# 03 Ionic Equilibria

## Subtopics

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- 3.2 Types of electrolyte
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Red cabbage as a pH indicator

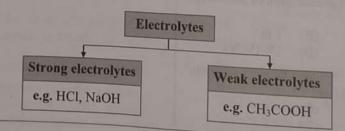


The red cabbage plant changes its colour according to the acidity or basicity of the soil in which is cultivated. In acidic soils, the leaves are reddisting neutral soils the leaves are purple, while in alkaling soils the leaves are greenish-yellow coloured. This because they contain chemicals from the natural coloured anthocyanin family of compounds.

The juice of red cabbage can be used as a home-made pH indicator. The juice is red, pink a magenta in acids, (pH < 7), purple in neuron solutions  $(pH \sim 7)$ , and ranges from blue to green yellow in alkaline solutions (pH > 7).

#### Quick Review

Types of electrolytes:



Various

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theory

Lewis

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Basic

Types

Salts of str

base
 Solution was neutral.

e.g. NaCl, KNO<sub>3</sub>, Na Various theories of acids and bases:

#### Chapter 03: Ionic Equilibria

Theory	Acid	The second second second second
Arrhenius theory	A substance that contains hydrogen and produces H <sup>+</sup> ions in aqueous solution.	Base A substance that contains OH group and
Bronsted-Lowry theory	Any substance that can donate a proton (H <sup>+</sup> ) i.e., proton donor.	produces OH ions in aqueous solution.  Any substance that can accept a proton i.e.,
Lewis theory	Any species that can accept a share in an electron pair.	Any species that can donate a share in an electron pair.

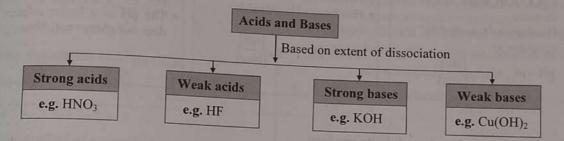
pH indicator

colour according I in which it is are reddish; in hile in alkaline oloured. This is n the naturally

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red, pink, or le in neutral ue to greenish

## Classification of acids and bases:



pH of solutions:

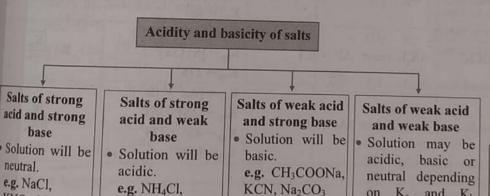
Acidic solutions	$[H^+] > 1.0 \times 10^{-7} M$	pH < 7.00
Basic solutions	$[H^{+}] < 1.0 \times 10^{-7} M$	pH > 7.00
Neutral solutions	$[H^+] = 1.0 \times 10^{-7} M$	pH = 7.00

Types of salts:

KNO<sub>3</sub>, Na<sub>2</sub>SO<sub>4</sub>

NH<sub>4</sub>NO<sub>3</sub>

(NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>

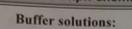


• If K<sub>a</sub> > K<sub>b</sub>, solution is acidic. e.g. NH<sub>4</sub>F neutral depending on Ka and Kb values of weak acid and weak base respectively. e.g. CH3COONH4, NH4CN,

(NH<sub>4</sub>)<sub>2</sub>CO<sub>3</sub>

 If K<sub>a</sub> < K<sub>b</sub>, solution is basic. e.g. NH<sub>4</sub>CN

 If K<sub>a</sub> = K<sub>b</sub>, solution is neutral. e.g. CH3COONH4



## **Buffer solutions**

A buffer solution resists drastic changes in pH upon the addition of a small amount of either an acid or a base.

### Types

#### Basic buffer solution Acidic buffer solution

- A solution containing a weak acid and its salt with a strong base.
  - e.g. CH3COOH and CH3COONa in water.
- Henderson-Hasselbalch equation is:

$$pH = pK_a + log_{10} \frac{[salt]}{[acid]}$$

- · A solution containing a weak base and its salt with a strong acid.
- e.g. NH<sub>4</sub>OH and NH<sub>4</sub>Cl in water. Henderson-Hasselbalch
  - equation is:  $pOH = pK_b + log_{10} \frac{[base]}{[base]}$

### **Properties**

- When a small amount of strong acid (or strong base) added to a buffer solution there is no significant change in the value of pH.
- The dilution of a buffer solution will not change its
- The pH of a buffer solution does not change with time.

#### Different expressions for solubility product:

Type of electrolyte	Example	Equation	K <sub>sp</sub> expression	Molar solu
AB (1:1 type salt)	AgCl	$AgCl \longrightarrow Ag^+ + Cl^-$	$K_{sp} = [Ag^{+}] [C\Gamma]$ $K_{sp} = S^{2}$	$S = \sqrt{K_s}$
AB <sub>2</sub> (1 : 2 type salt)	PbCl <sub>2</sub>	$PbCl_2 \Longrightarrow Pb^{2+} + 2Cl^{-}$	$K_{sp} = [Pb^{2+}] [Cl]^{2}$ $K_{sp} = [S] [2S]^{2}$ $K_{sp} = 4S^{3}$	$S = \sqrt[3]{\frac{K_1}{4}}$
A <sub>2</sub> B (2 : 1 type salt)	Ag <sub>2</sub> CrO <sub>4</sub>	$Ag_2CrO_4 \rightleftharpoons 2Ag^+ + CrO_4^{2-}$	$K_{sp} = [Ag^{+}]^{2} [CrO_{4}^{2-}]$ $K_{sp} = [2S]^{2} [S]$ $K_{sp} = 4S^{3}$	$S = \sqrt[3]{\frac{K_1}{4}}$
AB <sub>3</sub> (1:3 type salt)	AlCl <sub>3</sub>	$AlCl_3 \Longrightarrow Al^{3+} + 3Cl^{-}$	$K_{sp} = [AI^{3+}] [3CI^{-}]$ $K_{sp} = [S] [3S]^{3}$ $K_{sp} = 27S^{4}$	$S = \sqrt[4]{\frac{K_1}{2}}$
$A_2B_3$ (2 : 3 type salt)	As <sub>2</sub> S <sub>3</sub>	$As_2S_3 \iff 2As^{3+} + 3S^{2-}$	$K_{sp} = [As^{3+}]^2 [S^2]^3$ $K_{sp} = [2S]^2 [3S]^3$ $K_{sp} = 4S^2 \times 27S^3$ $K_{sp} = 108S^5$	$S = \sqrt[3]{\frac{K}{10}}$

## Condition for the formation of a precipitate:

Condition	Type of solution	THE PARTY STATES
Ionic product = K <sub>sp</sub>	Saturated solution	Result
Ionic product > K <sub>sp</sub>	Supersaturated solution	No precipitation
Ionic product < K <sub>sp</sub>	Unsaturated solution	Precipitation
-	solution	No precipitation

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- 1. Degree of dissociation ( $\alpha$ ):  $\alpha = \frac{\text{Number of moles dissociated}}{\text{Total number of moles}}$
- 2. Ostwald's dilution law:

$$\alpha \propto \frac{1}{\sqrt{C}}$$
 OR  $\alpha \propto \sqrt{V}$ 

where  $c = concentration in mol dm^{-3}$ V = volume in dm<sup>3</sup>

3. Acid dissociation constant (Ka):

For weak acid HA, 
$$K_a = \frac{[H^+][A^-]}{[HA]}$$

 $K_a$  =  $\alpha^{\,2}/V$  and  $K_a$  =  $\alpha^{\,2}\,c$ 

4. Base dissociation constant (K<sub>b</sub>):

For weak base BOH, 
$$K_b = \frac{[B^+][OH^-]}{[BOH]}$$
  
 $K_b = \alpha^2/V$  and  $K_b = \alpha^2 c$ 

- 5. Ionic product of water ( $K_w$ ):  $K_w = [H_3O^+][OH] = 1.0 \times 10^{-14}$
- 6. pH of solution:  $pH = -\log_{10}[H_3O^{+}]$
- 7. **pOH of solution:**  $pOH = -\log_{10}[OH]$
- 8. Relation between pH and pOH: pH + pOH = 14
- 9. Henderson-Hasselbalch equation:

$$pH = pK_a + log_{10} \frac{[salt]}{[acid]}$$

$$pK_a = -log_{10} K_a$$

Basic buffer:

$$pOH = pK_b + log_{10} \frac{[salt]}{[base]}$$

$$pK_b = -log_{10} K_b$$

10. Solubility product (K<sub>sp</sub>):

For salt 
$$B_x A_y$$
:  
 $K_{sp} = [B^{y+}]^x [A^{x-}]^y$ 

11. Molar solubility, S (mol/L):

$$S = \frac{Solubility in g/L}{Molar mass in g/mol}$$

12. Relation between K<sub>sp</sub> and S:

$$K_{sp} = x^x y^y S^{x+y}$$