



Rewarding Learning

**ADVANCED
General Certificate of Education
2019**

Chemistry
Assessment Unit A2 3
assessing
Further Practical Chemistry
Practical Booklet B (Theory)
[ACH32]

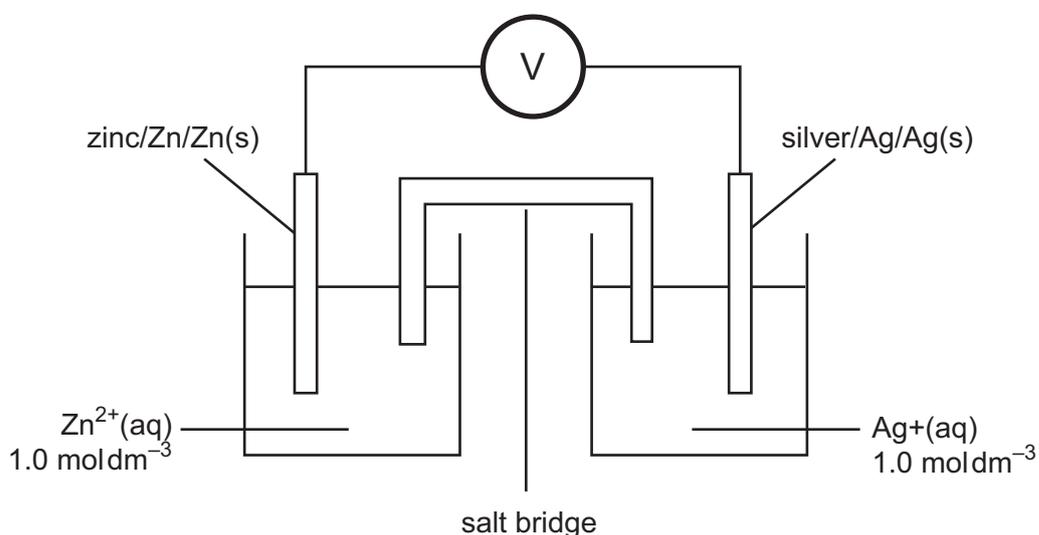
WEDNESDAY 19 JUNE, MORNING

**MARK
SCHEME**

- 1 (a) Run with solvent [1]
 Dry and mark the solvent front [1]
 Rotate by 90° (anti-clockwise) [1]
 Locate with, e.g. ninhydrin, iodine or UV light [1] [4]
- (b) Solvent 1 (6 ÷ 10 =) 0.6 Solvent 2 (9 ÷ 10 =) 0.9 [1]
- (c) Repeat the two-way chromatography using separate samples of pure leucine and serine [1]
 Second mark dependent on first
 Compare the positions of the spots/R_f values [1] [2]
- (d) Two-way chromatography improves separation/greater separation between the spots [1]
- (e) (i) Only one peak in the chromatogram [1]
 (ii) The retention time matches that of the pure drug [1]
 The mass spectrum matches that of the pure drug [1] [2]

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- 2 (a) The potential difference/voltage/emf measured when a half-cell is connected to the standard hydrogen electrode under standard conditions [2]
- (b) (i) 298K (or 25° C)



- each missing detail [-1] [4]
- (ii) $\text{emf} = 0.80 - (-0.76) = +1.56 \text{ V}$ [1]
- (c) $+1.61 = -0.76 - (x)$
 $x = -2.37 \text{ V}$ [1]

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- 3 (a) (i)** Aldehyde [1]
 The formation of the orange solid suggests that A is either an aldehyde or ketone/contains a carbonyl group [1] and the formation of the silver mirror means it is not a ketone [1] [3]
- (ii)** Propanal and $\text{CH}_3\text{CH}_2\text{CHO}^+$ [1]
- (b) (i)** Carboxylic acid [1]
 The vigorous reaction with phosphorus pentachloride suggests that B contains an $-\text{OH}$ group [1], the effervescence with sodium carbonate suggests that B is a carboxylic acid/acidic [1] [3]
- (ii)** $\text{CH}_3\text{CH}_2\text{COO}^-\text{H}^+$ [1]
 shift should be in the range 10.0–12.0 ppm [1]
 Second mark dependent on the first [2]
- (iii)** CH_3 triplet – two hydrogens on the adjacent carbon [1]
 CH_2 quartet – three hydrogens on the adjacent carbon [1] [2]

AVAILABLE
MARKS

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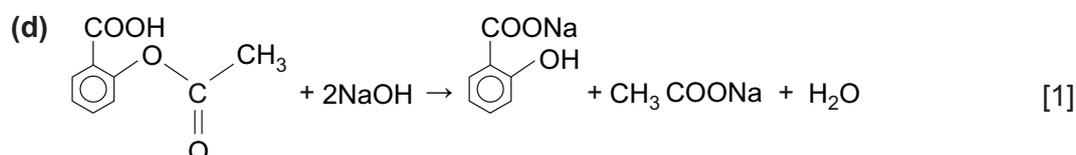
- 4 (a)** Either:

RMM of aspirin = $(9 \times 12) + (8 \times 1) + (4 \times 16) = 180$
 Actual yield of aspirin = $18.0/180 = 0.1$ mole
 Theoretical yield of aspirin = $(0.1/40) \times 100 = 0.25$ mole
 RMM of salicylic acid = $(7 \times 12) + (6 \times 1) + (3 \times 16) = 138$
 Mass of salicylic acid = $0.25 \times 138 = 34.5$ g
 Error [–1]

Or:

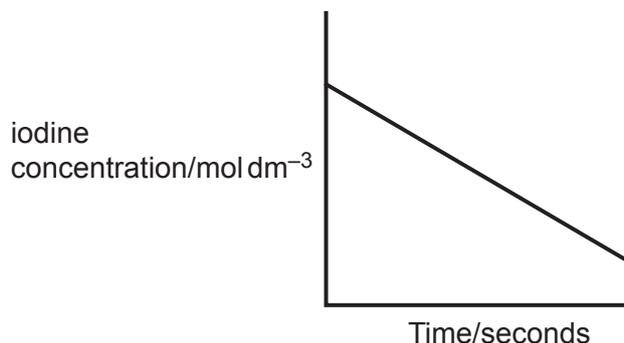
Theoretical yield of aspirin = $(18.0/40) \times 100 = 45$ g
 RMM of aspirin = $(9 \times 12) + (8 \times 1) + (4 \times 16) = 180$
 Theoretical yield of aspirin = $45/180 = 0.25$ mole
 RMM of salicylic acid = $(7 \times 12) + (6 \times 1) + (3 \times 16) = 138$
 Mass of salicylic acid = $0.25 \times 138 = 34.5$ g
 Error [–1] [3]

- (b)** Dissolve the crude product in the minimum volume of hot water/methanol/ethanol [1]
 Filter while hot [1]
 Allow filtrate to cool [1]
 Filter crystals under suction [1] [4]
- (c)** place some solid in a capillary tube sealed at one end [1]
 place the capillary tube in melting point apparatus/oil bath [1]
 heat slowly [1]
 record temperature at which melting begins and ends [1]
 a sharp melting point confirms the purity of the aspirin [1] [5]



- 5 (a) • plot a calibration curve/graph (absorption v $[I_2]$) [1]
 • using known concentrations of iodine [1]
 • place reaction tube mixture in colorimeter recording absorption v time [1]
 • convert absorbance to $[I_2]$ (using calibration graph) [1] [4]

(b) (i)



axes correctly labelled [1]
 straight line with negative gradient [1] [2]

(ii) value of the gradient = rate of reaction [1]

(c) so that concentrations of propanone and hydrogen ions remain (effectively) constant [1] 8

6 (a) (i) Acidified manganate(VII) acts as (its own) indicator/titration is self-indicating [1]

(ii) colourless [1] to pink [1] [2]

(b) Moles of manganate(VII) = $0.020 \times (18.0/1000) = 3.6 \times 10^{-4}$

1:5 ratio of (MnO_4^-) : Fe^{2+}

Moles of iron(II) = $5 \times 3.6 \times 10^{-4} = 1.8 \times 10^{-3}$ [2]

(c) (i) $Zn + 2 Fe^{3+} \rightarrow Zn^{2+} + 2 Fe^{2+}$ [1]

reducing agent/ reduces iron(III) to iron(II) [1] [2]

(ii) Moles of manganate(VII) = $0.02 \times (30.0/1000) = 6.0 \times 10^{-4}$

Moles of iron(II) = $5 \times 6.0 \times 10^{-4} = 3.0 \times 10^{-3}$

Increase in moles of iron(II) = $3.0 \times 10^{-3} - 1.8 \times 10^{-3} = 1.2 \times 10^{-3}$

% increase = $(1.2 \times 10^{-3}/3.0 \times 10^{-3}) \times 100 = 40\%$

or

$(30.0 - 18.0)/30.0 \times 100 = 40\%$ [2]

Total

60