Surname	Centre Number	Candidate Number
Other Names		2



GCE A LEVEL

A410U20-1





CHEMISTRY – A level component 2 Organic Chemistry and Analysis

TUESDAY, 12 JUNE 2018 – AFTERNOON 2 hours 30 minutes

Section A
Section B

For Exa	For Examiner's use only		
Question	Maximum Mark	Mark Awarded	
1. to 9.	15		
10.	19		
11.	18		
12	17		
13	16		
14	19		
15	16		
Total	120		

ADDITIONAL MATERIALS

In addition to this examination paper, you will need a:

- · calculator:
- Data Booklet supplied by WJEC.

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.

Section B Answer **all** questions in the spaces provided.

Candidates are advised to allocate their time appropriately between **Section A (15 marks)** and **Section B (105 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 120.

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

The assessment of the quality of extended response (QER) will take place in **Q.11**(*a*) and **Q.14**(*b*)(ii). If you run out of space, use the additional page(s) at the back of the booklet, taking care to number the question(s) correctly.

SECTION A

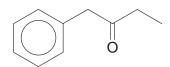
Answer all questions in the spaces provided.

1. Give the **name** of the monomer that is used to produce the polymer shown below. [1]

2. Give the structure of a product obtained when 1,4-dimethylbenzene is oxidised by alkaline potassium manganate(VII) and acidified. [1]

3. The compound below is reduced by sodium tetrahydridoborate(III).

State, giving a reason, whether a primary, secondary or tertiary alcohol is produced. [1]



4. The formulae for two aromatic compounds are given below.

$$\begin{array}{c} \mathsf{CH_3} \\ \mathsf{H_3C} \\ \mathsf{CH_3} \\ \mathsf{NO_2} \\ \\ \mathsf{mesitylene} \\ \end{array}$$

Complete the table which describes the ¹HNMR signals for these two compounds.

Compound	Number of peaks	Relative peak area ratio
mesitylene		
TNT		

5.	Draw the structure for a compound of formula C ₃ H ₈ O that shows an infrared absorption	n peak
	between 1000 and 1300 cm ⁻¹ but no peak between 2500 and 3550 cm ⁻¹ , other than t	that at
	2800-3100 cm ⁻¹ corresponding to C—H bonds. Give your reasoning.	[2]

[2]

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6.	(a)	Give the structure of the organic compound produced when 2-hydroxybenzenecarboxylic acid reacts with ethanoyl chloride.	[1]
	(b)	If the product in (a) was contaminated with unreacted 2-hydroxybenzenecar describe a chemical test that would show its presence. State the reagents upon the content of the	
	***********	observation made.	[2]
7.	Benz	zene-1,2-dicarboxylic acid (or its sodium salt) is heated with sodalime.	
	State	e the type of reaction that occurs and give the name of the product.	[2]

The boiling temperatures of three compounds of formula ${\rm C_3H_8O_2}$ are given in the table. 8.

Compound		Boiling temperature / °C
L	CH ₃ -CH ₂ -O-O-CH ₃	40
М	CH ₃ -O-CH ₂ -O-CH ₃	42
N	HO-CH ₂ -CH ₂ -CH ₂ -OH	214

Suggest why the boiling temperature of compound ${\bf N}$ is considerably higher than the boiling temperatures of the other two compounds, illustrating your answer by means of a diagram. [2]

Draw the structure of a compound that has an empirical formula CH₄N. 9.

[1]

SECTION B

Answer all questions in the spaces provided.

10. The willow tree has been used for hundreds of years to obtain substances with medical benefits. One of these substances is salicin, which shows analgesic and anti-inflammatory properties.

Salicin is a molecule that consists of two parts – one based on glucose and the other on (2-hydroxyphenyl)methanol, $C_6H_4(OH)CH_2OH$.

(a) The formula of glucose can be written as an open-chain formula.

Glucose is a reducing sugar and will react with Fehling's reagent.

State what is seen in the test with Fehling's reagent and identify the group present in glucose that is responsible for this reaction. [2]

(b) An outline of the method used to make glucose pentaethanoate is shown below.

- 0.015 mol of glucose is placed in a flask with a large excess of ethanoic anhydride.
- The mixture is heated under reflux at 100 °C for one hour.
- The product is cooled and poured into a beaker containing crushed ice.
- After stirring and allowing the ice to melt, the mixture is filtered.
- Ethanol is then used to recrystallise the glucose pentaethanoate.
- (i) In this reaction the —OH groups in glucose react with ethanoic anhydride to give ethanoate groups.

State and explain the minimum number of moles of ethanoic anhydride used in this reaction. [2]

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(ii)	Briefly outline how the mixture could be 'heated under reflux at 100 °C'.	[2]
(iii)	State how the method implies that glucose pentaethanoate is insoluble in water	.[1]
		•••••
(iv)	During recrystallisation glucose pentaethanoate is dissolved in the 'minim quantity of hot ethanol'.	um
	Why was the 'minimum quantity' used?	[1]
(v)	The melting temperature of glucose pentaethanoate is 134°C. In the experim the sample is found to melt at a lower temperature.	
	Suggest a possible cause for this lower value.	[2]
		•••••

(c) Ethanol can be made from glucose by fermentation.

$$C_6H_{12}O_6(aq) \longrightarrow 2C_2H_5OH(aq) + 2CO_2(g)$$

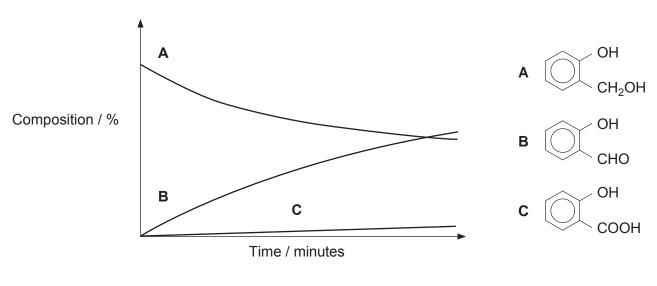
In an experiment 0.200 mol of glucose was dissolved in water and fermented using yeast. At the end of the reaction all the glucose had been converted into ethanol and carbon dioxide. If the volume of the resulting solution was 2.03 dm³, calculate the concentration of ethanol in g dm⁻³.

(d) The other product from the breakdown of salicin is (2-hydroxyphenyl)methanol – compound **A** below.

compound A

Compound **A** is both a phenol and an alcohol and oxidation of the alcohol group results in a number of other products, including compounds **B** and **C**. Both of these compounds have important commercial uses and a method that gives them in high yields is desirable.

In one method, compound **A** is oxidised at 50 °C using water as the solvent. The graph shows the composition of the resulting mixture.



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(i)	State how the quantity of compound A present in the mixture changes over time. [1]
(ii)	Comment on the proportion of compounds B and C in the mixture as the reactio	
	proceeds. [1]
•••••		
(iii)	The proportion of products present was found using HPLC.	
	Calculate the percentage of compound A oxidised after 15 minutes. [1	1

	Concentration / mol dm ⁻³		
Compound	Start	After 15 minutes	
Α	0.20	0.02	
В	0.00	0.10	
С	0.00	0.01	
other products	0.00	0.07	

Percentage oxidised = %

(iv) Compounds **A**, **B** and **C** are colourless and were detected during HPLC by using ultraviolet radiation.

Absorption maximum / nm	Compound
202.1	С
212.2	В
273.4	Α

	State, giving a reason, which of these three compounds has the absormaximum of the highest energy.	orption [2]
(v)	This oxidation can also be carried out using benzene as the solvent. Compound B is the only product and the rate of oxidation is higher than when water as the solvent.	using
	Give a reason why the method using water is seen as a 'greener' process.	[1]

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11. (a) The characteristic smell of some varieties of blue cheese is due mainly to heptan-2-one.

A teacher was asked whether heptan-2-one could be clearly distinguished from heptan-3-one by some suggested methods. His responses are shown in the table.

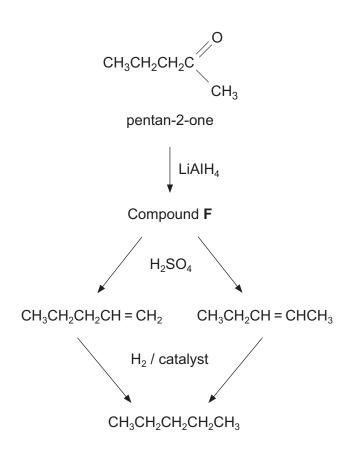
Method	Clearly identified?
mass spectroscopy	yes
gas-liquid chromatography	no
boiling temperature	yes
chemical analysis for C, H and O	no
reaction with alkaline iodine	yes
reaction with 2,4-dinitrophenylhydrazine	no
reaction with Tollens' reagent	no

Examiner only

Explain why each response is correct.	[6 QER]

(b) Pentan-2-one contributes to the smell of other blue cheese varieties.

A reaction sequence for obtaining pentane from pentan-2-one is shown below.



(i) Give the **skeletal** formula of compound **F**.

[1]

(ii) State the role of sulfuric acid when it reacts with compound **F** and explain why two products are formed. [2]

(iii)	The final stage involves catalytic hydrogenation. One method of carrying this out is to dissolve the alkene in a suitable solvent and react it with hydrogen at room temperature in the presence of a heterogeneous catalyst.				
	State a catalyst that can be used and why this is described as a <i>heterogeneous</i> catalyst. [2]				
•••••					
(iv)	One solvent that can be used for the hydrogenation described in (iii) above is hexane. The boiling temperatures of pentane and hexane are 35 °C and 69 °C respectively.				
	State the name of a method that can be used to separate these two compounds. [1]				
• • • • • • • • • • • • • • • • • • • •					

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(v) The boiling temperatures of some alkanes are shown in the table.

Alkane	Boiling temperature / °C
2,2-dimethylpropane	9
2-methylbutane	28
pentane	35
2,2-dimethylbutane	50
2-methylpentane	60
hexane	69
heptane	98

State two factors that affect the boiling temperature of an alkane that can be deduced from the formulae of these alkanes. Suggest reasons for these variations in boiling temperatures. [4]

(vi) A radical reaction occurs when alkanes are reacted with chlorine in the presence of ultraviolet light. For example in the chlorination of butane.

I. Give the **displayed** formula of the carbon-containing radical that leads to 2-chlorobutane. [1]

II. State the name of an alkane that can be made as a side product during the chlorination of butane. [1]

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12. (a) At room temperature propene reacts with chlorine by electrophilic addition.

Complete the mechanism for this reaction.

[2]

(b) At 500 °C propene reacts with chlorine via a radical mechanism to give 3-chloropropene together with hydrogen chloride.

3-chloropropene

Describe a simple chemical test to distinguish between 1,2-dichloropropane [formed in part (a)] and 3-chloropropene. Give the reagent and the result with both compounds. [2]

(c) (i) 3-Chloropropene can be converted to prop-2-en-1-ol by reaction with sodium hydroxide in a nucleophilic substitution reaction.

Give the formula of the nucleophile used in this reaction.

[1]

(ii) An alternative method of obtaining prop-2-en-1-ol is by the rearrangement of epoxypropane.

When choosing an appropriate method there are a number of factors to be considered.

Suggest **two** factors that should be considered if prop-2-en-1-ol is to be produced from epoxypropane rather than by the method described in part (i). [2]

1.	 	 	 	
2.				

- (d) Creosote is a pale yellow liquid that is used as a wood preservative. This material is largely a mixture of various phenols, including 4-methylphenol.
 - (i) In the laboratory 4-methylphenol can be made in several stages from methylbenzene.

First stage The nitration of methylbenzene

Second stage The reduction of 4-nitromethylbenzene

Third stage Making 4-methylphenol from the amine 4-methylphenylamine

I. Write the mechanism for the formation of 4-nitromethylbenzene, starting with methylbenzene and the nitronium ion. As part of your answer you should state the reagents used to generate the nitronium ion and name the type of reaction mechanism occurring. [4]

Reagents used

Type of reaction mechanism

II. Suggest a suitable reducing agent used to produce 4-methylphenylamine from 4-nitromethylbenzene. [1]

III. Complete the equation below which shows the production of 4-methylphenol from 4-methylphenylamine. [2]

(ii) pK_a is a term used to express the acidity of organic compounds. The lower the value, the greater the tendency for the compound to lose a hydrogen ion, making it more acidic.

The table shows the pK_a values for some compounds.

Compound	pK _a
ethanoic acid	4.8
4-nitrophenol	7.2
phenol	9.9
4-methylphenol	10.2

I.	Ethanoic acid is more acidic than phenol. State a chemical test that you couuse in the laboratory to show this difference in acidity. You are given aqueousolutions of each compound.	
II.	Suggest why 4-nitrophenol is much more acidic than phenol.	[1]
•••••		

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Examiner
only

13.	(a)	(a) Explain why amino acids are amphoteric compounds.							
 (b) If an amino acid is treated with methanal, the resulting compound can be titrated aga sodium hydroxide solution in a 1:1 ratio. 4.95 g of an amino acid was treated with methanal and the resulting solution made u 250 cm³. 25.0 cm³ of this solution was then titrated with sodium hydroxide of concentra 0.105 mol dm⁻³. The results are shown in the table below. 							made up to		
		Titration	1	2	3	4	5		
		NaOH(aq) used / cm ³	38.70	35.90	36.00	32.00	36.10		

(i)	Suggest a practical reason why the reading for titration 1 was too high.	[1]
•••••		

(ii) Use appropriate titration values to calculate the relative molecular mass of the amino acid. [5]

*M*_r =

(iii) Assuming that the amino acid in part (ii) is a straight chain aliphatic α -amino acid deduce its structure. [2]

(c) The formulae of three amino acids are shown below.

(i)	State which of the three compounds could be identified by its ability to r plane of plane polarised light. Give a reason for your answer.	otate the
(ii)	State how the infrared absorption spectrum of compounds S and T wo from each other in their significant functional group absorption(s).	uld differ
iii)	State which of these amino acids would not be able to form two different d with either of the other two amino acids. Explain your answer.	ipeptides [1]

(d) Nylon-11 is a bio-sourced polyamide which is made from castor oil. Undec-10-enoic acid is produced as an intermediate compound. This acid is reacted, under suitable conditions, to give 11-bromoundecanoic acid, which is then treated with ammonia to produce 11-aminoundecanoic acid. Polymerisation of this product gives nylon-11.

(ii) The bromo-compound reacts with ammonia to produce 11-aminoundecanoic acid.One step in the mechanism for this reaction is shown below.

I. Explain how partial charges (δ + and δ –) arise on the carbon and bromine atoms. [1]

II. State the role of ammonia in this reaction. [1]

(iii) Draw the structure of nylon-11, indicating the repeating unit present.

The formula of 11-aminoundecanoic acid is shown below. [1]

$$H_2N - (CH_2)_{10} - C$$

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14.	(a)	(i)	(i) Describe the structure and bonding in benzene. Explain why benzene is resistant to addition reactions and why its usual mode of reaction is by substitution.					
			You may include a diagram as part of your answer. [4	.]				

[2]

- (ii) Benzene is an example of an aromatic compound. There are a number of other aromatic systems. One of these is pyrrole.
 - I. Pyrrole can be made by reacting butan-1,4-dial with ammonia.

Calculate the atom economy of this reaction.

Atom economy = %

II. Pyrrole reacts in a similar way to benzene, although it is more reactive.

For example, pyrrole reacts readily with iodine to produce 2,3,4,5-tetraiodopyrrole.

$$\begin{bmatrix} \mathsf{C} & \mathsf{C} & \mathsf{C} \\ & \mathsf{C} & \mathsf{C} \\ \mathsf{C} & \mathsf{C} \\ \mathsf{N} & \mathsf{H} \end{bmatrix}$$

Suggest the balanced equation for the reaction between iodine and pyrrole, leading to 2,3,4,5-tetraiodopyrrole. [1]

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(b) Diseases caused by sucking insects are a major problem in hot countries. Extensive research has been carried out to find compounds that are deterrents against these insects. One of the these compounds is DEPA.

$$\begin{array}{c|c} CH_2 - C & \\ & CH_2CH_3 \\ \\ CH_2CH_3 \end{array}$$

DEPA

(i) In the first stage of a synthesis of DEPA, methylbenzene [M_r 92.1] reacts with chlorine until the increase in mass indicates that (chloromethyl)benzene, $C_6H_5CH_2CI$ [M_r 126.6] has been produced.

In an experiment 0.430 mol of methylbenzene was used. Calculate the increase in mass that will indicate that the conversion to $C_6H_5CH_2Cl$ is complete. [2]

Increase	in	mass	=	_	
IIICIEASE	1111	เมลออ		u	١.

(ii)	Suggest a three stage synthesis of phenylethanoic acid ($C_6H_5CH_2COOH$) starting from methylbenzene. In the first stage, methylbenzene reacts with chlorine in a similar way to the reaction of chlorine with methane.							
	For each stage you should state the reactants and the products as well as any important conditions, relevant equations and the type of reaction or mechanism occurring. [6 QER]							
•••••								
•••••								
•••••								
•••••								
(iii)	The overall yield of phenylethanoic acid, starting from methylbenzene, was 65%.							
	Calculate the mass of phenylethanoic acid $[M_{\rm r}\ 136]$ formed from 0.430 mol of methylbenzene. [1]							
	Mass = g							

(iv) The final stage to make DEPA is shown in the equation below.

$$CH_{2}-C = C + (CH_{3}CH_{2})_{2}NH$$

$$CI \qquad \qquad \text{diethylamine}$$

$$CH_{2}-C = C + HCI$$

$$N(CH_{2}CH_{3})_{2}$$

$$DEPA$$

A teacher asked some students for suggestions as to how this reaction should be carried out.

I. One student suggested that the reaction could be carried out using water as the solvent.

II. Another suggestion was that an excess of diethylamine should be used.

Suggest why this method might give a very poor yield of DEPA.

Explain why this method should be used. [1]

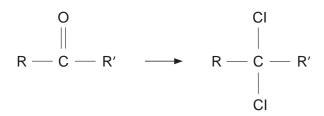
III. A further suggestion was that DEPA should be purified by distillation under reduced pressure as DEPA is a liquid with a high boiling temperature.

Explain why this process of separation should be used. [1]

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[1]

15. (a) Ketone **W** reacted with phosphorus(V) chloride to give the corresponding dichloro compound.



ketone W

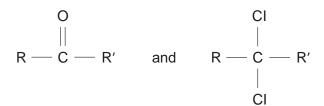
(i)	Describe how to show the presence of chlorine in this compound.	[4]

(ii) 6.35 g of the dichloro compound formed from ketone **W** contained 50.3 % of chlorine by mass, the remainder being carbon and hydrogen.

Calculate the relative molecular mass of the dichloro compound. [2]

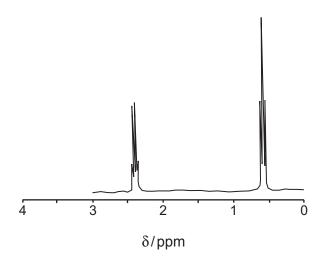
*M*_r =

(iii) Explain why both the molecules shown below have the same splitting pattern in their ¹H NMR high resolution spectra. [1]



.....

(iv) The ¹H NMR spectrum of ketone **W** is shown below.



Assuming that the alkyl groups R and R' are the same, use the information from the NMR spectrum to deduce the structure of ketone **W**. [3]

(b)		relative molecular mass of a volatile compound $f L$ can be found by weighing a sample measuring the volume of its vapour at a known temperature and pressure.
	(i)	In an experiment the following results were obtained.
		Mass of liquid $L = 0.222g$ Volume of vapour produced = 111 cm ³ Temperature = 423 K Pressure = 9.50×10^4 Pa
		Use this data to calculate the relative molecular mass of compound L . [2]
		$M_{r} = \dots$
	(ii)	 Tests on liquid L showed that it did not liberate carbon dioxide with sodium hydrogencarbonate solution. it contained 2 oxygen atoms in each molecule when it was heated with aqueous sodium hydroxide it did not produce ethanol as one of the products
		Discuss these results and then suggest a structure for liquid L . [4]

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