# Properties of Acids and Bases According to Boyle

- In *1661* **Robert Boyle** summarized the properties of **acids** as follows:
- 1. Acids have a sour taste.
- 2. Acids are corrosive.
- 3. Acids change the color of certain vegetable dyes, such as litmus, from blue to red.
- 4. Acids lose their acidity when they are combined with alkalies.

# **Robert Boyle**



Born: 25 Jan 1627 in Lismore, County Waterford, Ireland

Died: 30 Dec 1691 in London, England

# Properties of Acids and Bases According to Boyle

• The name "acid" comes from the Latin *acidus*, which means "*sour*", and refers to the sharp odor and sour taste of many acids.

#### Examples:

- Vinegar tastes sour because it is a dilute solution of acetic acid in water.
- Lemon juice tastes sour because it contains citric acid.
- Milk turns sour when it spoils because lactic acid is formed.
- The unpleasant, sour odor of **rancid butter** stems from **butyric acid** that form when fat spoils.

Properties of Acids and Bases According to Boyle

- In 1661 Boyle summarized the properties of *alkalies* as follows:
- Alkalies feel slippery.
- *Alkalies* change the color of litmus from red to blue.
- *Alkalies* become less alkaline when they are combined with **acids**.

# Properties of Acids and Bases According to Boyle

 In essence, Boyle defined alkalies as substances that

consume, or neutralize, acids:

- Acids lose their characteristic sour taste and ability to dissolve metals when they are mixed with alkalies.
- Alkalies even reverse the change in color that occurs when litmus comes in contact with an acid.

Properties of Acids and Bases According to Boyle

• Eventually *alkalies* became known as **bases** because they serve as the "base" for making certain salts.

# Definition of Acids and Bases According to Arrhenius

 In 1884 Svante Arrhenius suggested that salts such as NaCl dissociate when they dissolve in water to give particles he called ions.

 $H_2O$ NaCl(s)  $\rightarrow$  Na<sup>+</sup>(aq) + Cl<sup>-</sup>(aq)

# Svante Arrhenius (1859-1927)



Swedish chemist who explained the electrical conductivity of ionic solutions by presuming that **compounds dissociated into oppositely charged ions** whose motions constituted a current.

This conclusion was supported by observing that the freezing point depression of ionic solids were integer multiples larger than their concentrations would indicate according to Raoult's Law.

# Svante Arrhenius (1859-1927)



He described his theory in his 1884 thesis, which passed the defense with the lowest passing grade. However, it won him the Nobel Prize in chemistry in 1903. He also discovered the Arrhenius Rate Law, which describes the rate at which chemical reactions occur.

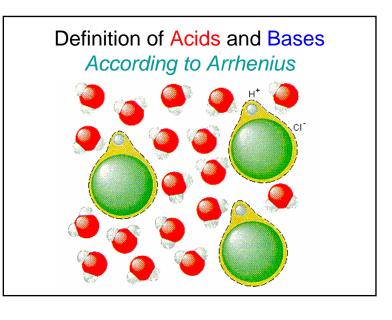
# Definition of Acids and Bases According to Arrhenius

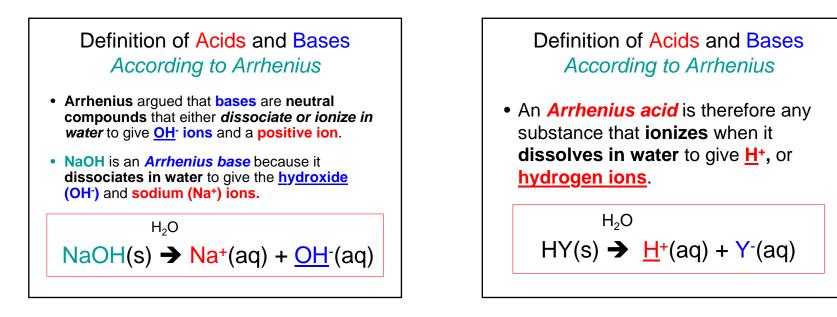
 Three years later Arrhenius extended this theory by suggesting that acids are neutral compounds that ionize when they dissolve in water to give H<sup>+</sup> ions and a corresponding negative ion.

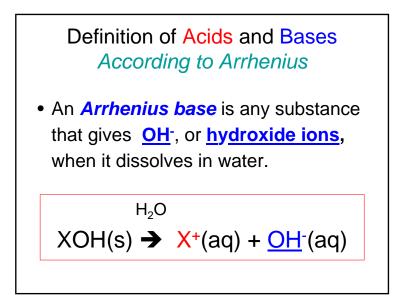


 According to his theory, hydrogen chloride is an acid because it ionizes when it dissolves in water to give hydrogen (H<sup>+</sup>) and chloride (CI<sup>-</sup>) ions as shown below:

H<sub>2</sub>O HCl(g)  $\rightarrow$  <u>H</u><sup>+</sup>(aq) + Cl<sup>-</sup>(aq)









 Arrhenius acids include compounds such as HCI, HCN, and H<sub>2</sub>SO<sub>4</sub> that ionize in water to give the <u>H</u><sup>+</sup> ion.

<sup>H₂O</sup> HY(s) **→** H⁺(aq) + <u>Y</u>⁻(aq)

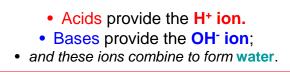


 Arrhenius bases include ionic compounds that contain the <u>OH</u><sup>-</sup> ion, such as NaOH, KOH, and Ca(OH)<sub>2</sub>.

 $XOH(s) \rightarrow X^{+}(aq) + OH^{-}(aq)$ 

# Definition of Acids and Bases According to Arrhenius

 This theory explains why all acids have similar properties: The characteristic properties of acids result from the presence of the H<sup>+</sup> ion generated when an acid dissolves in water. It also explains why acids neutralize bases and vice versa:



 $H^+(aq) + OH^-(aq) \rightarrow H_2O(I)$ 

# Definition of Acids and Bases According to Arrhenius

The Arrhenius theory has several disadvantages:

 It can be applied only to reactions that occur in water

because it defines **acids** and **bases** in terms of what happens when **compounds dissolve in water**.

# Definition of Acids and Bases According to Arrhenius

 The Arrhenius theory doesn't explain why some compounds, - in which hydrogen has an oxidation number of +1 (such as HCl), - dissolve in water to give acidic solutions,
whereas others (such as CH<sub>4</sub>) do not.

# Definition of Acids and Bases According to Arrhenius

• The *Arrhenius* theory doesn't explain why only the compounds that contain the **OH**<sup>-</sup> ion can be classified as Arrhenius bases.

## Definition of Acids and Bases According to Arrhenius

• The *Arrhenius* theory doesn't explain why only the compounds that contain the **OH**<sup>-</sup> ion can be classified as Arrhenius bases.

• The *Arrhenius* theory can't explain why other compounds (such as  $Na_2CO_3$ ) have the characteristic properties of bases.

# Definition of Acids and Bases According to **Brønsted**

The Brønsted, or Brønsted-Lowry, model is based on a simple assumption: Definition of Acids and Bases According to **Brønsted** 

The Brønsted, or Brønsted-Lowry, model is based on a simple assumption:

Acids donate H<sup>+</sup> ions to other ions or molecules, which act as a base.

# Some Remarks on the Concept of Acids and Bases

J. N. Brønsted Recueil des Travaux Chimiques des Pays-Bas (1923), Volume 42, Pages 718-728

We have held steadily to the idea that compound **A** is an acid if it is partly or completely broken down in solution according to the scheme

 $A \rightarrow B + H^+$  (1) However, there have been attempts from various sides to modify our concept of bases. P. Pfeiffer, has presented the view that bases form salts by addition of acids, which in terms of the theory of electrolytic dissociation must lead to the idea of

• a definition of bases as substances that can add hydrogen ions.

http://dbhs.wvusd.k12.ca.us/Chem-History/Bronsted-Article.html (May 14, 2003)

# Some Remarks on the Concept of Acids and Bases

J. N. Brønsted Recueil des Travaux Chimiques des Pays-Bas (1923), Volume 42, Pages 718-728

The formal advantages of the definition of a base as a substance that can bind a hydrogen ion are immediately apparent. In the scheme: A ---> B + H<sup>+</sup>, in which the concept of an acid (A) is established, the concept of a base (B) is equally defined and thus the reciprocal connection of acidic and basic properties is shown in the clearest and simplest way.

http://dbhs.wvusd.k12.ca.us/Chem-History/Bronsted-Article.html (May 14, 2003)

# Some Remarks on the Concept of Acids and Bases

J. N. Brønsted Recueil des Travaux Chimiques des Pays-Bas (1923), Volume 42, Pages 718-728 The advantages of scheme  $A \xrightarrow{} B + H^+$  (1) as a basis for an equal definition of bases and acids are not merely of a formal nature. For example, if we express the basic properties of ammonia in one case by the equation  $NH_4^+ <==> NH_3 + H^+$  (2) and in another by  $NH_3 + H_2O <==> NH_4^+ + OH^-$  (3)

we can derive identical equilibrium conditions.

http://dbhs.wvusd.k12.ca.us/Chem-History/Bronsted-Article.html (May 14, 2003)

# Some Remarks on the Concept of Acids and Bases

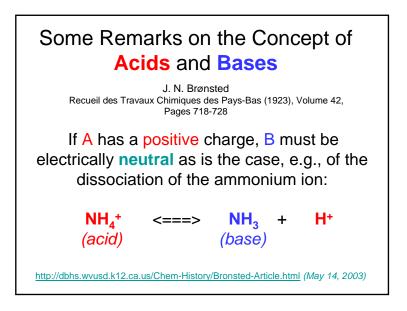
J. N. Brønsted Recueil des Travaux Chimiques des Pays-Bas (1923), Volume 42, Pages 718-728

It follows from the form of the expression that corresponding acids and bases do not have to have the same state of electrical charge. If A is electrically neutral, as an ordinary acid molecule is, B has a negative charge, e.g.

> CH<sub>3</sub>COOH <===> CH<sub>3</sub>COO<sup>-</sup> + H<sup>+</sup> (acid) (base)

where the acetate ion thus functions as a base.

http://dbhs.wvusd.k12.ca.us/Chem-History/Bronsted-Article.html (May 14, 2003)



		the Concept of Bases
	Table	I
Electrical charge	Naming of acid or base	Examples
Positive	cation acid	NH4 <sup>+</sup>
Positive	cation base	$NH_2 \cdot C_6H_4 \cdot C_6H_4 \cdot NH_3^+$
Neutral	neutral acid	CH <sub>3</sub> COOH
Neutral	neutral base	NH3
Negative	anion acid	COO · COOH [Note added: there is a bar above the second O from the left as in the graphic above]
Negative	anion base	CH <sub>3</sub> COO <sup>-</sup>

# Some Remarks on the Concept of Acids and Bases

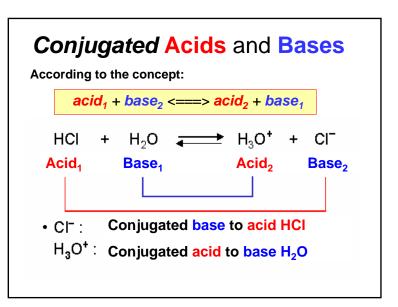
J. N. Brønsted Recueil des Travaux Chimiques des Pays-Bas (1923), Volume 42, Pages 718-728

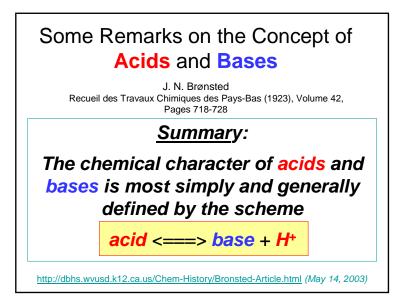
The equilibrium formulated in **scheme (1)** between hydrogen ion and the corresponding acid and base can be called **a simple acid-base equilibrium**.

By **mixing two simple systems**, a **double acidbase system** and an acid-base equilibrium result that can always be formulated as follows:

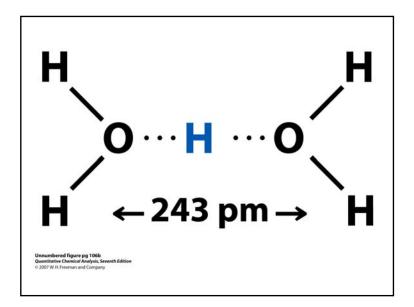
 $acid_1 + base_2 <==> acid_2 + base_1$  (4)

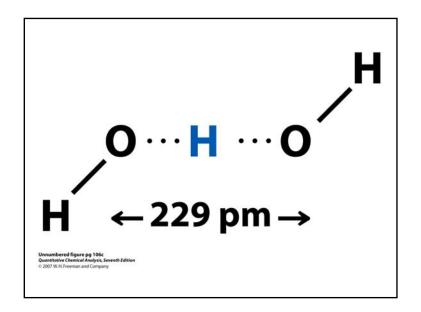
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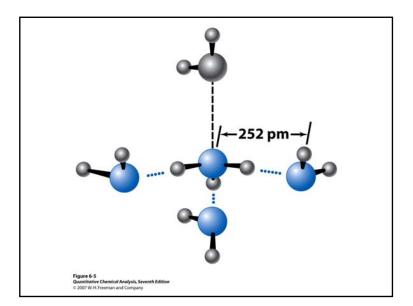


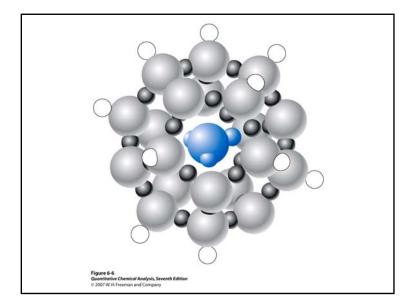


Definition of Acids and Bases According to Brønsted The dissociation of water, for example, involves the transfer of an  $H^+$  ion from one water molecule to another to form  $H_3O^+$  and  $OH^+$  ions.







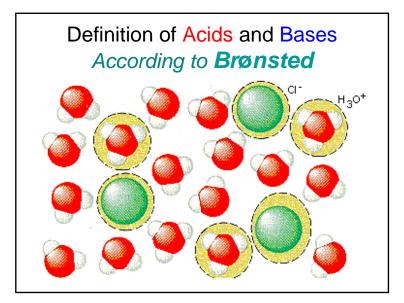


# Definition of Acids and Bases According to **Brønsted**

 According to the *Brønsted* model and definition of acids and bases, HCI doesn't dissociate in water to form H<sup>+</sup> and Cl<sup>-</sup> ions.

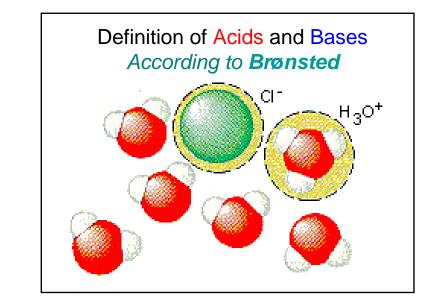
# Definition of Acids and Bases According to **Brønsted**

- According to the *Brønsted* model and definition of acids and bases, HCI doesn't dissociate in water to form H<sup>+</sup> and Cl<sup>-</sup> ions.
- Instead, an H<sup>+</sup> ion is transferred from HCl to a water molecule to form H<sub>3</sub>O<sup>+</sup> and Cl<sup>-</sup> ions, as shown in the following figure.



# The Role of H<sup>+</sup> and OH<sup>-</sup> lons in Aqueous Solutions

 Because oxygen (EN = 3.44) is much more electronegative than hydrogen (EN = 2.20), the electrons in the HO-bonds in water aren't shared equally by hydrogen and oxygen.



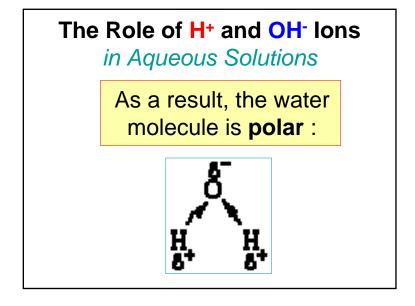
# The Role of H<sup>+</sup> and OH<sup>-</sup> lons in Aqueous Solutions

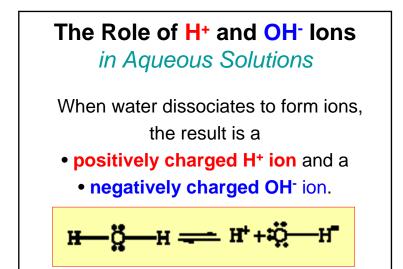
 Therefore, the electrons are drawn toward the oxygen atom in the center of the molecule and away from the hydrogen atoms on either end.

# The Role of H<sup>+</sup> and OH<sup>-</sup> lons in Aqueous Solutions

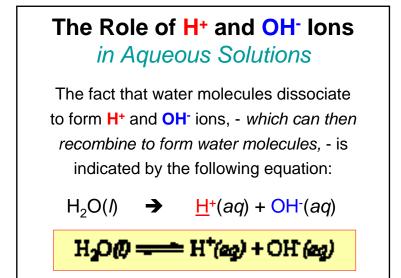
The **oxygen** atom carries a partial **negative** charge (-), and the **hydrogen** atoms carry a partial **positive** charge (+).







# The Role of H<sup>+</sup> and OH<sup>-</sup> lons in Aqueous Solutions The opposite can also occur: H<sup>+</sup> ions can combine with OH<sup>-</sup> ions to form neutral water molecules. $H^+ + OH^- \rightarrow H_2O$



# The Role of H<sup>+</sup> and OH<sup>-</sup> lons in Aqueous Solutions

To what extent does H<sub>2</sub>O dissociate to form ions?

At 25°C, the density of water is 0.9971 g/cm<sup>3</sup>, or 0.9971 g/mL.

 $\frac{0.9971 \text{ g } H_2O}{1 \text{ mL}} \times \frac{1000 \text{ mL}}{1 \text{ L}} \times \frac{1 \text{ mol } H_2O}{18.015 \text{ g } H_2O} = 55.35 \text{ mol } H_2O/L$ 

The concentration of  $H_2O$  is 55.35 molar.

# The Role of H<sup>+</sup> and OH<sup>-</sup> lons

in Aqueous Solutions

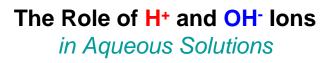
To what extent does H<sub>2</sub>O dissociate to form ions?

# The Role of H<sup>+</sup> and OH<sup>-</sup> lons

### in Aqueous Solutions

### To what extent does H<sub>2</sub>O dissociate to form ions?

The **concentration** of the H<sup>+</sup> and OH<sup>-</sup> **ions** formed by the dissociation of neutral H<sub>2</sub>O molecules at this temperature is only **1.0 x 10<sup>-7</sup> mol/L**.



To what extent does H<sub>2</sub>O dissociate to form ions?

The **concentration** of the  $H^+$  and  $OH^-$  **ions** formed by the dissociation of neutral  $H_2O$  molecules at this temperature is only **1.0 x 10<sup>-7</sup> mol/L**.

 $\frac{1.0 \times 10^{-7} M H^{+}}{55.35 M H_2 O} = 1.8 \times 10^{-9}$ 

# The Role of H<sup>+</sup> and OH<sup>-</sup> lons

in Aqueous Solutions

# To what extent does H<sub>2</sub>O dissociate to form ions?

At 25°C only about **2 parts per billion (ppb)** of the  $H_2O$ molecules **dissociate** into ions.

# The Role of H<sup>+</sup> and OH<sup>-</sup> lons in Aqueous Solutions

#### To what extent does H<sub>2</sub>O dissociate to form ions?

The **concentration** of the H<sup>+</sup> and OH<sup>-</sup> **ions** formed by the dissociation of neutral H<sub>2</sub>O molecules at this temperature is only **1.0 x 10<sup>-7</sup> mol/L**.

$$\frac{1.0 \times 10^{-7} M H^+}{55.35 MH_2O} = 1.8 \times 10^{-9}$$

Accordingly, the ratio of the concentration of the H<sup>+</sup> (or OH<sup>-</sup>) ion to that of the neutral H<sub>2</sub>O molecules is  $1.8 \times 10^{-9}$ .

## Definition of Acids and Bases The Operational Definition

The fact that water dissociates to form H<sup>+</sup> and OH<sup>-</sup> ions in a reversible reaction is the basis for an operational definition of acids and bases that is more powerful than the definitions proposed by Arrhenius.

# Definition of Acids and Bases The Operational Definition

In an operational sense:

- an <u>acid</u> is any substance that increases the concentration of the H<sup>+</sup> ion when it dissolves in water.
- a <u>base</u> is any substance that increases the concentration of the OH<sup>-</sup> ion when it dissolves in water.

# Definition of Acids and Bases The Operational Definition

These definitions tie the theory of **acids** and **bases** to a **simple laboratory test** for acids and bases:

To decide whether a compound is an acid or a base, we dissolve it in water and test the solution to see whether the H<sup>+</sup> or OH<sup>-</sup> ion concentration has increased.

# pH Meter Measures Acidity

 The U.S. Geological Survey analyzes hundreds of thousands of water samples every year. Many measurements are made right at the field site, and many more are made on water samples back at the lab. The pH is an important water measurement which is often measured both at the sampling site and in the lab. There are large and small models of pH meters. Portable models are available to take out in the field and larger models, such as this one, are used in the lab.



# pH Meter Measures Acidity

#### So, how does this contraption work?

• The water sample is placed in the little cup and the glass probe at the end of the retractable arm is placed in the water. The back of the probe is connected to the main box by electrical wires, and at the tip of the probe there is a thin glass bulb. Inside the probe there are two electrodes that measure voltage. One electrode is contained in a liquid that has a fixed acidity, or pH. The other electrode responds to the acidity of the water sample.

# pH Meter Measures Acidity

- In other words, the voltage of the second electrode responds to the amount of free hydrogen ions (the pH) in the sample.
- A voltmeter in the probe measures the difference between the voltages of the two electrodes.
- The meter then translates the voltage difference into pH and displays it on the little screen on the main box.

# pH Meter Measures Acidity

 Before taking a pH measurement, the meter must be "calibrated." The probe is immersed in a solution that has a known pH, such as pure water with a neutral pH of 7.0. The knobs on the box are used to adjust the displayed pH value to the known pH of the solution, thus calibrating the meter.