# Chemistry 3.6 Aqueous Systems

Acids and Bases

#### From Level 2

Water dissociation  $2H_2O \rightleftharpoons H_3O^+ + OH^ K_{w} = [H_3O^+][OH^-] = 10^{-14}$ "p" = -log  $pH = -log [H_3O^+]$  $10^{-pH} = [H_3O^+]$ pH + pOH = 14

# Dissociating

- Once a solid dissolve, each species can further under goes **dissociation** with **water**.
- If it is an acid species, it will donates a proton
   (H<sup>+</sup>) forming a conjugate base and H<sub>3</sub>O<sup>+</sup> ions
- If it is a base species, it will accepts a proton (H<sup>+</sup>) forming conjugate acid and OH<sup>-</sup> ions

# Strong acid and weak acid

Strong acid fully dissociate when it is dissolved

$$HCl_{(g)} \rightarrow H^+_{(aq)} + Cl^-_{(aq)}$$

 Weak acid first dissolve then partially dissociate

$$\begin{aligned} \mathsf{NH}_4\mathsf{Cl}_{(s)} &\xrightarrow{} \mathsf{NH}_4^+_{(aq)} + \mathsf{CI}^- \\ \mathsf{H}_2\mathsf{O}_{(\mathsf{I})} + \mathsf{NH}_4^+_{(aq)} &\rightleftharpoons \mathsf{NH}_{3(aq)} + \mathsf{H}_3\mathsf{O}^+_{(aq)} \end{aligned}$$

# Strong acid Calculation

Because strong acid fully dissociate, the concentration of H<sub>3</sub>O<sup>+</sup> (H<sup>+</sup>) ions are the same as displayed in the formula of acid

Example

[H<sup>+</sup>] in 0.1 molL<sup>-1</sup> of HCl = 0.1 molL<sup>-1</sup> [H<sup>+</sup>] in 0.1 molL<sup>-1</sup> of H<sub>2</sub>SO<sub>4</sub> = 0.2 molL<sup>-1</sup> pH =  $-\log[H^+]$ 

## Weak acid calculation

• K<sub>a</sub> is usually given  

$$K_a = \frac{[H_3O^+][A^-]}{[HA]}$$

- The assumption is that most of the acid molecules do not dissociate, therefore [HA] is very similar to the original concentration cHA
- Another assumption is  $[H_3O^+] = [A^-]$ . Rearranging the formula gives:  $[H_3O^+] = \sqrt{K_a \times cHA}$

#### Example

Calculate the pH of 0.05 molL<sup>-1</sup> of ethanoic acid solution.  $K_a = 1.74 \times 10^{-5}$  $CH_3COOH(aq) + H_2O(l)$   $CH_3COO^- + H_3O^+$  $K_a = \frac{[H_3O^+][CH_3COO^-]}{[CH_3COOH]} = 1.74 \times 10^{-5}$  $[H_3 O^+] = \sqrt{(1.74 \times 10^{-5}) \times 0.05}$  $[H_3O^+] = 9.33 \times 10^{-4}$  $pH = -log (9.33 \times 10^{-4}) = 3.03$ 

## Strong base and weak base

- Strong base fully dissociate when it dissolve.  $NaOH_{(s)} \rightarrow Na^{+}_{(aq)} + OH^{-}_{(aq)}$
- Weak base first dissolve then dissociate partially.

 $NH_{3(g)} \rightarrow NH_{3(aq)}$  $NH_{3(aq)} + H_2O_{(I)} \rightleftharpoons NH_4^+_{(aq)} + OH^-_{(aq)}$ 

## Strong base calculation

- Calculation the concentration of OH<sup>-</sup>
- $10^{-14} \div [OH^{-}] = [H_3O^{+}]$
- $-\log [H_3O^+] = pH$

#### Weak base calculation

 $A^- + H_2O \implies HA + OH^-$ 

• K<sub>b</sub> is given in terms of K<sub>a</sub> because...

$$K_b = \frac{K_w}{K_a} = \frac{[HA][OH^-]}{[A^-]}$$

- Same assumption with weak acid that the concentration of weak base [A<sup>-</sup>] ≈ cA<sup>-</sup>
- [OH<sup>-</sup>] = [HA] therefore

$$K_b = \frac{[HA][OH^-]}{[A^-]}$$
$$[OH^-] = \sqrt{K_b \times cA^-}$$

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