

Candidate Name	Centre Number	Candidate Number
		2



**General Certificate of Education
Advanced**

334/01

CHEMISTRY – CH4

A.M. TUESDAY, 22 January 2008

(1 hour 40 minutes)

ADDITIONAL MATERIALS

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.

INSTRUCTIONS TO CANDIDATES

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 **both** questions in **Section B** in a separate answer book which should then be placed inside this question-and-answer book.

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

ADVICE TO CANDIDATES

Candidates are advised to allocate their time appropriately between **Section A (35 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 75.

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

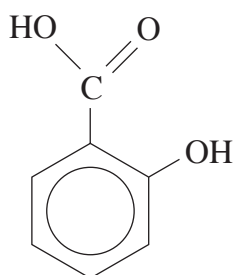
You are reminded that marking will take into account the Quality of Written Communication used in your answers.

No certificate will be awarded to a candidate detected in any unfair practice during the examination.

SECTION A

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

1. (a) Salicylic acid, which was originally derived from the bark of the Willow (*Salix*), has been used for centuries to treat a variety of complaints including fever and pain. It is now used extensively in the treatment of a range of skin conditions. It has the structure shown below.



- (i) **Name** the **two** functional groups in the molecule. [2]
- I.
- II.
- (ii) Calculate the relative molecular mass of salicylic acid. [1]
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- (iii) Draw the structure of the organic product formed when salicylic acid is treated with
- I. ethanol and concentrated sulphuric acid, [1]
- II. aqueous sodium hydroxide. [2]

- (b) 0.01 mole of a compound **A** reacts with bromine water to form 3.31g of a white compound **B**. Quantitative analysis of compound **B** shows that it contains 21.8% C, 0.90% H, 4.80% O and 72.5% Br by mass. Compound **A** also gives a purple colour with aqueous iron(III) chloride.

(It may be assumed that one mole of **A** reacts to give one mole of **B**.)

- (i) Calculate both the empirical and molecular formulae of compound **B**. [3]

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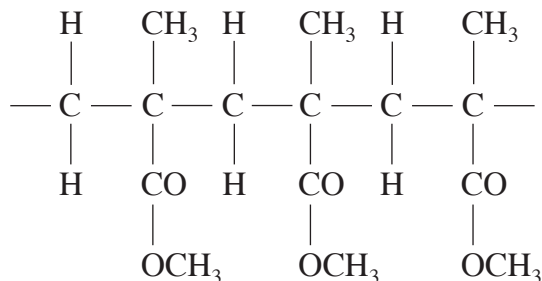
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- (ii) Give the structural formulae of compounds **A** and **B**. [2]

Total [11]

Turn over.

2. (a) Over the last 50 years, one of the great advances in chemistry has been in the field of polymers. A typical example of an addition polymer is perspex, a transparent plastic that can be used as a substitute for glass. Part of the structure is shown below.



Draw the structure of the monomer from which the plastic is formed. [1]

- (b) A typical example of a condensation polymer is nylon. Most of the nylon made in the UK is nylon-6,6 formed from hexanedioic acid and hexane-1,6-diamine.

(i) Draw the graphic (full structural) formula of hexanedioic acid. [1]

- (ii) Give a test, other than the use of indicators, to show that hexanedioic acid acts as a carboxylic acid.

State all the reagents and observations. [2]

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- (iii) Write a balanced equation for the ethanoylation of **one** amine group in hexane-1,6-diamine.
Use RNH₂ as the formula for hexane-1,6-diamine. [1]

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- (iv) Hexane-1,6-diamine is not very soluble in water. Explain why it dissolves readily in aqueous hydrochloric acid. [2]

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- (v) The repeating unit in nylon-6,6 is shown below.



- I. State the type of condensation polymer to which nylon-6,6 belongs. [1]

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- II. Give a reason for nylon made in this way being called nylon-6,6. [1]

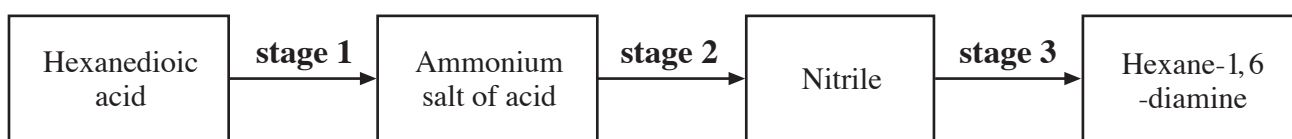
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- (vi) Give **two** large scale uses of nylon. [1]

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- (c) Hexane-1,6-diamine can be made from hexanedioic acid in a three stage process.



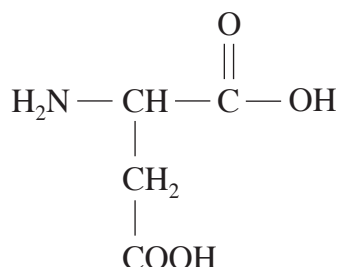
Classify the type of reaction which takes place in **stage 3** and name the reagent used for the conversion.

Type of reaction [1]

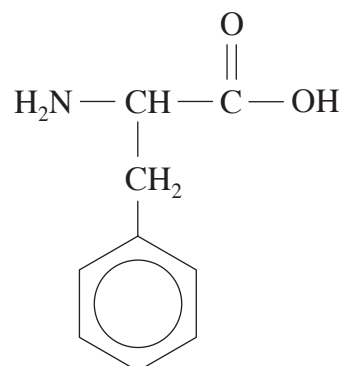
Reagent [1]

Total [12]

3. (a) Aspartame is an artificial sweetener which is about 200-times sweeter than sucrose. It is a methyl ester of a dipeptide formed from the following two α -amino acids.



aspartic acid



phenylalanine

- (i) Draw the structure of a dipeptide formed between these two α -amino acids. [1]
- (ii) On the diagram of the structure of **aspartic acid**, identify, using an asterisk (*), a chiral centre in the molecule. [1]
- (iii) Draw diagrams to show the two optical isomers of aspartic acid. [1]

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mirror

- (iv) State how the optical isomers of aspartic acid could be distinguished from each other. [1]

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- (v) In solution, phenylalanine exists largely in zwitterion form. Give the graphic formula of the zwitterion form of phenylalanine. [1]
- (vi) A disadvantage of using aspartame as a sweetener is that it breaks down quickly. In soft drinks stored in bottles, about 10% of the aspartame is destroyed each month. Suggest a reason, in terms of structure, for aspartame breaking down in this way. [1]
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- (b) When an electrical discharge is passed through hydrogen gas under suitable conditions, electromagnetic radiation is emitted.
- (i) Explain what processes cause this radiation to be emitted. [3]
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- (ii) The emission spectrum of atomic hydrogen is obtained as a series of discrete lines which converge.
- Explain why
- I. the lines are discrete, [1]
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- II. the lines converge. [1]
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- (iii) State what information can be derived from the convergence limit of the Lyman Series. [1]

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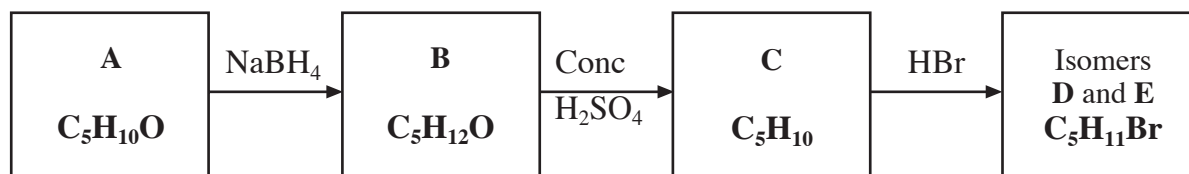
Total [12]

Section A Total [35]

SECTION B

Answer **both** questions in this section in the separate answer book provided.

4. (a) Study the reaction scheme shown below.



- (i) The infrared spectrum of **A** shows a strong absorption band at 1715 cm^{-1} .
 The mass spectrum of **A** shows a strong peak at *mass/charge* 57.
 The NMR spectrum of **A** has only two peaks, a triplet centered around $1.1\ \delta$, peak area 3 and a quartet centred around $2.4\ \delta$, peak area 2.
 The infrared spectrum of **B** shows a broad absorption at 3350 cm^{-1} .
 Use **all** the information to give the graphic (full structural) formulae for compounds **A**, **B** and **C**, explaining your reasoning. [8]
- (ii) State the type of reaction taking place for the conversion of
- I. **A** \longrightarrow **B**,
 - II. **B** \longrightarrow **C**. [2]
- (iii)
 - I. Name the type of isomerism shown by **C**. [1]
 - II. Explain why reacting HBr with **C** produces two isomers, **D** and **E**. [1]
 - III. Name the type of isomerism shown by isomers **D** and **E**. [1]
- (b) A sample of 1-bromobutane is divided into two portions. The first is warmed with dilute aqueous sodium hydroxide, whilst the second is warmed with concentrated sodium hydroxide in ethanol. In each case, a different organic product is formed.
- (i) For the first portion, name the organic product formed and give the mechanism for the reaction. [4]
 - (ii) For the second portion, name the organic product formed and write an equation for the reaction. [2]
- (c) Give a large scale use of a **named** organic halogen-containing compound of your own choice. [1]

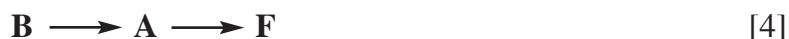
Total [20]

Turn over.

5. (a) Consider the following six compounds:



- (i) State, giving an explanation, which compound would have the highest boiling temperature. [2]
- (ii) Describe a reaction where compounds **D** and **E** show a similar chemical reaction. [1]
- (iii) For **each** of the following pairs of compounds, describe a **chemical** test that can be used to distinguish between them. The reagent(s) as well as the observation(s) for **each** compound, are required.
- I. Compounds **D** and **E** [2]
- II. Compounds **B** and **C** [2]
- (iv) State the reagent(s) and condition(s) needed for each stage in the following synthesis.



- (b) Although ethylamine can be prepared directly from chloroethane, phenylamine cannot be prepared directly from chlorobenzene. However, phenylamine can be synthesised from benzene.
- (i) State the type of reaction taking place when chloroethane is converted into ethylamine. [1]
- (ii) Give a reason why phenylamine cannot be prepared directly from chlorobenzene. [1]
- (iii) Outline a **two** step synthesis of phenylamine from benzene. Your answer should include reagents and conditions for **each** step. [4]
- (c) During a synthesis of phenylamine from benzene, 10.0 cm^3 of benzene produced 4.20 cm^3 of phenylamine. Given that the density of benzene is 0.878 g cm^{-3} , and that of phenylamine is 1.00 g cm^{-3} , calculate the percentage yield of the synthesis correct to 3 significant figures. [3]

Total [20]

Section B Total [40]