

No part of the candidate's evidence in this exemplar material may be presented in an external assessment for the purpose of gaining an NZQA qualification or award.

SUPERVISOR'S USE ONLY

3

91390



Draw a cross through the box (☒) if you have NOT written in this booklet

+



Mana Tohu Mātauranga o Aotearoa  
New Zealand Qualifications Authority

## Level 3 Chemistry 2025

### 91390 Demonstrate understanding of thermochemical principles and the properties of particles and substances

Credits: Five

Achievement	Achievement with Merit	Achievement with Excellence
Demonstrate understanding of thermochemical principles and the properties of particles and substances.	Demonstrate in-depth understanding of thermochemical principles and the properties of particles and substances.	Demonstrate comprehensive understanding of thermochemical principles and the properties of particles and substances.

Check that the National Student Number (NSN) on your admission slip is the same as the number at the top of this page.

**You should attempt ALL the questions in this booklet.**

A periodic table and other reference material are provided in the Resource Booklet L3-CHEMR.

If you need more room for any answer, use the extra space provided at the back of this booklet.

Check that this booklet has pages 2–12 in the correct order and that none of these pages is blank.

Do not write in the margins (✂). This area will be cut off when the booklet is marked.

**YOU MUST HAND THIS BOOKLET TO THE SUPERVISOR AT THE END OF THE EXAMINATION.**

Merit

**TOTAL 16**

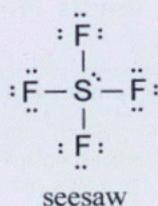
## QUESTION ONE

Sulfur forms a range of different fluoride-containing ions and molecules.

(a) Complete the table below:

	$SF_3^-$	$SBrF_5$
Lewis structure		
Name of shape	T-shaped	Octahedral

(b) The Lewis structure and shape for sulfur tetrafluoride,  $SF_4$ , are shown below.

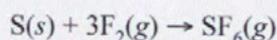


(i) Identify and explain the polarity of  $SF_4$ .

$SF_4$  has a seesaw shape.  $SF_4$  has 4 polar S-F bonds, which create bond dipoles due to the E.N difference between S and F. However, even though these bond dipoles are of the same strength, they are ~~unable~~ unable to cancel each other out due to the asymmetry of the seesaw shape of  $SF_4$ , making  $SF_4$  a polar molecule. The  $SF_4$  has bond angles of  $90^\circ$  and  $120^\circ/180^\circ$ .

- (ii) Justify why the F-S-F bond angles in SF<sub>4</sub> are different to the F-S-F bond angles in SBrF<sub>5</sub>. They have different bond angles of the F-S-F bond in each molecule due to the number of electron dense areas in each molecule. SBrF<sub>5</sub> has 6 areas of electron density meaning for maximum ~~repels~~ separation there are only bond angles of 90°, whereas SF<sub>4</sub> has both bond angles of 90° and 120° as there are only 5 areas of electron density, causing for more ~~less~~ space to separate the bonds.

- (c) Finely powdered sulfur, S(s), readily reacts with fluorine gas, F<sub>2</sub>(g), in an exothermic reaction to produce sulfur hexafluoride, SF<sub>6</sub>(g). The equation for the reaction is:



Justify, in terms of the entropy changes of the system and surroundings, why the reaction is spontaneous.

The reaction goes from 1 solid mol and 3 gaseous moles to only 1 gaseous mole, meaning the system's molecules are less able to move around and are less disordered which decreases the entropy of the system. However the reaction is exothermic, meaning it releases heat energy to the surroundings, this causing the surrounding gas molecules to move with more kinetic energy, meaning the surroundings are more disordered, increasing entropy. Since the reaction is spontaneous the increase in entropy from the surroundings must overcome the decrease in entropy of the system.

## QUESTION TWO

- (a) (i) Complete the table below.

Symbol	Electron configuration (use <i>s</i> , <i>p</i> , <i>d</i> notation)
Cl	$1s^2, 2s^2, 3p^6, 3s^2, 3p^5$
Cu	$[Ar], 4s^2, 3d^9$
$Fe^{3+}$	$[Ar], 3d^5$

- (ii) Iron, Fe, can form two different ions,
- $Fe^{2+}$
- and
- $Fe^{3+}$
- . Their ionic radii are given below:

 $Fe^{2+}$  92 pm $Fe^{3+}$  79 pmExplain why the radius of the  $Fe^{2+}$  ion is larger than the radius of the  $Fe^{3+}$  ion.

both  $Fe^{2+}$  and  $Fe^{3+}$  have the same nuclear charge but  $Fe^{2+}$  has an extra  $1e^-$  than  $Fe^{3+}$ . Since the nuclear charge is the same and the valence electrons are in the same energy level/shell it is due to electron - electron repulsion that causes  $Fe^{2+}$  the valence electrons to be pushed further from the nucleus, giving it a larger radius of 92 pm compared to  $Fe^{3+}$  79 pm

- (b) (i) Identify all the types of attractive forces between particles of the following substances in their liquid state.

Substance	$\Delta_{\text{vap}}H^\circ/\text{kJ mol}^{-1}$	Attractive forces
Hydrogen bromide, $HBr(l)$	17.3	temporary, permanent.
Hydrogen fluoride, $HF(l)$	25.2	<del>hydrogen</del> , permanent, temporary
Bromine, $Br_2(l)$	29.6	temporary

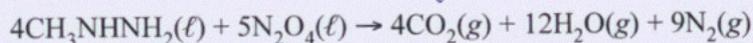
(ii) Explain why the  $\Delta_{\text{vap}}H^\circ$  of HF is higher than that of HBr.

HF has a higher  $\Delta_{\text{vap}}H^\circ$  of  $29.2 \text{ kJ mol}^{-1}$  compared to HBr with  $17.3 \text{ kJ mol}^{-1}$ . This is because HBr only has temporary and permanent dipoles intermolecular forces, whereas HF has hydrogen bonding which is a stronger intermolecular force than permanent and temporary. Both HF and HBr have permanent & temporary dipoles intermolecular forces but since F is the most electronegative the permanent dipole between H and F is stronger than H-Br, so will require more energy to overcome, so HF has a higher  $\Delta_{\text{vap}}H^\circ$  of  $29.2 \text{ kJ mol}^{-1}$  in comparison to HBr with only  $17.3 \text{ kJ mol}^{-1}$ .

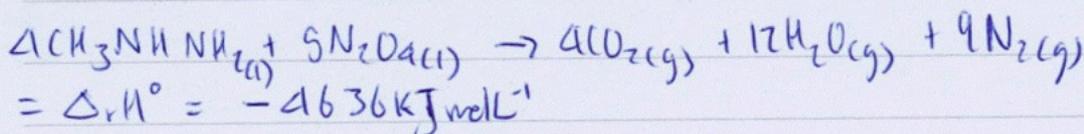
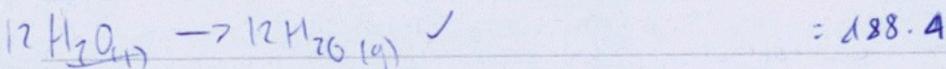
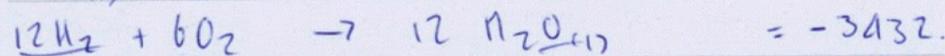
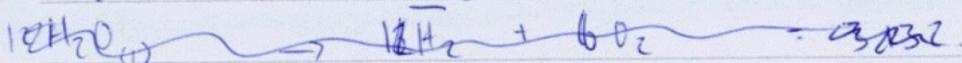
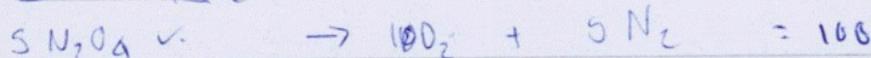
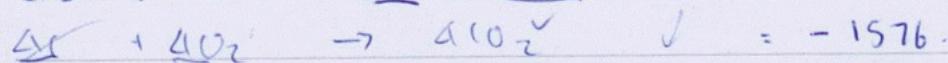
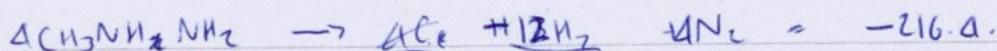
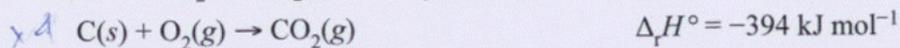
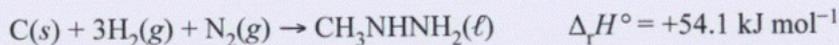
(iii) Justify why  $\text{Br}_2$  has the highest  $\Delta_{\text{vap}}H^\circ$  of the three substances.

$\text{Br}_2$  has the highest  $\Delta_{\text{vap}}H^\circ$  of  $29.6 \text{ kJ mol}^{-1}$  compared to HF and HBr even though it only has temporary dipoles, this is due to it having a much larger molar mass (159.8) meaning it has a larger electron cloud, meaning its temporary dipoles are much stronger than HF and HBr intermolecular forces.

- (c) Dinitrogen tetroxide,  $N_2O_4$ , was reacted with methylhydrazine,  $CH_3NHNH_2$ . The equation for the reaction is:



- (i) Calculate the standard enthalpy change of this reaction, using the following data.



2x0

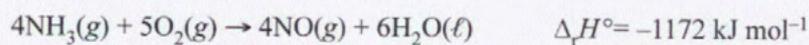
2x0

- (ii) Explain the effect on the standard enthalpy change calculated in part (c)(i) if the water was produced as a liquid.

if water were produced as a liquid as opposed to a gas, ~~the~~ the reaction would be more endothermic as less bonds would be broken, making  $\Delta_r H^\circ$  of the reaction less negative / more positive.

## QUESTION THREE

- (a) Nitric acid,  $\text{HNO}_3$ , is manufactured from ammonia,  $\text{NH}_3$ . The equation for one of the reactions in the process is:



- (i) Write the balanced equation to represent the standard enthalpy of formation of ammonia in the table below.

Substance	Equation	$\Delta_r H^\circ \text{ kJ mol}^{-1}$
$\text{NH}_3(\text{g})$	$\frac{1}{2}\text{N}_2(\text{g}) + \frac{3}{2}\text{H}_2(\text{g}) \rightarrow \text{NH}_3(\text{g})$	+45.7
$\text{NO}(\text{g})$	$\frac{1}{2}\text{N}_2(\text{g}) + \frac{1}{2}\text{O}_2(\text{g}) \rightarrow \text{NO}(\text{g})$	+90.3
$\text{H}_2\text{O}(\ell)$	$\text{H}_2(\text{g}) + \frac{1}{2}\text{O}_2(\text{g}) \rightarrow \text{H}_2\text{O}(\ell)$	-286

- (ii) Calculate the standard enthalpy of formation for ammonia,  $\Delta_f H^\circ(\text{NH}_3(\text{g}))$ , using the data given in part (i).

$$\begin{aligned} \text{products} - \text{reactants} &= \Delta_r H^\circ \\ (4(90.3) + 6(-286)) - (4(\text{NH}_3)) &= -1172 \\ (-1354.8) - (4\text{NH}_3) &= -1172 \\ -4\text{NH}_3 &= 182.8 \\ 4\text{NH}_3 &= -182.8 \\ \text{NH}_3 &= 45.7 \text{ kJ mol}^{-1} \end{aligned}$$

- (b) (i) Justify, using your knowledge of periodic trends, why magnesium, Mg, has a greater first ionisation energy than strontium, Sr.

Mg and Sr <sup>are</sup> ~~are~~ in the same group. Ionisation energy decreases when going down a group. Sr, while having a higher nuclear charge also has its valence electrons in shells/energy levels, further from its nucleus, meaning ~~therefore~~ the valence electrons experience shielding from the inner shells <sup>causing</sup> ~~causing~~ it to be less strongly held/ attracted by the nucleus and since the first ionisation energy is the energy required to remove an  $e^-$  from the valence shell in its gaseous state Sr will require less energy to do so. Mg has its valence electrons more ~~less~~ <sup>strongly</sup> attracted due to less shielding, meaning more energy is required, giving Mg a higher I.E than Sr.

- (ii) The atomic radii of five elements from Period 3 are shown in the table below.

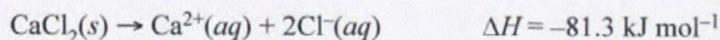
Period 3 element	Na	Mg	Al	Si	P
Atomic radius/nm	0.191	0.160	0.130	0.118	0.110

Explain the trend in atomic radius shown in the table.

The atomic radius decreases when moving across the period. This is due to the valence electrons being held in the same energy levels/shells but as you move down the period the nuclear charge increases. This increase in nuclear charge attracts the valence electrons more strongly, pulling them closer to the nucleus and therefore decreasing the atomic radius, which is why Na has the largest atomic radius of 0.191 nm in

period 3 and P has the smallest with  
0.110 nm.

- (c) When solid calcium chloride,  $\text{CaCl}_2(s)$ , dissolves in water, the temperature increases from  $20.4^\circ\text{C}$  to  $23.2^\circ\text{C}$ .



Calculate the mass of  $\text{CaCl}_2(s)$  that must dissolve to cause this temperature increase.

Assume the specific heat capacity of the calcium chloride solution is  $4.18 \text{ J g}^{-1} \text{ }^\circ\text{C}^{-1}$ .

Assume the mass of the calcium chloride solution is 68.0 g.

$$M(\text{CaCl}_2) = 111 \text{ g mol}^{-1}$$

$$\Delta T = 2.8^\circ\text{C}$$

$$4.18 \times 10^{-3} \text{ kJ g}^{-1} \text{ }^\circ\text{C}^{-1}$$

$$m(\text{CaCl}_2) = 68.0 \text{ g}$$

$$\Delta H = -81.3$$

$$q = 68 \times 4.18 \times 10^{-3} \times 2.8$$

$$q = 0.795872$$

$$n = \frac{-q}{\Delta H}$$

$$n = \frac{-0.795872}{-81.3}$$

$$n = 9.789 \times 10^{-3} \text{ mol}$$

$$m = n \times M$$

$$m = 9.789 \times 10^{-3} \times 111$$

$$m \approx m(\text{CaCl}_2(s)) = 1.09 \text{ g}$$





Extra space if required.  
Write the question number(s) if applicable.

QUESTION  
NUMBER

91390

## Merit

**Subject:** L3 Chemistry

**Standard:** 91390

**Total score:** 16

Q	Grade score	Marker commentary
One	M6	Correctly drew both Lewis diagrams and named the resulting shapes, explained the polarity of the molecule by recognising the arrangement of bond dipoles, and explained the spontaneity of the process by referring to the change in the surroundings' entropy.
Two	M5	Explained the difference in the ionic radius of the ions ( $\text{Fe}^{2+}$ and $\text{Fe}^{3+}$ ), correctly applied Hess' Law to calculate the enthalpy change for a reaction, and explained the differences in $\Delta_{\text{vap}}H^\ominus$ by linking the strength of temporary dipole attractions to a larger electron cloud.
Three	M5	Explained the difference in first ionisation energy and atomic radii in terms of the number of energy levels, nuclear charge, and the attraction between the nucleus and valence electrons. Calculated the mass of a solid dissolved correctly.