LATTICE ENTHALPY

WARNING

There can be two definitions - one is the opposite of the other! Make sure you know which one is being used.

Lattice Formation Enthalpy ($\Delta_{LE}H$)

OCR preferred

AQA can use either definition

Definition

The enthalpy change when ONE MOLE of an ionic crystal lattice is formed from its isolated gaseous ions.

Values

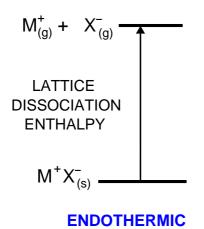
- highly exothermic strong electrostatic attraction between ions of opposite charge
- a lot of energy is released as the bond is formed
- relative values are governed by the charge density of the ions.

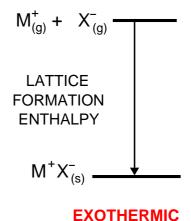
Example

$$Na^{+}(g)$$
 + $C\Gamma(g)$ ----> Na^{+} $C\Gamma(s)$

Notes

- one cannot measure this value directly; it is found using a Born-Haber cycle
- the greater the charge densities of the ions, the more they attract each other and the larger the lattice enthalpy.
- the more exothermic the lattice enthalpy, the higher the melting point





Lattice Dissociation Enthalpy ($\Delta_{LE}H$) Option for \boxed{AQA}

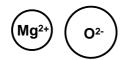
Definition The enthalpy change when ONE MOLE of an ionic lattice dissociates into isolated gaseous ions.

Values

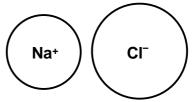
- **highly endothermic** there is a strong electrostatic attraction between ions of opposite charge
- a lot of energy must be put in to overcome the attraction

Example Na⁺ $C\Gamma(s)$ ----> Na⁺(g) + $C\Gamma(g)$

Consequences



HIGHER CHARGE DENSITY IONS GREATER ATTRACTION LARGER LATTICE ENTHALPY



LOWER CHARGE DENSITY IONS
LESS ATTRACTION
SMALLER LATTICE ENTHALPY

Q.1 Which substance in the the following pairs has the larger lattice enthalpy?

a) NaCl or KCl

b) NaF or NaCl

c) MgCl₂ or NaCl

d) MgO or MgCl₂

Thermal stability and Lattice Enthalpy

Oxides

• thermal stability of Group II oxides decreases down the group

	Mg ²⁺ O ²⁻	Ca ²⁺ O ²⁻	Sr ²⁺ O ²⁻	Ba ²⁺ O ²⁻
Lattice Enthalpy (kJ mol-1)	-3889	-3513	-3310	-3152
Melting Point (°C)	2853	decre	easing values	>

MgO

- magnesium oxide is used to line furnaces REFRACTORY LINING
- this is because of its high melting point (2853°C)
- the high melting point is a result of the large (highly exothermic) lattice enthalpy
- high lattice enthalpy due to the attraction between ions of high charge density

Carbonates

- thermal stability of Group II carbonates increases down the group
- MgCO₃ decomposes much easier than BaCO₃
- BUT the lattice enthalpy of MgCO₃ is HIGHER!

	$MgCO_3$	CaCO ₃	SrCO ₃	BaCO ₃
Decomposes at	350°C	832°C	1340°C	1450°C
Lattice Enthalpy (kJ mol-1)	-3123		>	-2556

- Mg²⁺ ions are SMALLER and have a HIGHER CHARGE DENSITY
- this makes them **MORE HIGHLY POLARISING**
- they **DISTORT THE CO₃²⁻** ion
- this WEAKENS THE ATTRACTION BETWEEN IONS
- the LATTICE IS NOT AS STRONG

Calculating Lattice Enthalpy

Introduction • you cannot measure lattice enthalpy directly

- values are found using a Born-Haber cycle
- Born-Haber cycles use Hess's Law
- The following enthalpy changes are part of a Born-Haber cycle.

Enthalpy Change of Formation ($\Delta_f H$)

Definition The enthalpy change when ONE MOLE of a compound is formed from its elements.

Standard Enthalpy Change of Formation ($\Delta_f H^{\circ}_{298}$)

Definition The enthalpy change when ONE MOLE of a compound is formed

in its standard state from its elements in their standard states.

Values Usually, but not exclusively, exothermic

Example(s) $Mg(s) + Cl_2(g)$ \longrightarrow $MgCl_2(s)$

 $Na(s) + \frac{1}{2}Cl_2(g) \longrightarrow NaCl(s)$

Notes
 Elements In their standard states have zero enthalpy of formation.

Q.2 Write equations representing the (standard) enthalpy changes of formation of

magnesium oxide

sodium oxide

potassium

potassium bromide

Enthalpy Change of Atomisation ($\Delta_{at}H$)

Definition The enthalpy change when ONE MOLE of gaseous atoms is formed from an element.

Standard Enthalpy Change of Atomisation ($\Delta_{\text{at}} H^{\circ}{}_{298}\!)$

Definition The enthalpy change when ONE MOLE of gaseous atoms is formed

in its standard state from an element in its standard state.

Values Always endothermic - you have to break the bonds holding the atoms together

Example(s) $\frac{1}{2}Cl_2(g)$ ---> Cl(g) and Na(s) ---> Na(g) (see note)

for solid elements, the change is known as ENTHALPY OF SUBLIMATION

Q.3 Write equations representing the (standard) enthalpy changes of atomisation of magnesium

oxygen

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First Ionisation Energy

Definition

The energy required to remove one mole of electrons (to infinity) from one mole of gaseous atoms to form one mole of gaseous positive ions.

Values

Always endothermic need to overcome the pull of the nucleus on the electron

Example(s)

$$Na(g)$$
 -----> $Na^+(g)$ + e^-

$$\mathbf{Na}^{+}(a) + \mathbf{e}$$

$$Mg(g)$$
 ----> $Mg^{+}(g)$ + e^{-}

Notes

- There is an ionisation energy for each successive electron removed.
 - e.g. SECOND IONISATION ENERGY

$$Mg^{+}(g)$$
 ----> $Mg^{2+}(g)$ + e^{-}

$$Mg^{2+}(g) + e$$

Look back in your notes to refresh your memory about the trends in I.E.'s

Q.4

Write equations representing the following ionisation energy changes;

1st IE of calcium

2nd IE of calcium

1st IE of lithium

1st IE of aluminium

Electron Affinity

Definition

The enthalpy change when ONE MOLE of gaseous atoms acquires ONE MOLE of electrons (from infinity) to form ONE MOLE of gaseous negative ions.

Values

Always exothermic - a favourable process due to the nucleus attracting the electron

Example

$$Cl(g) + e^- \longrightarrow Cl(g)$$

Notes

Do not confuse electron affinity with electronegativity.

0.5

Write equations representing the following electron affinity (EA) changes;

1st EA of bromine

1st EA of oxygen

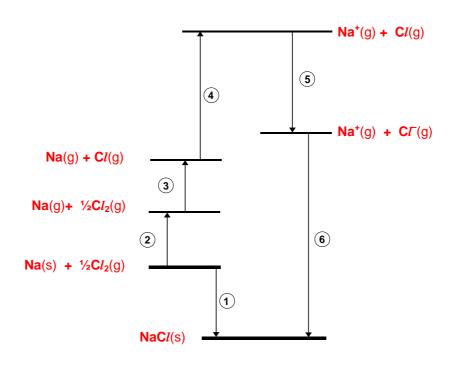
2nd EA of oxygen

BORN-HABER CYCLES

Theory

- involve the application of Hess's Law
- used to outline the thermodynamic changes during the formation of ionic salts
- used to calculate Lattice Enthalpy
- Lattice Enthalpy cannot be determined directly by experiment

BORN-HABER CYCLE FOR SODIUM CHLORIDE



STEPS (values are in kJ mol-1)

① Enthalpy change of formation of NaCl	$Na(s) + \frac{1}{2}Cl_2(g) \longrightarrow NaCl(s)$	-411
2 Enthalpy change of sublimation of sodium	Na(s) ——> Na(g)	+ 108
3 Enthalpy change of atomisation of chlorine	$^{1}/_{2}Cl_{2}(g)$ \longrightarrow $Cl(g)$	+ 121
4 Ist Ionisation Energy of sodium	$Na(g) \longrightarrow Na^+(g) + e^-$	+ 500
⑤ Electron Affinity of chlorine	$Cl(g) + e^{-} \longrightarrow Cl^{-}(g)$	- 364
6 Lattice Enthalpy of NaCl	$Na^{+}(g) + Cl^{-}(g) \longrightarrow NaCl(s)$	

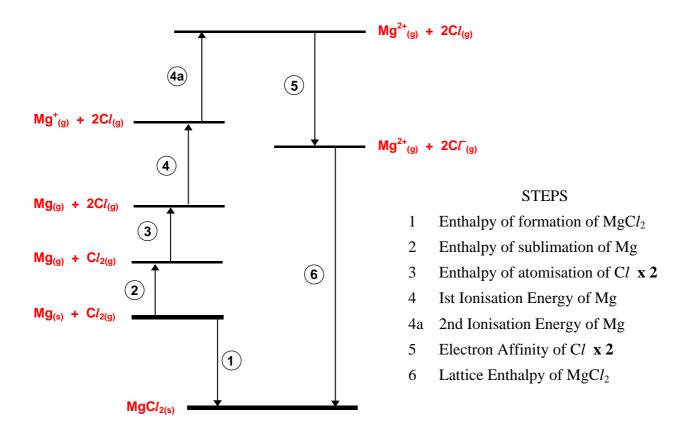
According to Hess's Law, the enthalpy change is independent of the path taken. Therefore...

STEP 6 = - (STEP 5) - (STEP 4) - (STEP 3) - (STEP 2) + (STEP 1)
-
$$(-364)$$
 - $(+500)$ - $(+121)$ - $(+108)$ + (-411) = -776 kJ mol⁻¹

Q.6 Construct a similar Born-Haber cycle for NaCl₂.

If the Lattice Enthalpy of NaCl₂ is -3360 kJ mol⁻¹, what is its enthalpy of formation? What does this tell you about the stability of NaCl₂?

BORN-HABER CYCLE FOR MAGNESIUM CHLORIDE



Q.7 If the Lattice Enthalpy of $MgCl_2$ is -2493 kJ mol⁻¹, what is its enthalpy of formation?

Will an ionic salt dissolve in water?

Introduction

If a pair of oppositely charged gaseous ions are placed together, they will attract each other. The energy change (LATTICE ENTHALPY) is highly exothermic.

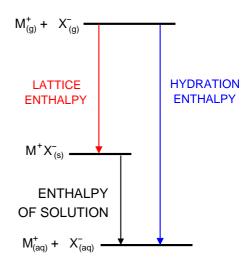
If the ions were put in water, they would be attracted to polar water molecules. the resulting energy change (HYDRATION ENTHALPY) is highly exothermic.

In both; the greater charge density of the ions = a more exothermic reaction

The missing stage of the cycle is known as the **ENTHALPY OF SOLUTION**.

The size and value of the enthalpy of solution depends on the **relative values** of the lattice enthalpy and the hydration enthalpy.

If HE >> LE then the salt will probably be soluble



Enthalpy Change of Hydration ($\Delta_{hyd}H$)

Definition

The enthalpy change when ONE MOLE of a gaseous ion dissolves in (an excess of) water (to give an infinitely dilute solution).

Values Exothermic

Example $Na^+(g)$ ----> $Na^+(ag)$

 $Cl^{-}(g)$ \longrightarrow $Cl^{-}(aq)$

Notes The polar nature of water stabilises the ions.

The greater the charge density of the ion, the greater the affinity for water and the more exothermic the process will be.

Enthalpy Change of Solution ($\Delta_{sol}H$)

Definition The enthalpy change when ONE MOLE of solute dissolves in (an excess of)

solvent (to give an infinitely dilute solution).

Values Mainly exothermic

Example NaCl(s) ——> NaCl(aq) [for ionic compounds, the ions will be dissociated]

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SOME USEFUL VALUES FOR THERMODYNAMIC CHANGES

Values, which may be slightly different in other books, are in kJ mol-1

Enthalpy	change
of format	ion

formation		Cl^-	Br^-	I^{-}	O^{2-}
	Na^+	-411	-381	-288	-414
	\mathbf{K}^{+}	-437	-398	-328	-361
	Mg^{2+}	-641	-524	-364	-602
	Ca^{2+}	-796	-683	-534	-635

Ionisation Energy

	1st I.E.	2nd I.E.
Na	+496	+4563
Mg	+738	+1451
Ca	+590	+1145
K	+419	+3051

2nd Electron Affinity

O +**798**

Lattice Enthalpy

Check which definition is being used and use appropriate sign for ΔH

	Cl^-	Br^-	F^{-}	O^{2-}
Na^{+}	-780	-742	-918	-2478
K^+	-711	-679	-817	-2232
Rb^+	-685	-656	-783	
Mg^{2+}	-2256			-3791
Ca^{2+}	-2259			

Hydration Enthalpy

 F^{-}

Cl

Br⁻

OH-

Ι-

-457

-384

-351

-307

-460