

Calorimetry 4

A Level

Essential Pre-Uni Chemistry F1.4

Specific heat capacity of water = $4.18 \text{ J g}^{-1} \text{ K}^{-1}$.

The enthalpy change of combustion of decane, $\text{C}_{10}\text{H}_{22}$, is $-6778 \text{ kJ mol}^{-1}$. Calculate the mass required to raise the temperature of 450 g of water by 80°C when burnt completely, with no heat losses from the water. Give your answer to 2 significant figures.

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Calorimetry 6

A Level

Essential Pre-Uni Chemistry F1.6

Specific heat capacity of water = $4.18 \text{ J g}^{-1} \text{ K}^{-1}$.

Calculate the enthalpy of combustion of propyne, C_3H_4 , given that complete combustion of 65 mg of propyne raises the temperature of 800 g of water from 20.15°C to 21.09°C .

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Calorimetry 8

A Level

Essential Pre-Uni Chemistry F1.8

Specific heat capacity of water = $4.18 \text{ J g}^{-1} \text{ K}^{-1}$

30.0 cm^3 of ethanoic acid at 1.60 mol dm^{-3} and 18.65°C is placed in an insulated polystyrene cup. When 40.0 cm^3 of sodium hydroxide at 1.00 mol dm^{-3} and 18.65°C is added, the temperature rises to 25.80°C .

Assuming that no heat is lost, that the specific heat capacity of water may be used, and that the solutions have a density of 1.00 g cm^{-3} at 18.65°C , find the enthalpy change of the reaction per mole of water produced by neutralisation.

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Calorimetry 9

A Level

Essential Pre-Uni Chemistry F1.9

Specific heat capacity of water = $4.18 \text{ J g}^{-1} \text{ K}^{-1}$.

When 5.0 g of ammonium nitrate dissolves in 100 g of water, the temperature of the water drops from 18°C to 14°C . Calculate the enthalpy of solution of ammonium nitrate in kJ mol^{-1} using the following scheme.

Part A Formula

Write down the formula of ammonium nitrate.

Part B Relative formula mass

Calculate the relative formula mass of ammonium nitrate. Give your answer to 3 significant figures.

Part C Number of moles

Calculate the number of moles of ammonium nitrate in 5.0 g.

Part D Heat loss of water

Calculate the heat lost from the 100 g of water. Give your answer to 2 significant figures.

Part E Molar heat loss of ammonium nitrate

Calculate the heat lost per mole of ammonium nitrate.

Part F Enthalpy of ammonium nitrate

Give the enthalpy of solution of ammonium nitrate.

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Bond Enthalpies 4

Essential Pre-Uni Chemistry F2.4

A Level

Given that the bond energy of H–H is 4.53 eV, D–D is 4.59 eV, and the energy change on reaction $\text{H}_2 + \text{D}_2 \longrightarrow 2 \text{HD}$ is +0.02 eV, find the bond energy of H–D. Give your answer to 3 significant figures.

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Formation and Combustion Enthalpies 3



Essential Pre-Uni Chemistry F3.3

Data (all in kJ mol^{-1}):

	$\Delta_f H^\ominus$		$\Delta_c H^\ominus$
$\text{CH}_4(\text{g})$	−74.8	$\text{C}_6\text{H}_6(\text{l})$	−3267.4
$\text{CCl}_4(\text{l})$	−129.6	$\text{H}_2(\text{g})$	−285.8
$\text{HCl}(\text{g})$	−92.3	$\text{C}_6\text{H}_{12}(\text{l})$	−3919.5
$\text{TiCl}_4(\text{l})$	−804.2	$\text{C}_2\text{H}_2(\text{g})$	−1300.8
$\text{TiCl}_3(\text{s})$	−720.9	$\text{C}_2\text{H}_6(\text{g})$	−1559.7
$\text{PCl}_3(\text{l})$	−319.7	$\text{C}_2\text{H}_5\text{OH}(\text{l})$	−1367.3
$\text{PCl}_5(\text{s})$	−443.5	$\text{C}_2\text{H}_4(\text{g})$	−1410.8
$\text{POCl}_3(\text{l})$	−597.1	$\text{CH}_3\text{COOH}(\text{l})$	−874.1
$\text{GeO}(\text{s})$	−212.1	$\text{C}_6\text{H}_{14}(\text{l})$	−4163.0
$\text{GeO}_2(\text{s})$	−551.0	$\text{CH}_3\text{COOC}_2\text{H}_5(\text{l})$	−2237.9
$\text{NH}_3(\text{g})$	−46.1	$\text{CO}(\text{g})$	−283.0
$\text{TiO}_2(\text{s})$	−939.7	$\text{Mg}(\text{s})$	−601.7

Use enthalpies of formation and combustion to calculate the reaction enthalpy for the reaction:

$\text{Ge}(\text{s}) + 2 \text{H}_2\text{O}(\text{l}) \longrightarrow \text{GeO}_2(\text{s}) + 2 \text{H}_2(\text{g})$ Give your answer to 3 significant figures.

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Formation and Combustion Enthalpies 4



Essential Pre-Uni Chemistry F3.4

Data (all in kJ mol^{-1}):

	$\Delta_f H^\ominus$		$\Delta_c H^\ominus$
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$\text{NH}_3(\text{g})$	-46.1	$\text{CO}(\text{g})$	-283.0
$\text{TiO}_2(\text{s})$	-939.7	$\text{Mg}(\text{s})$	-601.7

Use the reaction enthalpies given, and the combustion or formation enthalpies above to find the requested enthalpy change in each case:

Part A $\text{NH}_4\text{Cl}(\text{s})$

$\text{NH}_3(\text{g}) + \text{HCl}(\text{g}) \longrightarrow \text{NH}_4\text{Cl}(\text{s})$, $\Delta_r H^\ominus = -176 \text{ kJ mol}^{-1}$ find $\Delta_f H^\ominus$ of $\text{NH}_4\text{Cl}(\text{s})$

Part B $\text{MgCl}_2(\text{s})$

$\text{TiCl}_4(\text{l}) + 2 \text{Mg}(\text{s}) \longrightarrow 2 \text{MgCl}_2(\text{s}) + \text{Ti}(\text{s})$ $\Delta_r H^\ominus = -478.4 \text{ kJ mol}^{-1}$, find $\Delta_f H^\ominus$ of $\text{MgCl}_2(\text{s})$

Part C $\text{CH}_3\text{COOCOCH}_3(\text{l})$

$\text{CH}_3\text{COOCOCH}_3(\text{l}) + \text{H}_2\text{O}(\text{l}) \longrightarrow 2 \text{CH}_3\text{COOH}(\text{l})$ $\Delta_r H^\ominus = -46 \text{ kJ mol}^{-1}$, find $\Delta_c H^\ominus$ of $\text{CH}_3\text{COOCOCH}_3(\text{l})$ Give your answer to 4 significant figures.

Part D $\text{C}_6\text{H}_5\text{CHCH}_2$

$4 \text{C}_2\text{H}_2(\text{g}) \longrightarrow \text{C}_6\text{H}_5\text{CHCH}_2(\text{l})$, $\Delta_r H^\ominus = -808.2 \text{ kJ mol}^{-1}$, find $\Delta_c H^\ominus$ of $\text{C}_6\text{H}_5\text{CHCH}_2$ Give your answer to 4 significant figures.

Part E $\text{Al}_2\text{O}_3(\text{s})$

$4 \text{Al}(\text{s}) + 3 \text{GeO}_2(\text{s}) \longrightarrow 2 \text{Al}_2\text{O}_3(\text{s}) + 3 \text{Ge}(\text{s})$ $\Delta_r H^\ominus = -1698.4 \text{ kJ mol}^{-1}$, find $\Delta_f H^\ominus$ of $\text{Al}_2\text{O}_3(\text{s})$ Give your answer to 4 significant figures.

Part F Fe_2O_3

$\text{Fe}_2\text{O}_3(\text{s}) + 3 \text{CO}(\text{g}) \longrightarrow 2 \text{Fe}(\text{s}) + 3 \text{CO}_2(\text{g})$, $\Delta_r H^\ominus = -24.8 \text{ kJ mol}^{-1}$, find $\Delta_f H^\ominus$ of Fe_2O_3

Part G CuO (s)

$3 \text{ CuO (s)} + 2 \text{ NH}_3 \text{ (g)} \longrightarrow 3 \text{ Cu (s)} + \text{ N}_2 \text{ (g)} + 3 \text{ H}_2\text{O (l)}$, $\Delta_r H^\ominus = -293.3 \text{ kJ mol}^{-1}$, find $\Delta_f H^\ominus$ of CuO (s) Give your answer to 3 significant figures.

Part H H₃PO₄ (s)

$2 \text{ PCl}_5 \text{ (s)} + 8 \text{ H}_2\text{O (l)} \longrightarrow 2 \text{ H}_3\text{PO}_4 \text{ (s)} + 10 \text{ HCl (g)}$, $\Delta_r H^\ominus = -307.6 \text{ kJ mol}^{-1}$, find $\Delta_f H^\ominus$ of H₃PO₄ (s) Give your answer to 3 significant figures.

Part I Ga

$\text{Ga}_2\text{O}_3 \text{ (s)} + 3 \text{ Mg (s)} \longrightarrow 2 \text{ Ga (s)} + 3 \text{ MgO (s)}$, $\Delta_r H^\ominus = -716.1 \text{ kJ mol}^{-1}$, find $\Delta_c H^\ominus$ of Ga.

Part J HCl(g)

$\text{TiCl}_4 \text{ (l)} + 2 \text{ H}_2\text{O (l)} \longrightarrow \text{TiO}_2 \text{ (s)} + 4 \text{ HCl (aq)}$, $\Delta_r H^\ominus = -232.3 \text{ kJ mol}^{-1}$, find $\Delta_{\text{sol}} H^\ominus$ of HCl(g) Give your answer to 3 significant figures.

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C₃H₆ Combustion

A Level



A and **B** are two isomers with the molecular formula C₃H₆. The standard enthalpies of formation, $\Delta_f H^\ominus$, of both **A** and **B** have been found by first measuring the standard enthalpies of combustion, $\Delta_c H^\ominus$, of each. These values are given in the table below, together with the standard enthalpies of combustion of carbon and hydrogen.

	A	B	carbon	hydrogen
$\Delta_c H^\ominus / \text{kJ mol}^{-1}$	−2058	−2091	−393.5	−241.8

Part A Combustion equation

Give the equation for the complete combustion of C₃H₆. (Balance it for one mole of the hydrocarbon.)

Part B $\Delta_f H^\ominus$ of A

Calculate the standard enthalpy of formation of **A**.

Part C $\Delta_f H^\ominus$ of B

Calculate the standard enthalpy of formation of **B**.

Part D Isomerisation

Gaseous **B** needs to be stored carefully since it can convert explosively to the elements, to isomer **A**, or to other hydrocarbons. Calculate the standard enthalpy change for the reaction **B** \longrightarrow **A**.

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Homologous Series: Combustion

A Level

A student was studying the enthalpies of combustion of different alkanes, and thinking about the pattern seen when descending the homologous series.

Part A General equation

The student started by writing a balanced chemical equation for the combustion of a general alkane C_nH_{2n+2} , by balancing for one mole of the alkane:



Find the correct expressions for x , y and z as a function of n .

x :

The following symbols may be useful: n

y :

The following symbols may be useful: n

z :

The following symbols may be useful: n

Part B Bonds broken

Next, the student was thinking about all the bonds present on the left-hand side of the equation previously balanced, which need to be broken in order to form the products. How many of the different types of bond are present, per molecule of C_nH_{2n+2} , assuming a correctly balanced equation?

Number of C—C bonds:

The following symbols may be useful: n

Number of C—H bonds:

The following symbols may be useful: n

Number of O=O bonds:

Part C Bonds formed

The following consideration concerned all the bonds present on the right-hand side of the equation previously balanced, i.e. the bonds formed during the reaction. How many of the different types of bond are present, per molecule of C_nH_{2n+2} , assuming a correctly balanced equation?

Number of C=O bonds present:

The following symbols may be useful: n

Number of O—H bonds present:

Part D **Bond enthalpy assumption**

When using average bond enthalpies to estimate enthalpies of reaction, we must assume that the average values used are representative of the bonding present (we can use different average values, e.g. specific to hydrocarbons to improve estimates). What other assumption about the chemicals present must be met for a calculation using bond enthalpies alone to give a good estimate of the enthalpy of reaction?

Part E **Bond enthalpy calculation**

Using the following bond enthalpies, find an expression for the enthalpy change (in kJ mol^{-1}) of the reaction previously written out (corresponding to the complete combustion of $\text{C}_n\text{H}_{2n+2}$) assuming it is carried out under conditions so that the assumption from the previous part is met. Do not include the units in the expression, and quote numerical values in the expression to 3 s.f.

Bond	Bond enthalpy / kJ mol^{-1}
C–C	348
C–H	412
O=O	498
C=O in CO_2	805
O–H	463

The following symbols may be useful: n

Part F Deviation at RTP

If the combustion is instead carried out at RTP (room temperature and pressure), how would you expect the empirically obtained enthalpy change of combustion of ethane to compare to that calculated using the formula derived above?

The RTP enthalpy change will be exothermic. This is because is now present as a and energy is during the as bonds are formed.

Items:

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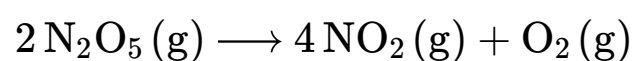
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Dinitrogen Pentoxide Formation

A Level



In the gas phase, N_2O_5 decomposes to oxygen and nitrogen dioxide. Use the data below (determined at 298 K) to calculate the standard enthalpy change at 298 K for the reaction:



	value/ kJ mol^{-1}
$\Delta_f H^\ominus$ of $\text{N}_2\text{O}_5(\text{g})$	11.3
$\Delta_r H^\ominus$ for $\text{NO}(\text{g}) + \frac{1}{2}\text{O}_2(\text{g}) \longrightarrow \text{NO}_2(\text{g})$	−58.1
bond strength in $\text{N}_2(\text{g})$	945
bond strength in $\text{O}_2(\text{g})$	498
bond strength in $\text{NO}(\text{g})$	631

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