## **AQA A2 CHEMISTRY**

# TOPIC 4.10 ORGANIC SYNTHESIS AND ANALYSIS

# TOPIC 4.11 STRUCTURE DETERMINATION

**BOOKLET OF PAST EXAMINATION QUESTIONS** 

1. Consider the following reaction sequence.

$$\begin{array}{c|cccc} CH_3 & CH_3 & CH_3 \\ \hline & Step 1 & Step 2 & \\ \hline & NO_2 & NH_2 \\ \hline \\ methylbenzene & E & F \\ \end{array}$$

	(a)		1 /1	1 '	1 .	the reagents	. 1 1
1	•	Hor Stan	I nama tha	machaniem	and atva	tha raggante	100000000
۱	а	i ron aten	1. Harrie the	писсианизни	and give	the reasems	miyoryed.

Name of mechanism	
Reagents	
	(2)
	(3)

(b) For Step 2, give a reagent or combination of reagents. Write an equation for this reaction using [H] to represent the reductant.

D ( - )	
Reagentist	

(c) Give the m/z value of a major peak which could appear in the mass spectrum of methylbenzene, but not in the spectrum of either **E** or **F**.

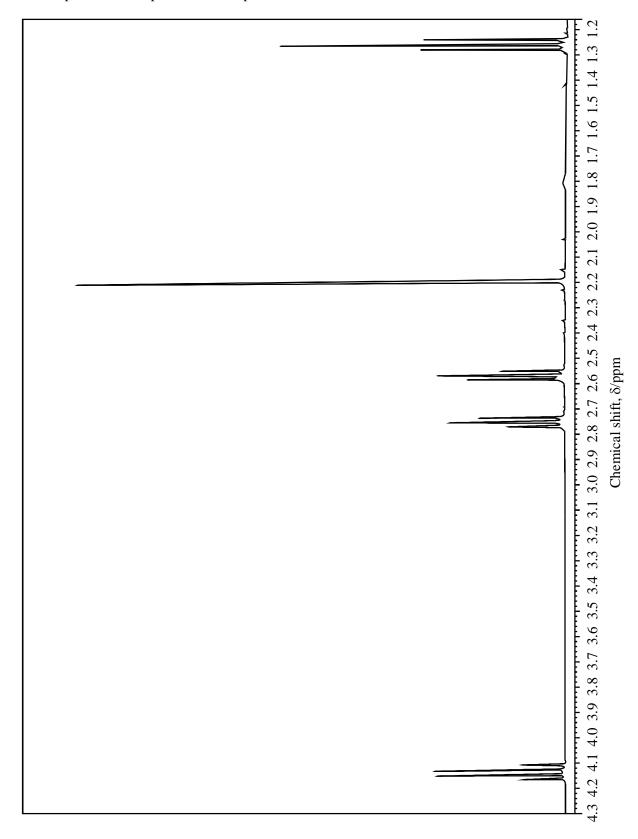
(d) Draw the structure of the species formed by  $\mathbf{F}$  in an excess of hydrochloric acid.

	(e)	(e) Compounds <b>G</b> and <b>H</b> are both monosubstituted benzenes and both are isomers of <b>F</b> . <b>G</b> is a primary amine and <b>H</b> is a secondary amine. Draw the structures of <b>G</b> and <b>H</b> below.					
		G					
		H					
		(Total 9 mar	(2) ks)				
2.	(a)	Write an equation for the formation of methyl propanoate, CH <sub>3</sub> CH <sub>2</sub> COOCH <sub>3</sub> , from methanol and propanoic acid.					
			(1)				
	(b)	Name and outline a mechanism for the reaction between methanol and propanoyl chloride to form methyl propanoate.					
		Name of mechanism					
		Mechanism					

(c)	Propanoic anhydride could be used instead of propanoyl chloride in the preparation of methyl propanoate from methanol. Draw the structure of propanoic anhydride.						
			(1)				
(d)	(i)	Give <b>one</b> advantage of the use of propanoyl chloride instead of propanoic acid in the laboratory preparation of methyl propanoate from methanol.					
	(ii)	Give <b>one</b> advantage of the use of propanoic anhydride instead of propanoyl chloride in the industrial manufacture of methyl propanoate from methanol.					
(e)		ester contains a benzene ring. The mass spectrum of this ester shows a molecular ion at $m/z = 136$ .	(2)				
	(i)	Deduce the molecular formula of this ester.					
	(ii)	Draw <b>two</b> possible structures for this ester.					
			(3)				
		(Total 12 ma	ırks)				

		Cl	
(a)	Nam	ne compound U.	
(b)	(i)	State why the mass spectrum of <b>U</b> contains two molecular ion peaks.	(1)
	(ii)	Give the $m/z$ values of these two peaks.	
			(2)
(c)	Nam	ne and outline a mechanism for the reaction of U with CH <sub>3</sub> OH	
	Nam	e of mechanism	
	Мес	hanism	

(5) (Total 8 marks) **4.** The proton n.m.r. spectrum of compound  $\mathbf{X}$  is shown below.



Compound X,  $C_7H_{12}O_3$ , contains both a ketone and an ester functional group. The measured integration trace for the peaks in the n.m.r. spectrum of X gives the ratio shown in the table below.

Chemical shift, δ/ppm	4.13	2.76	2.57	2.20	1.26
Integration ratio	0.8	0.8	0.8	1.2	1.2

Refer to the spectrum, the information given above and the data below the Periodic Table provided to answer the following questions.

W	That is the whole-number ratio of each type of proton in compound $\mathbf{X}$ ?
•••	
	raw the part of the structure of $\mathbf{X}$ which can be deduced from the presence of the peak at 2.20.
••••	
	ne peaks at $\delta 4.13$ and $\delta 1.26$ arise from the presence of an alkyl group. Identify the group ad explain the splitting pattern.
Al	kyl group
Ех	xplanation
•••	
•••	
	raw the part of the structure of $\mathbf{X}$ which can be deduced from the splitting of the peaks at 2.76 and $\delta 2.57$ .
•••	
D٤	educe the structure of compound $\mathbf{X}$ .

(a) The infra-red spectrum of $\mathbf{Q}$ contains a major absorption at 1724 cm <sup>-1</sup> . Identify responsible for this absorption.					<sup>1</sup> . Identify the bond		
b)		The mass spectrum of <b>Q</b> contains two molecular ion peaks at $m/z = 106$ and $m/z = 108$ . It also has a major peak at $m/z = 43$ .					
	(i)	Suggest why there are two	-				
	(ii)	A fragment ion produced frelements. Identify this frag	from <b>Q</b> has $m/z = 43$ from the ion and write	3 and contains ato	ms of <b>three</b> different		
		Fragment ion					
Equation							
c)	The proton n.m.r. spectrum of <b>Q</b> was recorded.						
	(i)	Suggest a suitable solvent	for use in recording	g this spectrum of	Q.		
	(ii)	Give the formula of the stan spectra.	dard reference con				
d)		proton n.m.r. spectrum of <b>Q</b> ber of adjacent, non-equivale	ent protons respons	Complete the tab	le below to show the		
			Peak 1	Peak 2	Peak 3		
Ir	Integration value		3	3	1		
S	Splitting pattern		doublet	singlet	quartet		
			1	· · · · · · · · · · · · · · · · · · ·			

5.

(e) Using the information in parts (a), (b) and (d), deduce the structure of compound  $\mathbf{Q}$ .

	(f)	A structural isomer of ${\bf Q}$ reacts with cold water to produce misty fumes. Suggest a structure for this isomer.	
		(Total 9 ma	(1) arks)
6.	(a)	Draw the structure of ethyl propanoate.	
			(1)
	(b)	Name and outline a mechanism for the formation of ethyl propanoate from propanoyl chloride and ethanol.	
		Name of mechanism	
		Mechanism	
			(5)
			(3)
	(c)	The mass spectrum of ethyl propanoate contains a major peak at $m/z = 57$ . Write an equation showing the fragmentation of the molecular ion to form the species responsible for the peak at $m/z = 57$ . Show the structure of this species in your answer.	
			(2)

	(d) Draw the structure of another ester which is an isomer of ethyl propanoate and whice a major peak at $m/z = 71$ in its mass spectrum.				
			(Total 9 ma	(1) arks)	
7.	(a)	(i)	Using mass spectrometry, what information in addition to $M_r$ can be obtained from the precise value of the mass of the molecular ion of an organic compound?		
		(ii)	Suggest why, for most organic compounds, it is possible to detect a peak at one mass unit higher than that of the molecular ion.		
				(2)	
	(b)	_	mentation of the molecular ion of methylbutanone, $(CH_3)_2CHCOCH_3$ , gives rise to mant peaks at $m/z = 71$ and $m/z = 43$ .		
		(i)	Construct a balanced equation to show how fragmentation of the molecular ion gives rise to the peak at $m/z = 71$ .		
		(ii)	In the fragmentation of $(CH_3)_2CHCOCH_3$ , two fragments with $m/z$ 43 are formed. Give the structural formula of each of these fragments.		
			Structure 1 Structure 2		
				(5)	
	(c)	and r	molecular ion peaks appear in the mass spectrum of 2-chloropropane at $m/z$ 78 $m/z = 80$ . Explain why two molecular ion peaks are found and why the relative sities of the peaks are approximately 3 to 1, respectively.		
				(2)	
			(Total 10 m	(3) arks)	

8. Three sections of the proton n.m.r. spectrum of CH<sub>3</sub>CHClCOOH are shown below. 12.0 11.0 10.0 4.6 4.5 4.4 4.3 1.8 1.7 1.6  $\delta \, / ppm$  $\delta\,/ppm$  $\delta\,/ppm$ Name the compound  $CH_3CHClCOOH$ (a) **(1)** Explain the splitting patterns in the peaks at  $\delta$  1.72 and  $\delta$  4.44 (b) **(2)** Predict the splitting pattern that would be seen in the proton n.m.r. spectrum of the isomeric (c) compound ClCH2CH2COOH

**(1)** 

(d) The amino acid *alanine* is formed by the reaction of CH<sub>3</sub>CHClCOOH with an excess of ammonia. The mechanism is nucleophilic substitution. Outline this mechanism, showing clearly the structure of *alanine*.

**(5)** 

(e) The amino acid *lysine* has the structure

$$H_2N - (CH_2)_4 - C - COOH$$
 $NH_2$ 

Draw structures to show the product formed in each case when lysine reacts with

(i) an excess of aqueous HCl,

(ii) an excess of aqueous NaOH,

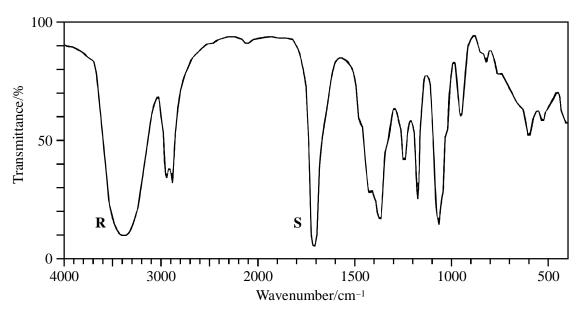
(iii) another molecule of lysine.

9.	(a)	(i)	Write an equation for the reduction of pentan-2-one by aqueous NaBH <sub>4</sub> to pentan-2-ol.	form
			Use [H] to represent the reductant.	
		(ii)	Name and outline a mechanism for this reduction.	
			Name of mechanism	
			Mechanism	
		(iii)	State why the pentan-2-ol produced in this reaction is not optically active.	
				(7)
	(b)		lict the $m/z$ values of the two most abundant fragments in the mass spectrum an-2-one.	
		Frag	gment 1	
		Frag	gment 2	
				(2) (Total 9 marks)

**10.** Spectral data for use in this question are provided below the Periodic Table (first item on the database).

Compound  $\mathbf{Q}$  has the molecular formula  $C_4H_8O_2$ 

(a) The infra-red spectrum of  $\mathbf{Q}$  is shown below.



Identify the type of bond causing the absorption labelled  ${\bf R}$  and that causing the absorption labelled  ${\bf S}$ .

R	
S	
	(2)

(b)  $\mathbf{Q}$  does not react with Tollens' reagent or Fehling's solution. Identify a functional group which would react with these reagents and therefore cannot be present in  $\mathbf{Q}$ .

(1)

- (c) Proton n.m.r. spectra are recorded using a solution of a substance to which tetramethylsilane (TMS) has been added.
  - (i) Give two reasons why TMS is a suitable standard.

Reason 2 .....

(ii) Give an example of a solvent which is suitable for use in recording an n.m.r. spectrum. Give a reason for your choice.

Solvent .....

Reason .....

**(4)** 

(d) The proton n.m.r. spectrum of  $\mathbf{Q}$  shows 4 peaks.

The table below gives  $\delta$  values for each of these peaks together with their splitting patterns and integration values.

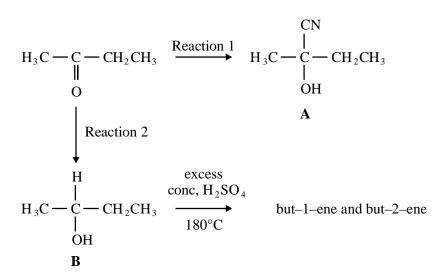
δ/ppm	2.20	2.69	3.40	3.84
Splitting pattern	singlet	triplet	singlet	triplet
Integration value	3	2	1	2

What can be deduced about the structure of  $\mathbf{Q}$  from the presence of the following in its n.m.r. spectrum?

(i)	The singlet peak at $\delta = 2.20$	
(ii)	The singlet peak at $\delta = 3.40$	
(iii)	Two triplet peaks	
		(3)

(e) Using your answers to parts (a), (b) and (d), deduce the structure of compound **Q**.

(1) (Total 11 marks) 11. Consider the reaction scheme below which starts from butanone.



(a) (i) Give a suitable reagent for Reaction I and state the type of mechanism involved.

Type of mechanism.

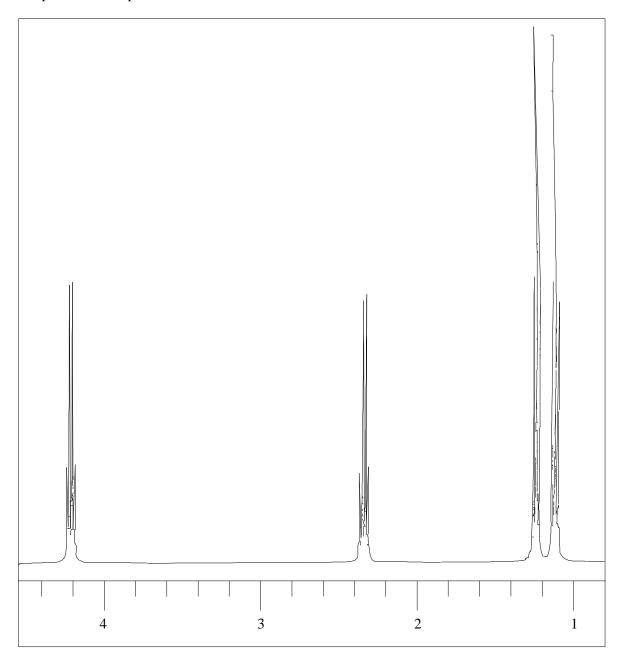
(ii) When 5.0 g of butanone were used to carry out Reaction 1, the yield was 64% of the theoretical maximum. Calculate the mass of compound A formed.

		•••••		••••••	
•••••	•••••	•••••	•••••	••••••	•••••
				•••••	

**(6)** 

(b)	(i)	Give a suitable reagent for Reaction 2.					
	(ii)	Product <b>B</b> is found to be optically inactive. Suggest why this is so.					
	(iii)	Given the table of infra-red absorption data below, describe the two major differences between the infra-red spectra of butanone and of <b>B</b> .					
(c)	Outli	ne a mechanism for the formation of but-1-ene by the dehydration of <b>B</b> .					
		(4					
(d)	Draw but-1	the structure of the repeating unit in the polymer which can be formed from ene.					
		(1 (Total 15 marks					

12. The proton n.m.r. spectrum of an ester,  $\mathbf{A}$ , is shown below.



Chemical shift,  $\delta/ppm$ 

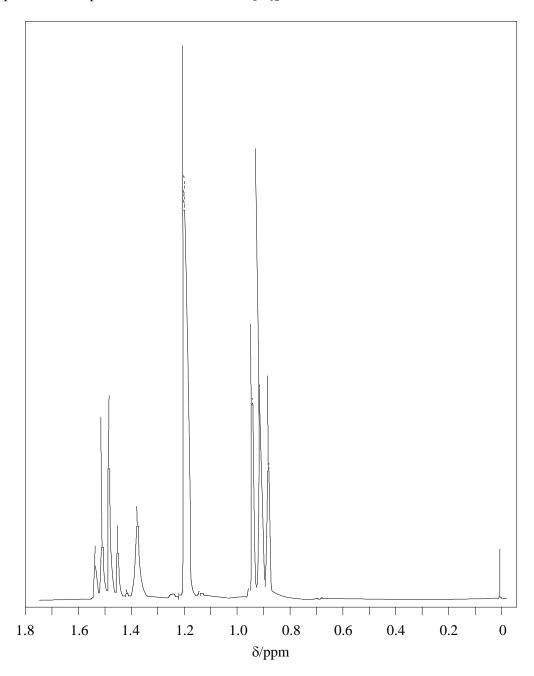
The measured integration trace gives the ratio 0.50 to 0.50 to 0.75 to 0.75 for the peaks at  $\delta$  4.13, 2.32, 1.33 and 1.09, respectively.

(a)	The mass spectrum of compound <b>A</b> has a molecular ion peak at $m/z = 102$ . Deduce the molecular formula of compound <b>A</b> .					

**(3)** 

(b)	What is the ratio of the no	umbers of eac	ch type of prot	ton?		
						(1)
(c)	What can be deduced aboun.m.r. spectrum?	out the arrang	ement of prote	ons from the s <sub>l</sub>	plitting pattern	s in the
						(1)
(d)	Deduce the structure of confequivalent protons.	ompound <b>A</b> a	nd label with	the letters a, b,	c and d the fo	ur groups
(e)	Assign the various signal		below to the §	groups of proto	ons in compou	(1) and ${f A}$
	previously labelled a, b, c	and d.				
	Chemical shift, δ/ppm	1.09	1.33	2.32	4.13	
	Label of group					
				J		(4)
					(	(Total 10 marks)

### 13. The proton n.m.r. spectrum of an alcohol, $\mathbf{A}$ , $C_5H_{12}O$ , is shown below



The measured integration trace gives the ratio 0.90 to 0.45 to 2.70 to 1.35 for the peaks at  $\delta$  1.52, 1.39, 1.21 and 0.93, respectively.

(a)	What compound is responsible for the signal at $\delta$ 0?	
		(1)
(b)	How many different types of proton are present in compound A?	
		(1)

	(c)	What	t is the ratio of the numbers of each type of proton?	
		•••••		(1)
	(d)		beaks at $\delta$ 1.52 and $\delta$ 0.93 arise from the presence of a single alkyl group. Identify this p and explain the splitting pattern.	
		Grou	p	
		Explo	anation	
				(3)
	(e)	What	t can be deduced from the single peak at $\delta$ 1.21 and its integration value?	
				(1)
	(f)	Give	the structure of compound A.	
			(Total 8 i	(1) marks)
14.			ester formed when ethanol reacts with ethanoic anhydride was analysed by resolution proton n.m.r. spectroscopy.	
		(i)	Give the structural formula of the ester.	
		(ii)	How many different types of proton are present in this ester?	
		(iii)	Describe the splitting pattern of the ethyl group in the n.m.r. spectrum of the ester.	
			(Total 3 r	(3) marks)

a)	Defi	ne the term structural isomerism.
b)	The	graph below shows part of the mass spectrum of an organic compound, ${f A}$ , which has
		nolecular formula C <sub>4</sub> H <sub>10</sub> .
		ative adance
	(i)	0 10 20 30 40 50 60 m/e Draw graphical formulae for the structural isomers of $C_4H_{10}$ .
	( )	Isomer 1 Isomer 2
	<i>(</i> '')	
	(ii)	Suggest the formula of the fragment that corresponds to each of the following m/e values shown in the mass spectrum of compound $\bf A$ .
		43
		29
		15
	(iii)	Deduce which of the isomers drawn in (i) corresponds to compound <b>A</b> . Give a reason for your answer.

(c)	(1)		gle functional group. The functional g	
		Isomer 3	Isomer 4	
				(2)
	(ii)	Give the name of the type of compo	and that each isomer represents.	
		Isomer 3		
		Isomer 4		(2)
				(Total 13 marks)

(a)  $\bf A$  and  $\bf B$  have the molecular formula  $C_2H_4O_2$ . An aqueous solution of  $\bf A$  reacts with calcium carbonate to liberate carbon dioxide, but  $\bf B$  does not.

A B

**(2)** 

(b)	<b>C</b> and <b>D</b> have the molecular formula $C_3H_8O$ . <b>C</b> has a broad absorption band at $3300 \text{ cm}^{-1}$ in its infra-red spectrum, but <b>D</b> does not.			
	C	D		
			(2)	
(2)	E and E have the male culor formula	C II C1  E has only one made in its law masslytics	(2)	
(c)	proton n.m.r. spectrum but <b>F</b> has tw	a $C_3H_6C1_2$ . <b>E</b> has only one peak in its low resolution vo.		
	E	${f F}$		
			(2)	
			(2)	
(d)	<b>G</b> and <b>H</b> have the molecular formul	a C <sub>3</sub> H <sub>6</sub> O and both have strong absorption bands at		
,		pectra. <b>G</b> turns acidified potassium dichromate(VI)		
	G	H		

**(2)** 

(e) <b>I</b> and <b>J</b> have the molecular formula $C_4H_9Br$ . <b>I</b> shows optical activities		${f I}$ and ${f J}$ have the molecular formula $C_4H_9Br.$ ${f I}$	hows optical activity, but ${f J}$ does not.
		I J	
			(2)
	(f)	<b>K</b> and <b>L</b> have the molecular formula $C_5H_{10}$ . <b>K</b> $1650 \text{ cm}^{-1}$ in its infra-red spectrum, but <b>L</b> does	
		K L	
			(2) (Total 12 marks)
17.	(a)	Name and outline a mechanism for the reaction	between propanone and hydrogen cyanide.
		Name	
		Mechanism	

**(5)** 

(i)		tion of propanone to the corresponding alcohol and nt. In the equation you may use the symbol [H] for
	Equation	
	Reducing agent	
(ii)	Two isomeric carbonyl compou have 3 peaks in their low-resolu	nds, $\bf A$ and $\bf B$ , with molecular formula $C_4H_8O$ both tion proton n.m.r. spectra.
		which has 5 peaks in its low-resolution proton n.m.r. <b>B</b> forms alcohol <b>D</b> which has 4 peaks in its extrum.
	Draw one possible structure for	each of these 4 compounds.
	Carbonyl compound <b>A</b>	Alcohol C
	Carbonyl compound <b>B</b>	Alcohol <b>D</b>
		(6) (Total 11 marks)

Carbonyl compounds can be reduced to alcohols.

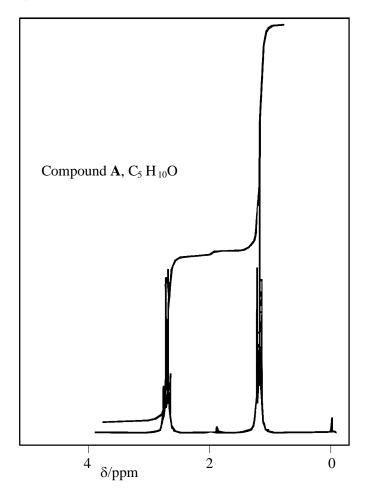
(b)

**18.** Compound **A**, C<sub>5</sub>H<sub>10</sub>O, reacts with NaBH<sub>4</sub> to give **B**, C<sub>5</sub>H<sub>12</sub>O. Treatment of **B** with concentrated sulphuric acid yields compound **C**, C<sub>5</sub>H<sub>10</sub>. Acid-catalysed hydration of **C** gives a mixture of isomers, **B** and **D**.

Fragmentation of the molecular ion of  $\mathbf{A}$ ,  $[C_5H_{10}O]^{+\bullet}$ , leads to a mass spectrum with a major peak at m/z 57. The infra-red spectrum of compound  $\mathbf{A}$  has a strong band at 1715 cm<sup>-1</sup> and the infra-red spectrum of compound  $\mathbf{B}$  has a broad absorption at 3350 cm<sup>-1</sup> (Table). The proton n.m.r. spectrum of  $\mathbf{A}$  has two signals at  $\delta$  1.06 (triplet) and 2.42 (quartet), respectively (Spectrum).

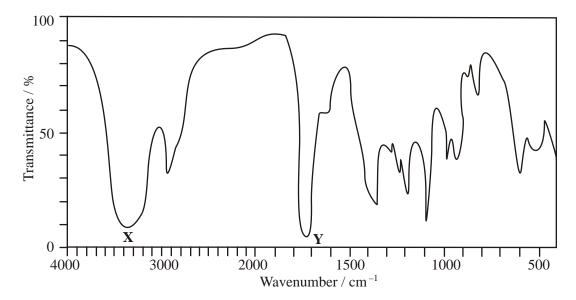
Use the analytical and chemical information provided to deduce structures for compounds A, B, C and D, respectively. Include in your answer an equation for the fragmentation of the molecular ion of A and account for the appearance of the proton n.m.r. spectrum of A. Explain why isomers B and D are formed from compound C.

#### **Spectrum**



(Total 20 marks)

### **19.** (a) The infra-red spectrum of compound A, $C_3H_6O_2$ , is shown below.



Identify the functional groups which cause the absorptions labelled X and Y.

Using this information draw the structures of the three possible structural isomers for A.

Label as **A** the structure which represents a pair of optical isomers.

**(6)** 

(b) Draw the structures of the three **branched-chain** alkenes with molecular formula  $C_5H_{10}$ 

Draw the structures of the three dibromoalkanes, C<sub>5</sub>H<sub>10</sub>Br<sub>2</sub>, formed when these three alkenes react with bromine.

One of these dibromoalkanes has only three peaks in its proton n.m.r. spectrum. Deduce the integration ratio and the splitting patterns of these three peaks.

(10)

(Total 16 marks)

- **20.** (a) The reaction of but-1-ene with chlorine produces 1,2-dichlorobutane,  $C_4H_8Cl_2$ 
  - (i) Given that chlorine exists as a mixture of two isotopes,  $^{35}$ Cl and  $^{37}$ Cl, predict the number of molecular ion peaks and their m/z values in the mass spectrum of  $C_4H_8Cl_2$
  - (ii) The mass spectrum of 1,2-dichlorobutane contains peaks at m/z = 77 and 79. Draw the structure of the fragment ion which produces the peak at m/z = 77 and write an equation showing its formation from the molecular ion.

**(6)** 

- (b) The reaction of but-2-ene with hydrogen chloride forms a racemic mixture of the stereoisomers of 2-chlorobutane.
  - (i) Name the type of stereoisomerism shown by 2-chlorobutane and give the meaning of the term *racemic mixture*. State how separate samples of the stereoisomers could be distinguished.
  - (ii) By considering the shape of the reactive intermediate involved in the mechanism of this reaction, explain how a racemic mixture of the two stereoisomers of 2-chlorobutane is formed.

**(7)** 

- (c) The reaction of but-2-ene with chlorine produces 2,3-dichlorobutane, C<sub>4</sub>H<sub>8</sub>Cl<sub>2</sub>
  - (i) State the number of peaks, their integration ratio and any splitting of peaks in the proton n.m.r. spectrum of 2,3-dichlorobutane.
  - (ii) Compound **S**, an isomer of C<sub>4</sub>H<sub>8</sub>Cl<sub>2</sub>, produces a proton n.m.r. spectrum which consists only of a singlet, a triplet and a quartet with an integration ratio of 3:3:2 respectively.

Compound **T**, also an isomer of C<sub>4</sub>H<sub>8</sub>Cl<sub>2</sub>, produces a proton n.m.r. spectrum which consists only of two singlets with an integration ratio of 3:1

Draw the structures of **S** and of **T**.

(6)

(Total 19 marks)

- 21. Butenedioic acid, HOOCCH=CHCOOH, occurs as two stereoisomers. One of the isomers readily forms the acid anhydride  $C_4H_2O_3$  when warmed.
  - (a) Draw the structures of the two isomers of butenedioic acid and name the type of isomerism shown.
    - Use the structures of the two isomeric acids to suggest why only one of them readily forms an acid anhydride when warmed. Draw the structure of the acid anhydride formed.

**(6)** 

(b) Identify one electrophile which will react with butenedioic acid and outline a mechanism for this reaction.

**(4)** 

(c) Write an equation for a reaction which occurs when butenedioic acid is treated with an excess of aqueous sodium hydroxide.

**(2)** 

(d) Describe and explain the appearance of the proton n.m.r. spectrum of butenedioic acid.

(3)

(Total 15 marks)

**22.** Propanoyl chloride can be used, together with a catalyst, in Step 1 of the synthesis of 1-phenylpropene from benzene via compounds **P** and **Q** as shown below.

(a) The mechanism of Step 1 is an electrophilic substitution. Write an equation to show the formation of the electrophile from propanoyl chloride. Outline the mechanism of the reaction of this electrophile with benzene in Step 1.

(b) The mass spectrum of **P** contains a molecular ion peak at m/z = 134 and major fragmentation peaks at m/z = 105 and 77. Identify the species responsible for the peak at m/z = 105 and also that responsible for the peak at m/z = 77. Write an equation for the formation, from the molecular ion, of the species responsible for the peak at m/z = 105.

- (c) NaBH<sub>4</sub> can be used in the reaction in Step 2. Name the mechanism involved in this reaction. Molecules of **Q** show optical isomerism but the sample of **Q** formed in Step 2 is optically inactive. State, in terms of their structure, why molecules of **Q** show optical isomerism. Explain, by reference to the mechanism, why the sample of **Q** obtained in Step 2 is not optically active.
- (d) Identify a suitable reagent for the reaction in Step 3.

  Name the type of stereoisomerism shown by the product of this reaction. State what is required in the structure of molecules to allow them to show this type of stereoisomerism.

  (4)

  (Total 20 marks)

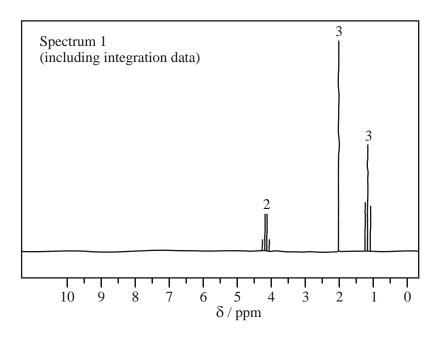
**(5)** 

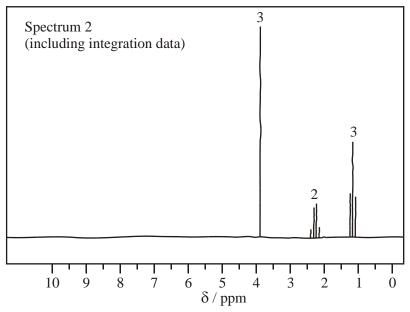
**(7)** 

**23.** (a) Ester **X**, CH<sub>3</sub>CH<sub>2</sub>COOCH<sub>3</sub>, can be produced by the reaction between propanoyl chloride and methanol. Name **X** and outline a mechanism for this reaction. Name the mechanism involved.

**(6)** 

(b) The proton n.m.r. spectrum of **X** is shown below together with that of an isomeric ester, **Y**. Deduce which of Spectrum 1 and Spectrum 2 is that obtained from **X**. Use **Table 1** on the Data Sheetand the integration data on the spectra to help you to explain your deduction. Suggest a structure for **Y**.





(4) (Total 10 marks)

24.		three compounds CH <sub>3</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH, (CH <sub>3</sub> ) <sub>3</sub> COH and CH <sub>3</sub> CH <sub>2</sub> CH <sub>2</sub> CHO can be aguished by use of the following three reagents	
		<ol> <li>potassium dichromate(VI) acidified with dilute sulphuric acid</li> <li>Tollens' reagent</li> <li>ethanoic acid, together with a small amount of concentrated sulphuric acid.</li> </ol>	
	(a)	Identify which of these three organic compounds would reduce acidified potassium dichromate(VI). Give the structures of the organic products formed. Write a half-equation for the reduction of dichromate(VI) ions in acidic solution.	(6)
	(b)	Identify which one of these three organic compounds would reduce Tollens' reagent. Give the structure of the organic product formed. Write a half-equation for the reduction of Tollens' reagent.	(3)
	(c)	Identify which of these three organic compounds would react with ethanoic acid in the presence of concentrated sulphuric acid. In each case, give the structure of the organic product formed.	
	(d)	State the number of peaks in the proton n.m.r. spectra of CH <sub>3</sub> CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> OH and of (CH <sub>3</sub> ) <sub>3</sub> COH. (Analysis of peak splitting is not required.)	(4)
		(Total 15 ma	(2) irks)
25.	(a)	Draw the structure of each of the three ketones which have the molecular formula $C_5H_{10}O$ . For each compound give the ratio of the areas under each peak in its low-resolution proton n.m.r. spectrum.	(6)
	(b)	Draw the structure of each of the four aldehydes which also have the molecular formula $C_5H_{10}O$ . Label with the letter $\mathbf{X}$ the compound which has only two peaks in its low-resolution proton n.m.r. spectrum. Label with the letter $\mathbf{Y}$ the compound which has five peaks with the ratios of the areas under each peak 3:3:2:1:1 in its low-resolution proton n.m.r. spectrum. Label with the letter $\mathbf{Z}$ the compound which shows optical isomerism.	(7)
	(c)	When carbonyl compounds react with HCN, racemic mixtures are often produced. Name the type of mechanism involved and explain what is meant by the term <i>racemic mixture</i> . Choose any carbonyl compound which does <b>not</b> form a racemic mixture when it reacts with HCN and draw the structure of the product formed by the reaction of this carbonyl compound with HCN.	
			<b>(4)</b>

Explain why aldehydes react with Tollen's (or Fehling's) reagent but ketones do not.

(d)

32

(3) (Total 20 marks) **26.** (a) Describe, by giving reagents and stating observations, how you could distinguish between the compounds in the following pairs using a simple test-tube reaction for each pair.

(i) 
$$\begin{array}{c} CH_3 \\ H_3C-C-OH \\ H \\ \end{array}$$
 and  $\begin{array}{c} CH_3 \\ H_3C-C-C+OHO \\ H \\ \end{array}$   $\begin{array}{c} CH_3 \\ H \\ H \\ \end{array}$   $\begin{array}{c} CH_3 \\ H \\ \end{array}$ 

(ii) and 
$$H_3C$$
  $C = C < CH_3$   $C = C$   $CH_3$ 

(b) State how compounds **E** and **F** in part (a)(iii) above could be distinguished by their infra-red spectra, without using the fingerprint region. Explain how *fingerprinting* is used to identify a compound.

(c) Suggest the structure of the fragment responsible for the major peak in the mass spectrum of compound  $\mathbf{E}$  and state its m/z value. Write an equation showing the formation of this fragment from the molecular ion.

(4) (Total 15 marks)

**(8)** 

**(3)** 

27. Use the data given on the back of the Periodic Table (PT) to help you answer this question. Compounds  $\bf A$  to  $\bf G$  are all isomers with the molecular formula  $C_6H_{12}O_2$ 

Isomer A, C<sub>6</sub>H<sub>12</sub>O<sub>2</sub>, is a neutral compound and is formed by the reaction between compounds X and Y in the presence of a small amount of concentrated sulphuric acid. X and Y can both be formed from propanal by different redox reactions.
 X has an absorption in its infra-red spectrum at 1750 cm<sup>-1</sup>.

Deduce the structural formulae of A, X and Y. Give suitable reagents, in each case, for the formation of X and Y from propanal and state the role of concentrated sulphuric acid in the formation of A.

(b) Isomers **B**, **C**, **D** and **E** all react with aqueous sodium carbonate to produce carbon dioxide. Deduce the structural formulae of the three isomers that contain an asymmetric carbon atom.

The fourth isomer has only three singlet peaks in its proton n.m.r. spectrum. Deduce the structural formula of this isomer and label it **E**.

**(4)** 

**(7)** 

(c) Isomer **F**, C<sub>6</sub>H<sub>12</sub>O<sub>2</sub>, has the structural formula shown below, on which some of the protons have been labelled.

$$a$$
 $CH_3$ — $CH_2$ — $O$ — $CH_2$ — $CH_2$ — $C$ — $CH_3$ 

A proton n.m.r. spectrum is obtained for **F**. Using Table 1 at the back of the Periodic Table (PT), predict a value of  $\delta$  for the protons labelled a and also for those labelled b. State and account for the splitting patterns of the peaks assigned to the protons a and b.

**(6)** 

(d) Isomer  $\mathbf{G}$ ,  $C_6H_{12}O_2$ , contains six carbon atoms in a ring. It has an absorption in its infra-red spectrum at 3270 cm<sup>-1</sup> and shows only three different proton environments in its proton n.m.r. spectrum. Deduce a structural formula for  $\mathbf{G}$ .

**(2)** 

(Total 19 marks)

28. The conversion of compound A into compound B can be achieved in two steps as shown below.

The intermediate compound, **X**, has an absorption at 1650 cm<sup>-1</sup> in its infra-red spectrum.

(a) Identify compound **X**. Explain your answer.

**(2)** 

(b) For each step in this conversion, give the reagents and essential conditions required and outline a mechanism.

(11)

(c) Show how the number of peaks in their proton n.m.r. spectra would enable you to distinguish between compounds **A** and **B**.

**(2)** 

(Total 15 marks)

29.	sing	le com	pound $\mathbf{B}$ , $C_{10}H_{12}O$ . Com	pound <b>B</b> is converted into <b>C</b>	cylbenzene $\mathbf{A}$ , $C_8H_{10}$ , to form a $C$ , $C_{10}H_{14}O$ , on treatment with d with concentrated $H_2SO_4$ .	
				on band in the infra-red at 168 bund <b>D</b> has an absorption ba	$35 \text{ cm}^{-1}$ , compound $\mathbb{C}$ has a broad and close to $1630 \text{ cm}^{-1}$ .	1
	(a)	used	to deduce structures for	rovided in the question and t compounds <b>A</b> , <b>B</b> , <b>C</b> and <b>D</b> . he fact that B is obtained as a	he data in the Table below can be	e
		Lxpi	an the significance of the	ic fact that B is obtained as a	i single compound.	(9)
	(b)			aking place in each step and om compound C and concen	outline a mechanism for the trated sulphuric acid.	(7)
	(c)	Expl	ain briefly why compou	nd C can exist in stereoisom	eric forms.	(2)
					(Total 18	
30.	Drote	on n m	r spectroscopy can be	used to distinguish between t	ha fallowing isomore:	
30.	F100	JII II.III		-	-	
			HOCH <sub>2</sub> CH <sub>2</sub> OCH <sub>3</sub> <b>A</b>	HOCH <sub>2</sub> OCH <sub>2</sub> CH <sub>3</sub> <b>B</b>	CH <sub>3</sub> OCH <sub>2</sub> OCH <sub>3</sub>	
	and	accoun			resolution proton n.m.r. spectrum ons are not coupled to any other	1
	prote	ons.			(Total 15	marks)
31.	(a)	nitra		phuric acid and concentrate	abstitution. The reagents for this d nitric acid in a 1:1 ratio and the	
		(i)	Explain what is meant	by the term electrophile and	I the term substitution.	(2)
		(ii)		to show its formation from t	olved in this reaction and write and he reagents. Write a mechanism	
						(6)
		(iii)	Explain why the tempo 60°C.	erature of the reaction should	d not be allowed to rise above	
						(1)
	(b)		robenzene is also prepar ition(s) required for this	-	ion. State the reagent(s) and	

(3)

	OI III	olecular formula	C <sub>4</sub> H <sub>9</sub> Cl, are given below	•		
	CH <sub>3</sub>	CH <sub>2</sub> CH <sub>2</sub> CH <sub>2</sub> Cl	CH <sub>3</sub> CH(Cl)CH <sub>2</sub> CH <sub>3</sub>	(CH <sub>3</sub> ) <sub>2</sub> CHCH <sub>2</sub> Cl	$(CH_3)_2C(Cl)CH_3$	
		A	В	C	D	
			IR analysis of compound with 3 peaks and one sp			
	(i)	Give the name of	of compound <b>D</b> .			(1)
	(ii)	Identify the con Explain your re	npound ( <b>A</b> , <b>B</b> , <b>C</b> or <b>D</b> ) wasoning.	hose NMR spectrum s	shows a single peak.	(2)
	(iii)	•	appound (A, B, C or D) who of the areas under these	•	-	(3)
	(iv)	elimination read	with an alcoholic solution ction. Two structurally is lae of these two structura	someric products are f		(2)
	(v)		cal formula for the produce an outline of the mech			(4)
(a)	spec	trometer, can be r $(M-R)^{+\bullet} \to M^+$	a molecular ion $(M-R)^{+\bullet}$ , epresented by the equation $+R^{\bullet}$ s of species shown in the	on		
		conversion.		<b>1</b>	F	(4)
(b)	66. T	The peak at $m/z = 0$ observation and sh	chloroethane shows two 64 is approximately three now, by means of an equa to a peak at an $m/z$ value	times as intense as thation, how the molecul	at at $m/z = 66$ . Explain	(4)
(c)	Sugg and	•	spectrum of 1,2-dichloro	ethane shows peaks at	m/z values of 98, 100	
	anu .	111/.				

32.

33. A molecular ion peak at mlz = 86 is present in the mass spectrum of each of three organic compounds, **A**, **B** and **C**. Each compound has only two peaks in its low-resolution proton n.m.r. spectrum.

A is a hydrocarbon.

 $\bf B$  and  $\bf C$  are isomers and both have a strong absorption near 1700 cm<sup>-1</sup> in their infra-red spectra.

A table of infra-red absorption data is provided below.

(a) For each compound, give the molecular formula and draw a possible structure. Deduce the ratio of the peak areas in the low-resolution proton n.m.r. spectrum of each compound.

**(9)** 

(b) Describe how you could distinguish between **B** and **C** using a chemical test. How could **B** or **C** be identified using its infra-red spectrum?

**(5)** 

(c)  $\mathbf{D}$  and  $\mathbf{E}$  are isomers of  $\mathbf{B}$  and  $\mathbf{C}$ .

**D** has infra-red absorptions at 3300 cm<sup>-1</sup> and also at 1650 cm<sup>-1</sup>.

**E** has an infra-red absorption at 3300 cm<sup>-1</sup> but none near 1650 cm<sup>-1</sup>.

Draw and label one possible structure each for **D** and **E**. Explain your answer.

**(4)** 

(Total 18 marks)

**34.** (a) Explain what is meant by the fingerprint region of an infra-red spectrum and describe how it is used to confirm the identity of organic molecules.

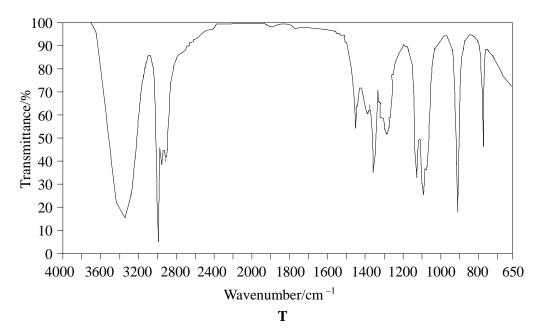
**(4)** 

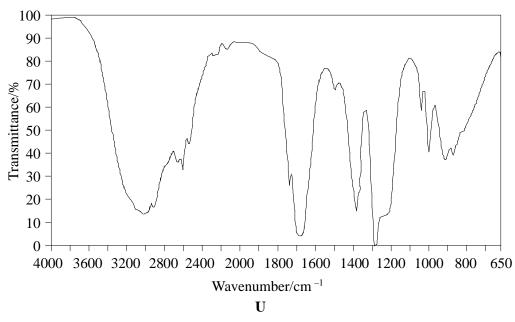
(b) Each of three isomeric esters,  $\mathbf{Q}$ ,  $\mathbf{R}$  and  $\mathbf{S}$ , of molecular formula  $C_5H_{10}O_2$  has a branched structure.

 $\bf Q$  has only two peaks in its low resolution proton n.m.r. spectrum, but  $\bf R$  and  $\bf S$  each have three peaks.

Hydrolysis of  ${\bf R}$  in acid conditions forms compounds  ${\bf T}$  and  ${\bf U}$ . The low-resolution proton n.m.r. spectrum of  ${\bf T}$  has three peaks and that of  ${\bf U}$  has two peaks. Infra-red spectra of  ${\bf T}$  and  ${\bf U}$  are shown below.

Draw structures of the compounds **Q**, **R**, **S**, **T** and **U** and explain how the n.m.r. and i.r. data support your conclusions. A table of infra-red absorption data is provided below.





(11) (Total 15 marks)

<b>35.</b>	Simple organic compounds, including structural isomers, can be distinguished from each other by
	chemical tests, or by spectroscopic methods, or by a combination of both. This question concerns
	the four compounds whose structural formulae are shown below.

 $CH_3(CH_2)_2CHO$   $(CH_3)_2CHCHO$   $CH_3COC_2H_5$   $CH_3COOC_2H_5$  **A B C D** 

(a) Draw graphical formulae for **A**, **B**, **C** and **D**, clearly indicating which is which. Give the names of **A**, **B**, **C** and **D**.

(8)

(b) Compound **D** can be made by a reaction called esterification. State the names of **two** compounds needed for the reaction and give a condition used in the reaction.

**(3)** 

(c) Give the name and draw the graphical formula of an isomer of compound **D** that is **not** an ester.

**(2)** 

(d) Describe **one** chemical test that would distinguish between **B** and **C**, and state what you would observe in each case.

**(3)** 

(e) The <sup>1</sup>H NMR spectra of **A**, **B**, **C** and **D** can be used to distinguish between some of the structures by considering the numbers of peaks and the ratios of the areas under them. The following table shows some results of a <sup>1</sup>H NMR investigation into **A**, **B**, **C** and **D**. Each of the four spectra is referred to by a number.

Spectrum	Number of peaks	Ratio of areas under the peaks
1	3	6:1:1
2	4	3:2:2:1
3	3	3:2:3
4	3	3:2:3

(i) Which compound (**A**, **B**, **C** or **D**) corresponds to spectrum 1? Give a reason for your choice.

**(2)** 

(ii) Which compound (**A**, **B**, **C** or **D**) corresponds to spectrum 2? Give a reason for your choice.

**(2)** 

(iii) Explain why spectrum 3 and spectrum 4 have the same number of peaks and the same ratio of areas under them.

**(1)** 

(f) Compound C reacts with HCN. Give the name of the type of mechanism, and an outline of the mechanism for this reaction.

**(5)** 

(Total 26 marks)