# MARKSCHEME 

November 2001

## CHEMISTRY

## Higher Level

## Paper 3

## OPTION C - HUMAN BIOCHEMISTRY

C1. (a) $\mathrm{CH}_{2} \mathrm{OH}-\mathrm{CHOH}-\mathrm{CH}_{2} \mathrm{OH}$
(b) -COOH

(Accept $\mathrm{CH}_{3}\left(\mathrm{CH}_{2}\right)_{6} \mathrm{COOH}$ or any correct alternative, including branched structures or alkenoic acids.)
(c) Molecules of saturated fats contain only single $\mathrm{C}-\mathrm{C}$ bonds in the carbon chains / contain no double bonds.
Molecules of unsaturated fats contain at least one $\mathrm{C}=\mathrm{C}$ double bond.
The degree of unsaturation can be found by determining the number of moles of iodine that react with one mole (or a stated mass) of fat.
Iodine adds across the $\mathrm{C}=\mathrm{C}$ double bond in a $1: 1$ stoichiometric ratio.
Total [7 marks]
C2. (a) $-\mathrm{NH}_{2} /$ amino group / amine.
(b) Peptide bond (accept amide bond)

(c) First hydrolyse the peptide bonds to release individual amino acids
then use chromatography (comparison of $\mathrm{R}_{\mathrm{f}}$ values) (accept electrophoresis / mass spectrometry).
(Award both marks if X-ray crystallography is given.)
(d) The secondary structure describes the type of coil or sheet / folding of polypeptide / $\alpha$-helix and $\beta$-pleated sheet.
Tertiary structure describes the interactions between the R groups of the amino acid residues.
(e) Hydrogen bond.
C3. (a) Enzymes provide an alternative pathway with a lower activation energy. ..... [1]
Enzymes posses an active site where the substrate binds to the enzyme. ..... [1]
Mention of 'lock and key' and/or induced fit hypothesis. ..... [1]
Competitive inhibitors (resemble the substrate in shape) compete with the ..... [1]
substrate for the active site (and therefore slow the rate of reaction).Non-competitive inhibitors bind to the enzyme away from the active site.[1]
(b) $K_{\mathrm{m}}$ for uninhibited enzyme $\left(=\frac{1}{2} V_{\max }\right)=7 \pm 1 \mu \mathrm{moldm}^{-3}$ ..... [1]
With a competitive inhibitor $K_{\mathrm{m}}$ is higher, $V_{\text {max }}$ is the same. ..... [1]At high [S] effect of inhibitor is negligible so no change to $V_{\max }$ but more substrateneeded to reach $\frac{1}{2} V_{\text {max }}$ so $K_{\mathrm{m}}$ is higher.[1]
With a non-competitive inhibitor $K_{\mathrm{m}}$ is the same but $V_{\text {max }}$ is lower. ..... [1]
Adding a non-competitive inhibitor has the same effect as lowering [enzyme](hence $V_{\text {max }}$ is lower), but does not affect the way the substrate interacts with thoseenzyme molecules that are uninhibited (hence $K_{\mathrm{m}}$ is unchanged).[1]

## OPTION D - ENVIRONMENTAL CHEMISTRY

D1. (a) (i) $2 \mathrm{CO}+2 \mathrm{NO} \rightarrow 2 \mathrm{CO}_{2}+\mathrm{N}_{2}$
(Award [1] for correct products and [1] for balanced equation.)
(ii) $\mathrm{CaO}+\mathrm{SO}_{2} \rightarrow \mathrm{CaSO}_{3}\left(\right.$ or $\left.2 \mathrm{CaO}+2 \mathrm{SO}_{2}+\mathrm{O}_{2} \rightarrow 2 \mathrm{CaSO}_{4}\right)$
(Award [1] for reactants and [1] for product.)
(b) $2 \mathrm{NO}+\mathrm{O}_{2} \rightarrow 2 \mathrm{NO}_{2}$ [1]
$4 \mathrm{NO}_{2}+2 \mathrm{H}_{2} \mathrm{O}+\mathrm{O}_{2} \rightarrow 4 \mathrm{HNO}_{3}[1]$
OR
$2 \mathrm{SO}_{2}+\mathrm{O}_{2} \rightleftharpoons 2 \mathrm{SO}_{3}[1]$
$\mathrm{SO}_{3}+\mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{H}_{2} \mathrm{SO}_{4}$ (accept $\mathrm{SO}_{2}+\mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{H}_{2} \mathrm{SO}_{3}$ ) [1]
(c) Irritation of the mucous membranes / fatigue / weakness / confusion (e.g. from exposure to $\mathrm{C}_{6} \mathrm{H}_{5} \mathrm{CH}_{3}$ ) / cancer forming / respiratory problems.

D2. (a) Polychlorinated biphenyls
(b) (i) $\mathrm{LD}_{50}$ - lethal dose in $50 \%$ of the population (the amount of poison that kills half the organisms in a randomly chosen batch of a named species). Maximum daily tolerance - the highest amount or concentration that has 'no effect' on any of the organisms in a specified batch.
(ii) $\mathrm{LD}_{50}$ :

Advantage: clearly measurable value.
Disadvantage: not done on human population so effect on humans 'unknown'.
OR
Maximum daily tolerance value:
Advantage: gives idea of 'safe amount'.
Disadvantage: same as for $\mathrm{LD}_{50}$ so has to be divided by 'safety factor'.
(c) (i) Batteries / seed dressings to prevent mould / chlor-alkali process
(do not accept "mercury fillings in teeth").
(ii) (Award [1] for any of the following:)
inflammation of the mouth / muscle spasms / nausea / diarrhoea / kidney damage / blindness / deafness / damage to CNS / personality change.
(iii) The $\mathrm{Hg}^{+}$ions interfere with the normal functioning of enzymes (e.g. by replacing the $\mathrm{Ca}^{2+}$ or $\mathrm{Mg}^{2+}$ ions).
D3. Water is a highly polar substance (due to shape and high electronegativity of oxygen). ..... [1] ..... [1]Capable of hydrogen bonding (that allows it to dissolve many chemicals).
Fresh water not available uniformly around the world / 'locked up' in glaciers and icebergs. ..... [1]Where the consumption is necessarily high it is easily contaminated with water bornediseases / by micro-organisms from human waste / from flooding / due to inadequatechemical treatment of water supplies.
Reverse osmosis:
Uses high pressure ..... [1]
to force water from salt-water through partially- (semi)-permeable membrane; ..... [1]
the partially-permeable membrane does not allow the passage of dissolved ions. ..... [1]
OR Osmosis is the net movement of water molecules from a region of highconcentration, i.e. pure water to one of lower concentration, i.e. less pure waterthrough a partially-permeable membrane / osmosis is the tendency to equaliseconcentrations.[1]Due to osmosis, pure water will move through a partially-permeable membrane intosalt water, thus diluting it.[1]If pressure greater than osmotic pressure is applied, the flow of solvent takes placein the opposite direction, called reverse osmosis.[1]
Ion exchange:
Requires the use of both a positive ion exchange (which can replace metal ions in sea water with $\mathrm{H}^{+}$ions) ..... [1]
and a negative ion exchange (which can replace anions with $\mathrm{OH}^{-}$ions). ..... [1]
The $\mathrm{H}^{+}$and $\mathrm{OH}^{-}$ions combine to form fresh/pure water. ..... [1]

## OPTION E - CHEMICAL INDUSTRIES

E1. (a) (i) $\mathrm{N}_{2}$ obtained from the fractional distillation of liquid air.
(Not enough to just state 'from air'.)
(ii) $\begin{aligned} & \mathrm{H}_{2} \text { obtained from cracking of petroleum products / from water using } \\ & \text { reduction with methane / from water using reduction with naptha / catalytic } \\ & \text { reforming / electrolysis of sodium chloride solution. }\end{aligned}$ [1]
(b) There are four volumes (moles) of gas on LHS and only two on RHS [1]
so increasing the pressure will move the position of equilibrium to the right [1]
Increasing the pressure increases the concentration of the gases [1]
So reaction rate increases. [1]
(c) The yield of ammonia is low
so most of the $\mathrm{N}_{2}$ and $\mathrm{H}_{2}$ needs to go round again (to save waste/cost).

E2. (a) The free radical electron $\left(\mathrm{HO}-\mathrm{CH}_{2}-\mathrm{CH}_{2}\right)$ attacks one of the C atoms in $\mathrm{H}_{2} \mathrm{C}=\mathrm{CH}_{2}$.
An electron from one of the bonds in the double bond joins this electron to form a single C-C bond.
The double bond is now a single bond with the electron on the end carbon.
(b) $\mathrm{HO}-\mathrm{CH}_{2}-\mathrm{CH}_{2}-\mathrm{CH}_{2}-\mathrm{CH}_{2}+\mathrm{H}_{2} \mathrm{C}=\mathrm{CH}_{2} \rightarrow \mathrm{HO}-\mathrm{CH}_{2}-\mathrm{CH}_{2}-\mathrm{CH}_{2}-\mathrm{CH}_{2}-\mathrm{CH}_{2}-\mathrm{CH}_{2}$
(c) High density is ionic.
(d) (Award [1] each for two conditions from the following:)

- Temperature about $60^{\circ} \mathrm{C}$;
- Pressure 2-6 atmospheres;
- Ziegler (Natta) catalyst/triethylaluminium with titanium(IV) chloride.
(e) $-\mathrm{O}-\stackrel{\text { O}}{\mathrm{C}}-\mathrm{O}-\mathrm{H} / \mathrm{O}-\stackrel{\text { O}}{\mathrm{C}}-\mathrm{Cl}$ and $\mathrm{H}_{2} \mathrm{~N}-$

OR


E3. Tabulated form of answer.
(For each, give [1] for the process, [1] for the reactants, [1] for the products, and [1] for explaining the conditions.)

Choose from the following:

| Temperature |  |
| :--- | :--- |
| Thermal cracking / coking | Thermal decomposition of very heavy fractions to give <br> mainly lower alkanes and alkenes and a high coke yield. |
| Steam cracking | Thermal cracking of $\mathbf{C}_{2}+$ hydrocarbons $\left(\mathbf{C}_{4}-\mathbf{C}_{12}\right)$ to <br> alkenes (ethene, 1,3-butadiene, etc.) in the presence of <br> steam. Temperature around $800{ }^{\circ} \mathrm{C}$. |
| Catalyst | Accelerated decomposition, with some aromatisation, of <br> middle/higher fractions over solid acidic <br> catalysts/zeolites. Gives lower alkanes, alkenes, <br> aromatics. |
| Catalytic cracking | Accelerated hydrogenolysis / decomposition of heavy <br> fractions to naptha over metal/acid catalysts e.g. <br> palladium on zeolite. |
| Hydrocracking | Metal/acid-catalysed e.g. platinum on acidic alumina <br> rearrangement / cyclisation and dehydrogenisation of <br> alkanes and aromatisation of napthas. |
| Catalytic reforming <br> (platforming etc.) | More general term, including methane to ethene - cracking <br> autothermal, with partial combustion. |
| Thermal / autothermal |  |

By choosing a suitable catalyst, the temperature required is lower which makes it more efficient (and less costly) / helps to give the desired products by dictating the mechanism.

## OPTION F - FUELS AND ENERGY

F1. (a) Anode: zinc [1]
Cathode: graphite (carbon) [1]
Electrolyte: ammonium chloride OR zinc chloride and ammonium chloride and water.
(b) Anode: $\quad \mathrm{Zn} \rightarrow \mathrm{Zn}^{2+}+2 \mathrm{e}^{-} \quad$ [1]

Cathode: $\quad 2 \mathrm{NH}_{4}^{+}+2 \mathrm{e}^{-} \rightarrow 2 \mathrm{NH}_{3}+\mathrm{H}_{2}$ [1]
OR $2 \mathrm{MnO}_{2}+2 \mathrm{NH}_{4}^{+}+2 \mathrm{e}^{-} \rightarrow \mathrm{Mn}_{2} \mathrm{O}_{3}+2 \mathrm{NH}_{3}+\mathrm{H}_{2} \mathrm{O}$
(State symbols are not required.)
(c) (Award [1] each for any two from the following:)

No decline in performance under high loads / no gases formed at cathode / longer shelf life / able to produce more current for a longer time / good for emergency lighting.
(d) Voltage does not change [1]
voltage depends primarily on materials used.

F2. (a) (i) ${ }_{0}^{1} \mathrm{n}+{ }_{92}^{235} \mathrm{U} \rightarrow{ }_{37}^{90} \mathrm{Rb}+{ }_{55}^{144} \mathrm{Cs}+2{ }_{0}^{1} \mathrm{n}$
(Give [1] for correct isotopes and [1] for two neutrons.)
(ii) $\begin{array}{rlrl}\mathrm{E} & =\mathrm{mc}^{2}=1 \times 0.001 \times\left(3.0 \times 10^{8}\right)^{2} & & \text { [1] } \\ & =9.0 \times 10^{13} \mathrm{~J}\left(9.0 \times 10^{10} \mathrm{~kJ}\right) & \text { [1] }\end{array}$
(b) Four half-lives [1]
so $t_{1 / 2}=\frac{57.2}{4}=14.3$ days

F3. (a) The crude oil is heated;
Fed into the base of a fractionating column;
Temperature gradient across column / lower boiling fractions distil first;
Fractions emerging in boiling range not pure compounds.
(Award [1] each for any three correct fractions from: (any two from number of C atoms, boiling range, and uses must be correct for each mark).)

| Fraction | Number of C atoms | Boiling range $/{ }^{\circ} \mathbf{C}$ | Uses |
| :--- | :---: | :---: | :---: |
| Gases | $1-4$ | $0-200$ | cooking, heating |
| Naptha | $5-7$ | $20-100$ | solvents |
| Gasoline (petrol) | $5-12$ | $40-175$ | motor car fuel |
| Kerosene | $12-18$ | $175-300$ | jets / diesel |
| Gas oil | $18-24$ | $300-400$ | diesel fuel |
| Lubricating oil | $20-30$ | non volatile | lubricants |
| Paraffin wax | $25-40$ | solid | candles |
| Bitumen / tar | $>30$ | solid | road surfaces |

(b) $\mathrm{C}_{12} \mathrm{H}_{26} \rightarrow \mathrm{C}_{8} \mathrm{H}_{18}+2 \mathrm{C}_{2} \mathrm{H}_{4}\left(\right.$ or $\mathrm{C}_{4} \mathrm{H}_{8}$ )
(must show an alkene for mark).
Octane for car engines and alkenes for polymers.

## OPTION G - MODERN ANALYTICAL CHEMISTRY

G1. (a) (i) $\begin{aligned} & \text { The positive molecular ion or } \mathrm{M}^{+} / \text {parent ion. } \\ & \text { The molecular formula of compound } \mathbf{A} \text { is } \mathrm{C}_{9} \mathrm{H}_{10} \mathrm{O} \\ & \text { (Accept "the relative molecular mass of compound } \boldsymbol{A} \text { is } 134 \text { ".) }\end{aligned}$
(ii) Due to the presence of ${ }^{13} \mathrm{C}$ isotopes in the compound. [1]
(iii) 105 due to $\left(\mathrm{M}-\mathrm{C}_{2} \mathrm{H}_{5}\right)^{+} /\left(\mathrm{C}_{7} \mathrm{H}_{5} \mathrm{O}^{+}\right) \quad$ [1]
(also accept $(\mathrm{M}-\mathrm{CHO})^{+} /\left(\mathrm{C}_{8} \mathrm{H}_{9}^{+}\right)$)
77 due to $\left(\mathrm{M}-\mathrm{C}_{3} \mathrm{H}_{5} \mathrm{O}\right)^{+} / \mathrm{C}_{6} \mathrm{H}_{5}^{+}$
(must have + sign to gain mark, but only penalize once if omitted)
(b) (i) It contains a carbonyl / either alkanal or alkanone / $\mathrm{C}=\mathrm{O}$ group. [1]
(Do not accept acid or ester.)
(ii) [1]
(iii) It does not contain an alkanol or phenol group / no - OH group present. [1]
(c) (i) Molecular formula: $\mathrm{SiC}_{4} \mathrm{H}_{12}\left(\right.$ accept $\left.\operatorname{Si}\left(\mathrm{CH}_{3}\right)_{4}\right)$ [1]
(Award [1] each for any two from the following:)

- Gives a sharp single peak;
- Strong peak as 12 protons in identical environment;
- Volatile so can be easily removed from the sample afterwards;
- Occurs well away from other peaks (which are shifted downfield from it).
(ii) They are in the ratio $3: 2: 5$ (since there are only 10 protons this is the actual
number).
$\begin{array}{ll}\text { (iii) It contains }\left(-\mathrm{C}_{2} \mathrm{H}_{5}\right) \text { group / a }-\mathrm{CH}_{3} \text { group next to a }-\mathrm{CH}_{2}-\text { group } & \text { [1] } \\ \text { The }-\mathrm{C}_{2} \mathrm{H}_{5}-\text { group is attached to a } \mathrm{C} \text { atom containing no other } \mathrm{H} \text { atoms } & \\ \text { bonded to it. } & \text { [1] }\end{array}$
(iv) It contains an aromatic (benzene) ring. [1]


## Question G1 continued

(d)

(Give [1] for correctly showing aromatic ring and [1] for $-\overbrace{C}^{\mathrm{O}}-\mathrm{C}_{2} \mathrm{H}_{5}$ group.)
G2. (a) The five d orbitals are split into different levels. ..... [1]
When an electron is excited from a lower to a higher level it absorbs light/energy. ..... [1]The wavelength of the transmitted light / complementary colour lies in the visibleregion of the spectrum.[1]
(b) A spectrum is run with the copper(II) sulfate solution to determine the wavelength of maximum absorption / $\lambda_{\text {max }}$. ..... [1]
The standard solution is diluted to give several solutions of different known concentrations. ..... [1]
The absorbance for each of these solutions at a fixed wavelength / $\lambda_{\text {max }}$ is recorded. ..... [1]
A calibration curve of absorption versus concentration is plotted. ..... [1]
The absorption of the unknown solution is recorded at the same wavelength and the concentration obtained from the graph. ..... [1]

## OPTION H - FURTHER ORGANIC CHEMISTRY

H1. (a) $\mathrm{C}_{6} \mathrm{H}_{6}+\mathrm{Br}_{2} \rightarrow \mathrm{C}_{6} \mathrm{H}_{5} \mathrm{Br}+\mathrm{HBr}$

(b) Addition of a halogen carrier / $\mathrm{AlBr}_{3} / \mathrm{Fe} / \mathrm{FeBr}_{3}$

Heat / reflux [1]
(c) Electrophile: $\mathrm{Br}^{+}$

$$
\mathrm{Br}-\mathrm{Br} \mathrm{AlBr}_{3} \rightarrow \mathrm{Br}^{+}+\mathrm{AlBr}_{4}^{-}
$$

(d)

(e) Phenol is more reactive than benzene.

No halogen carrier necessary for the reaction to proceed.
A pair of electrons on the oxygen atom delocalises with the $\pi$ electrons in the ring which increases the electron density on the ring (thus attracting electrophiles more readily).


H2. (a) The $\pi$ bond prevents the double bond from rotating in both cases.
In but-2-ene the atoms in the two isomers are arranged differently in space / contain two different groups attached to each carbon atom.
(b) (i)


(ii) They rotate the plane of polarised light in opposite directions.

Due to the presence of an asymmetric carbon atom / enantiomers / chirality.

H3. (a)


$$
\left(\text { accept } \mathrm{C}\left(\mathrm{CH}_{3}\right)_{3} \mathrm{Br}\right)
$$

(b) Electrophilic addition



OR primary carbocation

tertiary carbocation
Tertiary more stable
due to electron donating / positive inductive effect of the alkyl group.
OR R $\rightarrow \underset{\substack{\mathrm{C}}}{\stackrel{+}{\mathrm{C}}} \leftarrow \mathrm{R} \quad>\quad \mathrm{R} \rightarrow \underset{+}{\stackrel{+}{\mathrm{C}}-\mathrm{H}}$

