



# **GCE MARKING SCHEME**

**CHEMISTRY  
AS/Advanced**

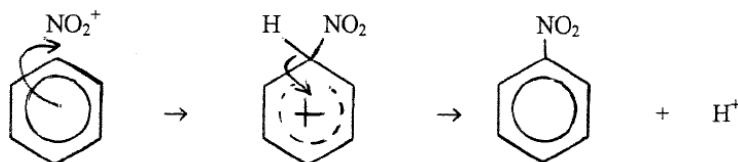
**SUMMER 2011**

## CHEMISTRY - CH4

Q.1 (a) (i) Chromophore [1]

(ii) Yellow transmitted (or reflected) / other colours (e.g. blue and red) absorbed [1]

(b) (i)



(arrow must come from the bond)

[1]

(ii)  $C_3H_2NO_2$  [1]

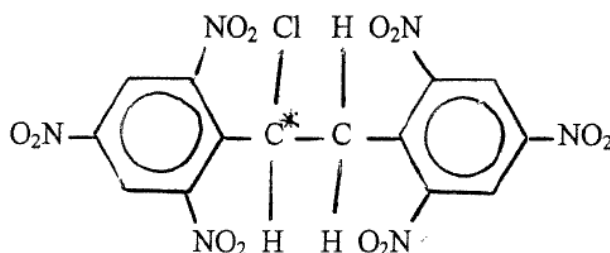
(iii)  $H_2SO_4$  is losing a proton (to another species and becoming an  $HSO_4^-$  ion, acids are proton donors). [1]

(c) The benzene ring is more stable than an alkene because of its delocalised electron structure /  $\pi$  electron system / OWTTE (1)  
If benzene underwent addition this would mean disrupting this stable electron system and this would require relative more energy / activation energy would be (much) higher. (1) [2]

QWC Legibility of text; accuracy of spelling, punctuation and grammar; clarity of meaning. [1]

(d) (i) There is no rotation about a double bond / each 'end' of the double bond has two different 'groups' attached to it [1]

(ii) I

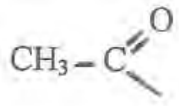


[1]

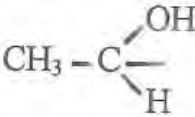
II They are mirror image forms (1) that rotate plane polarised light in opposite directions (1) [2]

III Elimination [1]

**Total [13]**

- Q.2** (a) (i) Sodium borohydride / sodium tetrahydridoborate(III) / lithium aluminium hydride / sodium and ethanol / zinc and ethanoic acid (accept correct formulae) [1]
- (ii) The absorption at  $\sim 1700 \text{ cm}^{-1}$  is due to the C = O bond (1)  
As the reaction proceeds the intensity of this absorption becomes smaller because the butanal is being used up / butan-1-ol does not contain a C = O bond (1) [2]
- (b) (i) butan-1-ol > propanone > ethanol [1]
- (ii) Propanone [1]
- (iii) *Compounds* propanone / ethanol / propan-2-ol  
*Any two for one mark* [1]
- Explanation* the compounds that give a positive iodoform test have to contain a
- 

or the



grouping
- [1]
- (iv) So that a valid comparison can be made between results from other teams / OWTTE [1]  
(do not accept 'fair test')
- (c) There is a balance between the 'carbon' produced by burning and the 'carbon' absorbed by the plant (1)  
When butan-1-ol is burnt carbon dioxide is produced, but this is used by plants / in photosynthesis to produce cellulose (1) [2]
- (d) (i)  $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{OH} + \text{CH}_3\text{COOH} \rightarrow \text{CH}_3\text{COOCH}_2\text{CH}_2\text{CH}_2\text{CH}_3 + \text{H}_2\text{O}$  [1]  
accept  $\text{C}_4\text{H}_9\text{OH}$  but not  $\text{C}_4\text{H}_{10}\text{O}$  - functional groups must be present
- (ii) (concentrated) sulphuric acid /  $\text{H}_2\text{SO}_4$  / hydrogen chloride (gas) /  $\text{HCl(g)}$  [1]  
do not accept  $\text{H}_2\text{SO}_4(\text{aq})$  /  $\text{HCl}$

**Total [12]**

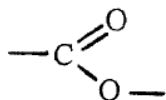
- Q.3** (a) (Free) radical [1]
- (b)  $2\text{C}_3\text{H}_6 + 9\text{O}_2 \rightarrow 6\text{CO}_2 + 6\text{H}_2\text{O}$  [1]
- (c) (i) It is providing a pair of electrons to bond to a proton / acting as a lone pair donor / proton acceptor [1]
- (ii) I A process of boiling / evaporation and condensation without loss (of reactants) [1]
- II By using an electrical heater / or a suitable heating bath / heating mantle - do not accept 'water bath' [1]
- (d) Halothane would cause the most damage as it contains a weaker C-Cl / C-Br bond (1), which is broken in the upper atmosphere (1) (producing radicals that attack ozone).  
Desflurane does not contain C-Cl / C-Br bonds, only the more stable C-F bonds. [2]
- (e) (i) Purple colour / solution / complex - do not accept 'precipitate' [1]
- (ii) I

<b>Compound</b>	<b>Colour given with Universal Indicator paper</b>	<b>Reaction with sodium hydrogencarbonate solution</b>
propofol	yellow / orange	no reaction
compound <b>L</b>	~~~~~	~~~~~
compound <b>M</b>	orange / red	fizzing

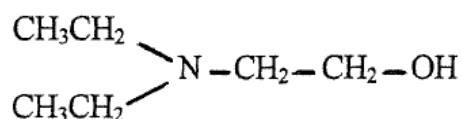
One mark for each correct column [2]

II Gas evolved turns 'lime water' milky [1]

- (f) (i) [1]



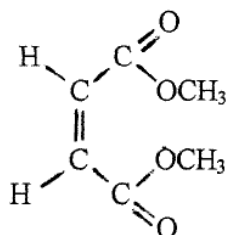
- (ii) [1]



- (g) It would melt at a lower temperature (than 89 °C) / below 89 °C (1)  
and over a range of temperature / not a sharp melting temperature (1) [2]

**Total [15]**

- Q.4** (a) (i) Stereoisomerism is where the compound has the same structural formula but whose atoms / groups take up different positions in space / in three dimensions (1)



[2]

- (ii) The signal at 3.8  $\delta$  due to the methoxy protons (1) would disappear and be replaced by a signal at 11.0  $\delta$  (1) due to the OH protons (1). These protons would have peak area 2 (rather than peak area 6 for the methoxy protons) (1)  
The signal at 6.9  $\delta$  would be (largely) unchanged (1) as the C–H bond is not affected by the hydrolysis of the ester. [5]

QWC Selection of a form and style of writing appropriate to purpose and to complexity of subject matter [1]

- (iii)  $C_6H_8O_4 \rightarrow 144$   
113 is 31 less, could be  $CH_3O$  (1)  
ion could be  $C_5H_5O_3^+$  (1) [2]

- (b) (i) Raw material prices become cheaper / reduce the reaction temperature / use a method where the % yield is increased [1]
- (ii) Use a different (more economic) starting material / find a way of reducing the time taken for fermentation / effect an easier separation method.  
Do not accept reference to increased amounts of enzyme /bigger batch. [1]
- (iii) Number of moles of fumaric acid expected = 140 (1)

Actual number of moles of fumaric acid obtained =  $\frac{13.0 \times 1000}{116} = 112$  (1)

$$\% \text{ Yield} = \frac{112 \times 100}{140} = 80 \quad (1)$$

Alternatively

180 g / kg of glucose give 2 x 116 g / kg of fumaric acid (1)

$\therefore$  1 g / kg of glucose gives  $\frac{2 \times 116}{180}$  g / kg of fumaric acid

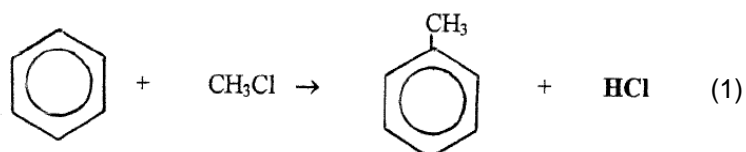
$\therefore$  12.6 kg of glucose gives  $\frac{2 \times 116 \times 12.6}{180}$  kg of fumaric acid = 16.2(4) kg (1)

$$\% \text{ Yield} = \frac{13.0 \times 100}{16.2} = 80 \quad (1)$$

[3]

- (iv) I starting material (1) e.g. ethanol / ethanal OR ethyl ethanoate OR ethanoyl chloride
- reagent (1)  $\text{Cr}_2\text{O}_7^{2-} / \text{H}^+$  acid(aq) / base (aq) followed by acidification water
- type of reaction (1) oxidation / redox hydrolysis hydrolysis [3]
- II platinum / nickel [1]
- (c) e.g. Tollens reagent - silver mirror
- OR 2,4 - dinitrophenylhydrazine - yellow/ orange / red precipitate
- OR Fehling's / Benedict's reagent - brown / red precipitate [1]
- Total [20]**

Q.5 (a) (i)



catalyst - aluminium chloride (1) [2]

(ii) Mass of methylbenzene = 27.6 g (1)

$$\text{Moles of methylbenzene} = \frac{27.6}{92.1} = 0.30(0) \quad (1)$$

$\therefore$  0.30 mole of  $\text{C}_6\text{H}_5\text{CH}_2\text{Cl}$  should be made  
this will have a mass of  $0.30 \times 126.6 = 38.0$  g (1)

$\therefore$  Mass of flask + product needs to be  $120.4 + 38.0 = 158.4$  g (1) [4]

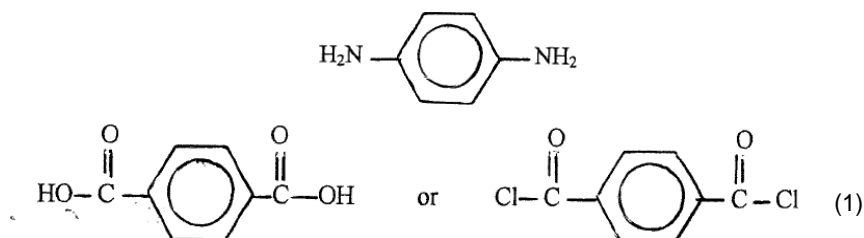
(iii) I potassium cyanide [1]

II lithium tetrahydridoaluminate(III) / lithium aluminium hydride  
(accept correct formulae) [1]

(b) The nitrogen atom is electron rich / has a lone pair (1) and will act as a proton acceptor / electron pair donor (1) [2]

(c) 2-Phenylethylamine reacts with nitrous acid giving an alcohol (1) and evolving nitrogen gas as bubbles (1)  
4-Ethylphenylamine gives a diazonium compound (1) [3]

(d)



[2]

(e) (i) 2-amino-3-hydroxypropanoic acid [1]

(ii) Hydrogen bonding occurs because of the difference in electronegativity between hydrogen and oxygen / nitrogen (in O-H and N-H bonds), (1), leading to polar covalent bonds /  $\delta^+$ ,  $\delta^-$  (1)  
There are attractive forces between the oxygen or nitrogen of one molecule and the hydrogen atom bonded to an oxygen or nitrogen atom of another molecule (1) [3]

(Marks can be obtained from a suitable diagram)

QWC Information organised clearly and coherently, using specialist vocabulary when appropriate [1]

**Total [20]**