## **WJEC Chemistry A-level**

## OA2.2: Aldehydes and Ketones

**Practice Questions** 

**England Specification** 

Ibuprofen is a common drug taken as an analgesic and anti-inflammatory treatment.

A possible route to the synthesis of ibuprofen is shown below.

(a) Step 1 is a Friedel-Crafts alkylation reaction. Give the reagent(s) and condition(s) required for this step.

[3]

(b) Compounds <b>B</b> and <b>C</b> can be analysed using chemical tests.	
(i) Give a chemical test that would give a positive result for <b>both</b> compound <b>B</b> and compound <b>C</b> . Include reagent(s) and the observation(s) expected for a positive result.	
	[2]
(ii) Give a chemical test that would give a positive result for compound <b>C</b> but <b>not</b> for compound <b>B</b> Include reagent(s) and the observation(s) for both compounds.	٠.
	[2]
<ul> <li>(c) Compound C shows optical isomerism. Discuss this statement. Your answer should include:</li> <li>What is meant by optical isomerism.</li> <li>What feature of compound C allows it to exhibit optical isomerism.</li> </ul>	
<ul> <li>What feature of compound C allows it to exhibit optical isomerism.</li> <li>Diagrams to show the two optical isomers of compound C.</li> <li>How the two optical isomers of compound C can be distinguished.</li> </ul>	
[4] QW0	) [1] —
(d) Give the reagent(s) and condition(s) required for step 5 and classify the reaction that occurs.	
	[3]

(e) A student investigating alternative methods of producing ibuprofen suggests that it would be better to convert compound **C** into ibuprofen in a one-step process. Discuss whether this is correct. Your answer should include: • The reagent(s) and condition(s) for a reaction expected to convert compound C directly into ibuprofen. • Why it is generally better to use one step rather than two or more steps when producing a desired compound. A suggestion of why a two-step process is chosen for the synthesis of ibuprofen from compound **C** rather than a one-step process. [4] QWC [1] (Total 20) 2. Mauveine is a purple dye that was developed by Perkin in 1856 and was one of the first organic compounds to be synthesised on a large scale. He is credited with launching the synthetic chemical industry. (a) Give the name for the part of a molecule that causes it to be coloured. [1] (b) The dye mauveine often contains a mixture of impurities. Iwan and Georgia wanted to confirm that a sample of the dye was impure. (i) Iwan used the melting temperature of the sample to confirm that the sample was impure. Give **one** way that the melting temperature would show this. [1] (ii) Georgia used gas chromatography to confirm that the sample was impure. State what information she obtained using this method that Iwan could not obtain from the melting temperature.

			[2

(c) Another compound synthesised by Perkin was cinnamic acid. Cinnamic acid can be produced in two steps from phenylmethanol as shown below.

(i) Give the reagent(s) and condition(s) required to obtain a sample of benzenecarbaldehyde from phenylmethanol.

	 ,
Reagent(s)	
Condition(s)	

(ii) The conversion of phenylmethanol to benzenecarbaldehyde has a yield of 86 %. Calculate the mass of benzenecarbaldehyde that could be produced from 10.0 g of phenylmethanol.

[3]

[2]

(iii)	The <sup>1</sup> H NMR high resolution spectrum of cinnamic acid contains peaks in the area 7.0-7.5 with an area of 5 due to the benzene ring. Describe what other features you would expect to see in the spectrum. [4]	1
15.55		
232		
	Total [13]	
	From the information given, draw the displayed formula of each compound.In parts (i)-(iii) the unds consist of molecules that have <b>three</b> carbon In part (iv) the compound has <b>four</b> carbon	
(i) A co	empound that is oxidised to a ketone	
	[1	1]
(ii) A n	eutral sweet-smelling compound	
	[1	IJ
(iii) An	α-amino acid	
	[1	۱]
(iv) A ł	nydrocarbon that exhibits E-Z isomerism	
	[1	1]

(b) The active compound in  $Ventolin^{\textcircled{R}}$  inhalers used by asthma sufferers is salbutamol, which shows optical isomerism.

salbutamol

(i)	Indicate a chiral centre in this molecule by labelling it with an asterisk (*).
(ii)	State how the optical isomers of salbutamol could be distinguished from each other.
(iii)	Suggest a reason why only one optical isomer of salbutamol is used as a pharmaceutical.
(iv)	Draw the displayed formula of the likely organic product formed when salbutamol is refluxed with acidified $K_2Cr_2O_7$ . [2]
(c)(i) Arrange	the following molecules in order of increasing acidity
	[1]
	ethanoic acid ethanol ethylamine phenol
least acidic	most acidic

(ii) Explain the difference in acid-base properties of ethylamine and phenol.
[4]
(Total 14)
4. (a) 1-Chloropentane can be made by the free radical chlorination of pentane, in a similar way to the reaction of methane with chlorine.
(i) Give the equation for the reaction of pentane with chlorine, showing the displayed formula of 1-chloropentane as part of your answer.
[1]
(ii) The free radical reaction of pentane with chlorine gives other chlorinated organic products. Give the structural formula of the carbon-containing free radical that leads to the formation of 2-chloropentane.
[1]
(b) Pentylbenzene can be produced by the reaction of 1-chloropentane and benzene in a Friedel-Crafts reaction. State the name of a catalyst that can be used in this reaction.
[1]

(c) A Friedel-Crafts reaction can be carried out with ethanoyl chloride in place of 1-chloropentane. This reaction gives phenylethanone as the main organic product.

- (i) State what is seen when a few drops of phenylethanone are added to a solution of 2,4-dinitrophenylhydrazine.
  - [1]
    - (ii) This preparation of phenylethanone also gives small traces of an impurity. This impurity has a molecular formula C<sub>10</sub>H<sub>10</sub>O<sub>2</sub> and reacts in a similar way to phenylethanone when it is treated with 2,4-dinitrophenylhydrazine. It does not react with Tollens' reagent. Suggest a displayed formula for this impurity, giving a reason for your choice.
      [2]

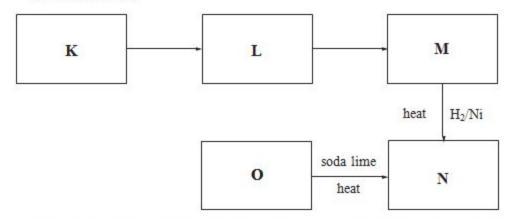
\_\_\_\_\_

(d) Methylbenzene can be oxidised to benzoic acid by heating it strongly with an alkaline solution of reagent R followed by treatment with reagent S. The benzoic acid can then be used to produce a number of other compounds. A reaction sequence is shown below.

PhysicsAndMathsTutor.com

(i) State the name of reagent <b>R</b>	[1]
(ii) State the name of reagent <b>S</b>	[1]
(iii) State the name of reagent <b>T</b> .	[1]
(iv) Give the displayed formula of the organic compound <b>U</b> .	[1]
(e) State and explain how the infrared spectrum of benzoic acid would differ from that of phenylmethanol.	
	[2]
	(Total 12)

 (a) Study the reaction scheme shown below and the other information about compounds K-O that follows:



Compound K has a relative molecular mass of 58.06. It gives an orange-yellow solid with 2, 4-dinitrophenylhydrazine and gives a positive triiodomethane (iodoform) test.

0.500 g of compound **O** in aqueous solution requires 56.75 cm<sup>3</sup> of sodium hydroxide solution of concentration 0.100 mol dm<sup>-3</sup> for complete neutralisation. Compound **O** reacts with sodium hydroxide in a 1:1 molar ratio.

Compound L cannot be oxidised to compound O.

(b) Rhodri prepared benzenecarboxylic acid, C<sub>6</sub>H<sub>5</sub>COOH, by hydrolysing ethyl benzenecarboxylate, C<sub>6</sub>H<sub>5</sub>COOC<sub>2</sub>H<sub>5</sub>.

The overall equation for this hydrolysis is:

$$C_6H_5COOC_2H_5 + H_2O \longrightarrow C_6H_5COOH + C_2H_5OH$$

He used the following method.

- Dissolve 3.20 g of sodium hydroxide in water and make up to 40.0 cm<sup>3</sup>.
- Add the aqueous sodium hydroxide to 2.90 cm<sup>3</sup> of ethyl benzenecarboxylate in a round bottomed flask and reflux for 30 minutes.
- Transfer the mixture into a beaker and add dilute sulfuric acid until the solution is acidic.
- Filter the crystals obtained and recrystallise the benzenecarboxylic acid by dissolving in the minimum amount of hot water.

At the end of the experiment Rhodri's yield of benzenecarboxylic acid was 1.45 g.	
(i) Suggest why Rhodri had to add sulfuric acid before recrystallising.	
	[1]
(ii) State why water is a suitable solvent for the recrystallisation.	
	[1]
(iii) Calculate the concentration, in mol dm <sup>-3</sup> , of the aqueous sodium hydroxide used.	
	[2]
	_
(iv) The density of ethyl benzenecarboxylate is 1.06 g cm-3. Calculate how many moles of ethyl benzenecarboxylate were used.	
	[2]
(v) Calculate the percentage yield obtained by Rhodri.	
	[2]
	_
(vi) Give a reason why the percentage yield was substantially lower than 100 %.	
	[1]
(Total	20)

6.

(a) 2,4-Dinitrophenylhydrazine reagent (2,4-DNP), Tollens' reagent and iodine in sodium hydroxide solution can all be used in the laboratory to identify unknown compounds. Complete the table below by giving any observations made (or writing 'no reaction' as appropriate) when these reagents are added to the compounds listed. [4]

	butan-2-ol	ethanal	ethanol	propanone
2,4-DNP	no reaction			
Tollens' reagent			no reaction	
I <sub>2</sub> /NaOH		8		

(b) Under certain conditions ethanol can be formed from ethene and water. A possible mechanism for this reaction is shown below.

$$CH_2 = CH_2 + H^+ \longrightarrow CH_3CH_2^+ \xrightarrow{H_2O} C_2H_5OH + H^+$$

- (i) Classify this type of mechanism. [1]
- (ii) State the name given to species such as the intermediate ion CH<sub>3</sub>CH<sub>2</sub><sup>+</sup>. [1]
- (iii) Give another reaction of ethene that follows this type of mechanism.

[1]

(iv) Give a reason why the main product of the reaction between propene and water under similar conditions is propan-2-ol.

[1]

(c) Propanone can react with hydrogen cyanide.	
(i) Classify the type of reaction taking place when propanone reacts in this way.	
	[1]
(ii) Draw the mechanism for this reaction	
	[3]
	(Total 12)
	, ,