

5.1 Enthalpy change

Name: _____ Class: _____ Date: _____

Total: 13 marks

Objective

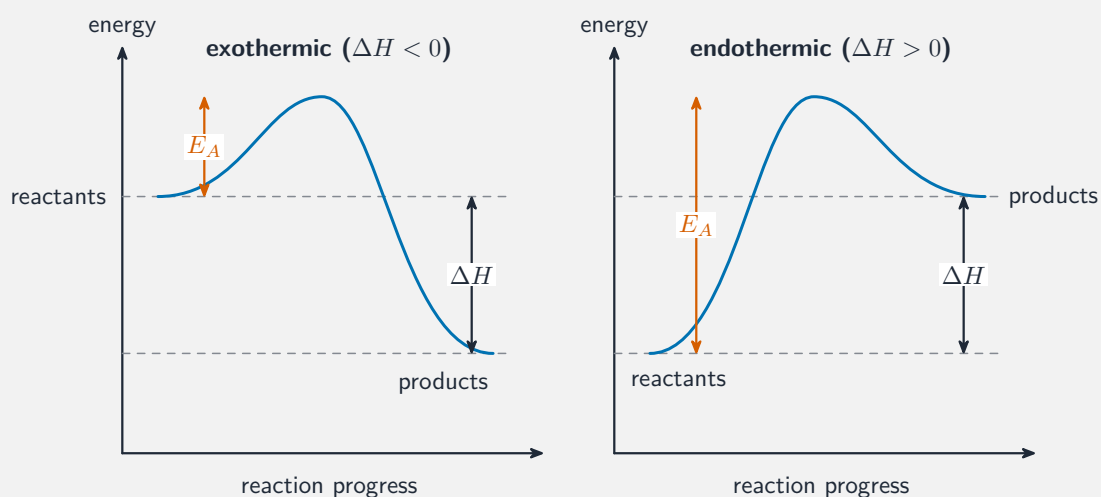
Build the skills to answer exam questions on **enthalpy change** 焓变—**exothermic** 放热 and **endothermic** 吸热 reactions, the standard enthalpy changes, **bond energy** 键能 calculations, and $q = mc\Delta T$.

You must be able to:

- interpret reaction pathway diagrams and define the standard enthalpy changes
- calculate ΔH_r from bond energies
- calculate enthalpy changes from experiment using $q = mc\Delta T$ and $\Delta H = -mc\Delta T/n$

1 Worked examples

■ Exothermic and endothermic

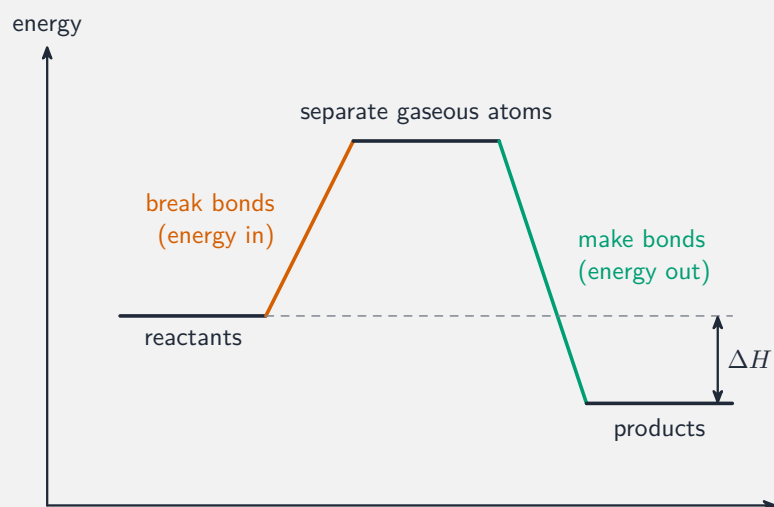


Exothermic: products lower, $\Delta H < 0$. Endothermic: products higher, $\Delta H > 0$. The hill height is the activation energy

Standard conditions: 298 K, 101 kPa ($^\ominus$). Know ΔH_f (formation), ΔH_c (combustion), ΔH_{neut} (neutralisation).

■ Bond energies

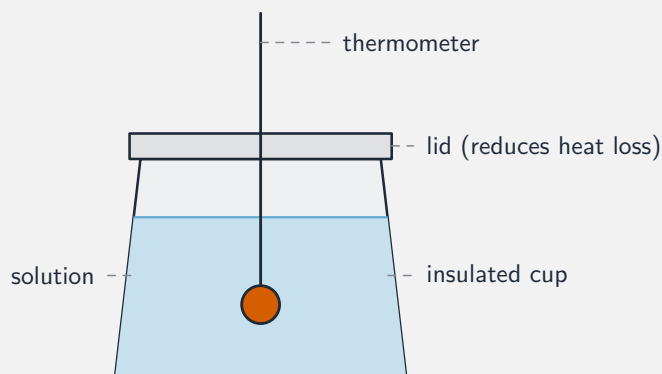
$$\Delta H_r = \sum(\text{bonds broken}) - \sum(\text{bonds made})$$



Breaking bonds takes energy in; making bonds gives energy out

Example. $\text{H}_2 + \text{Cl}_2 \rightarrow 2\text{HCl}$ (H–H 436, Cl–Cl 242, H–Cl 431): $\Delta H = (436 + 242) - (2 \times 431) = -184 \text{ kJ mol}^{-1}$.

■ **Measuring ΔH** ($q = mc\Delta T$)



Measure the temperature change of a known mass of solution

$q = mc\Delta T$; $\Delta H = -q/n$. **Example.** 0.50 g methanol (M_r 32) heats 100 g water by 18 °C ($c = 4.18$): $q = 100 \times 4.18 \times 18 = 7520 \text{ J}$; $n = 0.50/32 = 0.0156$; $\Delta H_c = -7520/0.0156 \approx -480 \text{ kJ mol}^{-1}$.

2 Practice

2.1 State whether ΔH is positive or negative for an exothermic reaction, and what this means for the products' energy. [2]

2.2 Define the **enthalpy change of combustion**.

[2]

3 Exam-style questions

3.1 In an endothermic reaction:

[1]

- **A** ΔH is negative and heat is given out
 - **B** ΔH is positive and heat is taken in
 - **C** the products are lower in energy
 - **D** no bonds are broken
-

3.2 Breaking a chemical bond is:

[1]

- **A** exothermic (releases energy)
 - **B** endothermic (takes in energy)
 - **C** neither
 - **D** always $+431 \text{ kJ mol}^{-1}$
-

3.3 For $\text{N}_2 + 3\text{H}_2 \rightarrow 2\text{NH}_3$, use bond energies (kJ mol^{-1} : N–N 945, H–H 436, N–H 391) to calculate ΔH_r .

[3]

3.4 Burning 0.32 g of a fuel ($M_r = 32$) raises the temperature of 200 g of water by 9.0°C ($c = 4.18$).

(a) Calculate the heat released, q , in J.

[1]

(b) Calculate the enthalpy change of combustion per mole.

[3]

4 Go further

You are now ready for the real exam questions on this subtopic. Open the **5.1 Enthalpy change** past-paper sheet in the Library, or try these in **Practice mode**:

- 9701/11 N25 —Q7 (enthalpy)
- 9701/12 N25 —Q10 (bond energies)
- 9701/33 N25 —Q2 (calorimetry, calculation)

Solutions

2.1 ΔH is negative; the products have less energy than the reactants.

2.2 the enthalpy change when one mole of a substance is burned completely in oxygen under standard conditions.

3.1 B. Endothermic = ΔH positive, heat taken in.

3.2 B. Breaking a bond requires energy (endothermic).

3.3 broken = $945 + 3(436) = 2253$; made = $6 \times 391 = 2346$; $\Delta H = 2253 - 2346 = -93 \text{ kJ mol}^{-1}$.

3.4 (a) $q = 200 \times 4.18 \times 9.0 = 7524 \text{ J}$.

(b) $n = 0.32/32 = 0.010 \text{ mol}$; $\Delta H_c = -q/n = -7524/0.010 = -752 \text{ kJ mol}^{-1}$ (≈ -750).