#### **Medical Chemistry**

# Solutions & Colloids

### Definitions

- □ A **SOLUTION** (soln): is a mixture of 2 or more substances in a single phase.
- One constituent is usually regarded as the SOLVENT and the others as SOLUTES.
- □ SOLUTE: the part of a solution that is being dissolved (usually the lesser amount).
- **SOLVENT:** the part of a solution that dissolves the solute (usually the greater amount).
- Solutions in which the solvent is WATER are called AQUEOUS SOLUTIONS.



## **Types of Solutions**

Solute	Solvent	Appearance of solution	Example
Solid	Solid	Solid	14-carat gold (Cu/Ag/Au) Brass alloy (Zn/Cu)
Solid	Liquid	Liquid	Salt water
Liquid	Liquid	Liquid	Alcohol in water
Gas	Liquid	Liquid	Soda (CO <sub>2</sub> in water)
Gas	Gas	Gas	Air (N <sub>2</sub> , O <sub>2</sub> ,)

**CAN A SOLUTION BE SOLID?** 

## **Characteristics of Solutions**

- 1. Distribution of particles is uniform.
- 2. Components do not separate on standing.
- 3. Components cannot be separated by filtration.
- 4. It is possible to make solutions of many different solute/solvent compositions.
- 5. Solutions are transparent (even if colored).
- 6. Solutions can be separated into pure components (e.g., distillation, chromatography). This separation is a physical change.

#### **Concentration Units 1**

- Concentration: is the amount of solute in a given amount of solution (rarely "in amount of solvent"). <u>UNITS:</u>
- **1. Percent Composition:**
- a. % mass (w/w) = (mass of solute/mass of soln) x 100
- b. % volume (v/v) = (volume of solute/volume of soln) x 100
- c. % mass/volume (w/v) = (mass of solute/volume of soln) x 100
- 2. Molarity (M) = moles of solute/liter of soln (v)

#### **Concentration Units 2**

- 3. Molality (m) = moles of solute/kg of SOLVENT
- 4. Parts per million (ppm) = (mass of solute/mass of soln) x 10<sup>6</sup>
- 5. mole fraction (x) = moles of solute/total moles of soln
- 6. Mass per volume (mg/L) = mass of solute/liter of soln
- 7. Normality (N) = equivalents of solute/liter of soln

# **Concentration Units 3**

- □ **M** = **m** when the solvent is distilled  $H_2O$  since its density = 1 then, 1 L = 1 kg (NOT salt  $H_2O$ ).
- **D** ppm =  $10^3$  ppb (part per billion) =  $10^6$  ppt (part per trillion)
- Mass (moles) of soln = mass (moles) of solute + mass (moles) of solvent
- Common mass ratios for solutions and solids are:

Units	Solutions		Solids	
ppm	mg/L	µg/mL	mg/kg	µg/g
ppb	μg/L	ng/mL	µg/kg	ng/g
ppt	ng/L	pg/mL	ng/kg	pg/g

Example 1: An IV soln is prepared by dissolving 5.0 g glucose (C<sub>6</sub>H<sub>12</sub>O<sub>6</sub>) in dist. H<sub>2</sub>O to make 100 mL soln. Calculate (a) molarity M, (b) % w/v, and (c) ppm of the IV soln.

#### **Solution:**

(a) Convert:  $g \rightarrow moles of glucose.$ Since, molar mass of  $C_6H_{12}O_6 = 180.0 \text{ g/mol}.$ Then, 5.0 g x (1 mol/180.0 g) = 2.78 x 10<sup>-2</sup> mol. Thus, M = moles of solute/L of soln = 2.78 x 10<sup>-2</sup> mol/1.00 X 10<sup>-2</sup> L = 2.78 M



(c) ppm = (mass of solute/mass of soln) x  $10^6$  = (5.0 g/100 g) x  $10^6$ = 5.0 x  $10^4$  [since d(H<sub>2</sub>O) = 1, 100 mL = 100 g]

#### **Preparing Solutions (1.0 M NaCl)**



# Dilution

Dilution: is adding extra solvent to decrease the concentration of a soln.

The amount of solute remains constant before and after dilution, but the concentration decreases.

Before dilution After dilution  $Conc_1 \times Vol_1 = Conc_2 \times Vol_2$   $M_1 \times V_1 = M_2 \times V_2$  $\%_1 \times V_1 = \%_2 \times V_2$ 

Concentrations and volumes can be most units as long as they are consistent. **Example:** How do we prepare 200 mL of a 3.5 M soln of acetic acid if we have a bottle of conc acetic acid (6.0 M) ?

**Given:** 

		Initial soln	Final soln	
<b>Concentration:</b>		6.0 M	3.5 M	
	Volume:	? L	0.20 L	
Find:	L of initial acetic acid			
Solve:	$M_1 \times V_1$	$= M_2 \times V_2$		
	6.0 M	x V <sub>1</sub> = 3.5 M x 0.2	0 L	
	$V_1 = 3.5 M x$	0.20 L/ 6.0 M = 0.1	Í2 L	
	• 0 1 2 I (1 2 0 )	ml) of conclusion	acid in a 200 m	

Put 0.12 L (120 mL) of conc acetic acid in a 200-mL volumetric flask, add some water and mix, and then fill to the mark with water.

#### How H<sub>2</sub>O Dissolves Ionic Compounds

- □ Consider NaCl (solute) dissolving in water (solvent).
- □ The water H-bonds have to be interrupted,
- □ NaCl dissociates into Na<sup>+</sup> and Cl<sup>-</sup>,
- □ Cat/Anions attract oppositely charged ends of H<sub>2</sub>O molecules (Na<sup>+</sup>....<sup>δ</sup>-OH<sub>2</sub> and Cl<sup>-</sup>....<sup>δ+</sup>H<sub>2</sub>O).
- When attraction forces of ions to H<sub>2</sub>O molecules is greater than ionic bond (keeping ion in crystal), the ion will be completely removed from the crystal and surrounded by H<sub>2</sub>O molecules (HYDRATED ions).
- Such interaction between solute and solvent is generally called SOLVATION.

#### How H<sub>2</sub>O Dissolves Covalent Compounds

- Molecules should have no more than 3 C atoms for each O, N, or F atom.
- **Examples:** Acetic Acid  $CH_3COOH$  is soluble but benzoic acid  $C_6H_5COOH$  is not.
- Although table sugar, C<sub>12</sub>H<sub>22</sub>O<sub>11</sub>, contains a large number of C atoms, it is very soluble in H<sub>2</sub>O because it has many O atoms and O-H bonds that can form many H-bonds with H<sub>2</sub>O.
- 2. Compounds rarely react with H<sub>2</sub>O giving ions

**E.g. 1**  $HCl(g) + H_2O(I) \rightarrow Cl^{-}(aq) + H_3O^{+}(aq)$ 

**E.g. 2** SO<sub>3</sub>(g) + 2H<sub>2</sub>O(I)  $\rightarrow$  HSO<sub>4</sub><sup>-</sup>(aq) + H<sub>3</sub>O<sup>+</sup>(aq)