

A Level Chem 33 EQ 22m to 09w Paper 4 Carboxylic acids and derivatives

85 marks

As you start and work through this worksheet you can tick off your progress to show yourself how much you have done, and what you need to do next. The first task is just to read the first question and should take you less than 3 minutes to complete.

Paper 4 Topic Checklist

Tick each task off as you go along

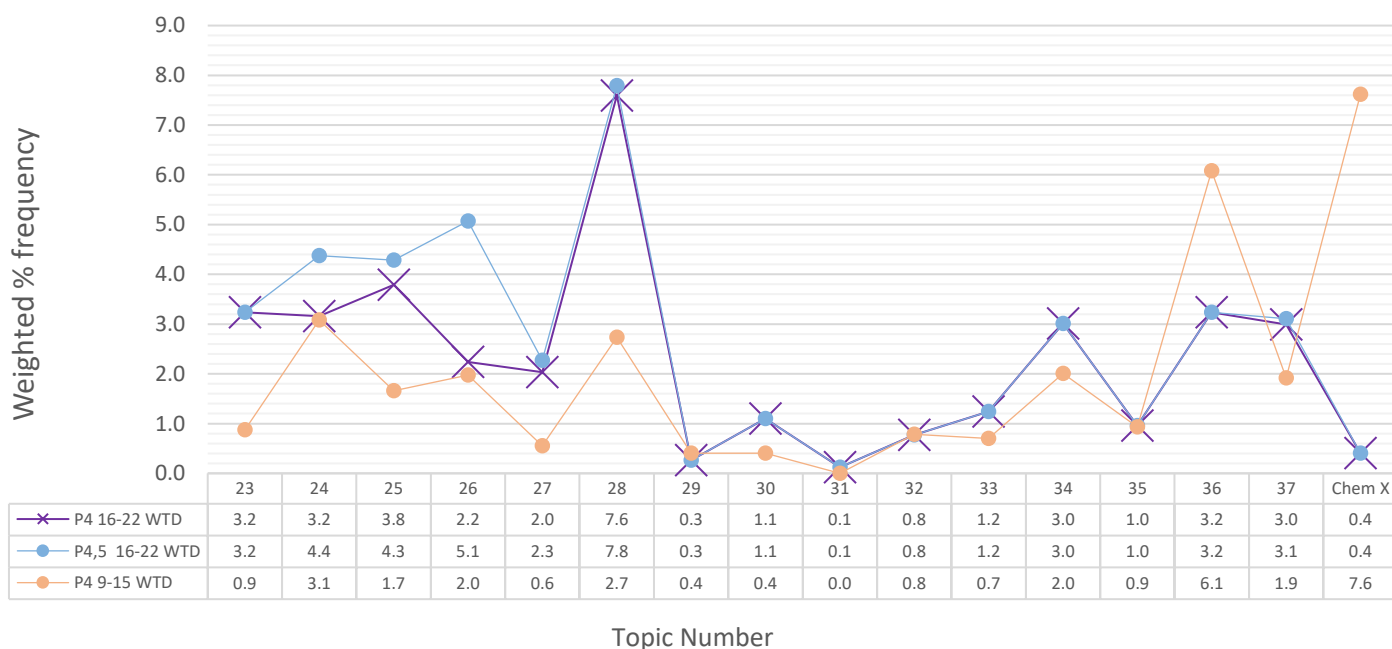
RANK:

Marks

		P4 Noob	P4 Novice	P4 Bronze	P4 Silver	P4 Gold	P4 ¹ Winner	P4 Hero	P4 Legend
		1 Q Started	1 Q done	10% of marks	25% of marks	40% of marks	50% of marks	75% of marks	100% of marks
Topic 33 (marks)	85		5	9	21	34	43	64	85
Time - 72 seconds per mark (minutes)	103		6	10	26	41	52	78	103

9701 Chemistry Weighted Mark Frequency: Paper 4 (A2 Focus)

2022w to 2016m in purple crosses, compared to earlier and all A2 exams combined



What the most thoughtful students will get out of their extensive studying will be a capacity to do meaningful brain-based work even under stressful conditions, which is a part of the self-mastery skillset that will continue to deliver value for the whole of their lives. Outstanding grades will also happen, but the most important outcome from skillful action in study is being better at any important tasks even if circumstances are do not feel ideal.

Learning how to manage oneself so we can more reliably get ambitious and successful outcomes out of our challenges in a productive and positive way is one aspect of life's most valuable pursuit summarised and inscribed on the Temple of Apollo at Delphi: "know thyself".

1. To complete these questions, as important as your answer, is checking your answer against the mark scheme.
2. For each question, or page, convert your mark score into a percentage. This will allow you to see (and feel) your progress as you get more experience and understanding with each topic.
3. If you find you get a higher percentage answering short answer questions than multiple choice questions that often means you are using the marking scheme correctly; your correct answer might not be fully complete. The marks easiest to miss rely on providing more details fully described.

¹ DO NOT work on these higher levels of completion unless you have also achieved at least a "Silver" (25%) in the same topic in Paper 5, if it exists.

33 Carboxylic acids and derivatives

33.1 Carboxylic acids

Learning outcomes

Candidates should be able to:

- 1 recall the reaction by which benzoic acid can be produced:
 - (a) reaction of an alkylbenzene with hot alkaline KMnO_4 and then dilute acid, exemplified by methylbenzene
- 2 describe the reaction of carboxylic acids with PCl_3 and heat, PCl_5 , or SOCl_2 to form acyl chlorides
- 3 recognise that some carboxylic acids can be further oxidised:
 - (a) the oxidation of methanoic acid, HCOOH , with Fehling's reagent or Tollens' reagent or acidified KMnO_4 or acidified $\text{K}_2\text{Cr}_2\text{O}_7$ to carbon dioxide and water
 - (b) the oxidation of ethanedioic acid, HOOC-COOH , with warm acidified KMnO_4 to carbon dioxide
- 4 describe and explain the relative acidities of carboxylic acids, phenols and alcohols
- 5 describe and explain the relative acidities of chlorine-substituted carboxylic acids

33.2 Esters

Learning outcomes

Candidates should be able to:

- 1 recall the reaction by which esters can be produced:
 - (a) reaction of alcohols with acyl chlorides using the formation of ethyl ethanoate and phenyl benzoate as examples

33.3 Acyl chlorides

Learning outcomes

Candidates should be able to:

- 1 recall the reactions (reagents and conditions) by which acyl chlorides can be produced:
 - (a) reaction of carboxylic acids with PCl_3 and heat, PCl_5 , or SOCl_2
- 2 describe the following reactions of acyl chlorides:
 - (a) hydrolysis on addition of water at room temperature to give the carboxylic acid and HCl
 - (b) reaction with an alcohol at room temperature to produce an ester and HCl
 - (c) reaction with phenol at room temperature to produce an ester and HCl
 - (d) reaction with ammonia at room temperature to produce an amide and HCl
 - (e) reaction with a primary or secondary amine at room temperature to produce an amide and HCl
- 3 describe the addition-elimination mechanism of acyl chlorides in reactions in 33.3.2(a) – (e)
- 4 explain the relative ease of hydrolysis of acyl chlorides, alkyl chlorides and halogenoarenes (aryl chlorides)

6 Lidocaine is used as an anaesthetic. A synthesis of lidocaine is shown in Fig. 6.1.

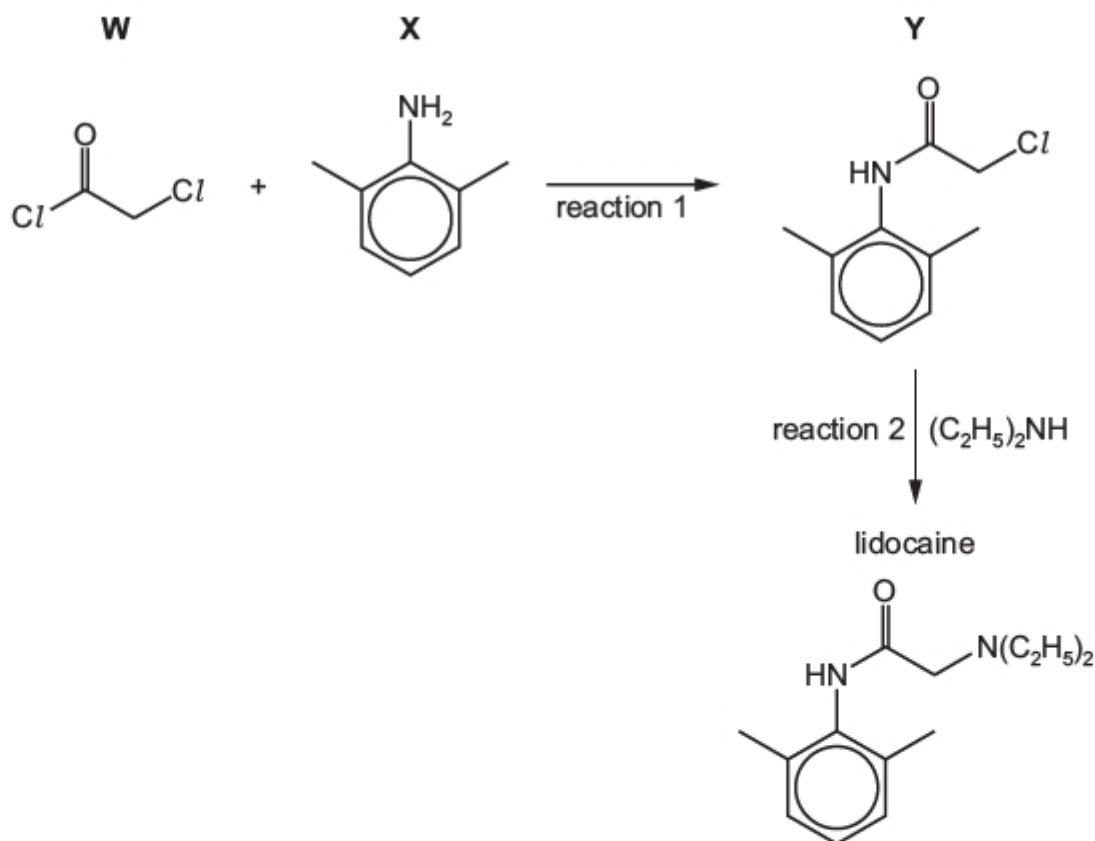


Fig. 6.1

(a) **W** can be formed by reacting HOCH_2COOH with an excess of SOCl_2 .

Write an equation for this reaction.

..... [1]

(b) After **W** and **X** have reacted together, an excess of $\text{CH}_3\text{COONa(aq)}$ is added to the reaction mixture.

Suggest why.

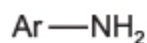
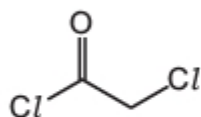
..... [1]

(c) The reaction of **W** with **X**, reaction 1, follows an addition–elimination mechanism.

Complete the mechanism for the reaction of **W** with **X**.

Include all relevant curly arrows, lone pairs of electrons, charges and partial charges.

Use Ar-NH₂ to represent **X**.



[4]

(d) (C₂H₅)₂NH reacts with **Y** in reaction 2.

Explain why (C₂H₅)₂NH can act as a nucleophile.

[1]

Q# 369/ ALvl Chemistry/2022/m/TZ 1/Paper 4/Q# 4 /www.SmashingScience.org :o)

(d) **K** can also be synthesised from phenol, C₆H₅OH.

Fig. 4.4 shows several reactions of phenol.

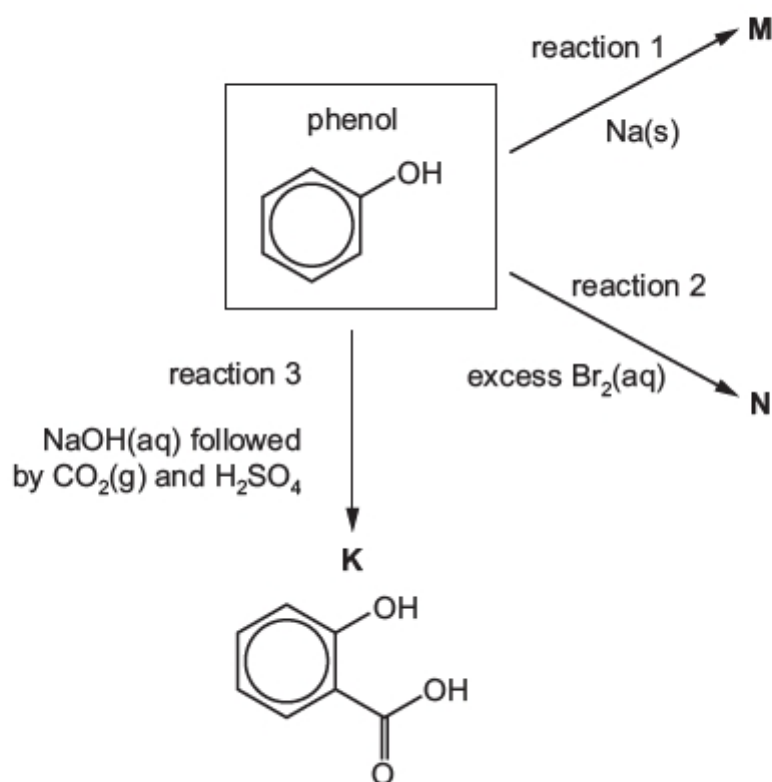


Fig. 4.4

(iii) Explain why phenol is a weaker acid than K.

.....

.....

.....

..... [2]

Q# 370/ ALvl Chemistry/2021/s/TZ 1/Paper 4/Q# 6 /www.SmashingScience.org :o)

6 (a) Compare and explain the relative acidities of butanoic acid, ethanol, ethanoic acid and water.

..... > > >

most acidic least acidic

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..... [4]

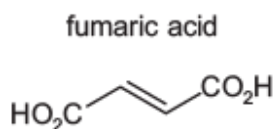
(b) Three carboxylic acids, methanoic acid, HCO_2H , ethanedioic acid, $\text{HO}_2\text{CCO}_2\text{H}$, and butanedioic acid, $\text{HO}_2\text{CCH}_2\text{CH}_2\text{CO}_2\text{H}$, are compared. Two tests were carried out on separate samples of each organic acid, as shown.

The following results were obtained. ✓ = observed change X = no observed reaction

test	reagents and conditions	HCO_2H	$\text{HO}_2\text{CCO}_2\text{H}$	$\text{HO}_2\text{CCH}_2\text{CH}_2\text{CO}_2\text{H}$	observed change
1		✓	X	X	
2		✓	✓	X	

(i) Complete the table with the reagents and conditions and the observed change for a positive test.
Assume these organic acids all have a similar acid strength. [3]

6 Fumaric acid is a naturally occurring dicarboxylic acid.

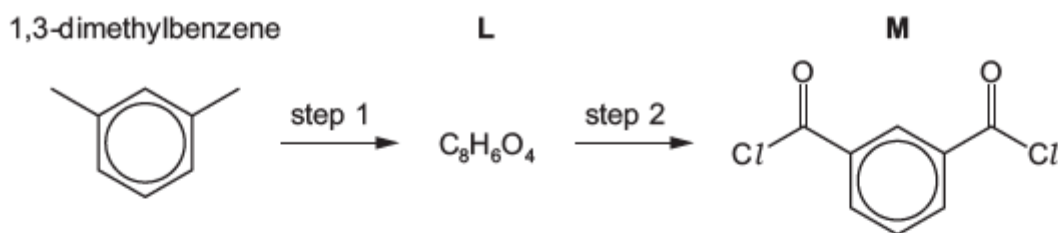


(a) Identify the products of the reaction between fumaric acid and an excess of hot, concentrated, acidified manganate(VII).

..... [1]

Q# 372/ ALvl Chemistry/2021/m/TZ 2/Paper 4/Q# 5 /www.SmashingScience.org :o)

(b) Compound **M** is made from 1,3-dimethylbenzene in a two-step synthesis.



(i) Draw the structure of **L**.

[1]

(ii) Suggest reactants and conditions for each step of this synthesis.

step 1

step 2

[2]

(iii) Write an equation for step 2.

..... [1]

- 6 (a) Compare and explain the relative acidities of 2-chloropropanoic acid, 3-chloropropanoic acid, and propanoic acid. Explain your answer.

..... > >
 most acidic least acidic

explanation

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.....

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.....

.....

[3]

- (c) Three tests were carried out on separate samples of the organic acids shown in the table. The following results were obtained.
 ✓ = observed change
 X = no observed reaction

test	reagent(s) and conditions	HCO ₂ H	CH ₃ COCO ₂ H	HO ₂ CCO ₂ H	observed change
1		✓	X	X	
2		X	✓	X	
3		✓	X	✓	

Complete the table with the reagent(s) and conditions and the observed change for each test. Assume these organic acids all have a similar acid strength. [5]



(iii) $[\text{Fe}(\text{C}_2\text{O}_4)_2\text{Cl}_2]^{4-}$ contains ligands which are anions of ethanedioic acid, $\text{HO}_2\text{CCO}_2\text{H}$.

Complete the table to show any observations for the reactions of $\text{HO}_2\text{CCO}_2\text{H}$ with the named reagents.

Where no change is observed, write 'none'.

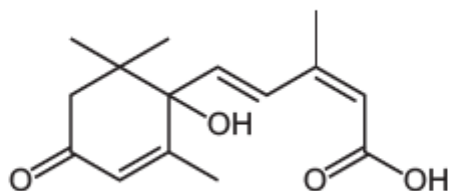
reagent	observations with $\text{HO}_2\text{CCO}_2\text{H}$
warm acidified manganate(VII)	
2,4-dinitrophenylhydrazine	
warm Tollens' reagent	

[2]

[Total: 20]

Q# 375/ ALVl Chemistry/2018/s/TZ 1/Paper 4/Q# 8 /www.SmashingScience.org :o)

8 Absciscic acid, $\text{C}_{15}\text{H}_{20}\text{O}_4$, is a plant hormone.



abscisic acid, $\text{C}_{15}\text{H}_{20}\text{O}_4$

(b) Absciscic acid is reacted with an excess of NaBH_4 .

Give the molecular formula of the organic product formed.

..... [1]

(c) If abscisic acid is treated with an excess of hot, concentrated, acidified KMnO_4 , three different carbon-containing products are formed.

(i) Draw the skeletal formula of the carbon-containing product with the **largest** molecular mass.

[1]

- (ii) Identify the carbon-containing product with the **smallest** molecular mass. Explain how this product arises.

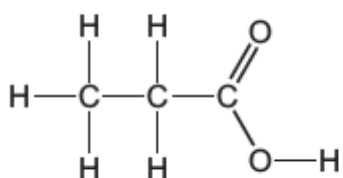
.....
.....
..... [2]

- (iii) Identify the third carbon-containing product of this reaction by giving its displayed or structural formula.

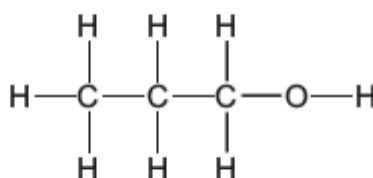
[1]

Q# 376/ ALvl Chemistry/2018/s/TZ 1/Paper 4/Q# 7 /www.SmashingScience.org :o)

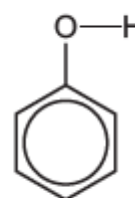
7 The three substances shown all have some acidic properties.



propanoic acid



propan-1-ol



phenol

- (b) (i) Give the order of the relative acidities of propanoic acid, propan-1-ol and phenol, stating the most acidic first.

..... [1]

- (ii) Explain your answer to (i).

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..... [2]

(c) Methanoic acid, HCO_2H , has a similar acid strength to propanoic acid.

Describe a chemical test to distinguish between these two acids. Name the acid which gives a positive result in this test and describe the observations that would be made.

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.....

..... [2]

(d) The ester phenyl propanoate, $\text{C}_2\text{H}_5\text{CO}_2\text{C}_6\text{H}_5$, can be made from phenol and propanoic acid in a **two-step** synthesis. The first step produces an acyl chloride.

For this **two-step** synthesis,

- draw the structure of the product of the first step,
- state the reagents and conditions needed for each step of the synthesis.

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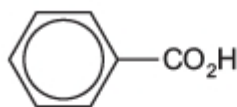
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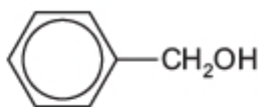
..... [3]

Q# 377/ ALvI Chemistry/2018/m/TZ 2/Paper 4/Q# 9 /www.SmashingScience.org :o)

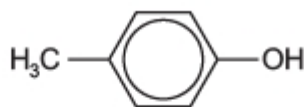
(b) Describe and explain the relative acidities of benzoic acid, phenylmethanol and 4-methylphenol.



benzoic acid



phenylmethanol



4-methylphenol

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..... [3]

Q# 378/ ALvI Chemistry/2018/m/TZ 2/Paper 4/Q# 1 /www.SmashingScience.org :o)

- (d) (i) The ethanedioates of the Group 2 elements, MC_2O_4 , decompose on heating to produce a mixture of two different gases and the solid oxide, MO , only.
- (ii) Describe **two** observations you would make during the reaction when ethanedioic acid, $H_2C_2O_4$, is warmed with acidified manganate(VII) ions.

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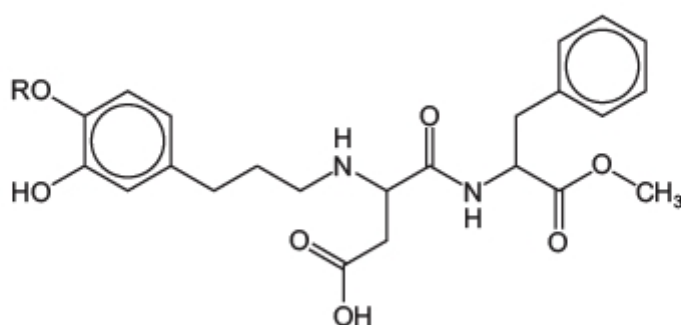
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..... [2]

[Total: 14]

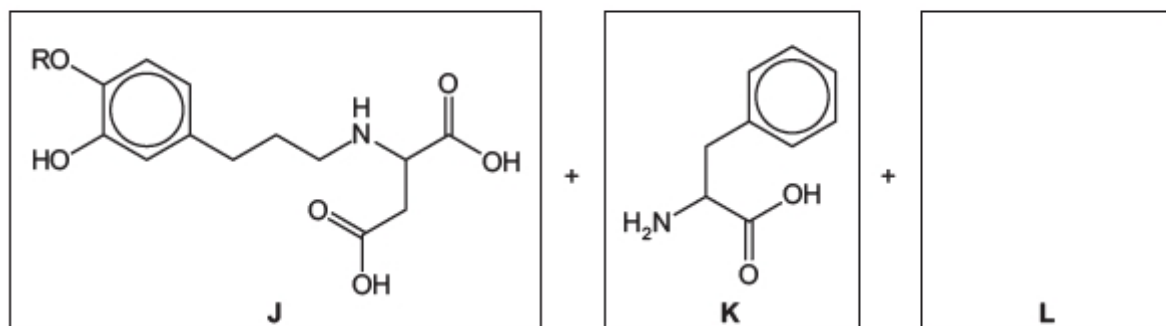
Q# 379/ ALVl Chemistry/2017/m/TZ 2/Paper 4/Q# 7 /www.SmashingScience.org :o)

- 7 The compound *Advantame* is a sweetener that tastes approximately 25 000 times sweeter than sucrose.



Advantame

- (b) The decomposition of *Advantame* produces three molecules, J, K and L. The RO– group in *Advantame* is unreactive.



- (i) Suggest possible reagents and conditions for this decomposition.

..... [1]

- (ii) Name the *type of reaction* occurring.

..... [1]

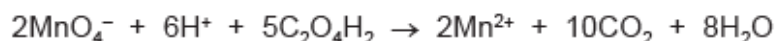
- (iii) Draw the structure of L in the box above.

[1]

10 (a) Ethanedioic acid, $C_2O_4H_2$, occurs in many vegetables. The amount that occurs in spinach can be estimated as follows.

- 40.0 g of spinach leaves are crushed and mixed with distilled water, using a mortar and pestle.
- The mixture is filtered, and the leaves are washed with a little more water.
- The combined filtrate and washings are made up to 100.0 cm^3 with water.
- A 25.0 cm^3 portion of the resulting solution is added to a conical flask, along with an excess of dilute sulfuric acid.
- The acidified solution is warmed, and then titrated with $0.0200\text{ mol dm}^{-3}\text{ KMnO}_4$.

The equation for the reaction between ethanedioic acid and acidified manganate(VII) ions is shown.

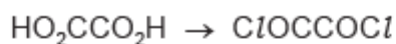


In the titration, 15.20 cm^3 of KMnO_4 was required to reach the end-point.

Calculate the percentage by mass of ethanedioic acid in the spinach leaves.

percentage of ethanedioic acid = % [3]

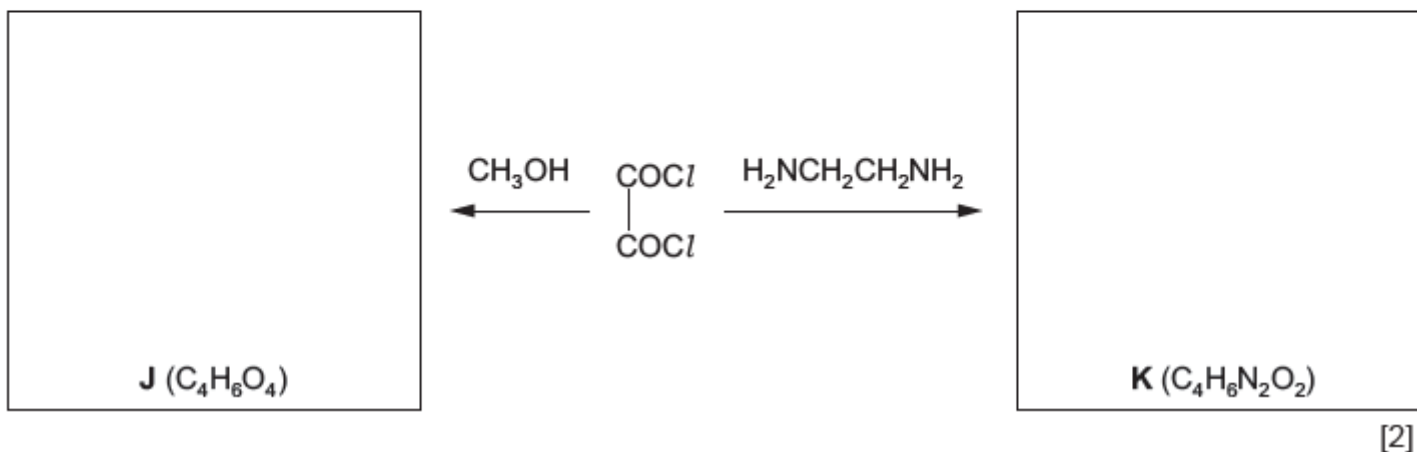
(b) Ethanedioic acid can be converted into ethanedioyl chloride:



(i) State a suitable reagent for this reaction.

..... [1]

(ii) For the reactions of ethanedioyl chloride below, suggest the structures of compounds **J** and **K** and draw them in the boxes.



Q# 381/ ALvl Chemistry/2015/s/TZ 1/Paper 4/Q# 2 /www.SmashingScience.org :o)

2 (a) Both chloroalkanes and acyl chlorides react with water, but only acyl chlorides fume in moist air.

(i) State which product causes the fumes in this reaction.

..... [1]

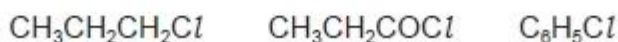
(ii) Explain why the reactivities of chloroalkanes and acyl chlorides differ.

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.....
..... [1]

5 (a) Organohalogen compounds can undergo hydrolysis.



State the relative rates of hydrolysis of the following compounds.



Explain your answer.

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..... [3]

Q# 383/ ALvl Chemistry/2014/w/TZ 1/Paper 4/Q# 4 /www.SmashingScience.org :o)

(b) (i) Describe and explain the relative acidities of chloroethanoic acid and ethanoic acid.

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(ii) Describe and explain the relative acidities of phenol and ethanol.

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[3]

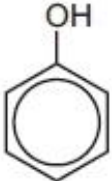
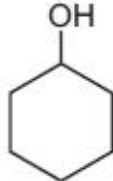
Q# 384/ ALvl Chemistry/2013/s/TZ 1/Paper 4/Q# 5 /www.SmashingScience.org :o)

5 (a) A series of experiments is carried out in which the reagent shown at the top of the column of the table is mixed, in turn, with each of the reagents at the side.

Complete the following table by writing in each box the formula of any gas produced.

Write x in the box if no gas is produced.

The first column has been completed as an illustration.

		OH 	CO ₂ H 	OH 
	H ₂ O			
Na	H ₂			
KOH(aq)	x			
Na ₂ CO ₃ (aq)	x			

[5]

6 Acyl chlorides are useful intermediates in organic syntheses.

(a) (i) State a suitable reagent for converting carboxylic acids into acyl chlorides.

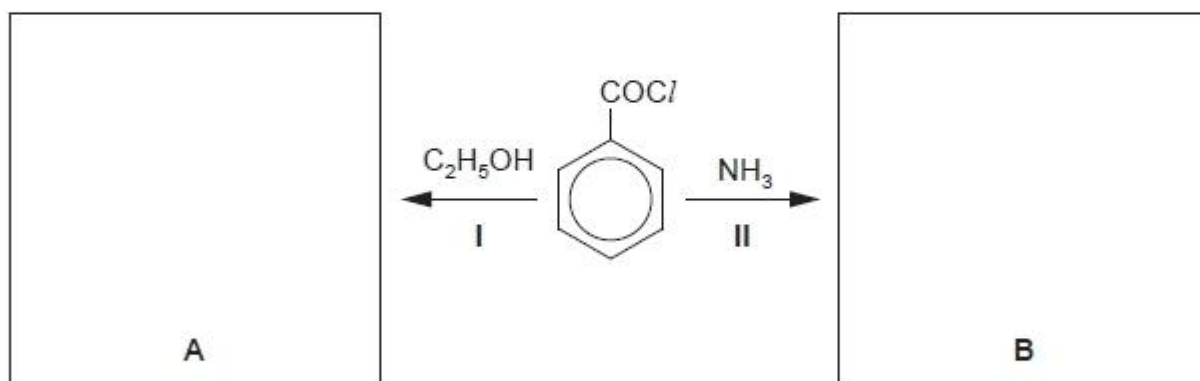
.....

(ii) Construct an equation for the reaction between ethanoic acid, $\text{CH}_3\text{CO}_2\text{H}$, and the reagent you have stated in (i).

.....

[2]

(b) (i) In the boxes provided draw the structures of the compounds formed when benzoyl chloride undergoes the following reactions.



(ii) Name the functional group in

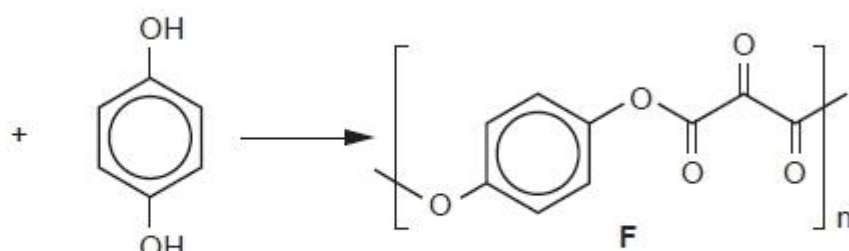
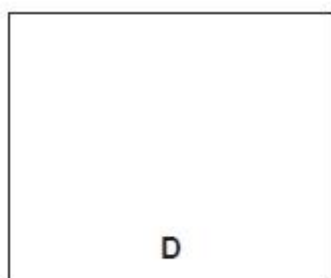
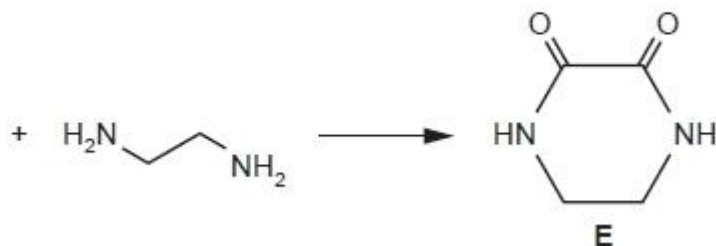
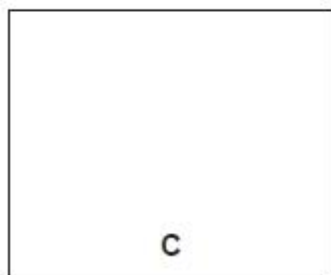
- compound A
- compound B

(iii) What *type of reaction* is reaction II?

.....

[5]

- (c) (i) Suggest suitable acyl chlorides to use in the following reaction. Draw their structures in the boxes provided.



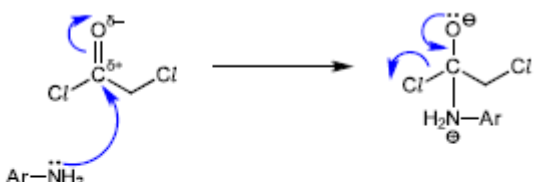
Compound **E** dissolves in, but does not react with, cold water.

- (iv) What type of polymer is compound **F**?

..... [3]

Mark Scheme A Level Chem 33 EQ 22m to 09w Paper 4 Carboxylic acids and derivatives 85 marks

Q# 368/ ALvI Chemistry/2022/m/TZ 1/Paper 4/Q# 6 /www.SmashingScience.org :o)

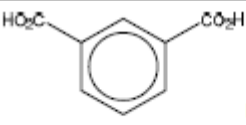
6(a)	$\text{HOCH}_2\text{COOH} + 2\text{SOCl}_2 \rightarrow \text{ClCH}_2\text{COCl} + 2\text{SO}_2 + 2\text{HCl}$	1
6(b)	to remove / neutralise excess H^+ / acid produced OR to react with any acidic by-products / HCl / SO_2 OR to react with any unreacted W	1
6(c)	 M1: curly arrow from lone pair on $:\text{NH}_2$ to carbonyl $\text{C}(=\text{O})$ M2: correct dipole on $^{\delta+}\text{C}=\text{O}^{\delta-}$ AND curly arrow from bond $\text{C}=\text{O}$ to $\text{O}^{\delta-}$ M3: correct structure of the intermediate (inc. charges) M4: curly arrow from lone pair on $:\text{O}^-$ to $\text{C}=\text{O}$ AND curly arrow from $\text{C}-\text{Cl}$ to Cl	4
6(d)	N / nitrogen can donate its lone pair / LP / pair of electrons	1

Q# 369/ ALvI Chemistry/2022/m/TZ 1/Paper 4/Q# 4 /www.SmashingScience.org :o)

4(d)(iii)	(CO)O—H bond weaker / more easy to donate H^+ in K - owing to negative inductive / electron withdrawing effect of $\text{C}=\text{O}$ / COOH group - carboxylate anion stabilised / phenoxide anion is less stabilised All three for two marks	2
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6(a)	M1: ethanoic acid > butanoic acid > water > ethanol M2: a reason given in terms of an electron donating or an electron withdrawing group for one of: strengthening of O–H bond OR weakening of O–H bond OR stability of anion Two out of the three alternatives <i>M3</i>, <i>M4</i> and <i>M5</i>: M3: <u>ethanol</u>: positive inductive effect / electron donating effect of ethyl / alkyl / R group M4: <u>butanoic acid</u>: positive inductive effect / electron donating effect of propyl / alkyl / R group M5: (either ethanoic or butanoic) <u>acid</u>: negative inductive effect of either C=O or carbonyl OR negative charge delocalised over COO⁻	4									
6(b)(i)	<table><tr><td></td><td>reagents and conditions</td><td>observed change</td></tr><tr><td>test 1</td><td>Tollen's reagent, warm OR Fehling's solution, warm</td><td>silver mirror (brick) red ppt / solid</td></tr><tr><td>test 2</td><td>acidified MnO₄⁻, warm</td><td>decolourises OR bubbles</td></tr></table> M1 / M2: reagents and conditions × 2 M3: observations both correct		reagents and conditions	observed change	test 1	Tollen's reagent, warm OR Fehling's solution, warm	silver mirror (brick) red ppt / solid	test 2	acidified MnO ₄ ⁻ , warm	decolourises OR bubbles	3
	reagents and conditions	observed change									
test 1	Tollen's reagent, warm OR Fehling's solution, warm	silver mirror (brick) red ppt / solid									
test 2	acidified MnO ₄ ⁻ , warm	decolourises OR bubbles									

6(a)	CO ₂ and H ₂ O / in words	1
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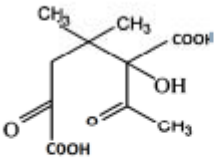
5(b)(i)	 L	1
5(b)(ii)	M1: heat / reflux with acidified / alkaline KMnO₄ (then acidify) M2: PCl₅ OR SOCl₂ / (heat with) PCl₅	2
5(b)(iii)	C₆H₅O₄ + 2PCl₅ → C₆H₄O₂Cl₂ + 2POCl₃ + 2HCl OR C₆H₅O₄ + 2SOCl₂ → C₆H₄O₂Cl₂ + 2SO₂ + 2HCl OR 3C₆H₅O₄ + 2PCl₅ → 3 C₆H₄O₂Cl₂ + 2H₃PO₃	1

6(a)	M1 2-chloropropanoic acid > 3-chloropropanoic acid > propanoic acid [1] M2 CH₃CHClCO₂H / ClCH₂CH₂CO₂H (are more acidic) as they contain an electronegative Cl atom so weaken O-H bond / stabilise carboxylate anion [1] M3 CH₃CHClCO₂H (is more acidic than ClCH₂CH₂CO₂H) as the Cl atom is closer to CO₂H so weaken O-H bond more / stabilise carboxylate anion more [1]			3												
6(c)	<table><tr><td></td><td>reagents and conditions</td><td>observed change</td></tr><tr><td>test 1</td><td>M1 Tollen's reagent, warm OR Fehling's solution, warm</td><td>silver mirror (brick)-red ppt.</td></tr><tr><td>test 2</td><td>M2 aqueous alkaline iodine OR 2,4-DNPH</td><td>yellow ppt. orange ppt.</td></tr><tr><td>test 3</td><td>M3 acidified MnO₄⁻, warm</td><td>decolourises (and bubbles)</td></tr></table> Two correct observations = 1 mark Three correct observations = 2 marks				reagents and conditions	observed change	test 1	M1 Tollen's reagent, warm OR Fehling's solution, warm	silver mirror (brick)-red ppt.	test 2	M2 aqueous alkaline iodine OR 2,4-DNPH	yellow ppt. orange ppt.	test 3	M3 acidified MnO ₄ ⁻ , warm	decolourises (and bubbles)	5
	reagents and conditions	observed change														
test 1	M1 Tollen's reagent, warm OR Fehling's solution, warm	silver mirror (brick)-red ppt.														
test 2	M2 aqueous alkaline iodine OR 2,4-DNPH	yellow ppt. orange ppt.														
test 3	M3 acidified MnO ₄ ⁻ , warm	decolourises (and bubbles)														

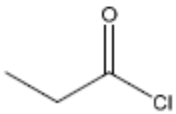
Q# 374/ ALvl Chemistry/2020/m/TZ 2/Paper 4/Q# 1 /www.SmashingScience.org :o)

1(c)(iii)		reactant	observation with (CO ₂ H) ₂	2
		warm H ⁺ /MnO ₄ ⁻	decolourised OR effervescence / bubbling / fizzing	
		2,4-DNPH	none / no reaction	
		warm Tollens' reagent	none / no reaction	

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8(b)	C ₁₅ H ₂₂ O ₄	1
8(c)(i)		1
8(c)(ii)	CO ₂	1
	oxidation / oxidative cleavage	1
8(c)(iii)	CH ₃ COCO ₂ H	1

Q# 376/ ALvl Chemistry/2018/s/TZ 1/Paper 4/Q# 7 /www.SmashingScience.org :o)

7(b)(i)	propanoic acid, phenol, propan-1-ol	1
7(b)(ii)	- propan-1-ol: O-H bond strengthened by positive inductive effect of alkyl group OR propoxide ion is destabilised by positive inductive effect of alkyl group - phenol: O-H bond weakened by negative inductive effect of ring OR phenoxide ion is stabilised by delocalisation of oxygen lone pair into ring - propanoic acid: O-H bond weakened by negative inductive effect of C=O OR propanoate ion is stabilised by delocalisation of minus charge by C=O 1 mark for a correct explanation, max 2 marks	2
7(c)	Tollens' reagent or Fehling's reagent	1
	methanoic acid gives a silver mirror/solid with Tollen's reagent OR red / orange ppt / solid with Fehlings' reagent	1
7(d)	PCl ₅ or PCl ₃ (+heat) or SOCl ₂ (added to propanoic acid)	1
	product of first step: 	1
	add product of first step to phenol in NaOH	1

Q# 377/ ALvl Chemistry/2018/m/TZ 2/Paper 4/Q# 9 /www.SmashingScience.org :o)

9(b)	benzoic acid > methylphenol > phenylmethanol methylphenoxide anion has delocalisation of the lone pair on oxygen over the ring benzoic acid has an (extra) electronegative oxygen or electron withdrawing C=O	3
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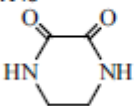
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1(d)(ii)	the KMnO ₄ would decolourise bubbles / gas evolution would be seen	2
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Q# 379/ ALvl Chemistry/2017/m/TZ 2/Paper 4/Q# 7 /www.SmashingScience.org :o)

7(b)(i)	H ⁺ (aq) + heat	1
7(b)(ii)	hydrolysis	1
7(b)(iii)	CH ₃ OH	1

Q# 380/ ALVl Chemistry/2016/s/TZ 1/Paper 4/Q# 10 /www.SmashingScience.org :o)

10 (a)	$n(\text{MnO}_4^-) = 0.02 \times 15.2/1000 = 3.04 \times 10^{-4} \text{ mol}$ $n(\text{C}_2\text{O}_4\text{H}_2) = 3.04 \times 10^{-4} \times 5/2 = 7.6 \times 10^{-4} \text{ (in } 25\text{cm}^3) = 3.04 \times 10^{-3} \text{ mol in } 100\text{cm}^3$ $M_r = 24 + 64 + 2 = 90$ $\text{mass of C}_2\text{O}_4\text{H}_2 = 3.04 \times 10^{-3} \times 90 = 0.2736 \text{ g (0.274)}$ $\text{percentage} = 0.2736 \times 100/40 = 0.68\%$	[1] [1] [1]
(b) (i)	SOCl_2 or PCl_5 or PCl_3	[1]
(ii)	J is $\text{CH}_3\text{OCO}-\text{COOCH}_3$ K is 	[1] [1]

Q# 381/ ALVl Chemistry/2015/s/TZ 1/Paper 4/Q# 2 /www.SmashingScience.org :o)

2 (a) (i)	hydrogen chloride or HCl	1
(ii)	either (RCOCl) has two electron-withdrawing groups/atoms, making the more δ^+ /electron deficient or (RCOCl) has an oxygen, making the carbon more δ^+ /electron deficient or (RCOCl) has two electron-withdrawing groups, weakening the C-Cl bond	1

Q# 382/ ALVl Chemistry/2014/w/TZ 1/Paper 4/Q# 5 /www.SmashingScience.org :o)

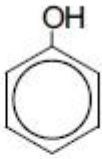
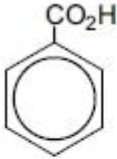
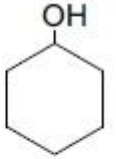
5 (a)	$\text{CH}_3\text{CH}_2\text{COCl} > \text{CH}_3\text{CH}_2\text{CH}_2\text{Cl} > \text{C}_6\text{H}_5\text{Cl}$ any two of: - C-Cl bond strength is weakest in $\text{CH}_3\text{CH}_2\text{COCl}$ ora - In $\text{C}_6\text{H}_5\text{Cl}$ (no hydrolysis) C-Cl bond is part of delocalised system OR p-orbital on Cl overlaps with π system OR electrons from Cl overlap with π system $\text{CH}_3\text{CH}_2\text{COCl}$ carbon in C-Cl bond is more electron deficient since it is also attached to an oxygen atom ora	1 1+1	[3]
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Q# 383/ ALVl Chemistry/2014/w/TZ 1/Paper 4/Q# 4 /www.SmashingScience.org :o)

(b) (i)	acidity of $\text{ClCH}_2\text{CO}_2\text{H} > \text{CH}_3\text{CO}_2\text{H}$ AND $(\text{ClCH}_2\text{CO}_2\text{H})$ as an electronegative/electron withdrawing Cl	1	
(ii)	acidity of phenol $> \text{CH}_3\text{CH}_2\text{OH}$ AND electrons on oxygen (on phenol) delocalised into ring OR benzene ring withdraws electrons from oxygen stronger acid linked to weakening O-H bond/anion being stabilised	1 1	[3]

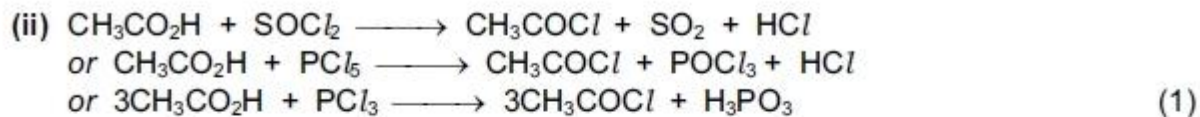
Q# 384/ ALVl Chemistry/2013/s/TZ 1/Paper 4/Q# 5 /www.SmashingScience.org :o)

5 (a)

	H_2O			
Na	H_2	H_2	H_2	H_2
KOH(aq)	X	X	X	X
$\text{Na}_2\text{CO}_3(\text{aq})$	X	X	CO_2	X

[5]

6 (a) (i) SOCl_2 or PCl_5 or PCl_3 (1)



[2]

(b) (i) A is $\text{C}_6\text{H}_5\text{CO}_2\text{C}_2\text{H}_5$ (1)
B is $\text{C}_6\text{H}_5\text{CONH}_2$ (1)

(ii) ester	(1)
amide	(1)

(iii) nucleophilic substitution / condensation (1) [5]

(c) (i) **C** is ClCOCOC l (1)
D is ClCOCOCOC l (1)

