

CHEM 206 section 01

LECTURE #8

Wed. Jan.30, 2008

## ASSIGNED READINGS:

**TODAY'S CLASS:** Ch.14: section 14.3

**NEXT CLASS:** finish Ch.14  
(14.5 - detergents only - Lect.07)

(1)

## How much solute is dissolved?

### Qualitative terminology...

- **Saturated solution** = contains maximum concentration of solute that will dissolve in that solvent at that temperature

NOTE: in a sat'd sol'n, have equal rates of dissolving and precipitating (an equilibrium situation...)

- **Solubility** = the quantity of solute dissolved in a saturated sol'n (often expressed as: g solute per 100mL water)
- **Unsaturated solution:** concentration less than saturated... more solute can dissolve
- **Supersaturated solution** (see Closer Look p.563)  
= a metastable state where too much solute is dissolved...  
...any disturbance  $\Rightarrow$  solute will precipitate suddenly

(2)

## 14.3 Environmental factors affecting solubility:

### temperature (T) & pressure (P)

Important for dissolving solids, liquids & gases

- Is the dissolution process enthalpically favourable? (*requires heat, or not?*)
- entropically favourable? (*will  $\uparrow T$  make  $\Delta G$   $\uparrow$  or  $\downarrow$ ?*)

Important for dissolving gaseous solutes only

- Partial pressure of gas determines frequency of collisions between gas & solution's surface
- $\Rightarrow$  determines [gas] at which have balance of rates of gas dissolving & leaving solution
- = a saturated sol'n, at eqm, where [gas] = "solubility"

Entropic thinking important here...

(3)

### Effect of Temperature

$\uparrow T \Rightarrow \uparrow$  Rate of dissolution (*always*)  
 $\uparrow$  Solubility? *...not always!*

### TRENDS FOR AQUEOUS SOL'NS: for most cases...

**Dissolved solids:** solubility  $\uparrow$  with  $\uparrow T$

- WHY? Dissolving is entropically favourable...
- Thus:  $\Delta G_{\text{dissolution}}$  becomes more -ve as  $T \uparrow$
- So: To crystallize out solutes: cool down

*Exceptions:  
sodium sulfate,  
cerium sulfate*

**Dissolved gases:** solubility  $\downarrow$  with  $\uparrow T$

- WHY? Dissolving gas is entropically unfavourable...
- Thus: To release gas (entropically favourable): warm up

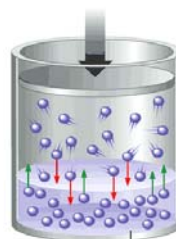
To predict effect of T case by case: consider both  $\Delta H$  &  $\Delta S$ ...

- textbook discusses enthalpy only, because it's only Ch.14
- you should use Ch.19's entropy thinking - other way isn't always right!

(4)

## Effect of pressure: on gas solubility

- Pressure related to # gas molecules per unit volume
- higher P ⇒ more COLLISIONS with liquid
  - ⇒ ↑ rate of entry of gas into solvent
  - exit unaffected



↑ P: ↑ Solubility

**Henry's law:**  $C = k_H P$   
 concentration of dissolved gas  
 IS DIRECTLY PROPORTIONAL TO  
 partial P of that gas above the solution

In Kotz textbook, written:

$$S_g = k_H P_g$$

↑  
**Solubility:** conc. of gas  
 in a sol'n saturated with  
 that gas at a given T & P

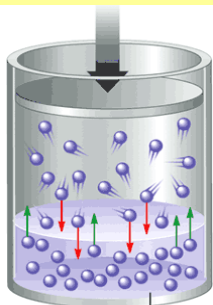
Table 14.3 • Henry's Law Constants (25 °C)

Gas	$k_H$ (M/mm Hg)
N <sub>2</sub>	$8.42 \times 10^{-7}$
O <sub>2</sub>	$1.66 \times 10^{-6}$
CO <sub>2</sub>	$4.48 \times 10^{-5}$

**IMPORTANT: \*\* unit analysis crucial here, especially P units! \*\***  
 often given solubility data at given partial P of gas (calculate own  $k_H$ )

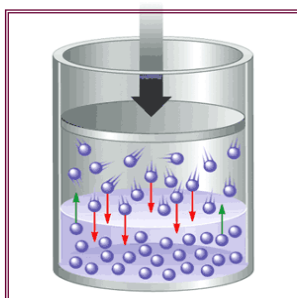
Zumdahl's  
 Figure 11.5

## GASES DISSOLVE BECAUSE MOLECULES COLLIDE WITH THE LIQUID'S SURFACE...



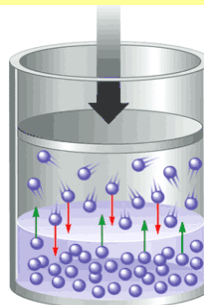
### SOLUTION AT EQUILIBRIUM

Balance between  
 gas molecules  
 entering (*dissolving*)  
 & escaping from  
 solution



### CHANGE:

↑ PRESSURE  
 ⇒ ↑ collisions  
 with surface  
 ⇒ ↑ rate of  
 dissolution



### RESULT:

↑ [dissolved gas]  
 ⇒ ↑ rate of escape  
 ⇒ new equilibrium  
 reached

## Example: supersaturated solutions containing GAS

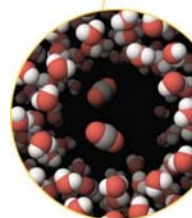
Sudden ↓ in pressure ⇒ gas erupts from solution!

### Tragedy at Lake Nyos, Cameroon (K&T p.556)

- Lake in deep volcanic crater  
⇒ active vents beneath, emitting  $\text{CO}_2(\text{g})$
- Poor turnover of water  
⇒ depths sat'd with  $\text{CO}_2(\text{g})$  at high  $P_{\text{CO}_2}$



- Aug. 21, 1986: Tremor?  $\Delta$  in temperature?  
⇒ sudden turnover of water in lake  
⇒ high- $[\text{CO}_2]$  water now seeing low  $P_{\text{CO}_2}$  of air
- 260 ft geyser of  $\text{CO}_2$
  - dense gas cloud hugged ground; 45 mph
  - 1700 people died, 12 miles away



Now: PREVENTION: 200 m pipes vent depths

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## Example: $\text{CO}_2$ in your soda pop (from an old exam)

The solubility of gaseous  $\text{CO}_2$  in water at  $10^\circ\text{C}$  is 0.240 g per 100.0 mL, under a pressure of 1.0 atm of carbon dioxide. During manufacturing, a soft drink at  $10.0^\circ\text{C}$  is saturated with  $\text{CO}_2$  under a pressure ( $P_{\text{CO}_2}$ ) of 4.0 atm, and then sealed.

- a) What mass of  $\text{CO}_2$  is dissolved in a 355 mL can of this drink?

Ans:  $k = 0.0024 \text{ g}/(\text{mL}\cdot\text{atm})$   
 $m_{\text{CO}_2} = 3.4 \text{ g dissolved}$

- b) Imagine you open a can of this beverage and leave it open to the atmosphere ( $P_{\text{TOTAL}} = 1 \text{ atm}$ ) at  $10^\circ\text{C}$  to go "flat".

[Air is ~ 0.031%  $\text{CO}_2$  by mole; thus,  $P_{\text{CO}_2} = \dots$ ]

What volume of  $\text{CO}_2$  will be released from the beverage?

Ans:  $m_{\text{CO}_2\text{-dissolved}} = 0.000264 \text{ g}$   
 $m_{\text{CO}_2\text{-released}} = 3.4077 \text{ g (all...)}$   
 $V_{\text{CO}_2 \text{ at } 1 \text{ atm}} = 1.8 \text{ L}$

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## ASSIGNED READINGS:

### BEFORE NEXT CLASS:

**Read:** rest of Ch.14

**Practice:** Converting between concentration units  
& solving gas solubility problems

**Next class:** finish Ch.14