



Topic 3.4 Theories of acids and bases. Ionic equilibria in electrolyte solutions. Buffer solutions.

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Outline

- **♦** Introduction
- Main part
- 1. Theories of acids and bases.
- 2. Ionic equilibria in electrolyte solutions.
- ♦ 3. Buffer solutions.
- Conclusion
- Literature

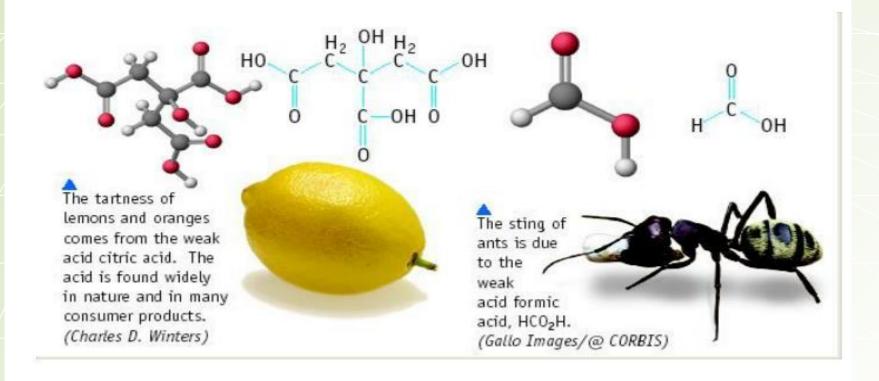




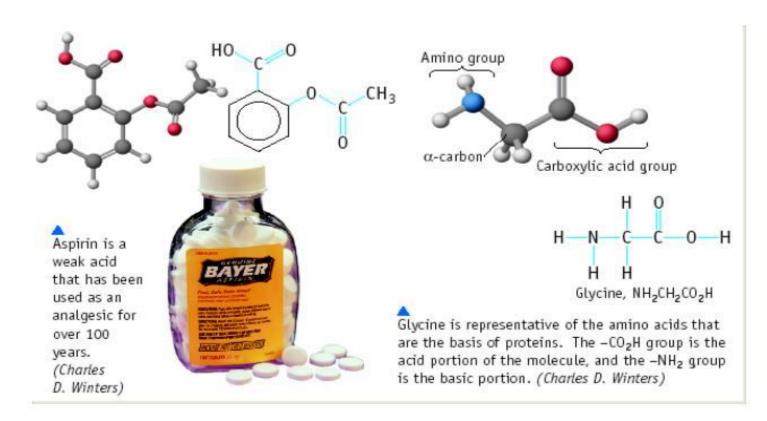
Interesting Background (not required)

- concepts acids and bases were loosely defined as substances that change some properties of water
- criteria that was often used was taste
 - substances were classified
 - · salty-tasting
 - · sour-tasting
 - · sweet-tasting
 - · bitter-tasting
- sour-tasting substances would give rise to the word 'acid', which is derived from the Greek word oxein, which mutated into the Latin verb acere, which means 'to make sour'

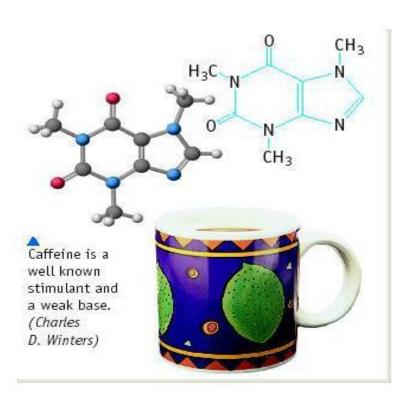


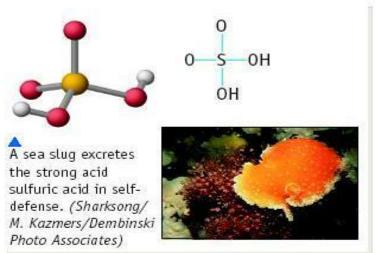






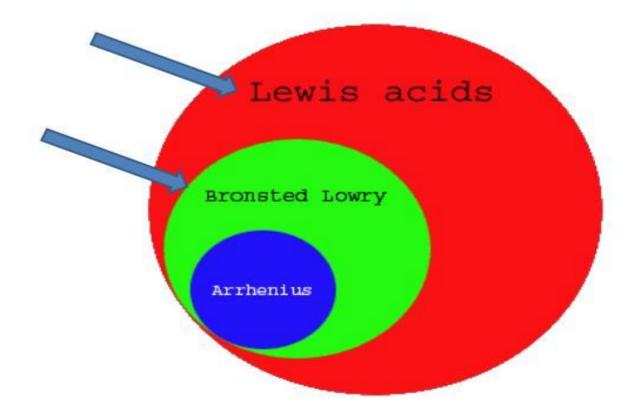








Three Acid-Base Definitions





Arrhenius definition





$$HNO_3(aq) \xrightarrow{H_2O} H^+(aq) + NO_3^-(aq)$$

with Arrhenius bases forming hydroxide ions (OH⁻) in solution



Each of these compounds increases the amount of H⁺ in water when dissolved in water.

$$HCI \xrightarrow{H_2O} H + CI$$

Notice that this acid produces twice the amount of H⁺ per mole of acid.

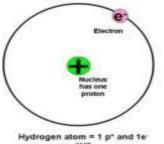
To identify Arrhenius acids look for compounds that dissociate or ionize in water forming H⁺

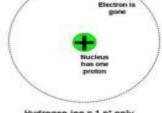


Brønsted – Lowry definition

- involves the transfer of a proton (H⁺)
 - a "proton" is really just a hydrogen atom that has lost its electron
 - -acids: molecule or ion that acts as proton (H+) donor
 - -bases: molecule or ion that acts as proton (H⁺) acceptor







Hydrogen ion = 1 p* only



Conjugate Pairs

- an acid-base reaction always involves (at least) two conjugate pairs that differ by an H⁺
- conjugate acids and conjugate bases are compounds formed when a H+ ion is gained and a H+ is lost
- a conjugate pair is:
 - an acid and its conjugate base
 - a base an its conjugate acid



Conjugate Pair

Conjugate Pair

HA(aq) +
$$H_2O(1)$$
 --> $H_3O^+(aq)$ + $A^-(aq)$

Acid Base Conjugate Conjugate

Acid Base

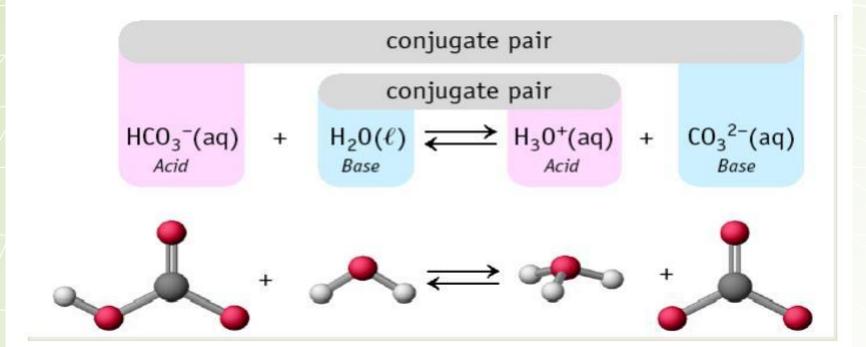
HF H_2O H_3O^+ F



acid base acid base

- HCl $(g) + H_2O(I) \rightleftharpoons H_3O^+(aq) + Cl^-(aq)$
 - -in the forward reaction,
 - HCl is the acid (will donate H⁺)
 - H₂O is the base (will accept H⁺)
 - -in the reverse reaction,
 - H₃O⁺ is the acid (will donate H⁺)
 - Cl⁻ is the base (will accept H⁺)

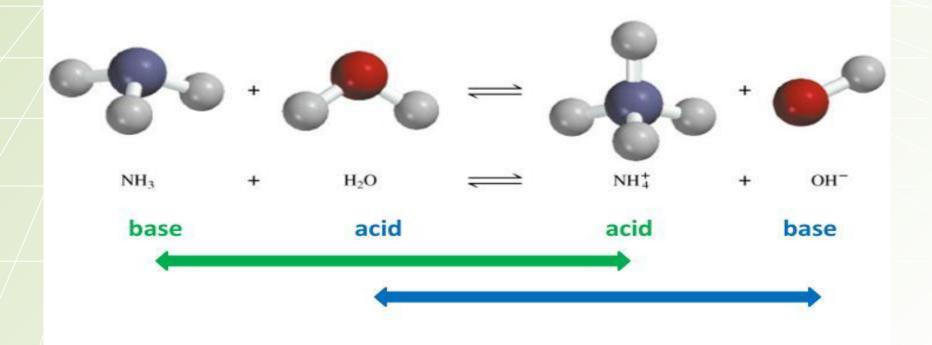








Brønsted-Lowry conjugate pairs





- Strong acid → Weak conjugate base
- Weak acid

 Strong conjugate base
- Weak base

 Strong conjugate acid

Acid	Strength	Base	Strength
H ₂ SO ₄	Very Strong	HSO ₄ -	Very Weak
HCI	Fairly Strong	Cl ⁻	Less Weak Fairly Strong
HNO ₃		NO ₃ -	
H₃O+		H ₂ O	
HSO ₄ -		SO ₄ ²⁻	
CH ₃ COOH	Weak	CH ₃ COO-	
H ₂ CO ₃	vveak	HCO ₃ -	
NH ₄ +		NH ₃	
HCO ₃ ·	Very Weak	CO ₃ ² ·	
H₂O ←		—— ОН-	



Water is amphiprotic/amphoteric (can act as acid or base)

Acid

HCl +

 H_2PO_4 +

 NH_4^+ +

Base

H₂O

H₂O

H₂O

Conjugate Acid Conjugate Base

→ H₃O+ + Cl

 \rightarrow H₃O⁺ + HPO₄²⁻

 \rightarrow H₃O⁺ + :NH₃

Base

:NH₃ +

PO₄3- +

Acid

H₂O

H₂O

Conjugate Acid Conjugate Base

 \rightarrow_{\leftarrow} NH₄⁺ + OH⁻

 \rightarrow_{\leftarrow} HPO₄²⁻ + OH⁻



Donating protons...

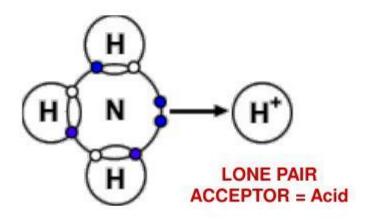
- monoprotic acids contain a single proton that can be donated (HCl, HNO₃, HNO₂, CH₃COOH)
- diprotic acids contain two protons that can be donated (H₂CO₃, H₂SO₄, H₂SO₃)
- triprotic acids contain three protons that can be donated (H₃PO₄)
- for a substance to be an acid, the hydrogen usually has to be attached to oxygen or a halogen
 - for example, in CH₃COOH, only the H on "OH" is able to be donated, the three hydrogens on carbon are non-acidic (do not write this C₂H₄O₂)

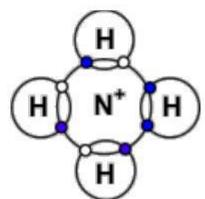


Lewis Acid and Base Definitions

- most general/encompassing definition
 - must have lone pairs (ligands) available to donate
- Lewis acids accept a pair of electrons to form a dative covalent bond
- Lewis bases donate a pair of electrons to form a dative covalent bond



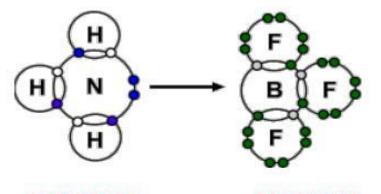


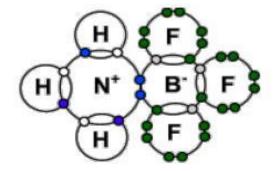


LONE PAIR
DONOR = Base









LONE PAIR DONOR = Base LONE PAIR ACCEPTOR = Acid

Ionic equilibria in electrolyte solutions.



Ionic Equilibrium in Solutions

The equilibrium established between the unionized molecules and the ions in the solution of weak electrolytes is called **ionic equilibrium**. For example, take acetic acid breaking up into acetate ions and hydrogen ions:

 $CH_3 COOH \rightarrow CH_3 COO^- + H^+$

Chemical substances that can conduct electricity in their aqueous state or in molten state are called **electrolytes**. In pure water or in an aqueous solution, the product of concentrations of hydrogen and hydroxyl ions is a constant at a given temperature. This is called **ionic product of water** and is conventionally designated by *Kw*.

2. Ionic equilibria in electrolyte solutions.



The idea of the ionic product of water can be understood by looking at the autoionization reaction of water that may be expressed as:

$$H_2 O + H_2 O = H_3 O^+ + OH^-$$

 $K_{W} = C_{H3 O+} C_{OH-}$

The value of K_w at 25°C is 1*10⁻¹⁴.

In the study of acid base equilibria in aqueous solutions, we're primarily interested in the hydrogen ion concentration of a solution. Solutions that we deal with are usually dilute, and the hydrogen ion concentrations are some negative power of 10.

2. Ionic equilibria in electrolyte solutions.



pH is a measure of hydrogen ion concentration. It's a measure of the acidity or alkalinity of a solution. Aqueous solutions at 25°C with a pH less than seven are acidic, while those with a pH greater than seven are basic or alkaline. pH can be calculated using the following formula.

$$pH = -log C_{H3 O+}$$

2. Ionic equilibria in electrolyte solutions.



Let's discuss how to calculate pH with an example.

What will be the concentration of C_{H30}^+ in a solution having pH = 5.6?

Solution: $pH = -log C_{H3 O+}$

- • $\log [H_3 O^+] = -pH$
- $-\log [H_3 O^+] = -5.6$
- •[$H_3 O^+$] = antilog (-5.6) = 2.512*10⁻⁶

Buffer solutions. Buffers



Definition

"Buffers are compounds or mixtures of compounds that by their presence in the solution resist changes in the pH upon the addition of small quantities of acid or alkali."



Necessity of a buffer system:

Sometimes it is necessary that a solution of a definite pH be prepared and stored. The preservation of such a solution is even more difficult than its preparation. If solution comes in contact with air, it will absorb CO₂ and becomes acidic. On the other hand, if solution is stored in a glass bottle, alkaline impurities from the glass may alter its pH. Due to these reasons, pharmaceutical solutions are buffered as the buffer solutions are capable of maintaining pH at some fairly constant value when even small amounts of acid or base are added.



Types of buffers:

Generally buffers are of two types;

- Acidic buffers
- Basic buffers



Acidic Buffers:

An acidic buffer is a combination of weak acid and its salt with a strong base.

i.e. Weak acid & salt with strong base (conjugate base).

- EXAMPLES:
- CH₃COOH / CH₃COONa
- H₂CO₃ / NaHCO₃
- H₃PO₄ / NaH₂PO₄
- HCOOH / HCOONa



Basic Buffers:

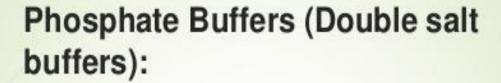
A basic buffer is a combination of weak base and its salt with a strong acid.

i.e. Weak base & salt with strong acid (conjugate acid).

EXAMPLES:

- NH₄OH / NH₄CI
- NH₃ / NH₄CI
- NH₃ / (NH₄)₂CO₃





Besides the two general types of buffers (i.e. acidic & basic), a third appears to exist. This is buffer system composed of two salts:

- Monobasic potassium phosphate (KH₂PO₄)
- Dibasic potassium phosphate (K₂HPO₄).



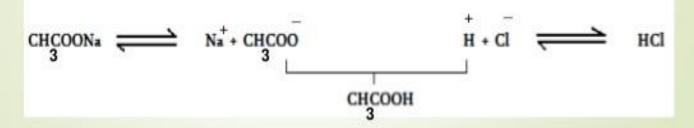
Buffer action

The resistance of a buffer solution to a change in pH is known as buffer action.



Mechanism of Action of acidic buffers:

- Consider a buffer system of CH₃COOH (Weak electrolyte) and CH₃COONa (Strong electrolyte). There will be a large concentration of Na+ ions, CH₃COONa ions, and undissociated CH₃COOH molecules.
- When an acid is added
- If a strong acid (Hcl) is added in CH₃COOH / CH₃COONa buffer, the changes that will occur may be represented as:





The hydrogen ions yielded by the Hcl are quickly removed as unionized acetic acid, and the hydrogen ion concentration is therefore only slightly affected (because acetic acid produced is very weak as compared to Hcl added).



When a base is added

If a strong base (NaoH) is added in CH₃COOH / CH₃COONa buffer, the changes that will occur may be represented as:

The hydroxyl ions yielded by the NaoH are therefore removed as water. The supply of hydrogen ions needed for this purpose being constantly provided by the dissociation of acetic acid.



Questions for self control



1. Whose definition of acids and bases emphasizes the role of protons?

- a. Brønsted and Lowry
- b. Arrhenius
- c. Lewis
- d. Faraday

2. An electron-pair acceptor is a

- a. Brønsted-Lowry base.
- b. Lewis base.
- c. Lewis acid.
- d. traditional acid.

3. Which statement about Arrhenius acids is FALSE?

- a. Their water solutions are called aqueous acids.
- b. They are molecular compounds with ionizable hydrogen atoms.
- c. Their pure aqueous solutions are electrolytes.
- d. They increase the concentration of hydroxide ions in aqueous solution.

Questions for self control



4.A buffer solution comprises which of the following?

- a) A weak acid in solution
- b) A strong acid in solution
- c) A weak base in solution
- d) A weak acid and its conjugate base in solution

5. Which of the following structures represents the conjugate acid of HPO₄²⁻?

- a) $H_{2}PO_{4}^{-1}$
- b) H₃PO₄
- c) H₄PO₄⁺
- d) PO₄³⁻

Question for self control



6.If NaOH is added in CH₃COOH/CH₃COONa buffer than hydroxyl ions yielded by NaOH removed as:

- A) Water
- B) Acetic acid
- C) Water and acetic acid
- 7.If HCl is added in CH₃COOH/CH₃COONa buffer than hydrogen ions yielded by HCl removed as:
- A) Water
- B) Acetic acid
- C) Water and acetic acid





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- 8. Эектронные ресурсы с www.bilimland.kz





Do you have any questions?

