No part of the candidate evidence in this exemplar material may be presented in an external assessment for the purpose

of gaining credits towards an NCEA qualification.



91392



Level 3 Chemistry, 2017

91392 Demonstrate understanding of equilibrium principles in aqueous systems

2.00 p.m. Wednesday 15 November 2017 Credits: Five

Achievement	Achievement with Merit	Achievement with Excellence
Demonstrate understanding of equilibrium principles in aqueous	Demonstrate in-depth understanding of equilibrium principles in aqueous	Demonstrate comprehensive understanding of equilibrium principles
systems.	systems.	in aqueous systems.

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 is provided on the Resource Sheet L3–CHEMR.

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

Check that this booklet has pages 2–11 in the correct order and that none of these pages is blank.

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

Achievement

TOTAL

- (a) Hydrogen fluoride, HF, and hydrogen bromide, HBr, both form acidic solutions when added to water.
 - (i) Write an equation for the reaction of each acid with water.

Hydrogen fluoride, HF, with water:

Hydrogen bromide, HBr, with water:

(ii) Compare and contrast the electrical conductivity of 0.150 mol L⁻¹ solutions of hydrofluoric acid, HF, and hydrobromic acid, HBr.

In your answer, you should:

- include the requirements for a solution to conduct electricity
- identify the species present AND their relative concentrations.

No calculations are necessary.

- There must be ions present. A solution with a high concentration at long will be a good conductor whereas a solution with a law concentration will be appear conductor at electricity.
- "Hydrofluoric acid is a very strong acid, which means it fully dissociates into its ions. This results in high concentrations of 450+ and FT, which therefore makes HF a good conductor of electricity.
- · In contrast, hydrobromic and is a weak and, and only partially dissociates into its ions. This results in low concentrations of HyOt and Br- which means fewer particles are present to carry a charge, and means that It Br is not a good conductor.

(b) 40.0 mL of 0.150 mol L⁻¹ HBr solution was added to 25.0 mL of a saturated silver bromide, AgBr, solution.

(i) Write an equation for the equilibrium occurring in a saturated solution of AgBr.

AgBr = Agt + Br

(ii) Explain the changes that occur to the concentrations of the species in the saturated solution of AgBr on the addition of the HBr solution.

One to the addition of HBT, more Brampe ions will be present in the solution. An increase in Br results in the equilibrium favoring the reaction (AgBr) side. And as the solution of a contrated this will result in a precipitate of AgBr forming.

(iii) Calculate the concentration of the silver ions, Ag⁺, after the HBr solution has been added.

$$K_{\rm s}({\rm AgBr}) = 5.00 \times 10^{-13}$$

Assume the concentration of Br⁻ in the original saturated solution of AgBr is insignificant.

$$K_{S} = A_{g} + B_{c} - 1$$
 1:1 roho

 $K_{S} = S^{2}$
 $S = \sqrt{5 \times 10^{-13}}$
 $S = 7.071 \times 10^{-7}$

$$n = 1.768 \times 10^{-8}$$

$$C = \frac{n}{V} \quad C = \frac{1.768 \times 10^{-8}}{0.065} \quad C = 2.72 \times 10^{-7}$$

$$= > [Ag^{+}] = 2.72 \times 10^{-7}$$

A4

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QUESTION TWO

ASSESSOR'S

(a) Ammonia, NH₃, is a weak base.

$$pK_a(NH_4^+) = 9.24$$
 $K_a(NH_4^+) = 5.75 \times 10^{-10}$

(i) Calculate the pH of a $0.105 \text{ mol } L^{-1} \text{ NH}_3 \text{ solution}$.

NH₃ + H₂0 = NH₄⁺ + OH⁻

$$K_{0} = [NH_{4}^{+}][OH^{-}]$$

[NH₃]

Assuming that $[NH_{4}^{+}] = [OH^{-}]$
 $CH_{3} = [OH^{-}]^{2} = [OH^{-}] = [OH^{-}]^{-1}$
 $CH_{3} = [OH^{-}]^{2} = [OH^{-}] = [OH^{-}]^{-1}$
 $CH_{3} = [OH^{-}]^{2} = [OH^{-}]^{2} = [OH^{-}]^{-1}$
 $CH_{3} = [OH^{-}]^{2} = [OH^{-}]^{2} = [OH^{-}]^{2}$
 $CH_{3} = [OH^{-}]^{2} = [OH^{-}]^{2} = [OH^{-}]^{2}$

(ii) Dilute hydrochloric acid, HCl, is added to the NH_3 solution until the ratio of NH_3 to NH_4^+ in the solution is 5:1.

Determine the pH of this solution, and evaluate its ability to resist a change in pH when small volumes of strong acid or base are added.

Write the equation for the equilibrium occurring in a saturated solution of copper(II)* (b) (i) hydroxide, Cu(OH)₂.

ASSESSOR'S USE ONLY

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Write the expression for $K_{c}(Cu(OH)_{2})$. (ii)

Ks = [cu2+] COH-]2

(iii) Calculate the solubility of Cu(OH)₂ in water at 25°C.

 $K_{\rm s}({\rm Cu(OH)}_2) = 4.80 \times 10^{-20}$

s = 2.29 x10-7 molt-1

Explain why the solubility of Cu(OH)₂ increases when dilute hydrochloric acid is added.

reach with the OHT ions present to form

When It30+ ions are added they will

voter. As the product side at the equalibrium

experiences a decreve, to product side

will be fevered in order to oppose

the change. This result in a decrease

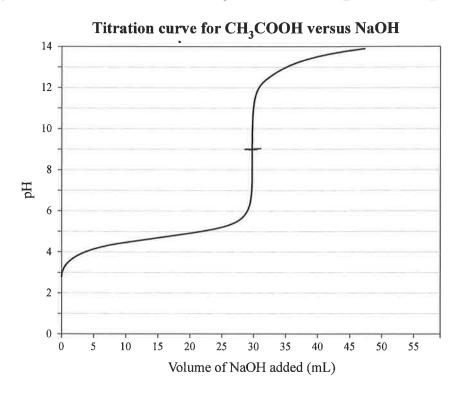
of cu (OH) 2 as more is dissolved.

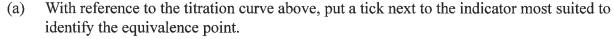
A titration was carried out by adding $0.112 \text{ mol } L^{-1}$ sodium hydroxide solution, NaOH(aq), to 20.0 mL of ethanoic acid solution, CH₃COOH(aq).

The equation for the reaction is:

$$\mathsf{CH_3COOH}(aq) + \mathsf{NaOH}(aq) \to \mathsf{CH_3COONa}(aq) + \mathsf{H_2O}(\ell)$$

$$K_a(\text{CH}_3\text{COOH}) = 1.74 \times 10^{-5}$$





Indicator	pK _a	Tick ONE box below,	
Methyl yellow	3.1		
Bromocresol purple	6.3		
Phenolphthalein	9.6	/	

		7	
(b)	(i)	The ethanoic acid solution, $CH_3COOH(aq)$, has a pH of 2.77 before any NaOH is added.	SESSOR'S SE ONLY
		Show by calculation that the concentration of the CH ₃ COOH solution is 0.166 mol L ⁻¹ .	
	10	CH3COOH FRO = CH3COO + H3O+	
		[Hyo+] = 10-2.77 = 1.70 ×10-3	
		M3 ED Wartson 2 x0005	
		$\kappa_{\alpha} = \frac{\Gamma \text{CH}_{3} \text{COOH}}{\Gamma \text{CH}_{3} \text{COOH}}$	
		7	
		[clt3cooH] = [clt3coo-][H30+] Ka	
		Ascence (H30+] = EcH300-] ECH300HJ = [1.70 ×10-3) = 0.166 mol L-1 1.74×10-5	
		[ctt3coott] = 1.74x10=5	°О
	(ii)	Calculate the pH of the solution in the flask after 10.0 mL of 0.112 mol L ⁻¹ NaOH has been added to 20.0 mL of ethanoic acid solution, CH ₃ COOH(aq).	
		Na OH + CH3 COOH -> CH3 COON9 + H20	
		n = ev n = 0.01 × 0.112 = 0.00112 mel	
		Onlobus oraza 0.0012 = 0.03 0.0373 m	nol L
		n f 0.166 × 6.02 n=3.32 × 10-3 3.32 × 10-3 0.117 × mole	<u>-</u> \
		$n = 0.01 \times 0.112$ $n = 1.12 \times 10^{-3}$ mol	
		1.12×10 = 0.0373 mel L-1	
		1:1 retto => [OH-] = 0.0373 moll-1	
		[H30+] for 20m1 = 1.70 x10-3	
		$n = 1.7 \times 10^{-3} \times 0.02 = 3.4 \times 10^{-5}$	0
¥		3.4×10 = 1.13 × 10 3 mold 1	
		$0.0373 \times 1.13 \times 10^{-3} = 4.23 \times 10^{-5}$	

Question Three continues on the following page.

- log 4.23 × 10-5 = 41.318

(c) The equivalence point pH for the titration of ethanoic acid with sodium hydroxide is 8.79.

(i) Identify the chemical species present at the equivalence point, other than water.

EOH- CH3COONS WOOH

(ii) In a second titration, a $0.166 \text{ mol } L^{-1}$ methanoic acid solution, HCOOH(aq), is titrated with the NaOH solution. The equivalence point pH for this titration is 8.28.

The equivalence point pH for the CH₃COOH titration is 8.79.

Compare and contrast the pH values at the equivalence point for both titrations.

 $K_{\rm a}({\rm HCOOH}) = 1.82 \times 10^{-4}$ $K_{\rm a}({\rm CH_3COOH}) = 1.74 \times 10^{-5}$

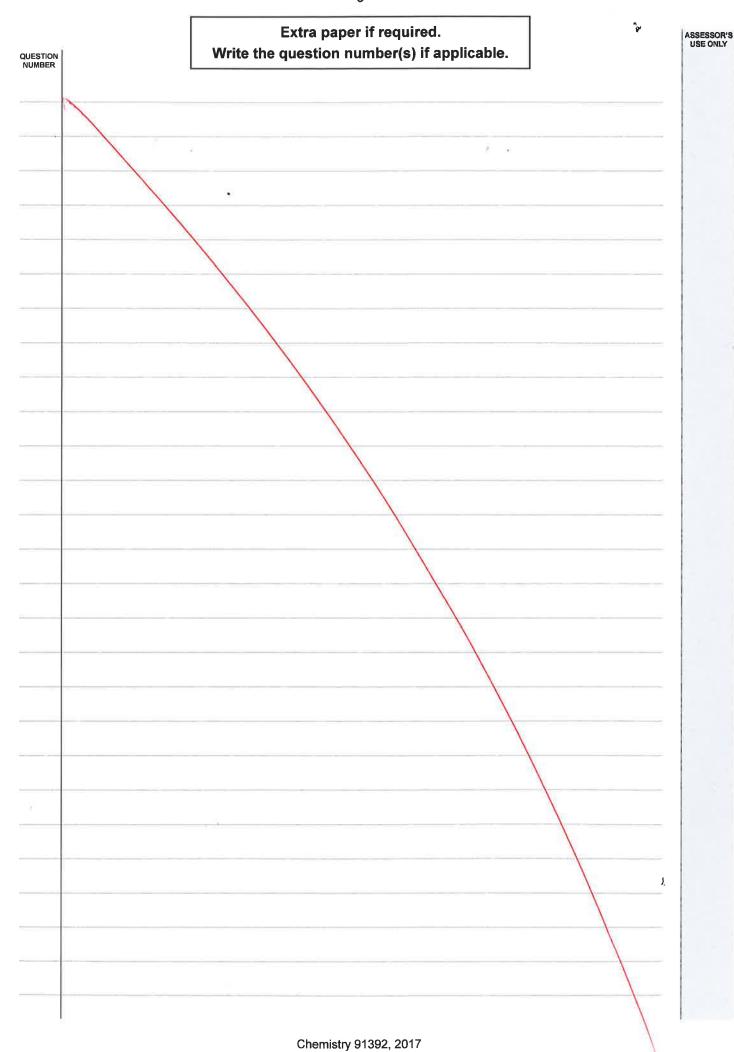
No calculations are necessary.

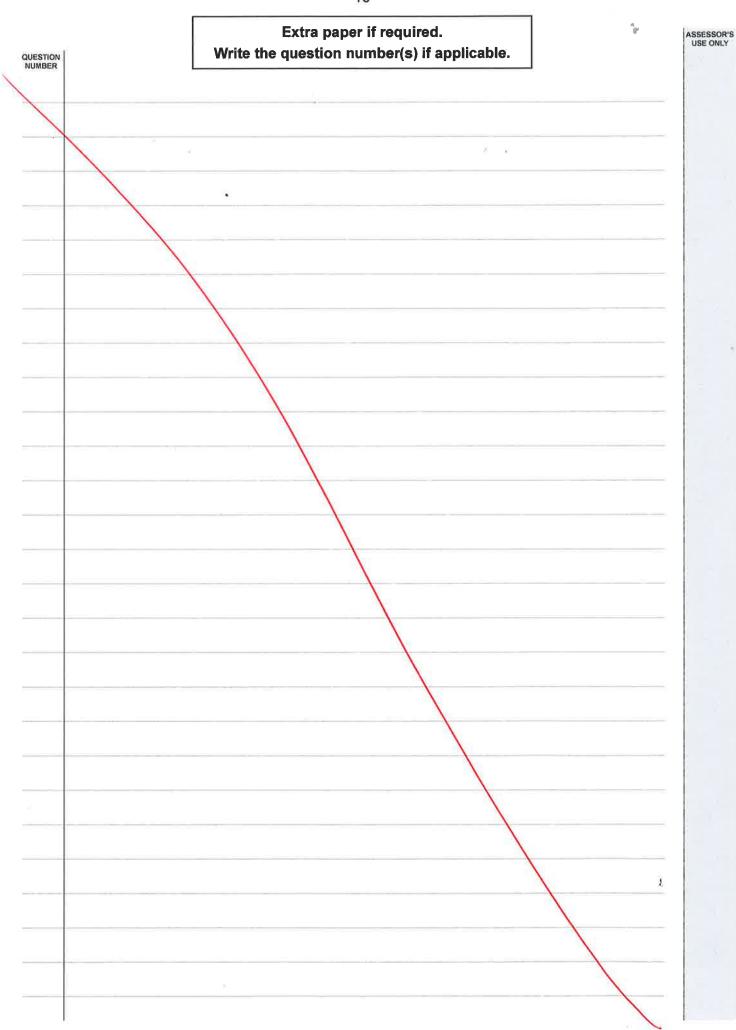
methonoic acid has a lower equividence point pht better obetter into its ions then extended acid. This can be seen by the Kay values as HCOOH has a higher Kay values as HCOOH has a higher Kay means more consentation of the southern. This means HCOOH has a laver pit then ctt 200H, and results in the equivalence point being beaut lower as well. As a stronger acid with the same base results in the equivalence point.

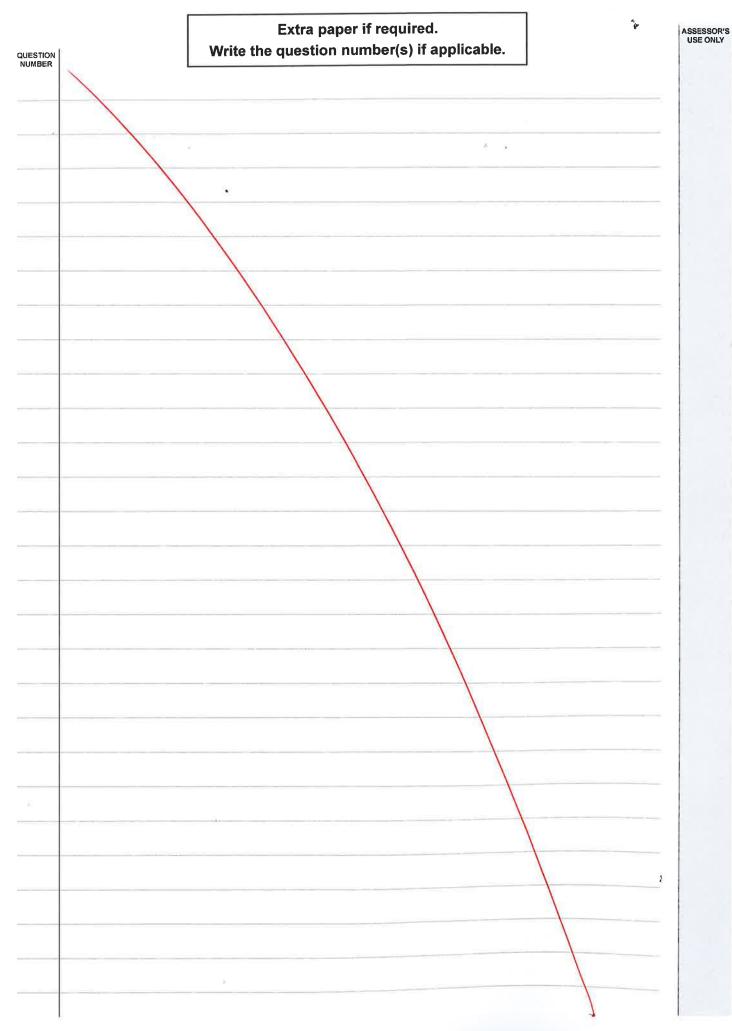


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Achievement exemplar for 91392 2017		Total score	11			
Q	Grade score	Annotation				
1	A4	The candidate was awarded A4 for the following	reasons:			
		In part (a)(i), both equations had incorrect arrows. In part (a)(ii), the candidate had the correct ideas on both conductivity and strength.				
		In part (b), the candidate wrote the correct equation; had a contradictory statement in the discussion on solubility; had a correct K_s expression, but incorrectly calculated the concentration of the silver ions present in the dilution.				
2	А3	The candidate was awarded A3 for the following reasons:				
		In part (a), the candidate calculated the pH incorrectly and did not answer part (a)(ii). Unfortunately, this response was required for the candidate to move up into Merit or higher.				
		In part (b), the correct equation and expression vused to correctly calculate the solubility of Cu(Ohunit.				
		In part (c), a full discussion on solubility relating t acid-base reaction was given.	o both equilibriu	ım and		
3	A4	The candidate was awarded A4 for the following	reasons:			
		In part (a), the correct indicator was chosen.				
		In part (b), the calculation of concentration was juthe three steps in the calculation of the pH was c		ne of		
		In part (c), the relative pH's related to acid streng merit, the candidate's response need to relate to e.g. the conjugate base strengths.	•			