# BUFFER SOLUTIONS

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## **Buffer solutions**

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## **Buffer solutions**

Before you start it would be helpful to ...

- know that weak acids and bases are only partly ionised in solution
- be able to calculate pH from hydrogen ion concentration
- be able to construct an equation for the dissociation constant of a weak acid

### **Buffer solutions - Brief introduction**

Definition

"Solutions which resist changes in pH when small quantities of acid or alkali are added."

Acidic Buffer (pH < 7) made from a weak acid + its sodium or potassium salt ethanoic acid sodium ethanoate

Alkaline Buffer (pH > 7) made from a

weak base ammonia

+ its chloride ammonium chloride

Uses Standardising pH meters Buffering biological systems (eg in blood) Maintaining the pH of shampoos

### **Buffer solutions - uses**

Definition

"Solutions which resist changes in pH when small quantities of acid or alkali are added."

#### **Biological Uses**

In biological systems (saliva, stomach, and blood) it is essential that the pH stays 'constant' in order for any processes to work properly. e.g. If the pH of blood varies by 0.5 it can lead to unconsciousness and coma

Most enzymes work best at particular pH values.

Other UsesMany household and cosmetic products need to control their pH values.ShampooBuffer solutions counteract the alkalinity of the soap and prevent irritationBaby lotionBuffer solutions maintain a pH of about 6 to prevent bacteria multiplyingOthersWashing powder, eye drops, fizzy lemonade

### **Buffer solutions - action**

It is essential to have a weak acid for an equilibrium to be present so that ions can be removed and produced. The dissociation is small and there are few ions.

$$CH_3COOH(aq)$$
 $CH_3COO^-(aq)$ + H+(aq)relative concs.HIGH $\leftarrow$ LOW

NB A strong acid can't be used as it is fully dissociated and cannot remove H<sup>+</sup>(aq)

 $HCl(aq) \longrightarrow Cl(aq) + H^{+}(aq)$ 

#### **Adding acid**

Any H<sup>+</sup> is removed by reacting with  $CH_3COO^-$  ions to form  $CH_3COOH$  via the equilibrium. Unfortunately, the concentration of  $CH_3COO^-$  is small and only a few H<sup>+</sup> can be "mopped up". A much larger concentration of  $CH_3COO^-$  is required.

To build up the concentration of CH<sub>3</sub>COO<sup>-</sup> ions, sodium ethanoate is added, which dissociates completely.

### **Buffer solutions - action**

We have now got an equilibrium mixture which contains a high concentration of <u>both</u> the undissociated weak acid,  $CH_3COOH(aq)$ , and its conjugate base,  $CH_3COO^-(aq)$ .

HIGH	LOW	+	LOW
CH <sub>3</sub> COO <sup>−</sup> Na <sup>+</sup> (aq) –	$-> CH_3COO^{-}(aq)$	+	Na+(aq)
	HIGH HIGH $CH_3COO^-Na^+ (aq) - 10W$	$\begin{array}{c} CH_3COOH(aq) & \longleftarrow & CH_3COO(aq) \\ HIGH & LOW \end{array}$ $\begin{array}{c} CH_3COO^-Na^+(aq) & \longrightarrow & CH_3COO^-(aq) \\ LOW & & HIGH \end{array}$	$\begin{array}{c} CH_{3}COOH(aq) & \longleftarrow & CH_{3}COO(aq) & + \\ HIGH & LOW \\ \\ CH_{3}COO^{-}Na^{+}(aq) & \longrightarrow CH_{3}COO^{-}(aq) & + \\ LOW & HIGH \end{array}$

#### Adding alkali

This adds OH ions

These react with the small concentration of H<sup>+</sup> ions: H<sup>+</sup>(aq) + OH<sup>-</sup>(aq) H<sub>2</sub>O(I) Removal of H<sup>+</sup> from the weak acid equilibrium means that more CH<sub>3</sub>COOH will dissociate to form ions to replace those being removed.

 $CH_3COOH(aq)$   $CH_3COO^{-}(aq)$  +  $H^+(aq)$ 

As the added OH<sup>-</sup> ions remove the H<sup>+</sup> from the weak acid system, the equilibrium moves to the right to produce more H<sup>+</sup> ions.

(There needs to be a large concentration of undissociated acid molecules to be available)

### **Buffer solutions - ideal concentration**

The concentration of a buffer solution is important

If the concentration is too low, there won't be enough  $CH_3COOH$  and  $CH_3COO^-$  to cope with the ions added.

#### Summary

For a buffer solution one needs ...

large [CH3COOH(aq)]for dissociating into H\*(aq) when alkali is addedweak acid(equilibrium shifts to the right)

large [CH3COO (aq)]- for removing H+(aq) as it is addedconjugate base(equilibrium shifts to the left)

This situation can't exist if only acid is present; a mixture of the acid and salt is used.

The weak acid provides the equilibrium and the large  $CH_3COOH(aq)$  concentration. The sodium salt provides the large  $CH_3COO^-(aq)$  concentration.

One uses a WEAK ACID + its SODIUM OR POTASSIUM SALT

### **Alkaline buffer solutions - action**

#### **Alkaline buffer**

Very similar but is based on the equilibrium surrounding a weak base; AMMONIA

	$NH_3(aq) + H_2O(l)$	 OH <sup>-</sup> (aq) +	NH <sub>4</sub> +(aq)
relative concs.	HIGH	LOW	LOW

but one needs ; a large conc. of  $OH^{-}(aq)$  to react with any  $H^{+}(aq)$  added a large conc of  $NH_4^{+}(aq)$  to react with any  $OH^{-}(aq)$  added

There is enough NH<sub>3</sub> to act as a source of OH<sup>-</sup> but one needs to increase the concentration of ammonium ions by adding an ammonium salt.

Use AMMONIA (a weak base) + AMMONIUM CHLORIDE (one of its salts)

Calculate the pH of a buffer whose [HA] is 0.1 mol dm<sup>-3</sup> and [A<sup>-</sup>] of 0.1 mol dm<sup>-3</sup>. The K<sub>a</sub> of the weak acid HA is 2 x 10<sup>-4</sup> mol dm<sup>-3</sup>

Calculate the pH of a buffer whose [HA] is 0.1 mol dm<sup>-3</sup> and [A<sup>-</sup>] of 0.1 mol dm<sup>-3</sup>. The K<sub>a</sub> of the weak acid HA is 2 x 10<sup>-4</sup> mol dm<sup>-3</sup>

Calculate the pH of a buffer whose [HA] is 0.1 mol dm<sup>-3</sup> and [A<sup>-</sup>] of 0.1 mol dm<sup>-3</sup>.

$$K_{a} = [H^{+}(aq)] [A^{-}(aq)]$$
$$[HA(aq)]$$
$$[H^{+}(aq)] = [HA(aq)] \times K_{a}$$

[A (aq)]

re-arrange

Calculate the pH of a buffer whose [HA] is 0.1 mol dm<sup>-3</sup> and [A<sup>-</sup>] of 0.1 mol dm<sup>-3</sup>.

	$K_a = [H^+(aq)] [A^-(aq)]$		
	[HA(aq)]		
re-arrange [H+(ac	$[A] = [HA(aq)] \times K_a$ $[A^{-}(aq)]$		
rom information given	[A <sup>-</sup> ] = 0.1 mol dm <sup>-3</sup> [HA] = 0.1 mol dm <sup>-3</sup>		

Calculate the pH of a buffer whose [HA] is 0.1 mol dm<sup>-3</sup> and [A<sup>-</sup>] of 0.1 mol dm<sup>-3</sup>.

from information given  $[A^-] = 0.1 \text{ mol } dm^{-3}$ [HA] = 0.1 mol  $dm^{-3}$ 

If the  $K_a$  of the weak acid HA is 2 x 10<sup>-4</sup> mol dm<sup>-3</sup>.

$$[H^+(aq)] = \underbrace{0.1 \times 2 \times 10^{-4}}_{0.1} = 2 \times 10^{-4} \text{ mol dm}^{-3}$$



Calculate the pH of a buffer whose [HA] is 0.1 mol dm<sup>-3</sup> and [A<sup>-</sup>] of 0.1 mol dm<sup>-3</sup>.

$$K_a = [H^+(aq)] [A^-(aq)]$$
  
[HA(aq)]

re-arrange 
$$[H^+(aq)] = [HA(aq)] \times K_a$$
  
 $[A^-(aq)]$ 

from information given  $[A^-] = 0.1 \text{ mol } dm^{-3}$ [HA] = 0.1 mol  $dm^{-3}$ 

If the  $K_a$  of the weak acid HA is 2 x 10<sup>-4</sup> mol dm<sup>-3</sup>.

$$[H^+(aq)] = \underbrace{0.1 \times 2 \times 10^{-4}}_{0.1} = 2 \times 10^{-4} \text{ mol dm}^{-3}$$

 $pH = -log_{10} [H^+(aq)] = 3.699$ 

Calculate the pH of the solution formed when  $500 \text{ cm}^3$  of 0.1 mol dm<sup>-3</sup> of weak acid HX is mixed with  $500 \text{ cm}^3$  of a 0.2 mol dm<sup>-3</sup> solution of its salt NaX. K<sub>a</sub> = 4 x 10<sup>-5</sup> mol dm<sup>-3</sup>.

Calculate the pH of the solution formed when 500cm<sup>3</sup> of 0.1 mol dm<sup>-3</sup> of weak acid HX is mixed with 500cm<sup>3</sup> of a 0.2 mol dm<sup>-3</sup> solution of its salt NaX.  $K_a = 4 \times 10^{-5}$  mol dm<sup>-3</sup>.

$$K_{a} = [H^{+}(aq)] [X^{-}(aq)]$$
  
[HX(aq)]

Calculate the pH of the solution formed when 500cm<sup>3</sup> of 0.1 mol dm<sup>-3</sup> of weak acid HX is mixed with 500cm<sup>3</sup> of a 0.2 mol dm<sup>-3</sup> solution of its salt NaX.  $K_a = 4 \times 10^{-5}$  mol dm<sup>-3</sup>.

$$K_{a} = [H^{+}(aq)] [X^{-}(aq)$$
[HX(aq)]
  
re-arrange
$$[H^{+}(aq)] = [HX(aq)] K_{a}$$

$$[X^{-}(aq)]$$

Calculate the pH of the solution formed when  $500 \text{ cm}^3$  of 0.1 mol dm<sup>-3</sup> of weak acid HX is mixed with  $500 \text{ cm}^3$  of a 0.2 mol dm<sup>-3</sup> solution of its salt NaX. K<sub>a</sub> = 4 x 10<sup>-5</sup> mol dm<sup>-3</sup>.

$$K_{a} = [H^{+}(aq)] [X^{-}(aq)]$$

$$[HX(aq)]$$
re-arrange
$$[H^{+}(aq)] = [HX(aq)] K_{a}$$

$$[X^{-}(aq)]$$

The solutions have been mixed; the volume is now 1 dm<sup>3</sup>

therefore [HX] =  $0.05 \text{ mol dm}^{-3}$  and [X<sup>-</sup>] =  $0.10 \text{ mol dm}^{-3}$ 



Calculate the pH of the solution formed when  $500 \text{ cm}^3$  of 0.1 mol dm<sup>-3</sup> of weak acid HX is mixed with  $500 \text{ cm}^3$  of a 0.2 mol dm<sup>-3</sup> solution of its salt NaX. K<sub>a</sub> = 4 x 10<sup>-5</sup> mol dm<sup>-3</sup>.

$$K_{a} = [H^{+}(aq)] [X^{-}(aq)]$$
[HX(aq)]
  
re-arrange
$$[H^{+}(aq)] = [HX(aq)] K_{a}$$

$$[X^{-}(aq)]$$

The solutions have been mixed; the volume is now 1 dm<sup>3</sup>

pH

=

therefore 
$$[HX] = 0.05 \text{ mol dm}^{-3}$$
 and  
 $[X^-] = 0.10 \text{ mol dm}^{-3}$   
Substituting  $[H^+(aq)] = \underbrace{0.05 \times 4 \times 10^{-5}}_{0.1} = 2 \times 10^{-5} \text{ mol dm}^{-3}$ 

- log<sub>10</sub> [H<sup>+</sup>(aq)]

4.699