Q1.The table below shows data for the four hydrocarbons ethyne, propyne, propene and propane. ΔHc is the standard enthalpy of combustion of these hydrocarbons.

| Compound | Name | M , | −Δ <i>H</i> c → / kJ mol ⁻¹ |
|---|---------|------------|---|
| HC≡CH | ethyne | 26 | 1300 |
| HC≡CCH₃ | propyne | 40 | 1940 |
| H ₂ C=CHCH ₃ | propene | 42 | 2060 |
| CH ₃ CH ₂ CH ₃ | propane | 44 | 2220 |

The complete combustion of 2.0 g of one of the above hydrocarbons releases exactly 100 kJ of heat energy.

This hydrocarbon is

- A ethyne
- B propyne
- **C** propene
- D propane

(Total 1 mark)

Q2.When 0.10 g of propane was burned the quantity of heat evolved was 5.0 kJ. The enthalpy of combustion of propane in kJ mol⁻¹ is

- **A** -800
- **B** -1500
- **c** –2200
- **D** –2900

Q3.This question is about the reaction given below.

$$CO(g) + H_2O(g) \implies CO_2(g) + H_2(g)$$

Enthalpy data for the reacting species are given in the table below.

| Substance | CO(g) | H ₂ O(g) | CO ₂ (g) | H₂(g) |
|-----------------------------------|-------|---------------------|---------------------|-------|
| Δ <i>H</i> / kJ mol ⁻¹ | -110 | -242 | -394 | 0 |

The standard enthalpy change for this reaction of carbon monoxide and steam is

- **A** +42 kJ mol⁻¹
- B −42 kJ mol⁻¹
- **C** +262 kJ mol⁻¹
- D –262 kJ mol⁻¹

(Total 1 mark)

Q4.Use the information below to answer this question.

| $C(s) + O_2(g) \rightarrow CO_2(g)$ | $\Delta H^{\bullet} = -394 \text{ kJ mol}^{-1}$ |
|--|---|
| $H_2(g) + \frac{1}{2}O_2(g) \rightarrow H_2O(I)$ | Δ <i>H</i> = −286 kJ mol ⁻¹ |
| $4C(s) + 5H_2(g) \rightarrow C_4H_{10}(g)$ | $\Delta H^{\bullet} = -126 \text{ kJ mol}^{-1}$ |

The standard enthalpy of combustion of butane, in kJ mol⁻¹, is

A -2880

B –2590

C –806

D –554

Q5.Use the information below to answer this question.

| $C(s) + O_2(g) \rightarrow CO_2(g)$ | ΔH = -393.5 kJ mol ⁻¹ |
|--|---|
| $H_2(g) + \frac{1}{2}O_2(g) \rightarrow H_2O(I)$ | $\Delta H^{\bullet} = -285.8 \text{ kJ mol}^{-1}$ |
| $3C(s) + 4H_2(g) \rightarrow C_3H_8(g)$ | Δ <i>H</i> = −104.0 kJ mol ⁻¹ |
| $4C(s) + 5H_2(g) \rightarrow C_4H_{10}(g)$ | Δ <i>H</i> = -125.2 kJ mol ⁻¹ |

The value in kJ mol $^{\mbox{\tiny -1}}$ of the enthalpy of thermal dissociation when but ane forms propane, hydrogen and carbon is

- A −26.3B −17.5
- **C** +17.5
- **C** +21.2

(Total 1 mark)

Q6.Use the information below to answer this question.

| $C(s) + O_2(g) \rightarrow CO_2(g)$ | Δ <i>H</i> = −393.5 kJ mol ⁻¹ |
|---|---|
| $H_2(g) + \frac{1}{2}O_2(g) \rightarrow H_2O(I)$ | $\Delta H = -285.8 \text{ kJ mol}^{-1}$ |
| $3C(s) + 4H_2(g) \rightarrow C_3H_8(g)$ | $\Delta H^{•}$ = -104.0 kJ mol ⁻¹ |
| $4C(s) + 5H_{\scriptscriptstyle 2}(g) \to C_{\scriptscriptstyle 4}H_{\scriptscriptstyle 10}(g)$ | $\Delta H^{\bullet} = -125.2 \text{ kJ mol}^{-1}$ |

The value in kJ mol⁻¹ for the enthalpy of combustion of propane is

- **A** –211.7
- **B** -419.7
- **C** –2220
- **C** –2878

Q7.The data below refer to the industrial production of nitric acid from ammonia.

Reaction 1
$$4NH_3(g) + 5O_2(g)$$
 $4NO(g) + 6H_2O(g)$ $\Delta H^{•} = -909 \text{ kJ mol}^{-1}$ Reaction 2 $2NO(g) + O_2(g)$ $2NO_2(g)$ $\Delta H^{•} = -115 \text{ kJ mol}^{-1}$ Reaction 3 $3NO_2(g) + H_2O(I)$ $2HNO_3(aq) + NO(g)$ $\Delta H^{•} = -117 \text{ kJ mol}^{-1}$

The direct oxidation of ammonia to nitrogen dioxide can be represented by the equation

$$4NH_3(g) + 7O_2(g) \rightarrow 4NO_2(g) + 6H_2O(g)$$

for which the standard enthalpy change, in kJ mol⁻¹, is

- **A** -1139
- **B** -1024
- **C** –794
- **D** -679

Q8.Using the information below, answer this question.

 $Fe_2O_3(s) + 3H_2(g) \rightarrow 2Fe(s) + 3H_2O(g) \quad \Delta H \stackrel{•}{=} +96 \text{ kJ mol}^{-1}, \ \Delta S \stackrel{\bullet}{=} +138 \text{ J } \text{K}^{-1} \text{ mol}^{-1}$ $Fe_2O_3(s) \qquad H_2(g) \qquad Fe(s)$

| 0 | 022.0 | 0 | |
|--|--------|-------|------|
| Δ <i>H</i> r / kJ mol ⁻¹ | -822.0 | 0 | 0 |
| Δ 5 / J K ⁻¹ mol ⁻¹ | 90.0 | 131.0 | 27.0 |

The standard enthalpy of formation of steam is

- A +286 kJ mol⁻¹
- **B** +242 kJ mol⁻¹
- C −242 KJ mol⁻¹
- D –286 kJ mol⁻¹

(Total 1 mark)

Q9. Using the data below, which is the correct value for the standard enthalpy of formation for TiCl₄(I)?

| $C(s) + TiO_2(s) + 2CI_2(g) \rightarrow TiCI_4(I) + CO_2(g)$ | ΔH = −232 kJ mol ⁻¹ |
|--|---|
| $Ti(s) + O_2(g) \rightarrow TiO_2(s)$ | $\Delta H_{\mathbf{f}}^{\mathbf{P}} = -912 \text{ kJ mol}^{-1}$ |
| $C(s) + O_2(g) \rightarrow CO_2(g)$ | $\Delta H_{\rm f}^{\rm e}$ = -394 kJ mol ⁻¹ |
| A −1538 kJ mol ⁻¹ | |

B –1094 kJ mol⁻¹

C −750 kJ mol⁻¹

D +286 kJ mol⁻¹

Q10.When ethanamide (CH₃CONH₂) burns in oxygen the carbon is converted into carbon dioxide, the hydrogen is converted into water and the nitrogen forms nitrogen gas.

| Substance | ethanamide | carbon dioxide | water |
|--|------------|----------------|-------|
| Enthalpy of formation ($\Delta H^{m{	au}}$) / kJ mol $^{-1}$ | -320 | -394 | -286 |

Using the data above, which one of the following is a correct value for the enthalpy of combustion of ethanamide?

- A −1823 kJ mol⁻¹
- B −1183 kJ mol⁻¹
- **C** –1000 kJ mol⁻¹
- **D** -360 kJ mo1⁻¹

(Total 1 mark)

- **Q11.**In which one of the following reactions is the standard enthalpy change equal to the standard enthalpy of formation of lithium fluoride?
 - A $Li(g) + F(g) \rightarrow LiF(s)$
 - **B** $Li^{\dagger}(g) + F^{-}(g) \rightarrow LiF(s)$
 - **C** $Li^{+}(aq) + F^{-}(g) \rightarrow LiF(s)$
 - **D** Li(s) + $\frac{1}{2}F_2(g) \rightarrow \text{LiF}(s)$

Q12.Consider the reactions

$$C_{2}H_{4}(g) + 2O_{2}(g) \rightarrow 2CO(g) + 2H_{2}O(g) \qquad \qquad \Delta H^{\bigoplus} = -758 \text{ kJ mol}^{-1}$$

$$2C(s) + 2H_{2}(g) \rightarrow C_{2}H_{4}(g) \qquad \qquad \Delta H^{\bigoplus} = +52 \text{ kJ mol}^{-1}$$

$$H_{2}(g) + \frac{1}{2} O_{2}(g) \rightarrow H_{2}O(g) \qquad \qquad \Delta H^{\bigoplus} = -242 \text{ kJ mol}^{-1}$$

The enthalpy of formation of carbon monoxide is

- A –111 kJ mol⁻¹
- B −163 kJ mol⁻¹
- C −222 kJ mol⁻¹
- **D** -464 kJ mol⁻¹

(Total 1 mark)

Q13. Given the following data

| $C(s) + 2H_2(g) \rightarrow CH_4(g)$ | <i>∆H</i> = −75 kJ mol ⁻¹ | |
|--------------------------------------|---------------------------------------|--|
| $H_2(g) \rightarrow 2H(g)$ | <i>ΔH</i> = +436 kJ mol ⁻¹ | |

which one of the following is the enthalpy change, in kJ mol⁻¹, of the reaction below?

| $CH_4(g) \rightarrow C(s) + 4H(g)$ | |
|------------------------------------|--|
| -947 | |
| +511 | |

C +797

Α

В

D +947

Q14.Nitric acid is produced industrially from ammonia, air and water using the following sequence of reactions:

| (1) | $4NH_3(g) + 5O_2(g) \rightarrow 4NO(g) + 6H_2O(g)$ | $\Delta H = -909 \text{ kJ mol}^{-1}$ |
|-----|---|--|
| (2) | $2NO(g) + O_2(g) \rightarrow 2NO_2(g)$ | $\Delta H = -115 \text{ kJ mol}^{-1}$ |
| (3) | $3NO_2(g) + H_2O(I) \rightarrow 2HNO_3(aq) + NO(g)$ | Δ <i>H</i> = –117 kJ mol ⁻¹ |

Which is the enthalpy change (in kJ mol⁻¹) for the following reaction?

 $4NH_3(g) + 7O_2(g) \rightarrow 4NO_2(g) + 6H_2O(g)$

- **A** -679
- **B** −794
- **C** -1024
- **D** -1139