

CAMBRIDGE INTERNATIONAL EXAMINATIONS

Cambridge Pre-U Certificate

MARK SCHEME for the May/June 2015 series

9791 CHEMISTRY

9791/02

Paper 1 (Part A Written), maximum raw mark 100

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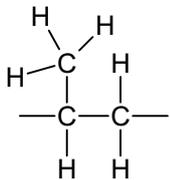
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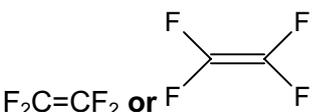
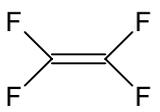
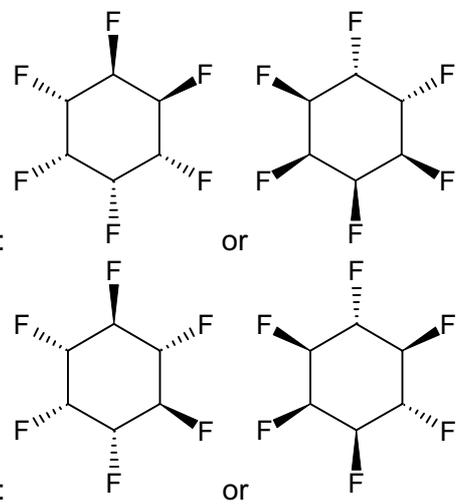
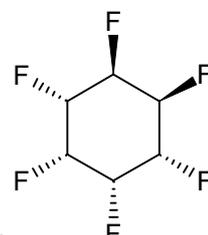
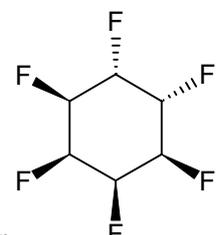
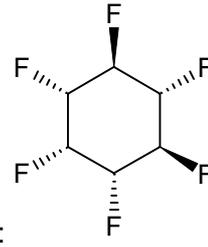
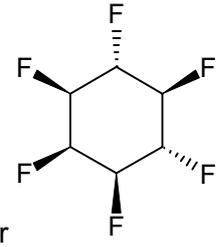
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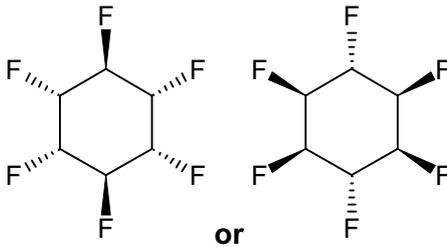
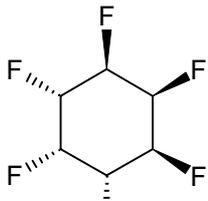
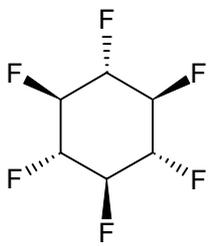
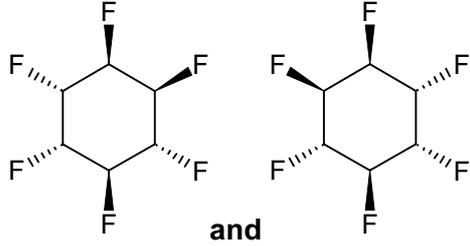
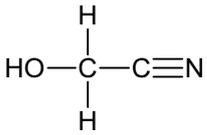
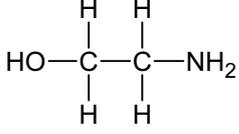
Page 2	Mark Scheme	Syllabus	Paper
	Cambridge Pre-U – May/June 2015	9791	02

1	(a) (i)	Forming <u>one mole</u> of substance/compound From its <u>elements</u> In their standard states or under standard conditions	1 1 1
	(ii)	$\Delta_c H^\ominus$ (298 K) = $\{-393.5 + 2 \times (-285.8) - (-74.8)\} \text{ kJ mol}^{-1} = -890.3 \text{ kJ mol}^{-1}$ Doubling the enthalpy for water Subtraction of reactant enthalpies from product enthalpies	1 1
	(b) (i)	$\Delta_r H^\ominus = \{(4 \times 435) + (2 \times 464) - 1077 - (3 \times 436)\} \text{ kJ mol}^{-1} = +283 \text{ kJ mol}^{-1}$ Correctly multiplying the energies by stoichiometric ratios. Subtracting product energies from reactant energies.	1 1
	(ii)	Either: large activation energy (1) as strong bonds must be broken (1) Or: forward reaction promoted (1) as it is endothermic (1)	1 1
	(c) (i)	$3\text{CO} + 6\text{H}_2 \rightarrow \text{C}_3\text{H}_6 + 3\text{H}_2\text{O}$	1
	(ii)	 <p>Dangling bonds required, but penalise if on the methyl group.</p>	1
	(d) (i)	S ₈ (The 8 must be subscripted.)	1
	(ii)	Oxidising agent	1
	(iii)	(2 × methane equation) minus ethene equation gives +101 kJ mol ⁻¹ Subtracting ethene equation/energy from methane equation/energy Doubling methane equation/energy	1 1
	(iv)	CS ₂ is linear H ₂ S is bent/V-shaped/non-linear	1 1
			[Total: 17]
2	(a) (i)	Mass = $0.50 \text{ g} \times (58.5/23) = 1.3 \text{ g} (1.27 \text{ g})$	1
	(ii)	Mass conc of Na ⁺ = $23 \text{ g mol}^{-1} \times 0.50 \text{ mol dm}^{-3} = 11.5 \text{ g dm}^{-3}$ Volume = $0.50 \text{ g} / 11.5 \text{ g dm}^{-3} = 43 \text{ cm}^3 (43.5 \text{ cm}^3)$	1 1
	(b) (i)	Volume = $\frac{1}{2} \times 24 \text{ dm}^3 \text{ mol}^{-1} \times 1.50 \text{ g} / 58.5 \text{ g mol}^{-1} = 0.31 \text{ dm}^3$ 2:1 stoichiometry for sodium chloride: chlorine taken into account. Correct use of molar volume and molar mass. Mark not awarded if answer not given to 2 sig figs.	1 1
	(ii)	pH 7	1
	(iii)	Blue/purple colour and $2\text{Na} + 2\text{H}_2\text{O} \rightarrow 2\text{NaOH} + \text{H}_2$	1
	(c)	Red colour <u>Polarising</u> Al ³⁺ (aq) weakens O–H bond OR stabilises coordinated OH ⁻ ion or	1

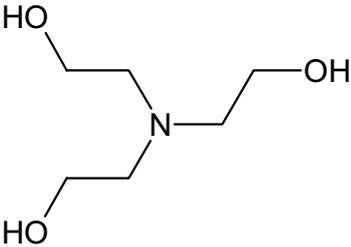
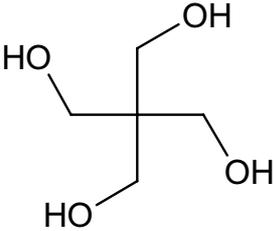
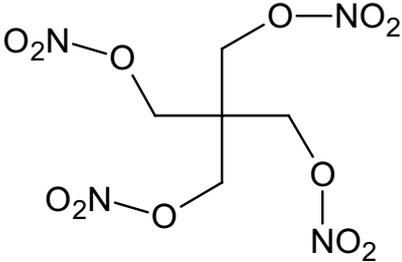
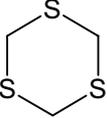
Page 3	Mark Scheme	Syllabus	Paper
	Cambridge Pre-U – May/June 2015	9791	02

	leads to dissociation of $H^+(aq)$ (from coordinated water molecules). $[Al(H_2O)_6]^{3+}(aq) \rightarrow [Al(H_2O)_5OH]^{2+}(aq) + H^+(aq)$	1 1
(d) (i)	$2NaOH + Cl_2 \rightarrow NaClO + NaCl + H_2O$ Chlorate(I)	1 1
(ii)	$Cl_2O + H_2O \rightarrow 2HOCl$	1
[Total: 13]		
3 (a) (i)	No. protons = 92. No. neutrons = 143. No. electrons = 92	1
(ii)	(Weighted) average mass Masses are relative to 1/12 of (the mass of a neutral atom of) carbon-12.	1 1
(iii)	Relative atomic mass = $(0.007 \times 235) + (0.993 \times 238) = 238.0$ Final answer given to 1 decimal place	1 1
(b) (i)	$1s^2 2s^2 2p^5$ The superscripting must be correct.	1
(ii)	Either: U–F dipoles cancel out (in an octahedral molecule) Or: δ^- charges of F atoms cause repulsion between molecules Or: fluorine has low polarisability.	1
(iii)	 $F_2C=CF_2$ or 	1
(c) (i)	Consistent: F_2 Inconsistent: HF	1 1
(ii)	Consistent: H–F or C–F Inconsistent: F_2	1 1
(d) (i)	Addition or reduction	1
(ii)	Pentanal	1
(e) (i)	 <p>1,2 isomer:  or </p> <p>1,3 isomer:  or </p>	

Page 4	Mark Scheme	Syllabus	Paper
	Cambridge Pre-U – May/June 2015	9791	02

	 <p>1,4 isomer: or</p> <p>Two marks for all three structures correct; one mark for two structures correct.</p>	2
(ii)	 <p>1,2,3 isomer:</p>  <p>1,3,5 isomer:</p>  <p>1,2,4 isomer and its enantiomer: and</p> <p>Enantiomers correctly identified.</p>	1 1 1 1
(iii)	9 isomers altogether.	1
[Total: 21]		
4 (a) (i)	Carbonyl, accept FGL2	1
(ii)	Methanol / CH ₃ OH	1
(b) (i)	 <p>C is</p>	1
(ii)	 <p>D is</p>	1

Page 5	Mark Scheme	Syllabus	Paper
	Cambridge Pre-U – May/June 2015	9791	02

(c) (i)	E is 	1
(ii)		1
(d)	B is 	1
(e) (i)	PETN is 	1
(ii)	Water	1
(f) (i)	$3\text{HCHO} + 3\text{H}_2\text{S} \rightarrow \text{C}_3\text{H}_6\text{S}_3 + 3\text{H}_2\text{O}$	1
(ii)	F is  Allow cyclopropane with an SH on each carbon. Allow a triangle of sulfurs, each bonded to =CH ₂ . Allow alternating single and double bonds around ring and an H bonded to each sulfur.	1
(g)	$\text{C}_2\text{H}_5\text{Br} + \text{Mg} \rightarrow \text{C}_2\text{H}_5\text{MgBr}$ (1) $\text{C}_2\text{H}_5\text{MgBr} + \text{HCHO} \rightarrow \text{C}_2\text{H}_5\text{CH}_2\text{OMgBr}$ (1) $\text{C}_2\text{H}_5\text{CH}_2\text{OMgBr} + \text{H}^+ \rightarrow \text{C}_2\text{H}_5\text{CH}_2\text{OH} + \text{Mg}^{2+} + \text{Br}^-$ (1) $\text{C}_2\text{H}_5\text{CH}_2\text{OH} + [\text{O}] \rightarrow \text{C}_2\text{H}_5\text{CHO} + \text{H}_2\text{O}$ (1) Suitable oxidising agent, eg acidified sodium or potassium dichromate, (to oxidise propan-1-ol to propanal). (1) <u>Dry ether</u> solvent for preparing or reacting Grignard reaction. (1) Oxidising agent added dropwise. (1) (Propanal recovered by) <u>distilling</u> (oxidation reaction). (1)	[Max: 7] 7
[Total: 18]		

Page 6	Mark Scheme	Syllabus	Paper
	Cambridge Pre-U – May/June 2015	9791	02

5 (a) (i)	C ₅ H ₅ N	1
(ii)		1
(b) (i)	3 (mol)	1
(ii)	Acid-base (as pyridine is a base and HI is an acid)/neutralisation	1
(iii)	Yellow indicates (a trace of) iodine past the end-point	1
(iv)	To ensure reaction with water goes to completion.	1
(c)	Atom oxidised is S and atom reduced is I. S goes from ox no. +4 to +6. I goes from ox no. 0 to -1.	1 1 1
(d) (i)	<p>M1: Mass of I₂ required = 0.2500 dm³ × 50.0 g dm⁻³ = 12.5 g (1) M2: Stoichiometric amount of Py = 3 × 0.2500 dm³ × 0.197 mol dm⁻³ = 0.148 mol(1) M3: Stoichiometric volume of Py = 0.148 mol × 80 cm³ mol⁻¹ = 12 cm³ (1) M4: Actual volume of Py chosen to be at least 12 cm³ (1) M5: Measure I₂ and Py with scales/a (mass) balance and a measuring cylinder, respectively. (1) M6: Dissolve I₂ and Py in the methanol(/SO₂) solution in a beaker. (1) M7: Transfer to a 250 cm³ volumetric flask with washings.(1) M8: Make up to the mark/line with the methanol/SO₂ solution. (1) M9: Invert / mix.(1) [Max 7 marks from M1 – M9]</p> <p>M10: Prepare the Fischer reagent in a fume cupboard.(1) M11: Avoid naked flames/Bunsen burners(1) [Max 8 marks total]</p>	8
(ii)	React iodine with (excess sodium) thiosulfate. Transfer methanol to a solvent residues bottle Or pour methanol down the sink with (plenty of) water	1 1
(iii)	<p>M1: Transfer sample solution to a conical flask. (1) M2: Use a 25 cm³ pipette to transfer the sample solution. (1) M3: Transfer the Fischer reagent to a burette. (1) M4: Place a white tile beneath the conical flask or swirling conical flask during addition from burette. (1) M5: Titrate <u>dropwise approaching endpoint</u> until end-point is reached or until (the first permanent) pale yellow colour is seen. (1) M6: Repeat until concordant/consistent titres are obtained. (1) [Max 5]</p>	5
(iv)	The methanol wasn't dry or methanol is hygroscopic or methanol absorbed water from the air.	1
(v)	<p>Conc of I₂ = 50.0 g dm⁻³ / 254 g mol⁻¹ = 0.197 mol dm⁻³ or Amount of I₂ = 0.05 g / 254 g mol⁻¹ = 1.97 × 10⁻⁴ mol. Consumption of water = 18000 mg mol⁻¹ × 0.000197 mol cm⁻³ = 3.54 mg cm⁻³</p>	1

Page 7	Mark Scheme	Syllabus	Paper
	Cambridge Pre-U – May/June 2015	9791	02

	<p>or Mass of water = $1.97 \times 10^{-4} \text{ mol} \times 18 \text{ g mol}^{-1} = 0.00354 \text{ g} = 3.54 \text{ mg}$</p> <p>Alternatively, the calculation may be done in reverse to show that 3.54 mg of water corresponds to 1 cm^3 of $50.0 \text{ g dm}^{-3} \text{ I}_2 \text{ (aq)}$:</p> <p>Amount of water = $3.54 \text{ mg} / 18 \text{ g mol}^{-1} = 1.97 \times 10^{-4} \text{ mol}$</p> <p>Mass of iodine = $1.97 \times 10^{-4} \text{ mol} \times 254 \text{ g mol}^{-1} = 0.05 \text{ g}$</p> <p>Conc of $\text{I}_2 = 0.05 \text{ g} / 0.001 \text{ dm}^3 = 50 \text{ g dm}^{-3}$</p>	1
(vi)	<p>M1: Vol of Fischer reagent that reacts with sample = $12.55 \text{ cm}^3 - 2.20 \text{ cm}^3 = 10.35 \text{ cm}^3$.</p> <p>M2: Mass of water in tartrate sample in titration = $3.54 \text{ mg cm}^{-3} \times 10.35 \text{ cm}^3 = 36.67 \text{ mg}$.</p> <p>M3: Mass of anhydrous sodium tartrate in total tartrate sample = $2.344 \text{ g} - (10 \times 0.03667 \text{ g}) = 1.977 \text{ g}$.</p> <p>M4: $x = \text{amount of water} / \text{amount of anhydrous sodium tartrate in total sample} = (0.3667 \text{ g} / 18 \text{ g mol}^{-1}) / (1.977 \text{ g} / 194 \text{ g mol}^{-1}) = 2.00 = 2$ (<u>integer required</u>)</p> <p>M3 & M4 could be based on the quantities in the titration, in which case the 2.344 g would need to be divided by 10 instead of the 0.03667 g being multiplied by 10.</p> <p>Alternative (longer) method:</p> <p>Mass of $\text{I}_2 = 50.0 \text{ g dm}^{-3} \times 0.01035 \text{ dm}^3 = 0.5175 \text{ g}$</p> <p>Amount of $\text{I}_2 = 0.5175 \text{ g} / 254 \text{ g mol}^{-1} = 0.0020374 \text{ mol}$</p> <p>M2: Amount of water = amount of $\text{I}_2 = 0.0020374 \text{ mol}$ (1)</p> <p>Mass of water in titration = $0.0020374 \text{ mol} \times 18 \text{ g mol}^{-1} = 0.03667 \text{ g}$</p> <p>M3: Mass of total tartrate = $2.344 \text{ g} - (10 \times 0.03667 \text{ g}) = 1.977 \text{ g}$ (1)</p> <p>M1 & M4 are the same as in the first method.</p>	1 1 1 1
[Total: 31]		