## Assessment Schedule – 2011

## Chemistry: Describe properties of aqueous systems (90700)

## Evidence Statement

Question	Evidence	Achievement	Merit	Excellence
ONE (a)	NH <sub>3</sub> weak base NaCl neutral NH <sub>4</sub> Cl weak acid HF weak acid	TWO of: • THREE from part (a) correct.	THREE from part (a) correct.	
(b)(i) (ii)	$NH_3 + H_2O \rightleftharpoons NH_4^+ + OH^-$ Equilibrium is to the left, so the greatest concentration of a species is $NH_3$ . For each $NH_3$ that reacts equal amounts of $NH_4^+$ and $OH^-$ are formed and are greater than the $OH^-$ and $H_3O^+$ formed by the dissociation of water. $NH_3 > OH^- \ge NH_4^+ > H_3O^+$ $HF + H_2O \rightleftharpoons F^- + H_3O^+$	<ul> <li>Correct equation.</li> <li>OR</li> <li>Correct rank for b(i) or b(ii)</li> </ul>	<ul> <li>Correct equation AND correct order of species for BOTH (b)(i) and (b)(ii).</li> </ul>	ONE explanation to Merit level. AND ONE full explanation. (all 4 species).
	Equilibrium is to the left, so the greatest concentration of a species is HF. For each HF that reacts equal amounts of $F^-$ and $H_3O^+$ are formed and are greater than the OH <sup>-</sup> and $H_3O^+$ formed by the dissociation of water. HF > $H_3O^+ \ge F^- > OH^-$		<ul> <li>Correct equation, order of species AND full explanation (all 4 species) for EITHER (b)(i) or (b)(ii).</li> </ul>	AND Correct answer with units, and appropriate number of sig. fig.
(c)	$K_{a} = 6.76 \times 10^{-4}$ HF + H <sub>2</sub> O $\rightleftharpoons$ H <sub>3</sub> O <sup>+</sup> + F <sup>-</sup> Assume [H <sub>3</sub> O <sup>+</sup> ] = [F <sup>-</sup> ] $K_{a} = \frac{[H_{3}O^{+}]^{2}}{[HF]} \implies [HF] = \frac{[H_{3}O^{+}]^{2}}{K_{a}}$ [H <sub>3</sub> O <sup>+</sup> ] = 4.57×10 <sup>-3</sup> mol L <sup>-1</sup> [HF] = 0.0309 mol L <sup>-1</sup>	EITHER • $K_a$ correct. OR • Correct [H <sub>3</sub> O <sup>+</sup> ].	AND • Correct answer with minor error (incorrect sig. fig. or units).	

TWO (a)(i) (ii)	$Zn(OH)_2(s) \rightleftharpoons Zn^{2+}(aq) + 2OH^{-}(aq)$ $K_s = [Zn^{2+}][OH^{-}]^2$	TWO of: • Part (a) correct.		
(b)	Let s be solubility $K_s = 4s^3$ $s = \sqrt[3]{\frac{K_s}{4}}$ $s = 1.96 \times 10^{-6} \text{ mol } \text{L}^{-1}$	• Method correct, but error in calculation. (Allow $s^2$ follow on from part (a) or $2s^3$ error but if so, must have calculated <i>s</i> value correctly according to the candidates	Solubility calculated correctly, (incorrect sig. fig.). AND ONE of: • Recognises that a complex ion	Solubility calculated correctly, 3 sig. fig. and <i>s</i> is defined. AND
(c)	Raising the pH will increase the concentration of $OH^-$ ions. This will initially cause additional precipitate to form. Once the pH has been increased sufficiently (enough $OH^-$ has been added) the formation of a complex ion with $Zn^{2^+}$ will occur, lowering $OH^-$ ion concentration in solution. Thus the precipitate will redissolve as a complex ion and less precipitate will be at the bottom of the test tube.	<ul> <li>follow on.)</li> <li>Recognises that [OH<sup>-</sup>] has increased.</li> <li>Recognises equilibrium will shift to the left.</li> </ul>	<ul> <li>will form and links this to either less solid remaining or equilibrium shifting to the right.</li> <li>Identifies equilibrium shifting to the left due to additional OH<sup>-</sup>.</li> <li>Explains equilibrium shifting to the left in terms of the I.P. now exceeding K<sub>s</sub>.</li> </ul>	re-dissolves, as equilibrium shifts in the forwards direction / to RHS. This shift to the right will occur so more $Zn^{2+}$ and $OH^-$ will dissolve into solution so that the solution becomes saturated again.

THREE (a)	$HG + H_2O \rightleftharpoons G^- + H_3O^+$	TWO of:		
	$OR \\ HOCH_2COOH + H_2O \rightleftharpoons HOCH_2COO^- + H_3O^+$	• Part (a) and (b) correct.		
(b)	$K_{\rm a} = \frac{[{\rm G}^-][{\rm H}_3{\rm O}^+]}{[{\rm HG}]}$ (must have equilibrium arrow)	•EITHER		
(c)	$[\mathrm{H}_{3}\mathrm{O}^{+}] = \sqrt{K_{\mathrm{a}}} \times [\mathrm{HG}]$	Correct value for $K_a$ OR	Correct answer with minor error.	Correct answer with appropriate number of sig. fig.
	$K_{\rm a} = 1.50 \times 10^{-4}$	Correct rearrangement of $K_a$ expression to make $[H_30^+]$ subject.		
	$[H_3O^*] = 9.99 \times 10^{-6} \text{ mol } L^{-6}$ pH = 2.00		AND	
(d)	$[H_3O^+] = 1.00 \times 10^{-4} \text{ mol } L^{-1}$		Correct [G <sup>-</sup> ].	AND
	$[G^{-}] = \frac{K_a \times [HG]}{[H_3O^{+}]} = 1.48 \text{ mol } L^{-1}$	• EITHER Correct [H <sub>3</sub> O <sup>+</sup> ].	OR	
	Thus in 200 mL = $0.2 \times 1.48 = 0.296$ mol	OR $K_{a}$ expression rearranged for [G <sup>-</sup> ] or	Correct method for [G <sup>-</sup> ] and n(G <sup>-</sup> ) calculation but incorrect answer.	Correct $n(G^{-})$ to 3 sig. fig.
	$pH = pK_{a} + \log_{10} \frac{[\text{weak base}]}{[\text{weak acid}]}$	other appropriate method for [G <sup>-</sup> ] stated and rearranged for [G <sup>-</sup> ].		
	$4.00 = 3.83 + \log_{10} \frac{[\text{base}]}{[\text{acid}]}$			
	$\log_{10}[\text{base}] = 0.17$			
	$[base] = 1.48 \text{ mol } L^{-1}$ Thus, in 200 mL = 00.2×1.48 = 0.296 mol			

FOUR (a)	A At point A, there is an equi-molar mixture of HEt and Et <sup>-</sup> . On addition of OH <sup>-</sup> ions, the acid part of the buffer neutralises the OH <sup>-</sup> ions, by donating a proton. The acid reacts with the base: HEt + OH <sup>-</sup> → Et <sup>-</sup> + H <sub>2</sub> O On addition of H <sub>3</sub> O <sup>+</sup> , the ethanoate will accept a proton from the hydronium ion: Et <sup>-</sup> + H <sub>3</sub> O <sup>+</sup> → HEt + H <sub>2</sub> O Candidate may discuss equilibrium shift. $pK_a = pH = 4.76$ (accept 4.5 – 4.9)	<ul> <li>ONE of:</li> <li>Recognises that at point A there is a buffer solution.</li> <li>States that equimolar amounts of acid / base conjugate are present at A.</li> <li>States that pH will not change when small amounts of acid or base are added.</li> <li>Correct pK<sub>2</sub> / K<sub>2</sub></li> </ul>	Describes how a buffer works (for when both acid AND base are added) by: EITHER • Giving equations for the specific buffer OR • Writing about how a buffer works in general terms OR • Links that due to equimolar	Shows recognition of equimolar HEt and Et <sup>-</sup> thus $pK_a = pH$ and discusses how the buffer solution works and links to equations.
(b)	B At the equivalence point all the HEt has been neutralised by NaOH. HEt + NaOH → EtNa + H <sub>2</sub> O The Et <sup>-</sup> reacts further to a small extent with water. Et <sup>-</sup> + H <sub>2</sub> O $\rightleftharpoons$ HEt + OH <sup>-</sup> Thus the pH of the equivalence point is above 7 due to presence of OH <sup>-</sup> .	<ul> <li>AND</li> <li>ONE of:</li> <li>Recognises that all the HEt has been used up at B.</li> <li>That the pH of equivalence point is greater than 7. (must have clearly indicated that point B is the equivalence point)</li> </ul>	<ul> <li>Entry that due to equinional HEt and Et<sup>-</sup> thus pK<sub>a</sub> = pH</li> <li>AND</li> <li>Recognises that none of the original HEt remains as ithas all reacted with NaOH OR</li> <li>That the pH of equivalence point is greater than 7 with a valid reason.</li> </ul>	AND Uses two equations to explain why the pH is above 7. (One equation may be implied in the candidate's written answer.)

## Judgement Statement

Achievement	Achievement with Merit	Achievement with Excellence
3 A	2 <b>M</b> + 1 <b>A</b>	2 <b>E</b> + 1 <b>A</b>