

YAKEEN NEET 2.0

2026

Chemical Equilibrium

Physical Chemistry


Lecture -03

By- Amit Mahajan Sir





Topics to be covered

- 1 Revision of Last Class
- 2 Equilibrium constant numericals
- 3 Homogeneous and heterogeneous equilibrium
- 4 Degree of dissociation  Trick
- 5 Magarmach Practice Questions (MPQ) & Home work from modules



Rules to Attend Class




- 1. Always sit in a peaceful environment with headphone and be ready with your copy and pen.**
- 2. Never ever attend a class from in between or don't join a live class in the middle of the chapter.**
- 3. Make sure to revise the last class before attending the next class & always complete your Magarmach Practice Questions.**
- 4. Never ever engage in chat whether live or recorded on the topic which is not being discussed in current class as by doing so u can be blocked by the admin team or your subscription can be cancelled.**



Rules to Attend Class



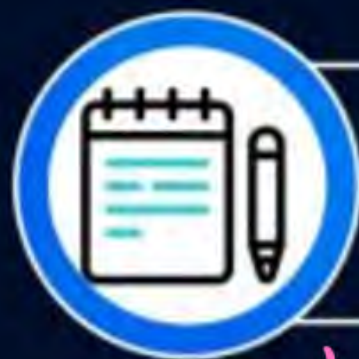
5. Try to make maximum notes during the class if something is left then u can use the notes pdf after the class to complete the remaining class.
6. Always ask your doubts in doubt section to get answer from faculty. Before asking any doubt please check whether same doubt has been asked by someone or not.



There is one big flaw in your Preparation that's name is Backlog ? What do we say to Backlog ?



NOT TODAY !!!



Revision of Last Class

LOMA

$$\underline{K_f [A]^a [B]^b} = \underline{K_b [C]^c [D]^d}$$

$$\frac{K_f}{K_b} = \underline{K_c} = \frac{[C]^c [D]^d}{[A]^a [B]^b}$$

$$K_p = \frac{(P_C)^c (P_D)^d}{(P_A)^a (P_B)^b}$$

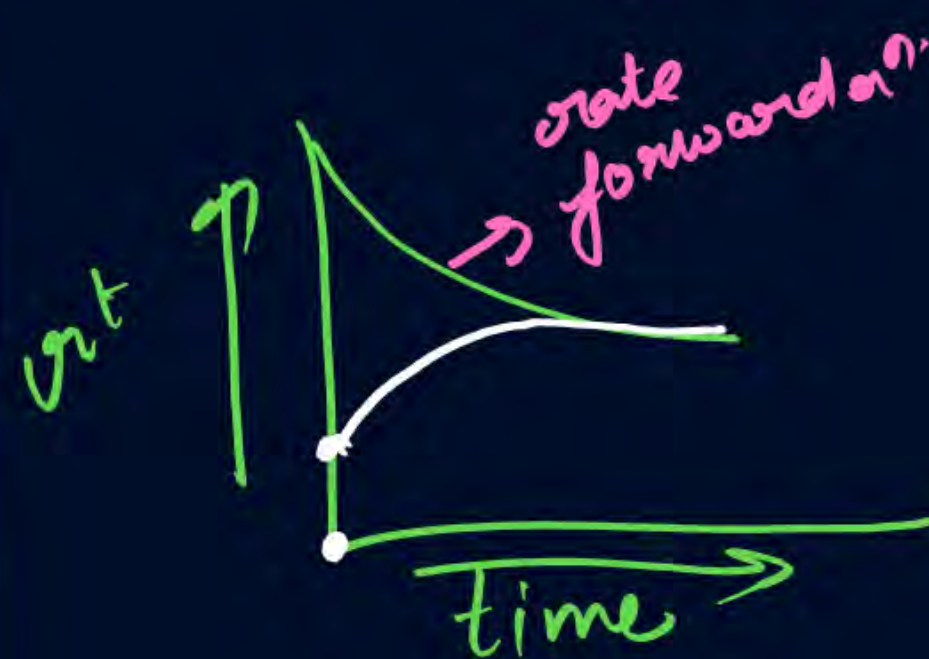


$$K_p = K_c (RT)^{\Delta n_g}$$

\downarrow
 $\Delta n_g = 0$
 $K_p = K_c$
 \downarrow
 $P = 1 \text{ atm}$
 $RT \approx 1$
 $K_p = K_c$

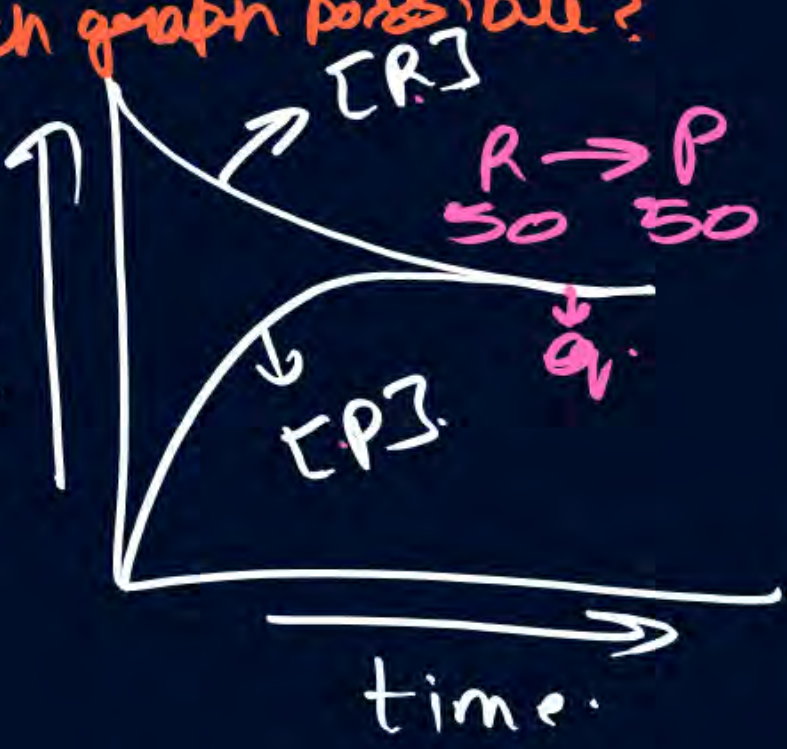
\downarrow
 $(+)\text{ve}$
 $K_p > K_c$

\downarrow
 $(-)\text{ve}$
 $K_p < K_c$

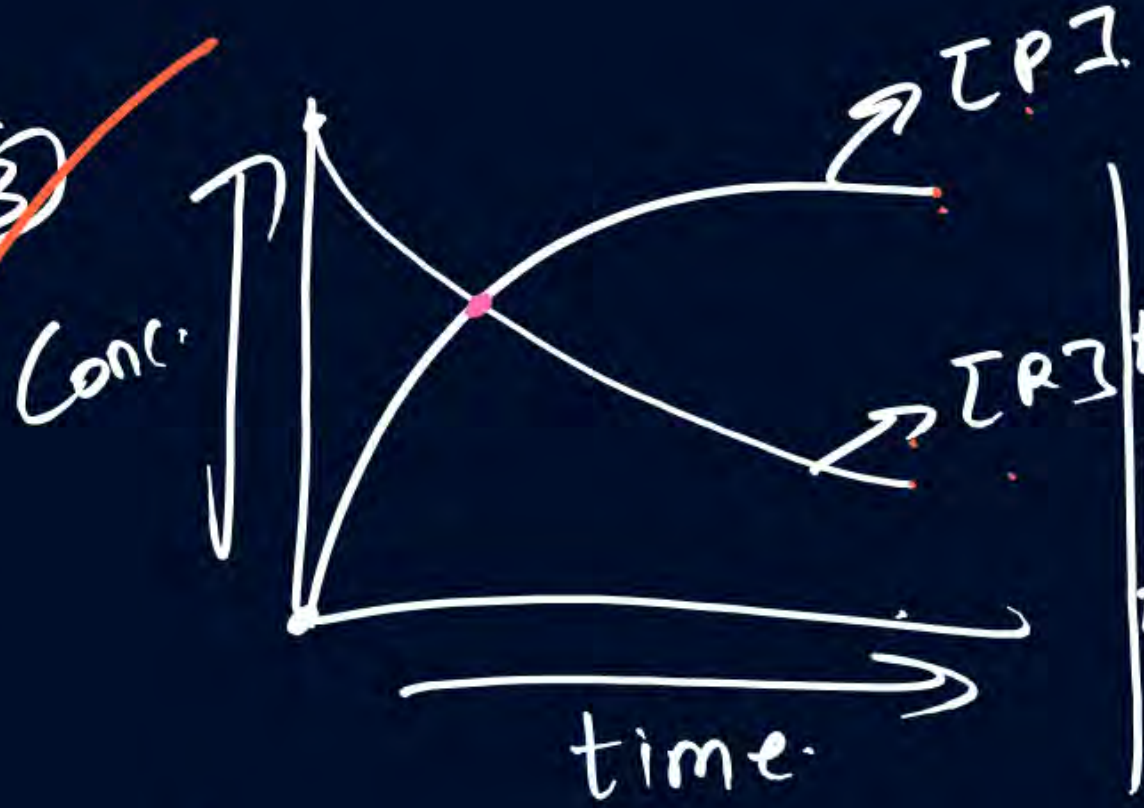


Q. 2 which graph possible?

Conc.
~~①~~



③

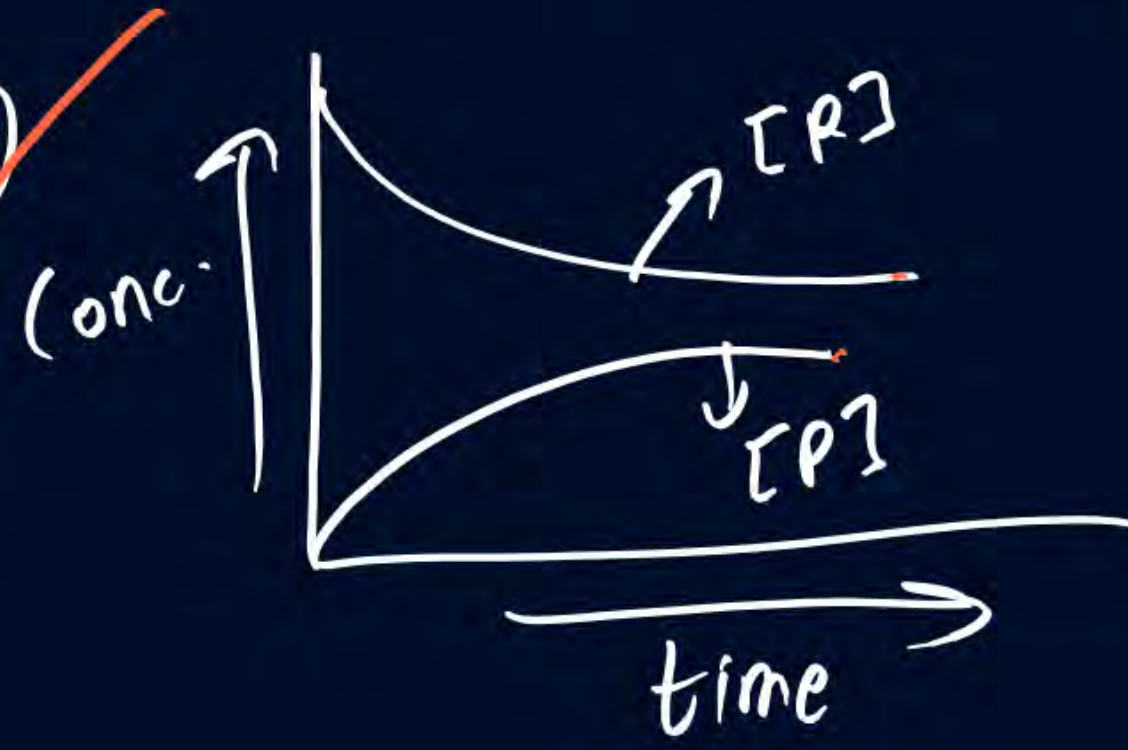


$t=0$	$R \rightarrow 100$	$P \rightarrow 0$
$t=t_{eq}$	$100 - 90 = 10$	90
	\downarrow	
	5 hr	

Conc.
~~②~~



④



Q2 which oxide more stable?



$K_C \uparrow$ Prod. stable



2.5 kcal/mole
↓
1 kcal

Q $A \rightleftharpoons B$ $\Delta H = 0$
if we inc. T from T_1 to T_2
 \downarrow \downarrow
 K_1 K_2

(a) $K_1 > K_2$

(b) $K_1 < K_2$

(c) $K_1 = K_2$

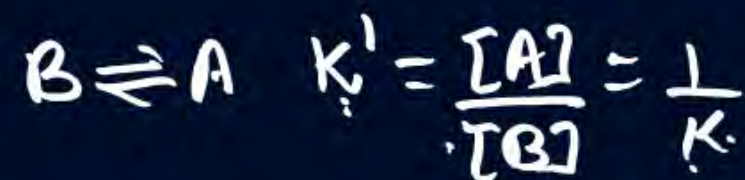
(d) Cannot be predicted

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

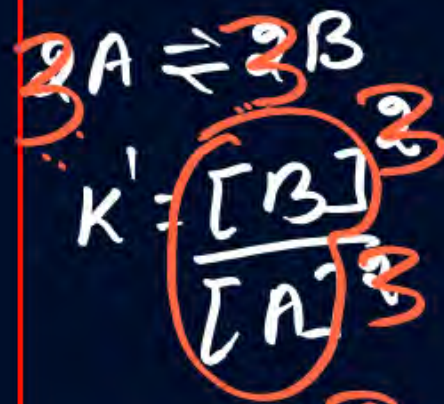
$$\cancel{\log K_2} = \cancel{\log K_1}$$



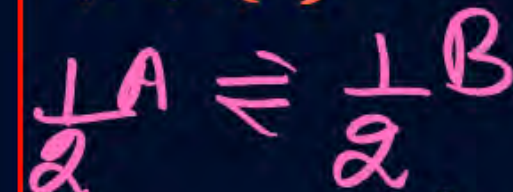
Characteristics of Equilibrium Constant



$$K' = \frac{[A]}{[B]}$$



$$K' = (K)^2$$



$$K' = \frac{[B]^{\frac{1}{2}}}{[A]^{\frac{1}{2}}} = (K)^{\frac{1}{2}}$$

#MIT

①

a^n is

effect on eq. Constt.

reverse.

$$K' = \frac{1}{K}$$

S.C. $\times x$

$$K' = (K)^x$$

$\frac{1}{y}$ \times S.C.

$$K' = (K)^{\frac{1}{y}}$$

a^n add.

$$K' = K_1 \times K_2 \times K_3$$

$$A \rightleftharpoons B \quad K = \frac{[B]}{[A]}$$

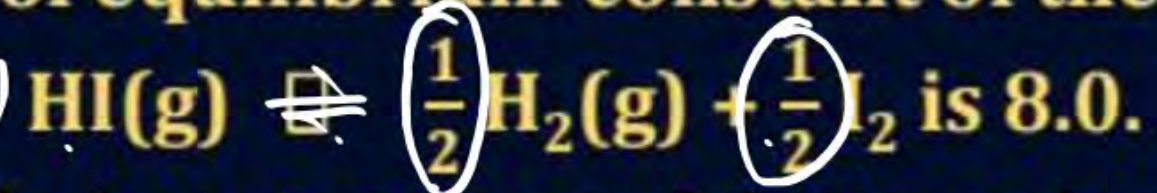


$$K = K_1 \times K_2 \times K_3 = \frac{\cancel{[C]}}{[A]} \times \frac{\cancel{[D]}}{\cancel{[C]}} \times \frac{[B]}{\cancel{[D]}} = \frac{[B]}{[A]}$$

QUESTION



The value of equilibrium constant of the reaction:



The equilibrium constant of the reaction
 $\text{H}_2\text{(g)} + \text{I}_2\text{(g)} \rightleftharpoons 2\text{HI(g)}$ will be:

A 16

B 1/8

C 1/16

D 1/64

or reverse \times S.C. = 2

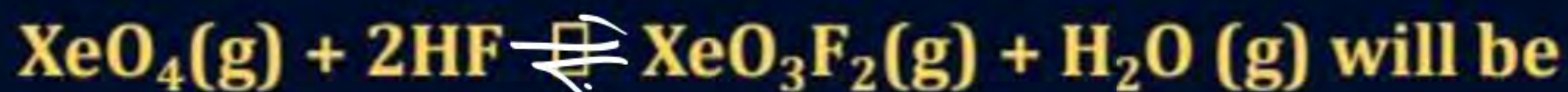
$$\left(\frac{1}{8}\right)^2 = \frac{1}{64}$$

QUESTION – (AIIMS 2013)

If K_1 and K_2 are respective equilibrium constants for the two reactions



The equilibrium constant for the reaction



A $\frac{K_1}{K_2^2}$

B $K_1 \cdot K_2$

C $\frac{K_1}{K_2}$

D $\frac{K_2}{K_1}$

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

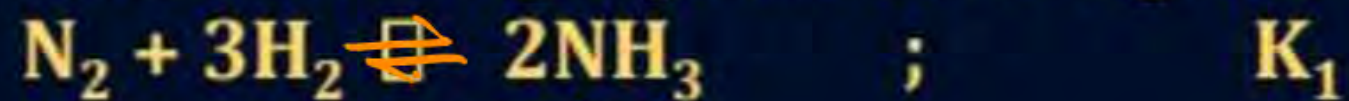
eq. (2) + (-eq. (1))

= eq. (2) + reverse eq. (1)

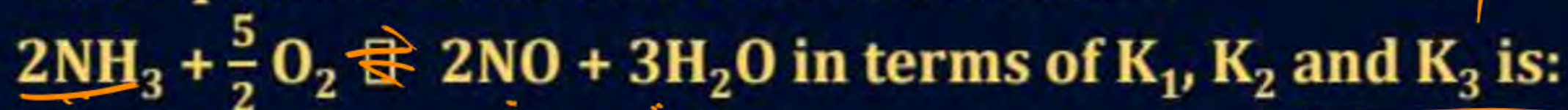
$$K = K_2 \times \frac{1}{K_1}$$

QUESTION – (AIIMS 2012)

The following equilibria are given:



The equilibrium constant of the reaction



A $\frac{K_1 K_2}{K_3}$

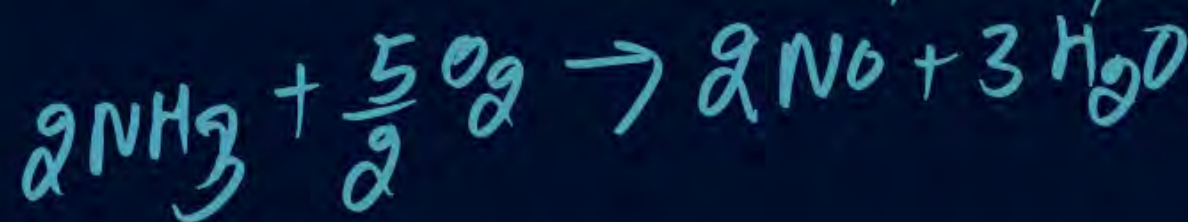
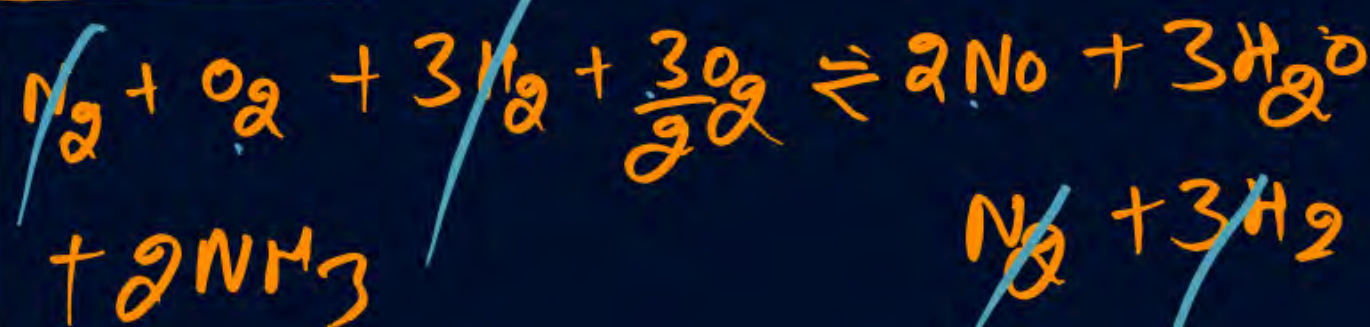
B $\frac{K_1 K_3^2}{K_2}$

C $\frac{K_2 K_3^3}{K_1}$

D $K_1 K_2 K_3$

reverse eq. ① + eq. ② + 3x eq. ③

$$K = \frac{1}{K_1} \times K_2 \times (K_3)^3$$



QUESTION



If the value of equilibrium constant for a particular reaction is 1.6×10^{12} , then at the equilibrium system will contain:

- ☒ **A** Mostly products
- ☐ **B** Similar amounts of reactants and products
- ☐ **C** All reactants
- ☐ **D** Mostly reactants

QUESTION – (AIIMS 2008)

In which of the following reactions, the concentration of the product is higher than the concentration of ~~the~~ ^{the} reactant at equilibrium? (K = equilibrium constant)

- ☐ A $A \rightleftharpoons B$; $K = 0.001$
- ☒ B $M \rightleftharpoons N$; $K = 10$
- ☐ C $X \rightleftharpoons Y$; $K = 0.005$
- ☐ D $R \rightleftharpoons P$; $K = 0.01$

QUESTION



On doubling P and V with constant temperature, the equilibrium constant will:

- ☒ **A** Remain constant
- ☐ **B** Become double
- ☐ **C** Become one-fourth
- ☐ **D** None of these

QUESTION



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: $\text{CN}^- + \text{CH}_3\text{COOH} \rightleftharpoons \text{HCN} + \text{CH}_3\text{COO}^-$ will be:

A 3.0×10^5

B 3.0×10^{-5}

C 3.0×10^{-4}

D 3.33×10^4



$\frac{1}{3} \times 10^5 = 0.33 \times 10^5$
 $= 3.3 \times 10^4$

eq (1) + reverse eq (2)

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

$$= \frac{1.5 \times 10^{-5}}{4.5 \times 10^{-10}} = 3.33 \times 10^4$$

QUESTION



In the figure shown below, reactant A (represented by square) is in equilibrium with product B (represented by circle). The equilibrium constant is:

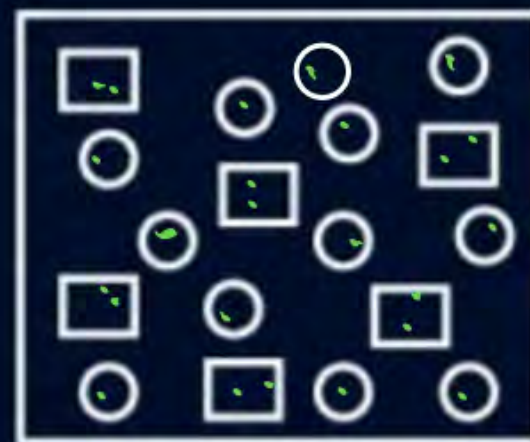
A 4

☒ **B** 2

C 8

D 1

$$K = \frac{12}{6} = 2.$$



QUESTION – (AIIMS 2013)

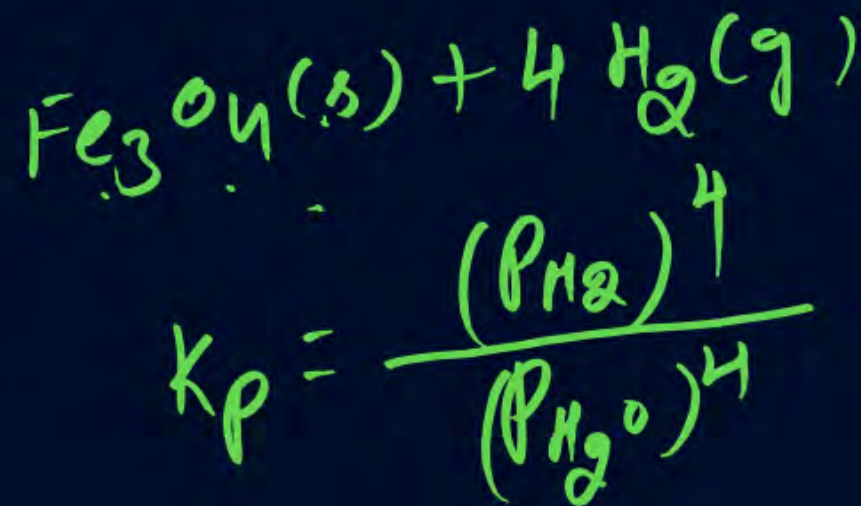
Steam reacts with iron at high temperature to give hydrogen gas and $\text{Fe}_3\text{O}_4(\text{s})$. The correct expression for the equilibrium constant is:

A $\frac{P_{\text{H}_2}^2}{P_{\text{H}_2\text{O}}^2}$

C $\frac{(P_{\text{H}_2})^4 [\text{Fe}_3\text{O}_4]}{(P_{\text{H}_2\text{O}})^4 [\text{Fe}]}$

B $\frac{(P_{\text{H}_2})^4}{(P_{\text{H}_2\text{O}})^4}$ $3\text{Fe}(\text{s}) + 4\text{H}_2\text{O}(\text{g})$

D $\frac{[\text{Fe}_3\text{O}_4]}{[\text{Fe}]}$



QUESTION



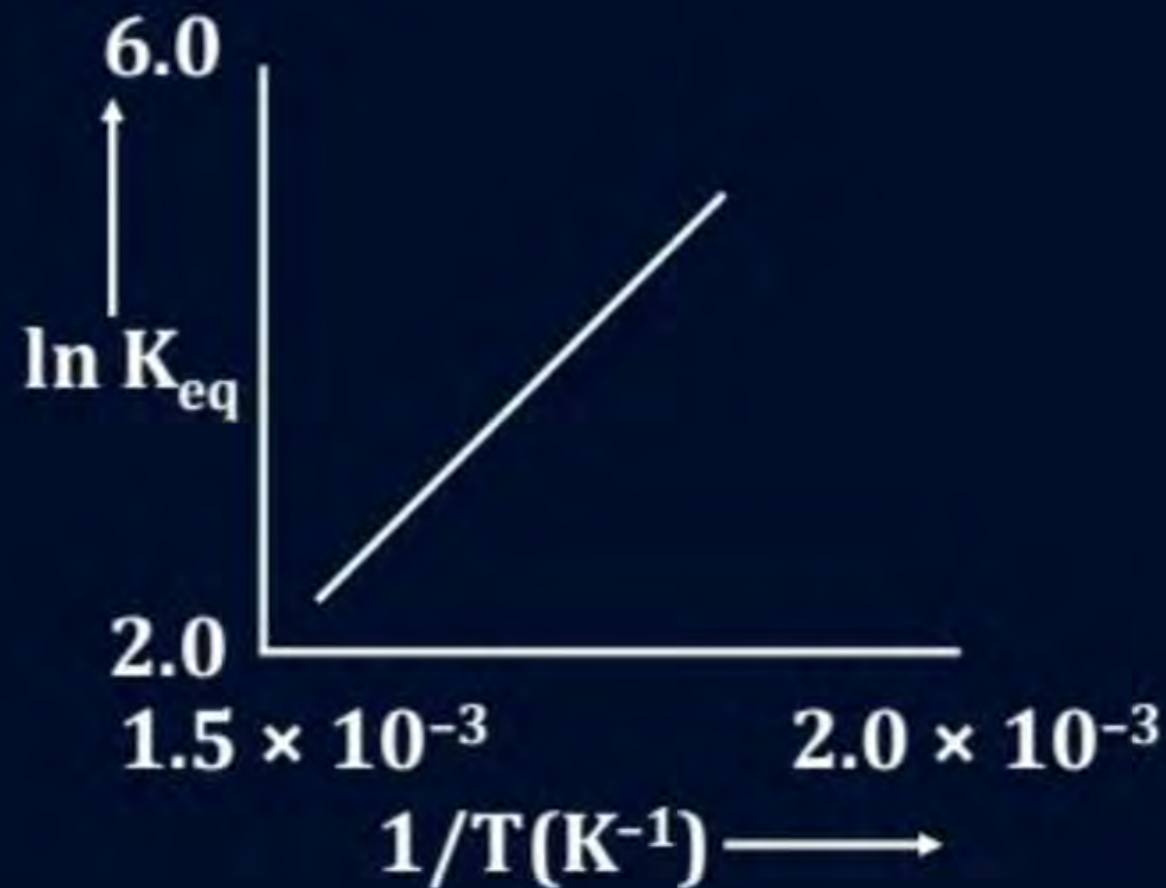
A schematic plot of $\ln K_{eq}$ versus inverse of temperature for a reaction is shown in the figure. The reaction must be:

- ☒ **A** Exothermic
- ☐ **B** Endothermic
- ☐ **C** One with negligible enthalpy change
- ☐ **D** Highly spontaneous at ordinary temperature

$$\ln K = \ln A - \frac{\Delta H}{2.303RT}$$

Handwritten note: ΔH is negative = exo.

$$y = c + m x$$





Reaction Quotient (Q_c)



Stoichiometric coefficients

$$a \cdot 0 + b \cdot 0 = c \cdot 0 + d \cdot 0$$

$$K_c = \frac{[C]^c [D]^d}{[A]^a [B]^b}$$

at eq

#

$$Q_c = \frac{[C]^c [D]^d}{[A]^a [B]^b}$$

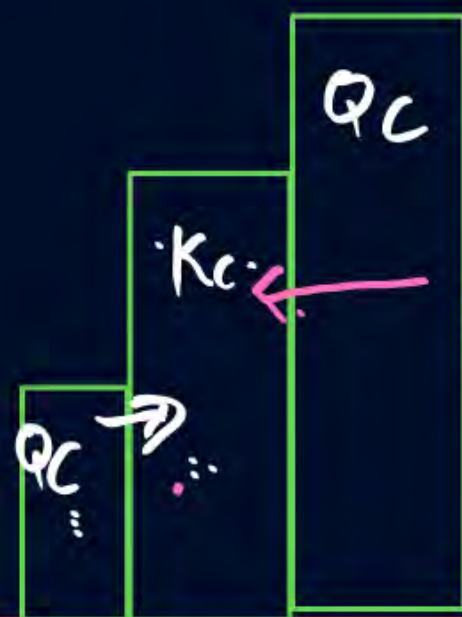
at any instant

$$② \quad t < t_{eq} \quad Q_c < K_c$$

$$t = t_{eq} \quad \text{or} \quad t > t_{eq}$$

$$Q_c = K_c$$

③



$Q_c < K_c$ eq. \rightarrow shift forward
 $Q_c > K_c$ eq. \rightarrow shift backward

QUESTION – (NEET 2024)

For the reaction $2A \rightleftharpoons B + C$, $K_c = 4 \times 10^{-3}$. At a given time, the composition of reaction mixture is:

$$[A] = [B] = [C] = 2 \times 10^{-3} \text{ M}$$

Then, which of the following is correct?

$$Q_c = \frac{2 \times 10^{-3} \times 2 \times 10^{-3}}{(2 \times 10^{-3})^2} = 1$$

$$Q_c > K_c$$

- ☐ A Reaction is at equilibrium.
- ☐ B Reaction has a tendency to go in forward direction.
- ☒ C Reaction has a tendency to go in backward direction.
- ☐ D Reaction has gone to completion in forward direction.

The equilibrium constant for a reaction is K and the reaction quotient is Q . For a particular reaction mixture, the ratio K/Q is 0.3. This means that

- ☒ **A** The reaction mixture will equilibrate to form more reactant species
- ☐ **B** The reaction mixture will equilibrate to form more product species
- ☐ **C** The equilibrium ratio of reactant to product concentrations will be 3
- ☐ **D** The equilibrium ratio of reactant to product concentrations will be 0.33

$$\frac{K}{Q} = 0.3$$

$$K = 0.3Q$$

$$K < Q$$

$$\frac{Q_c}{K_c} > 1$$

↓

QUESTION – (AIIMS 2011)

✓ **Assertion:** Reaction quotient is defined in the same way as equilibrium constant at any state of the reaction.

✗ **Reason:** If Q_c (reaction quotient) $<$ K_c (equilibrium constant) reaction moves in direction of reactants.

- ☐ **A** If both Assertion and Reason are correct and the Reason is the correct explanation of Assertion.
- ☐ **B** If both Assertion and Reason are correct but Reason is not the correct explanation of Assertion.
- ✓ ☒ **C** If Assertion is correct but Reason is incorrect.
- ☐ **D** If both the Assertion and Reason are incorrect.



Homogeneous Equilibrium

↓
reactants & products are in
same phase.

↓
both
gaseous.

↓
miscible
liquids.



an equilibrium condition
for a reaction in which the
reactants and products are
in the same state



homogeneous equilibrium



Heterogeneous Equilibrium

reactants & product are in diff. phase.

immiscible liquids.

different solid.

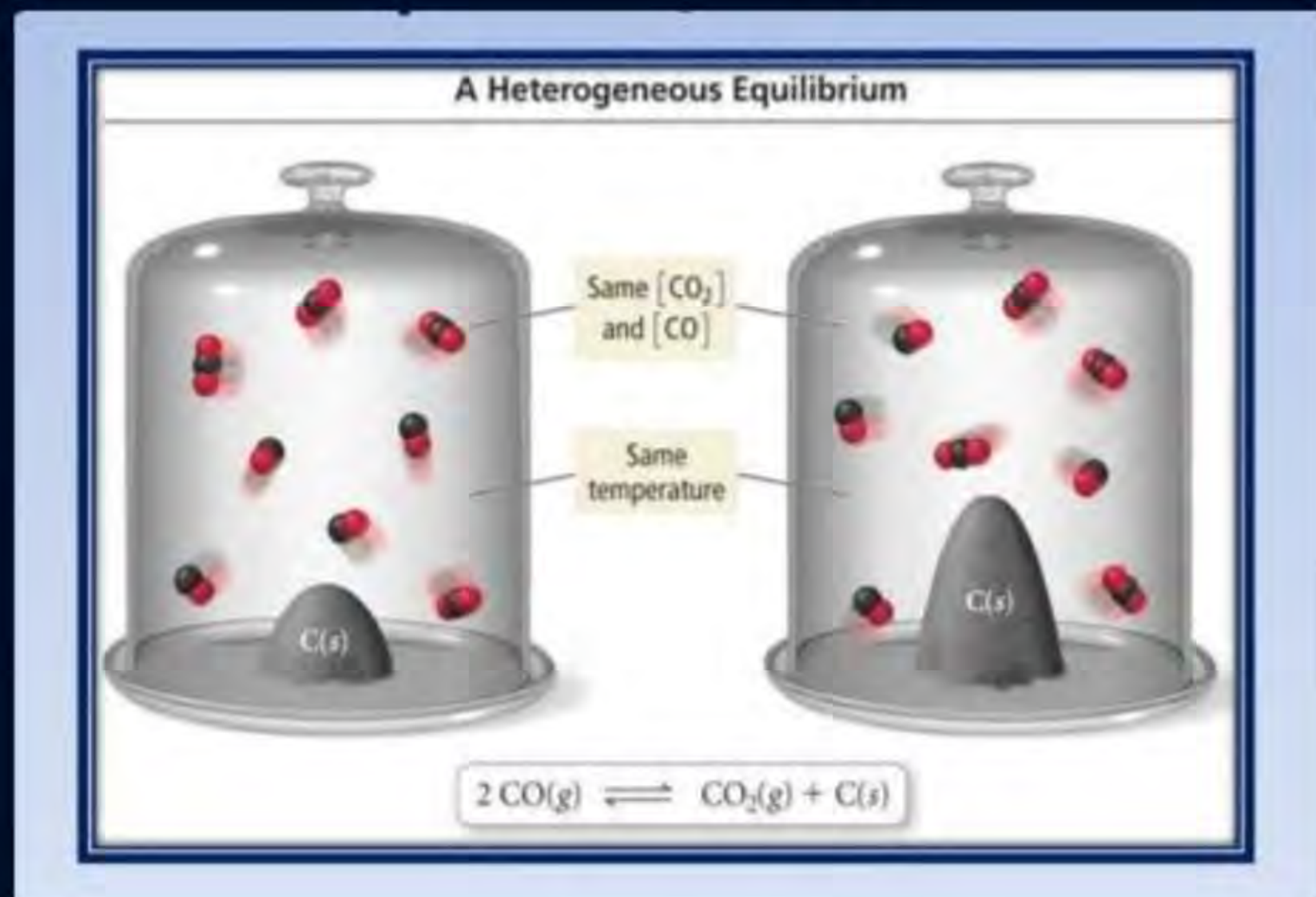
different physical state.



Phases = 2



Phases = 2 | Heterogeneous eq.





Degree of Dissociation

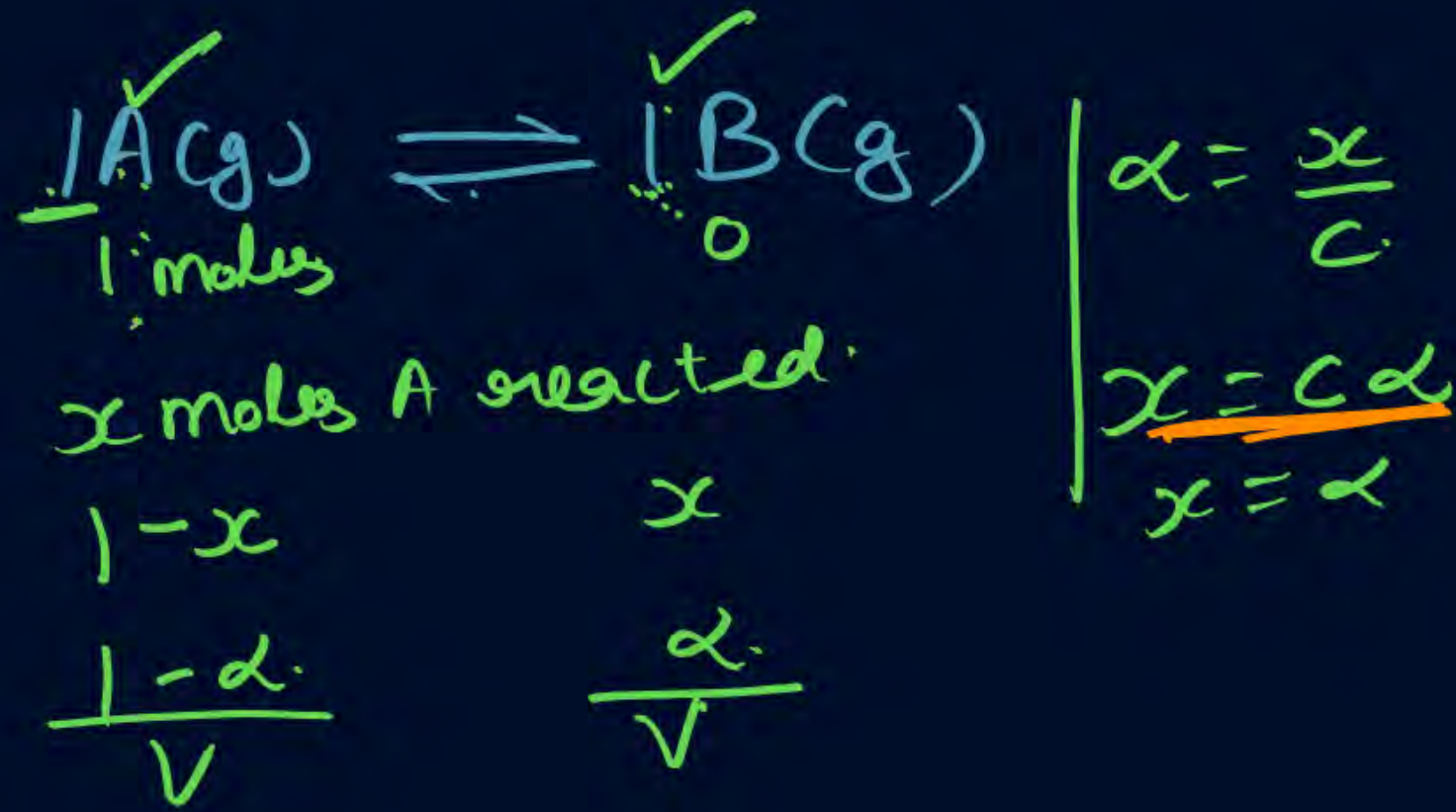
(α)

MIT

①
$$\alpha = \frac{\text{moles dissociated (x)}}{\text{moles taken (C)}}$$

②
$$\% \text{ age dissociation} = \alpha \times 100$$

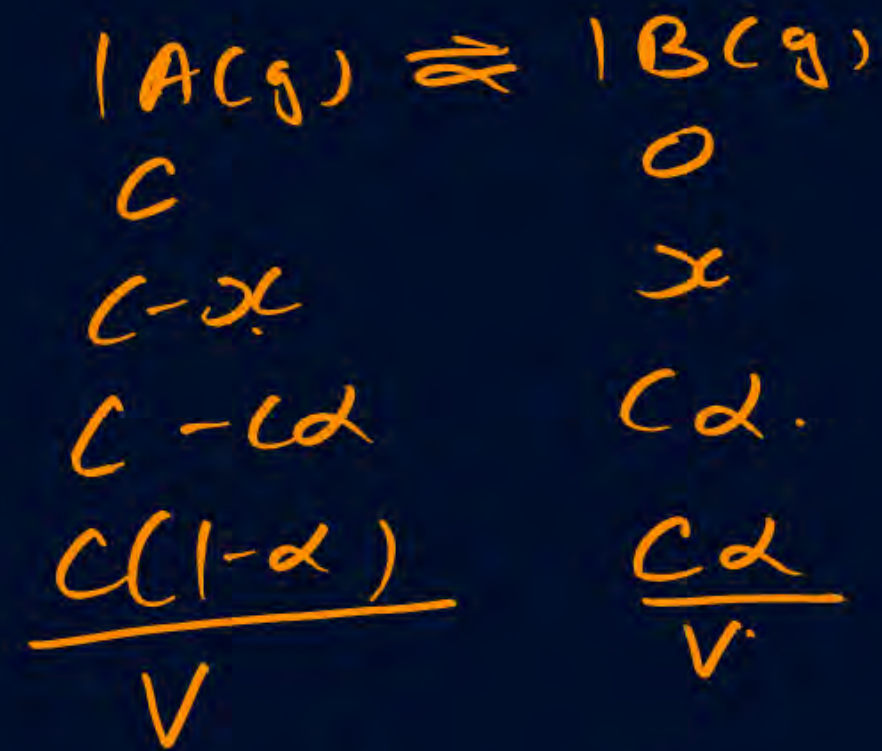
$t=0$
 $t=t$



$$K_c = \frac{[B]}{[A]} = \frac{\alpha \times V}{V(1-\alpha)}$$

$$K_c = \frac{\alpha}{1-\alpha}$$

$t=0$
 $t=t$



$$K_c = \frac{\cancel{C} \alpha \times \cancel{V}}{\cancel{C} (1-\alpha) 1-\alpha} = \frac{\alpha}{1-\alpha}$$



$t=0$

C

0

$t=t$

$C-x$

$2x$

$$\frac{C-x}{V}$$

$$\frac{2x}{V}$$

$$\frac{C(1-x)}{V}$$

$$\frac{2x}{V}$$

$$K_c = \frac{(2x)^2}{V} \times V$$

$$= \frac{4C^2 x^2}{V(1-x)}$$



C

0

$C-x$

$4x$

$C-x$

$4x$

$$\frac{C-x}{V}$$

$$\frac{4x}{V}$$

$$K_c = \frac{(4x)^4}{V^3} \times V = \frac{256C^3 x^4}{V^3(1-x)}$$



$$K_c = \frac{(2C\alpha)^2 (3C\alpha)^3 \times V}{V^4 C(1-\alpha)} = \frac{108 C^4 \alpha^5}{V^4 (1-\alpha)}$$

3 Page blank

QUESTION



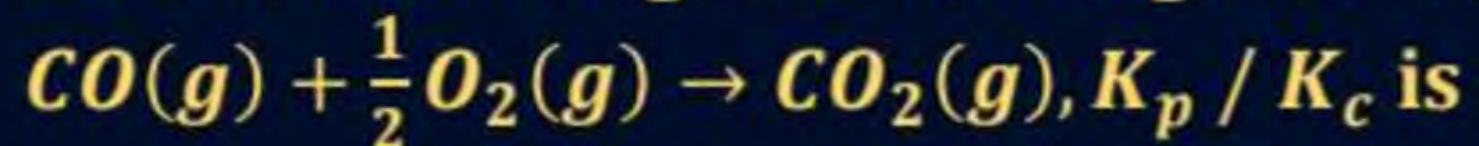
For PCl_5 , dissociation, molar mass is 120 g. Find α and %age of dissociation.

At a fixed temperature, the volume of the reaction container is halved. For this change, which of the following statements hold true regarding the equilibrium constant (K_p) and degree of dissociation (α)?

- A** Neither K_p nor α changes
- B** Both K_p and α change
- C** K_p changes but α does not change
- D** K_p does not change but α changes

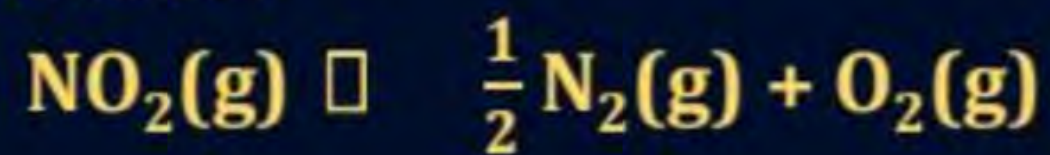
QUESTION – (AIIMS 2017)

For the following reaction in gaseous phase



- A** $(RT)^{1/2}$
- B** $(RT)^{-1/2}$
- C** (RT)
- D** $(RT)^{-1}$

For the reaction, $\text{N}_2(\text{g}) + \text{O}_2(\text{g}) \rightleftharpoons 2\text{NO}(\text{g})$, the equilibrium constant is K_1 . The equilibrium constant is K_2 for the reaction, $2\text{NO}(\text{g}) + \text{O}_2(\text{g}) \rightleftharpoons 2\text{NO}_2(\text{g})$. What is K for the reaction:



- A** $\frac{1}{2K_1K_2}$
- B** $\frac{1}{4K_1K_2}$
- C** $\left[\frac{1}{K_1K_2}\right]^{1/2}$
- D** $\frac{1}{K_1K_2}$

QUESTION – (AIIMS 2016)

Two equilibria, $AB \rightleftharpoons A^+ + B^-$ and $AB + B^- \rightleftharpoons AB_2^-$ are simultaneously maintained in a solution with equilibrium constants, K_1 and K_2 respectively. The ratio of $[A^+]$ to $[AB_2^-]$ in the solution is:

- A** directly proportional to $[B^-]$
- B** inversely proportional to $[B^-]$
- C** directly proportional to the square of $[B^-]$
- D** inversely proportional to the square of $[B^-]$

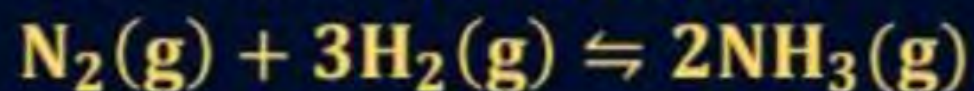
QUESTION – (AIIMS 2015)

The reaction $2A(g) + B(g) \rightleftharpoons 3C(g) + D(g)$ is begun with the concentrations of A and B both at an initial value of 1.00 M. When equilibrium is reached, the concentration of D is measured and found to be 0.25 M. The value for the equilibrium constant for this reaction is given by the expression

- A** $[(0.75)^3(0.25)] \div [(0.75)^2(0.25)]$
- B** $[(0.75)^3(0.25)] \div [(1.00)^2(1.00)]$
- C** $[(0.75)^3(0.25)] \div [(0.50)^2(0.75)]$
- D** $[(0.75)^3(0.25)] \div [(0.50)^2(0.25)]$

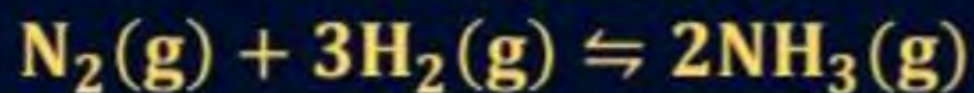
QUESTION – (AIIMS 2008)

Assertion: For reaction



Unit of $K_c = \text{L}^2 \text{mol}^{-2}$

Reason: For the reaction



Equilibrium constant, $K_c = \frac{[\text{NH}_3]^2}{[\text{N}_2][\text{H}_2]^3}$

- A** If both Assertion and Reason are correct and the Reason is a correct explanation of the Assertion.
- B** If both Assertion and Reason are correct but Reason is not a correct explanation of the Assertion.
- C** If the Assertion is correct but Reason is incorrect.
- D** If both the Assertion and Reason are incorrect.
- E** If the Assertion is incorrect but the Reason is correct.

QUESTION



For the reaction



- A** $K_c = K_p(RT)$
- B** $K_c = K_p(RT)^{-1/2}$
- C** $K_c = K_p(RT)^{1/2}$
- D** $K_c = K_p(RT)^{3/2}$

QUESTION – (NCERT Exemplar)

We know that the relationship between K_c and K_p is

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

What would be the value of Δn for the reaction



- A** 1
- B** 0.5
- C** 1.5
- D** 2

QUESTION – (NCERT Exemplar)

PCl_5 , PCl_3 and Cl_2 are at equilibrium at 500K in a closed container and their concentrations are $0.8 \times 10^{-3} \text{ mol L}^{-1}$, $1.2 \times 10^{-3} \text{ mol L}^{-1}$ and $1.2 \times 10^{-3} \text{ mol L}^{-1}$ respectively. The value of K_c for the reaction $\text{PCl}_5 (\text{g}) \rightleftharpoons \text{PCl}_3 (\text{g}) + \text{Cl}_2 (\text{g})$ will be

- A** $1.8 \times 10^3 \text{ mol L}^{-1}$
- B** 1.8×10^{-3}
- C** $1.8 \times 10^{-3} \text{ L mol}^{-1}$
- D** 0.55×10^4

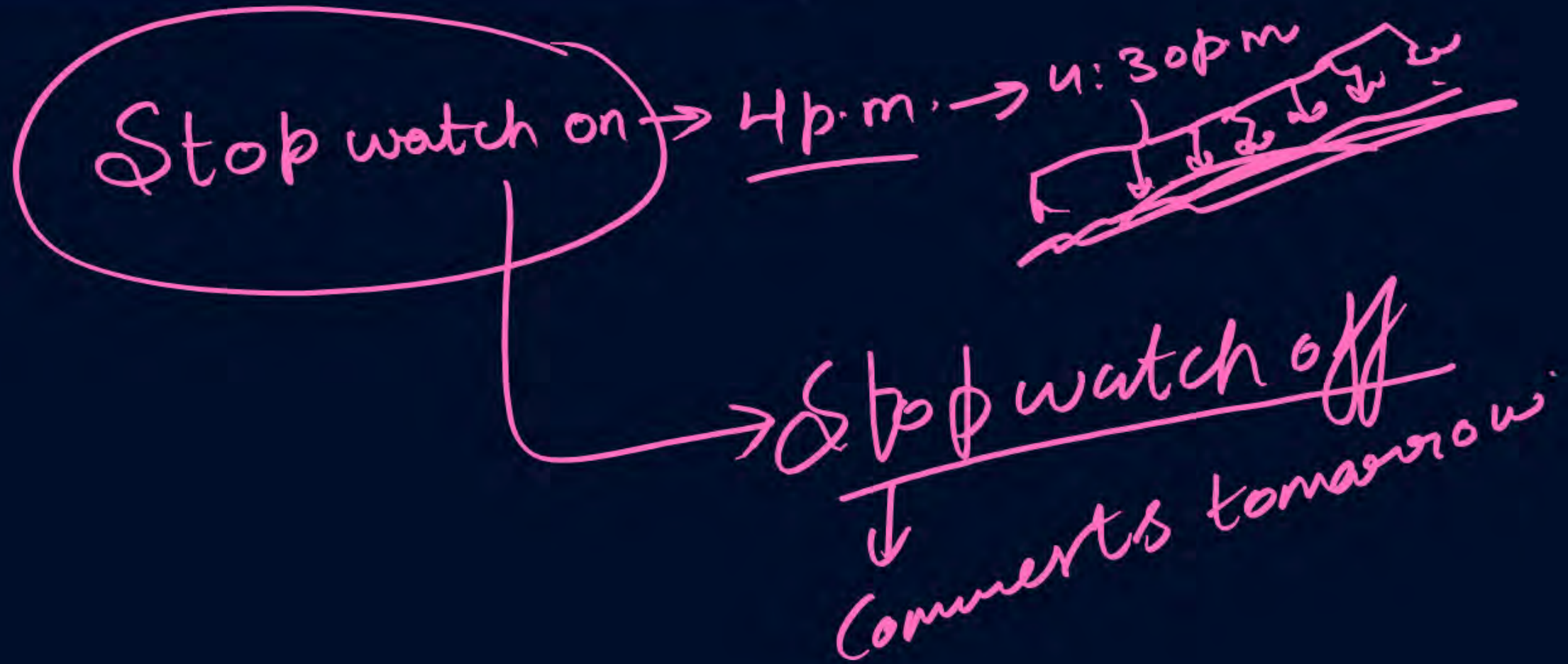
QUESTION – (NEET 2022)

$3\text{O}_2(\text{g}) \rightleftharpoons 2\text{O}_3(\text{g})$ for the given reaction at 298 K, K_c is found to be 3.0×10^{-59} . If the concentration of O_2 at equilibrium is 0.040 M, then concentration of O_3 in M is:

- A** 1.20×10^{21}
- B** 4.38×10^{-32}
- C** 1.90×10^{-63}
- D** 2.40×10^{31}



Trick

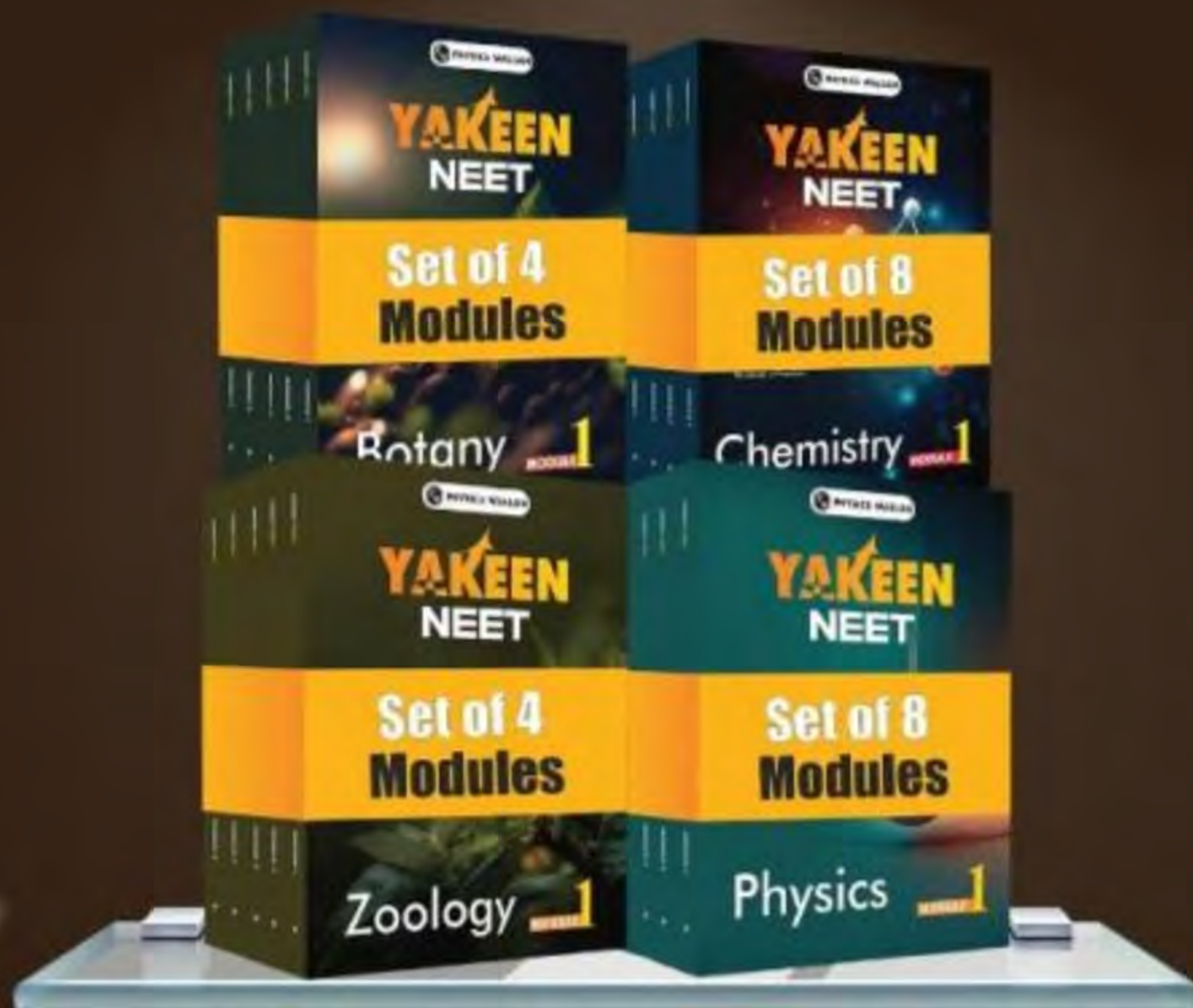


MED ICS Test → Thermodynamics
↳ Reg ←
↳ Lec-1 to Lec-3
↳ Moderate level.



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