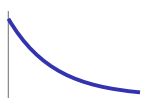


Reaction Kinetics

A-Level Chemistry • Topic 26

A-Level Chemistry



Reaction Kinetics

What we will cover

- 1 Rate equations & orders
- 2 Order graphs
- 3 Half-life & rate constant
- 4 Reaction mechanisms
- 5 Catalysts



measures gas made over time

1

Rate equations

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Rate equations

Rate equations and orders

$$\text{rate} = k[A]^m[B]^n$$

Worked example

doubling $[A]$ doubles the rate (order 1); doubling $[B]$ quadruples it (order 2) $\Rightarrow \text{rate} = k[A][B]^2$, overall order 3

The **order** 反应级数 and **rate constant** 速率常数 k are found by **experiment**, not from the equation.



Catalyst pellets

Reaction Kinetics

Rate equations

Order graphs

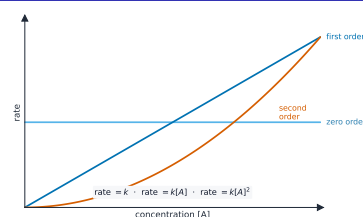
Rate against concentration:

- **zero order** – a flat line
- **first order** – a straight line through the origin
- **second order** – an upward curve

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Rate equations

Order graphs, in detail



Rate against concentration: zero order is a flat line, first order a straight line through the origin, second order an upward curve

Half-life and the rate constant

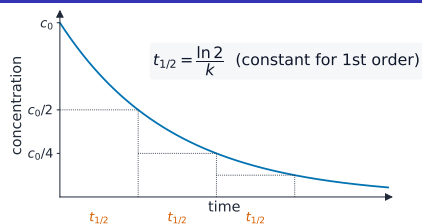
A **first-order** reaction has a **constant half-life** 恒定半衰期 $t_{1/2}$.

$$k = \frac{0.693}{t_{1/2}}$$

$$t_{1/2} = 120 \text{ s}$$

$$k = \frac{0.693}{120} = 5.8 \times 10^{-3} \text{ s}^{-1}$$

A first-order reaction has a constant half-life



A first-order reaction has a constant half-life: the concentration halves in the same time $t_{1/2}$ again and again, whatever the starting value

Half-life and the rate constant, continued

For a first-order reaction the **half-life** 半衰期 is constant:

$$t_{1/2} = \frac{\ln 2}{k} \approx \frac{0.693}{k}$$

A constant half-life on a concentration–time graph **proves** first order. Zero-order half-life gets shorter as concentration falls; second-order half-life gets longer.

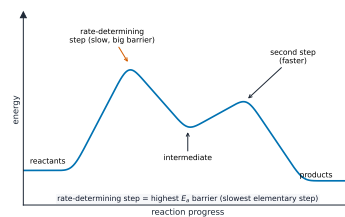
2 Mechanisms and catalysts

Reaction mechanisms

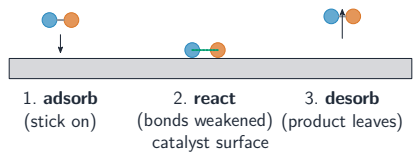
The slowest step is the **rate-determining step** 决速步骤 – it controls the overall rate.

Only species up to & including it appear in the rate equation. An **intermediate** 中间体 is made then used up – not in the overall equation.

Reaction mechanisms, in detail



The slowest step is the **rate-determining step** 决速步骤 – it controls the overall rate.

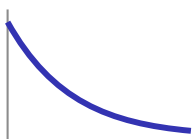
Reaction Kinetics ↳ Mechanisms and catalysts Effect of temperature Raising the temperature increases the rate constant k (more molecules pass the activation energy), so the rate goes up.	Reaction Kinetics ↳ Mechanisms and catalysts Catalysts A heterogeneous catalyst 多相催化剂 works in three stages: ■ adsorption 吸附 onto the surface ■ weakened bonds react ■ desorption 脱附 of the product A homogeneous 均相 one is reformed unchanged.  1. adsorb (stick on) 2. react (bonds weakened) 3. desorb (product leaves) catalyst surface
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Reaction Kinetics ↳ Mechanisms and catalysts The two kinds of catalyst, and the rate vocabulary <table border="1"> <tr> <td>heterogeneous catalyst 多相催化剂</td> <td>a different physical state from the reactants – iron in the Haber process, and platinum, palladium and rhodium in a catalytic converter</td> </tr> <tr> <td>how it works</td> <td>reactants adsorb onto the surface, react there, and desorb</td> </tr> <tr> <td>homogeneous catalyst 均相催化剂</td> <td>the same physical state as the reactants; it is used up in one step and remade in a later one</td> </tr> <tr> <td>order of reaction 反应级数</td> <td>the power to which a concentration is raised in the rate equation</td> </tr> <tr> <td>the overall order 总级数</td> <td>the sum of the individual orders</td> </tr> <tr> <td>the initial rate 初始速率</td> <td>measured at $t = 0$, before any product can interfere – which is why it is used to find orders</td> </tr> </table> Order is found by experiment, never from the balanced equation. That is the single most-tested point in this topic.	heterogeneous catalyst 多相催化剂	a different physical state from the reactants – iron in the Haber process, and platinum, palladium and rhodium in a catalytic converter	how it works	reactants adsorb onto the surface, react there, and desorb	homogeneous catalyst 均相催化剂	the same physical state as the reactants; it is used up in one step and remade in a later one	order of reaction 反应级数	the power to which a concentration is raised in the rate equation	the overall order 总级数	the sum of the individual orders	the initial rate 初始速率	measured at $t = 0$, before any product can interfere – which is why it is used to find orders	Reaction Kinetics ↳ Mechanisms and catalysts Exam tips ■ Find orders from initial-rate data: doubling a concentration that doubles the rate is first order, quadruples it is second order, no change is zero order. ■ Write rate = $k[A]^m[B]^n$ and work out the units of k from it. ■ The rate-determining step contains the species (and orders) in the rate equation – use this to test a mechanism. ■ A constant half-life means first order.
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3 Recap	Reaction Kinetics ↳ Recap Topic 26 – key takeaways <table border="1"> <tr> <td> 1 Rate equation rate = $k[A]^m[B]^n$; found by experiment </td><td> 2 Orders 0 flat, 1 straight, 2 curved </td></tr> <tr> <td> 3 Half-life first order: constant $t_{1/2}$; $k = 0.693/t_{1/2}$ </td><td> 4 Mechanism the rate-determining step controls the rate </td></tr> </table>	1 Rate equation rate = $k[A]^m[B]^n$; found by experiment	2 Orders 0 flat, 1 straight, 2 curved	3 Half-life first order: constant $t_{1/2}$; $k = 0.693/t_{1/2}$	4 Mechanism the rate-determining step controls the rate
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Check yourself

- 1 How is a rate equation found?
- 2 What graph shape is a first-order rate-conc plot?
- 3 What is special about a first-order half-life?
- 4 Which step controls the overall rate?



Answers 1. by experiment 2. a straight line through the origin 3. it is constant
4. the rate-determining step