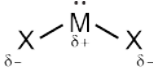
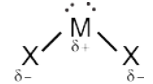
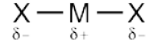


Assessment Schedule – 2013**Chemistry: Demonstrate understanding of bonding, structure, properties and energy changes (91164)****Assessment Criteria**

Achievement	Achievement with Merit	Achievement with Excellence
<i>Demonstrate understanding</i> involves describing, identifying, naming, drawing, calculating, or giving an account of bonding, structure and properties of different substances and the energy involved in physical and chemical changes. This requires the use of chemistry vocabulary, symbols and conventions.	<i>Demonstrate in-depth understanding</i> involves making and explaining links between the bonding, structure and properties of different substances and the energy involved in physical and chemical changes. This requires explanations that use chemistry vocabulary, symbols and conventions.	<i>Demonstrate comprehensive understanding</i> involves elaborating, justifying, relating, evaluating, comparing and contrasting, or analysing links between bonding, structure and properties of different substances and the energy involved in physical and chemical changes. This requires the consistent use of chemistry vocabulary, symbols and conventions.

Evidence Statement

Q	Evidence	Achievement	Merit	Excellence
ONE (a)	Lewis diagrams shown (Appendix One).	In (a) TWO Lewis structures correct.		
(b)	BF₃: trigonal planar: 120° bond angles. PF₃: trigonal pyramidal; ≈ / < 109.5° (107°). Shape is determined by the number of regions of electron density / electron clouds and whether they are bonding / non-bonding. BF₃ has three regions of electron density / electron clouds around the central B atom. The regions of electrons are arranged as far apart as possible from each other / to minimise repulsion, which results in a trigonal planar arrangement with a bond angle of 120°. All three regions of electrons are bonding, so the overall shape is trigonal planar. PF₃ has four regions of electron density / electron clouds around the central P atom. The regions of electrons make a tetrahedral arrangement with a bond angle of 109.5°. Only three regions of electrons are bonding and one is non-bonding, so the overall shape is trigonal pyramidal. <i>The non-bonding electrons have increased repulsion, therefore decreasing the bond angle to < 109.5°</i>	- In (b) TWO shapes correct. - In (b) TWO bond angles correct.	- In (b) the arrangement of electrons around the central atom is used to explain the shape of the molecule. - In (b) the arrangement of electrons around the central atom is used to explain the bond angle.	In (b) the arrangement of the electron density / electron clouds around the central atom is used to explain the shapes and angles of the molecules. Includes a comparison of the different shape and bond angles.

(c)(i)		The NH ₃ molecule is polar. The N–H bond is polar due to differences in electronegativity of N and H. The shape of the molecule is trigonal pyramidal, therefore the N–H polar bonds are not arranged symmetrically around the N atom. This means that the dipoles will not cancel. This results in a molecule which is polar.	• In (c) N–H bond is polar. • Predicts polarity of NH₃ correctly with one piece of supporting evidence.	• In (c)(i) the difference in electronegativities between N and H is used to explain the N–H bonds are polar. OR In (c)(i) links spread of charge to overall molecule polarity.				
		(c)(ii)				Polar: bent Non-polar: linear If MX ₂ is polar, this indicates that the polar M–X bonds are not spread symmetrically around the central M atom. There must be either three or four regions of negative charge with only two bonded atoms therefore the shape must be bent. Three regions of negative charge:  Four regions of negative charge:  If MX ₂ is non-polar this means that the polar M–X bonds are spread symmetrically around the central M atom. There must be only two regions of negative charge around the M atom, both bonded by X atoms in a linear shape. Two regions of negative charge: 	• Predicts one possible shape for MX₂. • Polarity depends upon the symmetry of the molecule.	• In (c)(ii) links the asymmetric spread of polar bonds to the shape.
NØ	N1	N2	A3	A4	M5	M6	E7	E8
No response or no relevant evidence.	1a	2a	4a	5a	3m	4m	3e with minor error / omission / additional information.	3e

Appendix One: Question One (a)

Molecule	Lewis structure
CH ₄	$\begin{array}{c} \text{H} \\ \\ \text{H} : \ddot{\text{C}} : \text{H} \\ \\ \text{H} \end{array} \quad \text{or} \quad \begin{array}{c} \text{H} \\ \\ \text{H} - \text{C} - \text{H} \\ \\ \text{H} \end{array}$
H ₂ O	$\begin{array}{c} \text{H} : \ddot{\text{O}} : \\ \\ \text{H} \end{array} \quad \text{or} \quad \begin{array}{c} \text{H} - \ddot{\text{O}} : \\ \\ \text{H} \end{array}$
N ₂	$:\text{N}::\text{N}: \quad \text{or} \quad :\text{N} \equiv \text{N}:$

Q	Evidence			Achievement	Merit	Excellence													
TWO (a)	<table><tr><th>Type of substance</th><th>Type of particle</th><th>Attractive forces between particles</th></tr><tr><td>Covalent network</td><td>Atom</td><td>Covalent (and weak intermolecular forces)</td></tr><tr><td>Molecular</td><td>Molecules</td><td>Weak intermolecular forces</td></tr><tr><td>Ionic</td><td>Ion</td><td>Ionic bonds / electrostatic attraction</td></tr><tr><td>Metal</td><td>Atom / cations and electrons</td><td>Metallic bonds / electrostatic attraction</td></tr></table>	Type of substance	Type of particle	Attractive forces between particles	Covalent network	Atom	Covalent (and weak intermolecular forces)	Molecular	Molecules	Weak intermolecular forces	Ionic	Ion	Ionic bonds / electrostatic attraction	Metal	Atom / cations and electrons	Metallic bonds / electrostatic attraction	• ONE row or ONE column correct. • Chlorine: low melting point OR is a gas at room temperature AND because it has weak intermolecular forces OR little energy is needed to turn it into a gas. • Copper chloride: High melting point OR is a solid at room temperature AND because it has strong ionic bonds OR a lot of energy would be needed to change it from a solid.	• Table completely correct. • Explains and links why chlorine is a gas and copper chloride is a solid at room temperature. Eg: Chlorine: has low melting point and is a gas at room temperature because it has weak intermolecular forces and little energy is needed to turn it into a gas Eg: CuCl₂: High melting point and is a solid at room temperature because it has strong ionic bonds and a lot of energy would be needed to change it from a solid.	Contrasts with reference to bonding and structure why chlorine is a gas at room temperature and copper chloride is a solid at room temperature.
Type of substance	Type of particle	Attractive forces between particles																	
Covalent network	Atom	Covalent (and weak intermolecular forces)																	
Molecular	Molecules	Weak intermolecular forces																	
Ionic	Ion	Ionic bonds / electrostatic attraction																	
Metal	Atom / cations and electrons	Metallic bonds / electrostatic attraction																	
(b)(i) Chlorine is a molecular substance composed of chlorine molecules held together by weak intermolecular forces. The weak intermolecular forces do not require much heat energy to break, so the boiling point is low (lower than room temperature); therefore chlorine is a gas at room temperature. Copper chloride is an ionic substance. It is composed of a lattice of positive copper ions and negative chloride ions held together by electrostatic attraction between these positive and negative ions. These are strong forces, therefore they require considerable energy to disrupt them and melt the copper chloride; hence copper chloride is a solid at room temperature. (b)(ii) For a substance to conduct electricity, it must have charged particles which are free to move. Graphite is a covalent network solid composed of layers of C atoms covalently bonded to three other C atoms. The remaining valence electron is delocalised (ie free to move) between layers; therefore these delocalised electrons are able to conduct electricity. Copper is a metallic substance composed of copper atoms packed together. Valence electrons are loosely held and are attracted to the nuclei of the neighbouring Cu atoms; ie the bonding is non-directional. These delocalised valence electrons are able to conduct an electrical current. For a substance to be made into wires, it needs to be stretched or drawn out without breaking.	• For something to conduct there must be free moving charged particles. • Graphite conducts because it has free moving electrons • Copper conducts because it has free moving electrons. • For something to be made into wires it needs to be able to be stretched without breaking / ductile	• Explains why both graphite and copper conduct electricity. • Explains why copper is ductile but graphite is not.	Contrasts with reference to bonding and structure why both graphite and copper can conduct electricity, however only copper is ductile.																

	In graphite, the attractive forces holding the layers together are very weak and are broken easily, so the layers easily slide over one another, but the attraction is not strong enough to hold the layers together and allow it to be drawn into wires or although the layers can slide due to weak forces, if graphite was to be made into a wire the very strong covalent bonds within the layers would have to be broken. Copper metal can easily be drawn into wires since, as it is stretched out, the non-directional metallic bonding holds the layers together, allowing it to be stretched without breaking.		- Graphite cannot be stretched since weak forces are easily broken or because the very strong covalent bonds have to be broken - Copper able to be stretched into wires because non directional bonding of valence electrons holds it together or because the metallic bonds can stretch without breaking.															
(c)			Identifies bonds broken and formed.		Process for calculating $\Delta_r H^\circ$ correct, however one minor error		Correctly calculates $\Delta_r H^\circ$, with units and negative sign.											
	<table><tr><td>Bonds broken:</td><td>Bonds formed:</td></tr><tr><td>C–H $\times$ 1</td><td>C–Cl $\times$ 1</td></tr><tr><td>Cl–Cl $\times$ 1</td><td>H–Cl $\times$ 1</td></tr><tr><td>414 + 242 = 656</td><td>324 + 431 = –755</td></tr></table>								Bonds broken:	Bonds formed:	C–H $\times$ 1	C–Cl $\times$ 1	Cl–Cl $\times$ 1	H–Cl $\times$ 1	414 + 242 = 656	324 + 431 = –755		
	Bonds broken:	Bonds formed:																
	C–H $\times$ 1	C–Cl $\times$ 1																
	Cl–Cl $\times$ 1	H–Cl $\times$ 1																
	414 + 242 = 656	324 + 431 = –755																
	656 kJ + (–755 kJ) = –99.0 kJ mol^{–1} OR																	
	<table><tr><td>Bonds broken:</td><td>Bonds formed:</td></tr><tr><td>C–H $\times$ 4</td><td>C–Cl $\times$ 1</td></tr><tr><td>Cl–Cl $\times$ 1</td><td>C–H $\times$ 3</td></tr><tr><td></td><td>H–Cl $\times$ 1</td></tr><tr><td>1656 + 242 = 1898</td><td>324 + 1242 + 431 = 1997</td></tr></table>								Bonds broken:	Bonds formed:	C–H $\times$ 4	C–Cl $\times$ 1	Cl–Cl $\times$ 1	C–H $\times$ 3		H–Cl $\times$ 1	1656 + 242 = 1898	324 + 1242 + 431 = 1997
	Bonds broken:	Bonds formed:																
	C–H $\times$ 4	C–Cl $\times$ 1																
Cl–Cl $\times$ 1	C–H $\times$ 3																	
	H–Cl $\times$ 1																	
1656 + 242 = 1898	324 + 1242 + 431 = 1997																	
1898 kJ + (–1997 kJ) = –99.0 kJ mol^{–1}																		
NØ	N1	N2	A3	A4	M5	M6	E7	E8										
No response or no relevant evidence.	1a	2a	5a	7a	3m	4m	2e	3e										

Q	Evidence	Achievement	Merit	Excellence
THREE (a)	Endothermic Gets colder The process is endothermic since the enthalpy change ($\Delta_r H^\circ$) is positive, which indicates that energy is absorbed by the system as the ammonium nitrate dissolves. Since heat energy is absorbed by the system from the surroundings (water & beaker), the water or beaker will get cooler as they lose heat energy.	- In (a) the reaction is endothermic because the value is positive OR because the ammonium nitrate is absorbing energy from the surroundings OR products have more energy than reactants. - In (a) beaker gets colder as heat energy is absorbed by ammonium nitrate.	Explains that since reaction is endothermic heat energy is absorbed by the system from the surroundings (water / beaker) so the beaker feels colder.	In (d) calculations correct with units and statement made about which iron oxide produces more heat energy. AND two bullet points from Merit.
(b)(i)	Exothermic The reaction is exothermic because the enthalpy change ($\Delta_r H^\circ$) is negative; indicating that heat energy is produced during the reaction.	- In (b)(i) exothermic since value is negative or because glucose reacting is releasing energy OR products have less energy than reactants. - In (b)(ii) calculation is correct.	- In (c)(i) explains the use of heat energy to break the weak intermolecular forces between butane molecules. - In (c)(ii) calculation is correct.	
(b)(ii)	9800 kJ / 2820 kJ mol ⁻¹ = 3.48 mol			
(c)(i)	Endothermic. Heat energy is needed to change the butane from a liquid to a gas; the energy is used to break the weak intermolecular forces between the butane molecules.			
(c)(ii)	n(C ₄ H ₁₀) = 100 g / 58.1 g mol ⁻¹ = 1.7212 mol -4960 kJ / 1.7212 mol = -2882 kJ mol ⁻¹	- In (c) the process is endothermic since energy is needed to boil butane. - In (c)(ii) one step correct in the calculation.		
(d)	n(Fe) = 2000 g / 55.9 g mol ⁻¹ = 35.78 mol Fe ₃ O ₄ : 3348 kJ / 9 = 372 kJ mol ⁻¹ 372 kJ mol ⁻¹ × 35.78 mol = 13 310.16 kJ = (-)1.33 × 10 ⁴ kJ	In (d) one step correct.	In (d) two steps correct	

	Fe_2O_3 : $851 \text{ kJ} / 2 = 425.5 \text{ kJ mol}^{-1}$ $425.5 \text{ kJ mol}^{-1} \times 35.78 \text{ mol} = 15\,224.4 \text{ kJ}$ $= (-)1.52 \times 10^4 \text{ kJ}$ Therefore Fe_2O_3 produces more heat energy when 2 kg iron is formed.							
NØ	N1	N2	A3	A4	M5	M6	E7	E8
No response or no relevant evidence.	1a	2a	4a	5a	2m	3m	e with minor error / incorrect unit / only 1m.	e

Judgement Statement

	Not Achieved	Achievement	Achievement with Merit	Achievement with Excellence
Score range	0 – 7	8 – 13	14 – 18	19 – 24