

Thinking Like a Chemist About Dissolution

Unit 5 Day 3

CH302 Vanden Bout/LaBrake Spring 2013

What are we going to learn today?

Thinking Like a Chemist in the Context of the
Dissolution Process.

- Macro Modeling
- Micro Modeling
- Energy of the change Modeling

CH302 Vanden Bout/LaBrake Spring 2013

Mathematical Relationship between VP and T

Vapor pressure of a liquid increases with increasing T

1

Mathematical Relationship between VP and T

Vapor pressure of a liquid increases with increasing T

Clausius-Clapeyron equation

$$\ln \left(\frac{P_2}{P_1} \right) = \frac{\Delta H_{vap}^\circ}{R} \left[\frac{1}{T_1} - \frac{1}{T_2} \right]$$

↑ vapor pressure any pressure unit

↑ gas constant J / K.mol

↑ temp in Kelvin

kJ/mol

watch units match all to gas const. J vs. kJ

Piazza!

CH302 Vanden Bout/LaBrake Spring 2013

IMPORTANT INFORMATION

HW1 due TODAY 9AM
 LM04 & LM05 due TODAY 9AM
 LM06 Solutions due Th 9AM
 LM07 Solutions – Concentrations due Th 9AM
 LM08 Henry's Law due Th 9AM

LUNCH WITH VDB or LaB?

NOTE: THIS CLASS HAS BEEN DESIGNED TO BE A "HYBRID" OR "BLENDED" STUDENT CENTERED LEARNING EXPERIENCE. THIS MEANS THAT SOME OF THE COURSE INFORMATION WILL BE PRESENTED IN A DIGITAL FORMAT ON THE WWW IN EITHER PRINT OR VIDEO FORMAT. SO LEARNING MODULE INDICATES NEW LEARNING. HOMEWORK INDICATES PRACTICING CONCEPTS/PROBLEM SOLVING THAT HAS BEEN FIRST INTRODUCED EITHER IN CLASS, ON THE WEBSITE OR IN A LEARNING MODULE.

16 dm

14 solⁿ & molarity

15 mass % vs density

16 soln dilution $m_1 V_1 = m_2 V_2$

20 solubility
 ↳ ALEXS

CH302 vanden bout/LaBrake Spring 2013

Quiz: CLICKER QUESTION 1

Which of the following has the highest vapor pressure?

- a) CH₃OH
- b) CH₂CH₂OH

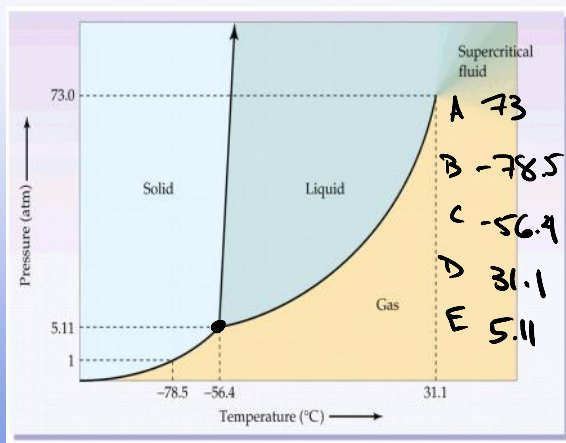
Which of the following has the highest vapor pressure?

- a) CH_3OH lowest IMF \rightarrow \uparrow VP
dispersion
b) $\text{CH}_3\text{CH}_2\text{OH}$
c) $\text{CH}_3\text{CH}_2\text{CH}_2\text{OH}$
d) $\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{OH}$

CH302 Vanden Bout/LaBrake Fall 2013

Quiz: CLICKER QUESTION 2

What is the temperature at the triple point?



A. 73.0
 $^{\circ}\text{C}$

B. -78.5
 $^{\circ}\text{C}$

C. -56.4
 $^{\circ}\text{C}$

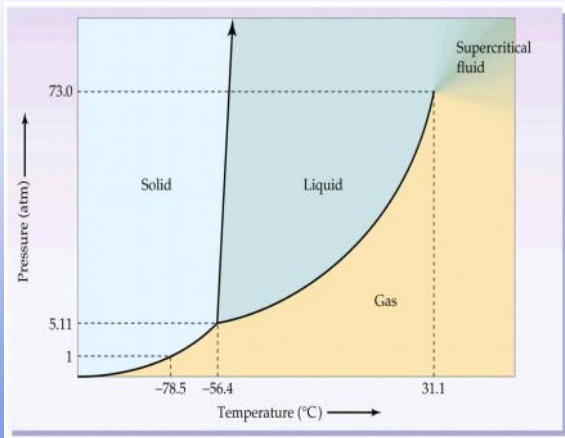
D. 31.1
 $^{\circ}\text{C}$

E. 5.11

CH302 Vanden Bout/LaBrake Spring 2013

Quiz: CLICKER QUESTION 3

At the triple point solid, liquid, and gas all have the same....



- A. number of moles
- B. free energy**
- C. volume
- D. density
- E. entropy

CH302 Vanden Bout/LaBrake Spring 2013

Solutions: Vocabulary Check!

Solution = Solvent + Solute
greater *less*

Solubility - *how much dissolves*
concentration

Dissolution *process of making solution,*
dissolving

Homogeneous Solution described by components
and concentrations of those components!

LM will instruct concentration units.

*Look at
NGM
if terms
are not
similar*

CH302 Vanden Bout/LaBrake Spring 2012

Get to work on the worksheet!

MACROSCOPIC DESCRIPTION OF DISSOLUTION

Get to work on the worksheet!

MACROSCOPIC DESCRIPTION OF DISSOLUTION

White solid/crystal dissolves
"vanishes" in liquid
no temperature change

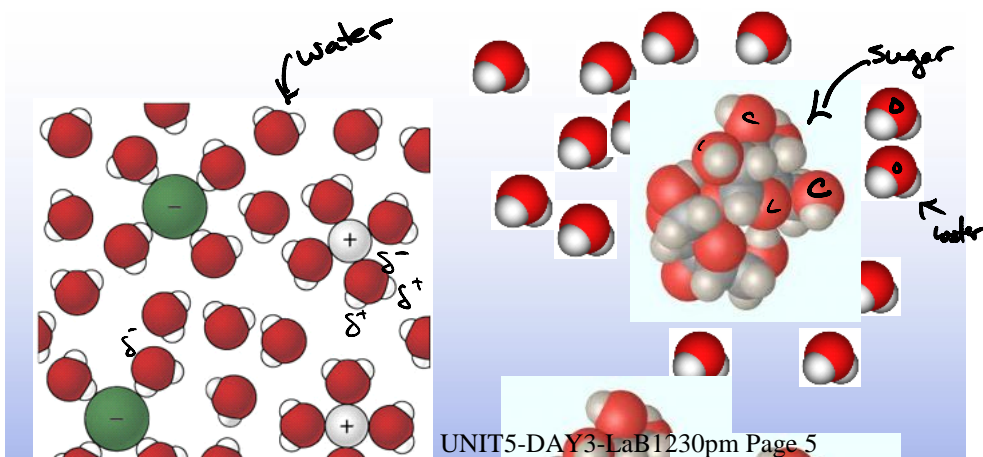
CH302 Vanden Bout/LaBrake Spring 2013

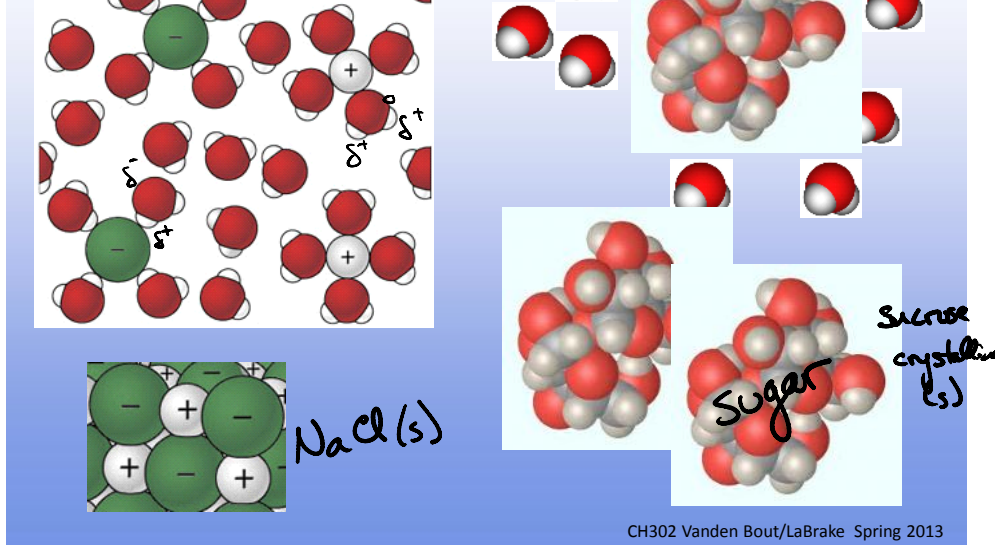
Polling: Clicker Question 4

A nice micro view of
the dissolution of a(n):

- A. ionic solid
- B. molecular solid
- C. metallic solid
- D. covalent solid

CH302 Vanden Bout/LaBrake Spring 2013





Dissolution Demonstration Observations

Sodium chloride in water

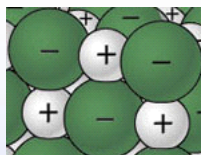
Sucrose in water

Ammonium nitrate in water

CH302 Vanden Bout/LaBrake Spring 2013

#7

How do we think about this in terms of energy? $\Delta H_{\text{solution}}$



1. break solute

- costs energy - Lattice Energy

$+\Delta H$ endothermic

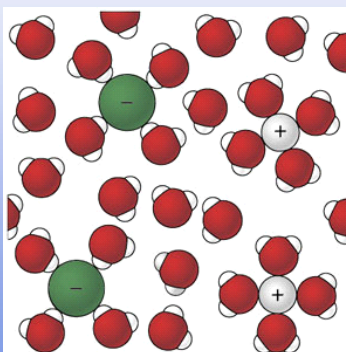
(ignore break of solvent) \rightarrow very small

2. form new

solute-solvent interactions

- give off energy - Solvation Energy

$-\Delta H$ exothermic



$$\Delta H_{\text{solution}} = \Delta H_{\text{Lattice Energy}} + \Delta H_{\text{solvation}}$$

\rightarrow hydration if water

CH302 Vanden Bout/LaBrake Spring 2013

Usually $|\Delta H_{\text{lattice}}| > |\Delta H_{\text{solvation}}|$
so $\Delta H_{\text{solution}}$ is typically endothermic

What did we learn just now?

$\Delta H_{\text{solution}}$ is hard to predict. Depends on energy needed to separate solute-solute and energy release when new solute-solvent attractions are formed.

$$\Delta H_{\text{solution}} > 0$$

Typical Situation Solute-solvent interactions are weaker than solute-solute (and solvent-solvent)

$$\Delta H_{\text{solution}} < 0$$

Unusual, but possible

Solute-solvent interactions are stronger than solute-solute and solvent-solvent

Exceptions: high charge density ^{ions}; $\Delta H_{\text{solvation}}$ very high
gas dissolving in liquid is exothermic

CH302 Vanden Bout/LaBrake Spring 2013

gas has no IMFs!

Polling: Clicker Question 5

What is going on with the entropy of dissolution for our example solutions?

\rightarrow solid solute

general

A. The entropy always increases.

B. The entropy always decreases.

What is going on with the entropy of dissolution for our example solutions?

- A. The entropy always increases.
- B. The entropy always decreases.
- C. It depends on the type of solid solute.
- D. It depends on the phase of the solute.

solid solute
in general solids dissolving in liquids increase entropy
gas dissolved in liquid, entropy dec

CH302 Vanden Bout/LaBrake Spring 2013

Entropy of Solution $\Delta S_{\text{solution}}$ usually easy to predict

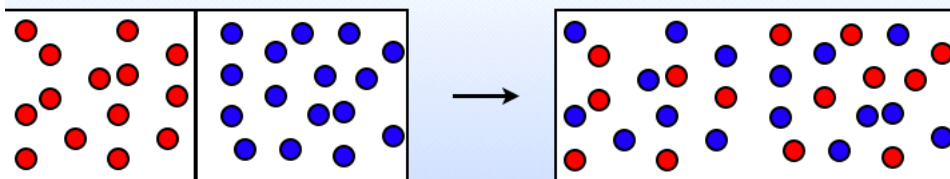
Solutions typically have a higher entropy than the unmixed compounds
Therefore $\Delta S_{\text{solution}} > 0$
For most cases

Since entropy almost always favors mixing, the differences between different substances are the result of enthalpy (intermolecular forces)

CH302 Vanden Bout/LaBrake Spring 2013

Mixtures

What is different than pure substances?



gas

gas

mix of gases

What is the free energy change for gases mixing?

ΔG < 0 - spontaneous

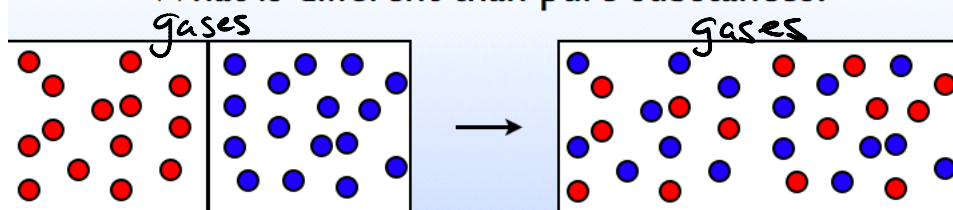
What is the free energy change for gases mixing?

$$\Delta G < 0, \text{ spontaneous}$$

POLLING CLICKER QUESTION 6

Mixtures

What is different than pure substances?



Why does the free energy decrease? ($\Delta G = \Delta H - T\Delta S$)

- for gases
- A. ΔH is positive, ΔS is zero
 - B. ΔH is zero, ΔS is positive
 - C. ΔH is negative, ΔS is positive
 - D. ΔH is negative, ΔS is zero

ΔH difference
in IMFs
gases have no
IMFs

POLLING CLICKER QUESTION 7

How can you explain the spontaneous dissolution of the endothermic solution?

$$\Delta G = \Delta H - T\Delta S$$

- A. It must be entropically driven
- B. It must not really be endothermic
- C. It must only occur at very low temperatures, because the solution does get cold
- D. It must be driven by the change in enthalpy
- E. It must not be as soluble as endothermic solutions

CH302 Vanden Bout/LaBrake Spring 2013

POLLING CLICKER QUESTION 8

What is the sign for the change in Free Energy for the dissolution process in which polystyrene peanuts were placed in water?

- A. Positive, nonspontaneous
- B. Negative
- C. $\Delta G = 0$,
- D. Need values for ΔH and ΔS to determine.

CH302 Vanden Bout/LaBrake Spring 2013

POLLING CLICKER QUESTION 9

What is the sign for the change in Entropy for the dissolution process in which polystyrene peanuts were placed in acetone?

- A. Positive
- B. Negative

were placed in acetone?

- A. Positive
- B. Negative
- C. ~~ΔG~~ ^{ΔS} = 0
- D. Need values for ΔH and ΔS to determine.

CH302 Vanden Bout/LaBrake Spring 2013

POLLING CLICKER QUESTION 10

What is the sign for the change in Enthalpy for the dissolution process in which biodegradable peanuts were placed in acetone?

$$\Delta G = \Delta H - T\Delta S$$

+ ? - (+)

- A. Positive
- B. Negative
- C. ~~ΔG~~ ^{ΔH} = 0
- D. Need values for ΔH and ΔS to determine.

CH302 Vanden Bout/LaBrake Spring 2013

What is the sign of change in free energy for mixing olive oil with water?

- A. Negative
- B. Positive
- C. 0
- D. None of the above

CH302 Vanden Bout/LaBrake Spring 2012

Talk about like dissolves like..

IMFs are same, so
 ΔH is near 0

CH302 Vanden Bout/LaBrake Spring 2012

What did we learn just now?

When a solute dissolves spontaneously in a solvent, the process is considered spontaneous.

For a spontaneous dissolution process, $\Delta G < 0$

In some cases non-spontaneous dissolution process can become spontaneous upon increasing the temperature.

CH302 Vanden Bout/LaBrake Spring 2012

What did we learn today?

Micro modeling of the dissolution process – noted the difference between molecular solids and ionic solids.

Energy modeling of the dissolution process – noted the changes in enthalpy (solute-solute interactions vs solute-solvent interactions).

Energy modeling of the dissolution process - noted changes in entropy – typically depends on the phase of the solute (solid dissolving in liquid vs gas dissolving in liquid).

Energy modeling of the dissolution process – predict based on sign of ΔG .

CH302 Vanden Bout/LaBrake Spring 2012

Learning Outcomes

Describe the factors that favor the dissolution process in terms of the intermolecular forces and thermodynamics (enthalpies of solution, solvation, lattice energy, entropies of solution, free energy of solution)

Describe how P (Henry's Law) affects solubility of a gas.

Define and perform calculations for common concentration units molarity, molality, and mole fraction.

CH302 Vanden Bout/LaBrake Spring 2012