Teaching and Learning Center
university of washington tacoma

## Solving Equilibrium Problems

We are able to group equilibrium problems into two types:

1) We have been given equilibrium concentrations (or partial pressures) and must solve for $K$ (equilibrium constant).
2) We have been given $K$ and the initial concentrations and must solve for the equilibrium concentrations.

For the first type of equilibrium problem, we can to solve for $K$ by directly substituting given equilibrium quantities into the reaction quotient:

For example, let's use the following reaction:
$\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \longleftrightarrow 2 \mathrm{NH}_{3}(\mathrm{~g})$

At equilibrium, the above reaction contains 2.25 M of $\mathrm{N}_{2}, 5.00 \mathrm{M}$ of $\mathrm{H}_{2}$, and 3.5 M of $\mathrm{NH}_{3}$. Calculate the equilibrium constant.

Because we have been given the equilibrium concentrations of each reactant AND product, we can simply substitute these quantities into the reaction quotient:

$$
Q_{c}=\frac{\left[N H_{3}\right]^{2}}{\left[N_{2}\right]\left[H_{2}\right]^{3}} \quad \text { therefore, } \quad K_{C}=\frac{(3.50)^{2}}{(2.25)(5.00)^{3}}=0.218
$$

We can also solve for $K$ if we have only been given some quantities, but not all. For instance, if you were given the initial concentrations and equilibrium concentrations you can set up a reaction table or 'ICE' table to help you calculate the equilibrium constant, $K$.

For example, let's use the following reaction and quantities to solve for $K$.

The decomposition of nitrogen oxide is shown by the reaction below:
$2 \mathrm{NO}(\mathrm{g}) \longleftrightarrow \mathrm{N}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g})$
This reaction was studied at 298 K with initial amount of 0.215 M of NO gas. At equilibrium, the concentration of NO was 0.083 M . Calculate $K_{c}$ for this reaction.

It is important to recognize that we have initial and equilibrium amounts for NO, but we don't know the equilibrium amounts of our products, $\mathrm{N}_{2}$ and $\mathrm{O}_{2}$. When we don't know some of our equilibrium amounts, we must set up a reaction or ICE table.

| Concentration (M) | $\mathbf{2 N O}(\mathbf{g})$ | $\longleftrightarrow$ | $\mathbf{N}_{2}(\mathbf{g})$ | + |
| :--- | :--- | :---: | :---: | :---: |
| Initial | 0.215 M | 0 | $\mathbf{O}_{2}(\mathbf{g})$ |  |
| Change | -2 x | +x | 0 |  |
| Equilibrium | $0.215-2 \mathrm{x}$ |  | x | +x |

In order to solve for $K_{c}$, we need equilibrium concentrations for all reactants and products. Based on the balanced equation, we know that when $2 x$ moles of NO reacts, x moles of $\mathrm{N}_{2}$ and $\mathrm{O}_{2}$ will form. We also were given the equilibrium concentration for NO ( 0.083 M ), so we can solve for x :
$0.215-2 x=0.083 \mathrm{M}$

Solve for x :
$x=(0.083 M-0.215 M) /-2$
$x=0.066 M$

Now that we have determined x, we can substitute the concentration into $Q_{c}$ :

$$
Q_{c}=\frac{\left[N_{2}\right]\left[O_{2}\right]}{[N O]^{2}} \quad \text { therefore, } \quad K_{C}=\frac{(0.066)(0.066)}{(0.083)^{2}}=0.632
$$

The second type of equilibrium problem you may encounter will give you both initial concentrations and $K$ and then ask you to solve for the equilibrium concentrations.

Let's examine the reaction involving the decomposition of HI :
$2 \mathrm{HI}(\mathrm{g}) \longleftrightarrow \mathrm{I}_{2}(\mathrm{~g})+\mathrm{H}_{2}(\mathrm{~g}) \quad K_{c}=0.67$

If 3.0 M of HI is placed in a flask, what is the equilibrium concentration of each product and reactant ?
Because we were given initial amounts, we must complete an ICE table:

| Concentration (M) | $\mathbf{2 H I}(\mathbf{g})$ |  | $\longrightarrow$ | $\mathbf{I}_{\mathbf{2}}(\mathbf{g})$ | $\mathbf{+}$ |
| :--- | :--- | :--- | :--- | :--- | :--- |
| Initial | 3.0 |  | 0 | $\mathbf{H}_{\mathbf{2}}(\mathbf{g})$ |  |
| Change | -2 x |  | x | 0 |  |
| Equilibrium | $3.0-2 \mathrm{x}$ |  | x | +x |  |

We were given the equilibrium constant for this reaction ( $K_{c}=0.67$ ), so we can set up our reaction quotient:
$Q_{c}=\frac{\left[I_{2}\right]\left[H_{2}\right]}{[H I]^{2}} \quad$ therefore $K_{c}=0.67=\frac{(x)^{2}}{(3.0-2 x)^{2}}$

We can take the square root of each side of the equation:

$$
\sqrt{0.67}=\frac{x}{3.0-2 x}
$$

Next, we can multiply each side by (3.0-2x):
$2.46-1.64 x=x$

So, $2.46=2.64 x$ therefore $x=0.93$

Now that we have solved for $x$, we can calculate the equilibrium concentrations.
$\left[I_{2}\right]=\left[H_{2}\right]=x=0.93 \mathrm{M}$
$[\mathrm{HI}]=3.0-2(0.93)=1.14 \mathrm{M}$

