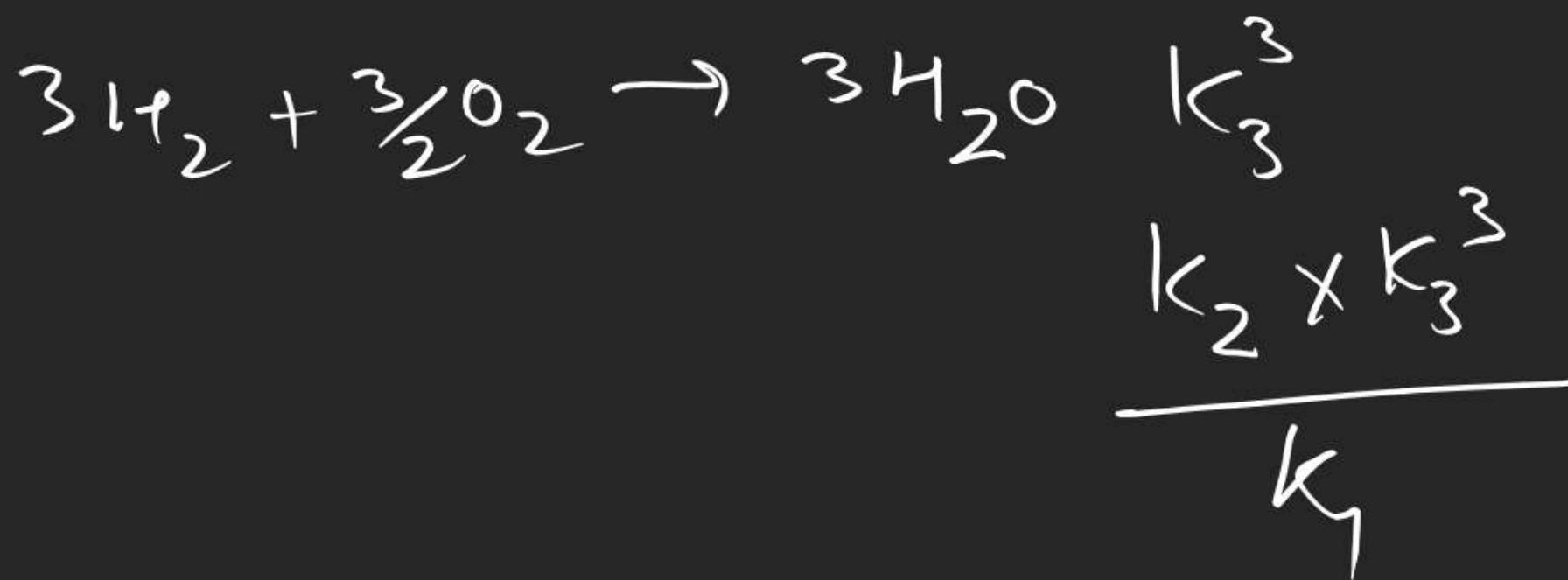
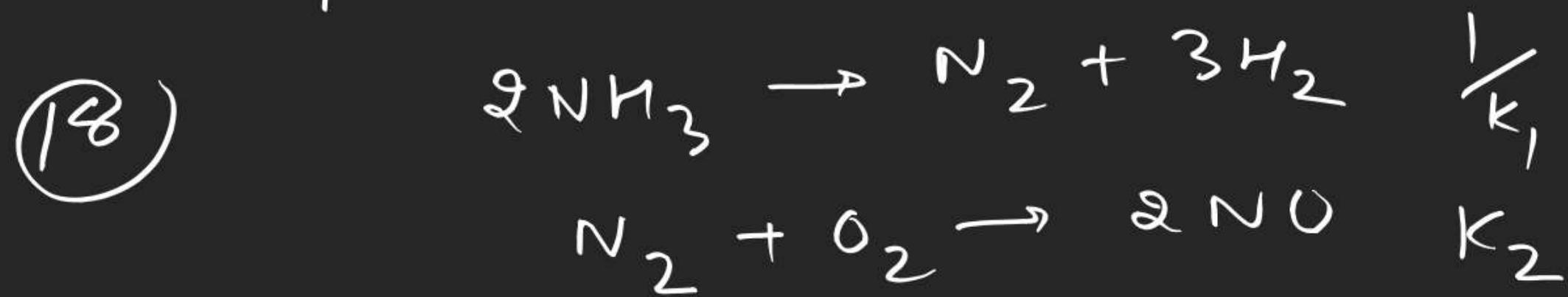
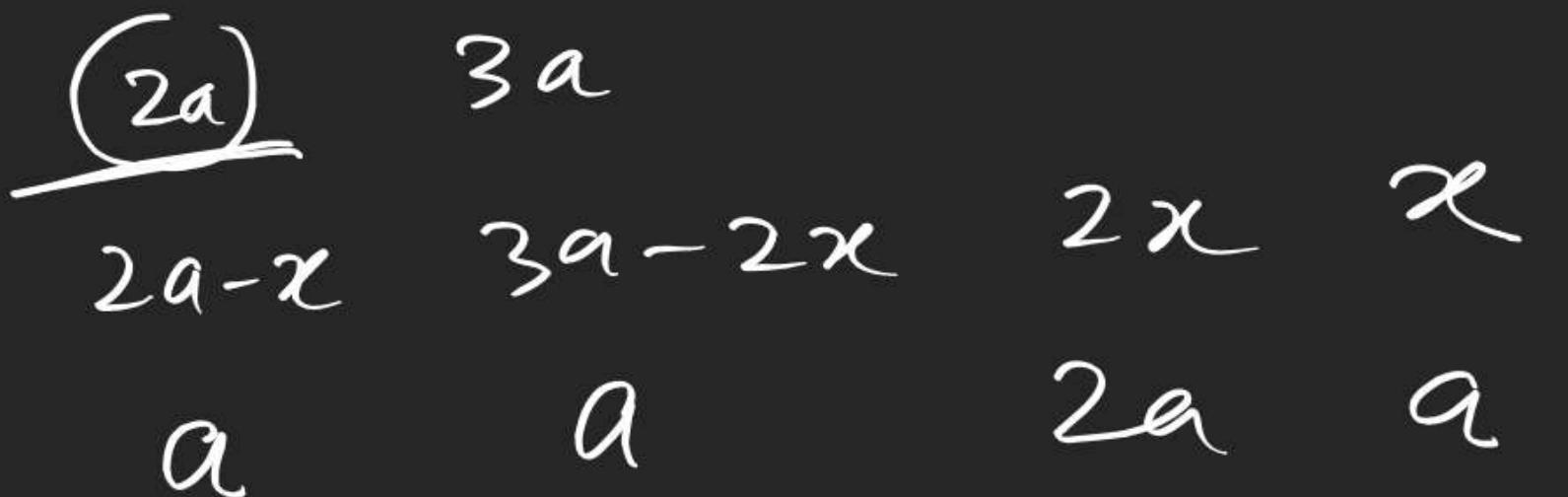


$$0 - \text{I} \quad 16 - 38$$

$$S - \text{I} \quad 11 - 16$$

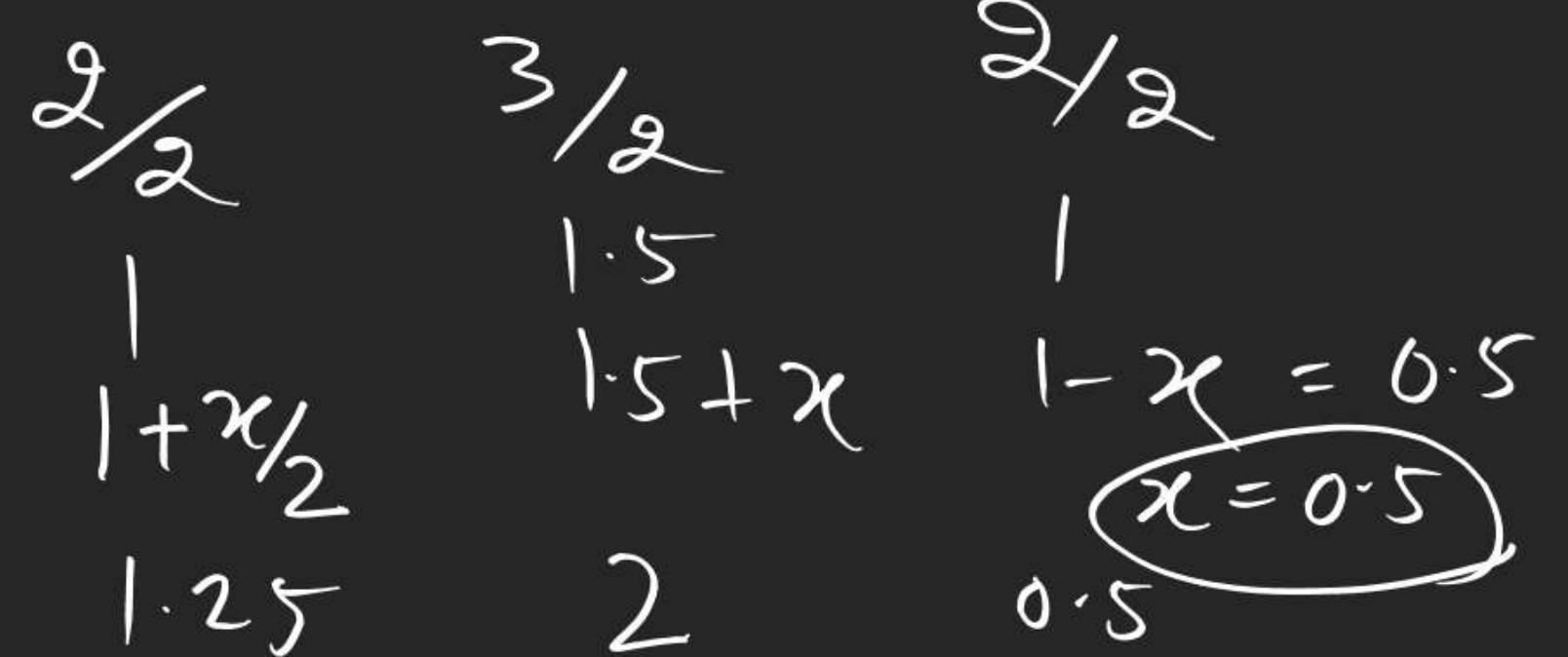
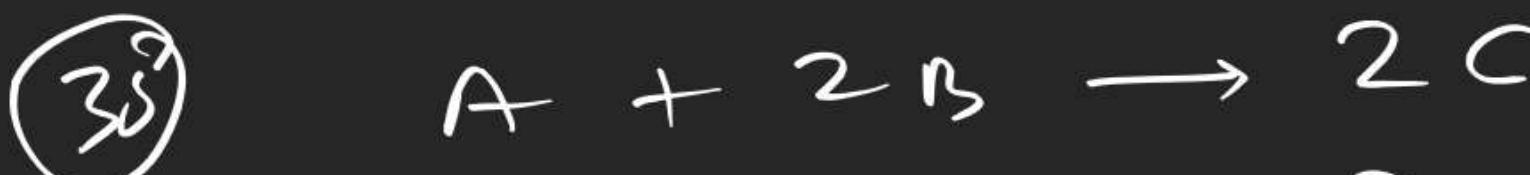
$$\textcircled{21} \quad K_p = k_c (RT)^{-1}$$





$$\begin{aligned} 2a-x &= x \\ a &= x \end{aligned}$$


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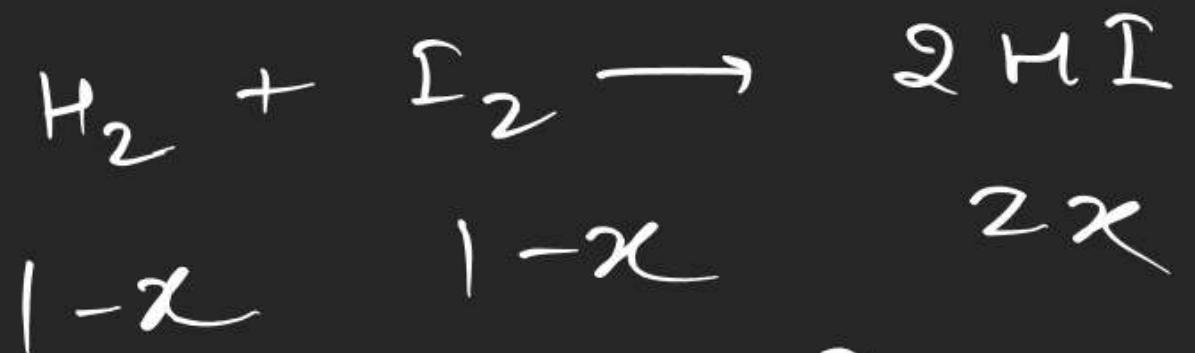


$$k_c = \frac{(0.5)^2}{1.25 \times 4} = \frac{1}{4}$$

$$= \frac{1}{20}$$

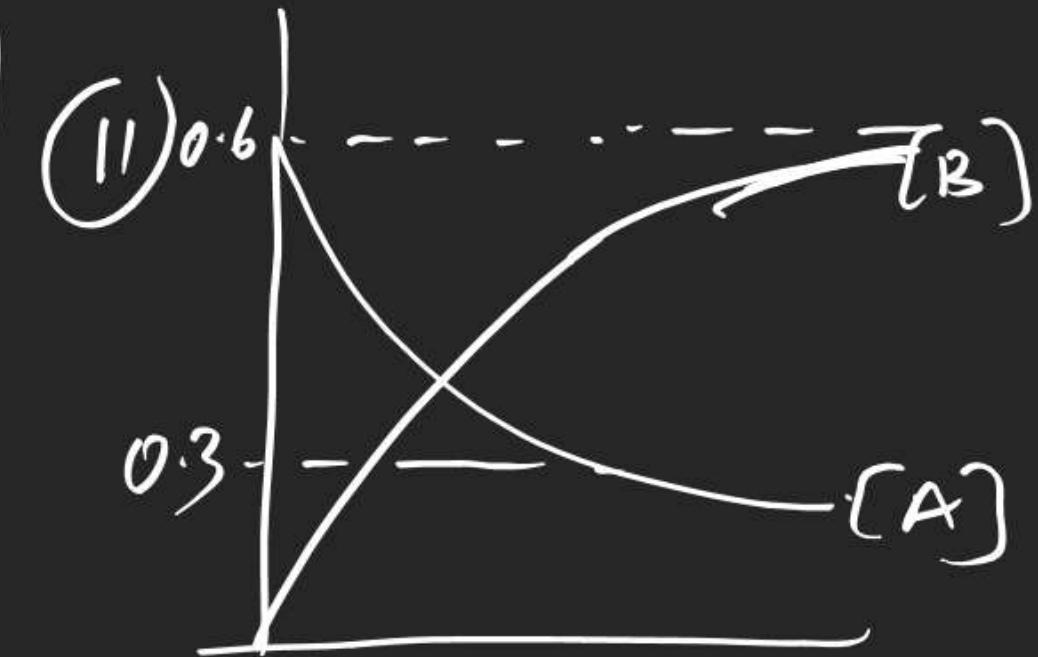
(37)

$$\frac{k_f}{k_{f0}} = 1$$



$$\frac{(2x)^2}{(1-x)^2} = 1$$

$$n = 1/3$$





$\begin{array}{ccc} 0.5 & 1 & 1 \\ \curvearrowleft & & \\ Q = \frac{1 \times 1}{0.5} = 2 & & Q < K_c \end{array}$

$0.5-x \quad 1+x \quad 1+x$

$$\frac{(1+x)(1+x)}{0.5-x} = 4$$

Q. find  $K_p$  if equilibrium mixture contains  
 5 mol  $NH_3$ , 3 mol  $N_2$  & 2 mol  $H_2$   
 at 20 atm.



$$P_{N_2} = \left(\frac{3}{10}\right) \times 20 \\ = 6 \text{ atm}$$

$$K_p = \frac{P_{NH_3}^2}{P_{N_2} \times P_{H_2}^3} =$$

$$= \frac{(5)^2}{3 \times (2)^3} \times \left(\frac{20}{10}\right)^{-2} \\ = \frac{25}{3 \times 8} \times \frac{1}{4} = \frac{25}{96}$$

$$K_p = \frac{(10)^2}{6 \times (4)^3}$$

$$\frac{5/6}{25/6} \cdot \frac{1}{1/20} = \frac{25/12}{1/20} = 25/96$$

Q. find  $K_p$  for the given rxn



if total pressure at  
1 atm obtained  
by the decomposition  
 $\text{NH}_2\text{COONH}_4(s)$   
in atm.

$$\begin{array}{lll} P & 2P & 3P = 9 \\ 3 \text{ atm} & 6 \text{ atm} & \underline{P = 3 \text{ atm}} \end{array}$$

$$\begin{aligned} K_p &= P_{\text{CO}_2} \times P_{\text{NH}_3}^2 \\ &= (3)(6)^2 \\ &= 108 \end{aligned}$$

$$\begin{array}{ccc} x & 2x & 9 \text{ atm} \end{array}$$

$$\begin{aligned} P_{\text{NH}_3} &= \frac{2x}{3x} \times 9 \\ &= \underline{6 \text{ atm}} \end{aligned}$$

Q.

P

$$K_p = 2.43 \times 10^{-13}$$

$$(\text{atm})^5$$

find Pressure at 2916m

$$K_p = (P_{\text{H}_2\text{O}})^5 = 243 \times 10^{-15}$$

$$P = \underline{P_{\text{H}_2\text{O}}} = 3 \times 10^{-3} \text{ atm}$$

Q. find vapour pressure  $\gamma$   $H_2O(l)$  at  $25^\circ C$

Given at  $25^\circ$   $\underline{H_2O(l) \rightarrow H_2O(s)}$   $K_p = 40$  torr.

$$40 = K_p = \underbrace{\rho_{H_2O}}$$

# Relationship bet" $K_p$ or $K_c$ and $\alpha$



$$\Rightarrow a(1-\alpha) \quad \alpha \quad \alpha$$

$$K_c = \frac{(a\alpha)(a\alpha)}{a(1-\alpha)} \left(\frac{1}{V}\right)^1 = \frac{a\alpha^2}{1-\alpha} \times \frac{1}{V}$$

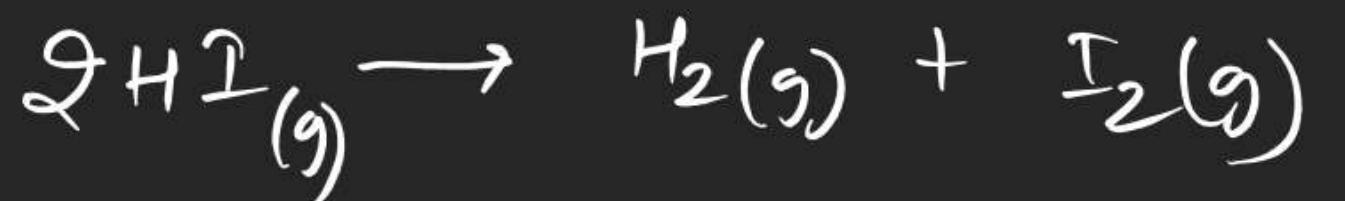
$$K_p = \frac{(a\alpha)(a\alpha)}{a(1-\alpha)} \times \left(\frac{P_T}{a(1+\alpha)}\right) = \frac{\alpha^2}{1-\alpha^2} P_T$$

$\alpha$  = degree of dissociation  
 = no of moles reacted  
 per mole of reactant

$$P_c = \frac{a\alpha}{a(1+\alpha)} \times P_T$$

If  $\Delta n_g = 0$

$$K_p = K_c$$



$$K_p = \frac{(\alpha/2)(\alpha/2)}{(1-\alpha)^2} \left( \frac{P_T}{T} \right)^0$$

$$K_c = K_p = \frac{(\alpha/2)^2}{(1-\alpha)^2}$$

$$\text{if } \Delta n_g = 0$$

$\gamma$  depends only on temperature.

find  $K_p$  for the Rxn if  $\text{PCl}_5$  decomposes 50% at 10 atm.

$$\alpha = 0.5$$



$$1 - \alpha \quad \alpha \quad \alpha$$

$$K_p = \frac{\alpha \cdot \alpha}{1 - \alpha} = \frac{\alpha^2}{1 - \alpha^2} \times 10 = \frac{0.25}{0.75} \times 10 = \frac{10}{3}$$

b) find the

pressure at  
which  $\alpha$  becomes

$$0.9.$$

$$K_p = \frac{10}{3} = \frac{\alpha^2}{1 - \alpha^2} P_T = \frac{0.04}{1 - 0.04} \times P_T$$

$$\frac{10}{3} = \frac{1}{24} P_T \quad P_T = 80$$