# <u>Chemical energetics – 2023 AS Chemistry 9701</u>

1. Nov/2023/Paper 9701/11/No.9

Which statement about enthalpy changes is correct?

A Enthalpy changes of reaction are always negative.

**B** Enthalpy changes of combustion are always positive.

C Enthalpy changes of formation are always positive.

D Enthalpy changes of neutralisation are always negative.

2. Nov/2023/Paper 9701/11/No.10

What is the definition of standard enthalpy change of neutralisation,  $\Delta H_{\text{neut}}^{\Theta}$ ?

A  $\Delta H_r^{\Theta}$  when one mole of an aqueous acid is neutralised by an aqueous alkali

**B**  $\Delta H_r^{\oplus}$  when one mole of an aqueous alkali is neutralised by an aqueous acid

 $\triangle H_r^{\ominus}$  when one mole of an aqueous acid is neutralised by one mole of an aqueous alkali

 ${\sf D}$   $\Delta H_{\rm r}^{\ominus}$  when an aqueous acid and an aqueous alkali react together to produce one mole of water

3. Nov/2023/Paper\_ 9701/12/No.11

Nitric acid is made industrially by the oxidation of ammonia. The overall equation for the process is shown.

equation 1 
$$NH_3 + 2O_2 \rightarrow HNO_3 + H_2O$$

The process happens in three stages. The equations and enthalpy changes for these stages are given.

1

stage 1 
$$4NH_3 + 5O_2 \rightarrow 4NO + 6H_2O$$
  $\Delta H = -904 \text{ kJ mol}^{-1}$   
stage 2  $2NO + O_2 \rightarrow 2NO_2$   $\Delta H = -114 \text{ kJ mol}^{-1}$   
stage 3  $4NO_2 + O_2 + 2H_2O \rightarrow 4HNO_3$   $\Delta H = -348 \text{ kJ mol}^{-1}$ 

What is the enthalpy change of the process shown in equation 1?

**A** -1480 kJ mol<sup>-1</sup>

**B** -370 kJ mol<sup>-1</sup>

C -341.5 kJ mol<sup>-1</sup>

**D** +82 kJ mol<sup>-1</sup>

4.	Nov/2023/Paper_	9701/21/No.1(e)
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(e) The structure of glucose,  $C_6H_{12}O_6$ , is shown in Fig. 3.2.

#### glucose

Fig. 3.2

(i) Complete Table 3.1 to identify the number of primary, secondary and tertiary alcohol groups present in the structure shown in Fig. 3.2.

Table 3.1

type of alcohol group	primary	secondary	tertiary
number of groups			

[1]

(ii) Separate samples of aqueous glucose are tested with the reagents shown in Table 3.2.

Complete Table 3.2 with the observation for each reaction.

Write "no reaction" if applicable.

Table 3.2

reagent and conditions	observation with glucose
acidified KMnO <sub>4</sub> (aq) and warm	
Fehling's reagent and warm	
alkaline I <sub>2</sub> (aq) and warm	

[3]

(iii) There are many structural isomers of C<sub>6</sub>H<sub>12</sub>O<sub>6</sub>.

Define structural isomers.

F4

# **5.** Nov/2023/Paper\_ 9701/21/No.3(b)

Compounds C and D are alkenes with the same molecular formula, C<sub>5</sub>H<sub>10</sub>.

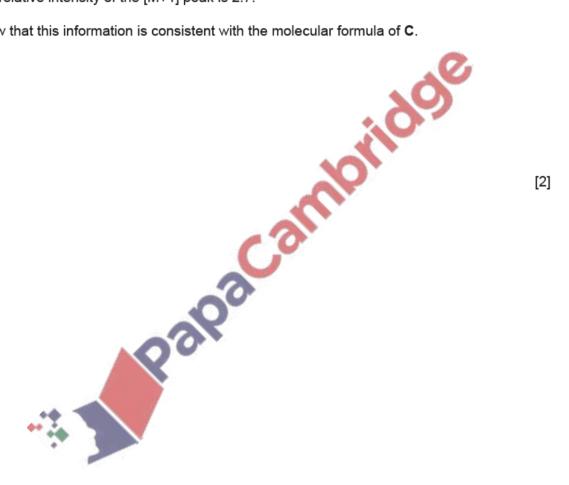


Fig. 4.1

(b) The mass spectrum of C shows a molecular ion peak at m/e = 70. This peak has a relative intensity of 48.7.

The relative intensity of the [M+1] peak is 2.7.

Show that this information is consistent with the molecular formula of C.



**6.** Nov/2023/Paper\_ 9701/22/No.3(b)

(ii)

Phosphoric(V) acid,  $H_3PO_4$ , is used in both inorganic and organic reactions.

(b)  $H_3PO_4$  is also formed in the process shown in reaction 1.

$$\text{reaction 1} \qquad \qquad 4\text{H}_3\text{PO}_3 \, \longrightarrow \, 3\text{H}_3\text{PO}_4 + \text{PH}_3$$

Table 3.1 shows some relevant thermodynamic data.

Table 3.1

compound	enthalpy change of formation, $\Delta H_{\rm f}/{\rm kJmol^{-1}}$
H <sub>3</sub> PO <sub>3</sub>	-972
H <sub>3</sub> PO <sub>4</sub>	-1281
PH <sub>3</sub>	+9

	1113	13	
(i)	Define enthalpy change of formation.	. 2	0
		107	[2
U	se the data in Table 3.1 to calculate the	ne enthalpy change, $\Delta H_{ m r}$ ,	of reaction 1.
		$\Delta H_{\rm r} = \dots$	kJ mol <sup>–1</sup> [2]

(iii) Explain why reaction 1 is a disproportionation reaction.

Explain your reasoning with reference to relevant oxidation numbers.

#### 7. June/2023/Paper 9701/11/No.5

 ${\sf Calcium\ carbide,\ CaC}_2,\ {\sf reacts\ with\ water,\ as\ shown.\ The\ data\ below\ the\ equation\ show,\ in\ kJ\ mol}^{-1},$ the standard enthalpies of formation of the compounds involved.

$$CaC_{2}(s)$$
 +  $H_{2}O(I)$   $\rightarrow$   $CaO(s)$  +  $C_{2}H_{2}(g)$   
-60 -286 -635 +228

What is the standard enthalpy change of the reaction shown?

- A -753 kJ mol<sup>-1</sup>
- **B** -61 kJ mol<sup>-1</sup>
- C +61 kJ mol<sup>-1</sup>
- **D** +753 kJ mol<sup>-1</sup>

# 8. June/2023/Paper\_9701/12/No.9

Vimol d'her A student mixed 25.0 cm<sup>3</sup> of 4.00 mol dm<sup>-3</sup> hydrochloric acid with an equal volume of 4.00 mol dm<sup>-3</sup> sodium hydroxide. The initial temperature of both solutions was 15.0 °C. The maximum temperature recorded was 30.0 °C. The heat capacity of the final solution can be assumed to be 4.18 J K<sup>-1</sup> g<sup>-1</sup> and the density of this solution can be assumed to be 1.00 g cm<sup>-3</sup>.

Using these results, what is the enthalpy change of neutralisation of hydrochloric acid?

- A -62.7 kJ mol<sup>-1</sup>
- B -31.4 kJ mol<sup>-1</sup>
- C -15.7 kJ mol<sup>-1</sup>
- **D** -3.14 kJ mol<sup>-1</sup>

# 9. June/2023/Paper 9701/12/No.10

Nitrogen monoxide is rapidly oxidised to nitrogen dioxide.

$$2NO(g) + O_2(g) \rightarrow 2NO_2(g)$$

Nitrogen dioxide can then dimerise to form dinitrogen tetroxide.

$$2NO_2(g) \rightarrow N_2O_4(g)$$
  $\Delta H^{\oplus} = -58 \text{ kJ mol}^{-1}$ 

$$\Delta H_{\rm f}^{\rm e}NO = +91 \, \rm kJ \, mol^{-1}$$
 and  $\Delta H_{\rm f}^{\rm e}NO_2 = +34 \, \rm kJ \, mol^{-1}$ 

What is the value of the standard enthalpy change for the reaction shown?

$$2NO(g) + O_2(g) \rightarrow N_2O_4(g)$$

- +56 kJ mol<sup>-1</sup>
- **B** −1 kJ mol<sup>-1</sup> **C** −115 kJ mol<sup>-1</sup> **D** −172 kJ mol<sup>-1</sup>

# 10. June/2023/Paper\_9701/13/No.10

Palpacalin Which equation represents the reaction whose standard enthalpy change is the standard enthalpy change of formation of water?

$$\textbf{A} \quad 2H_2(g) \ + \ O_2(g) \ \rightarrow \ 2H_2O(I)$$

$${\bm B} \quad {\bm H}_2(g) \; + \; {\textstyle \frac{1}{2}} {\bm O}_2(g) \; \to \; {\bm H}_2 {\bm O}({\bm I})$$

$$C ext{H}^+(aq) + OH^-(aq) \rightarrow H_2O(I)$$

$$\label{eq:definition} \textbf{D} \quad 2H(g) \ \textbf{+} \ O(g) \ \rightarrow \ H_2O(I)$$

#### 11. June/2023/Paper 9701/13/No.12

The enthalpy change for neutralisation of HNO<sub>3</sub>(aq) with NaOH(aq) is -57.0 kJ mol<sup>-1</sup>.

In an experiment, 20.0 cm<sup>3</sup> of 4.00 mol dm<sup>-3</sup> HNO<sub>3</sub> is mixed with 30.0 cm<sup>3</sup> of 2.00 mol dm<sup>-3</sup> NaOH in an insulated container. The initial temperature of both solutions is 25.0 °C.

It can be assumed that the heat capacity of the product mixture is  $4.2\,\mathrm{J\,cm}^{-3}\,^{\circ}\,\mathrm{C}^{-1}$  and that there are no heat losses.

What is the maximum final temperature of the mixture?

- **A** 41.3 °C
- **B** 44.0 °C
- **C** 46.7 ° C

6

D 52.1 °C

# 12. June/2023/Paper\_9701/13/No.13

Some bond energies are listed.

bond	bond energy /kJ mol <sup>-1</sup>
H–H	436
О-Н	463
0-0	146
0=0	496

One mole of hydrogen reacts with oxygen to give water vapour.

Palpa Calification Using the bond energy data, what is the value for the enthalpy change of this reaction?

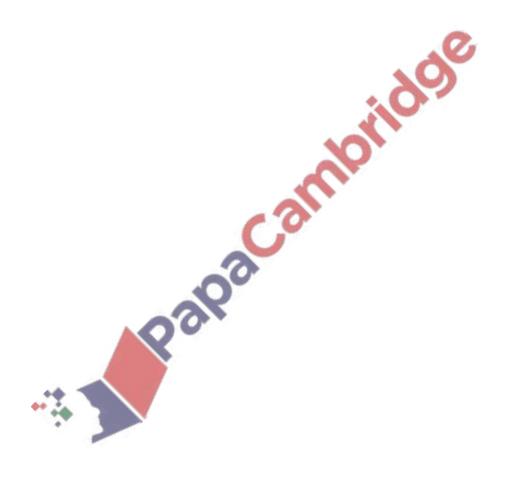
- +221 kJ mol<sup>-1</sup>
- $+6 \,\mathrm{kJ}\,\mathrm{mol}^{-1}$
- $-242 \, kJ \, mol^{-1}$
- $-417 \, \text{kJ} \, \text{mol}^{-1}$

# 13. June/2023/Paper 9701/21/No.2 A neutralisation reaction occurs when NaOH(aq) is added to H<sub>2</sub>SO<sub>4</sub>(aq). equation 1 $2NaOH(aq) + H_2SO_4(aq) \rightarrow Na_2SO_4(aq) + 2H_2O(I)$ (a) Define enthalpy change of neutralisation, $\Delta H_{\text{neut}}$ . (b) An experiment is carried out to calculate $\Delta H_{\text{neut}}$ for the reaction between NaOH(aq) and $H_2SO_4(aq)$ . $100\,\mathrm{cm^3}$ of $1.00\,\mathrm{mol\,dm^{-3}}$ NaOH(aq) is added to $75\,\mathrm{cm^3}$ of $1.00\,\mathrm{mol\,dm^{-3}}$ H<sub>2</sub>SO<sub>4</sub>(aq) in a polystyrene cup and stirred. Results from the experiment are shown in Table 2.1. Table 2.1 initial temperature of NaOH(aq)/°C 20.0 initial temperature of H<sub>2</sub>SO<sub>4</sub>(aq)/°C 20.0 maximum temperature of mixture/°C 27.8 Use equation 1 to calculate the amount, in mol, of H<sub>2</sub>SO<sub>4</sub>(aq) that is neutralised in the experiment. amount of H<sub>2</sub>SO<sub>4</sub>(aq) neutralised = ..... mol [1] Calculate $\Delta H_{neut}$ using the results in Table 2.1. Include units in your answer. Assume that: the specific heat capacity of the final solution is 4.18 J g<sup>-1</sup> K<sup>-1</sup> 1.00 cm<sup>3</sup> of the final solution has a mass of 1.00 g there is no heat loss to the surroundings full dissociation of H<sub>2</sub>SO<sub>4</sub>(aq) occurs the experiment takes place at constant pressure. Show your working.

[3]

c)	(i)	Complete the equation for the reaction that occurs when a solution of $Ba(OH)_2$ is added to aqueous sulfuric acid. Include state symbols.
		$H_2SO_4(aq) +Ba(OH)_2(aq) \rightarrow$ [2]
	(ii)	Suggest why the enthalpy change of neutralisation cannot be determined using the addition of dilute sulfuric acid to aqueous barium hydroxide.
		[1]

[Total: 9]



#### 14. June/2023/Paper\_9701/22/No.4

In industry, ethanol is made by reacting ethene with steam in the presence of H<sub>3</sub>PO<sub>4</sub>.

reaction 1 
$$C_2H_4(g) + H_2O(g) \rightleftharpoons C_2H_5OH(g)$$

(a) Use the bond energy values in Table 4.1 to calculate the enthalpy change,  $\Delta H_{\rm r}$ , for reaction 1.

Table 4.1

bond	bond energy/kJ mol <sup>-1</sup>	· 			
C–C	350	· 			
C=C	610	· 			
C≡C	840	· 			
C–H	410	0			
C-O	360	0			
C=O	740				
O–H	460				
A Palpa Califillo					
ΔH <sub>r</sub> =					
es equilibrium at constant temperature and pressure.					

$\Delta H_r =$	 kJ mol <sup>-1</sup>	[2]

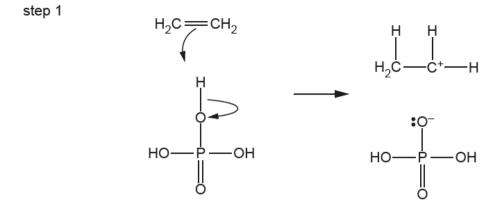
(b) Reaction 1 reaches equilibrium at constant temperature and pressure.

Deduce what effect increasing the pressure will have on the amount of ethanol in the new equilibrium mixture. Use Le Chatelier's principle to explain your answer.

effect of increasing pressure
explanation

[2]

(c) The mechanism for reaction 1 can be described in three steps. Steps 1 and 2 for reaction 1 are shown in Fig. 4.1.



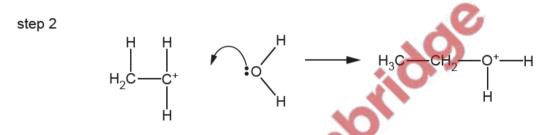


Fig. 4.1

Describe the behaviour of H<sub>3</sub>PO<sub>4</sub> in step 1 in Fig. 4.1. Explain your answer.

[1]

Identify the species that behaves as an electrophile in step 2 in Fig. 4.1. Explain your answer.

[1]

(iii) Complete Fig. 4.2 to show the mechanism for step 3 of reaction 1. Include charges, dipoles, lone pairs of electrons and curly arrows, as appropriate.

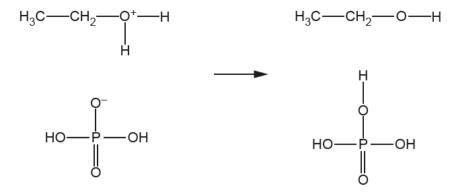


Fig. 4.2

(iv) Describe how a catalyst affects a reaction. Explain your answer.

[2]

(v) Use Fig. 4.1 and Fig. 4.2 to justify why H<sub>3</sub>PO<sub>4</sub> is described as a catalyst in reaction 1.

[1]

(vi) Propene also reacts with steam. A mixture of organic products is produced.

Explain why propan-2-ol is produced in the higher yield.

(d) Describe the covalent bonds present between the carbon atoms in an ethene molecule by completing Table 4.2.

Table 4.2

	sigma (σ)	pi (π)
type of orbitals involved in bond		
how the orbitals overlap		

[2]

[2]

[Total: 15]

			/Paper_9701 lisation read	1/23/No.3 ction occurs whe	n NaOH(aq)	is added to I	H <sub>2</sub> SO <sub>4</sub> (aq).		
е	qua	ation	1	$2NaOH(aq) + H_2SO_4(aq) \to Na_2SO_4(aq) + 2H_2O(I)$					
(a	a)	Defi	Define enthalpy change of		ralisation, Δ <i>F</i>	neut.			
									[2]
(k	b)	H <sub>2</sub> S	O₄(aq) in a	nt, 50.0 cm <sup>3</sup> of 2 a polystyrene cu The maximum te	p and stirred	l. Both solut	ions have a t	temperature o	
		(i)	Use equati experiment	tion 1 to calculat tt.	e the amoun	t, in mol, of	H <sub>2</sub> SO <sub>4</sub> (aq) th	at is neutralise	ed in the
					amount of	H <sub>2</sub> SO <sub>4</sub> neut	ralised =		. mol [1]
		(ii)	Assume the enthal is –57 full dis the sp 1.00 cr there i	Ipy change of ne 7.1 kJ mol <sup>-1</sup> ssociation of H <sub>2</sub> S pecific heat capacism <sup>3</sup> of the final so is no heat loss to operiment takes p	eutralisation, and a second security of the final of the surroun	ΔH <sub>neut</sub> , of Nars rs al solution is mass of 1.00 dings	OH(aq) and I 4.18Jg <sup>–1</sup> K <sup>–1</sup> )g	H <sub>2</sub> SO <sub>4</sub> (aq)	ıt.

(c) The enthalpy change of neutralisation of CH<sub>3</sub>COOH(aq) and NaOH(aq) is -55.2 kJ mol<sup>-1</sup>. Complete the equation for the reaction. Values for the enthalpy change of neutralisation,  $\Delta H_{\text{neut}}$ , are shown in Table 3.1. Table 3.1  $\Delta H_{\rm neut}/\,{\rm kJ\,mol^{-1}}$ reagents NaOH + HC1 -57.1 NaOH + CH<sub>3</sub>COOH -55.2Suggest why the value for  $\Delta H_{\text{neut}}$  of the weak acid, CH<sub>3</sub>COOH, reacting with NaOH is different to the value obtained using the strong acid, HCI. Assume that the values are determined under the same conditions. [Total: 8]

# 16. March/2023/Paper\_9701/12/No.10

The equation for a chemical reaction is shown. All substances are in their standard states.

$$XeF_6 + 3H_2O \rightarrow XeO_3 + 6HF$$

Which statement describes the standard enthalpy change of reaction for this reaction?

- A the enthalpy change when a total of one mole of products is produced
- B the enthalpy change when a total of one mole of reactants is reacted
- C the enthalpy change when one mole of water reacts
- **D** the enthalpy change when six moles of hydrogen fluoride are produced

#### 17. March/2023/Paper\_9701/12/No.16

The table shows bond energies for some diatomic molecules. Deuterium, D, is an isotope of hydrogen.

bond	energy /kJ mol <sup>-1</sup>	
F–F	158	
C1-C1	242	
Br–Br	193	
Į-Į	151	
H-H	436	
D-D	442	

Which statements are correct?

- 1 Diatomic molecules have exact values for their bond energies, which are always positive.
- 2 The trend in Group 7 bond energies can be explained by the variation in instantaneous dipole—induced dipole (id–id) forces.
- 3 A value for the enthalpy change for the reaction between deuterium and chlorine can be calculated using these data alone.
- A 1 only B 1 and 2 only C 1 and 3 only D 2 and 3 only

#### **18.** March/2023/Paper\_9701/22/No.3(d)

(d) The reaction of ethene, C<sub>2</sub>H<sub>4</sub>, with a 1:1 mixture of CO and H<sub>2</sub> is shown in equation 1.

equation 1 
$$C_2H_4(g) + CO(g) + H_2(g) \rightleftharpoons CH_3CH_2CHO(g)$$

At atmospheric pressure a cobalt-based catalyst is used in this reaction.

State and explain the effect of using a catalyst on this reaction.

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(ii)	Explain why the yield of $\mathrm{CH_3CH_2CHO}(g)$ increases when the overall pressure of the reaction mixture is increased.		
			[1]
(iii)	Use the information in Table 3.2 to calculate the enthalpy change, $\Delta H_{\rm r}$ , of the reaction equation 1.		
	equation	$C_2H_4(g) + CO(g)$	+ $H_2(g) \rightleftharpoons CH_3CH_2CHO(g)$
	Table 3.2		
		compound	enthalpy change of formation, Δ <i>H</i> <sub>f</sub> /kJ mol <sup>-1</sup>
		C <sub>2</sub> H <sub>4</sub> (g)	+52
		CO(g)	<b>-111</b>
		CH <sub>3</sub> CH <sub>2</sub> CHO(g)	-187
		ction mixture is cooled to colle	$\Delta H_r = \dots k J \text{ mol}^{-1}$ [2]
(iv)	The reaction mixture is cooled to collect CH CH CHO as a liquid		
(14)	The reaction mixture is cooled to collect CH <sub>3</sub> CH <sub>2</sub> CHO as a liquid.  Identify all types of van der Waals' forces that are present between molecules of CH <sub>3</sub> CH <sub>2</sub> CHO.		32
			[1]