## Assessment Schedule – 2024

## Chemistry: Demonstrate understanding of thermochemical principles and the properties of particles and substances (91390) Evidence Statement

Q		Evidence				Achievement			ment with erit	Ac	hievement with Excellence
ONE (a)	······································	F. F. Cl. Vramidal squar	2- ;⊂ci e .⊂i 		• TWO	correct.		All corre	ct.		
(b)(i)	$3N_{2}H_{4}(\ell) + 3O_{2}(g) \rightarrow 3N_{2}(g)$ $4ClF_{3}(g) + 4NH_{3}(g) \rightarrow 12HF(g)$ $2N_{2}(g) + 6H_{2}O(g) \rightarrow 4NH_{3}(g)$ $6H_{2}O(\ell) \rightarrow 6H_{2}O(g)$ $\Delta_{r}H = -1869 + -2400 + 1270 + 1$	$(g) + 2N_2(g) + 2Cl_2(g)$ + $3O_2(g)$ (reverse)	) -1: +1 +4	523 × 3 200 × 2 270 0.7 × 6	• Recog values	nises the need to mu	ltiply	Correct p     one error	process with	unit	rect answer with and significant res (accept 2–4).
(ii)	The entropy of the system inc gaseous molecules and 3 more randomised, disordered gaseo and energy in the system. Since the reaction is exotherm into the surroundings, so the p kinetic energy. As a result, the surroundings, so the entropy of The increase in entropy of the entropy change positive, so the	e ordered liquid moleco bus molecules. So, then nic (reaction is explosing particles in the surrour ere is greater dispersa- of the surroundings in e system and of the sur-	cules produce 17 high re is greater dispersal ive), heat energy is re adings gain heat energy l of matter and energy creases. rroundings makes the	of matter eleased gy / y in the	<ul> <li>Recognises that the entropy of the system increases due to the increase in moles/states change.</li> <li>Recognises that the entropy of the surroundings increases because</li> </ul>		OR	or system. ns entropy e for	why	justification of the process is ntaneous.	
N	IØ N1	N2	A3	A		M5		M6	E7		E8

NØ	N1	N2	A3	A4	M5	M6	E7	E8
No response; no relevant evidence.	1a	2a	3a	4a	2m	3m	2e, with minor error / omission.	2e

Q	Evidence	Achievement	Achievement with Merit	Achievement with Excellence
TWO (a)(i)	<ul> <li>3: third energy level / third electron shell</li> <li>p: the orbitals / subshell occupied</li> <li>6: there are a total of six electrons in the p orbitals / subshell.</li> </ul>	• TWO correct.		
(ii)	Ne > Ar > P	• Correct order.		
	Phosphorus and argon are in the same period. Ionisation energy increases across a period. Although the valence electron to be removed is in the same energy level, with the same repulsion / shielding from inner energy levels, the number of protons increases across a period. This means the electrostatic attraction between the positive nucleus and the valence electrons increases across a period, so Ar has a higher first ionisation energy than P. Neon and argon are in the same group. Ionisation energy decreases down a group because the valence electron to be removed is in an energy level further from the nucleus with greater repulsion / shielding from inner energy levels. Although the number of protons increases down a group, this effect is offset by the increasing distance between the nucleus and the valence electron to be removed. This means the electrostatic attraction between the positive nucleus and the valence electrons decreases down a group, so Ne has a higher first ionisation energy than Ar.	• Recognises one factor influencing ionisation energy trend.	• Explains 1st ionisation energy trend with reference to at least TWO of: energy levels, number of protons, repulsion from inner energy levels, strength of electrostatic attraction.	• Justifies the difference in first IE for all THREE elements.
(b)	There is an electronegativity difference between Br and Cl, so the Br–Cl bonds are polar covalent. There are five areas of electron density around the central Br atom, including three bond pairs and two lone pairs / non-bonding pairs. In the trigonal planar shape, the bond dipoles are arranged symmetrically around the central Br atom (due to lone pairs occupying axial positions), so the bond dipoles cancel out to make BrCl <sub>3</sub> a non-polar molecule. In contrast, the T-shaped arrangement has the bond dipoles arranged asymmetrically around the central Br atom (due to lone pairs occupying equatorial positions). As a result, the bond dipoles reinforce / add together / do not cancel out to make BrCl <sub>3</sub> a polar molecule.	<ul> <li>Identifies the correct polarity of each shape.</li> <li>Recognises the polarity of the Br–Cl bond due to the difference in electronegativity.</li> </ul>	<ul> <li>In a trigonal planar shape, the Br– Cl dipoles would cancel out as they are symmetrically arranged.</li> <li>In a T-shaped molecule, the Br– Cl dipoles would not cancel out, as they are asymmetrically arranged (due to the presence of lone pairs in equatorial positions).</li> </ul>	• Compares the polarity of both shapes of BrCl <sub>3</sub> , with reference to bond polarity and the arrangement of bond dipoles.

NØ	N1	N2	A3	A4	M5	M6	E7	E8
No response; no relevant evidence.	la	2a	3a	4a	2m	3m	2e, with minor error / omission.	2e

## NCEA Level 3 Chemistry (91390) 2024 — page 3 of 4

Q	Evidence	Achievement	Achievement with Merit	Achievement with Excellence
THREE (a)(i)	NH <sub>3</sub> : Hydrogen bonding, (permanent dipole), temporary dipole SO <sub>2</sub> : Permanent dipole, temporary dipole C <sub>5</sub> H <sub>12</sub> : Temporary dipole	• TWO rows correct.		
(ii)	Due to the large electronegativity difference between H and N, there is strong hydrogen bonding between $NH_3$ molecules. This is stronger than the permanent dipole attractions between the polar SO <sub>2</sub> molecules. However, SO <sub>2</sub> is a larger molecule than $NH_3$ , with a larger electron cloud. This means SO <sub>2</sub> has stronger / more temporary dipole attractions between the SO <sub>2</sub> molecules, and therefore a higher boiling point.	• Recognises hydrogen bonding is stronger than permanent /temporary dipole attractions.	• Explains why SO <sub>2</sub> has the higher boiling point by linking the larger electron cloud to stronger / more temporary dipole attractions.	• Justifies difference in boiling points for ALL three molecules in terms of strength of attractive forces.
(iii)	Despite the molecules having quite similar-sized e <sup>-</sup> clouds (molar mass), pentane has stronger intermolecular attractions. Pentane is a longer molecule when compared to the more spherical shape of polar SO <sub>2</sub> , so it has a greater surface area for interaction, meaning stronger temporary dipole attractions. Even though there are strong permanent dipole attractions between the polar SO <sub>2</sub> molecules, more energy is needed to break the attractions between the pentane molecules, so pentane has a higher boiling point.	<ul> <li>Recognises that molar mass / e- cloud size affects temporary dipole attractions /boiling point.</li> <li>Recognises that chain length /surface area affects temporary dipole attractions/ boiling point.</li> </ul>	• Explains that the linear shape / greater surface area of pentane means stronger attractive forces despite the permanent dipoles /polar nature of SO <sub>2</sub> .	
(b)(i)	$n(C_{2}H_{5}OH) = \frac{m}{M} - \frac{0.500}{46.0} = 0.0109 \text{ mol}$ -q = \Delta_{t}H \times n = -770 \times 0.0109 = -8.37 kJ Therefore q = 8.37 kJ = 8370 J $m = \frac{q}{c\Delta t} = \frac{8370}{4.18 \times 9.1} = 220 \text{ g}$	• One step of calculation correct.	• Correct process for calculation with one error.	<ul> <li>Correct final mass, including unit and significant figures (accept 2 – 4 significant figures for final answer).</li> </ul>
(ii)	Identifies temperature change of water. The temperature change (rise) as beaker is not insulated / no lid / heat escaping round the sides, therefore less heat is transferred to the water than should be. Heat loss to the gauze (and beaker), therefore less heat is transferred to the water than should be.	• Identifies temperature change of water and states ONE reason.	• Identifies temperature change of water and explains ONE reason.	

NØ	N1	N2	A3	A4	M5	M6	E7	E8
No response; no relevant evidence.	la	2a	3a	4a	2m	3m	2e, with minor error / omission.	2e

## Cut Scores

Not Achieved	Achievement	Achievement with Merit	Achievement with Excellence
0 - 06	07 – 13	14 – 18	19 – 24