



Al-Mustaqbal-College University
Chemical Engineering and Petroleum
Industry Department
Analytical chemistry
First class / first term
Lecture Three part 2

By

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Lecture three

3.4 The Equilibrium Constant

For a reaction: $aA + bB \leftrightarrow cC + dD$

equilibrium constant: $K = \frac{[C]^c[D]^d}{[A]^a[B]^b}$

The **equilibrium constant, K_c** , is the ratio of the equilibrium concentrations of products over the equilibrium concentrations of reactants each raised to the power of their stoichiometric coefficients.

What Does the Value of K Mean?



$K \gg 1$, equilibrium "lies to the right"

- If $K \gg 1$, the reaction is *product-favored*; product predominates at equilibrium.



$K \ll 1$, equilibrium "lies to the left"

- If $K \ll 1$, the reaction is *reactant-favored*; reactant predominates at equilibrium.

*When $10^{-3} < K < 10^3$, the reaction is considered to contain a significant amount of both reactants and products at equilibrium.

Note : if $K > 1$ the reaction is forward (product to right)
If $K < 1$ the reaction is reverse (reactant to left)

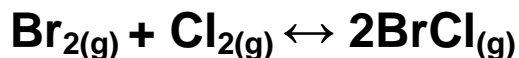
3.4.1 Calculating Equilibrium Concentrations

Use ICE Tables To Solve Equilibrium Problems For K_c Or Equilibrium Amounts:

- 1. I = initial concentration:** Initial concentration of reactants are usually given; initial [Product]'s are assumed to be 0 unless otherwise specified.
- 2. C = change in concentration:** Assign change as the variable x ; use the stoichiometry of the reaction to assign changes for all species.
- 3. E = equilibrium concentration:** $E = I + C$

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Example 1 \\ In an analysis of the following reaction at 100°C



the equilibrium concentrations are $[\text{Br}_2] = 2.3 \times 10^{-3} \text{M}$, $[\text{Cl}_2] = 1.2 \times 10^{-2} \text{M}$, $[\text{BrCl}] = 1.4 \times 10^{-2} \text{M}$. Write the equilibrium expression and calculate K_c for this reaction.

Solution:

$$K_c = \frac{[\text{BrCl}]^2}{[\text{Br}_2][\text{Cl}_2]}$$

$$K_c = \frac{[1.4 \times 10^{-2}]^2}{[2.3 \times 10^{-3}][1.2 \times 10^{-2}]} = 7.1$$

Example 2 \\ The chemicals A and B react as follows to produce C and D:

$A + B \leftrightarrow C + d$, $K = \frac{[D][C]}{[A][B]}$, The equilibrium constant K has a value of 0.30. Assume 0.20 mol of A and 0.50 mol of B are dissolved in 1.00 L, and the reaction proceeds. Calculate the concentrations of reactants and products at equilibrium.

Solve :

	[A]	[B]	[C]	[D]
Initial	0.20	0.50	0	0
Change ($x = \text{mmol/mL}$ reacting)	$-x$	$-x$	$+x$	$+x$
Equilibrium	$0.20 - x$	$0.50 - x$	x	x

We can substitute these values in the equilibrium constant expression and solve for x :

$$K = \frac{[D][C]}{[A][B]}$$

$$0.3 = \frac{(x)(x)}{(0.2-x)(0.5-x)}$$

$$x^2 = (0.10 - 0.70x + x^2)0.30$$

$$0.70x^2 + 0.21x - 0.030 = 0$$

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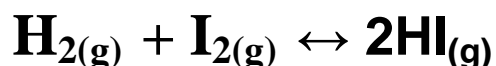
$$\begin{aligned}x &= \frac{-b \pm \sqrt{b^2 - 4ac}}{2a} \\&= \frac{-0.21 \pm \sqrt{(0.21)^2 - 4(0.70)(-0.030)}}{2(0.70)} \\&= \frac{-0.21 \pm \sqrt{0.044 + 0.084}}{1.40} = 0.11 M\end{aligned}$$

$$[A] = 0.20 - x = 0.09M$$

$$[B] = 0.50 - x = 0.39M$$

$$[C] = [D] = x = 0.11M$$

Example 3 \\ Determine the initial concentration of HI if the initial concentrations of H_2 and I_2 are both $0.10 M$ and their equilibrium concentrations are both $0.043 M$ at $430^\circ C$. The value of $K_c = 54.3$



Solution:

	$H_2(g)$	+	$I_2(g)$	\rightleftharpoons	$2HI(g)$
Initial	0.10		0.10		y
Change	-x		-x		+2x
Equilibrium	0.043		0.043		y + 2x

First solve for x: $0.10 - x = 0.043$; $x = 0.057$

Then solve for y: at equilibrium we have

$$K_c = 54.3 = \frac{[HI]^2}{[H_2][I_2]} = \frac{(y + 0.114)^2}{(0.043)(0.043)}$$

$$(y + 0.114)^2 = (54.3)(0.043)^2 \Rightarrow y = \sqrt{(54.3)(0.043)^2} - 0.114 = 0.203$$

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Example 4 A closed system initially containing 1×10^{-3} M H_2 and 2×10^{-3} M I_2 at 448°C is allowed to reach equilibrium. Analysis of the equilibrium mixture shows that the concentration of HI is 1.87×10^{-3} M. Calculate K_c at 448°C for the

reaction taking place, which is : $H_{2(g)} + I_{2(g)} \leftrightarrow 2HI_{(g)}$

solve :

1)	H_2	I_2	$2HI$
initial	1×10^{-3}	2×10^{-3}	0
change			
equilibrium			1.87×10^{-3}

2)	H_2	I_2	$2HI$
initial	1×10^{-3}	2×10^{-3}	0
change	-X	-X	+2X
equilibrium	$1 \times 10^{-3} - X$	$1 \times 10^{-3} - X$	1.87×10^{-3}

$$0 + 2X = 1.87 \times 10^{-3} \rightarrow X = 9.35 \times 10^{-4} \text{ M} = [HI]$$

3)	H_2	I_2	$2HI$
initial	1×10^{-3}	2×10^{-3}	0
change	-9.35×10^{-4}	-9.35×10^{-4}	1.87×10^{-3}
equilibrium	$1 \times 10^{-3} - 9.35 \times 10^{-4}$	$1.87 \times 10^{-3} - 9.35 \times 10^{-4}$	1.87×10^{-3}

$$[H_2] = 1 \times 10^{-3} - 9.35 \times 10^{-4} = 6.5 \times 10^{-5} \text{ M}$$

$$[I_2] = 1.87 \times 10^{-3} - 9.35 \times 10^{-4} = 1.065 \times 10^{-3} \text{ M}$$

$$K = \frac{[HI]^2}{[H_2][I_2]} \rightarrow \frac{[9.35 \times 10^{-4}]^2}{[6.5 \times 10^{-5}][1.065 \times 10^{-3}]} = 51$$