## WJEC Chemistry A-level

## 3.8: Equilibrium Constants Detailed Notes <br> Welsh Specification



## The Kc Constant

For reversible reactions, there is an equilibrium constant, Kc , that indicates the position of equilibrium for a reaction at a certain temperature. Kc for a given reaction always has the same value unless the reaction conditions, and therefore the position of equilibrium, is changed.

## Calculating Kc

In order to find the value of Kc , the concentrations of the species present in the reaction at the point of equilibrium needs to be known.
Kc is equal to the concentration of the products divided by the concentration of the reactants - and any variation in the number of moles, raises that species to a power with the same value as the number of moles.

## Example:

$$
\begin{gathered}
2 \mathrm{~A}+3 \mathrm{~B} \rightleftharpoons \mathrm{Y}+2 \mathrm{Z} \\
\mathrm{Kc}=\frac{[\mathrm{Y}][\mathrm{Z}]^{2}}{[\mathrm{~A}]^{2}[\mathrm{~B}]^{3}}
\end{gathered}
$$

This equilibrium constant has varying units depending on the chemical reaction. It can be calculated by substituting the units into the Kc expression. Some of these then cancel to give the overall units of Kc for that reaction. The units for the above reaction can be calculated to be $\mathrm{mol}^{-2} \mathrm{dm}^{6}$. If the units cancel completely then Kc has no units.

## Example:

Consider the following reaction:

$$
2 A+3 B \rightleftharpoons D+4 E
$$

Then we have:

$$
K c=\frac{[D][E]^{4}}{[A]^{2}[B]^{3}}
$$

To work out the units, we substitute units into each position of $A, B, D$ and $E$ :

$$
\mathrm{Kc}=\left[\left(\mathrm{moldm}^{-3}\right)\left(\mathrm{moldm}^{-3}\right)^{4}\right] /\left[\left(\mathrm{moldm}^{-3}\right)^{2}\left(\mathrm{moldm}^{-3}\right)^{3}\right]
$$

All the units cancel so Kc has no units.

## Temperature Changes and Kc

The value of Kc is not affected by concentration change or use of a catalyst, however it is affected by changing the reaction temperature. This is because the equilibrium position will change, resulting in different concentrations of reactants and products.

## Example:

$$
2 A+3 B \rightleftharpoons Y+2 Z \quad[\Delta \mathrm{H}=+30.7 \mathrm{KJ} / \mathrm{mol}]
$$

In this reaction, the forward reaction is endothermic. This means that an increase in temperature will favour the forward reaction, and will increase the production of products.

This will cause the value of Kc to increase as the concentration of the products at equilibrium will increase.


Similarly, if the temperature is changed so that the production of reactants is favoured, the value of Kc will decrease as the concentration of reactants increases (dividing by a bigger number will decrease the outcome number).

## The Kp Constant

Kp is the equilibrium constant used for gaseous equilibria. All reactants and products must be in a gaseous state in order for Kp to be calculated.

## Calculating Kp

Within a gaseous system, each gas has a partial pressure. These partial pressures allow the value of Kp for a gaseous equilibrium to be found. The sum of these partial pressures gives the total system pressure.

The molar fraction of a substance and the total pressure of the system are used to calculate partial pressures. The units for partial pressure are generally $\mathbf{k P a}$ - the same units of the total pressure.

$$
\underset{\underset{\text { of } A}{\text { Pressure }}}{\text { Partial }}=\frac{\text { Moles of } A}{\text { Total Moles }} \times \text { Total Pressure }
$$

Kp can then be calculated from the partial pressures of species present in the reaction:

| $2 \mathrm{~A}_{(\mathrm{g})}+3 \mathrm{~B}_{(\mathrm{g})}$ | $\rightleftharpoons \mathrm{Y}_{(\mathrm{g})}+2 \mathrm{Z}_{(\mathrm{g})}$ |
| ---: | :--- |
| Kp | $=\frac{(\mathrm{pY})(\mathrm{pZ})^{2}}{(\mathrm{pA})^{2}(\mathrm{pB})^{3}}$ |

It is similar to Kc in that any variation in moles raises the partial pressure to a power of equal quantity to the number of moles.

Kp also has varying units which are determined in the same way as for Kc - by substitution into the equation for Kp.

## Example:

Consider the following reaction:

$$
A+3 B \rightleftharpoons D+2 E
$$

Then we have:

$$
\mathrm{Kp}=\frac{(\mathrm{pD})(\mathrm{pE})^{2}}{(\mathrm{pA})(\mathrm{pB})^{3}}
$$

To work out the units, we substitute units into each position of $A, B, D$ and $E$ :

$$
\mathrm{Kp}=\left[(\mathrm{kPa})(\mathrm{kPa})^{2}\right] /\left[(\mathrm{kPa})(\mathrm{kPa})^{3}\right]=\mathrm{kPa}^{-1}
$$

The units for Kp are $\mathbf{k P a}^{-1}$.

## Temperature Changes and Kp

The value of Kp for a reaction can change depending on whether the reaction is endothermic or exothermic. This is because changing the temperature could favour the forwards or backwards reaction depending on which way around they are.

Example:
$2 \mathrm{~A}_{(\mathrm{g})}+3 \mathrm{~B}_{(\mathrm{g})} \rightleftharpoons \mathrm{Y}_{(\mathrm{g})}+2 \mathrm{Z}_{(\mathrm{g})}$
$[\Delta \mathrm{H}=-10.4 \mathrm{KJ} / \mathrm{mol}]$

In this reaction, the forwards reaction is exothermic.
This means that an increase in temperature will favour the backwards reaction, and will increase the production of reactants.

This will cause the value of Kp to decrease as the partial pressure of the reactants will increase
(dividing by a bigger number will decrease the outcome number).


Likewise, if the temperature is changed so that the production of products is favoured, the value of Kp will increase as partial pressures of products increase.

