

A-Level Chemistry

Energetics
(Multiple Choice)

Question Paper

Time available: 33 minutes Marks available: 30 marks

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Which reaction has an enthalpy change equal to the standard enthalpy of formation of lithium fluoride?

A Li(g) +
$$\frac{1}{2}$$
F₂(g) \rightarrow LiF(s)

$$\textbf{B} \qquad \text{Li}^+(g) + \text{F}^-(g) \rightarrow \text{LiF}(s)$$

C Li⁺(aq) + F⁻(aq)
$$\rightarrow$$
 LiF(s)

D Li(s) +
$$\frac{1}{2}$$
F₂(g) \rightarrow LiF(s)

(Total 1 mark)

2.

Two reactions of iron with oxygen are shown.

$$Fe(s) + \frac{1}{2}O_2(g) \rightarrow FeO(s)$$

$$\Delta H = -272 \text{ kJ mol}^{-1}$$

$$2\; Fe(s) + \frac{3}{2}O_2(g) \rightarrow Fe_2O_3(s)$$

$$\Delta H = -822 \text{ kJ mol}^{-1}$$

What is the enthalpy change, in kJ mol⁻¹, for this reaction?

2 FeO(s) +
$$\frac{1}{2}$$
O₂(g) \rightarrow Fe₂O₃(s)

Some enthalpy change data are shown.

$$C(s) + 2 H_2(g) \rightarrow CH_4(g) \qquad \Delta H = -75 \text{ kJ mol}^{-1}$$

$$\Delta H = -75 \text{ kJ mol}^{-1}$$

$$H_2(g) \rightarrow 2 H(g)$$

$$H_2(g) \to 2 H(g)$$
 $\Delta H = +436 \text{ kJ mol}^{-1}$

What is the enthalpy change, in kJ mol⁻¹, for the following reaction?

$$CH_4(g) \rightarrow C(s) + 4 H(g)$$

Α -947

В -361

C +361

D +947

(Total 1 mark)

4.

The temperature changed from 21.8 °C to 19.2 °C during a calorimetry experiment.

The uncertainty of each reading of the thermometer is ±0.1 °C

What is the percentage uncertainty in the temperature change?

Α 0.5% 0

1.0% В

C 3.8%

D 7.7% 0

	5.
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Which equation represents the reaction that has a standard enthalpy change equal to the standard enthalpy of formation for barium chloride?

- **A** Ba(g) + $Cl_2(g) \rightarrow BaCl_2(s)$
- 0
- **B** Ba²⁺(g) + 2 Cl⁻(g) \rightarrow BaCl₂(s)
- 0
- $\textbf{C} \qquad \text{Ba(s)} + \text{Cl}_2(\text{g}) \rightarrow \text{BaCl}_2(\text{s})$
- 0
- **D** Ba²⁺(s) + 2 Cl⁻(g) \rightarrow BaCl₂(s)
- 0

(Total 1 mark)



Some fuel in a spirit burner is burned, and the heat produced is used to heat a container of water. In this experiment:

The mass of water heated = m g

The temperature rise = y °C

The specific heat capacity of water = $c \, J \, K^{-1} \, g^{-1}$

What is the amount of heat energy absorbed by the water?

A mcy

0

B mc(y + 273)

0

C y/mc

- 0
- **D** (y + 273) / mc
- 0

Nitrogen dioxide is produced from ammonia and air as shown in these equations

$$4 \text{ NH}_3(g) + 5 \text{ O}_2(g) \rightarrow 4 \text{ NO}(g) + 6 \text{ H}_2\text{O}(g)$$
 $\Delta H = -909 \text{ kJ mol}^{-1}$

$$\Delta H = -909 \text{ kJ mol}^{-1}$$

$$2 \text{ NO(g)} + \text{O}_2 \text{ (g)} \rightarrow 2 \text{ NO}_2 \text{ (g)}$$

$$\Delta H = -115 \text{ kJ mol}^{-1}$$

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

4 NH₃(g) + 7 O₂ (g)
$$\rightarrow$$
 4 NO₂ (g) + 6 H₂O(g)

-679

0

-794

-1024

-1139

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(Total 1 mark)

What is the enthalpy of formation of buta-1,3-diene, $C_4H_6(g)$? 8.

Substance	Enthalpy of combustion / kJ mol ⁻¹
$C_4H_6(g)$	-2546
C(s)	-394
H ₂ (g)	-286

+112 kJ mol⁻¹

-112 kJ mol⁻¹

+746 kJ mol⁻¹

D -746 kJ mol⁻¹

0

The heat released when 1.00 g of ethanol ($M_r = 46.0$) undergoes complete combustion is 29.8 kJ

What is the heat released by each molecule, in joules, when ethanol undergoes complete combustion?

(the Avogadro constant $L = 6.022 \times 10^{23} \text{ mol}^{-1}$)

A $2.28 \times 10^{-18} \text{ J}$

0

B $4.95 \times 10^{-20} \text{ J}$

0

C $2.28 \times 10^{-21} \text{ J}$

0

D $4.95 \times 10^{-23} \text{ J}$

0

(Total 1 mark)

10.

Calculate the enthalpy change, in kJ, for this dissociation of mole of propan-1-ol.

$$C_3H_7OH(g) \longrightarrow 3C(g) + 8H(g) + O(g)$$

	C—H	С—С	C—O	0—Н
Mean bond dissociation enthalpy / kJ mol ⁻¹	412	348	360	463

- **A** -4751
- **B** -4403
- C +4403
- D +4751

What is the temperature rise, in K, when 504 J of heat energy are absorbed by 0.110 kg of solid iron?

Specific heat capacity of iron = 0.448 J K⁻¹ g⁻¹

0

- **A** 9.78×10^{-2}
- **B** 1.02×10^1
- **C** 2.83×10^2
- **D** 1.02×10^4

(Total 1 mark)

12.

Hydrogen is produced by the reaction of methane with steam. The reaction mixture reaches a state of dynamic equilibrium.

$$CH_4(g) + H_2O(g) \rightleftharpoons CO(g) + 3H_2(g)$$
 $\Delta H = +206 \text{ kJ mol}^{-1}$

Some enthalpy data is given in the table.

Bond	C–H	O–H	H–H	C≡H
Bond enthalpy / kJ mol ⁻¹	413	463	436	To be calculated

Use the information in the table and the stated enthalpy change to calculate the missing bond enthalpy.

A 234

0

B 1064

0

C 1476

0

D 1936

0

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

Compound	Name	M r	−Δ <i>H</i> c ⁻ / kJ mol ⁻¹
HC≡CH	ethyne	26	1300
HC≡CCH ₃	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)

14. This question is about the reaction given below.

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

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

Substance	CO(g)	H ₂ O(g)	CO ₂ (g)	H ₂ (g)
ΔH _r → / 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

Use the information below to answer this question.

$$C(s) + O_2(g) \rightarrow CO_2(g)$$
 $\Delta H^{\bullet} = -394 \text{ kJ mol}^{-1}$

$$\Delta H^{\bullet} = -394 \text{ kJ mol}^{-1}$$

$$H_2(g) + \frac{1}{2}O_2(g) \rightarrow H_2O(I)$$
 $\Delta H^{\bullet} = -286 \text{ kJ mol}^{-1}$

$$4C(s) + 5H_2(g) \rightarrow C_4H_{10}(g)$$
 $\Delta H^{\bullet} = -126 \text{ kJ mol}^{-1}$

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

- Α -2880
- В -2590
- C -806
- D -554

(Total 1 mark)

16.

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

$$C(s) + TiO_2(s) + 2CI_2(g) \rightarrow TiCI_4(I) + CO_2(g)$$
 $\Delta H^{\bullet} = -232 \text{ kJ mol}^{-1}$

$$\Delta H^{\bullet} = -232 \text{ kJ mol}^{-1}$$

$$Ti(s) + O_2(g) \rightarrow TiO_2(s)$$

$$\Delta H_{\mathbf{f}}^{\mathbf{F}} = -912 \text{ kJ mol}^{-1}$$

$$C(s) + O_2(g) \rightarrow CO_2(g)$$

$$\Delta H_{\mathbf{f}} = -394 \text{ kJ mol}^{-1}$$

- Α -1538 kJ mol⁻¹
- -1094 kJ mol⁻¹ В
- С -750 kJ mol⁻¹
- D +286 kJ mol⁻¹

Using the information below, answer this question.

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

	Fe ₂ O ₃ (s)	H₂(g)	Fe(s)
ΔH _f → / kJ mol ⁻¹	-822.0	0	0
ΔS ^Φ / 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)

18.

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

Reaction 1
$$4NH_3(g) + 5O_2(g) \implies 4NO(g) + 6H_2O(g)$$

$$\Delta H^{\bullet} = -909 \text{ kJ mol}^{-1}$$

Reaction 2
$$2NO(g) + O_2(g) \implies 2NO_2(g)$$

$$\Delta H^{\bullet} = -115 \text{ kJ mol}^{-1}$$

Reaction 3
$$3NO_2(g) + H_2O(l) \rightleftharpoons 2HNO_3(aq) + NO(g)$$

$$\Delta H^{\odot} = -117 \text{ kJ mol}^{-1}$$

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

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)
$$3NO_2(g) + H_2O(I) \rightarrow 2HNO_3(aq) + NO(g)$$

$$\Delta H = -117 \text{ kJ mol}^{-1}$$

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

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

(Total 1 mark)

20.

An experiment is done to determine the enthalpy of combustion of a fuel using a calorimeter containing water.

b = mass of fuel burned / g

w = mass of water heated / g

 ΔT = temperature rise of water / K

 $M_{\rm r}$ = relative molecular mass of fuel

c = specific heat capacity of water / J K⁻¹ g⁻¹

Which expression gives the enthalpy of combustion (in J mol⁻¹), assuming there is no heat loss?

$$A = -\frac{c w \Delta T M_r}{b}$$

$$\mathsf{B} = -\frac{c\,b\,\Delta T M_\mathsf{r}}{w}$$

$$C = -\frac{c b w M_r}{\Delta T}$$

$$D = -\frac{c b w \Delta T}{M_r}$$

Which reaction has an enthalpy change equal to the standard enthalpy of formation of silver iodide?

A Ag(g) + $\frac{1}{2}$ I₂(g) \rightarrow AgI(s)

0

 $\mathbf{B} \qquad \mathsf{Ag}(\mathsf{s}) + \tfrac{1}{2} \, \mathsf{I}_2(\mathsf{s}) \to \mathsf{AgI}(\mathsf{s})$

0

C $Ag^+(g) + I^-(g) \rightarrow AgI(s)$

- 0
- $\textbf{D} \qquad \text{Ag+(aq)} + \text{I-(aq)} \rightarrow \text{AgI(s)}$
- 0

(Total 1 mark)

22.

Which one of the following has the most covalent character?

- A MgF₂
- **B** MgBr₂
- C AIF₃
- D AlBr₃

(Total 1 mark)

23

The table shows the standard enthalpy of formation, $\Delta_f H^{\theta}$, for some of the substances in the reaction

$$C_2H_6(g) + 6F_2(g) \longrightarrow C_2F_6(g) + 6HF(g)$$
 $\Delta H^{\theta} = -2898 \text{ kJ mol} -1$

	C ₂ H ₆ (g)	C ₂ F ₆ (g)
$\Delta_{\mathrm{f}}H^{\mathrm{ heta}}$ /kJ mol $^{-1}$	-84	-1344

What is the standard enthalpy of formation, in kJ mol⁻¹, for HF(g)?

- **A** -1638
- 0
- **B** –273
- 0
- **C** +273
- 0
- **D** +1638
- 0

Use the information below to answer this question.

$$C(s) + O_2(g) \rightarrow CO_2(g)$$

$$C(s) + O_2(g) \rightarrow CO_2(g)$$
 $\Delta H^{\bullet} = -393.5 \text{ kJ mol}^{-1}$

$$H_2(g) + \frac{1}{2}O_2(g) \rightarrow H_2O(I)$$
 $\Delta H^{\Theta} = -285.8 \text{ kJ mol}^{-1}$

$$3C(s) + 4H_2(g) \rightarrow C_3H_8(g)$$
 $\Delta H^{\bullet} = -104.0 \text{ kJ mol}^{-1}$

$$\Delta H^{\bullet} = -104.0 \text{ kJ mol}^{-1}$$

$$4C(s) + 5H_2(g) \rightarrow C_4H_{10}(g)$$
 $\Delta H^{\bullet} = -125.2 \text{ kJ mol}^{-1}$

$$\Delta H^{\bullet} = -125.2 \text{ kJ mol}^{-1}$$

The value in kJ mol⁻¹ of the enthalpy of thermal dissociation when butane forms propane, hydrogen and carbon is

(Total 1 mark)

25.

Use the information below to answer this question.

$$C(s) + O_2(g) \rightarrow CO_2(g)$$

$$C(s) + O_2(g) \rightarrow CO_2(g) \qquad \qquad \Delta H^{\bullet} = -393.5 \text{ kJ mol}^{-1}$$

$$H_2(g) + \frac{1}{2}O_2(g) \rightarrow H_2O(I)$$
 $\Delta H^{\bullet} = -285.8 \text{ kJ mol}^{-1}$

$$3C(s) + 4H_2(g) \rightarrow C_3H_8(g)$$
 $\Delta H^{\bullet} = -104.0 \text{ kJ mol}^{-1}$

$$\Delta H^{\bullet} = -104.0 \text{ kJ mol}^{-1}$$

$$4C(s) + 5H_2(g) \rightarrow C_4H_{10}(g)$$
 $\Delta H^{\bullet} = -125.2 \text{ kJ mol}^{-1}$

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

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

(Total 1 mark)



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 $(\Delta H^{?})$ / 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** $-360 \text{ kJ mo}1^{-1}$

(Total 1 mark)

28.

In which one of the following reactions is the standard enthalpy change equal to the standard enthalpy of formation of lithium fluoride?

- **A** $Li(g) + F(g) \rightarrow LiF(s)$
- $\textbf{B} \qquad \text{Li}^+(g) + \text{F}^-(g) \rightarrow \text{LiF}(s)$
- $\textbf{C} \qquad \text{Li}^+(aq) \, + \, \text{F}^-(g) \rightarrow \text{LiF}(s)$
- **D** Li(s) + $\frac{1}{2}$ F₂(g) \rightarrow LiF(s)

Consider the reactions

$$\mathsf{C_2H_4(g)} + 2\mathsf{O_2(g)} \rightarrow 2\mathsf{CO(g)} + 2\mathsf{H_2O(g)}$$

$$\Delta H^{\bullet} = -758 \text{ kJ mol}^{-1}$$

$$2C(s) + 2H_2(g) \rightarrow C_2H_4(g)$$

$$\Delta H^{\bullet} = +52 \text{ kJ mol}^{-1}$$

$$H_2(g) + \frac{1}{2} O_2(g) \rightarrow H_2O(g)$$

$$\Delta H^{\odot} = -242 \text{ kJ mol}^{-1}$$

The enthalpy of formation of carbon monoxide is

(Total 1 mark)

30.

Given the following data

C(s) +
$$2H_2(g) \rightarrow CH_4(g)$$
 $\Delta H = -75 \text{ kJ mol}^{-1}$

$$\Delta H = -75 \text{ kJ mol}^{-1}$$

$$H_2(g) \rightarrow 2H(g)$$

$$\Delta H = +436 \text{ kJ mol}^{-1}$$

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

$$CH_4(g) \rightarrow C(s) + 4H(g)$$