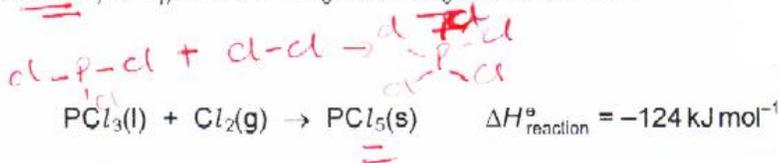


1. Nov/2022/Paper_11/No.9

The enthalpy changes of formation, ΔH_f^\ominus , of both PCl_3 and PCl_5 are exothermic.

PCl_3 reacts with chlorine.



Which pair of statements is correct?

	statement 1	statement 2
A	$\Delta H_{\text{reaction}}^\ominus$ is less negative than $\Delta H_f^\ominus(\text{PCl}_5)$.	The Cl_2 bond energy is needed in calculating $\Delta H_{\text{reaction}}^\ominus$ from enthalpies of formation.
B	$\Delta H_{\text{reaction}}^\ominus$ is more negative than $\Delta H_f^\ominus(\text{PCl}_5)$.	The Cl_2 bond energy is needed in calculating $\Delta H_{\text{reaction}}^\ominus$ from enthalpies of formation.
C	$\Delta H_{\text{reaction}}^\ominus$ is less negative than $\Delta H_f^\ominus(\text{PCl}_5)$.	The Cl_2 bond energy is not needed in calculating $\Delta H_{\text{reaction}}^\ominus$ from enthalpies of formation.
D	$\Delta H_{\text{reaction}}^\ominus$ is more negative than $\Delta H_f^\ominus(\text{PCl}_5)$.	The Cl_2 bond energy is not needed in calculating $\Delta H_{\text{reaction}}^\ominus$ from enthalpies of formation.

Can use Hess laws to compare enthalpy of reaction and formation



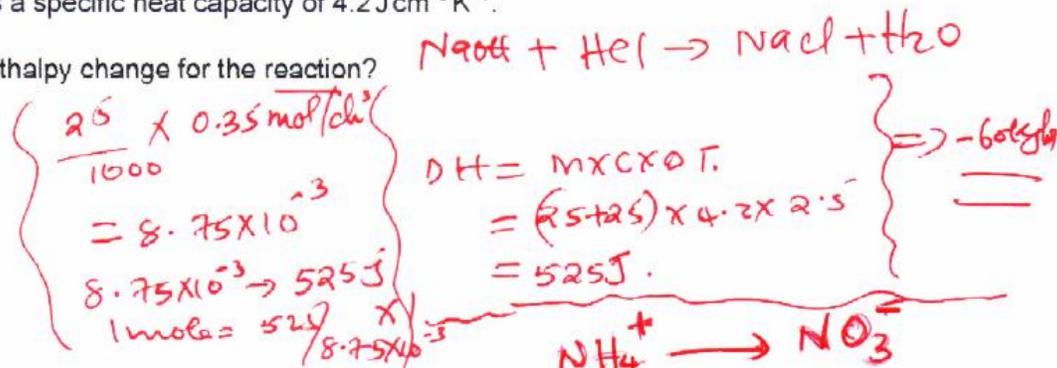
2. Nov/2022/Paper_11/No.10

A student mixes 25.0 cm^3 of $0.350 \text{ mol dm}^{-3}$ sodium hydroxide solution with 25.0 cm^3 of $0.350 \text{ mol dm}^{-3}$ hydrochloric acid. The temperature increases by 2.5°C . No heat is lost to the surroundings.

The final mixture has a specific heat capacity of $4.2 \text{ J cm}^{-3} \text{ K}^{-1}$.

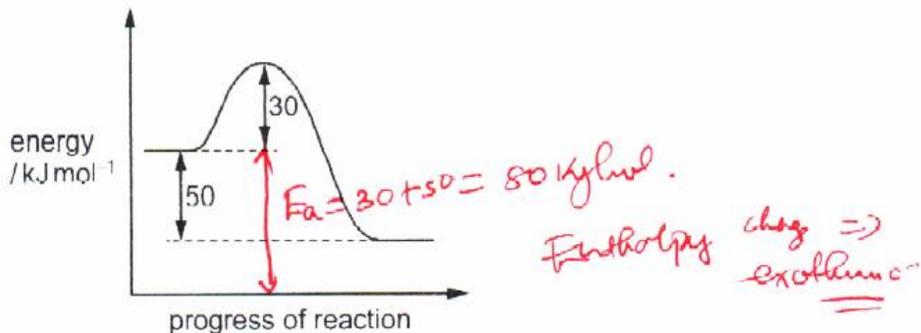
What is the molar enthalpy change for the reaction?

- A -150 kJ mol^{-1}
- B -60 kJ mol^{-1}
- C -30 kJ mol^{-1}
- D $-0.15 \text{ kJ mol}^{-1}$



3. Nov/2022/Paper_12/No.9

The reaction pathway for the forward reaction of a reversible reaction is shown.

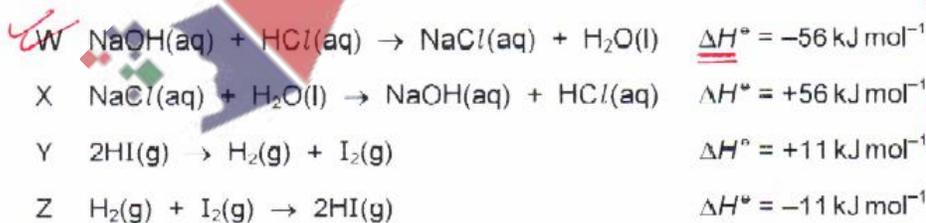


Which statement is correct?

- A The activation energy of the reverse reaction is $+80 \text{ kJ mol}^{-1}$.
- B The enthalpy change for the forward reaction is $+30 \text{ kJ mol}^{-1}$.
- C The enthalpy change for the forward reaction is $+50 \text{ kJ mol}^{-1}$.
- D The enthalpy change for the reverse reaction is $+30 \text{ kJ mol}^{-1}$.

4. Nov/2022/Paper_12/No.10

The enthalpy changes for the possible reactions W, X, Y and Z are given.



feasible reactions have low activation energies

Which statement about the activation energies of these reactions is correct?

- A X is greater than W; Z is greater than Y.
- B X is greater than W; Y is greater than Z.
- C W is greater than X; Z is greater than Y.
- D W is greater than X; Y is greater than Z.

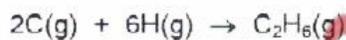
→ reactions that give out heat are exothermic and have a lower activation energy.

Use relevant enthalpy changes from the tables to answer this question.

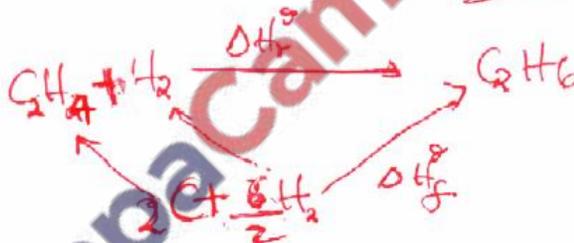
reaction	$\Delta H/\text{kJ mol}^{-1}$
$\text{C(s)} + 2\text{H}_2(\text{g}) \rightarrow \text{CH}_4(\text{g})$	-76
$\text{CH}_4(\text{g}) + 2\text{O}_2(\text{g}) \rightarrow \text{CO}_2(\text{g}) + 2\text{H}_2\text{O}(\text{g})$	-890
$\text{CH}_4(\text{g}) \rightarrow \text{C}(\text{g}) + 4\text{H}(\text{g})$	1648
$3\text{C}(\text{s}) + 4\text{H}_2(\text{g}) \rightarrow \text{C}_3\text{H}_8(\text{g})$	-105

bond	bond enthalpy / kJ mol^{-1}
H-H	436
C-C	350
C=C	610
C=O	805

Which value can be calculated for the enthalpy change for the following reaction?

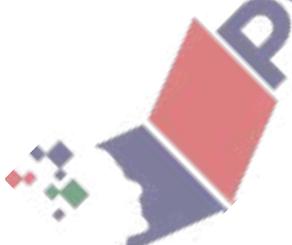


- A $-2822 \text{ kJ mol}^{-1}$
 B $-2122 \text{ kJ mol}^{-1}$
 C $-1998 \text{ kJ mol}^{-1}$
 D $-1772 \text{ kJ mol}^{-1}$



or use bond energies.
 $\text{CH}_4 \rightarrow \text{C} + 4\text{H}$
 and $\text{CH}_2\text{CH}_2 + \text{H}_2 \rightarrow \text{C}_2\text{H}_6$

Hess law. $1 \text{ C-H bond} = 412$
 $6 \text{ C-C} = 412 \times 6 = 2822$



6. Nov/2022/Paper_21/No.3(d)

(d) Sulfur, S_8 , reacts with chlorine to form several different chlorides. The most common are S_2Cl_2 and SCl_2 . SCl_2 forms when sulfur reacts with an excess of chlorine.



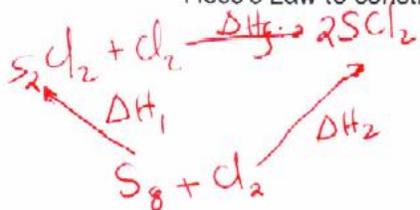
(i) SCl_2 is a cherry-red liquid that reacts vigorously with water to form an acidic solution.

Use this information to deduce the bonding and structure shown by SCl_2 .
Explain your answer.

(i) It forms a simple molecular structure, due to its low b.p.

(ii) The bonding is covalent; its hydrolysed. [2]

(ii) Calculate the enthalpy change of formation, ΔH_f , of $SCl_2(l)$. You may find it useful to use Hess's Law to construct an energy cycle.



$$(-58.2) + (-40.6)$$

$$-58.2 + 4(40.6) = 8\Delta H_f$$

$$\Delta H_f = 27.6$$

enthalpy change of formation of $SCl_2(l)$, $\Delta H_f = -27.6 \text{ kJ mol}^{-1}$ [2]

(iii) State the effect of a decrease in pressure on the position of equilibrium in reaction 2. Explain your answer.

A decrease in pressure will shift the equilibrium to the left, there are more molecules. [1]

Fig. 3.1 shows the two structural isomers of S_2Cl_2 .

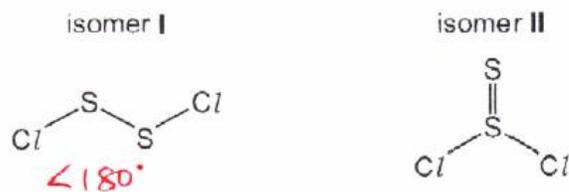


Fig. 3.1

(iv) Define the term structural isomer.

These are organic molecules with the same molecular formula; but different S.F. [2]

(v) Suggest a value for the Cl-S-S bond angle in isomer I. Explain your answer.

bond angle = 109°

explanation Sulfur has two lone pairs of electron and two bonding pairs

- The repulsion from the lone pairs is greater. [2]

(vi) Draw a dot-and-cross diagram to show the bonding in isomer II. Show outer shell electrons only.



[2]

7. June/2022/Paper_11/No.9

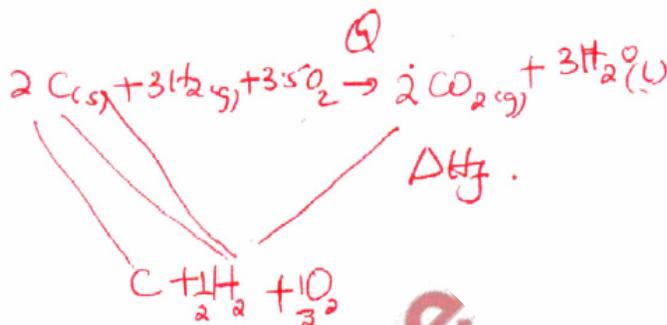
The equation for an enthalpy change is shown. The enthalpy change is Q.

(use Hess' law)



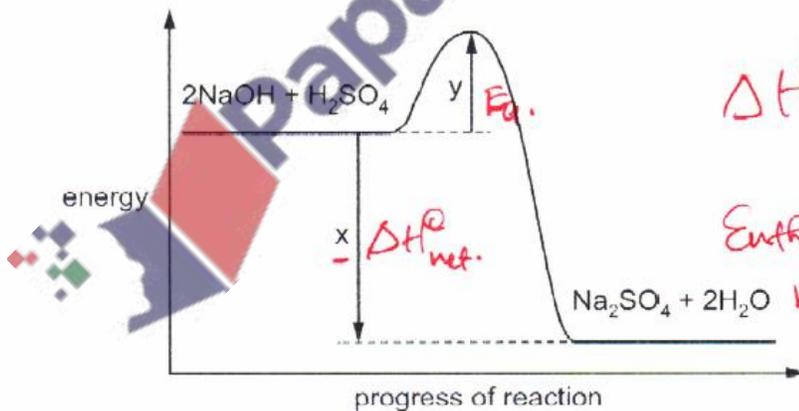
What is the correct expression to calculate Q?

- A $2 \times \Delta H_f^\circ [\text{CO}_2\text{(g)}] - 3 \times \Delta H_f^\circ [\text{H}_2\text{(g)}]$
- B $3 \times \Delta H_f^\circ [\text{H}_2\text{O(g)}] + 2 \times \Delta H_c^\circ [\text{CO}_2\text{(g)}]$
- C $2 \times \Delta H_f^\circ [\text{CO}_2\text{(g)}] - 3 \times \Delta H_f^\circ [\text{H}_2\text{(g)}]$
- D** $3 \times \Delta H_f^\circ [\text{H}_2\text{O(l)}] + 2 \times \Delta H_f^\circ [\text{CO}_2\text{(g)}]$



8. June/2022/Paper_11/No.10

A reaction pathway diagram for the reaction of aqueous sodium hydroxide and dilute sulfuric acid is shown.



ΔH_{neut} is exothermic

Enthalpy change of neutralisation is when one mole of H_2O is formed.

What is the value of the enthalpy change of neutralisation, ΔH_{neut} ?

- A x
- B x - y
- C** $\frac{x}{2}$
- D $\frac{(x - y)}{2}$

$\frac{x}{2}$

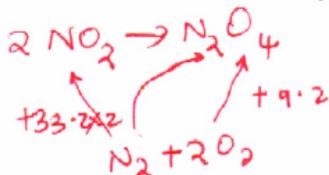
9. June/2022/Paper_12/No.9

The standard enthalpy of formation of $\text{NO}_2(\text{g})$ is $+33.2 \text{ kJ mol}^{-1}$.

The standard enthalpy of formation of $\text{N}_2\text{O}_4(\text{g})$ is $+9.2 \text{ kJ mol}^{-1}$.

What is the standard enthalpy change for the reaction $2\text{NO}_2(\text{g}) \rightarrow \text{N}_2\text{O}_4(\text{g})$?

- A $-57.2 \text{ kJ mol}^{-1}$
- B $-24.0 \text{ kJ mol}^{-1}$
- C $+42.4 \text{ kJ mol}^{-1}$
- D $+75.6 \text{ kJ mol}^{-1}$



$$\begin{aligned}
 (33.2 \times 2) \times \Delta H_f^\circ &= 9.2 \\
 \Delta H_r &= \underline{\underline{-57.2}}
 \end{aligned}$$

10. June/2022/Paper_12/No.10

Separate samples of 25.0 cm^3 of 0.1 mol dm^{-3} $\text{NaOH}(\text{aq})$ are added to each of three different acid solutions, as described. The temperature of each of the solutions was 298 K before mixing.

sample	acid	type of acid	concentration / mol dm^{-3}	volume / cm^3
1	H_2SO_4	strong	0.05	25.0
2	HCl	strong	0.05	25.0
3	$\text{CH}_3\text{CO}_2\text{H}$	weak	0.05	25.0

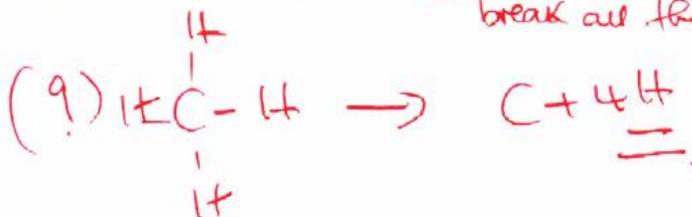
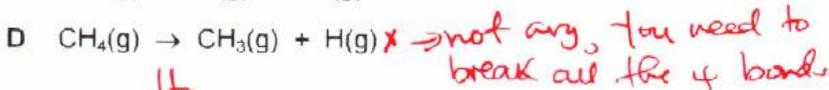
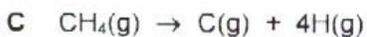
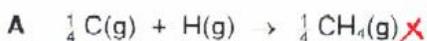
Which statement describes the temperature rises that occur on mixing each of these three acids separately with NaOH ?

- A The temperature rise in all three mixtures is the same.
- B The temperature rise using H_2SO_4 and HCl is the same.
- C The temperature rise using $\text{CH}_3\text{CO}_2\text{H}$ is greater than using HCl .
- D The greatest temperature rise occurs using H_2SO_4 .

\Rightarrow weak acids do not give out much heat.
 \Rightarrow write chemical eqns of each acid with NaOH .
 \Rightarrow calculate moles of each.

11. June/2022/Paper_13/No.9

Which equation represents an enthalpy change that is the average bond energy of the C-H bond in methane?



12. June/2022/Paper_13/No.10

Magnesium carbonate decomposes when heated in a Bunsen burner flame.

Values for the standard enthalpies of formation, ΔH_f° , of the species involved are shown.

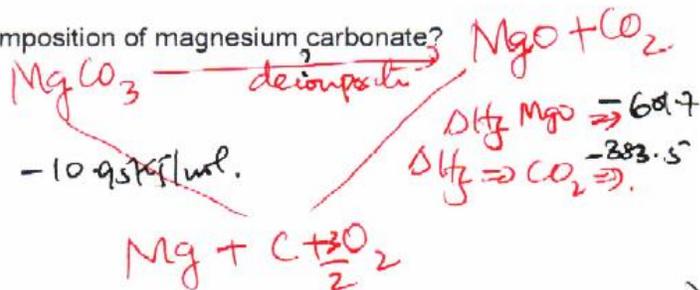
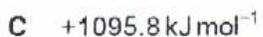
$\Delta H_f^\circ \text{MgCO}_3 = -1095.8 \text{ kJ mol}^{-1}$

$\Delta H_f^\circ \text{MgO} = -601.7 \text{ kJ mol}^{-1}$

$\Delta H_f^\circ \text{CO}_2 = -393.5 \text{ kJ mol}^{-1}$

use Hess diagram

What is the standard enthalpy change for the decomposition of magnesium carbonate?



13. June/2022/Paper_21/No.1(f)

(f) Magnesium, Mg, burns in oxygen, O_2 .
The activation energy, E_a , for this reaction is $+148 \text{ kJ mol}^{-1}$.

(i) State **one** observation when magnesium burns in oxygen.
Do **not** refer to temperature changes in your answer.

Burns with a bright white flame. [1]

(ii) On Fig. 1.1:

- sketch a reaction pathway diagram for the reaction that occurs when Mg burns in O_2
- label the diagram to show the enthalpy change, ΔH , and the activation energy, E_a , for the reaction.

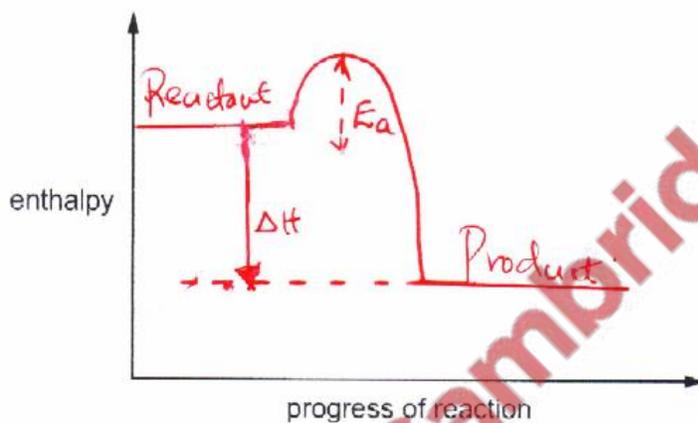
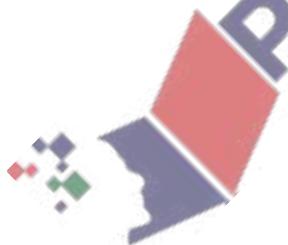


Fig. 1.1

[3]



14. June/2022/Paper_22/No.1(b)

(b) When magnesium is heated in air, magnesium oxide, MgO, is the major product. Smaller amounts of magnesium nitride, Mg₃N₂, are also made.

(i) Calculate the oxidation number for magnesium and for the nitrogen species in Mg₃N₂ to complete Table 1.1.

Table 1.1

species	magnesium in Mg ₃ N ₂	nitrogen in Mg ₃ N ₂
oxidation number	+ II or 2+	- III or 3-

[1]

(ii) Identify the type of reaction which takes place between magnesium and nitrogen. Explain your answer.

Redox reaction. Mg loses its electrons to Nitrogen.

[1]

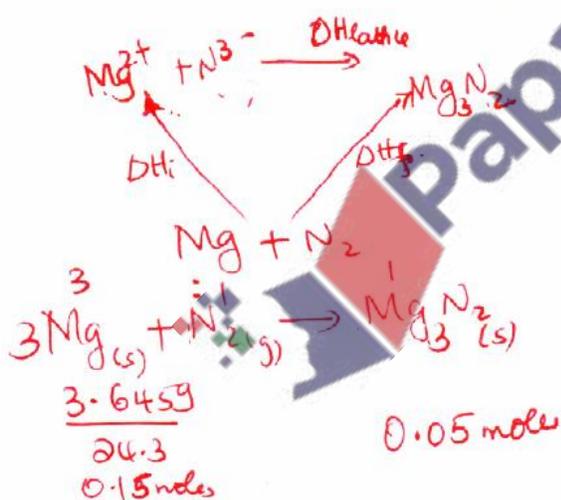
(iii) Define enthalpy change of formation.

Energy change when 1 mole of a substance is formed from its elements under standard conditions.

[2]

(iv) When 3.645g of Mg(s) burns in excess N₂(g) to form Mg₃N₂(s), 23.05kJ of energy is released.

Calculate the enthalpy change of formation, ΔH_f, of Mg₃N₂. Show your working.



$$\text{Mg} : \text{N}_2$$

$$3 : 1$$

$$\text{moles of N}_2 = 0.15 / 3 = 0.05$$

$$\frac{23.05 \text{ kJ}}{0.05} = 461 \text{ kJ/mol}$$

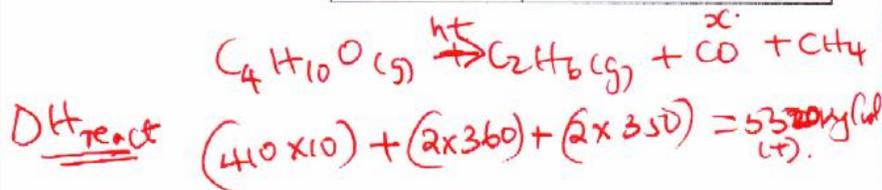
$$\Delta H_f(\text{Mg}_3\text{N}_2) = -461 \text{ kJ/mol}$$

[Total: 9]

- (ii) Calculate the bond energy of $\text{C}\equiv\text{O}$ using the bond energy values in Table 3.1 and the enthalpy change, ΔH , for the thermal decomposition of **G**. Show your working.

Table 3.1

bond	bond energy/kJ mol ⁻¹
C-C	350
C-O (in G)	360
C-H	410



$$-7 = (x + 4450) + (5520)$$

bond energy ($\text{C}\equiv\text{O}$) = +1077 kJ mol⁻¹ [2]

