

- 1. $\mathbf{B} 0.250 \text{ mol} \times 6.02 \times 10^{23} \text{ mol}^{-1} = 1.51 \times 10^{23} \text{ molecules}.$
- 2. **B** 6 protons + 7 neutrons = mass number 13, giving ¹³ C.
- 3. \mathbf{A} sulfur (Z = 16) fills orbitals to 3p⁴.
- 4. $\mathbf{C} n(C_2H_5OH) = 4.50 \text{ g} / 46 \text{ g mol}^{-1} = 0.0978 \text{ mol}; 2 \text{ mol } CO_2 \text{ per mol ethanol} \rightarrow 0.196 \text{ mol } CO_2.$
- 5. **B** The concentration falls from about 0.083 mol dm⁻³ to 0.060 mol dm⁻³ in 10 s (Δ [H₂O₂] \approx 0.023 mol dm⁻³). Rate = 0.023 / 10 \approx 2.3 \times 10⁻³ mol dm⁻³ s⁻¹.
- 6. **C** NH₃ has four electron domains (one lone pair) giving a trigonal pyramidal shape.
- 7. **B** mobile electrons allow charge flow, giving electrical conductivity.
- 8. **C** Raising the temperature spreads the energy distribution: more particles have higher energies, the peak moves right and lowers, but particle number (area) stays the same.
- 9. **A** delocalised electrons in graphite allow conduction; diamond has none and is extremely hard.
- 10. **A**. Liquid M contains four carbon atoms and a terminal -CHO group, so the correct name is butanal.
- 11. **D** low temperature and high pressure enhance intermolecular forces and particle volume effects.
- 12. **C** 0.12 g Mg ÷ 24.3 g mol⁻¹ = 4.94 × 10⁻³ mol. Mg + Cl₂ \rightarrow MgCl₂ (1 : 1). Mass MgCl₂ = 4.94 × 10⁻³ mol × 95.3 g mol⁻¹ ≈ 0.47 g.
- 13. **A –** Q = 500 g × 4.18 J g^{-1} °C⁻¹ × 10.9 °C = 2.28 × 10⁴ J = 22.8 kJ

Fuel mass burned = 20.24 g - 19.48 g = 0.76 g

 $\Delta H_{\text{(comb)}}$ per gram = 22.8 / 0.76 \approx 3.0 \times 10¹ kJ g⁻¹ (exothermic)

14. C - Applying Hess's law:

2
$$\Delta_a$$
 □ H(Ag) + 1462 + 249.2 - 141.1 + 798 - 2969 = -31 → 2 Δ_a □ H(Ag) ≈ 570 kJ mol⁻¹ → Δ_a □ H(Ag) ≈ 285 kJ mol⁻¹.

- 15. **A**. The H-O-N angle is determined by a bent arrangement around the oxygen (≈104°), while the nitrogen is trigonal-planar, giving an O-N-O angle of ≈120°.
- 16. **D** higher T favours the endothermic reverse reaction, shifting left and lowering Kc.



- 17. **C** Silicon forms a tetrahedral network of strong covalent bonds throughout the lattice, requiring very high energy to break.
- 18. **A** energy/CO₂: CH₄ = 890 kJ; ethanol ≈ 685 kJ; propane ≈ 740 kJ; octane ≈ 684 kJ highest for methane.
- 19. **C** The spectrum displays four distinct groups of multiplets (ignoring the TMS reference at 0 ppm), indicating four chemically non-equivalent sets of protons.
- 20. **C** NH₃ accepts H⁺ to form NH₄⁺, so is the base.
- 21. **B** − [H⁺] = $\sqrt{\text{(Ka} \times \text{c)}}$ = 4.24 × 10⁻⁴ mol dm⁻³; pH = 3.37 ≈ 3.38.
- 22. **A** combining the balanced half-equations in acidic conditions yields 2 ClO₂ + SO₃²⁻ + $H_2O \rightarrow 2$ Cl⁻ + SO₄²⁻ + 2 H⁺.
- 23. **B** Zinc is oxidised to Zn²⁺/ZnO at the anode, releasing electrons that travel through the external circuit from Zn to Aq. At the cathode, Aq₂O is reduced:

$$Ag_2O + H_2O + 2e^- \rightarrow 2Ag + 2OH^-$$

- 24. **A** electrophilic addition of liquid bromine across the C=C gives a di-bromo product.
- 25. **B** hydroxylation occurs by syn addition; the resultant diol is meso due to an internal plane of symmetry.
- 26. **C** NH₃ accepts H⁺ to form NH₄⁺, so is the base.
- 27. **C** CH₃COOCH₃ contains the –COO– linkage characteristic of esters, whereas A is a ketone, B is a carboxylic acid and D is an ether.
- 28. **B** bromine (group 17) is a halogen; the others are misclassified.

29. **B** –
$$\Delta$$
H° = [4(90)+6(-286)] – [4(-46)+0] = -1172 kJ ≈ -1.17 × 10³ kJ.

30. **D** – q =
$$(100 + 1.5)$$
 g × 4.18 J g⁻¹ K⁻¹ × 7.3 K ≈ 3.10 kJ. n(KOH) = 1.500 g / 56.1 g mol⁻¹ = 0.0268 mol. $\Delta H = -q/n \approx -116$ kJ mol⁻¹ (exothermic).