

GCE MARKING SCHEME

CHEMISTRY AS/Advanced

SUMMER 2010

CH4

SECTION A

1.	(a)	(i)	$C_{10}H_{16}$ (not shortened structural formula)	[1]
		(ii)		
				[1]
		(iii)	I Compound that contains no double bonds / single bonds only (Accept contains maximum number of hydrogens)	[1]
			II $C_{10}H_{16} + 3H_2 \longrightarrow C_{10}H_{22}$	[1]
			(accept structural formulae, consequential from (a)(i))	
		(iv)	Moles $H_2 = \frac{8.96}{22.4} = 0.4$	[1]
			4 double bonds	[1]
	(b)	(i)	Ethane (accept formula)	[1]
		(ii)	LiAlH ₄	[1]
		(iii)	H H Br H—C—C—H H H H	[1]
		(iv)	(1-)Propyl ethanoate	[1]

Total [10]

2. (a) In primary alcohols the –OH group is joined to a carbon atom bonded to two hydrogen atoms, in secondary alcohols the carbon atom is bonded to only one hydrogen. [1]

(b)	С	Н	0	
	68.1	13.7	18.2	
	12	1.01	16	
	5.675	13.56	1.1375	[1]
	4.99	11.92	1	

Empirical formula = $C_5H_{12}O$

[1]

Formula mass, 88.1(2), is same as M_r , therefore molecular formula is C₅H₁₂O [1]



(ii) I C [1]

II





CH₂OH

(iii) Add Tollens' reagent / Fehling's solution [1]
A no reaction, B gives silver mirror / red ppt. [1]
(iv) Add iodine in aqueous sodium hydroxide / KI and NaOCl [1]
C no reaction, D gives pale yellow ppt. [1]







Total [15]

(a)

3.

Iron(III) chloride / bromine water

Purple solution / white precipitate

[1] [1]

(b)	In phenol anion formed is stabilised by delocalisation of the negative chover the benzene ring (1) In alcohol anion formed is not stabilised by delocalisation (1) In carboxylic acids anion formed is stabilised by delocalisation of the n charge over the two oxygen atoms (1) Delocalisation is stronger in acid than in phenol (1) Phenol loses H+ more easily than alcohol and carboxylic acid does so m easily than phenol (1) (Accept diagrams)	narge egative nore
	max 4 marks	[4]
(c)	4-chloro-3,5-dimethylphenol	[1]
(d)	Atom economy = $\frac{94.06 \times 100}{122.07}$	[1]
	= 77%	[1]
(e)	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	[1]
(f)	Hexane-1,6-diamine / 1,6-diaminohexane	[1]
(g)		[1]
	$ \begin{array}{c} $	
(h)	In addition polymerisation only the polymer is formed (Accept description e.g. monomers join together with no water loss)	[1]
	In condensation polymerisation the polymer and a small molecule are for	ormed
	(Accept description e.g. water molecule lost as monomers join together (Accept for 1 mark only water is eliminated during condensation polymerisation but not during addition polymerisation))
(i)	Making aspirin / cellulose acetate / as an acylating agent	[1]
	То	tal [15]
	Total Section	n A [40]

SECTION B

4.	(a)	(i)				[3]
			H H H H H—C—C—C—C—Br H H H H	H Br	H ₃ C ¹ H ₅ H ₃ C ¹ H ₅ Br	
			(1)	(1)	(1)	
		(ii)	Use a polarimeter to dif One will rotate plane of left / opposite direction 1-bromobutane will hav	ferentiate between is plane polarised ligh s (1) we no effect on plane	somers (1) t to the right and one to polarised light (1)	o the [3]
			1-bromobutane (1) Answer gives peak pres isomers (1) (Accept converting to a	sent in 1-bromobutar	that is not in other out iodoform test)	
			QWC The information specialist vocabulary w	is organised clearly here appropriate	and coherently, using	[1]
	(b)	(i)	I $C_2H_5Cl + NH_3$ —	\rightarrow C ₂ H ₅ NH ₂ +	- HCl	[1]
			II Nucleophilic substit	tution		[1]
		(ii)	Nitrobenzene (1) Tin and concentrated hy	ydrochloric acid (1)		[2]
		(iii)	Sodium nitrite and hydr under 5 °C (1) No change phenylamine	ochloric acid, (1) e, bubbles (of nitroge	en) with ethylamine (1))
			Accept alternative – bro phenylamine, no change	omine (1), aqueous (e with ethylamine (1	1), white ppt with)	
						[3]

(c) (i)



(iii) Primary structure is the amino acid sequence (1) Secondary structure is the folding / coiling / twisting of the primary structure (held by hydrogen bonding) (1) Tertiary structure is the overall folding of the chain held by interactions (e.g. disulphide bridges, ionic bonding) between more distant amino acids / 3D structures (1) [3]

Total [20]

5.	(a)	Planar molecule (1) each carbon bonded covalently to 2 other carbons and 1 hydrogen.(1) Sideways overlap of p-orbitals. (1) Leading to delocalised π – bonds. (1) above and below plane of molecule (1) (Maximum 4 marks for bonding)					
		Alken attract In ben deloca	Alkenes have localised double bond /greater electron density / attract electrophiles more readily (1) In benzene stable delocalised system would be disturbed / has extra delocalisation energy / reacts by substitution (1)				
		<i>QWC</i> Legibility of text; accuracy of spelling, punctuation and grammar, clarity of meaning (1) Selection of a form and style of writing appropriate to purpose and to complexity of subject matter (1)					
	(b)	(i)	Benzene is too hazardous / methylbenzoate is less toxic	[1]			
		(ii)	(ii) The product is crude / impure				
		(iii)	Add crystals to minimum amount of hot ethanol (1) Filter if necessary and leave to cool (1) Filter and dry (in air) (1)				
		(iv)	Take a melting point and compare it with a book value	[1]			
		(v)	$\frac{2.75}{1.1} = 2.5 \text{ cm}^3$	[1]			
		(vi)	Moles methylbenzoate = $\frac{2.75}{136} = 0.0202$ (1)				
			Moles methyl 3-nitrobenzoate = $\frac{270}{181} = 0.0149$ (1)				
			% yield = 73.8 % (1)	[3]			
		(vii)	I Electrophilic substitution	[1]			
			II NO ₂ ⁺	[1]			
			Tota	l [20]			

Total Section B [40]