## A' Level Chemistry Year 2



## **Unit 20: Chromatography**

## **Summer Examination Revision Pack**

The questions in this pack should be attempted AFTER completing all other revision.



Grade Accelerator Recall Definitions Drawing Diagrams Using Equations Drawing Graphs



STIL

Condensed Notes Keywords & Definitions Key Concepts Application Key Skills



Quizlet Classes Flashcard Based Games Tests & Quizzes Keyword Spell Checker

# Veral A' Level Chemistry : Periodicity / Group II / Group VII

### **Online Forms**

Take Time to Answer Use Paper & Calculator Work It Out Review Missed Marks

### Use the 3 Wave Process when completing these revision packs.



 Complete the questions without assistance (Can't answer a question? Leave it and move on)
 Use your notes to fill any gaps after step 1
 Use the mark scheme to fill in any remaining gaps.

#### 1. Having gaps after step 1 is normal, that's why we are doing revision!

 If your notes don't help during step 2, they are not good enough! (Change your note taking method and try to understand the problem)
 If you don't understand why the mark scheme answer is correct, see Andy.

STOP If you struggle with the questions in the pack, STOP! and complete some more revision.

If you come to a complete dead-end, **STOP!** and speak to **Andy** asap.

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 $NO_2$ 





Figure 3

 $O_2N$ 



Use the chromatogram in Figure 3 to deduce the  $R_f$  value of 1,4-dinitrobenzene in this experiment.







[1 mark]

nes the distance travelled by a spot in TLC. [1 mark]
C plate was held by the edges and placed me cupboard. The lid was then replaced
t when placing the plate in the beaker. [1 mark]
ed out using 1,2-dinitrobenzene and te to that in Question <b>8.3</b> was used under
greater than that of 1,2-dinitrobenzene. 1,2-dinitrobenzene and 1,4-dinitrobenzene has the greater $R_f$ value.
[2 marks]



18

08.3	D	1	
08.4	(Balance between) solubility in moving phase and retention by stationary phase	1	OR (relative) affinity for stationary/solid and mobile/liquid/solvent (phase)
08.5	Solvent depth must be below start line	1	Ignore safety
	1,2- is more polar <b>OR</b> 1,4- is less polar	1	M2 dependent on correct M1
08.6	1,4- ( or Less/non polar is) less attracted to (polar) plate / stationary phase / solid	1	If M1 is blank then read explanation for possible M1 and M2
	<b>OR</b> (Less/non polar is) more attracted to / more soluble in (non-polar) solvent / mobile phase / hexane		Allow converse argument for 1,2
	No CE = 0 Yes - mark on but there is <b>NO MARK FOR YES</b>		
08.7	Solvent (more) polar or ethyl ethanoate is polar	1	Mark independently following yes
	Polar isomer more attracted to / more soluble in / stronger affinity to the solvent (than before)	1	Penalise bonded to mobile phase in M2
			<u></u>
Total		12	





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	Tu	rn over ►	
	Turn over for the next question		
			4
09.4	Suggest why it was necessary to use two different solvents.	[1 mark]	
09.3	Deduce the minimum number of amino acids present in the original mixture.	[1 mark]	
09.2	Suggest how the positions of the amino acids on the TLC plate were located.	[1 mark]	outside the box

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Question	Answers	Additional Comments/Guidelines	Mark
09.1	Conc HCI	Allow concentrations of 5M or higher Allow <u>conc</u> sulfuric or <u>conc</u> strong alkalis	1
09.2	Using ninhydrin or ultraviolet light	Allow I <sub>2</sub> (vapour)	1
<b>G</b> 09.3	7 or seven		1
09.4	Some of the amino acids did not separate/dissolve with the first/either solvent <b>OR</b> Some amino acids have the same Rf value or have the same affinity with the first/either solvent	Not amino acids have different Rf values in different solvents	1





	Tu	rn over ►
	Turn over for the next question	
02.3	State why each amino acid has a different R <sub>f</sub> value.	[1 mark]
	State how the amino acids can be made visible at the end of the experiment.	[1 mark]
02.2	The amino acids cannot be seen as they move during the experiment.	outside th box

Question	Answers	Additional comments/Guidelines	Mark
2.1	<b>M1</b> $\frac{27}{80} = 0.34$ <b>M2</b> glycine	<ul><li>M1 some relevant working is needed to arrive at 0.325 - 0.35</li><li>no ECF based on M1</li></ul>	1

Question	Answers	Additional comments/Guidelines	Mark
2.2	use uv lamp or ninhydrin	allow developing / locating agent / iodine	1

Question	Answers	Additional comments/Guidelines	Mark
2.3	each amino acid has different (relative) affinity/attraction to/solubility in stationary and mobile phases	<b>allow</b> reference to different solubility in solvent OR different affinity for stationary phase	1