



ACIDS & BASES

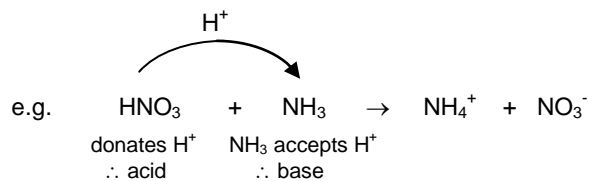
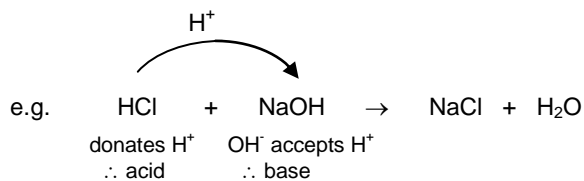


BRONSTED-LOWRY ACIDS & BASES

Bronsted-Lowry acid = **proton donor** (H^+ = proton)

Bronsted-Lowry base = **proton acceptor** (H^+ = proton)

Bronsted-Lowry acid-base reaction = **reaction involving the transfer of a proton**



TASK 1 – Bronsted-Lowry acids & bases

Identify the Bronsted-Lowry acid and base in each of the following reactions.

	acid	base
i) $\text{H}_2\text{O} + \text{NH}_3 \rightarrow \text{OH}^- + \text{NH}_4^+$		
ii) $\text{H}_2\text{O} + \text{HCl} \rightarrow \text{H}_3\text{O}^+ + \text{Cl}^-$		
iii) $\text{KOH} + \text{HCOOH} \rightarrow \text{HCOOK} + \text{H}_2\text{O}$		
iv) $\text{CH}_3\text{COOH} + \text{HCl} \rightarrow \text{CH}_3\text{COOH}_2^+ + \text{Cl}^-$		
v) $\text{NH}_3 + \text{HCl} \rightarrow \text{NH}_4\text{Cl}$		
vi) $\text{HCO}_3^- + \text{OH}^- \rightarrow \text{CO}_3^{2-} + \text{H}_2\text{O}$		
vii) $\text{HCO}_3^- + \text{H}^+ \rightarrow \text{CO}_2 + \text{H}_2\text{O}$		
viii) $\text{H}_2\text{SO}_4 + \text{HNO}_3 \rightarrow \text{HSO}_4^- + \text{H}_2\text{NO}_3^+$		

pH OF STRONG ACIDS

Monoprotic acid = **acid that releases one H⁺ ion per molecule**

e.g. HCl (hydrochloric acid), HNO₃ (nitric acid), CH₃COOH (ethanoic acid)

Diprotic acid = **acid that releases two H⁺ ions per molecule**

e.g. H₂SO₄ (sulfuric acid), H₂C₂O₄ (ethanedioic acid)

Moles of acid	Moles of H ⁺
3 moles of HNO ₃	
2 moles of HCl	
4 moles of H ₂ SO ₄	
0.3 moles of HNO ₃	
0.3 moles of H ₂ SO ₄	

Moles of acid	Moles of H ⁺
0.1 moles of H ₂ SO ₄	
0.2 moles of HCl	
0.08 moles of HNO ₃	
0.08 moles of H ₂ SO ₄	
0.35 moles of HCl	

Definition of pH

$$\text{pH} = -\log [\text{H}^+]$$

Useful rearrangement

$$[\text{H}^+] = 10^{-\text{pH}}$$

ALWAYS give pH to 2 DECIMAL PLACES

[H ⁺]	0.00100		1.50		2.5 x 10 ⁻⁴		4.5 x 10 ⁻¹²		
pH		2.75		3.30		13.70		1.85	-0.70

Calculating the pH of a strong acid

pH of 0.500 mol dm⁻³ HNO₃?

$$\begin{aligned} [\text{H}^+] &= 0.500 \\ \text{pH} &= -\log 0.500 \\ \text{pH} &= \underline{0.30} \end{aligned}$$

pH of 0.200 mol dm⁻³ H₂SO₄?

$$\begin{aligned} [\text{H}^+] &= 2 \times 0.200 = 0.400 \text{ (diprotic acid!)} \\ \text{pH} &= -\log 0.400 \\ \text{pH} &= \underline{0.40} \end{aligned}$$

[HCl] with pH 1.70?

$$\begin{aligned} [\text{H}^+] &= 10^{-1.70} = 0.0200 \\ [\text{HCl}] &= \underline{0.0200 \text{ mol dm}^{-3}} \end{aligned}$$

[H₂SO₄] with pH 1.30?

$$\begin{aligned} [\text{H}^+] &= 10^{-1.30} = 0.0501 \\ [\text{H}_2\text{SO}_4] &= 0.0501 / 2 = \underline{0.251 \text{ mol dm}^{-3}} \end{aligned}$$

Dilution of a strong acid

Calculate the pH of the solution formed when 100 cm³ of water is added to 50 cm³ of 0.100 mol dm⁻³ HNO₃.

$$[\text{H}^+] \text{ in original HNO}_3 \text{ solution} = 0.100$$

$$[\text{H}^+] \text{ in diluted solution} = 0.100 \times \frac{50}{150} = 0.0333$$

$$\text{pH} = -\log 0.0333 = \underline{1.48}$$

Calculate the pH of the solution formed when 250 cm³ of 0.300 mol dm⁻³ H₂SO₄ is made up to 1000 cm³ solution with water.

$$[\text{H}^+] \text{ in original H}_2\text{SO}_4 \text{ solution} = 2 \times 0.300 = 0.600$$

$$[\text{H}^+] \text{ in diluted solution} = 0.600 \times \frac{250}{1000} = 0.150$$

$$\text{pH} = -\log 0.150 = \underline{0.82}$$

TASK 2 – pH of strong acids

- 1) Calculate the pH of the following solutions.
 - a) 0.2 mol dm⁻³ HCl
 - b) 0.05 mol dm⁻³ HNO₃
 - c) 0.04 mol dm⁻³ H₂SO₄
 - d) 2.00 mol dm⁻³ HNO₃

- 2) Calculate the concentration of the following acids.
 - a) HCl with pH 3.55
 - b) H₂SO₄ with pH 1.70
 - c) HNO₃ with pH 1.30
 - d) H₂SO₄ with pH -0.50

- 3) Calculate the pH of the solutions formed in the following way.
 - a) addition of 250 cm³ of water to 50 cm³ of 0.200 mol dm⁻³ HNO₃
 - b) addition of 25 cm³ of water to 100 cm³ of 0.100 mol dm⁻³ H₂SO₄
 - c) adding water to 100 cm³ of 2.00 mol dm⁻³ H₂SO₄ to make 500 cm³ of solution
 - d) adding water to 25 cm³ of 1.50 mol dm⁻³ HCl to make 250 cm³ of solution

- 4) Calculate the pH of the following solutions.
 - a) 10 g dm⁻³ HCl
 - b) 20 g dm⁻³ H₂SO₄
 - c) 50 g dm⁻³ HNO₃
 - d) 100 g dm⁻³ H₂SO₄

THE IONIC PRODUCT OF WATER, K_w

The ionic product of water (K_w)



$$K_c = \frac{[\text{H}^+][\text{OH}^-]}{[\text{H}_2\text{O}]}$$

$$\therefore K_c [\text{H}_2\text{O}] = [\text{H}^+][\text{OH}^-]$$

As $[\text{H}_2\text{O}]$ is very much greater than $[\text{H}^+]$ and $[\text{OH}^-]$, then $[\text{H}_2\text{O}]$ is effectively a constant number

$$\therefore K_c [\text{H}_2\text{O}] = \text{a constant} = K_w$$

$$\boxed{K_w = [\text{H}^+][\text{OH}^-]}$$

The effect of temperature on the pH of water and the neutrality of water

As the temperature increases, the equilibrium moves right to lower the temperature

$$\therefore [\text{H}^+] \text{ and } [\text{OH}^-] \text{ increase}$$

$$\therefore K_w \text{ increases and } \therefore \text{pH increases}$$

However, the water is still neutral as $[\text{H}^+] = [\text{OH}^-]$ (and the definition of neutral is $[\text{H}^+] = [\text{OH}^-]$)

Calculating the pH of water

In pure water, $[\text{H}^+] = [\text{OH}^-]$

$$\therefore K_w = [\text{H}^+]^2$$

$$\therefore [\text{H}^+] = \sqrt{K_w}$$

Calculate the pH of water at 40°C when $K_w = 2.09 \times 10^{-14} \text{ mol}^2 \text{ dm}^{-6}$.

$$K_w = [\text{H}^+]^2$$

$$\therefore [\text{H}^+] = \sqrt{K_w} = \sqrt{(2.09 \times 10^{-14})} = 1.45 \times 10^{-7}$$

$$\therefore \text{pH} = -\log(1.45 \times 10^{-7}) = \underline{\underline{6.84}}$$

Calculate the pH of water at 100°C when $K_w = 51.3 \times 10^{-14} \text{ mol}^2 \text{ dm}^{-6}$.

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THE pH OF STRONG BASES

Monobasic base = base that releases one OH⁻ ion e.g. NaOH, KOH, NH₃

Dibasic base = base that releases two OH⁻ ions e.g. Ba(OH)₂, Ca(OH)₂

Calculating the pH of a strong base

pH of 0.200 mol dm⁻³ NaOH? [OH⁻] = 0.200

$$[H^+] = \frac{K_w}{[OH^-]} = \frac{10^{-14}}{0.200} = 5 \times 10^{-14}$$

$$pH = -\log[H^+] = -\log(5 \times 10^{-14}) = \underline{\underline{13.30}}$$

pH of 0.0500 mol dm⁻³ Ba(OH)₂? [OH⁻] = 2 x 0.0500 = 0.100

$$[H^+] = \frac{K_w}{[OH^-]} = \frac{10^{-14}}{0.100} = 1 \times 10^{-13}$$

$$pH = -\log[H^+] = -\log(1 \times 10^{-13}) = \underline{\underline{13.00}}$$

[KOH] with pH 12.70? [H⁺] = 10^{-pH} = 10^{-12.70} = 2.00 x 10⁻¹³

$$[OH^-] = \frac{K_w}{[H^+]} = \frac{10^{-14}}{2.00 \times 10^{-13}} = 0.05$$

$$[KOH] = \underline{\underline{0.05 \text{ mol dm}^{-3}}}$$

[Ba(OH)₂] with pH 13.30? [H⁺] = 10^{-pH} = 10^{-13.30} = 5.01 x 10⁻¹⁴

$$[OH^-] = \frac{K_w}{[H^+]} = \frac{10^{-14}}{5.01 \times 10^{-14}} = 0.200$$

$$[Ba(OH)_2] = \underline{\underline{0.100 \text{ mol dm}^{-3}}}$$

Dilution of a strong base

Calculate the pH of the solution formed when 50 cm³ of water is added to 100 cm³ of 0.200 mol dm⁻³ NaOH.

[OH⁻] in original NaOH solution = 0.200
 [OH⁻] in diluted solution = $0.200 \times \frac{100}{150} = 0.1333$

$$[H^+] = \frac{K_w}{[OH^-]} = \frac{10^{-14}}{0.1333} = 7.50 \times 10^{-14}$$

$$pH = -\log(7.50 \times 10^{-14}) = \underline{\underline{13.12}}$$

Calculate the pH of the solution formed when 50 cm³ of 0.250 mol dm⁻³ KOH is made up to 250 cm³ solution with water.

[OH⁻] in original KOH solution = 0.250
 [OH⁻] in diluted solution = $0.250 \times \frac{50}{250} = 0.0500$

$$[H^+] = \frac{K_w}{[OH^-]} = \frac{10^{-14}}{0.0500} = 2.00 \times 10^{-13}$$

$$pH = -\log(2.00 \times 10^{-13}) = \underline{\underline{12.70}}$$

TASK 3 – pH of strong bases

- 1) Calculate the pH of the following solutions.
 - a) 0.15 mol dm^{-3} KOH
 - b) 0.05 mol dm^{-3} NaOH
 - c) 0.20 mol dm^{-3} $\text{Ba}(\text{OH})_2$

- 2) Calculate the concentration of the following acids.
 - a) NaOH with pH 14.30
 - b) $\text{Ba}(\text{OH})_2$ with pH 12.50
 - c) KOH with pH 13.70

- 3) Calculate the pH of the solutions formed in the following way.
 - a) addition of 100 cm^3 of water to 25 cm^3 of $0.100 \text{ mol dm}^{-3}$ NaOH
 - b) addition of 25 cm^3 of water to 100 cm^3 of $0.100 \text{ mol dm}^{-3}$ $\text{Ba}(\text{OH})_2$
 - c) adding water to 100 cm^3 of 1.00 mol dm^{-3} KOH to make 1 dm^3 of solution

- 4) Calculate the pH of the following solutions.
 - a) 20 g dm^{-3} NaOH
 - b) 100 g dm^{-3} KOH
 - c) 1 g dm^{-3} $\text{Sr}(\text{OH})_2$

Reaction between a strong acid and a strong base

- 1) Calculate moles H^+
- 2) Calculate moles OH^-
- 3) Calculate moles XS H^+ or OH^-
- 4) Calculate XS $[\text{H}^+]$ or XS $[\text{OH}^-]$
- 5) Calculate pH

Calculate the pH of the solution formed when 50 cm^3 of $0.100 \text{ mol dm}^{-3}$ H_2SO_4 is added to 25 cm^3 of $0.150 \text{ mol dm}^{-3}$ NaOH.

$$\text{mol H}^+ = 2 \times \frac{50}{1000} \times 0.100 = 0.0100$$

$$\text{mol OH}^- = \frac{25}{1000} \times 0.150 = 0.00375$$

$$\therefore \text{XS mol H}^+ = 0.0100 - 0.00375 = 0.00625$$

$$\therefore \text{XS } [\text{H}^+] = \frac{0.00625}{\frac{75}{1000}} = 0.0833$$

$$\text{pH} = -\log(0.0833) = 1.08$$

Calculate the pH of the solution formed when 25 cm³ of 0.250 mol dm⁻³ H₂SO₄ is added to 100 cm³ of 0.200 mol dm⁻³ NaOH.

$$\text{mol H}^+ = 2 \times \frac{25}{1000} \times 0.250 = 0.0125$$

$$\text{mol OH}^- = \frac{100}{1000} \times 0.200 = 0.0200$$

$$\therefore \text{XS mol OH}^- = 0.0200 - 0.0125 = 0.0075$$

$$\therefore \text{XS [OH]} = \frac{0.0075}{\frac{125}{1000}} = 0.0600$$

$$\therefore \text{XS [H}^+] = \frac{K_w}{[\text{OH}^-]} = \frac{10^{-14}}{0.0600} = 1.67 \times 10^{-13}$$

$$\text{pH} = -\log(1.67 \times 10^{-13}) = \underline{12.78}$$

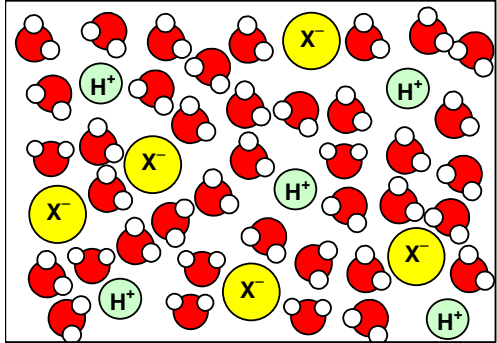
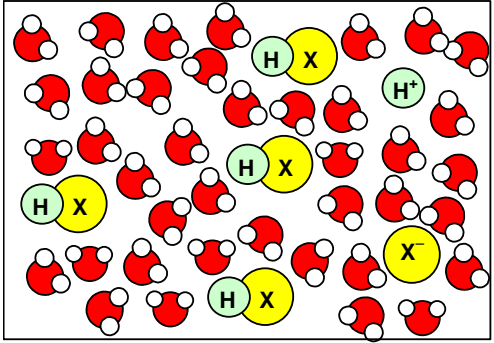
TASK 4 – strong acid + strong base

- 1) Calculate the pH of the solution formed when 20 cm³ of 0.100 mol dm⁻³ HNO₃ is added to 30 cm³ of 0.050 mol dm⁻³ KOH.
- 2) Calculate the pH of the solution formed when 25 cm³ of 0.150 mol dm⁻³ H₂SO₄ is added to 50 cm³ of 0.100 mol dm⁻³ NaOH.
- 3) Calculate the pH of the solution formed when 100 cm³ of 0.050 mol dm⁻³ HCl is added to 50 cm³ of 0.500 mol dm⁻³ KOH.
- 4) Calculate the pH of the solution formed when 10 cm³ of 1.00 mol dm⁻³ H₂SO₄ is added to 25 cm³ of 1.00 mol dm⁻³ NaOH.
- 5) Calculate the pH of the solution formed when 50 cm³ of 0.250 mol dm⁻³ HNO₃ is added to 50 cm³ of 0.100 mol dm⁻³ Ba(OH)₂.
- 6) Calculate the pH **change** to 100 cm³ of 0.200 mol dm⁻³ HCl solution in a flask if 50 cm³ of 0.100 mol dm⁻³ NaOH is added.
- 7) Calculate the pH **change** to 50 cm³ of 0.150 mol dm⁻³ KOH solution in a flask if 50 cm³ of 0.100 mol dm⁻³ H₂SO₄ is added.

TASK 5 – a mixture!

- 1) Calculate the pH of the following solutions:
 - a) 0.150 mol dm⁻³ Ba(OH)₂
 - b) 0.200 mol dm⁻³ HNO₃
 - c) 1.500 mol dm⁻³ H₂SO₄
 - d) 0.0500 mol dm⁻³ NaOH
- 2)
 - a) Calculate the pH of water at 50°C when $K_w = 5.48 \times 10^{-14} \text{ mol}^2 \text{ dm}^{-6}$.
 - b) is the water neutral? Explain your answer.
- 3)
 - a) 20 cm³ of 1.0 M H₂SO₄ with water added to make the volume up to 100 cm³.
 - b) 50 cm³ of 0.05 M KOH with 200 cm³ of water added.
- 4)
 - a) Calculate the pH of the solution formed when 100 cm³ of 0.100 mol dm⁻³ H₂SO₄ is added to 50 cm³ of 0.500 mol dm⁻³ NaOH.
 - b) Calculate the pH of the solution formed when 25 cm³ of 0.250 mol dm⁻³ HCl is added to 15 cm³ of 0.100 mol dm⁻³ KOH.
- 5) Calculate the pH of the solution formed when 3.5 g of impure sodium hydroxide (98.7 % purity) is dissolved in water and made up to 100 cm³, and then 25 cm³ of 0.35 mol dm⁻³ diprotic acid is added.

WEAK ACIDS

Strong acid	Weak acid
all the molecules break apart to form ions	only a small fraction of the molecules break apart to form ions
	
$\text{HX} \rightarrow \text{H}^+ + \text{X}^-$	$\text{HX} \rightleftharpoons \text{H}^+ + \text{X}^-$

Some common acids and bases

	Strong acids	Weak acids	Strong bases	Weak bases
Monoprotic / basic	HCl hydrochloric acid HNO ₃ nitric acid	carboxylic acids (e.g. ethanoic acid)	NaOH sodium hydroxide KOH potassium hydroxide	NH ₃ ammonia
Diprotic / basic	H ₂ SO ₄ sulphuric acid		Ba(OH) ₂ barium hydroxide	

The acid dissociation constant (K_a)



$$K_a = \frac{[\text{H}^+][\text{A}^-]}{[\text{HA}]} \quad \& \quad \text{p}K_a = -\log K_a \quad \& \quad K_a = 10^{-\text{p}K_a}$$

These expressions hold for weak acids at all times

Note

- K_a – has units mol dm⁻³
- K_a – the bigger the value, the stronger the acid
- $\text{p}K_a$ – the bigger the value, the weaker the acid

In a solution of a weak acid in water, with nothing else added:

a) $[\text{H}^+] = [\text{A}^-]$

- b) $[\text{HA}] \approx [\text{HA}]_{\text{initial}}$ (i.e. the concentration of HA at equilibrium is virtually the same as it was before any of it dissociated as so little dissociates, e.g. in a 0.100 mol dm⁻³ solution of HA, there is virtually 0.100 mol dm⁻³ of HA)

$$K_a = \frac{[\text{H}^+]^2}{[\text{HA}]}$$

This expression **ONLY** holds for weak acids in aqueous solution with nothing else added

Calculating the pH of a weak acid

Calculate the pH 0.100 mol dm⁻³ propanoic acid (pK_a = 4.87).

$$K_a = \frac{[H^+]^2}{[HA]}$$

$$[H^+]^2 = K_a [HA]$$

$$[H^+] = \sqrt{K_a [HA]} = \sqrt{(10^{-4.87} \times 0.100)} = 1.16 \times 10^{-3}$$

$$pH = -\log(1.16 \times 10^{-3}) = \underline{2.94}$$

Calculate the concentration of a solution of methanoic acid with pH 4.02 (K_a = 1.35 × 10⁻⁵ mol dm⁻³).

$$[H^+] = 10^{-4.02} = 9.55 \times 10^{-5}$$

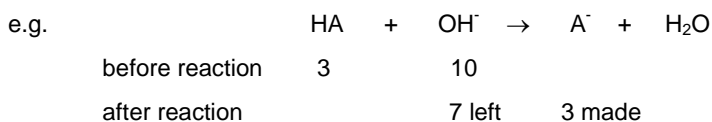
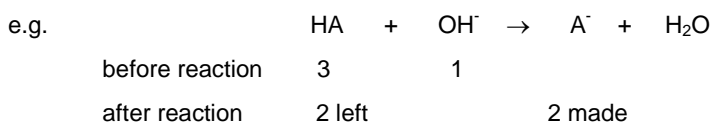
$$[HA] = \frac{[H^+]^2}{K_a} = \frac{(9.55 \times 10^{-5})^2}{1.35 \times 10^{-5}} = 6.76 \times 10^{-4} \text{ mol dm}^{-3}$$

TASK 6 – pH of weak acids

- 1) Calculate the pH of the following weak acids:
 - a) 0.150 mol dm⁻³ benenecarboxylic acid (pK_a = 4.20)
 - b) 0.200 mol dm⁻³ butanoic acid (K_a = 1.51 × 10⁻⁵ mol dm⁻³)
 - c) 1.00 mol dm⁻³ methanoic acid (K_a = 1.78 × 10⁻⁴ mol dm⁻³)
- 2) Calculate the concentration of the following weak acids.
 - a) ethanoic acid with pH 4.53 (K_a = 1.74 × 10⁻⁵ mol dm⁻³)
 - b) pentanoic acid with pH 3.56 (pK_a = 4.86)
- 3)
 - a) Which is the stronger acid, ethanoic acid (pK_a = 4.76) or propanoic acid (pK_a = 4.87)?
 - b) Which is the stronger acid, propanoic acid (1.35 × 10⁻⁵ mol dm⁻³) or propenoic acid (5.50 × 10⁻⁵ mol dm⁻³)?
- 4) Calculate the K_a value for phenylethanoic acid given that a 0.100 mol dm⁻³ solution has a pH of 2.66.

Reaction between a weak acid and a strong base

When a weak acid reacts with a strong base, for every mole of OH⁻ added, one mole of HA is used up and one mole of A⁻ is formed.



TASK 7 – Reaction of weak acids

When the following weak acids react with strong bases

- the moles of HA left after reaction
 - the moles of OH⁻ left after reaction
 - the moles of A⁻ formed in the reaction
- 1) 4 moles of HA with 2.5 moles of NaOH
 - 2) 6 moles of HA with 1.3 moles of Ba(OH)₂
 - 3) 0.15 moles of HA with 0.25 moles of KOH
 - 4) 0.30 moles of HA with 0.15 moles of NaOH
 - 5) 100 cm³ of 0.100 mol dm⁻³ HA with 50 cm³ 0.050 mol dm⁻³ NaOH
 - 6) 25 cm³ of 0.500 mol dm⁻³ HA with 40 cm³ of 1.0 mol dm⁻³ KOH
 - 7) 10 cm³ of 0.100 mol dm⁻³ HA with 10 cm³ of 0.080 mol dm⁻³ NaOH

Calculating the pH for the solution formed from reaction between a weak acid and a strong base

When a weak acid reacts with a strong base, for every mole of OH⁻ added, one mole of HA is used up and one mole of A⁻ is formed.

- 1) Calculate moles HA (it is still HA and not H⁺ as it is a weak acid)
- 2) Calculate moles OH⁻
- 3) Calculate moles XS HA or OH⁻

If XS HA

- 4) Calculate moles HA left and A⁻ formed
- 5) Calculate [HA] leftover and [A⁻] formed
- 6) Use K_a to find [H⁺]
- 7) Find pH

If XS OH⁻

- 4) Calculate [OH⁻]
- 5) Use K_w to find [H⁺]
- 6) Find pH

Note – if there is XS base, then in terms of working out the pH it is irrelevant whether it was a strong or weak acid as it has all reacted!

Calculate the pH of the solution formed when 30 cm³ of 0.200 mol dm⁻³ ethanoic acid (pK_a = 4.76) is added to 100 cm³ of 0.100 mol dm⁻³ NaOH.

$$\text{mol HA} = \frac{30}{1000} \times 0.200 = 0.00600$$

$$\text{mol OH}^- = \frac{100}{1000} \times 0.100 = 0.0100$$

OH⁻ is in XS

$$\text{XS mol OH}^- = 0.0100 - 0.00600 = 0.00400$$

$$\therefore \text{XS [OH}^-] = \frac{0.00400}{\frac{130}{1000}} = 0.0308$$

$$\therefore \text{XS [H}^+] = \frac{K_w}{[\text{OH}^-]} = \frac{10^{-14}}{0.0308} = 3.25 \times 10^{-13}$$

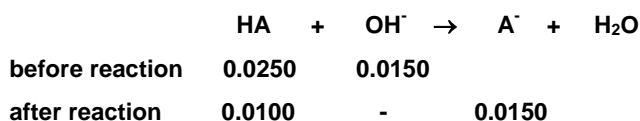
$$\text{pH} = -\log(3.25 \times 10^{-13}) = \underline{12.49}$$

Calculate the pH of the solution formed when 50 cm³ of 0.500 mol dm⁻³ ethanoic acid (pK_a = 4.76) is added to 75 cm³ of 0.200 mol dm⁻³ NaOH.

$$\text{mol HA} = \frac{50}{1000} \times 0.500 = 0.0250$$

$$\text{mol OH}^- = \frac{75}{1000} \times 0.200 = 0.0150$$

HA is in XS



$$\therefore \text{left over [HA]} = \frac{0.0100}{\frac{125}{1000}} = 0.0800$$

$$\therefore \text{formed [A}^-] = \frac{0.0150}{\frac{125}{1000}} = 0.120$$

$$K_a = \frac{[\text{H}^+][\text{A}^-]}{[\text{HA}]}$$

$$[\text{H}^+] = \frac{K_a [\text{HA}]}{[\text{A}^-]} = \frac{10^{-4.76} \times 0.0800}{0.120} = 1.16 \times 10^{-5}$$

$$\text{pH} = -\log(1.16 \times 10^{-5}) = 4.94$$

Half neutralisation of a weak acid

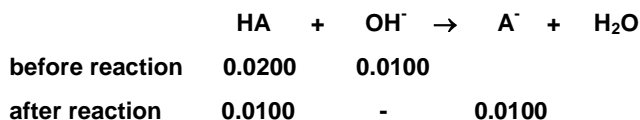
When half of the HA molecules have reacted with OH⁻, [HA] = [A⁻]. $\therefore K_a = [\text{H}^+]$ or $\text{pK}_a = \text{pH}$

Calculate the pH of the solution formed when 100 cm³ of 0.200 mol dm⁻³ ethanoic acid (pK_a = 4.76) is added to 40 cm³ of 0.250 mol dm⁻³ KOH.

$$\text{mol HA} = \frac{100}{1000} \times 0.200 = 0.0200$$

$$\text{mol OH}^- = \frac{40}{1000} \times 0.250 = 0.0100$$

HA is in XS



half neutralisation and so [HA] = [A⁻]

$$\therefore \text{pH} = \text{pK}_a = 4.76$$

TASK 8 – weak acid + strong base

- 1) Calculate the pH of the solution formed when 20 cm³ of 0.100 mol dm⁻³ methanoic acid (K_a = 1.7 × 10⁻⁴ mol dm⁻³) is added to 40 cm³ of 0.080 mol dm⁻³ KOH.
- 2) Calculate the pH of the solution formed when 50 cm³ of 0.500 mol dm⁻³ propanoic acid (pK_a = 4.87) is added to 100 cm³ of 0.080 mol dm⁻³ KOH.
- 3) Calculate the pH of the solution formed when 50 cm³ of 0.500 mol dm⁻³ ethanoic acid (pK_a = 4.76) is added to 50 cm³ of 0.250 mol dm⁻³ KOH.
- 4) Calculate the pH of the solution formed when 50 cm³ of 0.500 mol dm⁻³ chloroethanoic acid (pK_a = 2.86) is added to 25 cm³ of 0.100 mol dm⁻³ Ba(OH)₂.
- 5) Calculate the pH of the solution formed when 50 cm³ of 1.50 mol dm⁻³ dichloroethanoic acid (K_a = 0.0513 mol dm⁻³) is added to 100 cm³ of 2.00 mol dm⁻³ KOH.
- 6) Calculate the pH of the solution formed when 25 cm³ of 1.00 mol dm⁻³ benzenecarboxylic acid (pK_a = 4.20) is added to 50 cm³ of 0.0400 mol dm⁻³ NaOH.

TASK 9 – Another mixture

- 1) Calculate the pH of $0.100 \text{ mol dm}^{-3} \text{ H}_2\text{SO}_4$.
- 2) Calculate the pH of $0.250 \text{ mol dm}^{-3}$ methanoic acid ($K_a = 1.70 \times 10^{-4} \text{ mol dm}^{-3}$)
- 3) Calculate the pH of $0.20 \text{ mol dm}^{-3} \text{ Sr(OH)}_2$.
- 4) Calculate the pH of a mixture of 20 cm^3 of $0.500 \text{ mol dm}^{-3} \text{ NaOH}$ and 80 cm^3 of $0.200 \text{ mol dm}^{-3} \text{ HNO}_3$.
- 5) Calculate the pH of the solution formed when 100 cm^3 of water is added to 25 cm^3 of $0.100 \text{ mol dm}^{-3} \text{ NaOH}$.
- 6) Calculate the pH of a mixture of 25 cm^3 $0.200 \text{ mol dm}^{-3}$ ethanoic acid ($\text{p}K_a = 4.76$) and 25 cm^3 $0.100 \text{ mol dm}^{-3} \text{ NaOH}$.
- 7) Calculate the pH of a mixture of 100 cm^3 $0.100 \text{ mol dm}^{-3}$ ethanoic acid ($\text{p}K_a = 4.76$) and 50 cm^3 $0.150 \text{ mol dm}^{-3} \text{ NaOH}$.
- 8) Calculate the pH of a mixture of 50 cm^3 $0.200 \text{ mol dm}^{-3}$ propanoic acid ($\text{p}K_a = 4.87$) and 25 cm^3 $1.00 \text{ mol dm}^{-3} \text{ KOH}$.

pH CURVES & INDICATORS

- Indicators are weak acids where HA and A^- are different colours. $\text{HA} \rightleftharpoons \text{H}^+ + \text{A}^-$
- At low pH, HA is the main species present. At high pH, A^- is the main species present.
- The pH at which the colour changes varies from one indicator to another.
- Note that universal indicator is a mixture of indicators and so shows many colours at different pHs.

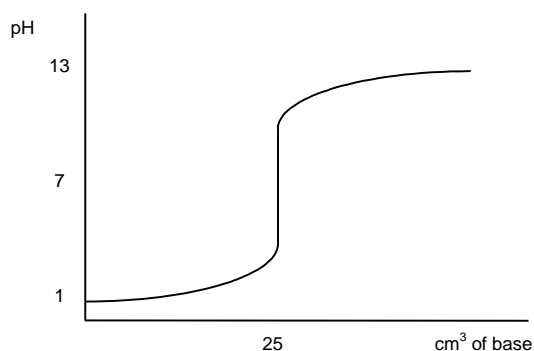
indicator	colour of HA	pH range of colour change	colour of A^-
methyl orange	red	3.2 - 4.4	yellow
phenolphthalein	colourless	8.2 - 10.0	pink

- In a titration, the pH changes rapidly at the end point as the last drop of acid/alkali is added. For an indicator to change colour at this moment where the moles of acid = moles of base, the indicator must change colour within the range of the rapid change in pH at the end point.

pH curves for monoprotic acids (e.g. HCl, HNO₃) with monoprotic bases

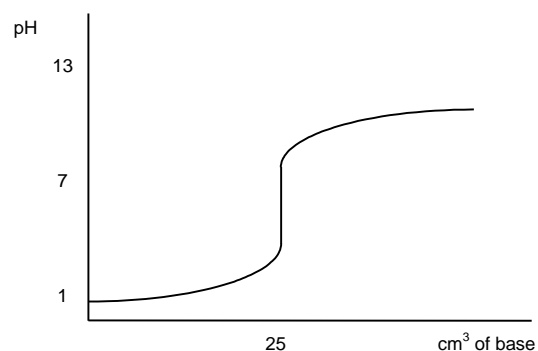
The curves below show the pH as 0.100 mol dm⁻³ base is added to 25.0 cm³ of 0.100 mol dm⁻³ acid:

a) strong acid - strong base



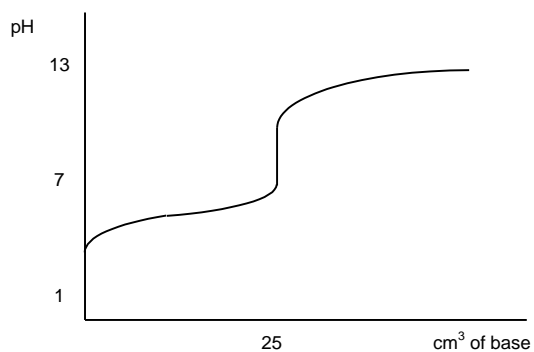
suitable indicators:

b) strong acid - weak base



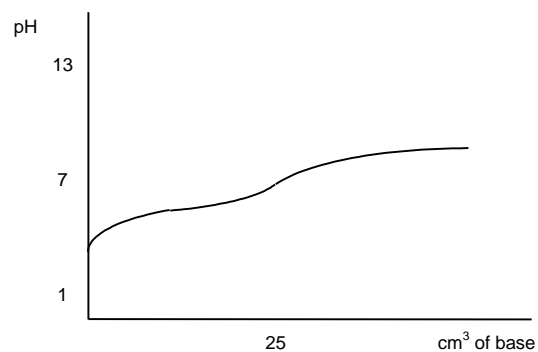
suitable indicators:

c) weak acid - strong base



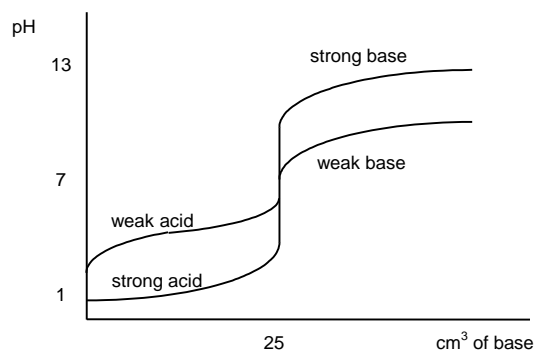
suitable indicators:

d) weak acid - weak base



suitable indicators:

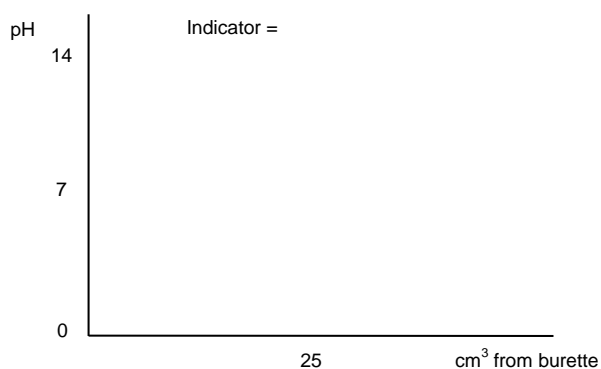
Summary:



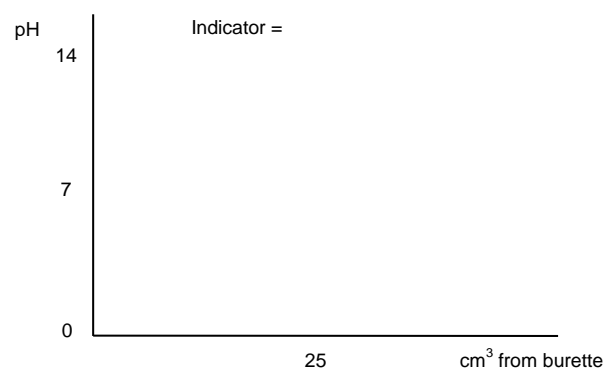
TASK 10 – sketching pH curves

Sketch each of the following pH curves on the grids shown, and name a suitable indicator.

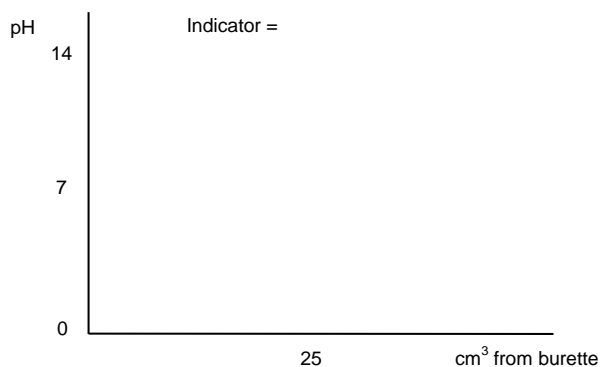
- (1)** Flask 25 cm³ 0.10 mol dm⁻³ HNO₃
Burette 50 cm³ 0.20 mol dm⁻³ NaOH



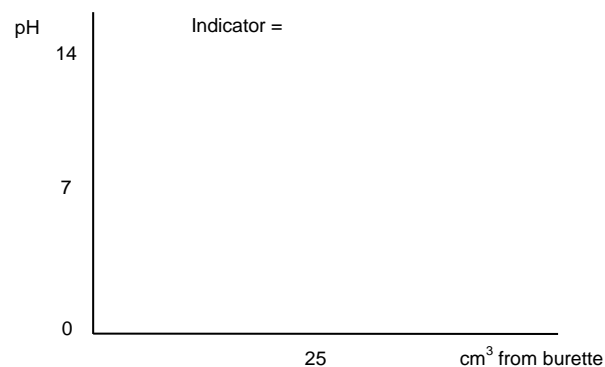
- (2)** Flask 20 cm³ 0.10 mol dm⁻³ NaOH
Burette 50 cm³ 0.10 mol dm⁻³ HCl



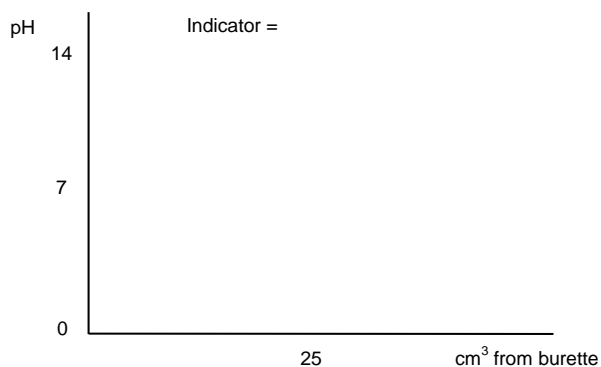
- (3)** Flask 10 cm³ 0.20 mol dm⁻³ HNO₃
Burette 50 cm³ 0.05 mol dm⁻³ NaOH



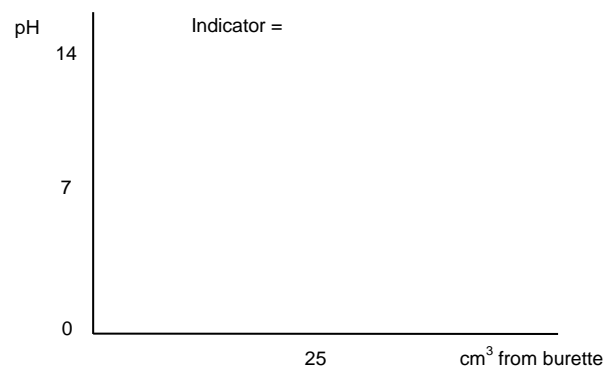
- (4)** Flask 30 cm³ 1.00 mol dm⁻³ NH₃
Burette 50 cm³ 1.00 mol dm⁻³ HCl



- (5)** Flask 20 cm³ 0.20 mol dm⁻³ CH₃COOH
Burette 50 cm³ 0.05 mol dm⁻³ NaOH

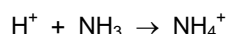
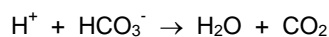
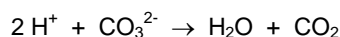
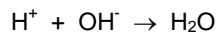


- (6)** Flask 50 cm³ 0.500 mol dm⁻³ NH₃
Burette 50 cm³ 1.00 mol dm⁻³ methanoic acid



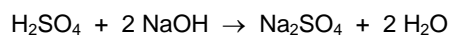
Titration calculations

Remember these ionic equations



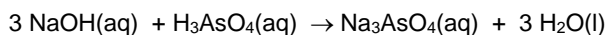
TASK 11 – Titration calculations

- 1) 25.0 cm³ of a solution of sodium hydroxide required 18.8 cm³ of 0.0500 mol dm⁻³ H₂SO₄.



- a) Find the concentration of the sodium hydroxide solution in mol dm⁻³.
b) Find the concentration of the sodium hydroxide solution in g dm⁻³.

- 2) 25.0 cm³ of arsenic acid, H₃AsO₄, required 37.5 cm³ of 0.100 mol dm⁻³ sodium hydroxide for neutralisation.



- a) Find the concentration of the acid in mol dm⁻³.
b) Find the concentration of the acid in g dm⁻³.

- 3) A 250 cm³ solution of NaOH was prepared. 25.0 cm³ of this solution required 28.2 cm³ of 0.100 mol dm⁻³ HCl for neutralisation. Calculate what mass of NaOH was dissolved to make up the original 250 cm³ solution.

- 4) 3.88 g of a monoprotic acid was dissolved in water and the solution made up to 250 cm³. 25.0 cm³ of this solution was titrated with 0.095 mol dm⁻³ NaOH solution, requiring 46.5 cm³. Calculate the relative molecular mass of the acid.

- 5) A 1.575 g sample of ethanedioic acid crystals, H₂C₂O₄.nH₂O, was dissolved in water and made up to 250 cm³. One mole of the acid reacts with two moles of NaOH. In a titration, 25.0 cm³ of this solution of acid reacted with exactly 15.6 cm³ of 0.160 mol dm⁻³ NaOH. Calculate the value of n.

- 6) A solution of a metal carbonate, M₂CO₃, was prepared by dissolving 7.46 g of the anhydrous solid in water to give 1000 cm³ of solution. 25.0 cm³ of this solution reacted with 27.0 cm³ of 0.100 mol dm⁻³ hydrochloric acid. Calculate the relative formula mass of M₂CO₃ and hence the relative atomic mass of the metal M.

- 7) A 1.00 g sample of limestone is allowed to react with 100 cm³ of 0.200 mol dm⁻³ HCl. The excess acid required 24.8 cm³ of 0.100 mol dm⁻³ NaOH solution. Calculate the percentage of calcium carbonate in the limestone.

- 8) An impure sample of barium hydroxide of mass 1.6524 g was allowed to react with 100 cm³ of 0.200 mol dm⁻³ hydrochloric acid. When the excess acid was titrated against sodium hydroxide, 10.9 cm³ of sodium hydroxide solution was required. 25.0 cm³ of the sodium hydroxide required 28.5 cm³ of the hydrochloric acid in a separate titration. Calculate the percentage purity of the sample of barium hydroxide.

BUFFER SOLUTIONS

What is a buffer solution?

- Buffer solution = solution that resists changes in pH when small amounts of acid or alkali are added.
- Note – the pH does change, just not by much!
- Acidic buffer solutions have a pH less than 7.
- Basic buffer solutions have a pH less than 7.

Examples of buffer solutions

- Acidic buffer solutions are made from a mixture of a weak acid and one of its salts (i.e. HA and A⁻) (e.g. ethanoic acid & sodium ethanoate).
- An acidic buffer solution can also be made by mixing an excess of a weak acid with a strong alkali as it results in a mixture of HA and A⁻.
- Basic buffer solutions are made from a mixture of a weak alkali and one of its salts (e.g. ammonia & ammonium chloride).
- A basic buffer solution can also be made by mixing an excess of a weak alkali with a strong acid

How acidic buffer solutions work

- This equilibrium exists in an acidic buffer solution: $HA \rightleftharpoons H^+ + A^-$
- There is a lot of HA and a lot of A⁻
- If a small amount of H⁺ is added: the equilibrium moves left to remove the H⁺ added; as the added H⁺ is removed, the pH remains roughly constant.
- If a small amount of OH⁻ is added: the OH⁻ reacts with and removes some H⁺; the equilibrium moves right to replace the OH⁻ removed; as the removed H⁺ is replaced, the pH remains roughly constant.

How basic buffer solutions work

- This equilibrium exists in a basic buffer solution: e.g. $NH_3 + H_2O \rightleftharpoons NH_4^+ + OH^-$
- There is a lot of NH₃ and a lot of NH₄⁺
- If a small amount of OH⁻ is added: the equilibrium moves left to remove the OH⁻ added; as the added OH⁻ is removed, the pH remains roughly constant.
- If a small amount of H⁺ is added: the H⁺ reacts with and removes some OH⁻; the equilibrium moves right to replace the OH⁻ removed; as the removed OH⁻ is replaced, the pH remains roughly constant.

Calculating the pH of acidic buffer solutions

A buffer solution was made by adding 2.05 g of sodium ethanoate to 0.500 dm³ of 0.01 mol dm⁻³ ethanoic acid. Calculate the pH of this solution (K_a for ethanoic acid = 1.74×10^{-5} mol dm⁻³).

$$M, \text{CH}_3\text{COONa} = 82.1$$

$$\text{mol CH}_3\text{COONa} = 2.05 / 82.1 = 0.0250$$

$$\text{mol CH}_3\text{COO}^- = 2.05 / 82.1 = 0.0250$$

$$[\text{A}^-] = \frac{0.0250}{0.500} = 0.0500$$

$$K_a = \frac{[\text{H}^+][\text{A}^-]}{[\text{HA}]}$$

$$[\text{H}^+] = \frac{K_a[\text{HA}]}{[\text{A}^-]} = \frac{1.74 \times 10^{-5} \times 0.01}{0.050} = 3.48 \times 10^{-6}$$

$$\text{pH} = -\log(3.48 \times 10^{-6}) = \underline{5.46}$$

TASK 12 – Buffer solution calculations

- 1) Calculate the pH of the following buffer solutions made by mixing weak acids with their salts.
 - a) 50 cm³ of 1.0 mol dm⁻³ methanoic acid ($K_a = 1.78 \times 10^{-4}$ mol dm⁻³) mixed with 20 cm³ of 1.0 mol dm⁻³ sodium methanoate.
 - b) 25 cm³ of 0.100 mol dm⁻³ butanoic acid ($\text{p}K_a = 4.82$) mixed with 20 cm³ of 0.100 mol dm⁻³ sodium butanoate.
 - c) 1.00 g of potassium ethanoate is dissolved in 50 cm³ of 0.200 mol dm⁻³ ethanoic acid ($K_a = 1.74 \times 10^{-5}$ mol dm⁻³).
- 2) Calculate the pH of the following buffer solutions made by mixing an excess of weak acids with strong bases.
 - a) 25 cm³ of 0.5 mol dm⁻³ methanoic acid ($K_a = 1.78 \times 10^{-4}$ mol dm⁻³) is mixed with 10 cm³ of 1.0 mol dm⁻³ sodium hydroxide.
 - b) 100 cm³ of 1.0 mol dm⁻³ ethanoic acid ($K_a = 1.78 \times 10^{-4}$ mol dm⁻³) is mixed with 50 cm³ of 0.8 mol dm⁻³ sodium hydroxide.
- 3)
 - a) What mass of sodium methanoate should be dissolved in 250 cm³ of 0.100 mol dm⁻³ methanoic acid to form a buffer solution with a pH of 5.20 (K_a for methanoic acid = 1.78×10^{-4} mol dm⁻³).
 - b) What mass of sodium ethanoate should be dissolved in 25 cm³ of 0.100 mol dm⁻³ ethanoic acid to form a buffer solution with a pH of 3.50 (K_a for ethanoic acid = 1.74×10^{-5} mol dm⁻³).
- 4)
 - a) 2 cm³ of 0.10 mol dm⁻³ NaOH is added to 100 cm³ of water. Calculate the **change** in pH of the water.
 - b) 2 cm³ of 0.10 mol dm⁻³ NaOH is added to 100 cm³ of a buffer solution containing 0.15 mol dm⁻³ ethanoic acid and 0.10 mol dm⁻³ sodium ethanoate (K_a ethanoic acid = 1.74×10^{-5} mol dm⁻³). Calculate the **change** in pH of the buffer solution.
 - c) Explain why the pH of the buffer solution only changes slightly compared to water.

GENERAL WORK

TASK 13 – A mixture of pH calculations

- 1) Calculate the pH of $0.100 \text{ mol dm}^{-3} \text{ H}_2\text{SO}_4$.
- 2) Calculate the pH of the solution formed when 200 cm^3 of water are added to 50 cm^3 of $0.500 \text{ mol dm}^{-3} \text{ HCl}$.
- 3) Calculate the pH of $0.500 \text{ mol dm}^{-3} \text{ NaOH}$ ($K_w = 1.00 \times 10^{-14} \text{ mol}^2 \text{ dm}^{-6}$).
- 4) Calculate the pH change when water is added to 25 cm^3 of $0.250 \text{ mol dm}^{-3} \text{ NaOH}$ to prepare 1.00 dm^3 of solution ($K_w = 1.00 \times 10^{-14} \text{ mol}^2 \text{ dm}^{-6}$).
- 5) Calculate the pH of $0.100 \text{ mol dm}^{-3}$ chloroethanoic acid given that $K_a = 1.38 \times 10^{-3} \text{ mol dm}^{-3}$.
- 6) Find the pH of $0.100 \text{ mol dm}^{-3}$ benzenecarboxylic acid ($K_a = 6.31 \times 10^{-5} \text{ mol dm}^{-3}$) when it has been half neutralised by NaOH.
- 7) Calculate the pH of water at 50°C given that $K_w = 5.476 \times 10^{-14} \text{ mol}^2 \text{ dm}^{-6}$ at 50°C and state and explain whether the water is still neutral.
- 8) Find the pH of the buffer solution prepared by adding 1.00 g of sodium ethanoate to 250 cm^3 of $0.100 \text{ mol dm}^{-3}$ ethanoic acid ($K_a = 1.74 \times 10^{-5} \text{ mol dm}^{-3}$).
- 9) Calculate the pH of the solution formed when 25 cm^3 of $0.100 \text{ mol dm}^{-3} \text{ NaOH}$ is added to 50 cm^3 of $0.250 \text{ mol dm}^{-3} \text{ HNO}_3$.
- 10) Calculate the pH of the solution formed when 100 cm^3 of $0.100 \text{ mol dm}^{-3} \text{ NaOH}$ is added to 20 cm^3 of $0.150 \text{ mol dm}^{-3} \text{ H}_2\text{SO}_4$.
- 11) Calculate the pH of the solution formed when 50 cm^3 of $0.100 \text{ mol dm}^{-3} \text{ NaOH}$ is added to 100 cm^3 of $0.300 \text{ mol dm}^{-3} \text{ CH}_3\text{COOH}$ ($K_a = 1.74 \times 10^{-5} \text{ mol dm}^{-3}$).
- 12) Calculate the pH of the solution formed when 50 cm^3 of $0.0500 \text{ mol dm}^{-3} \text{ Ba(OH)}_2$ is added to 20 cm^3 of $0.100 \text{ mol dm}^{-3} \text{ HCOOH}$ ($K_a = 1.78 \times 10^{-4} \text{ mol dm}^{-3}$).



FULL WORKED SOLUTIONS are available to subscribers of www.chemsheets.co.uk.

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TASK 1 – Bronsted-Lowry acids & bases

- | | | | | | |
|---|--|---|---|---|---|
| 1 | Acid = H ₂ O, base = NH ₃ | 2 | Acid = HCl, base = H ₂ O | 3 | Acid = HCOOH, base = KOH |
| 4 | Acid = HCl, base = CH ₃ COOH | 5 | Acid = HCl, base = NH ₃ | 6 | Acid = HCO ₃ ⁻ , base = OH ⁻ |
| 7 | Acid = H ⁺ , base = HCO ₃ ⁻ | 8 | Acid = H ₂ SO ₄ , base = HNO ₃ | | |

TASK 2 – pH of strong acids

- | | | | | | | | | |
|---|---|-------------------------|---|--------|---|--------|---|---------------------------|
| 1 | a | 0.70 | b | 1.30 | c | 1.10 | d | -0.30 |
| 2 | a | 2.82 x 10 ⁻⁴ | b | 0.0100 | c | 0.0501 | d | 1.58 mol dm ⁻³ |
| 3 | a | 1.48 | b | 0.80 | c | 0.10 | d | 0.82 |
| 4 | a | 0.56 | b | 0.39 | c | 0.10 | d | -0.31 |

TASK 3 – pH of strong bases

- | | | | | | | |
|---|---|-------|---|--------|---|----------------------------|
| 1 | a | 13.18 | b | 12.70 | c | 13.60 |
| 2 | a | 2.00 | b | 0.0158 | c | 0.501 mol dm ⁻³ |
| 3 | a | 12.30 | b | 13.20 | c | 13.00 |
| 4 | a | 13.70 | b | 14.25 | c | 12.22 |

TASK 4 – Strong acid + strong base

- | | | | | | | | | | |
|---|---------------------------------|---|------|---|----------------------------------|---|-------|---|------|
| 1 | 2.00 | 2 | 1.48 | 3 | 13.12 | 4 | 13.15 | 5 | 1.60 |
| 6 | new pH = 1.00, increase by 0.30 | | | 7 | new pH = 1.60, decrease by 11.58 | | | | |

TASK 5 – a mixture

- | | | | | | | | | |
|---|-------|-------|---|---|---|-------|---|-------|
| 1 | a | 13.48 | b | 0.70 | c | -0.48 | d | 12.70 |
| 2 | a | 6.63 | b | still neutral as [H ⁺] = [OH ⁻] | | | | |
| 3 | a | 0.40 | b | 12.00 | | | | |
| 4 | a | 12.52 | b | 0.93 | | | | |
| 5 | 13.74 | | | | | | | |

TASK 6 – pH of weak acids

- | | | | | | | |
|---|--|-------------------------|---|--|----------------|------|
| 1 | a | 2.51 | b | 2.76 | c | 1.87 |
| 2 | a | 5.00 x 10 ⁻⁵ | b | 5.50 x 10 ⁻³ mol dm ⁻³ | | |
| 3 | a | ethanoic acid | | b | propanoic acid | |
| 4 | 4.79 x 10 ⁻⁵ mol dm ⁻³ | | | | | |

TASK 7 – Reactions of weak acids

- 1) HA = 1.5, OH⁻ = 0, A⁻ = 2.5
- 2) HA = 3.4, OH⁻ = 0, A⁻ = 2.6
- 3) HA = 0, OH⁻ = 0.10, A⁻ = 0.15
- 4) HA = 0.15, OH⁻ = 0, A⁻ = 0.15
- 5) HA = 0.0075, OH⁻ = 0, A⁻ = 0.0025
- 6) HA = 0, OH⁻ = 0.0275, A⁻ = 0.0125
- 7) HA = 0.0002, OH⁻ = 0, A⁻ = 0.0008

TASK 8 – Weak acid + strong base

- | | | | | | | | | | | | |
|---|-------|---|------|---|------|---|------|---|-------|---|------|
| 1 | 12.30 | 2 | 4.54 | 3 | 4.76 | 4 | 2.26 | 5 | 13.92 | 6 | 3.14 |
|---|-------|---|------|---|------|---|------|---|-------|---|------|

TASK 9 – Another mixture

- | | | | | | | | | | |
|---|------|---|------|---|-------|---|------|---|-------|
| 1 | 0.70 | 2 | 2.19 | 3 | 13.60 | 4 | 1.22 | 5 | 12.30 |
| 6 | 4.76 | 7 | 5.24 | 8 | 13.30 | | | | |

TASK 11 – Titration calculations

- | | | | | | | | | | | | | |
|---|---|-----------------------------|---|-------------------------|---|---|---|---|---|-------|---|-------|
| 1 | a | 0.0752 mol dm ⁻³ | b | 3.01 g dm ⁻³ | | | | | | | | |
| 2 | a | 0.050 mol dm ⁻³ | b | 7.10 g dm ⁻³ | | | | | | | | |
| 3 | | 0.113 g | 4 | 87.8 | 5 | 2 | 6 | K | 7 | 87.7% | 8 | 90.8% |

TASK 12 – Buffer solution calculations

- | | | | | | | | | | | |
|---|---|---------------------------|---|--------------------------|---|------|--|--|--|--|
| 1 | a | 3.35 | b | 4.72 | c | 4.77 | | | | |
| 2 | a | 4.35 | b | 3.57 | | | | | | |
| 3 | a | 47.9 g | b | 0.0113 g | | | | | | |
| 4 | a | pH = 11.29, change = 4.29 | b | pH = 4.60, change = 0.02 | | | | | | |

TASK 13 – A mixture of pH calculations

- | | | | | | | | | | | | |
|----|---|---|------|---|-------|----|-------|----|------|---|------|
| 1 | 0.70 | 2 | 1.00 | 3 | 13.70 | 4 | 1.60 | 5 | 1.93 | 6 | 4.20 |
| 7 | 6.63, neutral as [H ⁺] = [OH ⁻] | 8 | 4.45 | 9 | 0.88 | 10 | 12.52 | 11 | 4.06 | | |
| 12 | 12.63 | | | | | | | | | | |