	–890 kJ mol ^{–1} .				
	•••••				
					(4) (Total 4 marks)
	•	₆ H ₁₁ OH, is comple	etely burned in oxyge	en, the products are	e carbon dioxide
	water.	annacont the meastic	n that a cours during	the measurement of	f the outhelms
	e an equation to re		n that occurs during	the measurement of	
					(Total 1 mark)
Γhe :	table below includ	les some values of	standard enthalpies o	of formation ($\Delta H_{\mathbf{f}}^{\mathbf{Q}}$).
Γhe 1).
	Substance	H ₂ O(l)	LiOH(s)	Li(s)).
).
Δ	Substance AHF /kJ mol ⁻¹	H ₂ O(1) -286	LiOH(s)	Li(s)).
Δ	Substance AHF /kJ mol ⁻¹	H ₂ O(l) -286 of solution of lithiu	LiOH(s)	Li(s) 0 n below.	
Δ	Substance $AH_{\mathbf{f}}^{\mathbf{G}}/kJ \text{ mol}^{-1}$ standard enthalpy $LiOH(s) \rightarrow Li^{+}(s)$	$H_2O(1)$ -286 of solution of lithin aq) + OH ⁻ (aq)	LiOH(s) -487 um hydroxide is give	Li(s) 0 n below.	
The:	Substance $AH_{\mathbf{f}}^{\mathbf{G}}/kJ \text{ mol}^{-1}$ standard enthalpy $LiOH(s) \rightarrow Li^{+}(s)$	$H_2O(1)$ -286 of solution of lithin aq) + OH ⁻ (aq)	LiOH(s) -487 $\Delta H^{\bullet} = 21 \text{ kJ mos}$	Li(s) 0 n below.	
<u>/</u> The :	Substance $AH_{\mathbf{f}}^{\bullet}/kJ \text{ mol}^{-1}$ standard enthalpy $LiOH(s) \rightarrow Li^{+}(s)$ State why the standard enthalpy	H ₂ O(1) -286 of solution of lithin aq) + OH ⁻ (aq) andard enthalpy of	LiOH(s) -487 am hydroxide is give $\Delta H^{\bullet} = 21 \text{ kJ mo}$ formation of lithium	Li(s) 0 n below. ol ⁻¹ is quoted as zero.	(1)
<u>/</u> The :	Substance $H_{\mathbf{f}}^{\bullet}$ /kJ mol ⁻¹ standard enthalpy LiOH(s) \rightarrow Li ⁺ (s State why the sta	H ₂ O(1) -286 of solution of lithin aq) + OH ⁻ (aq) and and enthalpy of on for the chemical	LiOH(s) -487 $\Delta H^{\bullet} = 21 \text{ kJ mos}$	Li(s) 0 n below. ol ⁻¹ is quoted as zero. sents the formation	
Δ The	Substance LHF /kJ mol ⁻¹ standard enthalpy LiOH(s) → Li ⁺ (s State why the standard enthalpy Write an equation hydroxide from it of formation.	H ₂ O(l) -286 of solution of lithin aq) + OH ⁻ (aq) andard enthalpy of the chemical its elements, in which	LiOH(s) -487 am hydroxide is give $\Delta H^{\bullet} = 21 \text{ kJ mo}$ formation of lithium reaction which repre	Li(s) 0 n below. ol ⁻¹ is quoted as zero. sents the formatio ge is equal to its st	n of lithium andard enthalpy

	(d)	Use the data given above to calcula lithium with water.	ate a value	e for the	enthalpy	change	for the i	reaction of	
									(3)
								(Total 8 m	arks)
4.	(a)	Explain the meaning of the terms n	nean bond	enthalp	y and <i>sta</i>	ındard e	nthalpy (of formation.	
		Mean bond enthalpy				•••••			
								•••••	
		Standard enthalpy of formation							
				•••••		••••••			
			••••••	•••••		••••••	•••••		
			•••••	•••••		•••••	•••••		(5)
	(b)	Some mean bond enthalpies are given	ven below	'.					
		Bond	N–H	N-N	N≡N	Н–О	O–O		
		Mean bond enthalpy/kJ mol ⁻¹	388	163	944	463	146		
		Use these data to calculate the enth between hydrazine, N ₂ H ₄ , and hydrazine				ing gas-	phase re	eaction	
		H N-N H + 2 H-O-	-О—Н		$N \equiv N$	+ 4 H	— 0—	Н	
								•••••	
								•••••	

(c) Some standard enthalpies of formation are given below.

	$N_2H_4(g)$	$H_2O_2(g)$	H ₂ O(g)
$\Delta H_{\rm f}^{ m{\Theta}} / { m kJ \ mol}^{-1}$	+75	-133	-242

These data can be used to calculate the enthalpy change for the reaction in part (b).

$$N_2H_4(g) + 2H_2O_2(g) \rightarrow N_2(g) + 4H_2O(g)$$

	(i)	State the value of $\Delta H_{\rm f}^{\theta}$ for N ₂ (g).	
	(ii)	Use the $\Delta H_{\rm f}^{ m e}$ values from the table to calculate the enthalpy change for this reaction.	
			(4)
(d)	Expl	ain why the value obtained in part (b) is different from that obtained in part (c)(ii).	
	•••••	(Total 13 ma	(1) arks)

- **5.** Methanol, CH₃OH, is a convenient liquid fuel.
 - (a) An experiment was conducted to determine the enthalpy of combustion of liquid methanol. The energy obtained from burning 2.12 g of methanol was used to heat 150 g of water. The temperature of the water rose from 298 K to 362 K. (The specific heat capacity of water is $4.18 \text{ J K}^{-1} \text{ g}^{-1}$)
 - (i) Define the term *standard enthalpy of combustion*.
 - (ii) Use the data above to calculate a value for the enthalpy of combustion of one mole of liquid methanol.

(b) Methanol can be synthesised from methane and steam by a process that occurs in two stages.

Stage 1
$$CH_4(g) + H_2O(g) \Longrightarrow 3H_2(g) + CO(g)$$
 is $\Delta H^{\oplus} = +206 \text{ kJ mol}^{-1}$
Stage 2 $CO(g) + 2H_2(g) \Longrightarrow CH_3OH(g)$ $\Delta H^{\oplus} = -91 \text{ kJ mol}^{-1}$

The standard enthalpies of combustion of carbon monoxide and of hydrogen are -283 kJ mol^{-1} and -286 kJ mol^{-1} , respectively. Use these data and the enthalpy change for *Stage 2* to calculate a value for the standard enthalpy of combustion of gaseous methanol.

(3) (Total 10 marks)

6. (a) Define the term *standard enthalpy of formation*.

(3)

(b) State Hess's Law and use it, together with the data given in the table below, to calculate the standard enthalpy change for the following reaction.

$$MgO(s) + 2HCl(g) \rightarrow MgCl_2(s) + H_2O(l)$$

	MgO	HCl(g)	MgCl ₂	H ₂ O
$\Delta H_{\rm f}^{\bullet}/{\rm kJ~mol}^{-1}$	-602	-92	-642	-286

(4)

(c) In an experiment, an excess of solid magnesium oxide was added to 50 cm^3 of 3.0 mol dm^{-3} hydrochloric acid. The initial temperature of the solution was 21 °C. After reaction, the temperature had risen to 53 °C. (The specific heat capacity of water is $4.2 \text{ J K}^{-1} \text{ g}^{-1}$)

Use this information to calculate the enthalpy change for the reaction of one mole of magnesium oxide with hydrochloric acid. For your calculation you should assume that all the heat from the reaction is used to raise the temperature of 50~g of water.

(8)

(Total 15 marks)

7. The table below contains some mean bond enthalpy data.

Bond	Н—Н	С—С	C=C	N≡N	N—H
Mean bond enthalpy / kJ mol ⁻¹	436	348	612	944	388

(a)	Expla	in the term mean bond enthalpy.	
			(2)
(b)	(i)	Write an equation for the formation of one mole of ammonia, NH ₃ , from its elements.	
	(ii)	Use data from the table above to calculate a value for the enthalpy of formation of ammonia.	
			(4)

(c) Use the following equation and data from the table above to calculate a value for the C–H bond enthalpy in ethane.

(3) (Total 9 marks)

8.	(a)	Write an equation for the complete combustion of propanone, C ₃ H ₆ O, to form carbon dioxide and water.				
		•••••				
	(b)	heat	laboratory experiment, 1.45 g of propanone were burned completely in oxygen. The from this combustion was used to raise the temperature of 100 g of water from 1 K to 351.2 K.			
		(i)	Calculate the number of moles of propanone in the 1.45 g.			
		(ii)	Calculate the heat energy required to raise the temperature of 100 g of water from			
			293.1 K to 351.2 K. (The specific heat capacity of water is $4.18 \text{ J K}^{-1} \text{ g}^{-1}$)			
		(iii)	Hence, calculate a value, in kJ mol ⁻¹ , for the enthalpy of combustion of propanone.			
	(c)		similar experiment, the enthalpy of combustion of butanone, C_4H_8O , was found to be 0 kJ mol^{-1} . A data book value for the same reaction is $\Delta H_c^{\bullet} = -2430 \text{ kJ mol}^{-1}$.			
		(i)	Suggest one reason why the experimental value is very different from the data book value.			
		(ii)	This data book value of ΔH_c^{\bullet} for butanone (-2430 kJ mol ⁻¹) refers to the formation of carbon dioxide gas and water in the gaseous state. How would this value differ if it referred to the formation of water in the liquid state? Explain your answer.			
			Difference			
			Explanation			

(d) Calculate a value for the standard enthalpy of formation for liquid ethanethiol, C_2H_5SH . Use the equation given below and enthalpy of combustion data from the following table.

Substance	C ₂ H ₅ SH(l)	C(s)	$H_2(g)$	S(s)
$\Delta H_{\rm c}^{-\Phi}$ / kJ mol ⁻¹	-1170	-394	-286	-297

	$2C(s) + 3H2(g) + S(s) \rightarrow C2H5SH(l)$		
(3)			
al 12 marks)	(Total		
	Define the term standard enthalpy of formation.	(a)	9.
(3)			
ige is	Write an equation, including state symbols, for a reaction for which the enthalpy chang the standard enthalpy of formation of liquid CH ₃ NO ₂	(b)	
(2)			
	Give the name of the principle or law which enables enthalpies of formation to be calculated from enthalpies of combustion.	(c)	
(1)			

(d) In the presence of a catalyst, gaseous hydrogen cyanide, HCN, burns in an excess of oxygen as shown by the equation below.

$$HCN(g) + 2\frac{1}{4}O_2(g) \rightarrow \frac{1}{2}H_2O(g) + CO_2(g) + NO_2(g)$$

Some standard enthalpies of combustion, $\Delta H_{\mathbf{c}}^{\mathbf{\Theta}}$ are given in the table below.

Substance	HCN(g)	$H_2(g)$	C(s)	N ₂ (g)
ΔH c /kJ mol ⁻¹	-611	-242	-394	+68

		Use these data to calculate a value for the standard enthalpy of formation for gas hydrogen cyanide.	eous
			(4) (Total 10 marks)
10.	(a)	State what is meant by the term <i>mean bond enthalpy</i> .	
			(2)

(b) Ethanal has the structure
$$H = C - C$$
 $H = H$ $H = H$

Gaseous ethanal burns as shown by the equation

$$CH_{3}CHO(g) \ + \ 21/_{2}O_{2}(g) \ \to \ 2H_{2}O(g) \ + \ 2CO_{2}(g)$$

Use the mean bond enthalpy data given below to answer the following questions.

Bond	Mean bond enthalpy/kJ mol ⁻¹
С—Н	+413
С—С	+347
C==O	+736
O==O	+498
О—Н	+464

(i)	Calculate the enthalpy change which occurs when all the bonds in the reactants shown in the above equation are broken.
(ii)	Calculate the enthalpy change which occurs when all the bonds in the products shown in the above equation are formed.
(iii)	Hence, calculate the enthalpy change for the complete combustion of ethanal as shown in the equation above.
	(5)
	(Total 7 marks)

11.	(a)	Define	e the term <i>standard m</i>	nolar enthalpy oj	formation, ΔH	. ⊕ f ·	
							••
							(3)
	(b)	State 1	Hess's law.				
							(1)
	(c)	Propa	none, CH ₃ COCH ₃ , bu	ırns in oxygen a	s shown by the	equation	()
		_	CH ₃ COCH ₃ (l)			_	
		Use th	ne data given below to				propanone.
				CO ₂ (g)	H ₂ O(l)	CH ₃ COCH ₃ (l)	•
			$\Delta H_{\rm f}^{\bullet}/{\rm kJ~mol}^{-1}$	-394	-286	-248	
			$\Delta H_{\rm f}$ /KJ mol	-374	-200	-240	
							(3) (Total 7 marks)

React	tion 1	$\frac{1}{2} N_2(g) + \frac{1}{2} O_2(g)$	=	NO(g)	$\Delta H_{\mathbf{f}}^{\mathbf{\Theta}} = +91 \text{ kJ mol}^{-1}$	
React	tion 2	$\tfrac{1}{2}N_2(g) + 1\tfrac{1}{2}H_2(g)$	\(\neq\)	NH ₃ (g)	$\Delta H_{\mathbf{f}}^{\mathbf{G}} = -46 \text{ kJ mol}^{-1}$	
React	tion 3	$H_2(g) + \frac{1}{2} O_2(g)$	=	$H_2O(g)$	$\Delta H_{\mathbf{f}}^{\mathbf{G}} = -242 \text{ kj mol}^{-1}$	
Use th	his info	ormation in answering the	questi	ons that follow.		
(a)	What	name is given to the symb	ol ΔH	f ? Explain fully	what it means.	
	Name	·				
	Expla	nation				
						(4)
(b)	(i)	ammonia and steam.		· ·	noxide, NO, with hydrogen to form	(4)
	(ii)	Calculate the value of ΔH Question 3.	oo for	this reaction, usin	ng the data given at the beginning of	
			•••••	•••••		
			•••••	•••••		
			•••••	•••••		
			••••••		(Total 8 ma	(4) arks)

Nitrogen, hydrogen and oxygen undergo the reactions shown below.

12.

13.	poly	A student added 50.0 cm ³ of hydrochloric acid to 50.0 cm ³ of sodium hydroxide solution in a polystyrene cup. The temperature rose by 6.5 °C. The initial concentration of each solution was 1.00 mol dm ⁻³ .					
	(a)	Write an ionic equation for the reaction occurring.					
			(1)				
	(b)	Calculate the number of moles of acid used in the reaction.					
			(1)				
	(c)	Calculate the heat energy evolved in the reaction. (Assume that the final solution has a specific heat capacity of $4.18~J~g^{-1}~K^{-1}$ and a density of $1.00~g~cm^3$.)					
			(2)				
	(d)	Calculate the molar enthalpy change for the reaction.					
		(Total	(2) l 6 marks)				
14.	(a)	Define the term standard enthalpy of combustion.					
	(b)	Using the data given below, calculate the standard enthalpy change for the following reaction.	(3)				
		$CH_4(g) + 2O_2(g) \rightarrow CO_2(g) + 2H_2O(1)$					
		$\Delta H_{\rm f} ^{\bullet} {\rm CO}_2(g) = -394 \text{ kJ mol}^{-1}$					
		$\Delta H_{\rm f}^{\bullet}$ $H_2{\rm O}(1) = -286~{\rm kJ~mol}^{-1}$					
		$\Delta H_{\rm f}^{\bullet}$ CH ₄ (g) = -75 kJ mol ⁻¹	(3)				

(c)	(i)	State what is meant by the term <i>mean bond enthalpy</i> .
	(ii)	Using the standard enthalpy of formation of methane given in part (b) and the data given below, calculate the mean bond enthalpy of the C-H bond in methane.
		$C(s) \rightarrow C(g)$ $\Delta H^{\bullet} = +715 \text{ kJ mol}^{-1}$
		$C(s) \rightarrow C(g)$ $\Delta H^{\bullet} = +715 \text{ kJ mol}^{-1}$ $H_2(g) \rightarrow 2H(g)$ $\Delta H^{\bullet} = +436 \text{ kJ mol}^{-1}$
	(iii)	Using the C–H bond enthalpy calculated in part (c)(ii) and the standard enthalpy change for the reaction given below, calculate the mean bond enthalpy of the C–C bond in propane.
		N.B. If you failed to complete part (c)(ii), you may assume that the mean bond enthalpy of the C–H bond is +390 kJ mol ⁻¹ . (This is not the correct value.)
		$H \longrightarrow H \longrightarrow H$ $H \longrightarrow C \longrightarrow C \longrightarrow C \longrightarrow H(g) \longrightarrow 3C(g) + 8H(g) \qquad \Delta H \stackrel{\Theta}{=} +4020 \text{ kJ mol}^{-1}$ $H \longrightarrow H \longrightarrow H$
		(Total 13 mar

(b)	Methanethiol, CH ₃ SH, is added to natural gas in order to give it a smell. When methanethiol burns in air it forms carbon dioxide, water and sulphur dioxide. Write a balanced equation for this combustion reaction.						
(c)		bustion. In this calc	ulation, which three e	etermined using Hess's Lenthalpy values would b			
(d)	Carbon disulphide, Carbon disulp	S_2 , burns in air as fol	lows:				
	C	$S_2(1) + 3O_2(g) \rightarrow CC$	$O_2(g) + 2SO_2(g)$				
	Some standard enthal	pies of formation, Δl	$H^{\bullet}_{\mathbf{f}}$ are given in the ta	able below.			
	Substance	CS ₂ (l)	CO ₂ (g)	SO ₂ (g)			
	$\Delta H_{\mathbf{f}}^{\mathbf{\Theta}}/\mathrm{kJ} \; \mathrm{mol}^{-1}$	+88	-394	-297			
	Use these data to calc disulphide.	ulate a value for the	enthalpy of combusti	on of liquid carbon			
				• • • • • • • • • • • • • • • • • • • •			

16. The tables below contain values of standard enthalpy of combustion, $\Delta H_{\mathbf{c}}^{\mathbf{c}}$ and standard enthalpy of formation, $\Delta H_{\mathbf{f}}^{\mathbf{c}}$.

	C(graphite)	S(s)
$\Delta H_{\rm c}^{\rm t}$ /kJ mol ⁻¹	-394	-297

	CS ₂ (1)	NO(g)	CO(g)	$SO_2(g)$
$\Delta H_{\rm f}^{\rm t}/{\rm kJ~mol^{-1}}$	88	90	-111	-297

(a)	State	Hess's Law.	
			(2)
(b)	Expla	ain the meaning of the term standard enthalpy of combustion of a compound.	
			(3)
(c)	(i)	Write equations, with state symbols, for the formation of liquid carbon disulphide (CS ₂) from its elements and for the combustion of carbon disulphide to form carbon dioxide and sulphur dioxide.	
		Equation for formation of CS ₂	
		Equation for combustion of CS ₂	
	(ii)	Calculate the enthalpy of combustion of carbon disulphide, using its enthalpy of formation and the enthalpies of combustion from the tables on above.	

$CS_2(1) + 5NO(g) \to CO(g) + 2SO_2(g) + \frac{5}{2}N_2(g)$
Use the enthalpies of formation given in the table to calculate the enthalpy change in this reaction.
(3) (Total 13 marks)

Carbon disulphide reacts explosively with nitrogen monoxide (NO) according to the following equation.

(d)

The	structu	re of hydrazine can be represented as shown below.	
		$H \longrightarrow N \longrightarrow N$	
		Н	
(a)	(i)	Write an equation for the formation of hydrazine from its elements.	
	(ii)	Steam and nitrogen gas are the combustion products of gaseous hydrazine. Write an equation for this reaction.	
	(iii)	Use the data given below to calculate the standard enthalpy of formation of gaseous hydrazine.	
		$\Delta H_{\mathbf{c}}^{\mathbf{\Theta}} N_2 H_4(g) = -568 \text{ kJ mol}^{-1}$	
		$\Delta H_{\mathbf{f}}^{\mathbf{G}} \operatorname{H}_2\operatorname{O}(g) = -242 \text{ kJ mol}^{-1}$	
			(4)
(b)	State	what is meant by the term mean bond enthalpy.	

(2)

17.

(c)	Use the enthalpy of formation of hydrazine calculated in part (a) and the mean bond
	enthalpies given below to calculate the enthalpy of the bond between the two nitrogen
	atoms in hydrazine.

NB If you failed to complete part (a) you may assume that $\Delta H_{\textbf{f}}^{\textbf{p}} \ N_2 H_4(g) = +50 \ kJ \ mol^{-1} \ (\text{not the correct value}).$

Bond	N–H	N≡N	Н–Н		
Mean bond enthalpy/kJ mol ⁻¹	389	945	436		
		••••••	•••••	••••••	(3)
The measured N-N bond enthalp enthalpy when calculated correct					
				(Tot	(1) tal 10 marks)

18. The table below gives some *standard enthalpies of formation*, $\Delta H_{\mathbf{f}}^{\mathbf{S}}$.

Substance	$C_2H_6(g)$	$F_2(g)$	CF ₄ (g)	HF(g)
$\Delta H_{\mathbf{f}}^{\mathbf{\Theta}}/\mathrm{kJ} \; \mathrm{mol}^{-1}$	-85	0	-680	-269

	ΔH† /KJ IIIOI	-03	O	-000	-207	
(a)	Define the term sta	ndard enthal _i	py of formati	on.		
		•••••			•••••	 (3)

(b)	Why is the value for the standard enthalpy of formation for fluorine equal to zero?				
(c)	Calc	ulate the standard enth	nalny change for the	following reaction	
(0)	Care		$2(g) \rightarrow 2CF_4(g) + 6$	· ·	
(c) (d)	The	table below shows sor	me bond enthalpies.		
		Bond	C–C	С–Н	F–F
I	Bond en	thalpy /kJ mol–1	348	412	158
	(ii)	Suggest the most lik for your answer.	ely first step in the	reaction shown in part ((c) and give a reason
		-			
		Reason			(Total 11 m
(a)				on and standard enthal	
(a)			ation	on and standard enthal	
(a)	Stan	dard enthalpy of form	ation		
(a)	Stan	dard enthalpy of form	ation		

(b) Use the standard enthalpies of formation, $\Delta H_{\mathbf{f}}^{\mathbf{e}}$, given below

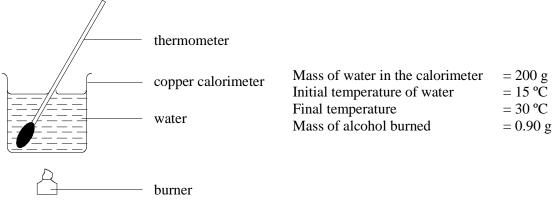
Compound	$\Delta H_{\mathbf{f}}^{\mathbf{e}} / \text{kJ mol}^{-1}$
CO ₂ (g)	-394
C ₃ H ₇ OH(l)	-304
H ₂ O(l)	-286

to calculate the standard enthalpy of combustion of an alcohol $\mathrm{C_3H_7OH}$, as shown by the equation:

<i>y</i> ,	4½O ₂ (g) → 3CC		

(3)

(c) A value for the enthalpy of combustion of the alcohol C_3H_7OH was determined in the laboratory using the apparatus shown below. The following results were obtained.



(i)	Calculate the heat energy required to raise the temperature of the water from 15 °C to 30 °C. The specific heat capacity of water is 4.2 J g^{-1} K ⁻¹ .
(ii)	Calculate the number of moles of the alcohol, C ₃ H ₇ OH, burned.
(iii)	Hence, calculate a value for the enthalpy of combustion of 1.0 mol of the alcohol.
(iv)	Give two reasons why you would expect your answer to part (c)(iii) to differ from that in part (b).
	Reason 1
	Reason 2

20.	An e	quation for the combustion of cyclopropane, C ₃ H ₆ , is shown below.	
		$C_3H_6(g) + 4\frac{1}{2}O_2(g) \rightarrow 3CO_2(g) + 3H_2O(g)$	
		standard enthalpy of combustion of cyclopropane can be calculated either from standard alpies of formation or by using mean bond enthalpies.	
	(a)	Use the standard enthalpies of formation given below to calculate the standard enthalpy of combustion of cyclopropane.	
		$\Delta H_{\mathbf{f}}^{\mathbf{G}} C_3 H_6(g) = +53 \text{ kJ mol}^{-1}$	
		$\Delta H_{\mathbf{f}}^{\mathbf{G}} \operatorname{CO}_{2}(g) = -393 \text{ kJ mol}^{-1}$	
		$\Delta H_{\mathbf{f}}^{\mathbf{G}} H_2 O(g) = -242 \text{ kJ mol}^{-1}$	
			(3)
	(b)	State what is meant by the term <i>mean bond enthalpy</i> .	
			(2)
	(c)	The following mean bond enthalpies have been obtained from a data book. Use these values to calculate the standard enthalpy of combustion of cyclopropane.	
		C–C 347 kJ mol ⁻¹ C–H 413 kJ mol ⁻¹	
		C=O 805 kJ mol ⁻¹ O=O 498 kJ mol ⁻¹	
		O–O 498 KJ IIIOI O–H 464 kJ mol ⁻¹	

(d)	Explain, by reference to the structure of cyclopropane, why the enthalpy of combustic calculated in part (a) is more exothermic than that calculated in part (c).	on
		(2)
	(Tota	l 10 marks)