

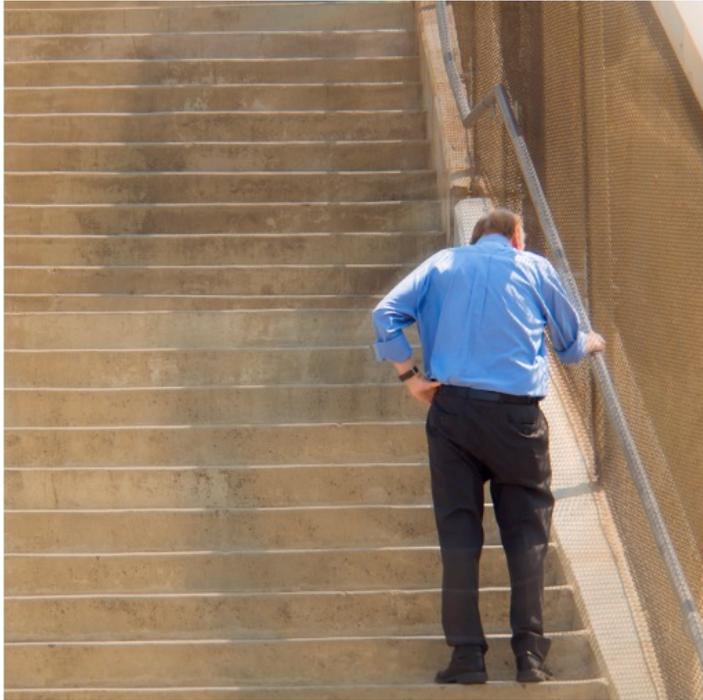


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# EQUILIBRIUM

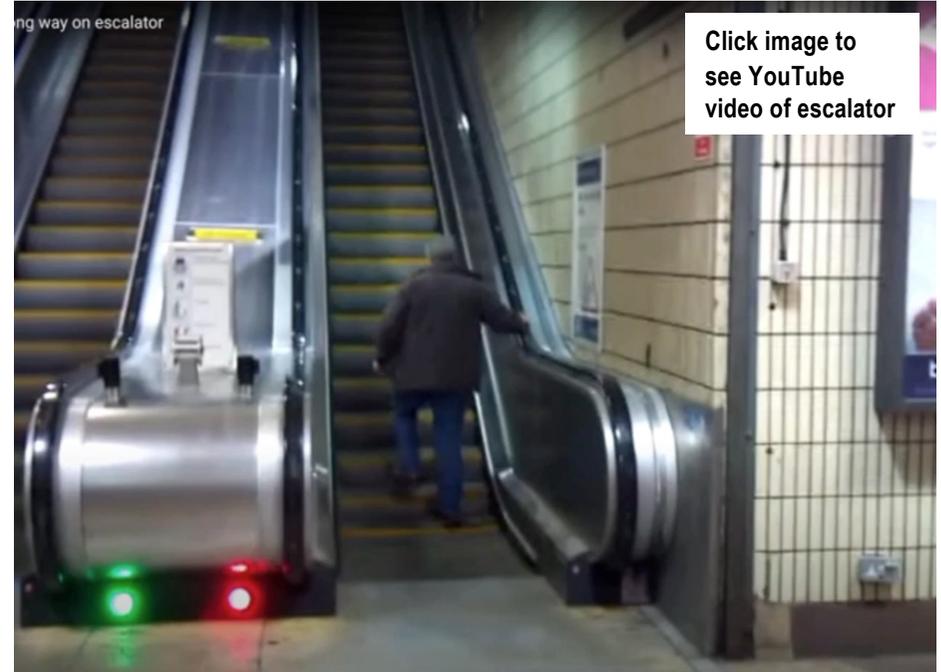
$K_p$

# static equilibrium



person standing still on stairs  
nothing is changing  
at **equilibrium**  
but **STATIC**

# dynamic equilibrium



person walking up a down escalator  
(at same speed as escalator)  
nothing is changing  
at **equilibrium**  
but **DYNAMIC**

# DYNAMIC EQUILIBRIUM



- Reversible reactions can reach a state of dynamic equilibrium in a closed system
- Forward and reverse reactions taking simultaneously and at the same rate
- The concentration of all reactants and products remains constant

# DYNAMIC EQUILIBRIUM



- If an equilibrium **lies to the left**:
  - there are more of the substances on the left side (P&Q) than those on the right side (R)
- If an equilibrium **moves to the left**:
  - more of the backwards reaction (relative to the forwards reaction) takes place until a new equilibrium is reached, meaning once this new equilibrium is reached the amount of the substances on the left (P&Q) have increased and the amount of the substances on the right (R) have decreased

# EQUILIBRIUM CONSTANT $K_c$

- $K_c$  is the equilibrium constant for a system at equilibrium which uses the concentrations (in mol dm<sup>-3</sup>) of the reactants and products.
- $K_c$  is a constant for an equilibrium. The only factor that changes  $K_c$  is temperature.



$$K_c = \frac{[C]^2}{[A]^3 [B]}$$

# EQUILIBRIUM CONSTANT $K_c$

Calculate  $K_c$  at temperature  $T$  for an equilibrium mixture containing 1.2 mol of A, 3.0 mol of B and 1.8 mol of C in a container with volume 2.0 dm<sup>3</sup>



$$K_c = \frac{[C]^2}{[A]^3 [B]} = \frac{\left[\frac{1.8}{2.0}\right]^2}{\left[\frac{1.2}{2.0}\right]^3 \left[\frac{3.0}{2.0}\right]} = 2.5 \text{ mol}^{-2} \text{ dm}^6$$

$$\frac{(\text{mol dm}^{-3})^2}{(\text{mol dm}^{-3})^4} = (\text{mol dm}^{-3})^{-2} = \text{mol}^{-2} \text{ dm}^6$$

# LE CHATELIER'S PRINCIPLE

In simple terms – if the conditions of a system in equilibrium are changed, then the position of the equilibrium moves to oppose that change.

*Gas pressure (if a reaction involves one or more gases)*

if  $P \uparrow$

equilibrium moves in the direction of the side with fewer gas molecules to lower  $P$

equilibrium moves, but  $K_c/K_p$  remains constant.

# LE CHATELIER'S PRINCIPLE

In simple terms – if the conditions of a system in equilibrium are changed, then the position of the equilibrium moves to oppose that change.

## *Temperature*

if  $T \uparrow$

equilibrium moves in the endothermic direction to lower  $T$

equilibrium moves, and  $K_c/K_p$  change

# LE CHATELIER'S PRINCIPLE

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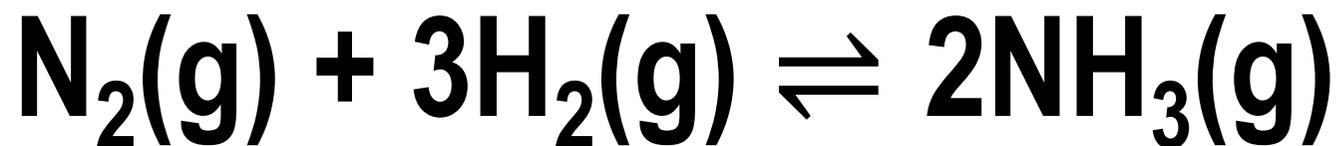
## *Concentration*

if more of a chemical is added

equilibrium moves to use up that chemical in reaction

equilibrium moves, but  $K_c/K_p$  remains constant

# $K_p$



$$K_p = \frac{(p\text{NH}_3)^2}{(p\text{N}_2) (p\text{H}_2)^3}$$

$$\frac{\text{kPa}^2}{\text{kPa}^4} = \text{kPa}^{-2}$$

# MOLE FRACTION

If you have 10 moles of a mixture of gases, of which 3 moles is O<sub>2</sub>, then the mole fraction of O<sub>2</sub> in the mixture is  $\frac{3}{10}$  or 0.3

(or even 30%, but it is not usually expressed as a %)

$$\text{mole fraction of gas A in a mixture of gases} = \frac{\text{moles of gas A}}{\text{total moles of gas in the mixture}}$$

Note that the sum of all the mole fractions in a mixture should add up to 1

# PARTIAL PRESSURE

Imagine a mixture of gases with a total pressure of 100 kPa.

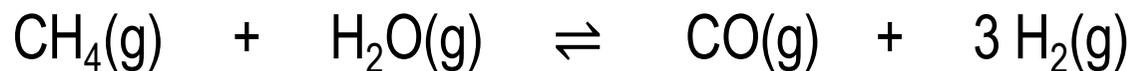
If the mole fraction of gas A in that mixture is  $\frac{3}{10}$ , then that gas makes up 30 kPa (i.e.  $\frac{3}{10}$ ) of the total pressure.

The contribution that each gas makes to the total pressure is called the partial pressure of that gas; therefore the partial pressure of gas A is 30 kPa.

$$\begin{array}{l} \text{partial pressure of gas A} \\ \text{in a mixture of gases} \end{array} = \text{mole fraction of gas A} \times \text{total pressure}$$

Note that the sum of all the partial pressures is the total pressure.

# PARTIAL PRESSURE



initial moles	2.00	2.00	0	0
change in moles	-1.20	-1.20	+1.20	+3.60
equilibrium moles	0.80	0.80	1.20	3.60
mole fraction	$\frac{0.80}{6.40} = 0.125$	$\frac{0.80}{6.40} = 0.125$	$\frac{1.20}{6.40} = 0.1875$	$\frac{3.60}{6.40} = 0.5625$
partial pressure	$0.125 P$ = 31.9 kPa	$0.125 P$ = 31.9 kPa	$0.1875 P$ = 47.8 kPa	$0.5625 P$ = 143.4 kPa

total moles = 6.40

total mole fractions = 1.00

total partial pressures = 255 kPa

# K<sub>p</sub>

- a) At equilibrium, there are 2.0 moles of A, 2.5 moles of B and 1.5 moles of C. The total pressure is 500 kPa. Calculate the value of K<sub>p</sub>



equilibrium moles	2.0	2.5	1.5	total = 6.0
mole fraction	$\frac{2.0}{6.0}$	$\frac{2.5}{6.0}$	$\frac{1.5}{6.0}$	total = 1.0
partial pressure	$\frac{2.0}{6.0} \times 500$ = 166.7 kPa	$\frac{2.5}{6.0} \times 500$ = 208.3 kPa	$\frac{1.5}{6.0} \times 500$ = 125 kPa	total = 500

$$K_p = \frac{(p_C)^2}{(p_A)^2 (p_B)} = \frac{125^2}{166.7^2 \times 208.3} = 0.0027 \text{ kPa}^{-1}$$

$$\frac{\text{kPa}^2}{\text{kPa}^3} = \text{kPa}^{-1}$$

# $K_p$

b) If the total pressure of this equilibrium was increased, explain what would happen to the yield of C and the value of  $K_p$



- equilibrium shifts right
- to side with fewer gas molecules
- to lower the pressure
- so yield of C increases
- no change to  $K_p$  as it is a constant only affected by temperature

# $K_p$

c) The forward reaction is exothermic. If the temperature of this equilibrium was increased, explain what would happen to the yield of C and the value of  $K_p$



- equilibrium shifts left
- in endothermic direction
- to lower the temperature
- so yield of C decreases
- $K_p$  decreases