**Disclaimer**: Although care has been taken to ensure the accuracy and reliability of the information provided, no responsibility is taken for lack of. Information provided is intended to aid rather than be a replacement

# Lecture 23

## Acids and bases

Lewis base: electron pair donor Lewis acid: electron pair acceptor Bronsted acid: proton donor

Bronsted acid: proton acceptor

• Degree of hydration with acids is uncertain (H<sub>3</sub>O<sup>+</sup>, H<sub>5</sub>O<sub>2</sub><sup>+</sup>, H<sub>7</sub>O<sub>3</sub><sup>+</sup>...)

# Conjugate acid-base pairs

B (base) + HA (acid)  $\rightleftharpoons$  HB<sup>+</sup> (conjugate acid) + A<sup>-</sup> (conjugate base) HB<sup>+</sup> (acid) + A<sup>-</sup> (base)  $\rightleftharpoons$  B (conjugate base) + HA (conjugate acid)

#### Water

Self ionizes however position of equilibrium lies towards the left

$$K_w = [H^+][OH^-] = 10^{-14}$$
 at 25°C

In pure water:  $[H^+] = [OH^-]$  hence solution is neutral

- $pH = -\log_{10}[H^+]$
- $pOH = -\log_{10}[OH^-]$
- pH + pOH = 14

## **Acid strength**

$$H_2O_{(1)} + HA_{(aq)} \rightleftharpoons H_3O^+_{(aq)} + A^-_{(aq)}$$
 $K_a = \frac{[H_3O^+][A^-]}{[HA]}$ 
 $pK_a = -\log_{10}(K_a)$ 

Strong acids have a high  $K_a$  but a small  $pK_a$ 

### **Base strength**

$$H_2O_{(1)} + B_{(aq)} \rightleftharpoons OH_{(aq)}^- + BH_{(aq)}^+$$
  
 $K_b = \frac{[OH^-][BH^+]}{[B]}$   
 $pK_b = -\log_{10}(K_b)$ 

Strong bases have high  $K_b$  but a small  $pK_b$ 

### Strong acids

- Ionises completely in aqueous solutions
- Anions are very weak bases
- Conjugate base of strong acid is a very weak base

#### Strong base

Conjugate acid of strong base is a very weak acid

$$K_a \times K_b = [H^+][OH^-]$$
  
 $K_a \times K_b = K_w$   
 $pK_a + pK_b = 14$ 

#### Weak acids

- Undergo a small degree of dissociation
- Conjugate base of a weak acid is also a weak base
- $pH = \frac{1}{2}pK_a \frac{1}{2}\log_{10}[HA]$
- Assumption: initial [HA] can be used as only a very small amount dissociates
- If not making assumption:  $K_a = \frac{[x]^2}{[initial-x]}$  where  $x = [H_3O^+]$  5% rule: if dissociation is < 5% ( $\frac{[dissociation]}{[initial]}$ ), then
- approximation/assumption is valid

### Weak bases

- Undergo a small degree of protonation
- $pOH = \frac{1}{2}pK_b \frac{1}{2}\log_{10}[B]$

# **Amines**

- · Bases that are like ammonia
- Reacts with water to produce OH<sup>-</sup>
- · Reacts with acids to form acid salts therefore enhancing its solubility