

**Further Exercises**

## Chapter 2

- W1. Calculate the change in work,  $w$ , for a system that releases 750 J of heat in a process for which the internal energy increases by 220 J.
- W2. Starting from the definition of enthalpy,  $H = U + pV$ , derive a relation between the two heat capacities,  $C_p$  and  $C_v$ , for an ideal gas.
- W3. A 8.0 g sample of methane occupies 10.0 dm<sup>3</sup> at 325 K. Calculate the work done when the gas expands isothermally against a constant external pressure of 2.0 bar until its volume has increased by 5.0 dm<sup>3</sup>. How much work would be done if the expansion occurred reversibly.
- W4. The reaction used to inflate car ‘air bags’ is the decomposition of sodium azide, NaN<sub>3</sub>
- $$\text{NaN}_3(\text{s}) \rightarrow \text{Na}(\text{s}) + 1.5 \text{N}_2(\text{g}) \quad \Delta_r H^\circ = -21.7 \text{ kJ mol}^{-1}$$
- Assuming an airbag needs 60 dm<sup>3</sup> of gas to inflate at 25 °C, calculate:
- the mass of NaN<sub>3</sub> required to produce this volume of gas.
  - the volume of gas produced by this mass on a cold day when the temperature in the car might be 10 °C.
  - the enthalpy change when the airbag is inflated at 298 K.
- W5. Calculate how much heat is required to convert 5 g of ice at 0 °C to steam at 100 °C. What happens to the H<sub>2</sub>O molecules at each stage?
- W6. Calcium carbonate occurs in several forms, including calcite and aragonite. If  $\Delta_f H = -1206.9 \text{ kJ mol}^{-1}$  for calcite and  $\Delta_f H = -1207.1 \text{ kJ mol}^{-1}$  for aragonite, calculate the enthalpy change for the transition from calcite to aragonite.
- W7. Using the data in Appendix 1, calculate the enthalpy change at 25 °C for:
- $$\text{CO}_{(\text{g})} + \text{H}_2\text{O}_{(\text{g})} \rightarrow \text{CO}_{2(\text{g})} + \text{H}_2(\text{g})$$
- W8. At 298 K, the standard enthalpy change of combustion of hydrogen is  $-285.8 \text{ kJ mol}^{-1}$ . The corresponding values for graphite and methanol are  $-393.5 \text{ kJ mol}^{-1}$  and  $-727.0 \text{ kJ mol}^{-1}$  respectively. Calculate the standard enthalpy change of formation of methanol.
- W9. The complete combustion of ethane releases  $1558.8 \text{ kJ mol}^{-1}$  at 25 °C. Calculate  $\Delta H^\circ(\text{combustion})$  at 100 °C.
- $C_p / \text{JK}^{-1} \text{ mol}^{-1}$ : C<sub>2</sub>H<sub>6(g)</sub> 52.6: O<sub>2(g)</sub> 29.4: CO<sub>2(g)</sub> 37.1: H<sub>2</sub>O<sub>(l)</sub> 75.3

- W10. A piece of apple weighing 2.5 g was burned in oxygen in a bomb calorimeter and produced a temperature rise of 2.05 K. In the same calorimeter, the combustion of 0.316 g of benzoic acid produced a temperature rise of 3.24 K.  $\Delta_c U$  for benzoic acid is  $-3251 \text{ kJ mol}^{-1}$ . If the average mass of an apple is 175 g, how much energy in (a) kJ and (b) Calories comes from an apple?
- W11. A bomb calorimeter was calibrated by igniting a 0.7807 g sample of benzoic acid in excess oxygen. The temperature of the calorimeter rose by 1.940 K from 298 K. The internal energy of combustion of benzoic acid is  $-3251 \text{ kJ mol}^{-1}$ . Combustion of 0.9008 g of  $\alpha$ -D-glucose gave a temperature rise of 1.311 K in the calorimeter.
- Using this information, together with data from Appendix 1, find the enthalpy of formation of  $\alpha$ -D-glucose at 298 K and the enthalpy of combustion at the human body temperature of 37 °C.