

Electrolysis, Electrode Potentials & Cells

Mark Scheme 6

Level	International A Level
Subject	Chemistry
Exam Board	CIE
Topic	Electrochemistry
Sub-Topic	Electrolysis, Electrode Potentials & Cells
Paper Type	Theory
Booklet	Mark Scheme 6

Time Allowed: 64 minutes

Score: /53

Percentage: /100

Grade Boundaries:

A*	A	B	C	D	E	U
>85%	77.5%	70%	62.5%	57.5%	45%	<45%

- 1 (a) $\text{O}_2 + 4\text{H}^+ + 4\text{e}^- \longrightarrow 2\text{H}_2\text{O}$ (or equation $\div 2$) [1]
- (b) \oplus [1]
- (c) 1.23 (V) (ignore sign) [1]
- (d) a better/larger salt bridge *or* a diaphragm *or* larger (area of) electrodes
or increase concentrations/pressure [1] 1
- (e) time = $400 \times 24 \times 60 \times 60 = 34\,560\,000$ seconds [1]
charge = current \times time = $0.01 \times 34\,560\,000 = 345\,600$ C ecf [1]
moles of H = $345\,600/96\,500 = 3.6$ mol \therefore mass of H = **3.6 g** ecf [1] 3
- (f) advantages: less pollution/ CO_2/NO_x etc. *or* cleaner by-products
less dependence on fossil fuels/finite resources any one [1]
- disadvantages: more expensive (to develop *or* to run)
takes up more space
poor power-to-volume ratio
hydrogen is difficult to store or to transport any one [1]
NOT hydrogen is explosive/flammable 2
- Total 9**

CHEMISTRY ONLINE
— TUITION —

- 2 (a) The EMF of a cell made up of the test electrode and a standard hydrogen electrode. [1]

EMF measured under standard conditions of T, P and concentration [1]

2

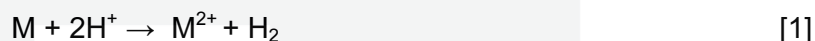
(b) (i) $E_{\text{left}} = E_{\text{right}} - E_{\text{cell}} = 0.34 - 0.76 = -0.42 \text{ (V)}$ [1]

(ii) \longrightarrow (arrow from left to right) [1]

(iii) I pink/red solid/ppt or copper will be formed or blue solution fades or M dissolves/corrodes [1]



II hydrogen/gas evolved or M dissolves (do not allow "M dissolves" for [2] marks in both I and II) [1]



6

(c) (i) polarity of d. c. source: \ominus is on the left, \oplus is on the right [1]

electrolyte is $\text{Cu}^{2+}(\text{aq})/\text{CuSO}_4/\text{CuCl}_2/\text{Cu}(\text{NO}_3)_2$ etc. or name [1]

(ii) moles of Cu = $0.5/63.5 = 7.87 \times 10^{-3}$ [1]

moles of $e^- = 2 \times 7.87 \times 10^{-3} = 1.57 \times 10^{-2}$

no. of coulombs = $96500 \times 1.57 \times 10^{-2} = 1517 \text{ (C)}$ [1]
ecf in $n(e^-)$

time = $1520/0.5 =$ [1]
ecf in coulombs

5

Total 13

Question	Scheme	Mark	Total																
3 (a)	<table><tr><td>name of particle</td><td>relative mass</td><td>relative charge</td></tr><tr><td>proton</td><td></td><td>+</td></tr><tr><td>electron</td><td>1/1836</td><td>–</td></tr><tr><td>neutron</td><td></td><td>0</td></tr></table>	name of particle	relative mass	relative charge	proton		+	electron	1/1836	–	neutron		0	[1] [1] [1]	[3]				
name of particle	relative mass	relative charge																	
proton		+																	
electron	1/1836	–																	
neutron		0																	
(b) (i)	Mass of an atom(s) relative to 1/12 th (the mass) of (an atom of) carbon-12 OR relative to carbon-12 which is (exactly) 12	[1] [1]	[2]																
(ii)	% of third isotope = 10 $\frac{(24 \times 79) + (26 \times 11.0) + 10x}{100} = 24.3$ 10x = 248 x = 24.8 (3s.f.)	[1] [1] [1]	[3]																
(c) (i)	anode $I^{-} \rightarrow Cl_2 + 2e^{-}$ cathode $Mg^{2+} + 2e^{-} \rightarrow Mg$	[1] [1]	[2]																
(ii)	<table><tr><td>Mg</td><td>O</td><td>H</td><td>Cl</td></tr><tr><td>31.65</td><td>20.84</td><td>1.31</td><td>46.2</td></tr><tr><td>24.3</td><td>16</td><td>1</td><td>35.5</td></tr><tr><td>1.30</td><td>1.30</td><td></td><td></td></tr></table> MgOHC ₂ Cl	Mg	O	H	Cl	31.65	20.84	1.31	46.2	24.3	16	1	35.5	1.30	1.30			[1] [1]	[2]
Mg	O	H	Cl																
31.65	20.84	1.31	46.2																
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1.30	1.30																		
(d) (i)	Na ₂ O basic/alkaline; Al ₂ O ₃ amphoteric/acidic and basic; SO ₃ acidic Na ₂ O (giant) ionic AND SO ₃ (simple/molecular) covalent	[1] [1]	[2]																
(ii)	Na ₂ O + 2HCl → 2NaCl + H ₂ O Al ₂ O ₃ + 6HCl → 2AlCl ₃ + 3H ₂ O Al ₂ O ₃ + 2NaOH + 7H ₂ O → 2NaAl(OH) ₄ (H ₂ O) ₂ OR Al ₂ O ₃ + 2NaOH + 3H ₂ O → 2NaAl(OH) ₄ OR Al ₂ O ₃ + 2NaOH → 2NaAlO ₂ + H ₂ O OR Al ₂ O ₃ + 2OH [–] + 7H ₂ O → 2[Al(OH) ₄ (H ₂ O) ₂] [–] OR Al ₂ O ₃ + 2OH [–] + 3H ₂ O → 2[Al(OH) ₄] [–] OR Al ₂ O ₃ + 2OH [–] → 2AlO ₂ [–] + H ₂ O SO ₃ + NaOH → NaHSO ₄ OR SO ₃ + 2NaOH → Na ₂ SO ₄ + H ₂ O	[1] [1] [1] [1] [1] [1] [1]	[4]																

4 (a) (i)	$K_{sp} = [Ag^+(aq)]^2[SO_4^{2-}(aq)]$ and units: mol^3dm^{-9}	1
(ii)	$K_{sp} = (2 \times 0.025)^2 \times (0.025) = 6.25 \times 10^{-5}$	1
(b)	<p>Diagram illustrating the enthalpy changes for the dissolution of silver sulfate:</p> <ul style="list-style-type: none"> $2Ag^+(g) + SO_4^{2-}(g)$ (Gaseous ions) $Ag_2SO_4(s)$ (Solid) $Ag_2SO_4(aq)$ or $2Ag^+(aq) + SO_4^{2-}(aq)$ (Aqueous ions) Enthalpy changes: ΔH°_{latt} (lattice enthalpy), ΔH°_{sol} (enthalpy of solution), ΔH°_{hyd} (enthalpy of hydration). 	1 1 1 1
(c) (i)	$E^\ominus_{cell} (= 0.80 - 0.77 =) (+)0.03V$ and Ag^+/Ag or $Ag/silver$ or right	1
(ii)	E_{cell} would be less positive/more negative because the $[Ag^+(aq)]$ (in the Ag electrode) is less than $1.0 mol dm^{-3}$	1
(iii)	<ul style="list-style-type: none"> no change 	1
	<ul style="list-style-type: none"> more negative/less positive 	1
(iv)	the $[Ag^+(aq)]$ will decrease $E_{electrode}$ becomes less positive or due to the common ion effect	1
(d)	$[Fe^{3+}(aq)] = 0.2 mol dm^{-3}$ $[H^+] = \sqrt{c.K_a} = \sqrt{0.2 \times 8.9 \times 10^{-4}}$ or $1.33 \times 10^{-2} (mol dm^{-3})$ $pH = -\log([H^+]) = 1.9$ (or 1.87–1.89)	1 1
[Total: 13]		