

Rate of Reaction

- **Rate of reaction measures** →
- How fast reactants are used up and how fast products are made.
- **Rate ($\text{Mol dm}^{-3} \text{ s}^{-1}$)** = $\frac{\text{change in concentration of reactant/product}}{\text{time for change to take place}}$
- Concentration is represented by square brackets around the molecule.
- **As the reaction proceeds, concentration decreases, leading to** →
- Fewer collisions per second between reactant particles.
- The rate slowing down.
- **Rate of reaction is determined by measuring the concentration of a substance at time intervals** →
- Rate is equal to the slope of the curve.
- Tangent is drawn – gradient is calculated.
- **Initial rate** → the change in concentration of a reactant, of product, per unit time at the start of the reaction when $t=0$.

Half-Lives

- **Exponential decay** →
- Many natural processes have a constant half-life – value halves every half-time.
- **Radioactive decay** →
- Constant half-life of a radioactive element is the time taken for half the element to decay.
- Used to measure element stability and in radio-carbon dating.

Equilibrium Constant

- A **dynamic equilibrium** is established in a closed system when →
- The rate of the forward reaction is the same as the reverse reaction.
- The concentrations of the reactants and products remain the same.
- **Equilibrium law** → the relative proportions of reactants and products present at equilibrium.



- **K_c** → the equilibrium constant for concentrations.

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

[A] = concentration of A in mol dm⁻³
a = number of moles of A

- **Homogeneous equilibrium** → all species making up the reactants and products are in the same physical states.
- **Heterogeneous equilibrium** → species are in different states.

Strong & Weak Acids

- The strength of an acid (HA) is the extent of its dissociation into H^+ and A^- ions.
- **Strong acids dissociate completely in aqueous solution.**
- **Examples** →
 - HCl , HNO_3 , H_2SO_4 , HBr , HI , HClO_4 .
- **Weak acids only partially dissociate.**
- **Example:** $\text{CH}_3\text{COOH} \leftrightarrow \text{H}^+ + \text{CH}_3\text{COO}^-$.
- Equilibrium lies well to the left.
- **Acid dissociation constant (K_a)** → measures the extent of acid dissociation.
- **Weak acid:** $\text{HA} \leftrightarrow \text{H}^+ + \text{A}^-$.
- A large K_a value → large extent of dissociation → a strong acid.

- $\text{p}K_a = -\log K_a$
- $K_a = 10^{-\text{p}K_a}$

$$K_a = \frac{[\text{H}^+][\text{A}^-]}{[\text{HA}]}$$

$$K_a = \frac{\overset{\text{simplified}}{[\text{H}^+]^2}}{[\text{HA}]}$$

Buffer Solutions

- **Buffer solution** → a mixture that minimises pH changes on addition of **small** amounts of acid or base.
- **A buffer solution is a mixture of** →
 - A weak acid, HA
 - Its conjugate base, A⁻.
- **A buffer can be made from a weak acid and the salt of a weak acid.**
- **Example** →
 - In the CH₃COOH/CH₃COONa buffer system →
 - The weak acid dissociates partially: $\text{CH}_3\text{COOH} \leftrightarrow \text{H}^+ + \text{CH}_3\text{COO}^-$.
 - The salt dissociates completely: $\text{CH}_3\text{COONa} \rightarrow \text{CH}_3\text{COO}^- + \text{Na}^+$.
- **The high concentration of the conjugate base pushes the equilibrium of the weak acid to the left.**
- Therefore, the concentration of H⁺ ions is very small.
- **Buffer equilibrium:** $\text{HA} \leftrightarrow \text{H}^+ + \text{A}^-$.
- The weak acid (HA) removes added alkali.
- The conjugate base (A⁻) removes added acid.

Born-Haber Cycles

- The cycle begins with the elements in their standard states at zero energy.
- This is known as the datum line.
- **Endothermic** → ΔH points upwards.
- **Exothermic** → ΔH points downwards.
- The ionic solid is on the lowest energy level.

Redox

- **Oxidation** → loss of electrons.
- **Reduction** → gain of electrons.
- **Oxidising agent** → is reduced and accepts electrons.
- **Reducing agent** → is oxidised and donates electrons.

- **Balancing redox equations** →
- Complete half equations.
- Sometimes add H^+ , OH^- or H_2O .
- In acidic conditions, add H^+ .
- In alkaline, add OH^- .

Storage & Fuel Cells

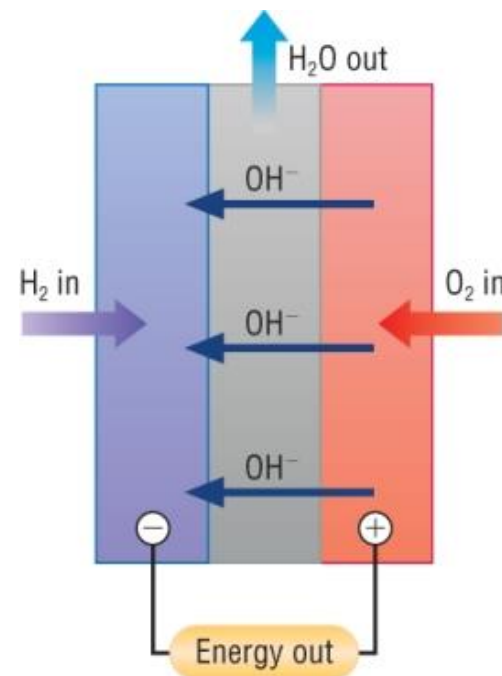
- **Fuel cells →**

- The hydrogen-oxygen fuel cell uses energy from the reaction of a fuel with oxygen to create a voltage.
- The reactants flow in and products flow out while the electrolyte remains in the cell.
- Fuel cells can operate continuously so long as the fuel and oxygen continue to flow into the cell.
- Electricity is generated when oxygen and hydrogen make water.

- **Hydrogen –Oxygen Fuel Cell Equations →**

- $\text{H}_2 + 2\text{OH}^- \rightarrow 2\text{H}_2\text{O} + 2\text{e}^-$
- $\frac{1}{2} \text{O}_2 + 2\text{H}_2\text{O} + 2\text{e}^- \rightarrow 2\text{OH}^-$

- **Overall Equation:** $\text{H}_2 + \frac{1}{2} \text{O}_2 \rightarrow \text{H}_2\text{O}$



Hydrogen for the Future

- **Fuel-cell vehicles (FVC's) →**
- Uses hydrogen gas or hydrogen-rich fuels.
- Hydrogen-rich fuels (e.g. methanol) are mixed with water and converted into hydrogen gas at 250-300 °C by an onboard “reformer”.
- **Using methanol instead of hydrogen gas →**
- Liquid fuel is easier to store than gas.
- Methanol can be generated from biomass.
- Carbon dioxide is waste product – greenhouse gas.
- **Advantages of fuel-cell vehicles →**
- Less pollution and carbon dioxide – hydrogen-rich fuels produce only small amounts of carbon dioxide and air pollutants.
- Greater efficiency – petrol engines are less than 20% efficient in converting chemical energy – hydrogen fuel-cell vehicles are 40-60% efficient.