

Exercise walk-through

Similarities and trends in the properties of the Group 2 metals, magnesium ■ 27.1

eXercise

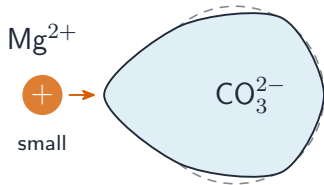
You must be able to

- explain the thermal-stability trend using ionic radius and **polarisation** 极化
- explain the solubility/ ΔH_{sol} trends using lattice energy and hydration enthalpy

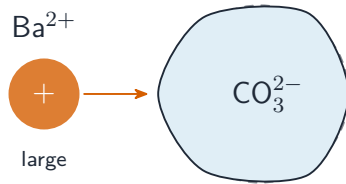
Method — Thermal stability and polarisation

Down the group the cation gets larger $\rightarrow$ lower charge density $\rightarrow$ polarises the anion **less** $\rightarrow$ the carbonate/nitrate is harder to decompose (more stable).

Method — Thermal stability and polarisation



high charge density —
distorts the anion a lot
less stable: decomposes easily



low charge density —
distorts the anion little
more stable: needs high heat

A small Mg^{2+} strongly distorting the carbonate anion; a large Ba^{2+} barely distorting it

Method — Solubility energetics

Hydroxides become **more soluble**, sulfates **less soluble** down the group — because the lattice energy and hydration enthalpies change at different rates as the cation grows.

Method — Solubility energetics

$$\Delta H_{\text{sol}} = -\Delta H_{\text{latt}} + \Delta H_{\text{hyd}}$$

hydroxides (small OH^-):

lattice energy falls *a lot* down the group
 $\Rightarrow$ outweighs the smaller hydration fall

more
soluble

sulfates (large SO_4^{2-}):

lattice energy stays almost constant
 $\Rightarrow$ the hydration fall then dominates

less
soluble

Comparing lattice energy and hydration enthalpy to explain ΔH_{sol} trends

Practice 2.1 ■ 1 mark

State the trend in thermal stability of the Group 2 carbonates down the group.

Solution 2.1

Method: Thermal stability and polarisation

Worked steps

it increases down the group **B1**.

Practice 2.2 ■ 1 mark

State the trend in solubility of the Group 2 sulfates down the group.

Solution 2.2

Method: Solubility energetics

Worked steps

it decreases down the group **B1**.

Exam-style 3.1 ■ 1 mark

Magnesium carbonate decomposes at a lower temperature than barium carbonate because the Mg^{2+} ion:

- A** is larger
- B** has a higher charge density and polarises the carbonate more
- C** has a lower charge
- D** is more soluble

Solution 3.1

Method: Thermal stability and polarisation

- A is larger
- B has a higher charge density and polarises the carbonate more 
- C has a lower charge
- D is more soluble

Worked steps

B. The small Mg^{2+} has a high charge density and polarises the carbonate more.

Exam-style 3.2 ■ 1 mark

The increasing thermal stability of the nitrates down Group 2 is explained by:

- A** increasing cation charge
- B** decreasing polarisation of the anion by the larger cation
- C** increasing anion size
- D** decreasing temperature

Solution 3.2

Method: Thermal stability and polarisation

- A increasing cation charge
- B decreasing polarisation of the anion by the larger cation 
- C increasing anion size
- D decreasing temperature

Worked steps

B. The larger cation polarises the anion less, so the nitrate is more stable.

Exam-style 3.3 ■ 3 marks

Explain, in terms of polarisation, why the thermal stability of the Group 2 nitrates increases down the group.

Solution 3.3

Method: Thermal stability and polarisation

Worked steps

down the group the cation gets larger, so its charge density falls (M1); it polarises/distorts the large nitrate ion less (M1); so the N–O bonds are weakened less and a higher temperature is needed to decompose it (A1).

Exam-style 3.4 ■ 2 marks

ΔH_{sol} depends on the lattice energy and the hydration enthalpies. Explain why both quantities become **less** exothermic down Group 2.

Solution 3.4

Method: Solubility energetics

Worked steps

down the group the cation is larger (M1); so both the lattice energy (ions further apart) and the hydration enthalpy (lower charge density, weaker attraction to water) become less exothermic (A1).