VICTORIAN CURRICULUM AND ASSESSMENT AUTHORITY



Victorian Certificate of Education 2012

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STUDENT NUMBER

Figures Words



CHEMISTRY

Written examination 2

Tuesday 13 November 2012

Reading time: 9.00 am to 9.15 am (15 minutes) Writing time: 9.15 am to 10.45 am (1 hour 30 minutes)

QUESTION AND ANSWER BOOK

Structure of book

Section	Number of questions	Number of questions to be answered	Number of marks
A	20	20	20
B	9	9	52
			Total 72

- Students are permitted to bring into the examination room: pens, pencils, highlighters, erasers, sharpeners, rulers and one scientific calculator.
- Students are NOT permitted to bring into the examination room: blank sheets of paper and/or white out liquid/tape.

Materials supplied

- Question and answer book of 25 pages.
- A data book.
- Answer sheet for multiple-choice questions.

Instructions

- Write your **student number** in the space provided above on this page.
- Check that your **name** and **student number** as printed on your answer sheet for multiple-choice questions are correct, **and** sign your name in the space provided to verify this.
- All written responses must be in English.

At the end of the examination

- Place the answer sheet for multiple-choice questions inside the front cover of this book.
- You may keep the data book.

Students are NOT permitted to bring mobile phones and/or any other unauthorised electronic devices into the examination room.

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SECTION A – Multiple-choice questions

Instructions for Section A

Answer **all** questions in pencil on the answer sheet provided for multiple-choice questions.

Choose the response that is **correct** or that **best answers** the question.

A correct answer scores 1, an incorrect answer scores 0.

Marks will not be deducted for incorrect answers.

No marks will be given if more than one answer is completed for any question.

Question 1

A solvent has the following risk statement printed on its label.

'Inhalation of fumes may cause dizziness.'

To minimise the risk associated with the effects of exposure when using this solvent, a student should

- A. use gloves.
- **B.** wear a laboratory coat.
- C. keep the solvent away from flames.
- **D.** use the solvent in a well-ventilated area.

Question 2

Which one of the following fuels is the most sustainable?

- A. biodiesel
- **B.** uranium
- C. brown coal
- **D.** natural gas

SECTION A - continued

The diagram below represents the distribution of the kinetic energy of reactant particles at two different temperatures. Assume that the areas under the curves are equal.



From this diagram it can be concluded that

A. at T_1 a greater number of particles have sufficient energy to react. T_1 is greater than T_2 .

B. at T_1 a greater number of particles have sufficient energy to react. T_2 is greater than T_1 .

C. at T_2 a greater number of particles have sufficient energy to react. T_1 is greater than T_2 .

D. at T_2 a greater number of particles have sufficient energy to react. T_2 is greater than T_1 .

Question 4

Enthalpy changes for the melting of iodine, I2, and for the sublimation of iodine are provided below.

 $I_2(s) \rightarrow I_2(l) \qquad \Delta H = +16 \text{ kJ mol}^{-1}$ $I_2(s) \rightarrow I_2(g) \qquad \Delta H = +62 \text{ kJ mol}^{-1}$

The enthalpy change for the vaporisation of iodine that is represented by the equation $I_2(l) \rightarrow I_2(g)$ is

A. −78 kJ mol⁻¹

B. −46 kJ mol^{−1}

C. $+46 \text{ kJ mol}^{-1}$

D. $+78 \text{ kJ mol}^{-1}$

SECTION A – continued TURN OVER

3

Nitrogen dioxide decomposes as follows.

 $2NO_2(g) \rightarrow N_2(g) + 2O_2(g)$ $\Delta H = -66 \text{ kJ mol}^{-1}$

The enthalpy change for the reaction represented by the equation $\frac{1}{2} N_2(g) + O_2(g) \rightarrow NO_2(g)$ is

A. −66 kJ mol^{−1}

- **B.** −33 kJ mol⁻¹
- **C.** +33 kJ mol⁻¹
- **D.** $+66 \text{ kJ mol}^{-1}$

Question 6

Pure water at 100 °C has a pH of 6.14.

This is because

- A. the self-ionisation of water is endothermic.
- **B.** pH measurements at this temperature are unreliable.
- C. pH measurements are affected by the bubbles of hydrogen gas that form in boiling water.
- **D.** the concentration of H_3O^+ ions is not equal to the concentration of OH^- ions at this temperature.

SECTION A - continued

Use the following information to answer Questions 7 and 8.

5

$$Fe^{3+}(aq) + SCN^{-}(aq) \rightleftharpoons Fe(SCN)^{2+}(aq)$$

yellow deep red

Question 7

The concentration profile below represents a change to the above equilibrium system at time t_1 .



Which one of the following would account for the changes in concentration at time t_1 ?

- A. the addition of SCN⁻
- **B.** the removal of $Fe(SCN)^{2+}$
- **C.** an increase in temperature
- **D.** a decrease in temperature

Question 8

Which one of the following best represents the changes in concentration when the equilibrium mixture is diluted at time t_2 ?



SECTION A – continued TURN OVER Use the following information to answer Questions 9–11.

The following reaction is used in some industries to produce hydrogen.

 $CO(g) + H_2O(g) \rightleftharpoons CO_2(g) + H_2(g) \qquad \Delta H = -41 \text{ kJ mol}^{-1}$

Question 9

Carbon monoxide, water vapour, carbon dioxide and hydrogen were pumped into a sealed container that was maintained at a constant temperature of 200 °C. After 30 seconds, the concentration of gases in the sealed container was found to be [CO] = 0.1 M, $[H_2O] = 0.1 \text{ M}$, $[H_2] = 2.0 \text{ M}$, $[CO_2] = 2.0 \text{ M}$.

The equilibrium constant at 200 °C for the above reaction is K = 210.

Which one of the following statements about the relative rates of the forward reaction and the reverse reaction at 30 seconds is true?

- A. The rate of the forward reaction is greater than the rate of the reverse reaction.
- **B.** The rate of the forward reaction is equal to the rate of the reverse reaction.
- C. The rate of the forward reaction is less than the rate of the reverse reaction.
- **D.** There is insufficient information to allow a statement to be made about the relative rates of the forward and reverse reactions.

Question 10

The reaction between carbon monoxide and water vapour is carried out in a sealed container.

The equilibrium yield of hydrogen will be increased by

- A. an increase in pressure at constant temperature.
- **B.** a decrease in temperature.
- C. the addition of an inert gas at constant temperature.
- **D.** the use of a suitable catalyst at constant temperature.

 $\textbf{SECTION}\, \textbf{A} - \textbf{continued}$

In trials, the reaction is carried out with and without a catalyst in the sealed container. All other conditions are unchanged. The change in hydrogen concentration with time between an uncatalysed and a catalysed reaction is represented by a graph.

Which graph is correct?



Question 12

Consider the following energy profile diagram for a reaction represented by the equation $X + Y \rightarrow Z$.



Which one of the following provides the correct values of the activation energy and enthalpy for the reaction $X + Y \rightarrow Z$?

	Activation energy (kJ mol ⁻¹)	Enthalpy (kJ mol ⁻¹)
A.	+75	+100
B.	+100	+175
C.	+175	+100
D.	+200	-125

SECTION A – continued TURN OVER

1.30 g of glucose (M = 180 g mol⁻¹) underwent complete combustion. The energy released was used to heat an unknown mass of water.

If the temperature of the water increased by 24.3 °C and it is assumed no heat was lost, the mass of the water heated was

A. 2.00×10^{-1} g

- **B.** 1.02×10^2 g
- **C.** 2.00×10^2 g
- **D.** 3.84×10^3 g

Question 14

When 50 g of water at 90 °C is added to a calorimeter containing 50 g of water at 15 °C, the temperature increases to 45 °C.

Assuming no energy is lost to the environment, the energy absorbed by the calorimeter is equal to the

- **A.** energy lost by the hot water.
- **B.** energy gained by the cold water.
- C. sum of the energy gained by the cold water and the energy lost by the hot water.
- **D.** difference between the energy lost by the hot water and the energy gained by the cold water.

Question 15

If 54.0 kJ of energy is required to convert 1.00 mol of liquid water to steam at 100 °C, the amount of heat energy, in kilojoule, required to convert 100 g of water at 20 °C to steam at 100 °C is

- **A.** $3.34 \times 10^1 \text{ kJ}$
- **B.** $2.67 \times 10^2 \text{ kJ}$
- **C.** $3.00 \times 10^2 \text{ kJ}$
- **D.** $3.33 \times 10^2 \, \text{kJ}$

SECTION A – continued

Use the following information to answer Questions 16–18.

9

A galvanic cell set up under standard conditions is shown below.



Question 16

Which one of the following is correct?

As the cell discharges

	electrons would flow from the	in the salt bridge
A.	zinc electrode to the silver electrode.	anions migrate to the Ag ⁺ /Ag half-cell.
B.	silver electrode to the zinc electrode.	cations migrate to the Zn^{2+}/Zn half-cell.
C.	silver electrode to the zinc electrode.	cations migrate to the Ag ⁺ /Ag half-cell.
D.	zinc electrode to the silver electrode.	anions migrate to the Zn^{2+}/Zn half-cell.

Question 17

In this cell

NO WRITING ALLOWED IN THIS AREA

- **A.** $Ag^+(aq)$ is reduced and the Zn(s) is oxidised.
- **B.** Ag(s) is oxidised and the $Zn^{2+}(aq)$ is reduced.
- C. Ag(s) is reduced and the $Zn^{2+}(aq)$ is oxidised.
- **D.** $Ag^+(aq)$ is oxidised and the Zn(s) is reduced.

Question 18

The cathode in this cell and the maximum voltage produced by this cell, under standard conditions, are respectively

- **A.** Ag and 0.16 V
- **B.** Ag and 1.56 V
- **C.** Zn and 0.16 V
- **D.** Zn and 1.56 V

SECTION A – continued TURN OVER

Which one of the following statements is true for both galvanic cells and electrolytic cells?

- A. Reduction occurs at the negative electrode in both cells.
- **B.** Reduction occurs at the cathode in both cells.
- C. Anions migrate to the cathode in both cells.
- **D.** The anode is positive in both cells.

Question 20

Fuel cells have a number of applications that offer advantages over conventional methods of electricity generation. Which one of the following is **not** a feature of modern fuel cells?

- A. They generate very little noise.
- **B.** They are a cheap source of electricity.
- **C.** They enable electricity to be generated on site.
- **D.** They have the potential to reduce emissions of carbon dioxide into the atmosphere.

END OF SECTION A

CONTINUES OVER PAGE

TURN OVER

SECTION B – Short answer questions

Instructions for Section B

Answer **all** questions in the spaces provided. Write using black or blue pen.

To obtain full marks for your responses you should

- give simplified answers with an appropriate number of significant figures to all numerical questions; unsimplified answers will not be given full marks.
- show all working in your answers to numerical questions. No marks will be given for an incorrect answer unless it is accompanied by details of the working.
- make sure chemical equations are balanced and that the formulas for individual substances include an
 indication of state; for example, H₂(g); NaCl(s)

Question 1

Two experiments were conducted to investigate various factors that affect the rate of reaction between calcium carbonate and dilute hydrochloric acid.

$$CaCO_3(s) + 2HCl(aq) \rightleftharpoons CO_2(g) + CaCl_2(aq) + H_2O(l)$$

The two experiments are summarised in the diagrams below.

experiment 1

experiment 2



a. How could the rate of this reaction be measured in these experiments?

1 mark

SECTION B – Question 1 – continued

12

c.

b.

	13 2012 CHEM EXA
i.	Identify the rate determining factor that is investigated in experiment 1.
i.	In experiment 2, will the rate of reaction be faster in beaker A or beaker B? Explain your selection in terms of collision theory.
Vhv	1 + 2 = 3 marks v is the following statement incorrect ?
Col	llision theory states that all collisions between reactant particles will result in a chemical reaction.'
	2 marks

SECTION B – continued TURN OVER

13

14

Question 2

The reaction between 2-bromo-2-methylpropane and hydroxide ions occurs in two steps.

step 1 $(CH_3)_3CBr(aq) \rightarrow (CH_3)_3C^+(aq) + Br^-(aq)$ step 2 $(CH_3)_3C^+(aq) + OH^-(aq) \rightarrow (CH_3)_3COH(aq)$

a. Write an equation that represents the overall reaction between 2-bromo-2-methylpropane and hydroxide ions.

The energy profile diagrams for step 1 and step 2 are shown below. Both are drawn to the same scale.



b. i. Which step involves an endothermic reaction? Provide a reason for your answer.

The reaction at step 1 occurs at a different rate to the reaction at step 2.

ii. Which step is slower? Justify your answer.

1 + 2 = 3 marks

SECTION B - continued

1 mark

CONTINUES OVER PAGE

SECTION B – continued TURN OVER

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Question 3

The following weak acids are used in the food industry.

Acid	Common use	Formula	Structure	K _a values
sorbic	preservative	C ₆ H ₈ O ₂	$ \begin{array}{c c} H & H & O \\ I & I & \parallel \\ H_{3}C & C & C & C \\ H_{3}C & H & H \\ H & H & H \end{array} $	1.73 × 10 ⁻⁵
malic	low-calorie fruit drinks	C ₄ H ₆ O ₅	$HO \xrightarrow{C} C \xrightarrow{C} C \xrightarrow{C} OH$ $HO \xrightarrow{C} OH$ $HO \xrightarrow{C} OH$ $HO \xrightarrow{C} OH$	3.98 × 10 ⁻⁴ 8.91 × 10 ⁻⁶

a. What does the term 'weak acid' mean?

b. i. Why are two K_a values listed for malic acid?

The equation related to the first K_a value of malic acid is

 $\mathrm{HOOCCH_2CH(OH)COOH(aq)} + \mathrm{H_2O(l)} \iff \mathrm{HOOCCH_2CH(OH)COO^{-}(aq)} + \mathrm{H_3O^{+}(aq)}$

ii. Write an appropriate chemical equation that relates to the second K_a of malic acid.

1 + 1 = 2 marks

1 mark

SECTION B – Question 3 – continued

	17	2012 CHEM EXAM
c.	Sorbic acid, $CH_3(CH)_4COOH$, has antimicrobial properties that are used to inhibit year However, its solubility is very low. The more soluble potassium sorbate is used instead activity of sorbic acid is retained because an equilibrium exists according to the equat	ist and mould growth. d. The antimicrobial ion
	$CH_3(CH)_4COO^-(aq) + H_2O(l) \rightleftharpoons CH_3(CH)_4COOH(aq) + OH^-(aq)$	
	sorbate ion sorbic acid	
	How would the addition of a small amount of 1.0 M hydrochloric acid affect the concession solution? Justify your answer in terms of equilibrium principles.	entration of sorbic acid in
		2 marks
d.	Calculate the percentage dissociation of sorbic acid when the $pH = 4.76$.	
		3 marks

SECTION B – continued **TURN OVER**

In an experiment, 1.0 mol of pure phosgene, COCl_2 , is placed in a 3.0 L flask where the following reaction takes place.

18

 $\text{COCl}_2(g) \rightleftharpoons \text{CO}(g) + \text{Cl}_2(g) \qquad \text{K} = 2.1 \times 10^{-8} \text{ M}$

a. It can be assumed that, at equilibrium, the amount of unreacted COCl_2 is approximately equal to 1.0 mol. On the basis of the data provided, explain why this assumption is justified.

2 marks

b. i. Calculate the equilibrium concentration, in mol L⁻¹, of carbon monoxide, CO. Assume that the amount of unreacted COCl₂ is approximately equal to 1.0 mol.

ii. What is the equilibrium concentration of chlorine gas?

3 + 1 = 4 marks

SECTION B - continued

Q Ci THIS AREA a. Str b. Z **NO WRITING ALLOWED**

			19		2012 CHEM EXA
restion	5				
rcle the	e industrial chemical th	nat you have st	udied in detail thi	s semester.	
	ammonia	ethene	nitric acid	sulfuric acid	
<u> </u>					
Stat	e one application of yo	our selected che	emical that is used	ul to society.	
					1 mark
rict env	vironmental guidelines	are attached to	the industrial pro	oduction of your selected chem	ical.
i.	State one undesirable	e effect that the	production of yo	ur selected chemical has on the	environment.
ii.	Outline one procedur	e that would be	e appropriate to a	void this damage to the enviror	ment during the
	production of your se	elected chemica	al.	-	-
					1 + 1 = 2 marks

SECTION B – continued **TURN OVER** 20

Question 6

Methanol, CH₃OH, undergoes combustion according to the equation

 $2CH_3OH(1) + 3O_2(g) \rightarrow 2CO_2(g) + 4H_2O(g)$

In an experiment to determine its suitability as a fuel, a sample of methanol underwent complete oxidation in a bomb calorimeter.

The calorimeter was first calibrated by passing a current through an electric heater placed in the water surrounding the reaction vessel. A potential of 5.25 volts was applied for 3.00 minutes. The measured current was 1.50 amperes and the temperature of the water and reaction vessel increased by 0.593 °C.

a. i. Determine the calibration constant, in kJ $^{\circ}$ C⁻¹, for the calorimeter and its contents.

A student then used this calorimeter to determine the molar heat of combustion of methanol.

0.934 g of methanol was placed in the reaction vessel and excess oxygen was added. An electric ignition heater provided the energy required to initiate the combustion reaction. On this occasion, the temperature of the water increased by 8.63 °C.

ii. Use this experimental data to determine the value of ΔH for the combustion of methanol given by the following equation.

 $2CH_3OH(l) + 3O_2(g) \rightarrow 2CO_2(g) + 4H_2O(g)$

Include appropriate units in your answer.

2 + 5 = 7 marks

SECTION B – Question 6 – continued

b. The value of ∠H, calculated using the enthalpy of combustion provided in the data book, is different from the value of ∠H calculated from the experimental data provided in part a.ii.
 Provide a reason for this difference.

1 mark

Methanol is suitable for use in a micro fuel cell that is used to power laptop computers and similar small electrical items. The methanol is oxidised to carbon dioxide and water. The half-equation for the anode reaction is

 $CH_3OH(l) + H_2O(l) \rightarrow 6H^+(aq) + CO_2(g) + 6e^-$

c. i. Write a balanced half-equation for the cathode reaction.

ii. A finely divided platinum/ruthenium catalyst is used in this cell.

Give a reason why it is important to have a catalyst that will significantly reduce the activation energy for the cell reaction.

1 + 1 = 2 marks

SECTION B – continued TURN OVER

21

Hydrogen gas is an energy source. Researchers are investigating the production of hydrogen gas in a microbial electrolysis cell.

The cell is made up of an anode half-cell and a cathode half-cell. The half-cells are separated by a proton exchange membrane, as shown in the diagram below.

22



A number of reactions take place in the cell, resulting in the production of hydrogen. These reactions are summarised below.

Anode half-cell

- The anode half-cell contains moist plant matter and electrochemically active bacteria that live on an inert anode.
- The gaseous mixture that is present in the half-cell does not contain oxygen.
- The moist plant matter ferments to produce ethanoic acid (CH₃COOH). Bacteria on the anode consume the ethanoic acid and release hydrogen ions, electrons and carbon dioxide gas. A small voltage is then applied to reduce the H⁺ ions.

Cathode half-cell

- The cathode half-cell contains an inert cathode.
- The gaseous mixture that is present in the half-cell does not contain oxygen.
- The released hydrogen ions and electrons react to form hydrogen gas, as shown in the equation below.

 $2\mathrm{H}^{+}(\mathrm{aq}) + 2\mathrm{e}^{-} \rightarrow \mathrm{H}_{2}(\mathrm{g})$

Ethanoic acid is converted to carbon dioxide gas and H⁺ ions at the anode.
 Write an equation for this reaction.

b. On the diagram above, use one arrow to indicate the direction of electron flow in the cell when an external voltage is supplied to the cell by the power supply.

1 mark

SECTION B - Question 7 - continued

d.

c.	Hydrogen gas is not produced at the cathode if oxygen is present in the half-cell.
	Write a balanced half-equation to show the product that would be produced at the cathode if oxygen were
	present in the half-cell.

Describe one difference between an electrolysis cell and a traditional fuel cell.

1 mark

1 mark

SECTION B – continued TURN OVER

23

Decisions about clean energy with reduced carbon dioxide emissions will have an impact on electricity generation from brown coal. However, there will be a much smaller impact on the use of black coal for electricity generation. The following table compares the energy and carbon content of three different coal samples.

	Percentage carbon* by mass	Energy content (kJ g ⁻¹)
Black coal	93	36.0
Brown coal (dried)	66	28.0
Brown coal (wet – as mined)	40	5.0

*Coal is not a pure substance and the composition of samples will vary even within one mine.

From the data in this table, it can be deduced that the complete combustion of 1 tonne of black coal will generate 3.6×10^7 kJ of energy.

a. i. Calculate the mass, in tonne, of wet brown coal that is required to generate 3.6×10^7 kJ of energy.

ii. Calculate the mass, in tonne, of carbon dioxide that is produced from the complete combustion of this mass of wet brown coal.

1 + 2 = 3 marks

b. What are the most likely reasons for the energy content of wet brown coal being so much lower than the energy content of dried brown coal? Justify your answer.

2 marks

SECTION B - continued

A teacher demonstrated the process of electrolysis of a molten salt using an unknown metal salt, XBr_2 . The apparatus was set up as shown below.



At the conclusion of the demonstration, the students were provided with the following information.

- A current of 1.50 amperes was applied for 30.0 minutes.
- 2.90 g of metal X was produced.
- **a.** Write a balanced half-equation for the anode reaction in this electrolytic cell.

1 mark

b. i. Determine the amount, in mol, of metal X that was deposited on the cathode.

ii. Identify metal X.

3 + 2 = 5 marks

END OF QUESTION AND ANSWER BOOK

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Victorian Certificate of Education 2012

CHEMISTRY Written examination

Tuesday 13 November 2012

Reading time: 9.00 am to 9.15 am (15 minutes) Writing time: 9.15 am to 10.45 am (1 hour 30 minutes)

DATA BOOK

Directions to students

• A question and answer book is provided with this data book.

Students are NOT permitted to bring mobile phones and/or any other unauthorised electronic devices into the examination room.

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2 He 4.0 Helium 10 Neon Neon 18 39.9 Ar Argon	36 Kr 83.8 Krypton	54 Xe 131.3 Xenon	86 Rn (222) Radon	118 Uuo (294)	
9 F F I 19.0 Fluorine 17 CI 35.5 CI	35 Br 79.9 Bromine	53 1 126.9 Iodine	85 At (210) Astatine	117 Uus (294)	
8 0 0 0 0 0 0 0 0 xygen 16 0 32.1 Sulfur Sulfur	34 Se Selenium	52 Te 127.6 Tellurium	84 Po (210) Polonium	116 Uuh (293)	71 Lu 175.0 Lutetium
7 N 14.0 Nitrogen 15 7 31.0 Phosphorus	33 As 74.9 Arsenic	51 Sb 121.8 Antimony	83 Bi 209.0 Bismuth	115 Uup (288)	70 Yb 173.1 Ytterbium
6 C Carbon Carbon 14 Si Silicon	32 Ge 72.6 Germanium	50 Sn 118.7 Tin	82 Pb 207.2 Lead	114 Uuq (289)	69 Tm 168.9 Thulium
5 B Boron Boron 13 Al 27.0 Aluminium	31 Ga 69.7 Gallium	49 In 114.8 Indium	81 T1 204.4 Thallium	113 Uut (284)	68 Er 167.3 Erbium
	30 Zn 65.4 Zinc	48 Cd 112.4 Cadmium	80 Hg 200.6 Mercury	112 Cn (285) Copernicium	67 H0 164.9 Holmium
ol of element of element	29 Cu 63.5 Copper	47 Ag 107.9 Silver	79 Au 197.0 Gold	111 Rg (272) Roentgenium	66 Dy 162.5 Dysprosium
79 symb 70.0 77.0 old name	28 Ni 58.7 Nickel	46 Pd 106.4 Palladium	78 Pt 195.1 Platinum	110 Ds (271) Darmstadtium	65 Tb 158.9 Terbium
ic mass 19	27 Co 58.9 Cobalt	45 Rh 102.9 Rhodium	77 Ir 192.2 Iridium	109 Mt (268) Meitnerium	64 Gd 157.3 Gadolinium
atomic 1 ielative atomi	26 Fe 55.8 Iron	44 Ru 101.1 Ruthenium	76 Os 190.2 Osmium	108 Hs (267) Hassium	63 Eu 152.0 Europium
_	25 Mn 54.9 Manganese	43 Tc (98) Technetium	75 Re 186.2 Rhenium	107 Bh (264) Bohrium	62 Sm 150.4 Samarium
	24 Cr 52.0 Chromium	42 Mo 96.0 Molybdenum	74 W 183.8 Tungsten	106 Sg (266) Seaborgium	61 Pm (145) Promethium
	23 V 50.9 Vanadium	41 Nb 92.9 Niobium	73 Ta 180.9 Tantalum	105 Db (262) Dubnium	60 Nd 144.2 Neodymium
	22 Ti 47.9 Titanium	40 Zr 91.2 Zirconium	72 Hf 178.5 Hafhium	104 Rf (261) Rutherfordium	59 Pr 140.9 Praseodymium
	21 Sc 45.0 Scandium	39 Y 88.9 Yttrium	57 La 138.9 Lanthanum	89 Ac (227) Actinium	58 Ce 140.1 Cerium
4 4 Be 9.0 Beryllium 12 Mg 24.3 Magnesium	20 Ca 40.1 Calcium	38 Sr 87.6 Strontium	56 Ba 137.3 Barium	88 Ra (226) Radium	
1 1 H H Hydrogen 3 J 1.0 J 1.0 J 1.0 J 1.0 J 1.1 J 1.1 J 1.1 J 1.1 J 1.1 Na 23.0 Z3.0 Sodium	19 K 39.1 Potassium	37 Rb 85.5 Rubidium	55 Cs 132.9 Caesium	87 Fr (223) Francium	www.thea

1. Periodic table of the elements

The value in brackets indicates the mass number of the longest-lived isotope.

	103	Lr	(262)	Lawrencium
	102	No	(259)	Nobelium
	101	Md	(258)	Mendelevium
	100	\mathbf{Fm}	(257)	Fermium
	66	Es	(252)	Einsteinium
	98	Cf	(251)	Californium
	97	Bk	(247)	Berkelium
	96	Cm	(247)	Curium
_	95	Am	(243)	Americium
	94	Pu	(244)	Plutonium
	93	Np	(237)	Neptunium
	92	n	238.0	Uranium
	91	Pa	231.0	Protactinium
	90	$\mathbf{T}\mathbf{h}$	232.0	Thorium

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2. The electrochemical series

	E° in volt
$F_2(g) + 2e^- \rightleftharpoons 2F^-(aq)$	+2.87
$H_2O_2(aq) + 2H^+(aq) + 2e^- \rightleftharpoons 2H_2O(l)$	+1.77
$\operatorname{Au}^+(\operatorname{aq}) + \operatorname{e}^- \rightleftharpoons \operatorname{Au}(s)$	+1.68
$Cl_2(g) + 2e^- \rightleftharpoons 2Cl^-(aq)$	+1.36
$O_2(g) + 4H^+(aq) + 4e^- \rightleftharpoons 2H_2O(1)$	+1.23
$Br_2(l) + 2e^- \Longrightarrow 2Br^-(aq)$	+1.09
$Ag^{+}(aq) + e^{-} \rightleftharpoons Ag(s)$	+0.80
$Fe^{3+}(aq) + e^{-} \rightleftharpoons Fe^{2+}(aq)$	+0.77
$O_2(g) + 2H^+(aq) + 2e^- \rightleftharpoons H_2O_2(aq)$	+0.68
$I_2(s) + 2e^- \rightleftharpoons 2I^-(aq)$	+0.54
$O_2(g) + 2H_2O(l) + 4e^- \rightleftharpoons 4OH^-(aq)$	+0.40
$Cu^{2+}(aq) + 2e^{-} \rightleftharpoons Cu(s)$	+0.34
$\operatorname{Sn}^{4+}(\operatorname{aq}) + 2e^{-} \rightleftharpoons \operatorname{Sn}^{2+}(\operatorname{aq})$	+0.15
$S(s) + 2H^+(aq) + 2e^- \Longrightarrow H_2S(g)$	+0.14
$2\mathrm{H}^{+}(\mathrm{aq}) + 2\mathrm{e}^{-} \rightleftharpoons \mathrm{H}_{2}(\mathrm{g})$	0.00
$Pb^{2+}(aq) + 2e^{-} \rightleftharpoons Pb(s)$	-0.13
$\operatorname{Sn}^{2+}(\operatorname{aq}) + 2e^{-} \rightleftharpoons \operatorname{Sn}(s)$	-0.14
$Ni^{2+}(aq) + 2e^{-} \rightleftharpoons Ni(s)$	-0.23
$\operatorname{Co}^{2+}(\operatorname{aq}) + 2e^{-} \rightleftharpoons \operatorname{Co}(s)$	-0.28
$Fe^{2+}(aq) + 2e^{-} \rightleftharpoons Fe(s)$	-0.44
$Zn^{2+}(aq) + 2e^{-} \rightleftharpoons Zn(s)$	-0.76
$2H_2O(l) + 2e^- \Longrightarrow H_2(g) + 2OH^-(aq)$	-0.83
$Mn^{2+}(aq) + 2e^{-} \Longrightarrow Mn(s)$	-1.03
$Al^{3+}(aq) + 3e^{-} \rightleftharpoons Al(s)$	-1.67
$Mg^{2+}(aq) + 2e^{-} \Longrightarrow Mg(s)$	-2.34
$Na^+(aq) + e^- \rightleftharpoons Na(s)$	-2.71
$Ca^{2+}(aq) + 2e^{-} \rightleftharpoons Ca(s)$	-2.87
$K^+(aq) + e^- \rightleftharpoons K(s)$	-2.93
$Li^+(aq) + e^- \rightleftharpoons Li(s)$	-3.02

3. Physical constants

Avogadro's constant $(N_A) = 6.02 \times 10^{23} \text{ mol}^{-1}$ Charge on one electron $= -1.60 \times 10^{-19} \text{ C}$ Faraday constant $(F) = 96500 \text{ C mol}^{-1}$ Gas constant $(R) = 8.31 \text{ J K}^{-1}\text{mol}^{-1}$ Ionic product for water $(K_w) = 1.00 \times 10^{-14} \text{ mol}^2 \text{ L}^{-2}$ at 298 K (Self ionisation constant) Molar volume (V_m) of an ideal gas at 273 K, 101.3 kPa (STP) $= 22.4 \text{ L mol}^{-1}$ Molar volume (V_m) of an ideal gas at 298 K, 101.3 kPa (SLC) $= 24.5 \text{ L mol}^{-1}$ Specific heat capacity (c) of water $= 4.18 \text{ J g}^{-1} \text{ K}^{-1}$ Density (d) of water at 25 °C $= 1.00 \text{ g mL}^{-1}$ 1 atm = 101.3 kPa = 760 mm Hg $0 ^{\circ}\text{C} = 273 \text{ K}$

4. SI prefixes, their symbols and values

SI prefix	Symbol	Value
giga	G	109
mega	М	106
kilo	k	10 ³
deci	d	10^{-1}
centi	с	10 ⁻²
milli	m	10 ⁻³
micro	μ	10-6
nano	n	10-9
pico	р	10^{-12}

5. ¹H NMR data

Typical proton shift values relative to TMS = 0

These can differ slightly in different solvents. Where more than one proton environment is shown in the formula, the shift refers to the ones in **bold** letters.

Type of proton	Chemical shift (ppm)
R–CH ₃	0.8–1.0
R-CH ₂ -R	1.2–1.4
$RCH = CH - CH_3$	1.6–1.9
R ₃ –CH	1.4–1.7
$CH_3 - C$ or $CH_3 - C$ O OR NHR	2.0

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Type of proton	Chemical shift (ppm)
R CH ₃	
	2.1–2.7
$R-CH_2-X$ (X = F, Cl, Br or I)	3.0-4.5
R–С H₂– ОН, R ₂ –С H –ОН	3.3–4.5
//0	
R—Ć	3.2
NHC H ₂ R	
R —O— CH_3 or R —O— CH_2R	3.3
О Ш —О—С—СН3	2.3
R—CO OCH ₂ R	4.1
R–О–Н	1–6 (varies considerably under different conditions)
R-NH ₂	1–5
$RHC = CH_2$	4.6-6.0
ОН	7.0
Н	7.3
R—C NHCH ₂ R	8.1
R—C H	9–10
R—CO O—H	9–13

6. ¹³C NMR data

Type of carbon	Chemical shift (ppm)
R–CH ₃	8–25
R-CH ₂ -R	20–45
R ₃ -CH	40–60
R ₄ -C	36–45
R-CH ₂ -X	15-80
R ₃ C–NH ₂	35–70
R-CH ₂ -OH	50–90
RC≡CR	75–95
$R_2C=CR_2$	110–150
RCOOH	160–185

7. Infrared absorption data

Characteristic range for infrared absorption

Bond	Wave number (cm ⁻¹)
C–Cl	700–800
C–C	750–1100
С-О	1000-1300
C=C	1610–1680
C=O	1670–1750
O-H (acids)	2500-3300
С–Н	2850-3300
O-H (alcohols)	3200-3550
N-H (primary amines)	3350-3500

8. 2-amino acids (α-amino acids)





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9.	Formulas	of some	e fatty	acids
- •		01 00111		

Name	Formula
Lauric	C ₁₁ H ₂₃ COOH
Myristic	C ₁₃ H ₂₇ COOH
Palmitic	C ₁₅ H ₃₁ COOH
Palmitoleic	C ₁₅ H ₂₉ COOH
Stearic	C ₁₇ H ₃₅ COOH
Oleic	C ₁₇ H ₃₃ COOH
Linoleic	C ₁₇ H ₃₁ COOH
Linolenic	C ₁₇ H ₂₉ COOH
Arachidic	C ₁₉ H ₃₉ COOH
Arachidonic	C ₁₉ H ₃₁ COOH

10. Structural formulas of some important biomolecules







deoxyribose



11. Acid-base indicators

Name	pH range	Colour change		K _a
		Acid	Base	
Thymol blue	1.2–2.8	red	yellow	2×10^{-2}
Methyl orange	3.1–4.4	red	yellow	2×10^{-4}
Bromophenol blue	3.0-4.6	yellow	blue	6×10^{-5}
Methyl red	4.2-6.3	red	yellow	8 × 10 ⁻⁶
Bromothymol blue	6.0–7.6	yellow	blue	1×10^{-7}
Phenol red	6.8-8.4	yellow	red	1×10^{-8}
Phenolphthalein	8.3-10.0	colourless	red	5×10^{-10}

12. Acidity constants, K_a , of some weak acids at 25 °C

Name	Formula	K _a
Ammonium ion	NH4 ⁺	5.6×10^{-10}
Benzoic	C ₆ H ₅ COOH	$6.4 imes 10^{-5}$
Boric	H ₃ BO ₃	$5.8 imes10^{-10}$
Ethanoic	СН ₃ СООН	1.7×10^{-5}
Hydrocyanic	HCN	$6.3 imes 10^{-10}$
Hydrofluoric	HF	$7.6 imes 10^{-4}$
Hypobromous	HOBr	2.4×10^{-9}
Hypochlorous	HOCI	$2.9 imes10^{-8}$
Lactic	HC ₃ H ₅ O ₃	$1.4 imes10^{-4}$
Methanoic	НСООН	$1.8 imes10^{-4}$
Nitrous	HNO ₂	7.2×10^{-4}
Propanoic	C ₂ H ₅ COOH	1.3×10^{-5}

13. Values of molar enthalpy of combustion of some common fuels at 298 K and 101.3 kPa

Substance	Formula	State	$\Delta H_{\rm c} (\rm kJ mol^{-1})$
hydrogen	H ₂	g	-286
carbon (graphite)	С	S	-394
methane	CH ₄	g	-889
ethane	C ₂ H ₆	g	-1557
propane	C ₃ H ₈	g	-2217
butane	C ₄ H ₁₀	g	-2874
pentane	C ₅ H ₁₂	1	-3509
hexane	C ₆ H ₁₄	1	-4158
octane	C ₈ H ₁₈	1	-5464
ethene	C ₂ H ₄	g	-1409
methanol	CH ₃ OH	1	-725
ethanol	C ₂ H ₅ OH	1	-1364
1-propanol	CH ₃ CH ₂ CH ₂ OH	1	-2016
2-propanol	CH ₃ CHOHCH ₃	1	-2003
glucose	C ₆ H ₁₂ O ₆	S	-2816

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