

YAKEEN NEET 2.0

2026

Chemical Equilibrium

Physical Chemistry

Lecture -02

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Topics to be covered

- ✓ 1 Revision of Last Class
- ✓ 2 Law of Mass Action
- ✓ 3 Equilibrium constants and their relation
- ✓ 4 Reaction Quotient
- ✓ 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





Physical Equilibrium

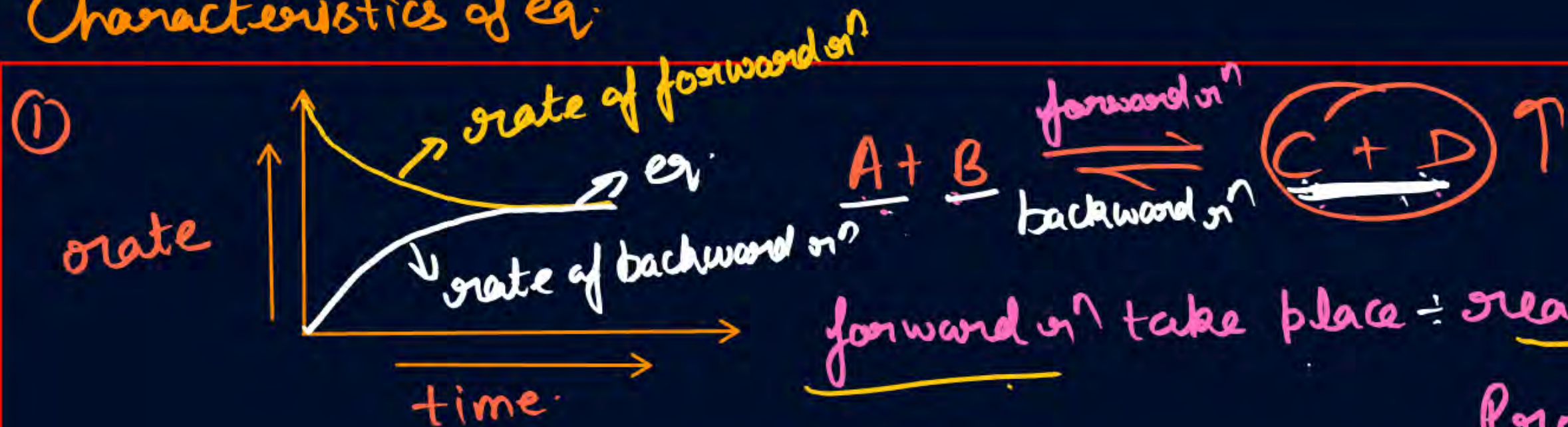




Solid-Gas Equilibrium

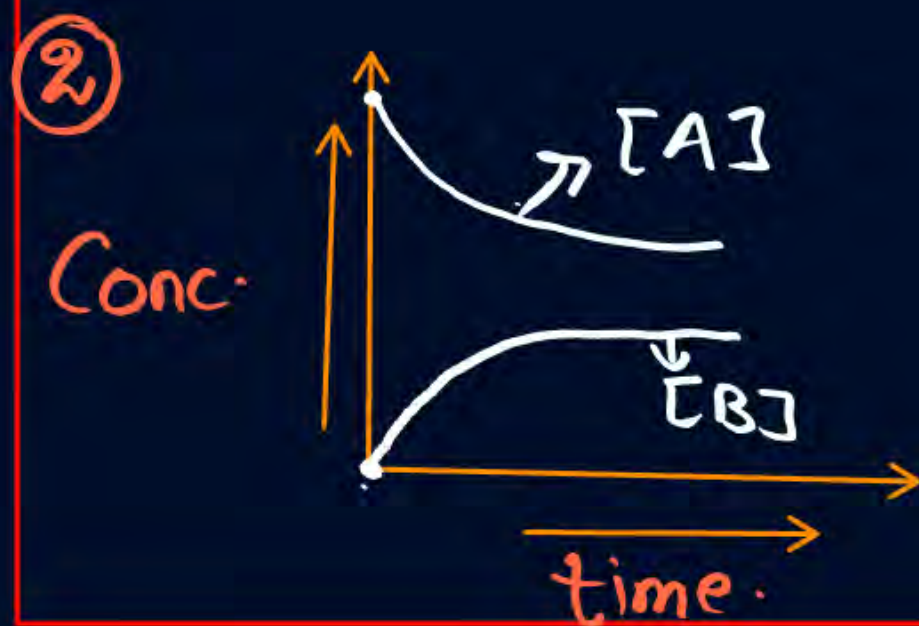


Characteristics of eq.



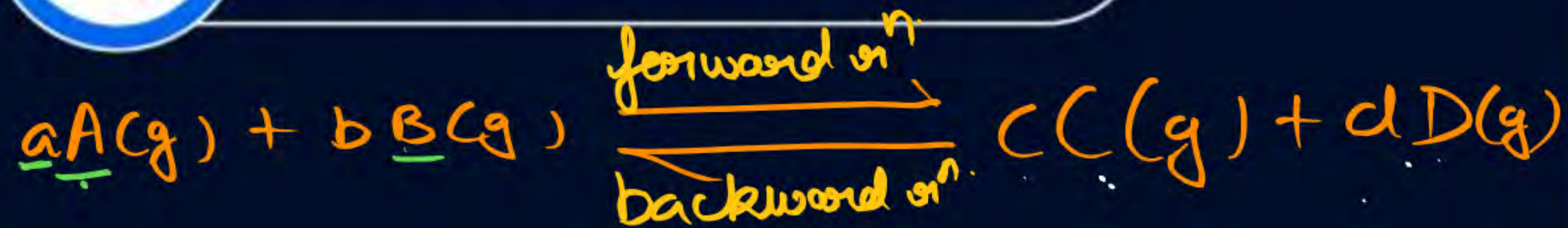
forward rxn take place \div reactant Conc. \downarrow
 product Conc. \uparrow

backward rxn take place \div product Conc. \downarrow
 reactant Conc. \uparrow





Law of Mass Action (LOMA)



Rate of r^n \propto Product of active mass of reacting species raise to

power S.C.
active mass = Conc.

$$\text{rate backward } r^n \propto [C]^c [D]^d$$

$$= K_b [C]^c [D]^d$$

K_b = rate constt backward r^n .

$$\text{Rate of forward } r^n \propto [A]^a [B]^b$$

$$\text{rate forward } r^n = K_f [A]^a [B]^b$$

K_f = rate constt forward r^n .



at eq.

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

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

Eq. Constt.
in Conc.

ratio of K_f & K_b \leftarrow ratio of $\frac{(\text{Conc. of Prod.})^x}{(\text{Conc. of react})^y}$

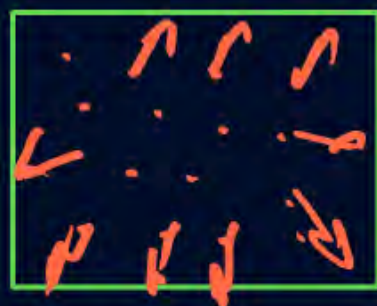


Relation Between K_p & K_c



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

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



$$P = \frac{nRT}{V}$$

$$P_A = \left(\frac{n_A}{V} \right) RT$$

$$P_A = [A] RT$$

$$P_B = [B] RT$$

$$P_C = [C] RT$$

$$P_D = [D] RT$$

$$K_p = \frac{([C]RT)^c ([D]RT)^d}{([A]RT)^a ([B]RT)^b}$$

$$K_p = \frac{[C]^c [D]^d}{[A]^a [B]^b} \frac{(RT)^{c+d}}{(RT)^{a+b}}$$

$$K_p = K_c (RT)^{\frac{(c+d) - (a+b)}{n_p(g) \quad n_r(g)}}$$

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

$$\textcircled{1} \quad K_p = \underline{K_c} \underline{(RT)}^{\Delta n_g}$$

$$\Delta n_g = \underline{n_p(g)} - \underline{n_r(g)}$$

$$\text{if } n_p(g) > n_r(g) \Rightarrow K_p > K_c$$

$$n_p(g) < n_r(g) \Rightarrow K_p < K_c$$

$$n_p(g) = n_r(g) \Rightarrow K_p = K_c$$

$$\text{if } T \approx 12 K \Rightarrow RT \approx 1 \Rightarrow K_p = K_c$$

In which one of the following equilibria, $K_p \neq K_c$?

- ☒ **A** $2\text{C(s)} + \text{O}_2\text{(g)} \rightleftharpoons 2\text{CO(g)}$ $\Delta n_g = 1$
- ☐ **B** $2\text{HI(g)} \rightleftharpoons \text{H}_2\text{(g)} + \text{I}_2\text{(g)}$ $\Delta n_g = 0$
- ☐ **C** $\text{NO}_2\text{(g)} + \text{SO}_2\text{(g)} \rightleftharpoons \text{NO(g)} + \text{SO}_3\text{(g)}$ $\Delta n_g = 0$
- ☐ **D** $2\text{NO(g)} \rightleftharpoons \text{N}_2\text{(g)} + \text{O}_2\text{(g)}$ $\Delta n_g = 0$

QUESTION – (AIIMS 2018, 27 May)

K_c for the reaction $\text{N}_2(\text{g}) + \text{O}_2(\text{g}) \rightleftharpoons 2\text{NO}(\text{g})$ at 300 K is 4.0×10^{-6} . K_p for the above reaction will be ($R = 2 \text{ cal mol}^{-1} \text{ K}^{-1}$)

- ☐ A 2.4×10^{-3}
- ☒ B 4×10^{-6}
- ☐ C $4 \times 10^{-6} (RT)^2$
- ☐ D 16×10^{-12}

$T = 300 \text{ K}$ $K_c = 4 \times 10^{-6}$
 $\Delta n_g = 2 - 2 = 0$ $K_p = K_c (RT)^{\Delta n_g}$
 $K_p = K_c$



Characteristics of Equilibrium Constant



MIT

① K_c high \Rightarrow Product conc. \uparrow \therefore product more stable
 K_c low \Rightarrow reactant \uparrow \therefore reactant more stable

② While writing eq. Constt. \Rightarrow we write only gaseous substance or aq. solⁿ, don't write pure solids & liquids \rightarrow active mass Constt.

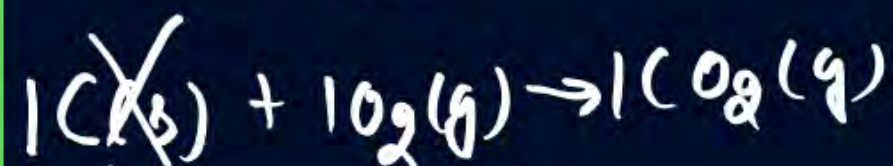
active mass = conc. = $\frac{n}{V} = \frac{m}{M \times V} = \frac{\text{density}}{M}$ \rightarrow pure solid or liquid Constt

③ unit of $K_c = (\text{mol L}^{-1})^{\Delta n_{\text{g or sol}^n}}$

$$K_p = (\text{atm})^{\Delta n_{\text{g or sol}^n}}$$



$$K_c = \frac{[B]}{[A]}$$



$$K_c = \frac{[CO_2]}{[O_2]}$$

$$K_p = \frac{(P_{CO_2})}{(P_{O_2})}$$

$$d = \frac{PM}{RT}$$



$$\Delta n_g = 2 - 4 = -2$$

$$K_c = (\text{mol L}^{-1})^{-2}$$

$$= \text{L}^2 / \text{mol}^2$$

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

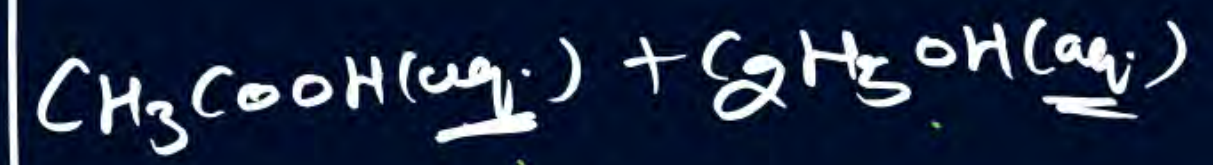
$$= \frac{(\text{mol L}^{-1})^2}{(\text{mol L}^{-1})^4}$$



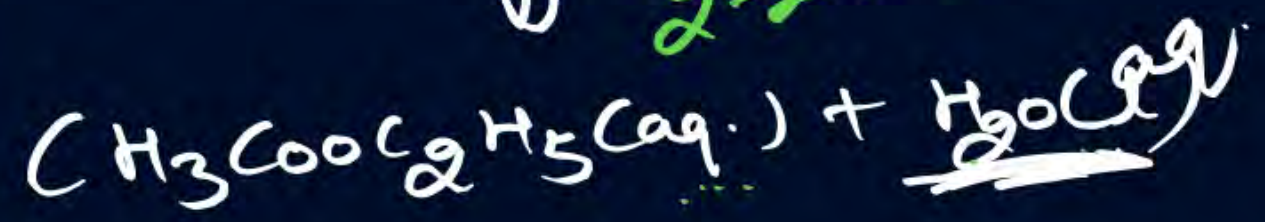
$$K_c = \frac{1}{[\text{H}_2]^2 [\text{O}_2]}$$

$$K_p = \frac{1}{(\text{P}_{\text{H}_2})^2 (\text{P}_{\text{O}_2})}$$

$$= (\text{mol L}^{-1})^{-2}$$



$$\downarrow 2-2=0$$



$$K_c = \frac{[\text{CH}_3\text{COOC}_2\text{H}_5][\text{H}_2\text{O}]}{[\text{CH}_3\text{COOH}][\text{C}_2\text{H}_5\text{OH}]^2}$$

↓
No unit

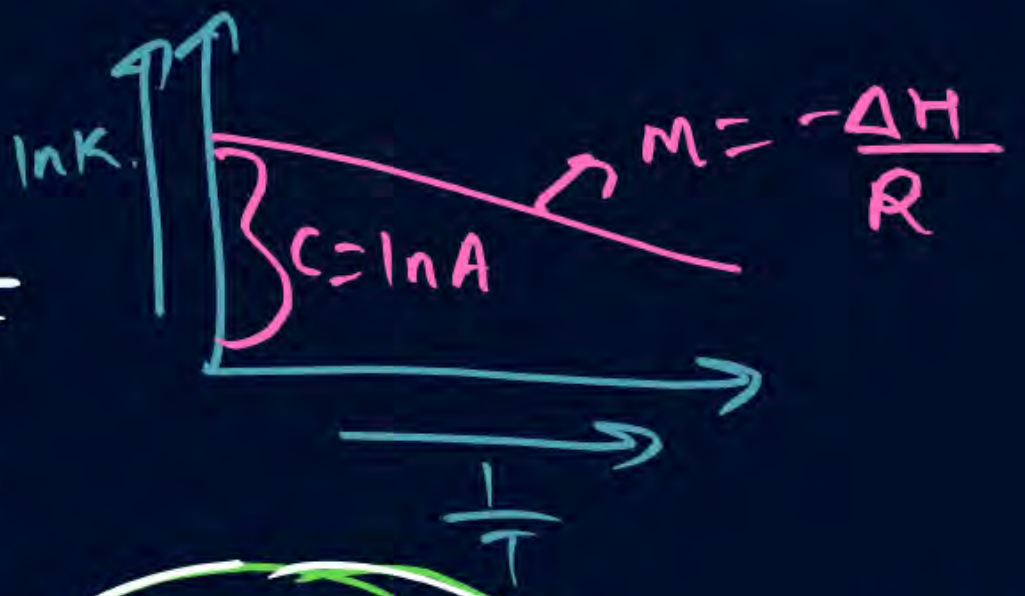
#MIT

④ K_p or K_c change only when T is changed.

$$K = A e^{-\frac{\Delta H}{RT}}$$

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

$$y = c + mx$$



$$\log_{10} \frac{K_2}{K_1} = \frac{\Delta H}{2.303 R} \left(\frac{T_2 - T_1}{T_1 T_2} \right)$$

$$\left(\frac{T_2 - T_1}{T_1 T_2} \right)$$

Van't Hoff eqⁿ.

⑤ (a) endothermic orⁿ.

$$K \propto T$$

$$\begin{matrix} T \uparrow & K \uparrow \\ T \downarrow & K \downarrow \end{matrix}$$

(b) exothermic orⁿ.

$$K \propto \frac{1}{T}$$

$$\begin{matrix} T \uparrow & K \downarrow \\ T \downarrow & K \uparrow \end{matrix}$$

~~endo~~ exo
 $\Delta H = \text{exothermic} (-)ve$

$$T_2 > T_1$$

$$\log \frac{K_2}{K_1} = -x$$

$$\log K_2 = \log K_1 - x$$

$$K_2 < K_1$$

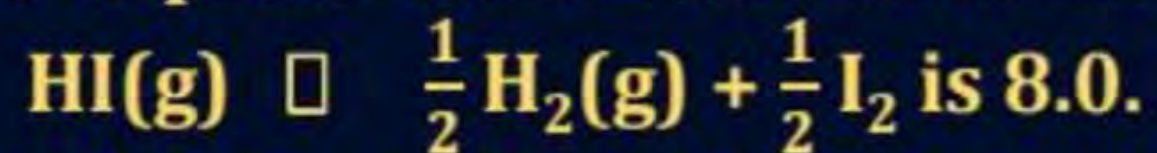


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QUESTION



The value of equilibrium constant of the reaction:



The equilibrium constant of the reaction



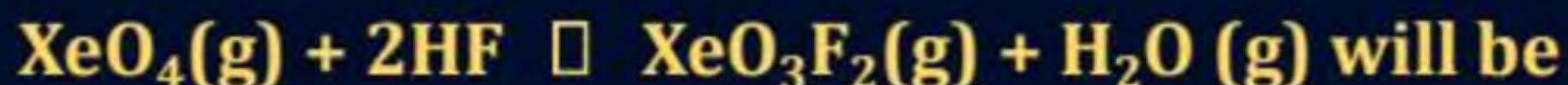
- A** 16
- B** 1/8
- C** 1/16
- D** 1/64

QUESTION – (AIIMS 2013)

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



The equilibrium constant for the reaction



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

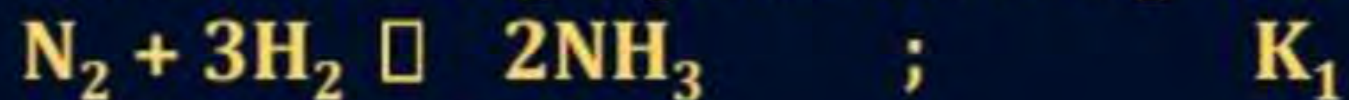
B $K_1 \cdot K_2$

C $\frac{K_1}{K_2}$

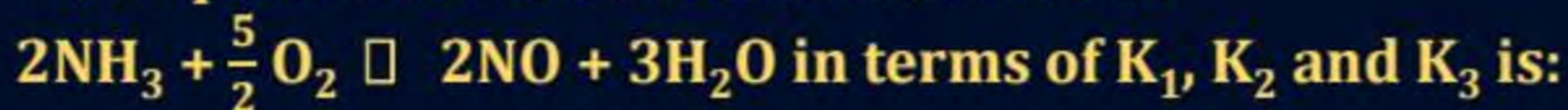
D $\frac{K_2}{K_1}$

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 $K_1 K_2 K_3$

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 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:

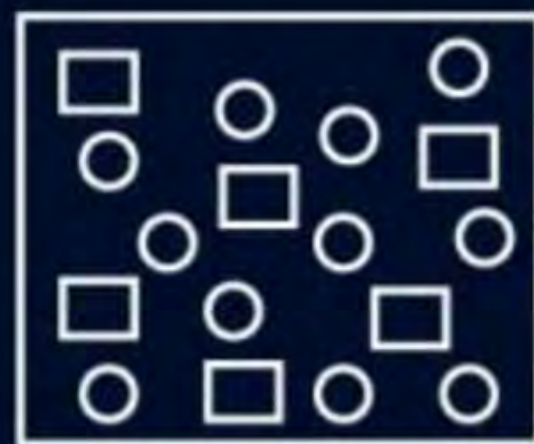
- A** 3.0×10^5
- B** 3.0×10^{-5}
- C** 3.0×10^{-4}
- D** 3.33×10^4

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



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** $\frac{P_{\text{H}_2}^2}{P_{\text{H}_2\text{O}}^2}$ **B** $\frac{(P_{\text{H}_2})^4}{(P_{\text{H}_2\text{O}})^4}$
- C** $\frac{(P_{\text{H}_2})^4 [\text{Fe}_3\text{O}_4]}{(P_{\text{H}_2\text{O}})^4 [\text{Fe}]}$ **D** $\frac{[\text{Fe}_3\text{O}_4]}{[\text{Fe}]}$

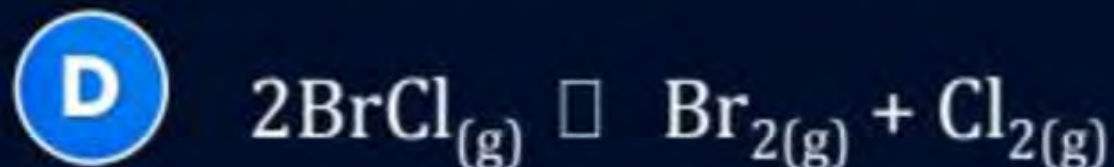
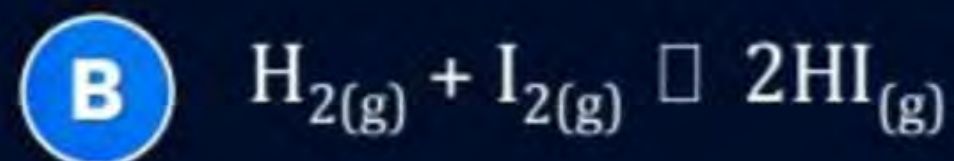


Magarmach Practice Questions (MPQ)



QUESTION – (NEET 2024)

In which of the following equilibria, K_p and K_c are NOT equal?



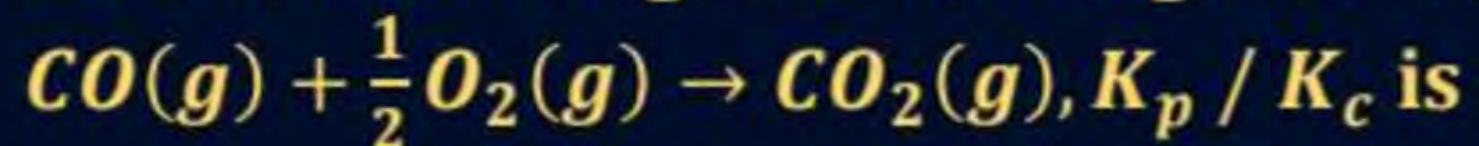
QUESTION – (AIIMS 2005)

For reaction, $2\text{NOCl(g)} \rightleftharpoons 2\text{NO(g)} + \text{Cl}_2\text{(g)}$; K_c at 427°C is $3 \times 10^{-6} \text{ L mol}^{-1}$. The value of K_p is nearly:

- A** 7.50×10^{-5}
- B** 2.50×10^{-5}
- C** 2.50×10^{-4}
- D** 1.75×10^{-4}

QUESTION – (AIIMS 2017)

For the following reaction in gaseous phase



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

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

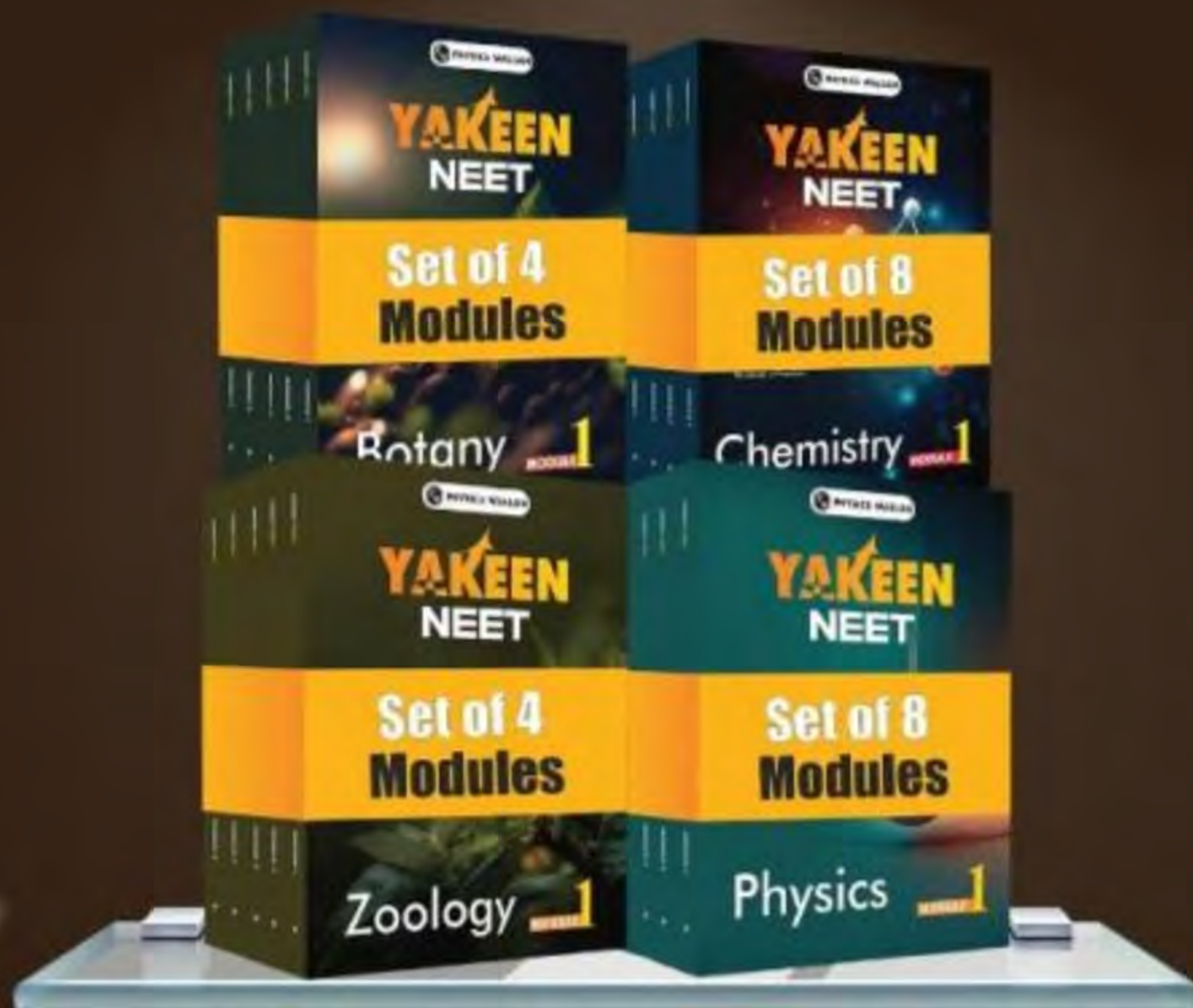


- A** 1
- B** 0.5
- C** 1.5
- D** 2



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