<b>Q1.</b> (a)	Define the term standard enthalpy of formation, $\Delta H_r^{\circ}$

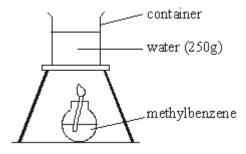
(b) Use the data in the table to calculate the standard enthalpy of formation of liquid methylbenzene,  $C_7H_8$ 

Substance	C(s)	H₂(g)	C <sub>7</sub> H <sub>8</sub> (I)
Standard enthalpy of combustion, $\Delta H_{c}^{\circ}$ /kJ mol <sup>-1</sup>	-394	-286	-3909

$$7C(s) + 4H_2(g) \rightarrow C_7H_8(I)$$
(3)

(3)

(c) An experiment was carried out to determine a value for the enthalpy of combustion of liquid methylbenzene using the apparatus shown in the diagram.

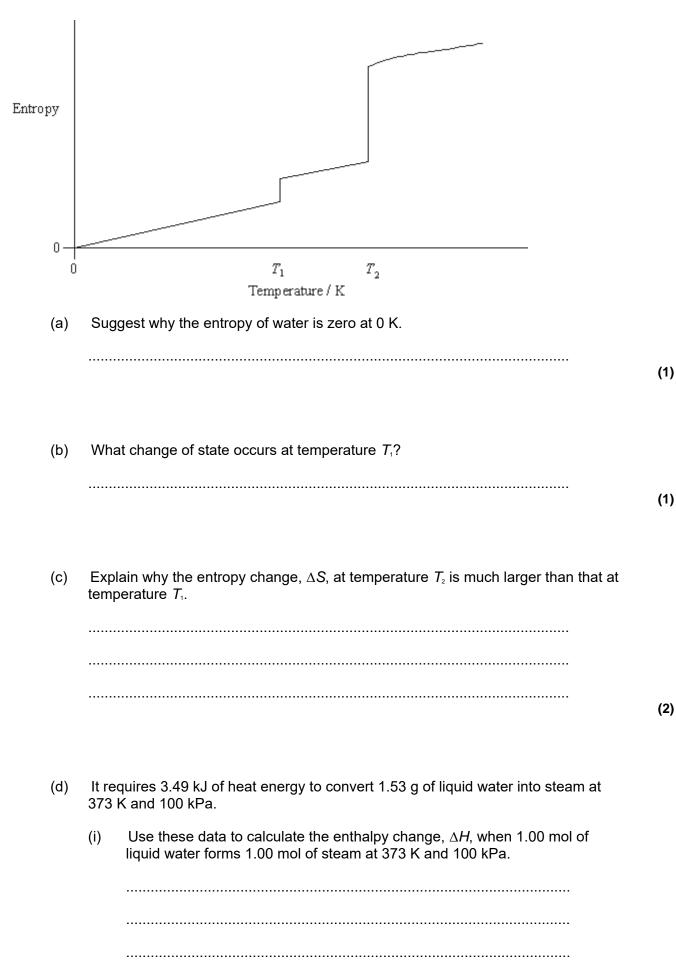


Burning 2.5 g of methylbenzene caused the temperature of 250 g of water to rise by 60°C. Use this information to calculate a value for the enthalpy of combustion of methylbenzene,  $C_7H_{\scriptscriptstyle 8}$ 

(The specific heat capacity of water is 4.18 J K⁻¹ g⁻¹. Ignore the container.)	heat capacity of the
container.)	

		(4)
(d)	A 25.0 cm³ sample of 2.00 mol dm⁻³ hydrochloric acid was mixed with 50.0 cm³ of a 1.00 mol dm⁻³ solution of sodium hydroxide. Both solutions were initially at 18.0 °C.	
	After mixing, the temperature of the final solution was 26.5°C.	
	Use this information to calculate a value for the standard enthalpy change for the following reaction.	
	$HCI(aq) + NaOH(aq) \rightarrow NaCI(aq) + H_2O(I)$	
	In your calculation, assume that the density of the final solution is 1.00 g cm <sup>-3</sup> and that its specific heat capacity is the same as that of water. (Ignore the heat capacity of the container.)	
		(4)
(e)	Give <b>one</b> reason why your answer to part (d) has a much smaller experimental error than your answer to part (c).	
	(Total 15 ma	(1) rks)

**Q2.** The sketch graph below shows how the entropy of a sample of water varies with temperature.



(ii)	Write an expression showing the relationship between free-energy c $\Delta G$ , enthalpy change, $\Delta H$ , and entropy change, $\Delta S$ .	hange,
	-, 1, 3, , 1, 3, -	
(iii)	For the conversion of liquid water into steam at 373 K and 100 kPa, $\Delta G$ = 0 kJ mol <sup>-1</sup>	
	Calculate the value of $\Delta S$ for the conversion of one mole of water integrated under these conditions. State the units.	o steam
	(If you have been unable to complete part (d)(i) you should assume that $\Delta H = 45.0 \text{ kJ mol}^{-1}$ . This is not the correct answer.)	
	Calculation	
	11.2	
	Units	(6) (Total 10 marks)
		( . J. cai i v iiiai ka)

- **Q3.** Methanol, CH<sub>3</sub>OH, 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 J K<sup>-1</sup> g<sup>-1</sup>)
    - (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.

**(7)** 

(Β)		stages.	
		Stage 1 $CH_4(g) + H_2O(g) = 3H_2(g) + CO(g)$ $\Delta H^e = +206 \text{ kJ mol}^{-1}$	
		Stage 2 $CO(g) + 2H_2(g) \longrightarrow CH_3OH(g)$ $\Delta H^0 = -91 \text{ kJ mol}^{-1}$	
	(i)	Explain why, in <i>Stage 1</i> , a higher yield of hydrogen and carbon monoxide is <b>not</b> obtained if the pressure is increased.	
	(ii)	Stage 2 is carried out at a compromise temperature of 500K. By considering what would happen at higher and lower temperatures, explain why 500 K is considered to be a compromise for Stage 2.	(5)
(c)	–283 chan	standard enthalpies of combustion of carbon monoxide and of hydrogen are kJ mol <sup>-1</sup> and –286 kJ mol <sup>-1</sup> , respectively. Use these data and the enthalpy ge for <i>Stage 2</i> to calculate a value for the standard enthalpy of combustion of ous methanol.	(3)

	(Total 15 marks	s)