

Multiple Choice Questions

1.	Calcium hydroxide is a strong base. Calculate the [Ca ²⁺] and [OH-] for a solution
	prepared by dissolving 0.60 g of Ca(OH), in enough water to make 1.50 L of
	solution.

- 5.4×10^{-3} ; 1.1×10^{-2} (a)
- (b) 5.4×10^{-3} ; 5.4×10^{-3}
- 5.4×10^{-3} ; 9.1×10^{-13} (c)
- (d) 8.1×10^{-3} ; 8.1×10^{-3}
- 8.1×10^{-3} ; 1.6×10^{-2} (e)

2. What is the conjugate acid of CH₃NH₂?

- CH,NH-
- (b) CH_2NH^+ (c)
 - $CH_2NH_3^+$ (d)

CH₂NH₂

CH,NH2+ (e)

What is the conjugate acid of H₂SO₄? 3.

- (a) SO₄2-
- (b)
- $H_{2}SO_{4}^{-}$ (c) $H_{3}SO_{4}^{+}$
- (d) HSO,

(e) HSO₄+

What is the pH of human muscle fluid with a hydronium ion concentration of 4. $1.6 \times 10^{-7} \text{ mol L}^{-1}$?

- (a) 7.20
- (b) 7.16
- 7.80 (c)
- (d) 6.60

6.20 (e)

The hydrogen ion concentration of the oceans is about 7.1×10^{-9} mol L⁻¹. What is 5. the pH of ocean water?

- (a) 5.85
- (b) 7.15
- 8.15 (c)
- (d) 8.85

(e) 9.71

A brand of vinegar has a hydroxide ion concentration of 1.3×10^{-12} mol L⁻¹. What 6. is the pH of the vinegar?

- 3.11 (a)
- (b) 2.89
- (c) 3.88
- (d) 2.11

11 (e)

What is the pH of a 0.012 mol L⁻¹ solution of calcium hydroxide? 7.

- (a) 12.00
- 1.62 (b)
- (c) 11.10
- (d) 12.38

12.20 (e)

The pH of a 0.10 mol L⁻¹ solution containing 0.10 mol L⁻¹ NH₄Cl is 9.20. What is 8. the concentration of the H₃O+ ions in it?

- 2.0×10^{-9} (a)
- (b)
- 6.3×10^{-10} (c) 1.7×10^{-10}
- (d) 1.0×10^{-1}

 1.6×10^{-5} (e)

9.	OH- ions in it?									
	(a) (e)	3.2×10^{-3} 3.2×10^{-4}	(b)	5.0 × 10) ⁻¹⁰	(c)	3.2 × 10) ⁻¹¹	(d)	5.0×10^{-11}
10.	In th	e reaction, I	HCO ₃ -($(aq) + H_2($	O(1) ≠	OH-(ac	1) + H ₂ CO) ₃ (aq), t	he conji	ıgate base is
	(a)	HCO ₃	(b)	H_2O		(c)	OH-		(d)	H_2CO_3
	(e)	none of the	ese							
11.	The it fo	llows that H	r HF a: IF is a	nd HNC	o ₂ are 6. acid	8×10 ⁻⁴ than H	and 4.5 > INO ₂ and	< 10 ⁻⁴ re d F⁻ is a	espective a	ely. Therefore base than
	(a)	weaker, st	rongei		(b)	weal	ker, weal	ker		
	(c)	stronger, v			(d)	stror	nger, stro	nger		
12.		ich of the fol Iagnesium 1,2,3		romium	3. Le	ad 4.	Sodium	5. Ti	n	hydrogen?
13.		at is the corere K _w is 2.5			OH- i	ons in	a neutra	al solu	tion of	water at 37°0
	(a)	1.5×10^{-8}		(b)	2.6 ×	10-8	(c)	2.6 ×	< 10 ⁻⁷	
	(d)	1.6 x 10 ⁻⁷		(e)		1				
14.	Wh	at is the con	centra	tion of a	Ba(OI	H), solı	ution tha	t has a	pH 9.30	0?
	(a)				(b)	$2.00 \times 10^{-5} \text{ mol L}^{-1}$				
	(c)	2.50×10^{-1}	o mol	L-1	(d)	5.01	× 10 ⁻¹⁰ r	nol L-1		
15.		w much wat e the pH to		st be ado	ded to	200 ml	L of a 0.0	10 M s	olution	of HCl to
	(a)	232 mL	(b)	332 m	L	(c)	432 mL	i		
	(d)	532 mL	(e)	632 m	L					

Longer Questions

1. What is the relationship between the strength of an acid and its numerical value of K_a ?

2. How will the following ions react with water? Give the equation, where applicable:

I-, NO₂-, K+, HSO₄-, CO₃²⁻?

3. Predict whether aqueous solutions of the following salts would be acidic, alkaline, or close to neutral: KCl, $\rm NH_4NO_2$, $\rm Na(HCOO)$.

4. Predict whether aqueous solutions of the following salts would be acidic, alkaline or close to neutral: NH_4CN , KI, $LiCH_3COO$.

5. Identify the acids and bases according to the Brønsted-Lowry theory of acids and bases in the following

$$\text{(a)} \quad HNO_{3}(\textbf{l}) + H_{2}O(\textbf{l}) \;\; \rightleftharpoons \;\; H_{3}O^{+}(\textbf{aq}) + NO_{3}^{-}(\textbf{aq})$$

(b)
$$HCl(g) + NH_3(g) \Rightarrow NH_4^+(aq) + Cl^-(aq)$$

$$\text{(c)} \hspace{0.5cm} H_2SO_4(l) + H_2O(l) \hspace{0.2cm} \rightleftharpoons \hspace{0.2cm} H_3O^+(aq) + HSO_4^-(aq)$$

 $\text{(d)} \quad H_2O({\rm l}) + H_2O({\rm l}) \quad \ \, \rightleftharpoons \ \, H_3O^+({\rm aq}) + OH^-({\rm aq})$

(a)	What is meant by hydrolysis?
List	three ions that hydrolyse to give
(b)	an acidic solution
(c)	a basic solution.
	*
If 27	
If 32 solu	$2.0 \mathrm{mL}$ of a $0.1 \mathrm{M} \mathrm{H_2SO_4}$ is required to precipitate all the barium ions tion of $\mathrm{BaCl_2}$, what mass of $\mathrm{BaCl_2}$ was present in the solution?
If 32 solu	2.0 mL of a $0.1 \text{M H}_2 \text{SO}_4$ is required to precipitate all the barium ions
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Calc (a) (b) (c)	2.0 mL of a 0.1M H ₂ SO ₄ is required to precipitate all the barium ions tion of BaCl ₂ , what mass of BaCl ₂ was present in the solution? Culate the pH of the following solutions 0.10 M HBr 0.00001 M HNO ₃ 0.010 M KOH
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	0.050 mol of $Ba(OH)_2$
(c)	0.100 mol of HCl
Dete solu	ermine the pH of the solution formed by mixing equal volumes of the two tions in each case.
(a)	0.10 M HCl and 0.10 M NaOH
(b)	0.20 M HCl and 0.10 M NaOH
(c)	0.40 M HCl and 0.20 M NaOH
(d)	$0.10~\mathrm{M}~\mathrm{HCl}$ and $0.10~\mathrm{M}~\mathrm{Ba(OH)}_2$
A 1.5	. 50 litre water solution contains 1.7 micrograms of pure HCl (1 microgram = grams).
(i)	Calculate the concentration of HCl in ppm.
(ii)	Calculate the concentration of hydrogen ions in the solution.
(iii)	The calculated pH of this solution appears to be 7.50 which indicates a basic solution of an acid - but this cannot be correct. What is wrong with the assumption here?
If 4.9 what	$90~{\rm g}$ of sulfuric acid is added to water so that the final volume is 100 mL, twill be the pH of the solution, if 1 mole of ${\rm H_2SO_4}$ gives 1.3 moles of ${\rm H_3O^+}$?
	0.0 mL of 0.200 M sodium hydroxide solution is added to 30.0 mL of

0.200 volu	mL of 0.0500 mol L^{-1} barium hydroxide solution is mixed with 400 ml $$ M nitric acid. The mixture is then diluted with water so that the f me is 6.00 L.
Wha	t is the pH of the final solution?
	$0.0~\mathrm{mL}$ sample of $0.005~\mathrm{mol}~\mathrm{L}^{\text{-1}}$ calcium hydroxide is diluted with water $0.0~\mathrm{L}$
(a)	What is the pH of the original solution?
(b)	How does the pH change due to the dilution?
(c)	What mass of calcium hydroxide is present in the undiluted and the diluted solution?
	\
(d)	What volume of CO_2 at 25°C and 110 kPa pressure must be bubbled through the solution above in order to precipitate all the Ba ²⁺ ions?
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	lain using equations, why aqueous solutions of sodium carbonate, sodide and sodium ethanoate all have a pH value greater than 7.
A 50	olution has a pH of 6. If you dilute it to 100 times its original volume its

20.

19.	Ammonium	carbonate	dissolves	in	water	as	follows
17.	Allillolliull	carbonate	dissolves	111	water	as	10110

$$(NH_4)_2CO_3(s) \rightarrow 2NH_4^+(aq) + CO_3^{2-}(aq)$$

(1)

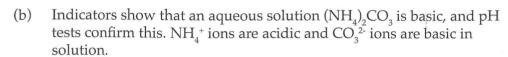
Following dissolution, two further reactions occur as follows:

$$NH_4^+(aq) + H_2O(l) \implies NH_3(aq) + H_3O^+(aq)$$

(2)

$$CO_3^{\ 2\text{-}}(\mathsf{aq}) + H_2O(\mathsf{l}) \ \rightleftharpoons \ HCO_3^{\ \text{-}}(\mathsf{aq}) + OH^\text{-}(\mathsf{aq})$$

(3)



Explain how a salt can produce a solution that is basic.

	*	
Suggest a reason why H ₂ I	${}^{2}O_{4}^{-}$ is a weaker acid t	han H ₃ PO ₄ .

- but for H_2SO_4 result is about $0.1 \times 1.3 = 1.3$, so less H⁺ and higher pH.
- Answer is D.
- Acidic compounds are CO₂, SO₃, OCl₂ and NO, (non-metal oxides).
- 8. Ba(OH), is a strong base with 3 ions per formula unit.
- OH-, HCO₃-, NH₃ (all can gain or lose a proton).
- 10. (i) CO, dissolves in water to produce H₂CO₃ which reacts with water: $H_2CO_3 + H_2O \rightarrow HCO_3^- + H_3O^+$ (acidic pH < 7). When boiled, this reaction reverses the produce neutral water again and pH 7. (ii) The K value for water goes up because the reaction is endothermic so more H^+ ions. (iii) Although there are more H⁺ ions if the pH is smaller, there is still an equal number of OH- ions so it is still neutral. N.B. The pH of water is 7 only at 25°C.
- 11. (ii) and (v) are not Brønsted-Lowry reactions.
- 12. (i) The ammonia solution must have a much higher concentration than the HCl as it a weak base and a 0.05 M solution would not supply enough OH ions to neutralise it. (ii) NH₃ ions would be most common as it "wants to" stay as a molecule.
- 13. A Brønsted-Lowry acid is a species that donates a H^+ ion. To react with water, a proton must be exchanged which means the reactant must be a Brønsted-Lowry acid or base.
- 14. $NH_3 + H^+ \rightarrow NH_4^+$ $H_2O + H^+ \rightarrow H_2O^+$ $HCO_3^- + H^+ \rightarrow H_2CO_3$ $CO_3^{2-} + H^+ \rightarrow HCO_3^{-}$ $CN^{-} + H^{+} \rightarrow HCN$ $OH^- + H^+ \rightarrow H_2O$
- 15. $HCO_3^- + H_2O \rightarrow H_3O^+ + CO_3^{2-}$ $HCO_3^- + H_3O^+ \rightarrow H_2CO_3 + H_2O$ CO_3^{2-} is the conjugate base of HCO_3^{-} and H_2CO_3 is the conjugate acid of HCO_3 .
- 16. Acids: NH₄⁺, H₂O, HBr Bases: SO₃²⁻, AlCl₃,
- 17. (i) HCl, HCN, HS-, NH₄+ (ii) O²⁻, HSe⁻, NH₂-, CO₃²⁻

Set 2 Acidity

Multiple Choice Answers

1. a, 2. c, 3. c, 4. d, 5. c, 6. d, 7. d, 8. b, 9. e, 10. a, 11. c, 12. b, 13. d. 14. a, 15. c.

Answers to Longer Questions

Acid strength increases with increasing K_{σ} , acid strength decreases with decreasing K_z.

I ion is the conjugate base of a strong acid. It will simply be hydrated and stay as I. NO, ion is the conjugate base of a weak acid, HNO_{2} , and would react to form a base. $NO_2(aq) + H_2O(l) \rightleftharpoons HNO_2(aq) + OH(aq).$ K⁺ ion does not react with water except to get hydrated because the ions stay as

 $CO_3^2 + H_2O \rightleftharpoons HCO_3^2 + OH^2$ $HSO_{4}^{-} + H_{2}O \Rightarrow SO_{4}^{2-} + H_{3}O^{+}$

- KCl is the salt of the cation K⁺, which is a weaker acid than water and the anion Cl-, which is a weaker base than water. Because neither ion reacts with water to any extent, a KCl solution will be neutral. NH_4NO_2 is the salt of a cation and anion both of which can react with water. The solution will be nearly neutral. The pH will depend on the relative degree of ionisation of the anion and cation in water. *Na(HCOO) is the salt of a cation that does* not react with water and an anion that is stronger base than water (because it is the anion of a weak acid). The solution will therefore be basic.
- *NH*₄*CN*: close to neutral, because both the parent acid and base are weaker than water KI: close to neutral, because both the parent acid and base are stronger than water. *LiCH*₃*COO*: alkaline since the parent base *Li(OH)*, *is a stronger base than water.*
- (a) Acids: HNO_3 , H_3O^+ Bases: H_2O , NO_3^- (b) Acids: HCl, NH₄⁺ Bases: NH₃, Cl⁻ (c) Acids: H₂SO₄, H₃O⁺ Bases: H₂O, HSO₄ (d) Acids: H_2O , H_3O^+ Bases: H_2O , OH^-

 $CO_2 + H_2O \rightarrow H_2CO_3$, and, $H_2CO_3 + H_2O$ $\rightarrow H_3O^+ + HCO_3^-$

 $Na_2O + H_2O \rightarrow 2Na^+ + 2OH^-$

- (a) Hydrolysis is a reaction in which a salt reacts with water to produce a H^+ (H_3O^+) or an OH^- ion. Acidic salts give H_3O^+ ions and basic salts give OH- ions with water.
 - (b) Cations of the following acidic salts give H_3O^+ ions:

 $Fe^{3+} + 2H_2O \rightarrow Fe(OH)^{2+} + H_3O^+$ $NH_4^+ + H_2O \rightarrow NH_3 + H_3O^+$

 $HSO_4^- + H_2O \rightarrow SO_4^{2-} + H_3O^+$ (c) Anions of the following basic salts give OH- ions:

 $NH_2^- + H_2O \rightarrow NH_3 + OH^ CO_3^{2-} + H_2O \rightarrow HCO_3^{-} + OH^{-}$ $PO_4^{3-} + H_2O \rightarrow HPO_4^{2-} + OH^{-}$

 $BaCl_2 + H_2SO_4 \rightarrow BaSO_4 + 2HCl$ 1 mol of H_2SO_4 reacts with 1 mole of $BaCl_2$. \rightarrow mass of BaCl₂ = [0.1 × (137.3 + 70.9) × [0.032] = 0.667 g.

All are strong acids and completely dissociated.

> (a) $[H^+] = 1 \times 10^{-1} M$ $\therefore pH = 1.0$

> (b) $[H^+] = 1 \times 10^{-5} M$:. pH = 5.0

(c) $[OH^{-}] = 1 \times 10^{-2} M$: $[H^{+}] = 1 \times 10^{-12} M$:. pH = 12.0

(d) $[OH^{-}] = 2 \times 10^{-2} M$: $[H^{+}] = 5 \times 10^{-13} M$ $\rightarrow pH = 12.3$

(e) $[H^+] = 1.72 \times 10^{-2} \,\mathrm{M} : [H^+] = 1 \times 10^{-1.76}$

M : pH = 1.76

10. 1.0 L of 0.1 M HCl contains 0.1 mol of HCl. (a) HCl left after reaction with 0.010 mol of $KOH = 0.09 \ mol$ $[H^+] = 0.09 \text{ mol } L^{-1} = 9.0 \times 10^{-2} \text{M},$ $\therefore pH = -\log 9 \times 10^{-2} = 1.05$

> (b) $[OH^{-}] = 2 \times 0.05 = 0.1 \text{ mol}$: HCl left *after reaction is none* :: pH = 7.

(c) $[OH^{-}] = 0.1 \text{ mol}$: HCl left after reaction is none :. pH = 7

11. Assume 0.5 L of each solution is mixed. (a) Since $[H^+] = [OH^-]$ in the mixture, the pH is 7.

(b) Before reaction, there are 0.20×0.50 (0.1) mol) of HCl, and $0.10 \times 0.500 = (0.05 \ mol)$ of NaOH.

After the reaction 0.05 mole of HCl is left. \therefore $[H^+] = 0.05/1.0 = 0.05 \text{ mol } L^{-1}$

 $\therefore pH = -\log 0.05 = 1.30$

(c) Before reaction, there are 0.40×0.50 (0.2) mol) of HCl, and $0.20 \times 0.500 = (0.10 \ mol)$ of NaOH.

After the reaction 0.10 mole of HCl is left. :: $[H^+] = 0.10/1.0 = 0.1 \text{ mol } L^{-1}$

 $\therefore pH = -\log 0.1 = 1.00$

(d) Before reaction, there are 0.10×0.50 $(0.05 \text{ mol}) \text{ of HCl, and } 0.20 \times 0.50 = (0.10)$ mol) of OH- ions.

After the reaction 0.05 mole of OH- ions left. $\therefore [OH^{-}] = 0.05/1.0 = 0.05 \text{ mol } L^{-1}$

 $\therefore pOH = -\log 0.05 = 1.30$

so pH = 14 - 1.3 = 12.7.

12.

(i) 1.7×10^{-6} g in 1500 g of solution $= 1.13 \times 10^{-9}$ grams per gram 1.13×10^{-9} multiply by a million $= 1.13 \times 10^{-3} \ ppm.$

(ii) $n(HCl) = \frac{1.7 \times 10^{-6}}{36.45} = 4.664 \times 10^{-8} \, \text{mol}$

$$[H^+] = \frac{4.664 \times 10^{-8}}{1.5} = 3.109 \times 10^{-8} \, mol \, L^{-1}$$

(iii) Calculated $pH = -log(3.109 \times 10^{-8}) =$ 7.50 but, this concentration of H^+ ions is lower than that for neutral water and so the contribution from water cannot be disregarded. Total concentration of $H^+ = 3.109 \times 10^{-8}$

 $(acid) + 1 \times 10^{-7} (water) = 1.31 \times 10^{-7} \text{ mol } L^{-1}$ $pH = -log(1.31 \times 10^{-7}) = 6.88 (acidic!)$

13. $n(H_2SO_4) = (4.90/98.086) = 0.050 \text{ mol},$ $[H_2SO_4] = (0.050/0.100) = 0.50 \text{ M}$ $[H^+] = 1.3 \times 0.5 M = 0.65 M$ $\therefore pH = -log [0.65] = 0.187$

14. $2 \text{ NaOH} + H_2SO_4 \rightarrow \text{Na}_2SO_4 + 2H_2O$ $n (NaOH) = 0.025 \times 0.20 = 0.0050 \ mol$ (limiting reagent) $n(H_2SO_4)$ required to use up all the NaOH = $\frac{1}{2} \times 0.005 = 0.0025 \ mol$ For H_2SO_4 , $n(H_2SO_4) = 0.030 \times 0.175 =$ 0.00525 mol (excess reagent) H₂SO₄ is left over and is in excess by $(0.00525 - 0.0025) = 0.00275 \ mol$

:. Final $[H_2SO_4] = \frac{0.00275}{0.05} = 0.055 \text{ M}$:. Final $[H^+] = 2 \times 0.055 = 0.959 M$

 $\therefore pH = -\log [0.182] = 0.74$

15. $Ba(OH)_2 + 2HNO_3 \rightarrow Ba(NO_3)_2 + 2H_2O$ For $Ba(OH)_2$, $n = 0.050 \times 0.200 = 0.010$ mol = (limiting reagent) For HNO₃, $n = 0.200 \times 0.400 = 0.080 \text{ mol} =$ (excess reagent) HNO₃ is left over and is in excess. :. H^+ is in excess by 0.080 mol - 0.020 = $0.060\ mol.$ After dilution $[H^+]=0.06/6.0=1\times 10^{-2} M$

pH = 2. 16. (a) $[OH^{-}] = 2 \times 0.00500 = 0.0100 M$ $= 1.00 \times 10^{-2} M$

 $\therefore pOH = 2 pH = 14 - 2 = 12$

 \therefore The pH of the original solution = 12. (b) After making the final volume 1.0 L, the

final concentration is calculated using the dilution relationship, $c_1V_1 = c_2V_2$.

 $0.010 L \times 1.0 \times 10^{-2} M = 1.99 L \times c_2$

 $c_2 = [(0.010 \times 1.0 \times 10^{-2})/1.0 \text{ L}]$

 $= 1.0 \times 10^{-4} M$

 $\therefore [OH^{-}] = 1.0 \times 10^{-4} M$, and $[H^{+}]$

 $= 1.0 \times 10^{-10} M$

 $\therefore pH = 10.$

pH changes from 12 to 10.

(c) $n(Ca(OH)_2 = 0.00500 \times 0.010$

 $= 5.00 \times 10^{-5} \text{ mol}$

 $m [(Ca(OH)_2] = 5.00 \times 10^{-5} \times 74.09$

= 0.00370 g.

 $(d) Ca(OH)_2 + CO_2(g) \rightarrow CaCO_3(s) +$ $H_2O(l)$

 $n(CO_2) = n[(Ca(OH)_2] = 5.00 \times 10^{-5} \text{ mol}$ Using the relationship pV = nRT and V = $nRT \div p$, $V = [(5.00 \times 10^{-5} \times 8.314 \times 298)]$

(110)] = $1.126 \times 10^{-3} L = 1.126 mL$. 17. $CO_3^{2-} + H_2O \rightarrow HCO_3^{-} + OH^{-}(CO_3^{2-} \text{ is a})$ stronger base than H₂O) $S^{2-} + H_2O \rightarrow HS^- + OH^- (S^{2-} \text{ is a stronger})$

base than H,O)

 $CH_3COO^- + H_2O \rightarrow CH_3COOH + OH^-$ (CH_3COO^- is a stronger base than H_2O)

- 18. pH of 6 means $[H^+] = 1 \times 10^{-6}$ mol L^{-1} and a pH of 8 means $[H^+] = 1 \times 10^{-8}$ mol L^{-1} which is 100 times less than the first concentration.
- 19. (a) $(NH_4)_2CO_3(s) + water \rightarrow 2NH_4^+(aq) + CO_3^{2-}(aq)$ This reaction (reaction 1) is an ionisation

reaction.

 NH_4^+ (aq) produces H_3O^+ ions (reaction 2). It is from an acidic salt. Anions of basic salts produce H_3O^+ ions.

 CO_3^{2-} (aq) produces OH⁻ ions (reaction 3). It is from a basic salt. Cations of acidic salts produce OH⁻ ions.

Reactions 2 and 3 are hydrolysis reactions. NH_4^+ ion is a weak acid. It does not produce many H_3O^+ ions. Although the carbonate ion is in the minority, it has a higher K value so it will react with water to give an excess of OH^- ions.

20. In a series, acids such as H_3PO_4 , $H_2PO_4^-$ and HPO_4^{2-} the extent of dissociation decreases as H^+ ions had to be removed from negative ions and consequently their strength decreases also. (H_3PO_4 is the relatively strongest one of the three).

Set 3 Acids/Base Reactions

Multiple Choice Answers

1. a, 2. e, 3. e, 4. e, 5. b, 6. c, 7. c, 8. e, 9. d, 10. e, 11. b, 12. b, 13. c, 14. c, 15. b, 16. c, 17. c, 18. b, 19. b, 20. c.

Acid Base Calculations

- 1. No. of moles of $Ca(OH)_2 = cV = 0.015 \times 0.02 = 3.0 \times 10^{-4} \, mol$ $n(OH^-) = 2 \times 3.0 \times 10^{-4} \, mol = 6.0 \times 10^{-4} \, mol$ $(Ca(OH)_2 \rightarrow 2OH^-).$ $n(H^+)$ from $HNO_3 = 0.010 \times 0.080 = 8.0 \times 10^{-4} \, mol.$ OH- is the limiting reagent. So $6.0 \times 10^{-4} \, moles$ of H_2O is formed and $n(H^+)$ left will be $8.0 \times 10^{-4} 6.0 \times 10^{-4} = 2.0 \times 10^{-4} \, mol.$ Total volume of mixed solution = $20 + 80 = 100 \, mL$ so $[OH^-] = n/V = 2.0 \times 10^{-4}/0.10 = 2.0 \times 10^{-3} \, mol \, L^{-1}$
- pH = 2.72. $NaOH(aq) + HNO_3(aq) \rightarrow NaNO_3(aq) + H_2O(l)$ $n(NaOH) = c \times V = 0.0250 \times 0.200$ $= 5.00 \times 10^{-3} \ mol$ $n(HNO_3) = c \times V = 0.030 \times 0.175$ $= 5.25 \times 10^{-3} \ mol$ From the equation 1 mol of NaOH reacts

with 1 mol of HNO₃. ∴ 5.00×10^{-3} mol of NaOH will react with 5100×10^{-3} mol HNO₃. → HNO₃ is in excess by 0.25×10^{-3} mol and $[H^+] = n/V = (0.25 \times 10^{-3}/0.055)$ = 4.55×10^{-3} mol L^{-1} ∴ pH = 2.35

- 3. $Ba(OH)_2 + 2HNO_3 \rightarrow Ba(NO_3)_2 + 2H_2O(l)$ $n (Ba(OH)_2) = 0.0500 \times 0.200 = 0.0100$ mol $n (HNO_3) = 0.200 \times 0.400 = 0.0800$ mol From the equation, 1 mole of $Ba(OH)_2$ reacts with 2 mols of HNO_3 . Accordingly, 0.0100 mol of $Ba(OH)_2$ reacts with 0.0200 mol of HNO_3 . HNO_3 is in excess by 0.0600 mol. The final concentration after diluting to 6.00 L is [0.0600/6.00 L] = 0.01 M $[H^+] = 1.0 \times 10^{-2} \text{ M}$ and, pH = 2.0
- (a) pH = 2 $\therefore [H^+] = 1.0 \times 10^{-2} M$; volume = 0.300 L $\therefore n (H^+) needed = 1.0 \times 10^{-2} \times 0.300$ $= 3.00 \times 10^{-3} mol$ $Hence, n (HCl) needed = 3.00 \times 10^{-3} mol$ $\therefore m (HCl) = 3.00 \times 10^{-3} \times 36.46 = 0.109 g$
- (b) $n (HCl) = 0.0730 / 36.46 = 2.00 \times 10^{-3} \text{ mol.}$ $\therefore n (H^+) = 2.00 \times 10^{-3} \text{ mol.}$ $[H^+] = (2.00 \times 10^{-3} / 2.00 \text{ L}) = 1.0 \times 10^{-3} \text{ M}$ $\therefore pH = 3.00$
- (a) pH = 13, and hence $[H^+] = 1.0 \times 10^{-3} M$ and $[OH^-] = 1.0 \times 10^{-1} M$ V = 0.600 L and, $n (NaOH) = c \times v = 1 \times 10^{-1} \times 0.600$ $= 6.00 \times 10^{-2} mol$ $m (NaOH) = 6.00 \times 10^{-2} \times 40.0 = 2.40 g$.
- (b) m (NaOH) = 0.600 g and n (NaOH)= 0.600/40.0 = 0.0150 mol[NaOH] = n/V = 0.0150/1.500 = 0.01 M, and [OH-] = $1.0 \times 10^{-2} \text{ M}$ $\therefore [H^+] = 1.0 \times 10^{-12} \text{ M}$, thus pH = 12.0
- 6. n (HCl) = $0.100 \times 0.020 = 2.00 \times 10^{-3}$ mol pH required is 3. Therefore, [H⁺] should be 1.0×10^{-3} M.

 Since the number of moles are the same

Since the number of moles are the same before and after dilution, $0.100 \times 0.020 = 1.0 \times 10^{-3} \times V$

 $V = [(0.100 \times 0.020) / 1.0 \times 10^{-3}] = 2.0 L.$

- \therefore Volume of water to be added = 2.0 L 0.020 L = 1.98 L
- 7. pH = 4 and so $[H^+] = 1.0 \times 10^{-4}$ M. V = 1.00 L n (HCl) = $n(H^+) = c \times V = 1.0 \times 10^{-4} \times 1.00$ = 1.0×10^{-4} mol V (HCl @ STP) = $n \times 22.71$ = $1.0 \times 10^{-4} \times 22.71 = 2.27$ mL
- 8. All these hydrolyse to produce OH ions.