Surname	Centre Number	Candidate Number
Other Names		2



GCE A level

1094/01

CHEMISTRY CH4

P.M. WEDNESDAY, 15 June 2011 $1^{3}/_{4}$ hours

ADD	ITION	A T	N/I A	TEDI	T A T
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In addition to this examination paper, you will need:

- a calculator:
- an 8 page answer book;
- a Data Sheet which contains a Periodic Table supplied by WJEC. Refer to it for any relative atomic masses you require.

FOR EXAMINER'S USE ONLY					
Section	Section Question Mark				
	1				
A	2				
	3				
D	4				
В	5				
TOTAL MARK					

INSTRUCTIONS TO CANDIDATES

Use black ink or black ball-point pen.

Write your name, centre number and candidate number in the spaces at the top of this page.

Section A Answer all questions in the spaces provided.

Answer both questions in Section B in a separate answer book which should then Section B

be placed inside this question-and-answer book.

Candidates are advised to allocate their time appropriately between Section A (40 marks) and Section B (40 marks)

INFORMATION FOR CANDIDATES

The number of marks is given in brackets at the end of each question or part-question.

The maximum mark for this paper is 80.

Your answers must be relevant and must make full use of the information given to be awarded full marks for a question.

You are reminded that marking will take into account the Quality of Written Communication in all written answers.

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SECTION A

Answer all questions in the spaces provided.

1. Nitrobenzene, C₆H₅NO₂, is a yellow oily liquid. (a)

> Give the general name of a group responsible for colour in organic compounds. (i)

> > [1]

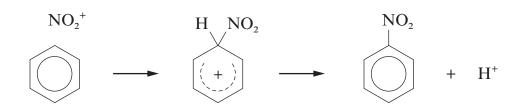
(ii) State why nitrobenzene is yellow in white light.

[1]

- Nitrobenzene is produced from benzene by reaction with the nitronium ion *(b)* (nitryl cation), NO_2^+ .
 - Complete the mechanism below by the use of the curly arrows (\frown)

[1]

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(ii) During this reaction to produce nitrobenzene small quantities of 1,3-dinitrobenzene are produced. [1]

Give the **empirical** formula of 1,3-dinitrobenzene.

In this reaction the nitronium ion is produced from nitric and sulfuric acids.

$$HNO_3 + 2H_2SO_4 \longrightarrow NO_2^+ + H_3O^+ + 2HSO_4^-$$

Use this equation to state why the sulfuric acid is acting as an acid. [1]

(c)	Explain why benzene compounds tend to react by electrophilic substitution	rather t	han
	undergo electrophilic addition.		[2]
		(OIII	Г.

(QWC) [1]

(d) Many explosives contain nitro-groups. The explosive hexanitrostilbene (HNS)

$$O_2N$$
 $\begin{array}{c|cccc}
NO_2 & H & O_2N \\
& & & \\
& & & \\
NO_2 & C = C \\
& & & \\
NO_2 & H & O_2N
\end{array}$
 $\begin{array}{c|cccc}
NO_2 & H & O_2N \\
& & & \\
& & & \\
NO_2 & H & O_2N \\
\end{array}$

hexanitrostilbene

has been used to separate different sections in space rockets and for seismic experiments on the Moon.

- (i) HNS is the E-isomer of a pair of E-Z isomers. State why HNS has both E- and Z-isomers. [1]
- (ii) The manufacture of HNS is believed to proceed via compound **R**.

$$O_{2}N \xrightarrow{NO_{2}} Cl H O_{2}N$$

$$C - C$$

$$NO_{2} H H O_{2}N$$

$$NO_{2} H H O_{2}N$$

compound R

I Compound **R** contains a chiral centre. Identify the chiral centre in the formula of compound **R** by using an asterisk (*).

11	enantiomers and how these affect plane-polarised light. [2]
III	State the type of reaction that occurs when compound \mathbf{R} is converted to
111	HNS by the use of a suitable base. [1]
	Total [13]

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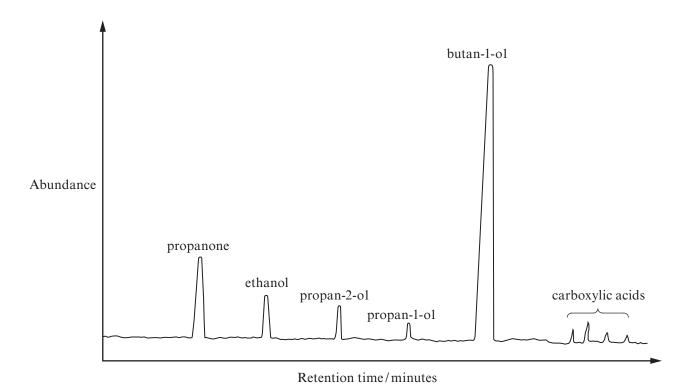
2. (a) Butan-1-ol can be produced by the reduction of butanal.

(i) State the name of a reducing agent that can be used for this reaction. [1]

(ii) The infrared spectrum of butanal shows an absorption at 1731 cm⁻¹. State which bond in butanal is responsible for this absorption and explain how the intensity of this absorption changes as the reduction proceeds. [2]

(b) A traditional route for making butan-1-o1 is by the fermentation of sugar cane residues and other starch-containing materials. One problem with this method is that a number

of other products are also obtained. The gas chromatogram shows the major products from a typical fermentation of starch.



Use the chromatogram to help you answer the questions below and opposite.

(i) State, in **decreasing** order of abundance, the three main products of this fermentation. [1]

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	(ii)	State which one of the products in (i) cannot normally be oxidised to a carboxylic acid. [1]
	(iii)	Select two compounds, from the chromatogram, that will give a positive result in the triiodomethane (iodoform) reaction giving an explanation for your answer. [2]
		Compounds and
		Explanation
	(iv)	Since there is a plentiful supply of cellulose from plants, scientists are using a new bacterium to ferment cellulose rather than starch. The first results of this research have been promising. If this new method is to be tested by other research groups why is it essential that the conditions are kept exactly the same? [1]
(c)	wou	re is interest in developing butan-1-ol as a fuel to replace petrol and diesel as this ld be a carbon neutral fuel. gest why this fuel is described as carbon neutral, giving a reason for your answer. [2]
(d)		rge proportion of the butan-1-ol produced is used to react with ethanoic acid to luce 1-butyl ethanoate.
	(i)	Give the equation for this reaction. [1]
	(ii)	State the name of the catalyst that is used. [1]
		Total [12]

Read the passage below and then answer parts (a)-(g) in the spaces provided.

3.

Anaesthetics

It is difficult to believe, in these days of modern medicine, that only 100 years ago tooth extractions were still being carried out without any form of anaesthetic. Modern anaesthetics are of two types – general anaesthetics, which have a whole body effect and local anaesthetics, which remove pain at the site of surgery.

The first demonstrations of anaesthesia were in the late 1840s when Morton, in the USA, used ethoxyethane, and Simpson, in Scotland, used chloroform (trichloromethane). Later that century nitrous oxide, N₂O, was successfully used. In more recent times, up to about 1960, the most commonly used anaesthetics were cyclopropane and ethoxyethane. In the 21st century a number of safer general anaesthetics are in use, the choice depending on the condition for which they are being used. The formulae of some general anaesthetics are show below, together with their common names.

$$\begin{array}{c} \text{CH(CH}_3)_2 \\ \text{CH}_3\text{CH}_2\text{OCH}_2\text{CH}_3 \\ \text{ethoxyethane} \end{array} \qquad \begin{array}{c} \text{CH}_2 \\ \text{CH}_2 \\ \text{CH}_2 \end{array}$$

In the twentieth century, the general anaesthetic nitrous oxide was still being used for routine dental extractions but in recent years the use of local anaesthetics has become the norm.

A popular choice for dentists is the ester procaine, although a number of other compounds are available; their use depending on the anaesthetic effect required and its duration of action. The formulae and common names of some local anaesthetics are shown below.

$$\begin{array}{c} O \\ \parallel \\ -C-O-CH_2-CH_2-N \\ \hline \\ CH_2CH_3 \end{array} \quad procaine \\ \\ CH_2CH_3 \end{array}$$

(1094-01)

Benzocaine, for external use only, has uses in skin creams for which a numbing action is required.

$$\begin{array}{c} O \\ \parallel \\ -C-O-CH_2-CH_3 & benzocaine \end{array}$$

The relative safety of the anaesthetic itself is an important factor in deciding which anaesthetic to use for each situation. However, another factor to be considered is the method by which the body metabolises the anaesthetic and the nature of the compounds that are produced.

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[1]

(<i>a</i>)		nitrogen monoxide.	gen
		$2N_2O \longrightarrow N_2 + 2NO$	
		ogen monoxide is a reactive molecule that contains an unpaired electron. e the general name for species that contain an unpaired electron.	[1]
<i>(b)</i>	The flam C ₃ H	use of cyclopropane as an anaesthetic causes concern because of its extremability. Give the balanced equation for the complete combustion of cyclopropa	eme ane, [1]
<i>(c)</i>	one	oxyethane, $CH_3CH_2OCH_2CH_3$, reacts with some aqueous acids to give ethano of the products. stage of this reaction can be represented as follows:	ıl as
CH ₃ CH ₂ -	-Ö-(i) H+	$CH_2CH_3 \longrightarrow CH_3CH_2 \stackrel{+}{\overset{\circ}{\bigcirc}} CH_2CH_3 \longrightarrow CH_3CH_2 \stackrel{\circ}{\overset{\circ}{\bigcirc}} + {}^+CH_2CH_3$ State why ethoxyethane is behaving as a nucleophile in this reaction stage.	CH ₃
	(ii)	I To reduce the danger of fire when carrying out this reaction, the reacta are heated together under reflux. State what is meant by the term heating under reflux.	ants
			•••••

The reactants need to be refluxed at a temperature of 130 °C.

The article mentions the use of halothane (line 12) and desflurane (line 12) as general anaesthetics. State and explain which of these two compounds could cause more damage

laboratory fume cupboard.

Suggest how this mixture could be safely heated at this temperature in a

Π

to the ozone layer.

(*d*)

- (e) Propofol, which allegedly caused the death of Michael Jackson in 2009, is an important intravenous anaesthetic.
 - (i) State what would be seen if a few drops of iron(III) chloride solution were added to a solution of propofol in a suitable solvent. [1]

$$CH(CH_3)_2$$
 $-OH$
 $CH(CH_3)_2$
propofol

(ii) The formulae of propofol, compound L and compound M are shown below.

I The three compounds are dissolved separately in a suitable solvent and each solution tested with universal indicator paper and with sodium hydrogenearbonate solution. Complete the table below giving any **observations** or writing 'no reaction' as appropriate.

Compound	Colour given with universal indicator paper	Observation with sodium hydrogencarbonate solution
propofol		
compound L	green	no reaction
compound M		

[2]

II Give the test for any gas produced with sodium hydrogencarbonate solution. [1]

.....

<i>(f)</i>	The	article describes procaine (line 19) as an ester.
	(i)	Draw the section of the formula that identifies procaine as an ester. [1]
	(ii)	Use the formula of procaine to help you draw the structural formula of the nitrogen-containing alcohol that will react with a suitable acid to give procaine. [1]
(g)	deter	benzocaine has a melting temperature of 89°C. A melting temperature rmination shows that a sample of benzocaine is impure. two observations that would indicate that this sample is impure. [2]
		Total [15] Total Section A [40]

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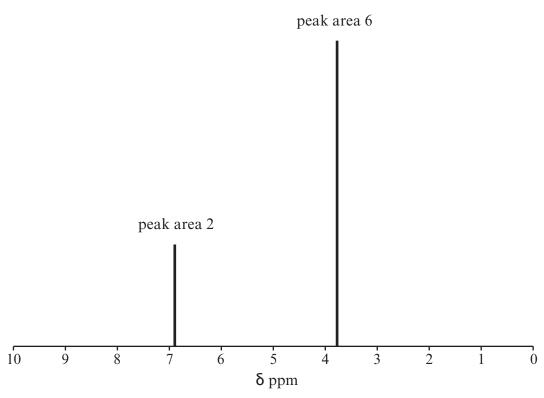
(1094-01) **Turn over.**

SECTION B

Answer both questions in the separate answer book provided.

4. (a) In 2009 a man received compensation after he received chemical burns from a compound used as an antifungal agent in an imported leather sofa. The compound causing this problem was the ester dimethyl fumarate (DMF).

- (i) DMF is one of a pair of E-Z stereoisomers. State what is meant by *stereoisomerism* and draw the displayed formula of the other stereoisomer of DMF. [2]
- (ii) The NMR spectrum of the ester DMF is shown below.



The hydrolysis of dimethyl fumarate produces the dicarboxylic acid, fumaric acid. Describe how the NMR spectrum of fumaric acid would be different from the NMR spectrum of dimethyl fumarate.

Your answer should identify the peaks involved and include reasons for any changes that occur. You should also identify any NMR signal that does not change and the reasons for this. [5]

(QWC) [1]

- (iii) The mass spectrum of DMF, C₆H₈O₄, shows only a weak signal for its molecular ion at m/z 144. The strongest signal is seen at m/z 113. Suggest a molecular formula for the positive ion at m/z 113, giving your reasoning. [2]
- (b) Fumaric acid (E-butenedioic acid) is an important starting compound for the manufacture of many other materials. The usual method for producing fumaric acid is from crude oil, but there is increasing interest in a fermentation route, using enzymes, from a sugar such as glucose. A comparison of these two methods is shown in the table.

	Crude oil route	Fermentation route
Raw material	Maleic anhydride	Glucose
Reaction temperature / °C	95	35
Raw material price / £ kg ⁻¹	1.00	0.30

- (i) Suggest **one** way that the crude oil route could be made more economic to operate. [1]
- (ii) At present the crude oil route is the more economic route from which to obtain fumaric acid.

 Suggest **one** factor by which the fermentation route could be modified to make it more competitive with the crude oil route, other than by simply increasing the yield. You are reminded that the optimum temperature for enzymes in this reaction is 35 °C.
- (iii) Fumaric acid was obtained in a pilot-scale experiment by the fermentation route, using glucose.A simplified equation for the reaction is shown below.

$$2CO_2 + C_6H_{12}O_6 \longrightarrow 2C_4H_4O_4 + 2H_2O$$
 $M_r 180 \qquad M_r 116$

(1094-01)

In this experiment 12.6 kg of glucose (70 moles) gave 13.0 kg of fumaric acid. Calculate the percentage yield of fumaric acid. [3]

- (iv) Small amounts of other organic acids are produced during the fermentation.
 - I One of these acids is ethanoic acid.
 Outline any **one** other reaction that produces ethanoic acid.
 Your answer should include
 - the name of your starting material,
 - any other reagent(s) used,

• the type of reaction occurring.

[3]

II A small amount of butanedioic acid is also produced.

butanedioic acid

This acid can also be produced by the hydrogenation of the unsaturated acid, fumaric acid.

fumaric acid

Suggest a suitable catalyst for this hydrogenation.

[1]

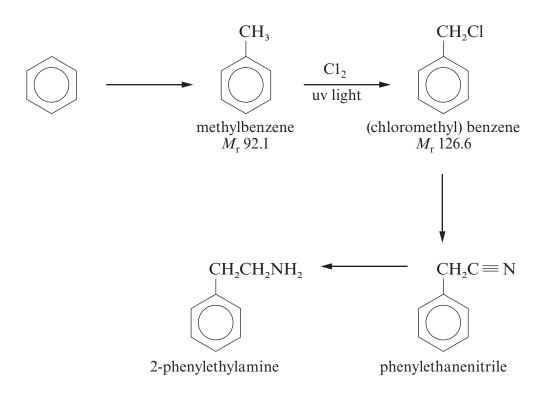
(c) A research student claimed to have made compound U, which is an isomer of fumaric acid.

State a reagent that would react with compound U and not fumaric acid, giving the result of the test. [1]

Total [20]

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5. (a) 2-Phenylethylamine, present in chocolate, can be made from benzene in four stages.



- (i) Give the equation, and the name of a suitable catalyst, for the Friedel-Crafts alkylation of benzene leading to methylbenzene. [2]
- (ii) (Chloromethyl)benzene is produced by passing chlorine gas into methylbenzene in the presence of ultraviolet light. In practice the substitution by chlorine can proceed further giving (dichloromethyl)benzene and (trichloromethyl)benzene. In order to prevent further chlorination the reaction is stopped when the increase in mass corresponds to (chloromethyl)benzene being produced. You should assume that the other product, gaseous hydrogen chloride, is lost from the mixture.

Mass of flask + product = 158.4 g Mass of flask + methylbenzene = 148.0 g Mass of flask = 120.4 g

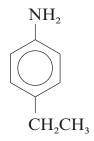
In an experiment the following results were obtained.

Show that the increase in mass corresponds to the conversion of all the methylbenzene into (chloromethyl)benzene. [4]

- (iii) State the names of the reagents necessary to convert
 - I (chloromethyl)benzene to phenylethanenitrile, [1]
 - II phenylethanenitrile to 2-phenylethylamine. [1]

(b) Explain why 2-phenylethylamine is a base.

- [2]
- (c) State how both 2-phenylethylamine and its isomer 4-ethylphenylamine react with nitric(III) (nitrous) acid at 5 °C.



4-ethylphenylamine

In **each** case you should state the type of compound produced and any relevant observations. [3]

(d) Kevlar \mathbb{R} is a polyamide that is used in bullet-proof vests.

Give the formula of two starting materials that can be reacted together to give Kevlar ®. [2]

(e) Silk is a naturally occurring material composed of polymerised serine molecules. Serine is an α -amino acid.

$$\begin{array}{c|c} H & NH_2 & O \\ & | & | & \\ & -C - C - C \\ & | & | \\ & H & H & O - H \end{array}$$
serine

- (i) Give the **systematic name** of serine, which is a derivative of propanoic acid. [1]
- (ii) Hydrogen bonding is largely responsible for the solubility of serine in water. Explain what is meant by hydrogen bonding, using serine to illustrate your answer.

 [3]

(QWC) [1]

Total [20]

Section B Total [40]



GCE A level

CHEMISTRY CH4 DATA SHEET

P.M. WEDNESDAY, 15 June 2011

Infrared Spectroscopy characteristic absorption values

Wavenumber/cm ⁻¹
500 to 600
650 to 800
1000 to 1300
1620 to 1670
1650 to 1750
2100 to 2250
2800 to 3100
2500 to 3550
3300 to 3500

Nuclear Magnetic Resonance Spectroscopy

Candidates are reminded that the splitting of any resonance into \mathbf{n} components indicates the presence of $\mathbf{n-1}$ hydrogen atoms on the **adjacent** carbon, oxygen or nitrogen atoms.

Typical proton chemical shift values (δ) relative to TMS = 0

Type of proton	Chemical shift (ppm)
$-CH_3$	0.1 to 2.0
R — CH_3	0.9
$R-CH_2-R$	1.3
CH₃—C≡N	2.0
CH_3-C O $-CH_2-C$	2.0 to 2.5
$-CH_2-C$	2.0 to 3.0
$-O-CH_3$, $-OCH_2-R$, $-O-CH=C$	3.5 to 4.0
R—OH	4.5 *
$CH_2=C$	4.8
$CH_2=C$ $R-C$ H	9.8 *
$R-C \bigcirc O$ OH	11.0 *

^{*}variable figure dependent on concentration and solvent

THE PERIODIC TABLE

							IHE	PEKI	THE PEKIODIC TABLE	IABLE								
		7						5	Group				n	4	w	9	_	0
Period		s Block															L	
*								1	V 23.									4.00
-	1.01 H							4	,									Helium
T	Hydrogen 1							A,	relative	ic		,			p Block	ock		7
C	6.94 Li	9.01 Be						Symbol Name					10.8 B	12.0 C	0.41 0. X	16.0 O	19.0 F	20.2 Ne
7	Lithium 3	Beryllium 4							number	ı			Boron 5	Carbon 6	Nitrogen 7	Oxygen 8	Fluorine 9	Neon 10
c	23.0 Na	24.3 Mg											27.0 Al	28.1 Si	31.0 P	32.1 S	35.5 CI	40.0 Ar
?	Sodium 11	Magnesium 12					d Block	ock				^	Aluminium 13	Silicon 14	Phosphorus 15	Sulfur 16	Chlorine 17	Argon 18
	39.1 K	40.1	45.0 SC	47.9 Ti	50.9 V	52.0 Cr	54.9 Mn	55.8 Fe	58.9	58.7 N:	63.5	65.4 Zn	69.7 Ga	72.6 Ge	74.9 As	79.0 Se	79.9 Br	83.8 Kr
4	Potassium 19	Calcium 20	Scandium 21	Titanium 22	Vanadium 23	Chromium 24	Manganese 25	Iron 26	Cobalt 27	Nickel 28	Copper 29	Zinc 30		Germanium 32	Arsenic 33	Selenium 34	Bromine 35	Krypton 36
	85.5 Pb	87.6	88.9	91.2 7 :	92.9 NB	95.9 Mo	98.9 7.7	101	103 D.b	106 Dd	108	7 7 7	115 La	911 S.	122 S	128 T.	127	131 X o
S	Rubidium	Strontium	Yttrium	Zirconium	u	Molybdenum	Technetium	Ruthenium	Rhodium	Palladium	Ag Silver	Cadmium	Indium	Tin S	Antimony	Tellurium	I Iodine	Xenon 54
	7.5	30	951	04	191	747	186	1 5	5	105	/+ [64 26	4 6	000	1000	25	6.6	+ CC
9	Cs Cs	Ba	La La	Hť	181 Ta	N 84	Re Re	OS O	l92 Ir	Pt 33	Au Au	Hg	707 E	Pb	Bi	(210) Po	(210) At	(222) Rn
	Caesium 55	Barium 56	Lanthanum 57	Hafnium 72	Tantalum 73	Tungsten 74	Rhenium 75	Osmium 76	Iridium 77	Platinum 78	Gold 79	Mercury 80	Thallium 81	Lead 82	Bismuth 83	Polonium 84	Astatine 85	Radon 86
7	(223) Fr Francium 87	(226) Radium 88	(227) Ac Actinium 89															
			'							f Block	ock							
		►Lan eler	► Lanthanoid elements	Ce Cerium 58	141	144 Nd Neodymium 60	(147) Pm Promethium 61	Samarium 62	(153) Eu Europium 63	157 Gd Gadolinium 64	159 Tb Terbium 65	163 Dy Dysprosium 66	165 Ho Holmium 67	167 Er Erbium 68	Tm Thulium	173 Yb Ytterbium 70	175 Lu Lutetium	
		►► Actinoid elements	Actinoid elements	232 Th Thorium	(231) Pa Protactinium 91	238 U Uranium 92	(237) Np Neptunium 93	Pu Putonium 94	(243) Am Americium 95	(247) Cm Curium 96	(245) Bk Berkelium 97	(251) (254) Cf Es Californium Einsteinium 98 99	(254) Es Einsteinium 99	(253) Fm Fermium 1	(256) Md Mendelevium 101	(254) No Nobelium 102	(257) Lr Lawrencium 103	