \frac{9}{11}$$

$$\text{So } [D] = 1+x = 1 + \frac{9}{11} = \frac{20}{11} = 1.818$$

$$\Rightarrow \boxed{[D] = 1.818 \text{ M}} \leftarrow \text{mole L}^{-1}$$

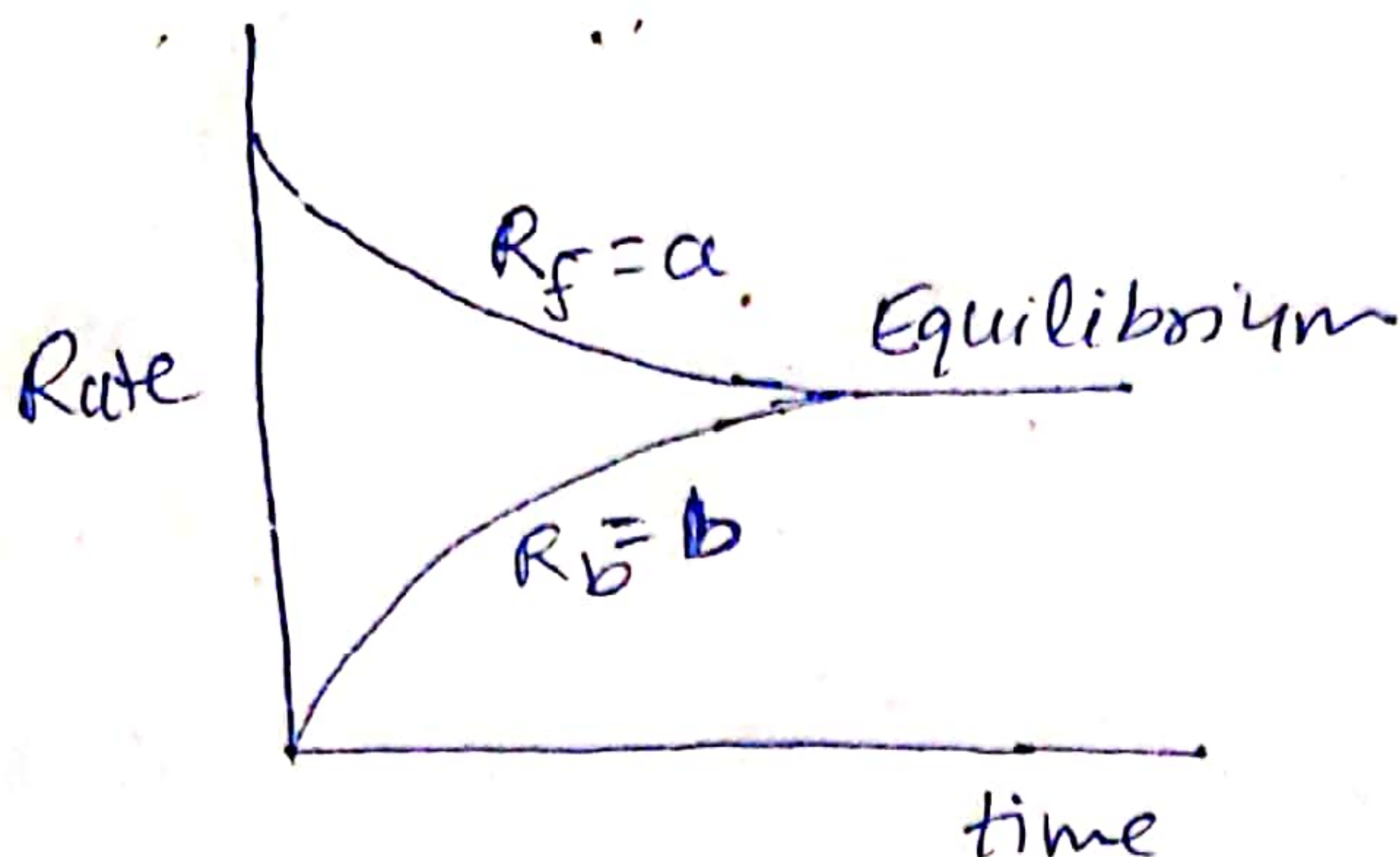
Solution-12



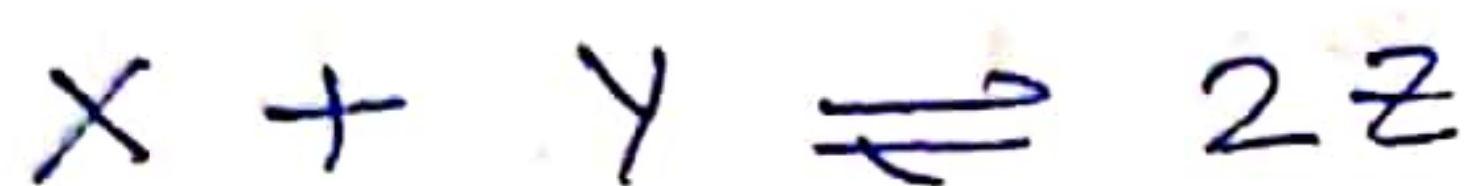
According Le-Chatelier's principle Addition of any solid matter does not affect equilibrium. So In given option Addition of Fe₂O₃ will not disturb the equilibrium.

(13).

We know



(14)



$$t=0 \quad 1 \quad 1.5 \quad 0.5$$

$$Eq \quad 1-x \quad 1.5-x \quad 0.5+2x$$

$$\text{Given } (0.5+2x) = 1$$

$$x = 0.25$$

$$K = \frac{(0.5+2x)^2}{(1-x)(1.5-x)}$$

$$K = \frac{1^2}{0.75 \times 1.25} = \frac{16}{15}$$

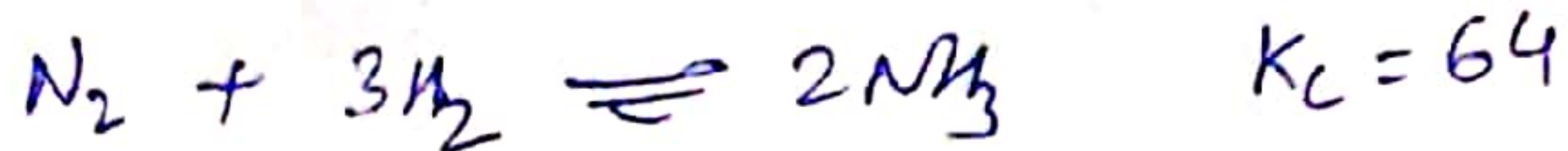
$$\Rightarrow x = 16$$

(15)

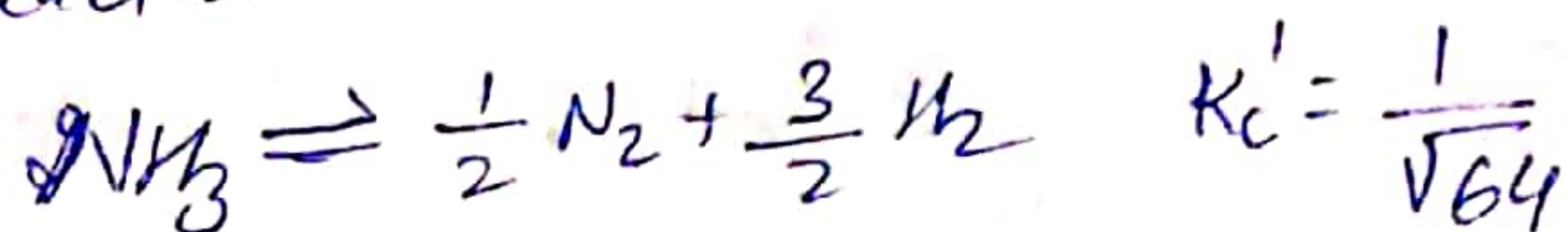
(a) Temperature is decreased
hence for endothermic reaction,
equilibrium shift towards reactant

(b) As N_2 gas (inert gas) is added
at constant volume, hence no
effect on equilibrium.

(16)

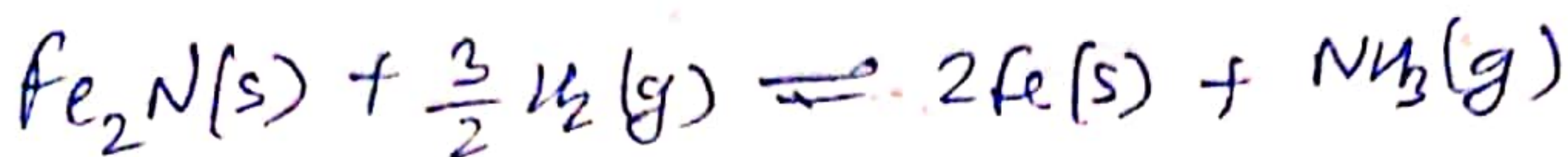


for reaction



$$K_c' = \frac{1}{8}$$

(17)

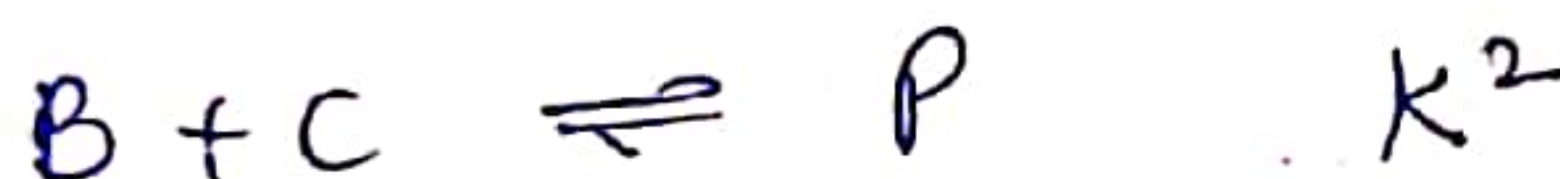
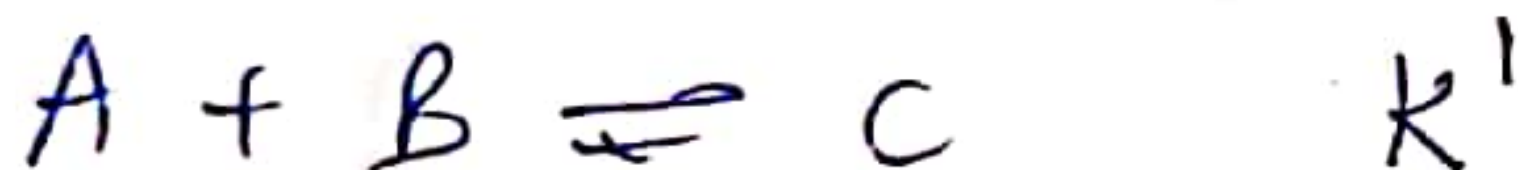


$$\Delta n_g = \left(1 - \frac{3}{2}\right) = -\frac{1}{2}$$

$$\Rightarrow K_p = K_c(RT)^{-1/2}$$

$$\therefore K_c = K_p(RT)^{1/2}$$

(18)



$$K = K^1 \cdot K^2$$

J. Adv.



(a) ΔH is dependent on temperature
as per ~~Kinetic~~ equation

$$\log \left(\frac{K_2}{K_1} \right) = \frac{\Delta H}{2.303R} \left(\frac{1}{T_1} - \frac{1}{T_2} \right)$$

(b) & (c) \rightarrow Equilibrium constant (K)
depends on temperature only.

(d) $\Delta H = (E_{af} - E_{ab})$ and it is
independent of catalyst