

Equilibria

A-Level Chemistry • Topic 25

A-Level Chemistry



Equilibria

What we will cover

- 1 Conjugate acids & bases
- 2 pH & the constants
- 3 Calculating pH
- 4 Buffer solutions
- 5 Solubility & partition



pH paper estimates acidity

1 Acids and pH

Equilibria

↳ Acids and pH

Conjugate acids and bases

An acid loses H^+ to give its **conjugate base** 共轭碱; a base gains H^+ to give its **conjugate acid** 共轭酸.

A **conjugate pair** 共轭酸碱对 differs by just one H^+ – e.g. CH_3COOH and CH_3COO^- .

conjugate pair: differ by one H^+



Equilibria

↳ Acids and pH

pH and the constants

$$pH = -\log[H^+]$$

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

$$K_w = [H^+][OH^-]$$

A larger K_a 酸解离常数 (smaller pK_a) is a stronger acid; $K_w = 1.0 \times 10^{-14}$ at 298 K.

Equilibria

↳ Acids and pH

Calculating pH

strong acid – $[H^+] = \text{conc.}$; HCl 0.050 $\Rightarrow$ $pH = 1.30$

strong base – $[H^+] = K_w/[OH^-]$; $NaOH$ 0.10 $\Rightarrow$ $pH = 13.0$

weak acid – $[H^+] = \sqrt{K_a[HA]}$; ethanoic 0.10 $\Rightarrow$ $pH = 2.87$

The four pH calculations

strong acid	fully ionised, so $[H^+]$ is the acid's concentration; $pH = -\log[H^+]$
weak acid	only partly ionised, so use $[H^+] = \sqrt{K_a \times [HA]}$, where K_a is the acid dissociation constant 酸解离常数
strong alkali 强碱	find $[OH^-]$, then use the ionic product of water 水的离子积 $K_w = [H^+][OH^-] = 10^{-14}$
a buffer	$[H^+] = K_a \frac{[HA]}{[A^-]}$
a conjugate pair 共轭酸碱对	an acid and the base left after it donates its proton – the two components of every buffer

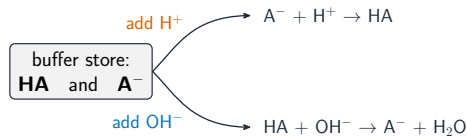
A buffer works because the mixture holds a **store of both partners**: the acid mops up added alkali, and the conjugate base mops up added acid.

2 Buffers and solubility

Buffer solutions

A **buffer** 缓冲溶液 (a weak acid + its conjugate base) resists pH change:

- added H^+ removed by A^-
 - added OH^- removed by HA
- HCO_3^- keeps blood near pH 7.4.



either way, the pH barely changes

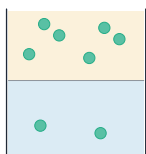
Solubility product



The **solubility product** 溶度积 $K_{sp} = [Ag^+][Cl^-]$ (each ion to the power of its number).

The **common ion effect** 同离子效应: a salt is **less** soluble in a solution already containing one of its ions (Le Chatelier).

Partition coefficients



solvent 1 (non-polar)
 $K_{pc} = \frac{[\text{solute in 1}]}{[\text{solute in 2}]}$
solvent 2 (water)

The **partition coefficient** 分配系数 is the ratio of a solute's concentrations in two immiscible solvents.

$$K_{pc} = \frac{[\text{solute in 1}]}{[\text{solute in 2}]}$$

It depends on the **polarity** 极性 of the solute & solvents.

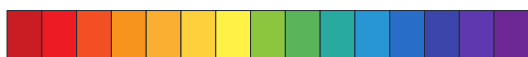
When a partition coefficient applies

what it is	the ratio of a solute 溶质's concentrations in two immiscible solvents at equilibrium
the condition	it holds only when the solute is in the same physical state in both solvents – no dimerising or dissociating
what it depends on	the polarity 极性 of the solute and of the two solvents
the rule	"like dissolves like" – a polar solute favours the polar solvent
where it is used	solvent extraction, and chromatography, where the same partition runs continuously along the plate

Repeated extraction with **small** portions removes more solute than one extraction with the same total volume. That is the calculation the exam sets.

pH and the equilibrium constants

$$\text{pH} = -\log[\text{H}^+]$$



0

acidic

7

neutral

14

alkaline

The pH scale: $\text{pH} = -\log$ of the hydrogen-ion concentration

Exam tips

- $\text{pH} = -\log[\text{H}^+]$; for a **strong acid** $[\text{H}^+] = \text{concentration}$, for a **weak acid** use $[\text{H}^+] = \sqrt{K_a[\text{HA}]}$.
- For a base, get $[\text{H}^+]$ from $K_w = [\text{H}^+][\text{OH}^-] = 1.0 \times 10^{-14}$.
- For a **buffer** use $[\text{H}^+] = K_a \times [\text{acid}]/[\text{salt}]$ and explain how it removes added H^+/OH^- .
- Write the K_{sp} expression with the correct powers; the number of decimal places in a pH equals the significant figures of $[\text{H}^+]$.

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Recap

Worked example – pH of a strong acid

Find the pH of $0.050 \text{ mol dm}^{-3}$ hydrochloric acid.

- HCl is a **strong acid**: it dissociates completely.
- So $[\text{H}^+] = 0.050 \text{ mol dm}^{-3}$ – the concentration itself.
- $\text{pH} = -\log_{10}[\text{H}^+] = -\log_{10}(0.050)$
- = 1.30.

Strong acid: $[\text{H}^+]$ is the concentration.
Weak acid: you must go through K_a – never assume full dissociation.

Topic 25 – key takeaways

1 Conjugate pairs

differ by one H^+

2 pH

$-\log[\text{H}^+]$; K_a , K_w for weak/strong

3 Buffers

weak acid + its salt resist pH change

4 Solubility

K_{sp} ; the common ion effect

Check yourself

- 1 Write the pH formula.
- 2 What does a buffer resist?
- 3 What two things make a buffer?
- 4 What does the common ion effect do to solubility?

Answers 1. $\text{pH} = -\log[\text{H}^+]$ 2. a change in pH 3. a weak acid & its conjugate base 4. lowers it