## TRIAL TEST 2: ACIDS AND BASES



Time allowed: 70 minutes

Section 1 - Multiple Choice

20 marks

Total marks:

80

Section 2 - Short & Extended Answer

60 marks

### SECTION 1 - MULTIPLE CHOICE (20 MARKS)

Section 1 – Multiple Choice (20 marks)

Which one of the following equations shows water behaving as a base? 1.

> $CO_{(g)} + H_2O_{(g)} \rightarrow CO_{2}(g) + H_{2}(g)$ (a)

(b)  $2Na(s) + 2H_2O(l) \rightarrow 2Na^+(aq) + 2OH^-(aq) + H_2(g)$ 

 $O^{2-}(aq) + H_2 O(1) \rightarrow 2OH^{-}(aq)$ (c)

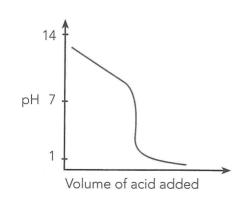
- (d)  $NH_4^+(aq) + H_2O(1) \rightarrow H_2O^+(aq) + NH_2(aq)$
- When phenolphthalein was added to a solution, a pink colour resulted. Which of the 2. following statements could be correct for that solution?
  - The solution was formed when 50.0 mL of 2.00 mol L-1 nitric acid was added to 50.0 mL of 2.00 mol L<sup>-1</sup> sodium hydroxide.
  - The solution was formed by the addition of 50.0 mL of 2.00 mol L-1 ethanoic (b) acid to 50.0 mL of 2.00 mol L<sup>-1</sup> potassium hydroxide.
  - The solution was formed at the end point of a titration between a strong acid and (c) a weak base.
  - The solution was formed when 3.75g of ammonium chloride was dissolved in (d)20.0 mL of water.
- 3. Which one of the following statements is true?
  - The hydrogen ion concentration of a neutral solution is dependent on the temperature of the solution.
  - The equivalence point of a titration occurs when equal numbers of moles of acid (b) and base have been mixed.
  - A buffer solution can be produced by mixing equal quantities of a strong acid and (c) strong base to ensure a plentiful supply of H<sup>+</sup> ions and OH<sup>-</sup> ions.
  - The salt produced when hydrochloric acid is reacted with sodium carbonate is (d) weakly acidic.
- A  $2.50 \times 10^{-2}$  mol L<sup>-1</sup> ethanoic acid solution will have a pH of: 4.
  - (a)
  - -2.50(b)
  - more than 1.60 but less than 7.00 (c)
  - dependent on the temperature of the water but always more than the pH of a (d) hydrochloric acid solution with the same hydrogen ion concentration
- 5. Which of the following lists contains one acidic, one basic, and one neutral salt?

(a)	ammonium chloride	sodium hydrogensulfate	barium chloride
(b)	calcium nitrate	sodium chloride	ammonium chloride
(c)	barium nitrate	sodium phosphate	ammonium nitrate
(d)	barium sulfide	sodium fluoride	ammonium chloride

6. A buffer solution with a pH of approximately 4.7 was made by mixing 500 mL of 0.500 mol L<sup>-1</sup> CH<sub>3</sub>COOH with 500 mL of 0.500 mol L<sup>-1</sup> NaCH<sub>3</sub>COO.

$$CH_{3}COOH_{(aq)} + H_{2}O_{(l)} \prod H_{3}O^{+}_{(aq)} + CH_{3}COO^{-}_{(aq)}$$

- (a) The addition of a 10 mL of 0.01 mol  $L^{-1}$  NaOH<sub>(aq)</sub> would raise the pH of the buffer solution by consuming the  $H_3O^+$  ions.
- (b) The additional of small quantities of hydrochloric acid will force the equilibrium to shift to the left by favouring the reaction that partially counteracts the increased pH.
- (c) The addition of any amount of sodium hydroxide solution will not greatly change the pH because this is a buffer solution.
- (d) The addition of 10.0 mL of 0.01 mol L-1 HNO<sub>3</sub> would not greatly alter the pH of the solution but the concentration of the CH<sub>3</sub>COO- ions would decrease.
- 7. Sulfuric acid is not suitable for use as a primary standard because:
  - In its manufacture, there is uncertainty about how much SO<sub>3</sub> is dissolved in each litre of water.
  - (b) It is a strong acid and so will react too rapidly.
  - (c) It does not have a sufficiently high molar mass.
  - (d) It produces sulfate salts which are sometimes insoluble.
- 8. A solution was produced by blending 500 g of celery, boiling it for 20 minutes in water and then straining the mixture. Analysis found it to have a pH of 4.2. The hydroxide ion concentration of this solution would be closest to which value stated below?
  - (a)  $3 \times 10^8 \text{ mol L}^{-1}$
  - (b)  $2 \times 10^{-10} \text{ mol L}^{-1}$
  - (c)  $4 \times 10^{-12} \text{ mol L}^{-1}$
  - (d)  $5 \times 10^{-13} \text{ mol L}^{-1}$
- 9. A student prepared a sodium hydroxide solution by dissolving 1.15 g of sodium hydroxide pellets in enough water to make exactly 50.0 mL of solution. This 50.0 mL of solution was then titrated against a 0.995 mol L<sup>-1</sup> HCl solution. The end point of the titration was noted when 22.5 mL of the acid had been added instead of the expected 28.9 mL. The most likely cause of this lower then expected result would be:
  - (a) the student chose methyl orange as an indicator instead of phenolphthalein.
  - (b) the sodium hydroxide pellets had absorbed water from the atmosphere prior to weighing.
  - (c) the student rinsed the burette with a small quantity of HCl solution before filling it with HCl solution.
  - (d) the 50.0 mL volumetric flash was rinsed with distilled water before the sodium hydroxide pellets were added.
- 10. The results of an acid base titration are shown in the graph to the right. Which of the following statements would be true about the titration?
  - (a) The titration was between 0.01 mol L<sup>-1</sup> NaOH and 0.01 mol L<sup>-1</sup> HCl.
  - (b) Phenolphthalein could be used as an indicator.
  - (c) Phenolphthalein or methyl orange could be used as an indicator.
  - (d) The titration involved a strong acid.



## SECTION 2 – SHORT AND EXTENDED ANSWER (60 MARKS)

Answer each question in the space provided.

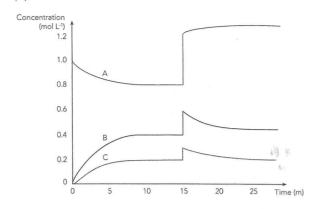
the fol	chemical test and subsequent observations that could be used to distinguish between llowing pairs of substances. You must clearly state the expected observation for ubstance tested. No equations need to be given.
(a)	1 mol $L^{-1}$ $H_2$ SO <sub>4</sub> and 1 mol $L^{-1}$ HNO <sub>3</sub>
TEST	
OBSE	RVATION
(b)	Two white powders: magnesium carbonate and magnesium hydroxide
OBSEI	RVATION
(c)	Two colourless solutions: potassium ethanoate and potassium chloride
OBSEI	RVATION
	[12 marks]
Write	balanced ionic equations for the following:
(a)	$Ba(OH)_{2}(aq) + H_{2}SO_{4}(aq)$
(b)	$KOH_{(aq)} + H_3PO_4^{(aq)}$
(c)	$CaCO_3(s) + HCl(aq)$
	[6 marks]
	quations to help explain why a 0.01 mol L <sup>-1</sup> HCl solution has a lower pH than a nol L <sup>-1</sup> CH <sub>3</sub> COOH solution
<u> </u>	

			[4 m
Write in wa		ess that occurs when the following acids a	re disso
(a)	carbonic acid:		
(b)	phosphoric acid:		
(c)	sulfuric acid:		
(a)	Explain what hydrolysis of	f a salt is.	[6 n
(b)	Write equations to show the is an acidic, basic or neutral	he hydrolysis of the following salts. State i al solution	f the ro
	(i) Na <sub>2</sub> CO <sub>3</sub>		
	(i) Na <sub>2</sub> CO <sub>3</sub> (ii) (CH <sub>3</sub> COO) <sub>2</sub> Ca		
(a)	(ii) (CH <sub>3</sub> COO) <sub>2</sub> Ca		
(a) 	(ii) (CH <sub>3</sub> COO) <sub>2</sub> Ca (iii) NH <sub>4</sub> NO <sub>3</sub>		[8 n
(a)  (b)	(ii) (CH <sub>3</sub> COO) <sub>2</sub> Ca (iii) NH <sub>4</sub> NO <sub>3</sub>	of a primary standard.	
	(iii) (CH <sub>3</sub> COO) <sub>2</sub> Ca (iii) NH <sub>4</sub> NO <sub>3</sub> State four characteristics o	of a primary standard.	[8 n
(b)	(iii) (CH <sub>3</sub> COO) <sub>2</sub> Ca (iii) NH <sub>4</sub> NO <sub>3</sub> State four characteristics of the NaOH is deliquescent. Exp	of a primary standard.  plain what this means.	[8 n
(b)	(iii) (CH <sub>3</sub> COO) <sub>2</sub> Ca (iii) NH <sub>4</sub> NO <sub>3</sub> State four characteristics of the NaOH is deliquescent. Exp	of a primary standard.  plain what this means.	[8 n

b)						
b)						
	Use Le Châtelie solution if 30 m				~ ~	-
						[,
c)	Explain (qualita 2 mol L <sup>-1</sup> hydro				pH of the so	olution if 20
using prepar to corr titrated	experiment, a stuthe primary started by dissolving rectly fill a 500.0 and against a solution in the table be	ndard, sodi 2.23 g of a mL volume on of hydro	ium carbon inhydrous so etric flask. 2	ate. The so odium carb .0.0 mL alio	odium carb onate in enquots of this	onate solu ough distill s solution v
using prepar to corr titrated	the primary star red by dissolving rectly fill a 500.0 d against a soluti	ndard, sodi 2.23 g of a mL volumo on of hydro	ium carbon inhydrous so etric flask. 2	ate. The so odium carb .0.0 mL alio	odium carb onate in enquots of this	of hydrochl onate solu ough distill s solution v
using prepar to correction show	the primary star red by dissolving rectly fill a 500.0 ed against a soluti wn in the table be	ndard, sodi 2.23 g of a mL volumo on of hydro	ium carbon inhydrous so etric flask. 2 ochloric acio	ate. The so odium carb 0.0 mL alio l. The volui	odium carb onate in enquots of this	of hydrochl onate solu ough distill s solution v used in the
using prepar to correction show	the primary started by dissolving rectly fill a 500.0 and against a solution with the table between tables are table to table the table between tables are tables as tables are tables as tables are tables as tab	ndard, sodi 2.23 g of a mL volume on of hydro elow:	ium carbon inhydrous so etric flask. 2 ochloric acid	ate. The so odium carb 0.0 mL alio l. The volui	odium carb onate in en- quots of this ne of HCl u	of hydrochl onate solu ough distill s solution v used in the

(d) Pressure was increased by reducing volume of the containing vessel.

(e)



(f) The equilibrium will shift so as to compensate for the greater imposed pressure. Moves right as there are less molecules. Concentration of the  $H_2$  affected most as there are two molecules of it. The other reactants affected equally (one molecule of each) but in opposite directions.

[16]

[20]

TRIAL TEST 2: Acids and Bases

#### Section 1

1. d 6. d 2. b 7. a 3. a 8. b 4. c 9. b 5. c 10. d

Section 2

11.

(a) Test: Add Ba(NO<sub>3</sub>)<sub>2</sub>(aq) to both solutions Observation: white precipitate forms in the H<sub>2</sub>SO<sub>4</sub>, no change in the HNO<sub>3</sub>

(b) Test:Add powders to HCl solutions Observation: MgCO<sub>3</sub> will fizz as bubbles of gas are produced, Mg(OH)<sub>2</sub> will simply dissolve

(c) Test: Add universal indicator to both Observation: KCl solution will turn green, KCH<sub>3</sub>COO will form orange/yellow.

[12]

12.

(a)  $Ba^{2+}(aq) + 2OH^{-}(aq) + 2H^{+}(aq) + SO_4^{2-}(aq) \rightarrow BaSO_4(s) + 2H_2O(l)$ 

 $(b)\,3\mathrm{OH^{\text{-}}}(aq) + H_{3}\bar{P}\mathrm{O}_{4}(aq) \rightarrow P\mathrm{O}_{4}^{3}\cdot(aq) + 3H_{2}\mathrm{O}(l)$ 

(c)  $CaCO_3(s) + 2H^+(aq) \rightarrow Ca^{2+}(aq) + CO_2(g) + H_2O(l)$ 

13.HCl is a strong acid and is completely ionized when in solution

 $HCl(aq) \rightarrow H^{+}(aq) + Cl^{-}(aq)$ For HCl, the  $[H^{+}] = [HCl]$   $CH_{3}COOH$  is a weak acid and so only a small percentage of molecules ionise  $CH_{3}COOH(aq) \Rightarrow H^{+}(aq) + CH_{3}COO^{-}(aq)$ 

For CH<sub>3</sub>COOH, the  $[H^+]$  <  $[CH_3COOH]$ Therefore,  $[H^+]$  in HCl is >  $[H^+]$  in CH<sub>3</sub>COOH and pH of 0.01 mol  $L^{-1}$  HCl is less

[4]

14.(a)  $H_2CO_3(aq) \rightleftharpoons H^+(aq) + HCO_3^-(aq)$ 

(b)  $H_3^2 PO_4(aq) \Rightarrow H^+(aq) + H_2^2 PO_4(aq)$ 

(c)  $H_2SO_4(aq) \rightleftharpoons H^+(aq) + HSO_4^-(aq)$ 

[6]

15

(a) Hydrolysis is the reaction between a salt and water to produce either  $H_3O^+$  ions or  $OH^-$  ions.

(b)

(i)  $CO_3^{2-}(aq) + H_2O(l) \Rightarrow HCO_3^{-}(aq) + OH^{-}(aq);$ basic

(ii)  $CH_3COO^-(aq) + H_2O(l) \rightleftharpoons CH_3COOH(aq) + OH(aq); basic$ 

(iii)  $NH_4^+(aq) + H_2O(l) \Rightarrow NH_3(aq) + H_3O^+(aq);$ 

[8]

16.

(a) be obtained pure; have a known formula; not react with surroundings; have a high molar mass

(b) deliquescent: absorbs water from the atmosphere and dissolves in the water

(c) end point: the point at which the titration is stopped because the desired colour change is observed

equivalence point: reactants have been mixed in stoichiometrically equivalent amounts

[8]

17.

(a)  $H_2PO_4(aq) + H_2O(l) \rightleftharpoons HPO_4(aq) + H_3O^+(aq)$ 

(b) The OH ions will reduce the concentration of the H<sub>3</sub>O+ ions. The forward reaction would be favoured to partially counteract this change and the pH would remain reasonably constant.

(c) The buffer capacity of the solution would be exceeded and the pH would drop

considerably.

[6]

$$n(Na_2CO_3)$$
 in 500  $mL = \frac{m}{M} = \frac{2.23}{105.99}$   
= 0.0210  $mol$   
 $c(Na_2CO_3) = \frac{n}{V} = 0.0421 \ mol \ L^{-1}$   
 $n(Na_2CO_3)$  used in titration =  $cV$   
= 0.0421 × 0.0200 = 8.42 × 10<sup>-4</sup>  $mol$   
 $n(HCl) = 2n(Na_2CO_3)$   
= 2 × 8.42 × 10<sup>-4</sup> = 1.68 × 10<sup>-3</sup>  
 $c(HCl) = \frac{n}{V} = \frac{1.68 \times 10^{-3}}{0.0413}$   
= 4.08 × 10<sup>-2</sup>  $mol \ L^{-1}$ 

# TRIAL TEST 3: Oxidation and Reduction

#### Section 1

1. d 6. d 2. c 7. a 3. d 8. a 4. a 9. b 5. c 10. a

Section 2

#### 11.

- (a) Equation:  $Br_{2}(aq) + 2I^{-}(aq) \rightarrow 2Br_{(aq)} + I_{2}(aq)$ Observation: straw yellow solution turns a red/brown colour
- (b) Equation:  $Zn(s) + Cu^{2+}(aq) \rightarrow Zn^{2+}(aq) + Cu(s)$ Observation: metal turns black and then black coloured crystals grow on it. Solution loses blue colour
- (c) Equation:  $2Na(s) + 2H_2O(l) \rightarrow 2Na^+(aq) + 2OH^-(aq) + H_2(g)$ Observation: silver coloured metal fizzes around on top of water, colourless, colourless gas produced
- (d) Equation:  $2MnO_4^{-(aq)} + 5H_2O_2^{-(aq)} + 6H^{+(aq)} \rightarrow 2Mn^{2+(aq)} + 5O_2^{-(g)} + 8H_2O_{(l)}$ Observation: purple solution goes colourless and bubbles of colourless odourless gas produced

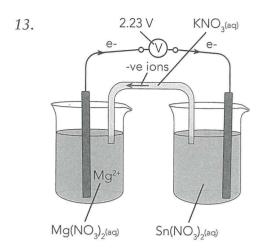
12.

(a)

- (i) Oxidation  $Zn(s) \rightarrow Zn^{2+}(aq) + 2e^{-}$ Reduction  $2H^{+}(aq) + 2e^{-} \rightarrow H^{2}(g)$
- (ii) Oxidation  $Mg(s) \rightarrow Mg^{2+}(aq) + 2e^{-}$   $Reduction 2H_{2}O(l) + 2e^{-} \rightarrow 2OH^{-}(aq)$  $+ H_{2}(g)$

(b)

- (i) Oxidation  $Zn(s) \rightarrow Zn^{2+}(aq) + 2e^{-}$ Reduction  $(Ag^{+}(aq) + e^{-} \rightarrow Ag(s)) \times 2$ Redox  $Zn(s) + 2Ag^{+}(aq) \rightarrow Zn^{2+}(aq) +$ Ag(s)
- (ii) Oxidation  $Mg(s) \rightarrow Mg^{2+}(s) + 2e$ Reduction  $Cl_2(g) \rightarrow 2Cl'(s)$ Redox  $Mg(s) + Cl_2(g) \rightarrow MgCl_2(s)$ [10]



ANODE:  $Mg \rightarrow Mg^{2+} + 2e^{-}$ CATHODE:  $Sn^{2+} + 2e^{-} \rightarrow Sn$ 

[20] 14.

- (a) Anode:  $Fe \rightarrow Fe^{2+} + 2e^{-}$ Cathode:  $O_2 + 2H_2O + 4e^{-} \rightarrow 4OH^{-}$ Rust formation:  $2Fe(OH)_3 \rightarrow Fe_2O_3.H_2O + 2H_2O$
- (b) (i) Coat the windmill with a paint to stop the oxygen and water coming in contact with the iron. This will prevent the cathodic reaction.
  - (ii) Connect another metal of higher oxidation potential to the windmill so that the iron acts as a cathode and the other metal an anode. For example if the other metal is zinc it will oxidise instead of the iron.

[12]

[12]

15.

(a)

[12]

(i)  $(Fe(s) \rightarrow Fe^{2+}(aq) + 2e^{-}) \times 2$  anodic reaction  $O_{2}(g) + 2H_{2}O(l) + 4e^{-} \rightarrow 4OH^{-}(aq) \text{ cathodic reaction}$   $2Fe(s) + O_{2}(g) + 2H_{2}O(l) \rightarrow 2Fe(OH)_{2}(s)$ 

 $\begin{array}{l} (ii)\,4Fe(\mathsf{OH})_2(s)+2H_2\mathsf{O}(l)+\mathcal{O}_2(g) \to 4Fe(\mathsf{OH})_3(s) \\ (iii)\,2Fe(\mathsf{OH})_3(s) \to Fe_2\mathsf{O}_3.\ H_2\mathsf{O}+2H_2\mathsf{O}(l) \end{array}$ 

(b) Any two of the following:

• Painting or plating the iron. This excludes air and/or water hence reaction prevented.

• Using a sacrificial anode such as galvanising