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## O-level chemistry

## Calculations involving heat in reactions

Reaction involve energy change; exothermic reactions such as burning fire woods produce heat while endothermic reactions such as dissolution of ammonium chloride in water absorb heat.

The amount of heat absorbed or released in a reaction depend on

- Amounts of the reactants
- Temperature and
- Pressure at which a reaction is carried out.

When molar quantities are involved at 298 K and 1 atmosphere, the resultant heat changes are referred to as standard heat/enthalpy changes given a symbol a symbol $\Delta \mathrm{H}$ (delta H ).

Enthalpy changes $(\Delta H)$ (for exothermic reactions carry a negative sign because heat is lost from the system while enthalpy changes $(\Delta H)$ for endothermic reaction carry a positive sign because heat is gained by the system.

Heat is measured in joules (J)

Each heat change is identified by names; the common ones are:

1. Heat of combustion or enthalpy of combustion of a substance is the heat change when 1 mole of substance is burnt completely in oxygen.

## Experiment to determine enthalpy of combustion

A given mass $\left(M_{1} g\right)$ of a substance of molecular mass, $M r$, is burnt completely in excess oxygen. Heat liberated raises the temperature of ( Mw g ) of water through a temperature change of $\theta^{0}$.
Assumption

$$
\begin{array}{ll}
\text { Heat liberated by a burning substance } & =\text { heat absorbed by water } \\
& =\mathrm{MwC} \mathrm{\theta} \text { (where } \mathrm{C}=\text { specific heat capacity of water) } \\
\Rightarrow \mathrm{M}_{1} \mathrm{~g} \text { of the substance produce } & \mathrm{MwC} \mathrm{\theta} \mathrm{~J} \\
\Rightarrow \mathrm{Mr} \text { g of substance produce } & \frac{M_{w} C \theta \times M r}{M_{1}} \mathrm{Jmol}^{-1} .
\end{array}
$$

Therefore, enthalpy of combustion of the substance $=\frac{M_{w} C \theta \times M r}{M_{1}} \mathrm{Jmol}^{-1}$

Experimental depends on whether a fuel is a liquid or a solid.

## Experimental method for finding enthalpy of combustion a liquid fuel

The figure below shows a simple method for obtaining approximate value for the enthalpy of combustion of a fuel.


## Calculations

Assumption
Heat produced by

combusting fuel $\quad$| Heat gained by |
| :--- |
| calorimeter and |
| water |

Heat gained by calorimeter and water = $\mathrm{C} \Theta$ joules

It implies that
$\mathrm{m}_{1} \mathrm{~g}$ of fuel produce $=\mathrm{C} \Theta$ joules

> Mr (mass equivalent to 1
> mole of fuel) produces $=\frac{M r C \theta}{m_{1}}$ joulesmol $^{-1}$

## Example 1

When 23 g of ethanol completely burnt, 13600 KJ of heat was produced.
Calculate the molar heat of combustion of ethanol $(\mathrm{C}=12, \mathrm{H}=1, \mathrm{O}=16)$

## solution

Formula mass of ethanol $\left(\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{OH}\right)=12 \times 2+6+16=46$
23 g of ethanol liberate 13600 KJ

Then, 46 g of ethanol liberate $\frac{13600 \times 46}{23}=27200 \mathrm{~kJ}$
Therefore, the enthalpy of combustion of ethanol $=27200 \mathrm{kJmol}^{-1}$.

## 2. Enthalpy of neutralization

This refers to enthalpy change for the formation 1 mole of water from hydrogen and hydroxide ions $\mathrm{H}^{+}(\mathrm{g})+{ }^{-} \mathrm{OH}(\mathrm{aq}) \rightarrow \mathrm{H}_{2} \mathrm{O}(\mathrm{I})$

## Measurement of standard enthalpy of neutralization

The heat released when a known amount of water is formed is found by measuring the temperature produced in a calorimeter and its contents.


## NB. A vacuum flask is used to minimize heat losses

Know volume of standard acid $\left(\mathrm{V}_{1}\right)$ and alkali $\left(\mathrm{V}_{2}\right)$ are added to calorimeter, and temperature change $\theta^{\circ} \mathrm{C}$ is noted. The number of moles of water formed, $M_{w}$, is calculated

## Calculations

Heat given out $=\quad$ Heat received by
By water calorimeter of capacity, C.

Total volume $=\left(\mathrm{V}_{1}+\mathrm{V}_{2}\right) \mathrm{cm}^{3}$
Given that the density of solution = density of water $\left(\right.$ or $1 \mathrm{gcm}^{-1}$ )
Mass of water, $m=\left(V_{1}+V_{2}\right) g$

Heat $=\mathrm{mc} \theta \mathrm{J}$ (where c is the specific heat capacity

Amount of water formed $=m_{w}$ moles

The standard enthalpy of neutralization is $(\mathrm{mc} \theta) / \mathrm{m}_{\mathrm{w}} \mathrm{Jmol}^{-1}$.

## Example 2

$250 \mathrm{~cm}^{3}$ of 0.40 M NaOH were added to $250 \mathrm{~cm}^{3}$ of 0.40 M HCl in the calorimeter. The temperature of the two solutions was $17.5^{\circ} \mathrm{C}$ and rose to $20.1^{\circ} \mathrm{C}$

Calculate the enthalpy of neutralization assuming that the specific heat capacities of solution are the same as that of water $=4180 \mathrm{~J} \mathrm{~kg}^{-1} \mathrm{~K}^{-1}$.
Solution
Temperature rise $\quad=20.1-17.5=2.6^{\circ} \mathrm{C}$
Mass of solution $\quad=$ total volume of solution

$$
=(250+250)=500 \mathrm{~g}
$$

Heat liberated

$$
=m c \theta
$$

$$
=500 \times 4.180 \times 2.6
$$

$$
=5434 \mathrm{~J}
$$

Mole of water produced $=$ moles of NaOH or moles of HCl

$$
=\frac{250 \times 0.4}{1000}=0.1 \mathrm{~mole}
$$

0.1 mole of water require $=5434 \mathrm{~J}$

1 mole of water require $=\frac{5434 \times 1}{0.1}=54340 \mathrm{~J}$
Therefore, enthalpy of neutralization of water $=54340 \mathrm{Jmol}^{-1}$

Exercise

| 1 | When 1 gram of methanol is burnt in excess air 22.6 kJ of heat was liberated. What is the quantity of heat in kJ liberated when 1 mole of methanol was burnt under similar conditions <br> A. 22.6 <br> B. 32 <br> C. 723.2 <br> D. 777.8 |
| :---: | :---: |
| 2 | 5.3 kJ of energy are required to vaporize 13 g of a liquid X (molecular mass of $X=78$ ) <br> The molar heat of vaporization of X in $\mathrm{kJ} /$ mole is <br> A. $\frac{5.3 \times 78}{13}$ <br> B. $\frac{13 \times 78}{5.3}$ <br> C. $\frac{5.3 \times 13}{78}$ <br> D. $5.3 \times 13 \times 78$ |
| 3 | When 0.4 g of ethanol was burnt, it raised the temperature of 0.1 kg of water by $20^{\circ} \mathrm{C}$. the heat of combustion of ethanol is (specific heat capacity of water $=4.2 \mathrm{~kJ} / \mathrm{kg} /{ }^{\circ} \mathrm{C}, \mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}=46$ ) <br> A. $\frac{4.2 \times 20 \times 46}{0.1 \times 0.4} \mathrm{kjmol}^{-1}$ <br> B. $\frac{0.14 \times 4.2 \times 20}{46 \times 0.1} \mathrm{kjmol}^{-1}$ <br> C. $\frac{0.1 \times 4.2 \times 20 \times 46}{0.4} \mathrm{kjmol}^{-1}$ <br> D. $\frac{0.1 \times 20 \times 46}{46 \times 0.4} \mathrm{kjmol}^{-1}$ |


| 4 | Glucose burn according to the following equation below giving out $2802 \mathrm{kJmol}^{-1}$ of heat energy. $\mathrm{C}_{6} \mathrm{H}_{12} \mathrm{O}_{6}(\mathrm{~s})+6 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow 6 \mathrm{CO}_{2}(\mathrm{~g})+6 \mathrm{H}_{2} \mathrm{O}(\mathrm{I})$ <br> The amount of heat produced when 18 g of glucose is burnt in oxygen at the same temperature is $(H=1, C=12, O=16)$ <br> A. $\frac{2802 \times 18}{180 \times 25}$ <br> B. $\frac{180}{2802 \times 18}$ <br> C. $\frac{180 \times 2518}{2802}$ <br> D. $\frac{2802 \times 18}{180}$ |
| :---: | :---: |
| 5 | Graphite burns in oxygen according to he following equation $\mathrm{C}(\mathrm{~s})+\mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{CO}_{2}(\mathrm{~g}) \quad \Delta \mathrm{H}=-390 \mathrm{kJmol}^{-1}$ <br> When 48 g of graphite is burnt in oxygen the heat produced is <br> A. -97.5 kJ <br> B. -195 kJ <br> C. -780 kJ <br> D. -1560 kJ $(C=12, O=16, H=1)$ |
| 6 | The formation of methanol from hydrogen and carbon monoxide is represented by equation $2 \mathrm{H}_{2}(\mathrm{~g})+\mathrm{CO}(\mathrm{~g}) \longrightarrow \mathrm{CH}_{3} \mathrm{OH}(\mathrm{~g}) \Delta \mathrm{H}=-92 \mathrm{kJmol}-1$ <br> What would be the energy released, in kJ , when 3.2 g of methanol is formed? <br> A. 2.9 <br> B. 3.6 <br> C. 9.2 <br> D. 10.2 |
| 7 | When 2.0 g of substance $X$ were burnt the heat produced raised the temperature of 1000 g of water by $15.6^{\circ} \mathrm{C}$. The molar heat of combustion of $X$ in joules is (the specific heat capacity of water is $4.2 \mathrm{Jg}^{-10} \mathrm{C}$, relative molecular mass of X is 60 ) <br> A. $\frac{1000 \times 4.2 \times 15.6 \times 20}{6 o}$ <br> B. $\frac{15.6 \times 60 \times 1000}{2.0 \times 4.2}$ <br> C. $\frac{15.6 \times 2.0 \times 1000}{4.2 \times 60}$ <br> D. $\frac{4.2 \times 15.6 \times 60}{2}$ |
| 8 | Methanol burns in excess air according to the equation $2 \mathrm{CH}_{3} \mathrm{OH}(\mathrm{I})+3 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow 2 \mathrm{CO}_{2}(\mathrm{~g})+4 \mathrm{H}_{2} \mathrm{O}(\mathrm{I}) \quad \Delta \mathrm{H}=-730 \mathrm{kJmol}$ <br> The amount of heat liberated when 3.2 g of methanol $(\mathrm{Mr}=32)$ is burnt completely is <br> A. 73 kJ <br> B. 730 kJ <br> C. 1416 kJ <br> D. 2929 kJ |
| 9 | Carbon reacts with sulphur according to the following equation $\mathrm{C}(\mathrm{~s})+2 \mathrm{~S}(\mathrm{~s}) \longrightarrow \mathrm{CS}_{2}(\mathrm{~s}) \quad \Delta \mathrm{H}=116 \mathrm{~kJ}$ <br> The amount of heat absorbed when 16 g of sulphur reacts with excess carbon is ( $\mathrm{C}=12, \mathrm{~S}=32$ ) <br> A. 7 kJ <br> B. 29 kJ <br> C.58kJ <br> D. 116 kJ |


| 10 | When 8 g of salt was dissolved in 100 g of water the temperature decreased by $10^{\circ} \mathrm{C}$. The drop in temperature when 2 g of the salt is dissolved in 100 g of water would be ( $\mathrm{H}=1, \mathrm{O}=16$ ) <br> A. $10^{\circ} \mathrm{C}$ <br> B. $8.5^{\circ} \mathrm{C}$ <br> C. $5.0^{\circ} \mathrm{C}$ <br> D. $2.5^{\circ} \mathrm{C}$ |
| :---: | :---: |
| 11 | Ethanol burns in oxygen according to the following equation $\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}(\mathrm{I})+\frac{7}{2} \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow 2 \mathrm{CO}_{2}(\mathrm{~g})+3 \mathrm{H}_{2} \mathrm{O}(\mathrm{I}) \Delta \mathrm{H}-1185 \mathrm{~kJ}$ <br> Calculate heat given out when 0.2 moles of ethanol is burnt completely <br> A. -237 kJ <br> B. -592 kJ <br> C. -1185 kJ <br> D. -2370 kJ |
| 12 | Carbon monoxide reacts with hydrogen according to the equation $\mathrm{CO}(\mathrm{~g})+2 \mathrm{H}_{2}(\mathrm{~g}) \longrightarrow \mathrm{CH}_{3} \mathrm{OH}(\mathrm{I}) \Delta \mathrm{H}=+91 \mathrm{kJmol}^{-1}$ <br> What mass of carbon monoxide would cause heat change of $+188.2 \mathrm{~kJ}(\mathrm{H}=1, \mathrm{C}=12, \mathrm{O}=16)$ <br> A. 2 g <br> B. 28 g <br> C. 58 g <br> D. 273 g |
| 13 | 13.7 kJ of heat was evolved when 4.0 g of copper was displaced from copper (II) sulphate solution by zinc. The amount heat evolved when one mole of copper was displaced is <br> A. $\frac{63.5 \times 4}{13.7}$ <br> B. $\frac{13.7 \times 63.5}{4}$ <br> C. $\frac{13.7 \times 4}{63.5} \quad$ D. $\frac{63.5}{13.7 \times 4}$ |
| 14 | Carbon burns in excess oxygen according to the equation $\mathrm{C}(\mathrm{~s})+\mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{CO}_{2}(\mathrm{~g}) \Delta H-393 \mathrm{~kJ}$ <br> What mass of carbon in grams would produce750kJ of energy <br> A. $\frac{393 \times 12}{750}$ <br> B. $\frac{750 \times 12}{393000}$ <br> C. $\frac{75012}{393}$ <br> D. $\frac{750 \times 393}{12}$ |
| 15 | 10 g of methanol, $\mathrm{CH}_{3} \mathrm{OH}$, burns in air to liberate 226 kJ of heat. The amount of heat liberated when 1 mole of methanol is burn in air is ( $H=1, C=12$ ) <br> A. $\frac{32 \times 226}{10}$ <br> B. $\frac{10 \times 32}{226}$ <br> C. $\frac{10}{32226}$ <br> D. $\frac{10 \times 226}{32}$ |
| 16 | When 1.0 g of carbon is burnt in excess oxygen, the heat produced raises the temperature of 400 g of water by $19^{\circ} \mathrm{C}$. the heat of combustion of carbon is ( $\mathrm{C}=12$, specific heat capacity of water is $4.2 \mathrm{kJkg}^{-1} \mathrm{~K}^{-1}$ ) <br> A. $0.4 \times 4.2 \times 19 \times 12 \mathrm{kJmol}^{-1}$ <br> B. $\frac{0.4 \times 4.2}{12 \times 19} \mathrm{kJmol}^{-1}$ <br> C. $400 \times 4.2 \times 19 \times 12 \mathrm{kJmol}^{-1}$ |


|  | D. $\frac{1219 \times 19}{0.4 \times 4.2} \mathrm{kJmol}^{-1}$ |
| :---: | :---: |
| 17 | Ethanol burns in oxygen according to the following equation $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{OH}(\mathrm{I})+3 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow 2 \mathrm{CO}_{2}(\mathrm{~g})+3 \mathrm{H}_{2} \mathrm{O}(\mathrm{I})$ <br> Calculate the amount of heat evolved when 45 g of oxygen is used for complete combustion of ethanol. ( $C=12, H=1, O+16$, the molar heat of combustion of ethanol is $1370.0 \mathrm{kJmol}^{-1}$ ) <br> A. 642.2 kJ <br> B. 1284.4 kJ <br> C. 1340.2 kJ <br> D. 1926.6 kJ |
| 18 | When 1 mole of ammonium chloride was dissolved in a certain volume of water, 2.94 kJ of heat was absorbed. The amount heat absorbed when 5.35 g of ammonium chloride is dissolved in the same volume of water is $\left(\mathrm{NH}_{4} \mathrm{Cl}=53.5\right)$ <br> A. $\frac{53.5}{2.94 \times 5.53} \mathrm{~kJ}$ <br> B. $\frac{2.94 \times 5.35}{53.5} \mathrm{~kJ}$ <br> C. $\frac{29.4 \times 53.5}{5.35}$ <br> D. $\frac{53.5 \times 5.35}{2.94}$ |
| 19 | When 2.3 g of ethanol was completely burnt in oxygen, heat evolved raised the temperature of 100 g of water by $30^{\circ} \mathrm{C}$. the molar heat of combustion f ethanol is [The molar heat of combustion of ethanol in joule is (the molar mass of ethanol $=46$ and the specific heat capacity of water $=4.2 \mathrm{Jg}^{-1} \mathrm{~K}^{-1}$ ) <br> A. $\frac{100 \times 4.2 \times 30 \times 46}{2.3}$ <br> B. $\frac{30 \times 4.2 \times 2.3 \times 100}{46}$ <br> C. $\frac{20 \times 4.22 .3 \times 100}{46}$ <br> D. $\frac{20 \times 4.2 \times 46100}{2.3}$ |
| 20 | When 2.4 gof magnesium was reacted with $200 \mathrm{~cm}^{3}$ of 2 M hydrochloric acid, 13.6 kJ of heat was evolved. The molar heat of reaction of magnesium with the acid is $(M g=24)$ <br> A. $\frac{13.6 \times 200}{24 \times 2.4} \mathrm{~kJ}$ <br> B. $\frac{13.6 \times 24}{2.4 \times 200}$ <br> c. $\frac{2.4 \times 24}{13.6} \mathrm{~kJ}$ <br> D. $\frac{24 \times 13.6}{2.4} \mathrm{~kJ}$ |
| 21 | Butane undergoes combustion according to the following equation: $2 \mathrm{C}_{4} \mathrm{H}_{10}(\mathrm{~g})+13 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow 8 \mathrm{CO}_{2}(\mathrm{~g})+10 \mathrm{H}_{2} \mathrm{O}(\mathrm{I})+\text { heat }$ <br> The mass of butane required to produce 950 kJ of heat is ( $H=1, C=12,1$ mole of butane produces 2877 kJ of heat) <br> A. $\frac{950 c 58}{2 \times 2877} g$ <br> B. $\frac{950 \times 58}{2877}$ <br> C. $\frac{950 \times 52 \times 2}{2877}$ <br> D. $\frac{2877 \times 58}{950}$ |

## Section B

| 22. | (a) | What is meant by enthalpy of neutralization? (02marks) |
| :---: | :---: | :---: |
|  | (b) | When $50.0 \mathrm{~cm}^{3}$ of 1 M sulphuric acid was added to $50 \mathrm{~cm}^{3}$ of 2 M sodium hydroxide, the temperature rose by $13.6^{\circ} \mathrm{C}$. <br> (i) Write ionic equation for the reaction that took place (01 mark) <br> (ii) Calculate the enthalpy of neutralization of sodium hydroxide. <br> (specific heat capacity of water $=4.2 \mathrm{Jmol}^{-1}$, density of water $1 \mathrm{~g} / \mathrm{cm}^{3}$ (3marks) |
| 23 |  | Biogas contain mainly methane. |
|  | (a) | Name two raw materials that can be used to produce biogas |
|  | (b) | Methane burns in oxygen according to the following equation $\mathrm{CH}_{4}(\mathrm{~g})+2 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{CO}_{2}(\mathrm{~g})+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{I}) \Delta \mathrm{H}=-890 \mathrm{kJmol}^{-1}$ <br> Calculate the volume of methane at s.t.p that will produce 2670 kJ . |
| 24 | (a) | $50 \mathrm{~cm}^{3}$ of 2 M HCl and $50 \mathrm{~cm}^{3}$ of 2 M NaOH each at $22^{\circ} \mathrm{C}$ were mixed in a plastic beaker. The mixture was stirred and its maximum temperature was $35^{\circ} \mathrm{C}$. [heat capacity of solution is $4.2 \mathrm{kkg}^{-10} \mathrm{C}$, density of water $=1 \mathrm{gcm}^{-3}$ ] <br> (i) Write ionic equation for the reaction. <br> (ii) Calculate the heat of reaction |
|  | (b) | $50 \mathrm{~cm}^{3}$ of 2 M ammonia was used instead of 2 M NaOH in (a). State whether the heat of the reaction was greater than, equal to or less than the value obtained in (a)(ii) above. Explain your answer. |
| 25 | (a) | Write equation for complete combustion of methane |
|  | (b) | A litre of methane gas costs 600/=. Calculate the cost of methane required to produce $1746 \times 10^{3}$ J of heat. ( 1 mole of a gas occupies $24 \mathrm{dm}^{3}$ at room temperature, heat of combustion of methane is $-882 \mathrm{kJmol}^{-1}$ ) |
| 26 | (a) | Define the term enthalpy of combustion (02marks) |
|  | (b) | With the aid of a diagram describe an experiment you would carry out in a laboratory to determine the enthalpy of combustion of propanol (9marks) |
|  | (c) | In an experiment to determine the enthalpy of combustion of propanol, 0.54 g of propanol was burnt and the heat evolved caused the temperature of $150 \mathrm{~cm}^{3}$ of water to rise by $21.5^{\circ} \mathrm{C}$ ( molar mass of propanol $=60$, density of water is $1 \mathrm{gcm}^{-3}$, specific heat capacity of |


|  |  | water $=4.1 \mathrm{Jg}^{-1}$ ) <br> Calculate heat capacity experiment value of enthalpy of combustion of propanol ( $31 / 2$ marks) |
| :---: | :---: | :---: |
| 27 |  | Ethane burns in oxygen according to the following equation $2 \mathrm{C}_{2} \mathrm{H}_{6}(\mathrm{~g})+7 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow 4 \mathrm{CO}_{2}(\mathrm{~g})+6 \mathrm{H}_{2} \mathrm{O}(\mathrm{I})$ <br> When 2.0 g of ethane was burnt in excess oxygen, 104 kJ of heat was produced. <br> Calculate |
|  | (a) | Mass of water formed (03marks) |
|  | (b) | Molar heat of combustion of ethane (02marks) |
| 28 | (a) | What is meant by the term enthalpy of combustion? (02mark) |
|  | (b) | Ethanol burns in oxygen according to the following equation $\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}(\mathrm{I})+3 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{CO}_{2}(\mathrm{~g})+3 \mathrm{H}_{2} \mathrm{O}(\mathrm{I}) \quad \Delta H-1360 \mathrm{kJmol}^{-1}$ <br> Calculate the mass of ethanol that is required to raise the temperature of $1000 \mathrm{~cm}^{3}$ of water by $10.0^{\circ} \mathrm{C}$ ( 03 marks ) (specific heat capacity of water $=4.2 \mathrm{Jg}^{-1} \mathrm{~K}^{-1}$ ) |


| 1 | C | Formula mass of methanol, $\mathrm{CH}_{3} \mathrm{OH}=12+3+16+1=32$ <br> 1 g of methanol produce 22.6 kJ <br> 32 g (1mole) produce $22.6 \times 32=723.2 \mathrm{~kJ}$ |
| :---: | :---: | :---: |
| 2 | B | 5.3 g of X require 5.3 kJ 78 g of X require $\frac{5,3 \times 78}{13} \mathrm{~kJ}$ |
| 3 | C | Heat $=\mathrm{mc} \theta=(0.1 \times 4.2 \times 20) \mathrm{kJ}$ <br> 0.4 g of ethanol produce $(0.1 \times 4.2 \times 20) \mathrm{kJ}$ <br> 46 g of ethanol produce $\frac{0.1 \times 4.2 \times 20 \times 46}{0.4}$ |
| 4 | D | Formula mass $\mathrm{C}_{6} \mathrm{H}_{12} \mathrm{O}_{6}$ weigh $6 \times 12+12 \times 1+16 \times 6=180$ 180g glucose produce 2802 kJ <br> 18 g of glucose produce $\frac{2802 \times 18}{180} \mathrm{~kJ}$ |
| 5 | D | 12g produce 390kJ <br> 48 g produce $\frac{390 \times 48}{12}=1560 \mathrm{~kJ}$ |
| 6 | C | Formula mass of methanol, $\mathrm{CH}_{3} \mathrm{OH}=32$ <br> 32 g of $\mathrm{CH}_{3} \mathrm{OH}$ produce 92 kJ <br> 3.2 g of $\mathrm{CH}_{3} \mathrm{OH}$ produce $\frac{92 \times 3.2}{32}=9.2 \mathrm{~kJ}$ |
| 7 | D | $\begin{aligned} & \text { Heat }=\mathrm{mc} \theta=(1000 \times 4.2 \times 15.6) \mathrm{J} \\ & 2 \mathrm{~g} \text { of } \mathrm{X} \text { produce }(1000 \times 4.2 \times 15.6) \mathrm{J} \\ & 60 \mathrm{~g} \text { of } \mathrm{X} \text { produce } \frac{1000 \times 4.2 \times 15.6 \times 60}{2} \mathrm{Jmol}^{-1} \end{aligned}$ |


| 8 | A | 32 g of methanol produce 730kJ <br> 3.2 g of methanol produce $\frac{730 \times 3.2}{32}=73 \mathrm{~kJ}$ |
| :---: | :---: | :---: |
| 9 | B | ( $2 \times 32$ )g of sulphur produce 116 kJ <br> 16 g of sulphur produce $\frac{116 \times 16}{64}=29 \mathrm{~kJ}$ |
| 10 | D | 8 g cause a drop in temperature of $10^{\circ} \mathrm{C}$ <br> 2 g will cause a drop in temperature of $\frac{2 \times 10^{0}}{8} 2.5^{\circ} \mathrm{C}$ |
| 11 | A | 1 mole of ethanol produce 1185 kJ <br> 0.2 mole produce $0.2 \times 1185$ |
| 12 | C | Formula mass of $\mathrm{CO}=12+16=28$ <br> 91 kJ is absorbed by 28 g of CO <br> 1882 kJ is absorbed by $\frac{1882 \times 28}{91}=57.9 \mathrm{~g}$ |
| 13 | B | 4.0 g of copper produce 13.7 kJ <br> 63.5 g of copper produce $\frac{63.5 \times 13.7}{4} \mathrm{~g}$ |
| 14 | C | 393 kJ are produced by 12 g of carbon 750 kJ require $\frac{750 \times 12}{393} \mathrm{~g}$ |
| 15 | D | Formula mass of $\mathrm{CH}_{3} \mathrm{OH}=12+3 \times 1+16+1=32 \mathrm{~g}$ <br> 10 g of methanol produce 226 kJ <br> 32 g (1mole) produce $\frac{226 \times 32}{10} \mathrm{~kJ}$ |
| 16 | A | $400 \mathrm{~g}=\frac{400}{1000}=0.4 \mathrm{~kg}$ <br> Heat produced $=m c \theta=(0.4 \times 4.2 \times 19) \mathrm{kJ}$ <br> 1 g of carbon produces $(0.4 \times 4.2 \times 19) \mathrm{kJ}$ $12 \mathrm{~g} \text { produce } \frac{(0.4 \times 4.2 \times 19 \times 12)}{1} k J$ |


| 17 | B | Formula mass of $\mathrm{O}_{2}=16 \times 2=32$ <br> $(3 \times 32) \mathrm{g}$ of oxygen produce 1370 kJ <br> 45 g of oxygen produce $=\frac{1370 \times 45}{3 \times 32}$ |
| :--- | :--- | :--- |
| 18 | B | Formula mass of $\mathrm{NH}_{4} \mathrm{Cl}=14+1 \times 4+35.5=53.5$ <br> $53.5 \mathrm{gof} \mathrm{NH}_{4} \mathrm{Cl}$ absorb 2.94 kJ <br> 5.35 g of $\mathrm{NH}_{4} \mathrm{Cl}$ absorb $\frac{5.35 \times 2.94}{53.5}$ |
| 19 | A | Formula mass of ethanol, $\mathrm{CH} 3 \mathrm{CH} 2 \mathrm{OH}=12+3+12+2+16+1=46$ <br> Heat $=$ mc $\theta=(100 \times 4.2 \times 30) \mathrm{J}$ <br> 2.3 g of ethanol produce $(100 \times 4.2 \times 30) \mathrm{J}$ <br> 46 g of ethanol produce $\frac{0.1 \times 4.2 \times 30 \times 46}{2.3} \mathrm{~J}$ |
| 20 | D | 2.4 g of magnesium produce 13.6 kJ <br> 24 g of magnesium produce $\frac{24 \times 13.6}{2.4}$ |
| 21 | B | Formula mass of butane, $\mathrm{C}_{4} \mathrm{H}_{10}=4 \times 12+1 \times 10=58$ <br> 2877 kJ is produced by 58 g of butane <br> 950 kg is produced by $\frac{58 \times 950}{2877}$ |

22 (a) Enthalpy of neutralization is enthalpy change when 1 mole of water is formed from aqueous hydrogen and hydroxyl ions.

|  | (b) | (i) $\mathrm{H}^{+}(\mathrm{aq})+\mathrm{OH}^{-}(\mathrm{aq}) \longrightarrow \mathrm{H}_{2} \mathrm{O}(\mathrm{I})$ <br> (ii) total volume of water $=50+50=100 \mathrm{~cm}^{3}$ <br> Mass of water $=100 \times 1=100 \mathrm{~g}$ <br> Heat liberated $=\mathrm{mc} \theta=100 \times 4.2 \times 13.6=5712 \mathrm{~J}$ <br> Mole of water formed $=$ moles $\mathrm{NaOH}=\frac{50 \times 2}{1000}=0.1$ mole <br> Production of 0.1 mole of water produce 5712J <br> 1 mole of water produced $\frac{5712 \times 1}{0.1} 57120 \mathrm{~J}=57.12 \mathrm{kJmol}^{-1}$ |
| :---: | :---: | :---: |
| 23 | (a) | Cow dung, water, plant remaining |
|  | (b) | Formula mass of $\mathrm{CH}_{4}=12+4=16$ 890 kJ areproduced by 16 g of methane 2670 kJ are produced by $\frac{16 \times 2670}{890}=48 \mathrm{~g}$ |
| 24 | (a) | $\mathrm{H}^{+}(\mathrm{aq})+\mathrm{OH}^{-}(\mathrm{aq}) \longrightarrow \mathrm{H}_{2} \mathrm{O}(\mathrm{l})$ |
|  | (b) | total volume of water $=50+50=100 \mathrm{~cm}^{3}$ <br> Mass of water $=100 \times 1=100 \mathrm{~g}$ <br> Temperature change $=35-22=13^{\circ} \mathrm{C}$ <br> Heat liberated $=m c \theta=100 \times 4.2 \times 13=5460 \mathrm{~J}$ <br> Mole of water formed $=$ moles $\mathrm{NaOH}=\frac{50 \times 2}{1000}=0.1$ mole <br> Production of 0.1 mole of water produce 5460 J <br> 1 mole of water produced $\frac{5460 \times 1}{0.1} 54600 \mathrm{~J}=54.6 \mathrm{kJmol}^{-1}$ |
| (b) |  | It would be less because ammonia is a weak base |


| 25 | (a) | $\mathrm{CH}_{4}(\mathrm{~g})+2 \mathrm{O}_{2}(\mathrm{~g}) \longrightarrow \mathrm{CO}_{2}(\mathrm{~g})+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{I})$ |
| :---: | :---: | :---: |
|  | (b) | Formula mass of $\mathrm{CH}_{4}=12+4=16$ <br> 882 kJ are produce by 24 I <br> 1746kJ require $\frac{1746 \times 24}{882}=47.5 l$ <br> Cost of 47.5I of methane $=47.5 \times 600=28500 /=$ |
| 26 |  | Ethnalpy of combustion is entalpy change when 1 mole of a substance is completely burnt in oxygen. |
|  |  | Experimental method for finding enthalpy of combustion a liquid fuel <br> The figure below shows a simple method for obtaining approximate value for the enthalpy of combustion of a propanol. <br> Thermometer record rise in temperature, $\theta^{0} \mathrm{C}$ <br> Heat shield reduces heat loss <br> Metal calorimeter and water both of known heat capacity $\mathrm{C}, \mathrm{Jk}$-1 <br> Spirit burner contains propanol, weight before and after gives mass of fuel burnt $=m_{1}$ g <br> Calculations <br> Assumption <br> Heat produced by = Heat gained by <br> combusting fuel - calorimeter and water <br> Heat gained by calorimeter and water $=\mathrm{C} \Theta$ joules <br> It implies that <br> $\mathrm{m}_{1} \mathrm{~g}$ of fuel produce $=\mathrm{C}$ joules <br> Mr (mass equivalent to 1 <br> mole of fuel) produces $=\frac{M r C \theta}{m_{1}}$ joulesmol $^{-1}$ |


|  | (b) | Mass of water $=150 \times 1=150 \mathrm{~g}$ <br> Heat $=\mathrm{mc} \theta=150 \times 4.2 \times 21.5=13545 \mathrm{~J}$ <br> 0.54 gof propanol produce 13545 J <br> 60 g of propanol produce $\frac{13545 \times 60}{0.54}=1505 \mathrm{~kJ}$ |
| :--- | :--- | :--- |
| 27 | (a)Formula mass of ethane $\mathrm{C}_{2} \mathrm{H}_{6}=2 \times 12+6 \times 1=30$ <br> Formula mass of water, $\mathrm{H}_{2} \mathrm{O}=1 \times 2+16=18$ <br> $(2 \times 30) \mathrm{g}$ of ethane produce $(6 \times 18) \mathrm{g}$ of water <br> 2 g of ethane produce $\frac{2 \times 6 \times 18}{2 \times 30}=3.6 \mathrm{~g}$ of water |  |
| (b)2 g of ethane produce 104 kJ <br> $30 \mathrm{~g}\left(1\right.$ mole) of ethane produce $\frac{30 \times 104}{2} 1560 \mathrm{~kJ}$ |  |  |
| 28 | (a)Enthalpy of combustion is the enthalpy change when 1 mole of a substance is burnt <br> completely in oxygen. |  |
| (b)Formula mass of ethanol $\mathrm{C}_{2} \mathrm{H}_{5} \mathrm{OH}=2 \times 12+6+16=46$ <br> Mass of water $=$ volume of water $=1000 \mathrm{~g}$ <br> Heat $=$ mc $\theta=1000 \times 10 \times 4.2=42000 \mathrm{~J}$ <br> 2 g of ethanol produce 42000 <br> 30 g of ethanol produce $\frac{420000 \times 30}{2} 630 \mathrm{~kJ}$ |  |  |

