

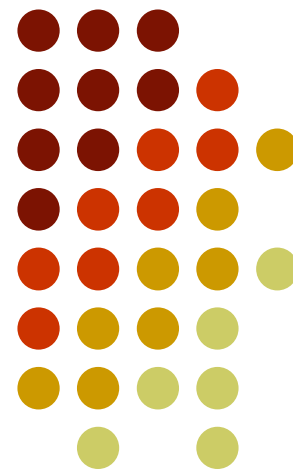
# LECTURE 3 SOLUTIONS. CONCENTRATIONS of SOLUTIONS



Senior teacher

PhD in physical chemistry

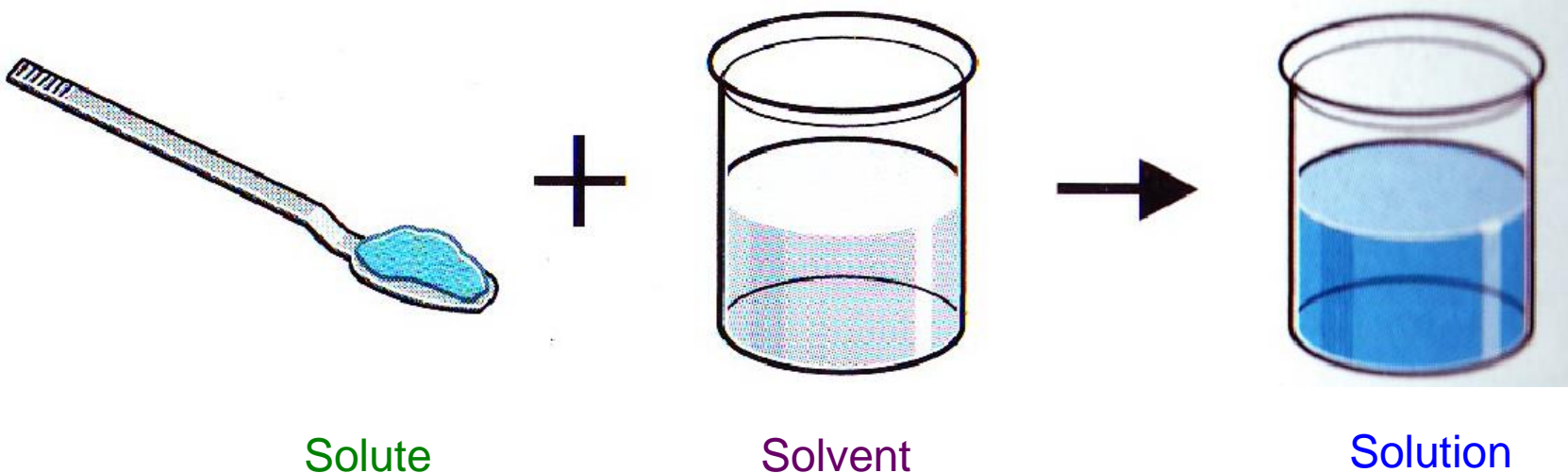
**Yanovska Anna Olexandrivna**





# Pure?!

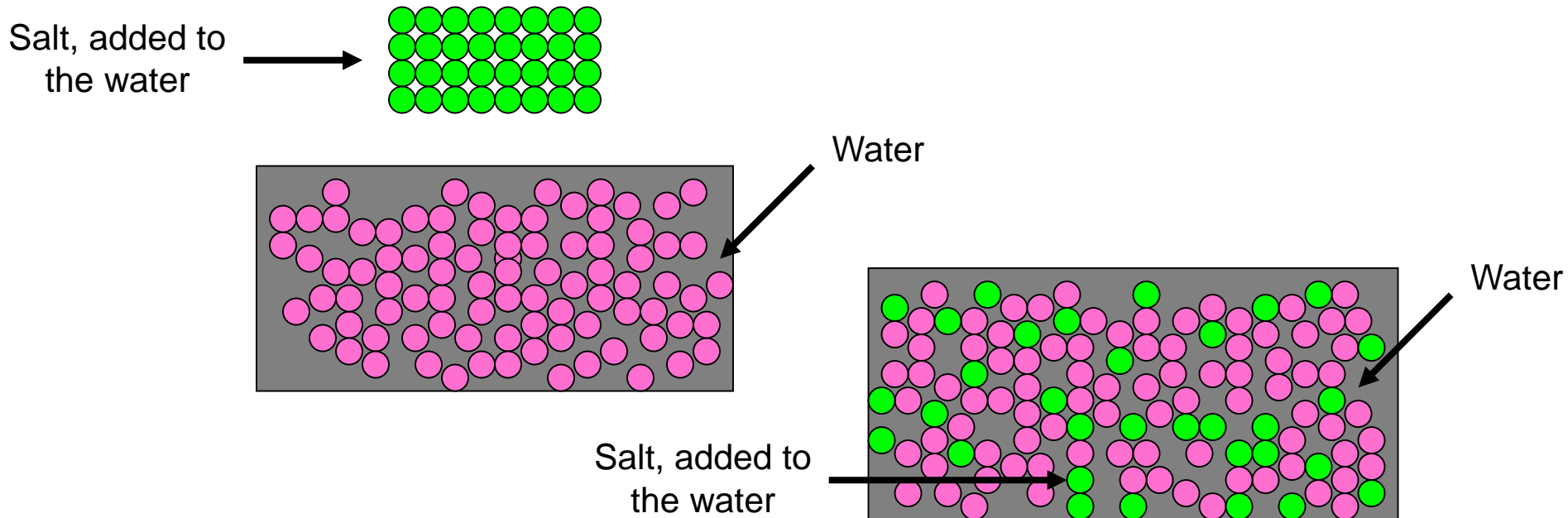
- Even though the water still looks ‘pure’, this term is wrong. Pure would mean there were only water particles, however we now have water particles + sugar particles!
- The sugar has not disappeared - instead it has dissolved in the water we would call the water + sugar a **solution**
- A solid dissolved in a liquid makes a **solution**
- In a solution the liquid is called the **solvent**, and the solid is called the **solute**



# The main terms of the topic



- **Solutions** are homogeneous systems consisting of two or more components and the products of their interaction.
- Compulsory components of the solution are the ***solvent*** and the ***solute***.
- The ***solvent*** is the solution component present in greatest quantity or the component that determines the state of matter in which a solution exist.

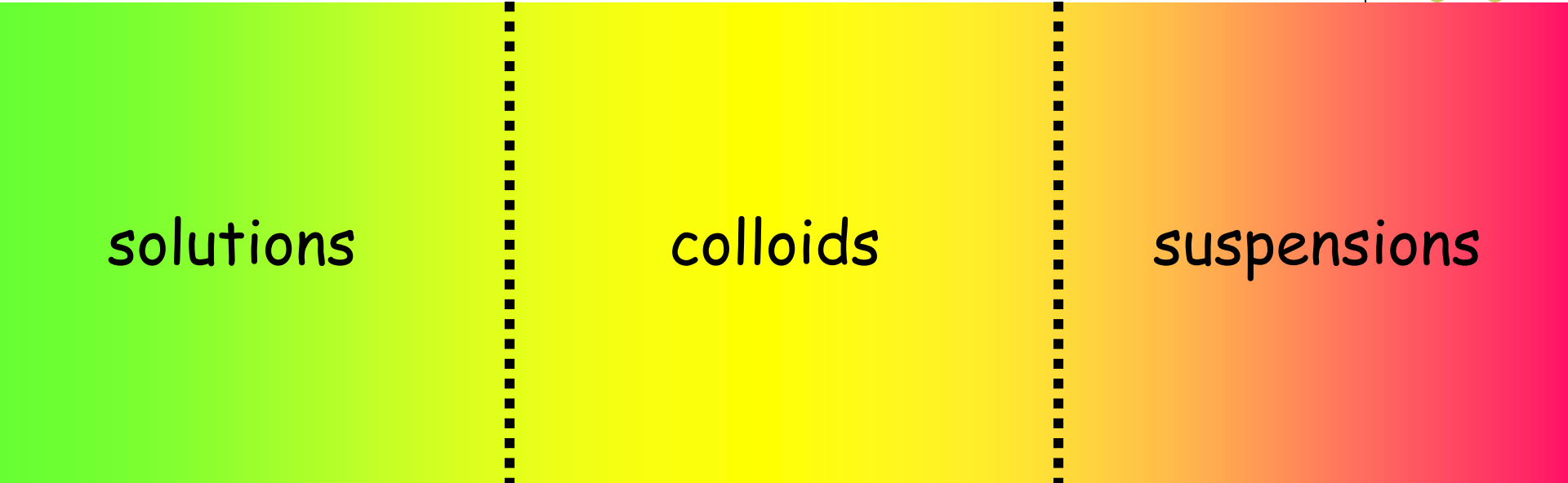




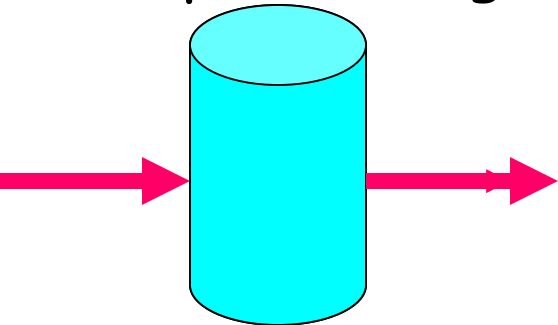
# Comparison of Solutions, Colloids, and Suspensions

< 1 nm

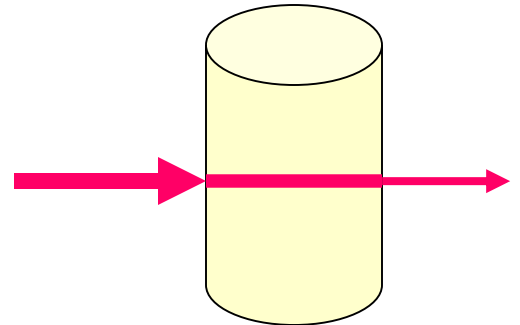
> 100 nm



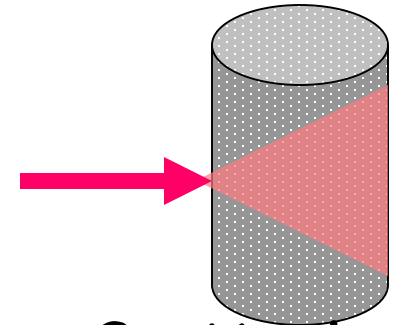
## Absorption of light



Passage of light



Scattering in beam



Scattering in all directions

# Solutions: homogeneous mixtures



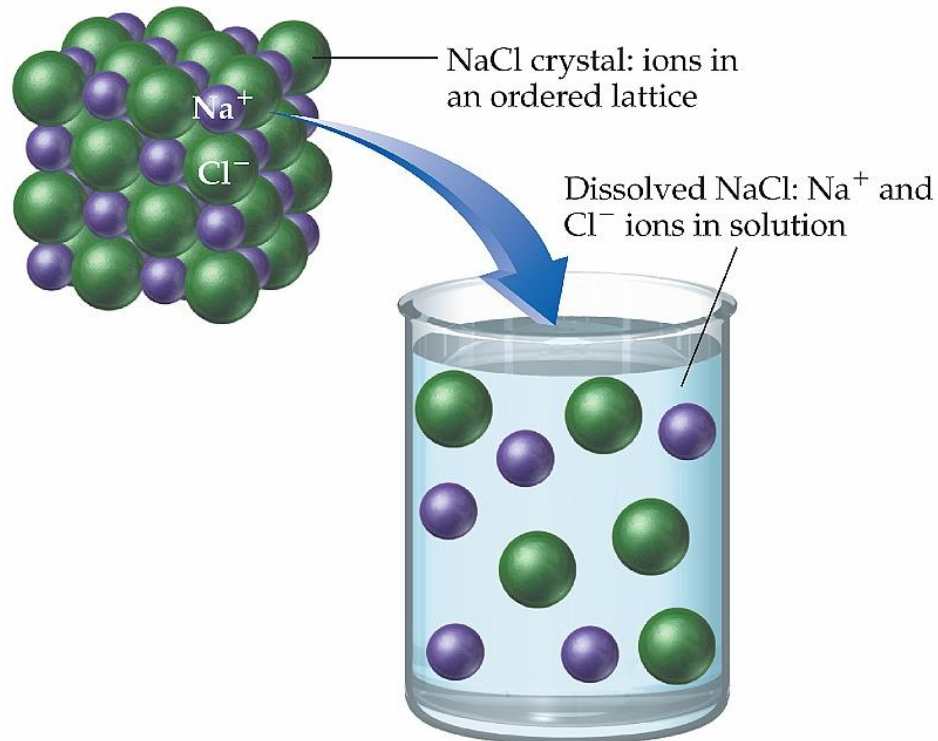
- Absence of settling
- Two components (at the least)-
  - **Solute** – the substance being dissolved (the dissolved particles)
  - **Solvent** – a substance which dissolves another substance (the dissolving medium)

A solution in which water is the solvent is called an **aqueous solution**.

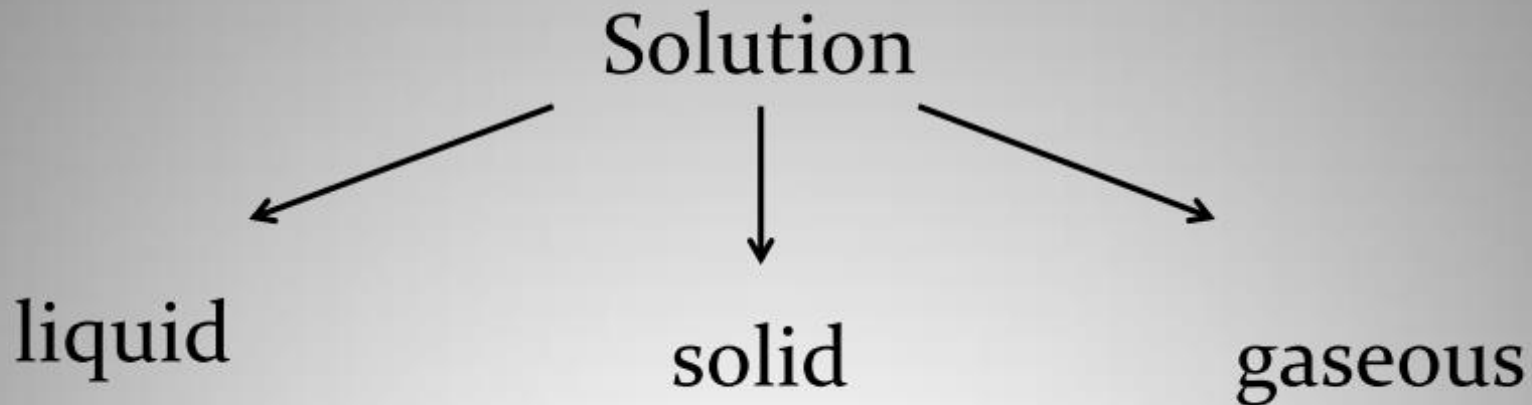
**Solution:** a homogeneous mixture of two or more substances

**Solution = Solute + Solvent**

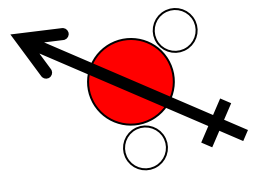
A solution containing a relatively large quantity of solute is said to be **concentrated**.  
If the quantity of solute is small,  
the solution is **dilute**.



# Types of solution



Gas	Gas	Air (oxygen in nitrogen)
Gas	Liquid	Soda water (carbon dioxide in water)
Solid	Liquid	Ocean water (salt in water)

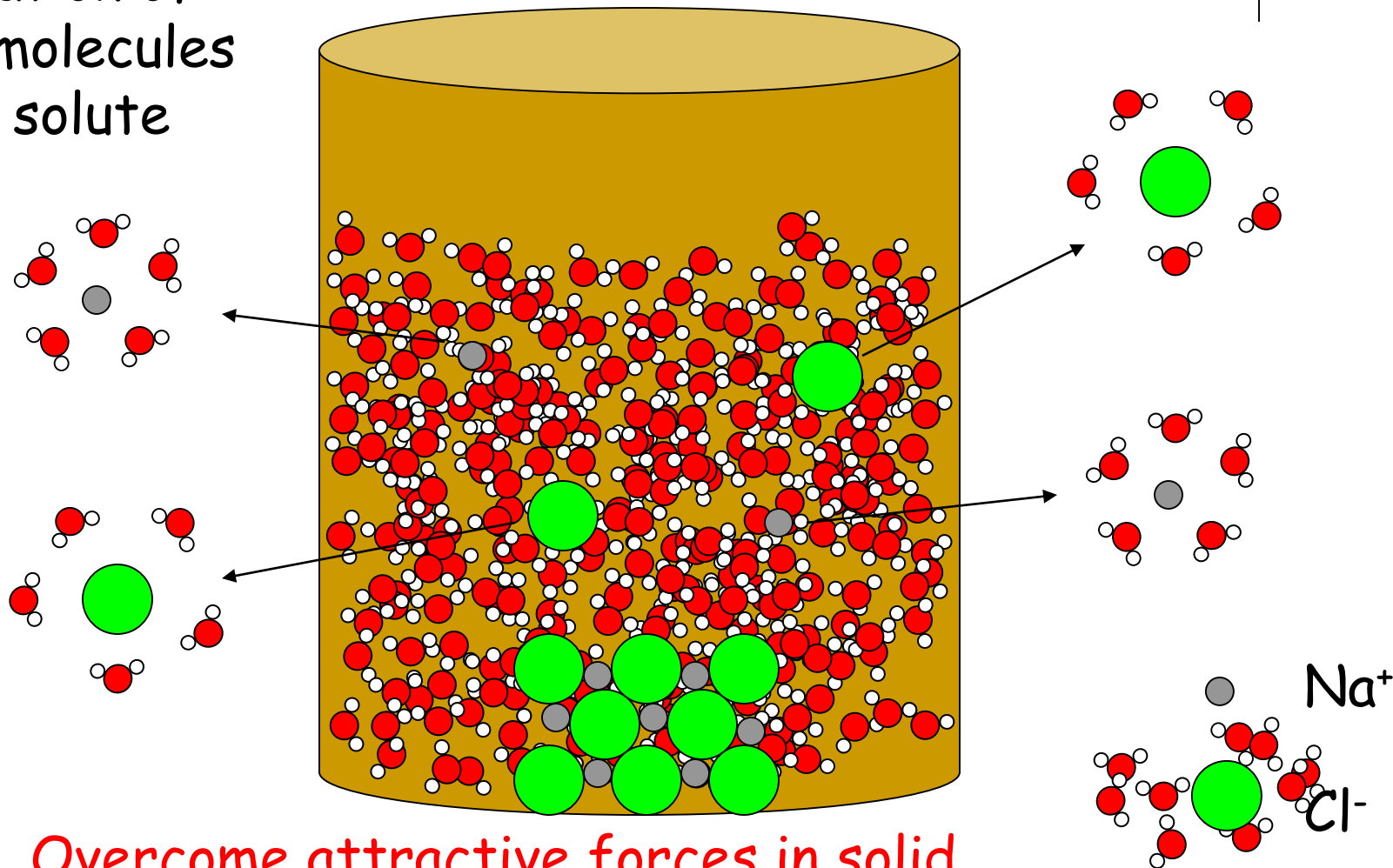


# Dissolving process in water



## 2. Hydration of solute

Orientation of water molecules around solute







In a **liquid solution** the **solvent** is a **liquid** substance.

For examples: gasoline is mixture of a number of liquid hydrocarbons.

Seawater is an aqueous solution of sodium chloride and other ionic solids.

Carbonated water is an aqueous solution of  $\text{CO}_2$ .

All gaseous mixtures are solutions.

The best known example of a gaseous solution is air, which consists of  $N_2$ ,  $O_2$ ,  $CO_2$  and other gases.

In a solid solution the solvent is a solid substance.

The ability to form solid solution is particularly common among metals, and such solid solutions are called **alloys**. For example: an alloy of nickel and copper, an alloy of gold and silver.



# Dissolving of a substances in solvent.

## Solubility

Dissolving of substances in solvent are:

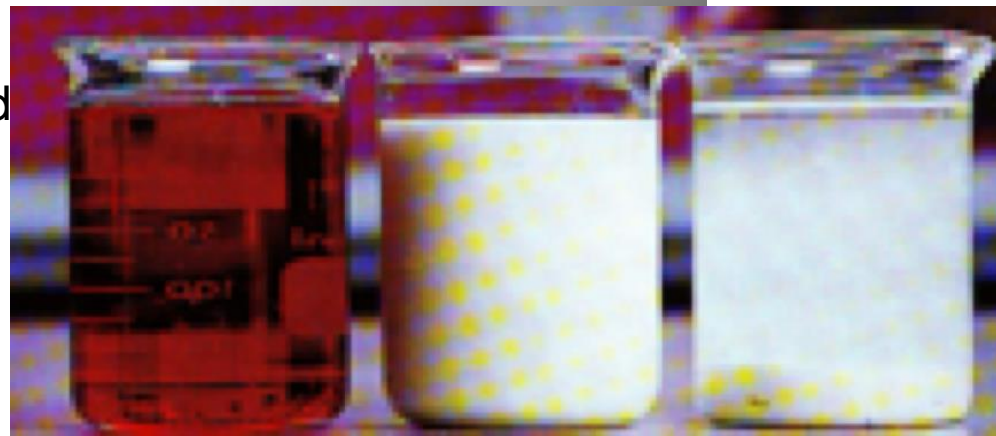
1. The destruction of crystalline lattice.
2. The interaction of the solvent with the particles of the solute.
3. The uniform distribution of one substance in the whole volume of another substance.



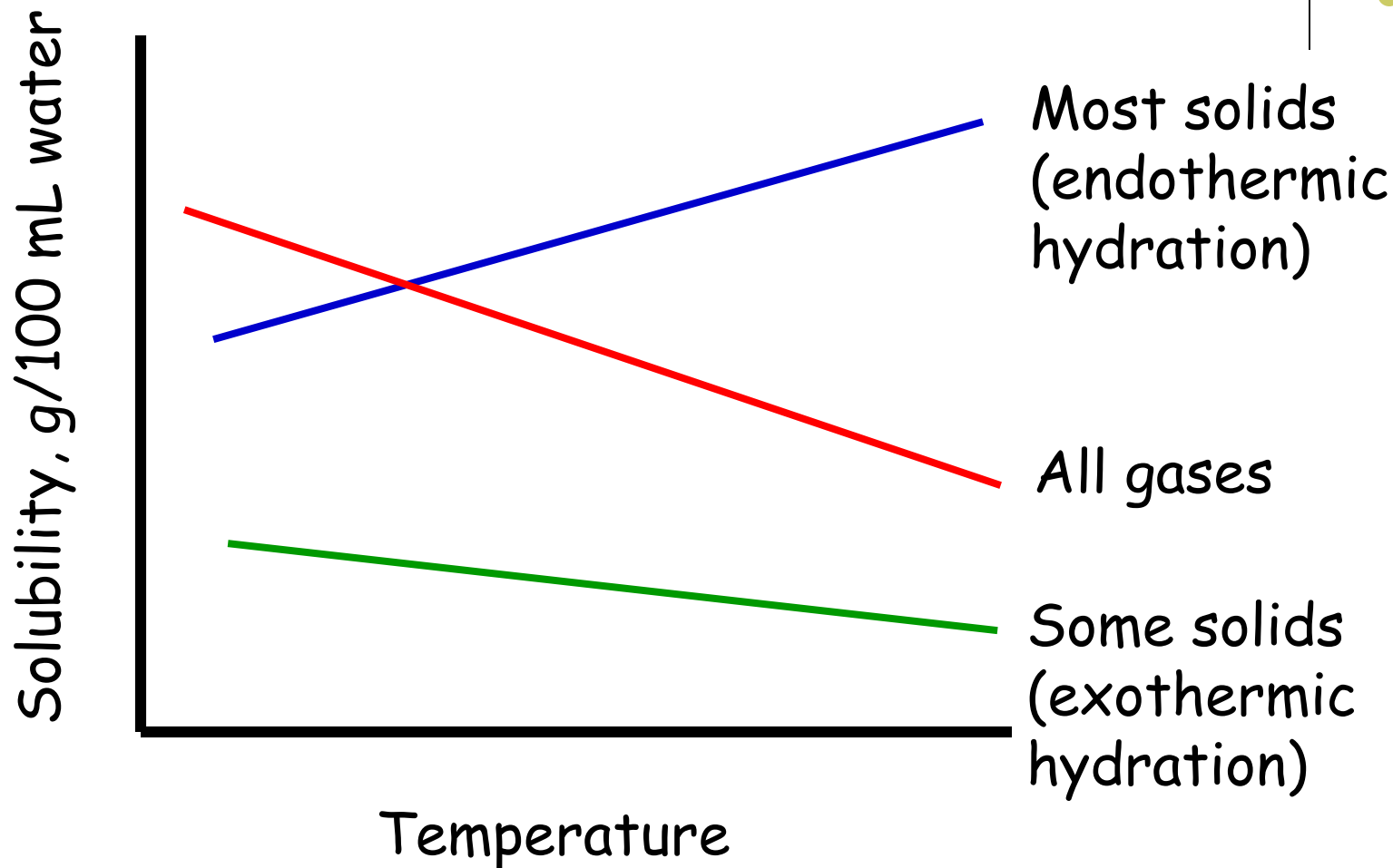
**Dissolve** - mixing of a substance in a liquid

**Soluble** - a substance which can dissolve  
(mix in a liquid)

**Insoluble** - a substance which cannot  
dissolve (mix in a liquid)



# Solubility of Solutes in Water



**Solubility** is the ability of the substance to uniformly distribute in the whole volume of another substance.

$$\frac{\text{g of solute}}{100 \text{ g water}}$$



**Solubility (s)** is the maximum amount of solute that dissolves in a specific amount of solvent.

- expressed as grams of solute in 100 grams of solvent water.

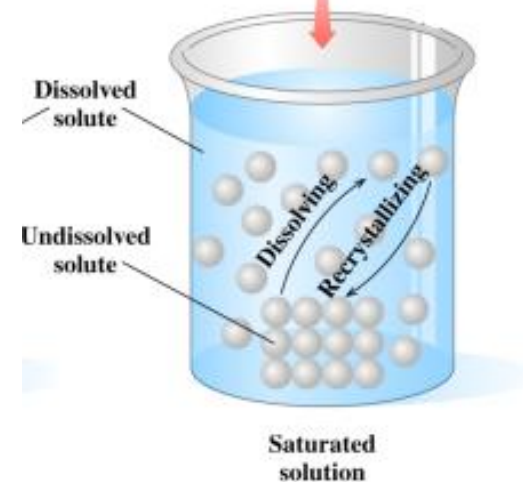
Solubility depends on the nature of substances, temperature and pressure. For example, solubility of solid substances increases and gases decreases with the raising of temperature. Solubility of gases increases with raising pressure.

A solution in which under the certain temperature the solute cannot be dissolved any more is called a **saturated** solution.



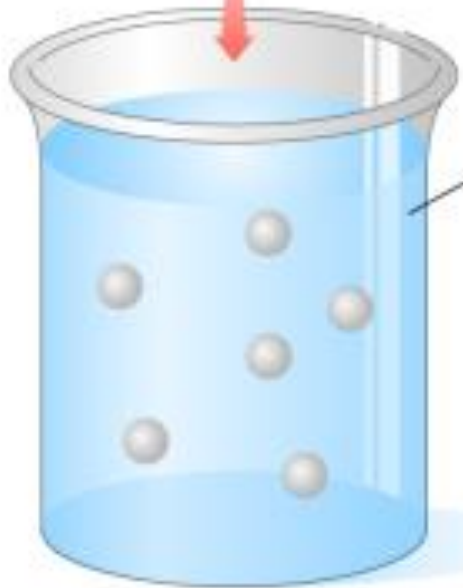
## Saturated solutions

- contain the maximum amount of solute that can dissolve.
- have undissolved solute at the bottom of the container.





A solution in which under the certain temperature more solute can be dissolved is called an **unsaturated** solution.



Dissolved  
solute

**Unsaturated  
solution**

contain less than the maximum amount of solute.

- can dissolve more solute.

# Supersaturated Solutions



An unstable solution that contains an amount of solute greater than the solute solubility.

- Also has undissolved solute at the bottom of the container.



(a) A seed crystal of  $\text{NaC}_2\text{H}_3\text{O}_2$  being added to the supersaturated solution.



(b) Excess  $\text{NaC}_2\text{H}_3\text{O}_2$  crystallizes from the solution.



(c) The solution arrives at saturation.





## Learning Check

At 40°C, the solubility of KBr is 80 g/100 g H<sub>2</sub>O. Identify the following solutions as either 1) saturated or (2) unsaturated. Explain.

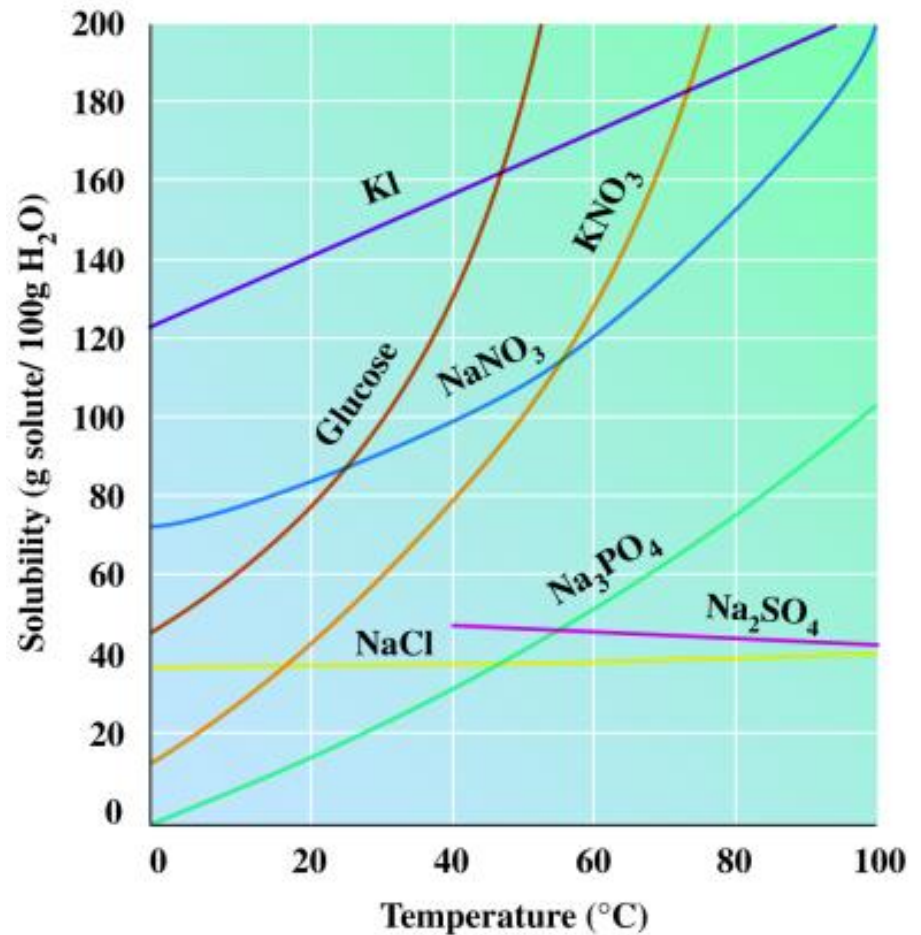
- A. 60 g KBr added to 100 g of water at 40°C.
- B. 200 g KBr added to 200 g of water at 40°C.
- C. 25 g KBr added to 50 g of water at 40°C.



# Effect of Temperature on Solubility

## Solubility

- Depends on temperature.
- For most solids increases with increase of temperature.
- For gases decreases as temperature increases.



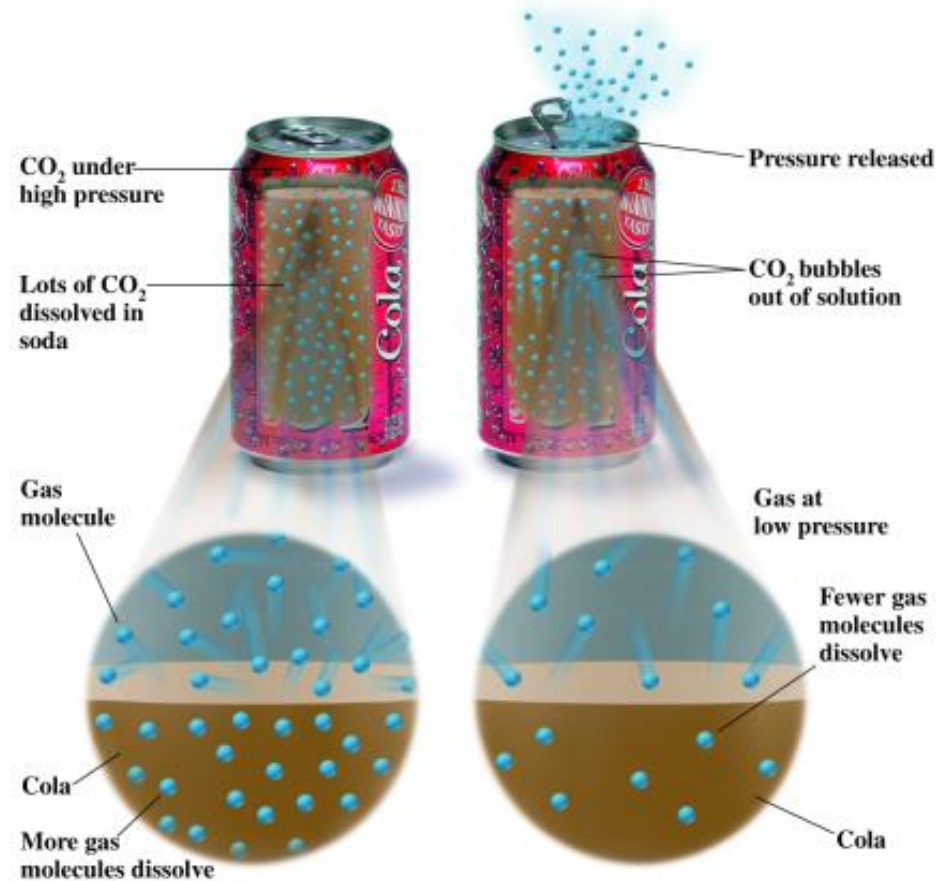
# Solubility and Pressure



## Henry's Law

states

- the solubility of a gas in a liquid is directly related to the pressure of that gas above the liquid.
- at higher pressures, more gas molecules dissolve in the liquid.

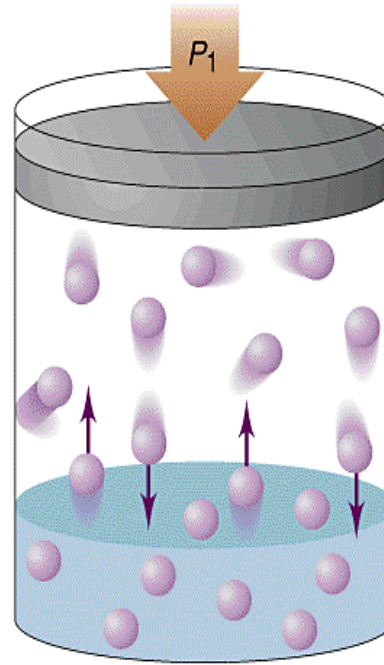


# Pressure affect on gases

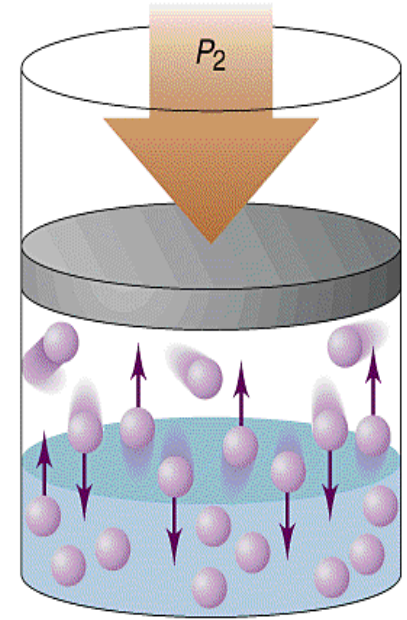


## Pressure:

- affects the solubility of gaseous solutes. The solubility of a gas in any solvent increases as the pressure above the solution increases



A



B

# 3 common ways to increase the collisions between solute and solvent

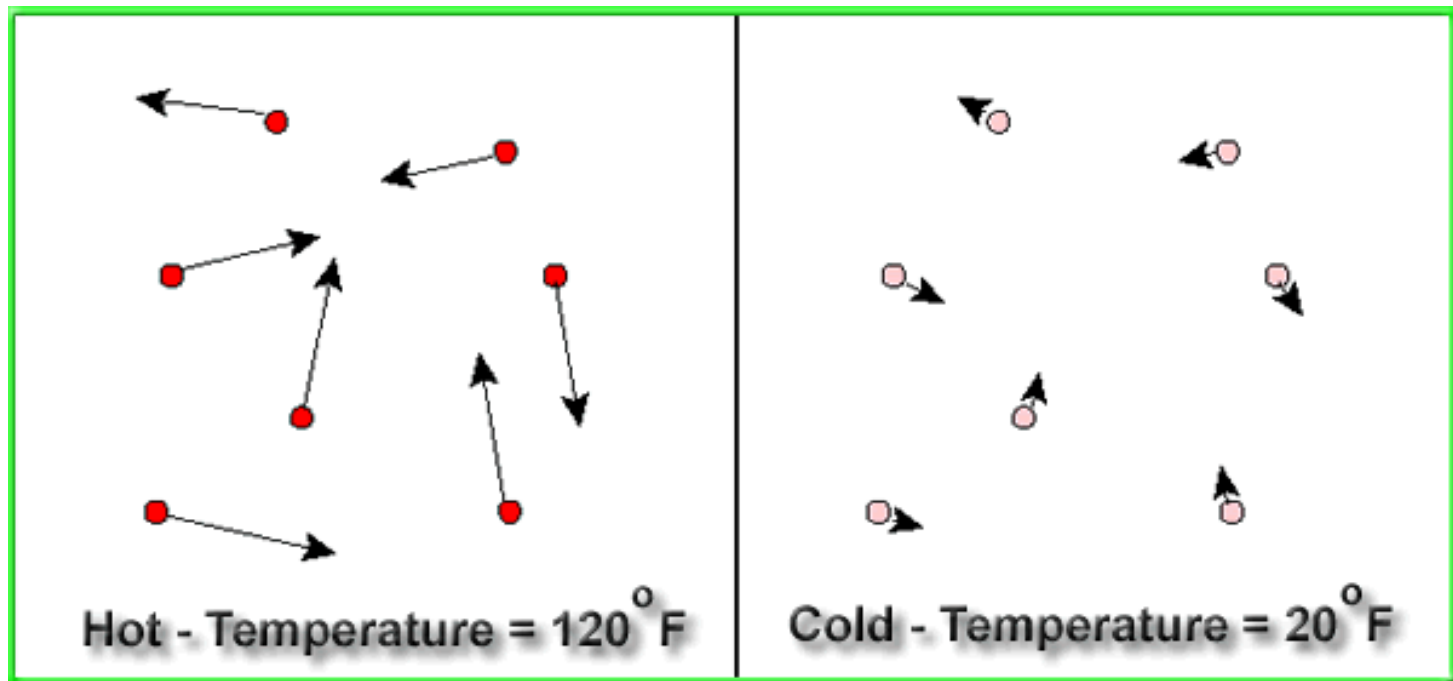


Temperature

Amount of Surface Area Exposed

Agitation

- Temperature: raising the temperature increases the kinetic energy of the particles, resulting in more frequent and forceful collisions.



# Amount of Surface Area Exposed



- Amount of Surface Area Exposed: breaking the solute into smaller pieces increases its surface area. A greater surface area allows more collisions to occur and therefore, faster

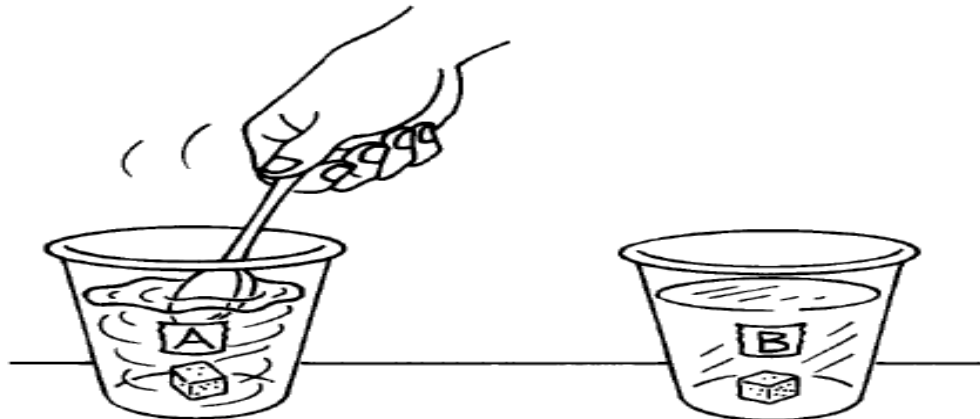


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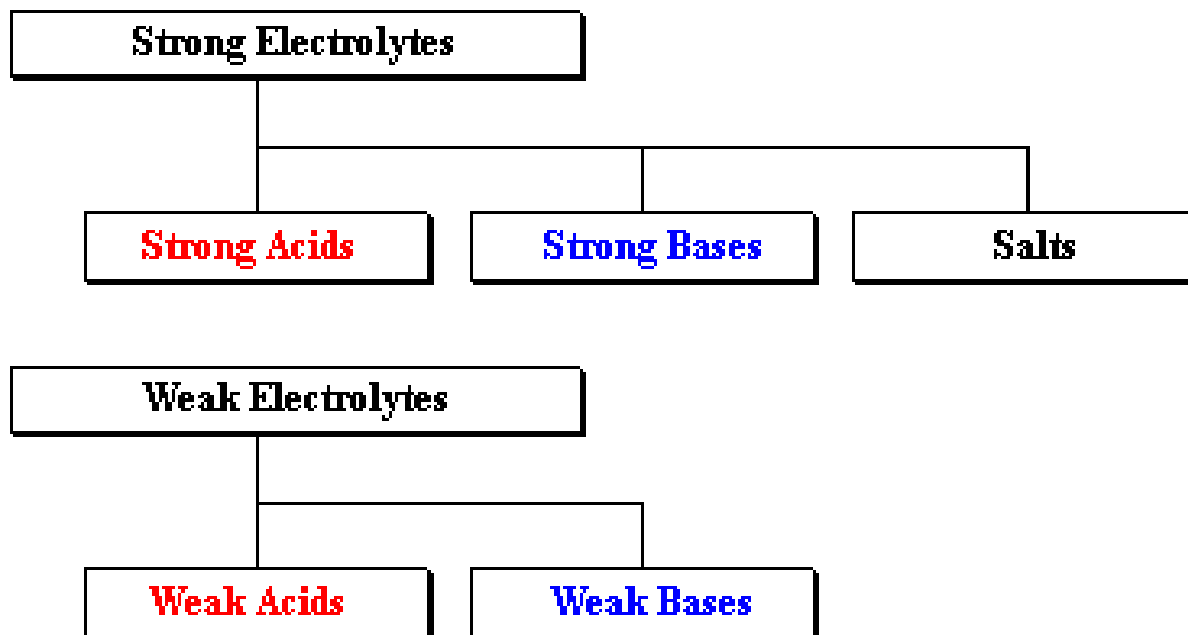
# Agitation (mixing, stirring, etc.)



- Agitation: stirring moves dissolved solute particles away from the contact surfaces more quickly and thereby allows new collisions between solute and solvent particles to occur. Without stirring, solvated particles move away from the contact areas slowly.



# Strong and weak electrolytes



## Strong Electrolytes

HCl, HBr, HI  
HClO<sub>4</sub>  
HNO<sub>3</sub>  
H<sub>2</sub>SO<sub>4</sub>  
KBr  
NaCl  
NaOH, KOH  
Other soluble ionic  
compounds

## Weak Electrolytes

CH<sub>3</sub>CO<sub>2</sub>H  
HF

## Nonelectrolytes

H<sub>2</sub>O  
CH<sub>3</sub>OH (methyl alcohol)  
C<sub>2</sub>H<sub>5</sub>OH (ethyl alcohol)  
C<sub>12</sub>H<sub>22</sub>O<sub>11</sub> (sucrose)  
Most compounds of carbon  
(organic compounds)



# Dissociation of weak electrolytes



For  $\text{CH}_3\text{COOH}$  we have  $\text{CH}_3\text{COOH} \leftrightarrow \text{CH}_3\text{COO}^- + \text{H}^+$

$$K_g = \frac{\bar{k}}{\bar{k}} = \frac{[\text{CH}_3\text{COO}^-][\text{H}^+]}{[\text{CH}_3\text{COOH}]}$$

if concentration of  $\text{CH}_3\text{COOH}$  is  $C$ ,  $\alpha$  – the degree of dissociation, so, concentrations of ions will be  $\alpha C$ , and concentration of non dissociated molecules is  $(1-\alpha)C$ , we have:

$$K_g = \frac{\alpha^2 C}{1-\alpha} \qquad K_g = \frac{(\alpha C)^2}{(1-\alpha)C}$$

for weak electrolytes  $\alpha$  is too low, so we have:

$$K_g = \alpha^2 C,$$

$$\alpha = \sqrt{\frac{K_g}{C}}$$

# For strong electrolytes we use a (activity) instead of concentration



$$a = f \cdot C$$

Where  $f$  is activity coefficient,  $C$  is molarity of electrolyte  
 $f$  of *some ion* depends on its charge ( $Z$ ) and ionic strength of the solution ( $I$ )

$$I = \frac{1}{2} \sum C_i Z_i^2$$

Concentration is a measure of the amount of solute dissolved in a given amount of solvent.



**PERCENT by MASS concentration :**  
the number of grams of solute per 100 grams of solution

- $W\% = \frac{\text{mass of solute}}{\text{mass of solution}} \times 100\%$

Parts of solute per 100 parts of solution.



*mass of solution = mass of solute + mass of solvent*

The statement “5% aqueous solution of NaCl”

has the following meaning:

the solution contains 5g of NaCl in 100g  
of solution.

The solution contains 5g of NaCl and

$100 - 5 = 95$ g of  $H_2O$ .



**MOLARITY (M): the number of moles of solute dissolved per liter of solution.**

Note: Here, volume refers to the total volume of the solution in L, not the volume of the solvent.

- Molarity (M) =  $\frac{\text{moles of solute}}{\text{volume of solution (L)}}$
- M is pronounced “molar”

# Molal concentration (Molality)

Molality ( $C_m$ ) is a concentration of a solution expressed as number of moles of solute per kilogram of solvent.

$$\begin{aligned} \text{molal concentration } (C_m) &= \\ &= \frac{\text{number of moles solute}}{\text{number of kilograms solvent}} \end{aligned}$$

$$C_m = \frac{\nu}{m_{\text{solvent}} (\text{kg})}$$





# Mole Fraction

- A mole fraction equals the number of moles of solute or solvent in a solution divided by the total number of moles of solute and solvent

$$X_A = \frac{n_A}{n_A + n_B}$$

$X_A$  = mole fraction of substance A

$n_A$  = moles of A

$n_B$  = moles of B

# $C_N$ - Normality


$$N = \frac{\text{Weight of solute in gram}}{\text{Equivalent mass} \times \text{Volume in litre}}$$

- Relation between normality and molarity.

$$N \times \text{Eq. wt} = \text{Molarity} \times \text{Molar mass}$$

$$N = \text{Molarity} \times \text{Valency}$$

$$N = \text{Molarity} \times \text{Number of H}^+ \text{ or OH}^- \text{ ion.}$$


$$C_N = \frac{w}{f_{eq} \bullet V_{solute} (L)}$$

$$f_{eq} = \frac{1}{n}$$

$$\text{Normality (N)} = \frac{\text{gram equivalent of the solute}}{\text{volume of solution (L)}}$$



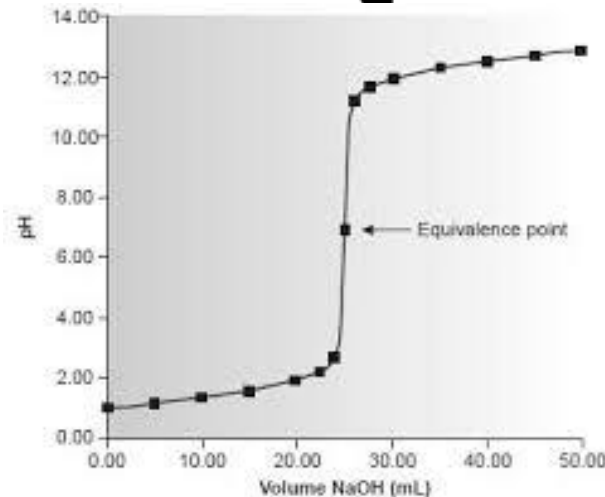
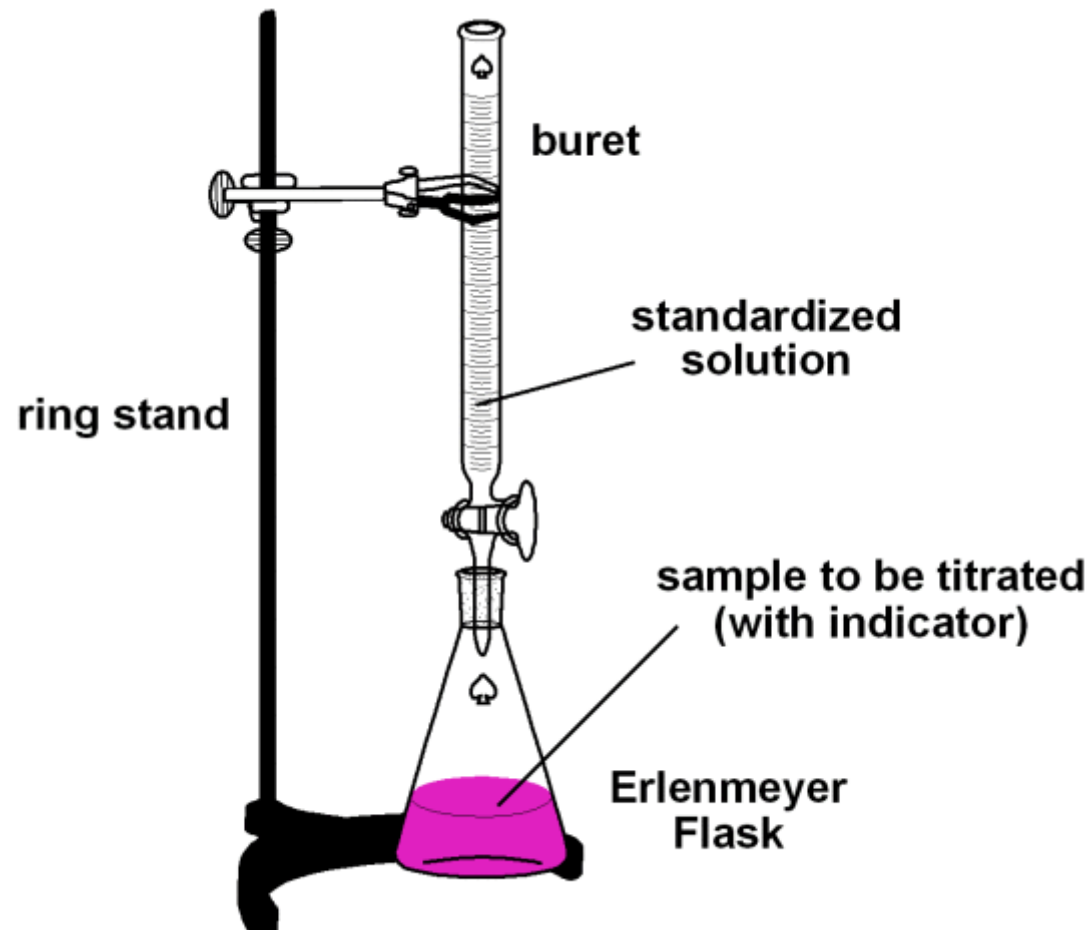
# Titration

Law of equivalents

$$V_1 \times C_{N1} = V_2 \times C_{N2}$$

Titer shows mass of solute in grams dissolved in volume of solution in mL

$$T = \frac{\text{mass in g}}{V \text{ in mL}}$$



# Conclusions



## MEASURING CONCENTRATION \*



**molarity = moles / liters**

SOLUTION  
VOLUME

**molality = moles / kg**

SOLVENT  
MASS

**mole fraction = moles / total moles**

**% solution = (grams / total grams) x 100 %**

**normality = equivalents / liters**

MOLES OF H<sup>+</sup> OR OH<sup>-</sup>

THE NUMERATORS IN ALL OF THE  
EQUATIONS REFER TO THE SOLUTE

More information you can find in our site:

[chem.teset.sumdu.edu.ua](http://chem.teset.sumdu.edu.ua)

Information for students. Teacher Yanovska A.

# Thank you for your attention!



If you're not part of  
the **solution**



you're part of the  
**precipitate.**