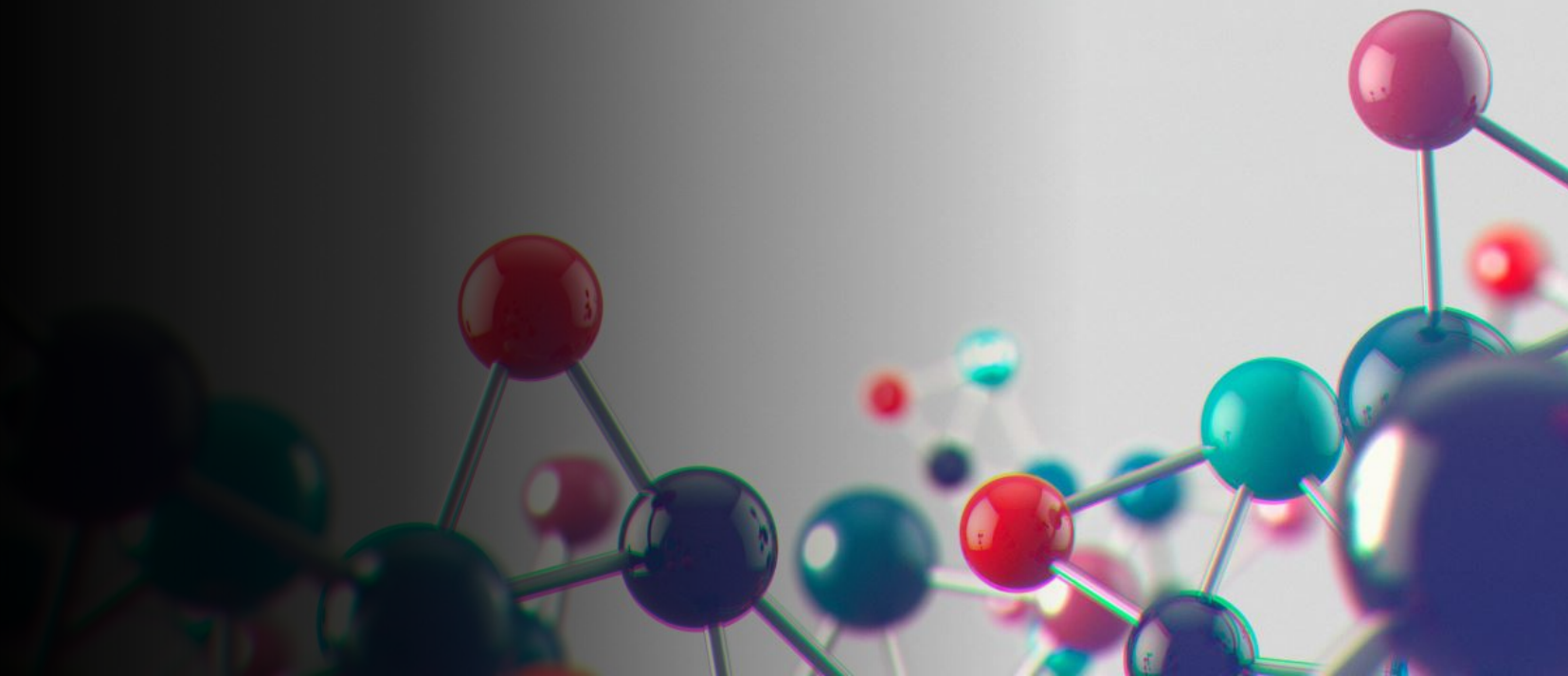


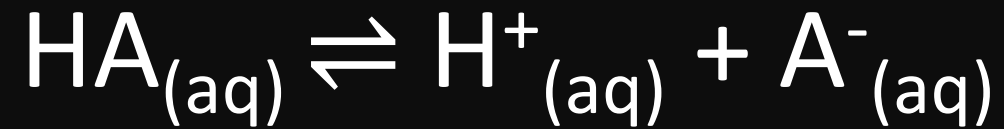
A2 Physical Chemistry

Buffer Calculations

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Acidic Buffers



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

$$[\text{H}^+_{(\text{aq})}] = K_a \times \frac{[\text{HA}_{(\text{aq})}]}{[\text{A}^-_{(\text{aq})}]}$$

Basic Buffers



$$K_b = \frac{[\text{HB}^+_{(\text{aq})}][\text{OH}^-_{(\text{aq})}]}{[\text{B}_{(\text{aq})}]}$$

$$[\text{OH}^-_{(\text{aq})}] = K_b \times \frac{[\text{B}^-_{(\text{aq})}]}{[\text{HB}^+_{(\text{aq})}]}$$

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).

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).

$$[H^+_{(aq)}] = K_a \times \frac{[HA_{(aq)}]}{[A^-_{(aq)}]}$$

$$\text{Moles of sodium ethanoate} = \frac{2.05}{82} = 0.025 \text{ moles}$$

$$\text{Conc of sodium ethanoate} = \frac{0.025}{0.500} = 0.050 \text{ mol dm}^{-3}$$

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$$[H^+_{(aq)}] = 1.74 \times 10^{-5} \times \frac{0.01}{0.050}$$

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$$[H^+_{(aq)}] = 3.48 \times 10^{-6} \text{ mol dm}^{-3}$$

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

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

$$\text{pH} = 5.46$$

A buffer solution was made by mixing 25.0 cm³ of 1.00 mol dm⁻³ ethanoic acid with 25 cm³ of 0.400 mol dm⁻³ sodium hydroxide.

(K_a for ethanoic acid = 1.74 × 10⁻⁵ mol dm⁻³).

Find the pH of this buffer.

A buffer solution was made by mixing 25.0 cm³ of 1.00 mol dm⁻³ ethanoic acid with 25 cm³ of 0.400 mol dm⁻³ sodium hydroxide. (K_a for ethanoic acid = 1.74×10^{-5} mol dm⁻³).

Find the pH of this buffer.

$$\begin{array}{l} \text{Moles of} \\ \text{Ethanoic acid} \end{array} = 0.025 \times 1.00 = 0.025 \text{ moles}$$

$$\begin{array}{l} \text{Moles of} \\ \text{NaOH} \end{array} = 0.025 \times 0.400 = 0.010 \text{ moles}$$



$$\text{Moles of ethanoic acid remaining} = 0.025 - 0.010 = 0.015 \text{ moles}$$

$$\text{Conc of ethanoic acid} = \frac{0.015}{0.050} = 0.300 \text{ mol dm}^{-3}$$

$$\begin{array}{l} \text{Moles of sodium} \\ \text{ethanoate formed} \end{array} = 0.010 \text{ moles} \qquad \begin{array}{l} \text{Conc of sodium} \\ \text{ethanoate} \end{array} = \frac{0.010}{0.050} = 0.200 \text{ mol dm}^{-3}$$

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Conc of ethanoic acid = 0.300 mol dm⁻³

Conc of sodium ethanoate = 0.200 mol dm⁻³

$$[H^+_{(aq)}] = K_a \times \frac{[HA_{(aq)}]}{[A^-_{(aq)}]}$$

$$[H^+_{(aq)}] = 1.74 \times 10^{-5} \times \frac{0.300}{0.200}$$

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Conc of sodium ethanoate = 0.200 mol dm⁻³

$$[H^+_{(aq)}] = K_a \times \frac{[HA_{(aq)}]}{[A^-_{(aq)}]}$$

$$[H^+_{(aq)}] = 2.61 \times 10^{-5} \text{ mol dm}^{-3}$$

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

$$\text{pH} = -\log(2.61 \times 10^{-5})$$

$$\text{pH} = 4.58$$

A buffer solution contains 0.20 mole of NH_3 and 0.60 mole NH_4Cl in 750 cm^3 . Calculate the pH of this solution (K_b for ammonia = $1.8 \times 10^{-5} \text{ mol dm}^{-3}$).

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$$[\text{NH}_{3(\text{aq})}] = \frac{0.20}{0.750} = 0.267 \text{ mol dm}^{-3}$$

$$[\text{NH}_4\text{Cl}] = \frac{0.60}{0.750} = 0.800 \text{ mol dm}^{-3}$$

$$[\text{OH}^-_{(\text{aq})}] = K_b \times \frac{[\text{B}_{(\text{aq})}]}{[\text{HB}^+_{(\text{aq})}]}$$

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$$[\text{OH}^-_{(\text{aq})}] = 1.8 \times 10^{-5} \times \frac{0.267}{0.800}$$

$$[\text{OH}^-_{(\text{aq})}] = 6.01 \times 10^{-6} \text{ mol dm}^{-3}$$

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

$$1 \times 10^{-14} = [\text{H}^+](6.01 \times 10^{-6})$$

$$[\text{H}^+] = 1.66 \times 10^{-9} \text{ mol dm}^{-3}$$

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

$$\text{pH} = -\log(1.66 \times 10^{-9})$$

$$\text{pH} = 8.78$$

Carbonic acid, H_2CO_3 , is a weak Brønsted–Lowry acid formed when carbon dioxide dissolves in water. Healthy blood at a pH of 7.40 has a hydrogencarbonate : carbonic acid ratio of 10.5 : 1.

A patient is admitted to hospital. The patient's blood pH is measured as 7.20. Calculate the hydrogencarbonate : carbonic acid ratio in the patient's blood.

$$[\text{H}^+_{(\text{aq})}] = 10^{-\text{pH}}$$

$$[\text{H}^+_{(\text{aq})}] = 10^{-7.40}$$

$$[\text{H}^+_{(\text{aq})}] = 3.98 \times 10^{-8}$$

$$[\text{H}^+_{(\text{aq})}] = K_a \times \frac{[\text{H}_2\text{CO}_{3(\text{aq})}]}{[\text{HCO}_3^-_{(\text{aq})}]}$$

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$$\frac{[\text{H}_2\text{CO}_{3(\text{aq})}]}{[\text{HCO}_3^-_{(\text{aq})}]} = \frac{6.31 \times 10^{-8}}{4.18 \times 10^{-7}} \quad \frac{[\text{HCO}_3^-_{(\text{aq})}]}{[\text{H}_2\text{CO}_{3(\text{aq})}]} = 6.6$$

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