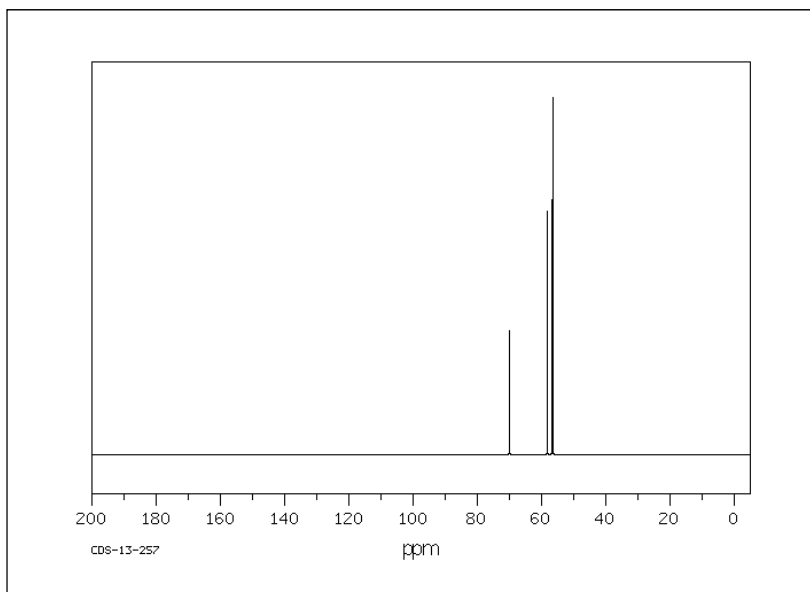


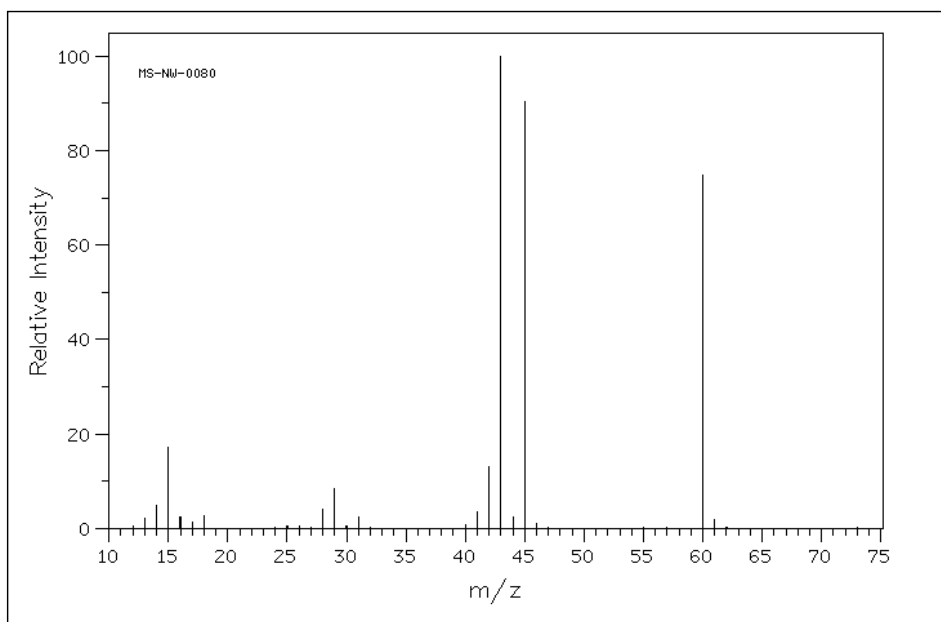
This is the ^1H NMR spectrum of acetylcholine – it is only species with **four different hydrogen environments**. ❶

Spectrum 2.



This is the ^{13}C NMR spectrum of choline – it is only species with **three different carbon environments**. ❶

Spectrum 3.



This is the mass spectrum of ethanoic acid (acetic acid) – the parent peak at $m/z = 60$ corresponds with the molecular mass of CH_3COOH . ❶

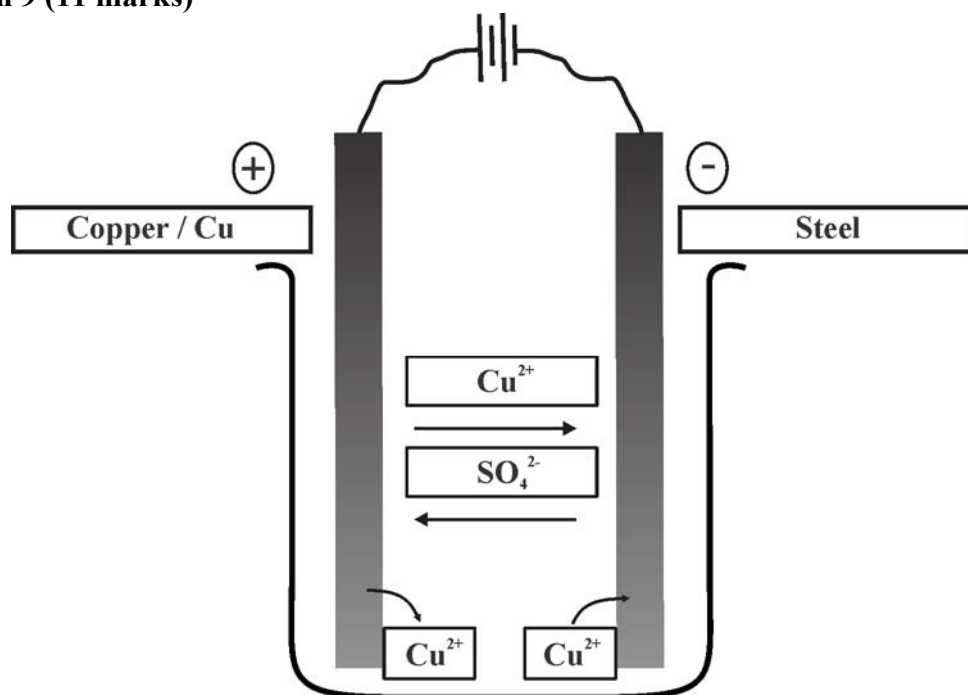
- ii. The peak with relative intensity at $m/z = 43$ is the **base peak** and is associated with the **most abundant species** resulting from fragmentation in the mass spectrometer.

Species formula – $[\text{CH}_3\text{CO}]^+$ ❶

- c. **Compounds in nerve agents binding to serine in *acetylcholinesterase* would disrupt its tertiary structure and so interfere with the active site on the enzyme.** ❶
This would reduce the ability of the enzyme to break down acetylcholine.
- d. Pralidoxime chloride: $C_7H_9N_2OCl$
 $m(C_7H_9N_2OCl)$ required in 20 minutes = $30 \text{ mg kg}^{-1} \times 82.5 \text{ kg}$
 $= 2.5 \times 10^3 \text{ mg} = 2.5 \text{ g}$
 $n(C_7H_9N_2OCl)$ required in 20 minutes = $2.5 \text{ g} / 172.5 \text{ g mol}^{-1}$
 $= 1.4 \times 10^{-2} \text{ mol}$ ❶
 $n(C_7H_9N_2OCl)$ required in 60 minutes = $1.4 \times 10^{-2} \text{ mol} \times 3$
 $= 4.3 \times 10^{-2} \text{ mol}$
 Rate administered = $4.3 \times 10^{-2} \text{ mol / hour}$ ❶

Question 9 (11 marks)

a.



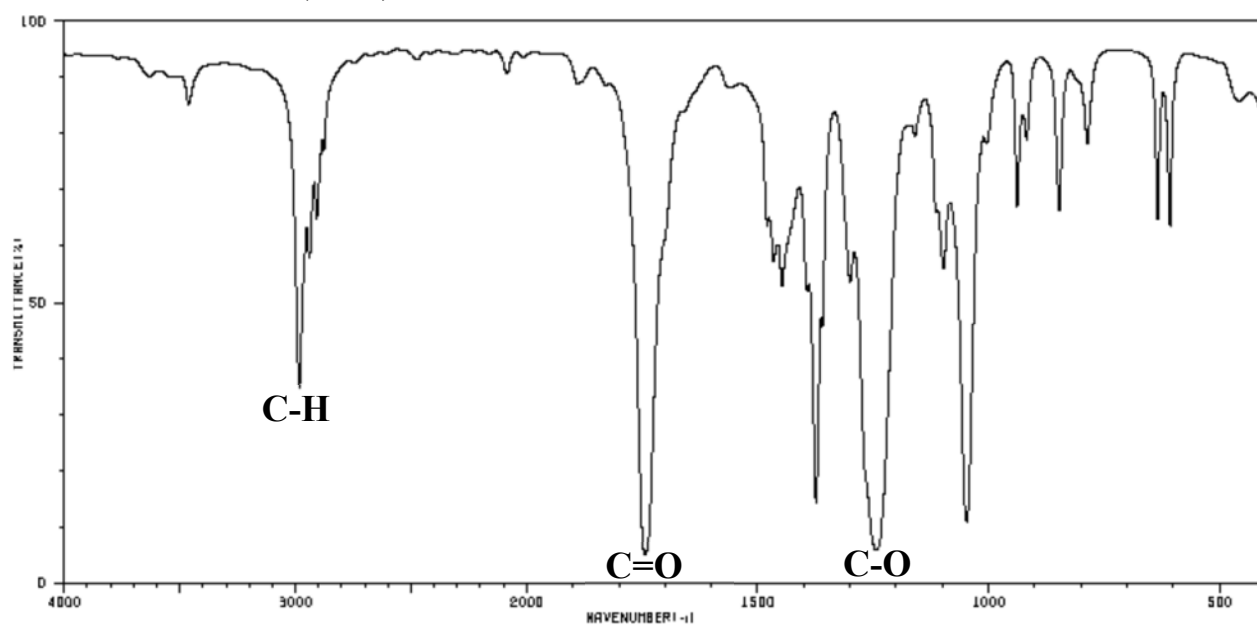
- ❶ for correct electrode signs.
 ❶ for correct labelling of electrodes.
 ❶ for correct species identification in electrolyte.
- b. Independent variable – **current**.
 Dependent variable – **mass of Cu deposited** / change in mass of plate. ❶
- c. Three from
 Electrolyte concentration, electrode surface area, distance between electrodes, voltage, temperature and time. ❶
- d. % efficiency = $[\text{actual } m(\text{Cu}) \text{ deposited} / \text{theoretical } m(\text{Cu}) \text{ deposited}] \times 100$
 Reduction half-equation = $\text{Cu}^{2+}(\text{aq}) + 2\text{e}^- \rightarrow \text{Cu}(\text{s})$
 Clarisse: Actual $m(\text{Cu})$ deposited = 0.395 g
 $Q = It = 1.45 \times 15 \times 60 = 1.3 \times 10^3 \text{ C}$
 $n(\text{e}^-) = Q/F = 1.3 \times 10^3 / 96500 = 0.014 \text{ mol}$ ❶
 Theoretical $m(\text{Cu})$ deposited = $\frac{1}{2} \times 0.014 \times 63.5$
 $= 0.43 \text{ g}$ ❶
 % efficiency = $(\text{actual mass} / \text{theoretical mass}) \times 100$
 $= (0.395 / 0.43) \times 100$
 $= 92 \%$ ❶

- e. Possible reasons.
- Masud did not dry the copper plated electrode prior to weighing. ❶
 - Clarisse accidentally removed some of the deposited copper when drying the copper plated electrode.
 - Some of the copper deposited on Masud's electrode was oxidised during drying.
- f. $\text{Cu}^{2+}(\text{aq})$ is simultaneously produced at the anode and consumed at the cathode. ❶
- g. In the original analysis, some of copper produced at the steel electrode may not stick to the electrode or may be accidentally removed during drying, making the recorded change in the electrode mass lower than the mass of copper oxidised from the anode. Whilst this would be less of an issue if the copper anode was weighed before and after the electrolysis the **results obtained would not be a true experimental outcome.** ❶

Question 10 (9 marks)

- a. During the discharge the strongest oxidant is reacting with the strongest reductant. Since $\text{Ce}^{3+}(\text{aq})$ is a weaker reductant than $\text{Zn}(\text{s})$ then $\text{Ce}^{4+}(\text{aq})$ must be a stronger oxidant than $\text{Zn}^{2+}(\text{aq})$.
During discharge $\text{Zn}(\text{s})$ is oxidised at the (-) **electrode – Zn**, whilst $\text{Ce}^{4+}(\text{aq})$ is reduced at the (+) **electrode – graphene oxide – graphite composite**. So $\text{Zn}^{2+}(\text{aq})$ is **pumped over the (-) electrode** and $\text{Ce}^{4+}(\text{aq})/\text{Ce}^{3+}(\text{aq})$ is **pumped over the (+) electrode.** ❶
- (-) $\text{Zn}(\text{s}) \rightarrow \text{Zn}^{2+}(\text{aq}) + 2\text{e}^-$
(+) $\text{Ce}^{4+}(\text{aq}) + \text{e}^- \rightarrow \text{Ce}^{3+}(\text{aq})$ ❶
- b. **During recharging Zn(s) is deposited on the (-) electrode** due to
 $\text{Zn}^{2+}(\text{aq}) + 2\text{e}^- \rightarrow \text{Zn}(\text{s})$ ❶
- c. $2.20 \text{ V} = E^\circ(\text{oxidant half-cell}) - E^\circ(\text{reductant half-cell}) = E^\circ(\text{Ce}^{4+}/\text{Ce}^{3+}) - E^\circ(\text{Zn}^{2+}/\text{Zn})$
 $= E^\circ(\text{Ce}^{4+}/\text{Ce}^{3+}) - (-0.76) - \text{from Table 2 of the VCE Chemistry Data Book}$
 $= E^\circ(\text{Ce}^{4+}/\text{Ce}^{3+}) + 0.76$
 $E^\circ(\text{Ce}^{4+}/\text{Ce}^{3+}) = 2.20 - 0.76$
 $= 1.44 \text{ V}$ ❶
- d. In a conventional rechargeable battery all the reactants are within the cell. However, in the **redox flow battery the reactants are stored outside the cell and pumped in as required as happens in fuel cells.** However, **electrical recharging is not characteristic of fuel cells.** ❶
- e. i. General reaction pathway from $\text{CH}_3\text{CH}_2\text{Cl}$ with no other organic compounds involved is
 $\text{CH}_3\text{CH}_2\text{Cl} \rightarrow \text{CH}_3\text{CH}_2\text{OH} \rightarrow \text{CH}_3\text{CHO} \rightarrow \text{CH}_3\text{COOH} \rightarrow \text{CH}_3\text{COOCH}_2\text{CH}_3$
 The mass spectrum shows **parent ion peak at $m/z = 88.0$ and base peak at $m/z = 43.0$.** ❶
 The ^1H NMR spectrum shows **three hydrogen environments** with a **quartet, singlet, triplet** splitting pattern.
 This data is consistent with the ester **ethyl ethanoate $\text{CH}_3\text{COOCH}_2\text{CH}_3$** ❶
 $M_r(\text{CH}_3\text{COOCH}_2\text{CH}_3) = 88.0$
 - $m/z = 43.0$ is consistent with **CH_3CO^+ fragment**
 - **three different hydrogen environments**
 - **quartet, triplet** is consistent with **$-\text{CH}_2\text{CH}_3$** ❶
 Also, signal at $\delta = 2.0$ ppm is consistent with CH_3COOR ,
 signal at $\delta = 4.0$ ppm is consistent with RCOOCH_2R .

- ii. Use **Table 14 in the VCE Chemistry Data Book** to identify peaks for C-H, C-O and C=O(esters) bonds.



① for all 3 correct

End of Suggested Answers