

## Mark Scheme - Electron Configuration (SL)

### Paper 1A

- A

- B

- A

- B

- B

- D

- D

- B

- B

- A

- A

- B

- B

- C

- B

- B

- C

- B

- A

- B

### Paper 1B

- 1. (a) 4 marks: Correct axes with labels (1), all points plotted correctly (1), smooth best-fit line (1), linear trend through origin (1).

- 1. (b) 0.40 absorbance corresponds to a concentration of roughly  $0.08 \text{ mol dm}^{-3}$  (reading from graph) [1].

- 1. (c) At  $\lambda_{\text{max}}$ , the absorbance is highest, which minimizes the relative error in measurements and provides maximum sensitivity [1].

- 2. (a)  $E = hc/\lambda$ .  $E = (6.63 \times 10^{-34} \times 3.00 \times 10^8) / (486 \times 10^{-9}) = 4.09 \times 10^{-19} \text{ J}$  [3].

- 2. (b) Visible Balmer series:  $n_{\text{initial}} > 2$ ,  $n_{\text{final}} = 2$  [1].

## Paper 2

- 1. (a) Degenerate: Orbitals that have the same energy level [1].

- 1. (b)  $\Delta E = hc/\lambda$ ; Wavelength is inversely proportional to splitting energy [1].

- 1. (c) Different ligands exert different electrostatic fields on the d-orbitals [1], causing different splitting gaps ( $\Delta E$ ) and thus absorption of different wavelengths [1].

- 1. (d) 410 nm absorbs violet light. Observed color is the complementary color: Yellow [1].

- 2. (a)  $[\text{Ar}] 3d^{10} 4s^1$  has a fully-filled 3d subshell, which offers extra stability compared to the partially filled subshell in  $[\text{Ar}] 3d^9 4s^2$  [2].

- 2. (b) (i)  $\text{Fe}^{3+}$ :  $[\text{Ar}] 3d^5$  [1]. (ii) Zn:  $[\text{Ar}] 3d^{10} 4s^2$  [1].

- 3. (a) Correct order:  $1s < 2s < 2p < 3s < 3p < 4s < 3d$  [3].

- 3. (b) As the principal quantum number (n) increases, the energy differences between successive levels decrease, causing the emission lines to merge at the high-frequency series limit [2].

- 3. (c) Ionization energy at convergence limit (91.2 nm):  $E = hc/\lambda = (6.63 \times 10^{-34} \times 3.00 \times 10^8) / (91.2 \times 10^{-9}) = 2.18 \times 10^{-18} \text{ J}$  [3].