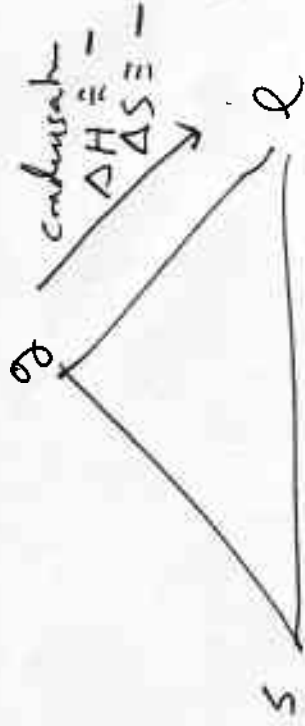


30 question types for the exam

Chapter 8

1 Theory: temperature and physical equilibria (a phase change where you have to predict H and S)

review from CH301 w/ ~~lecture~~ ^{equation} $\Delta G = \Delta H - T\Delta S$



given a transition, you tell me the sign of $\Delta H, \Delta S$

(signs are same for all phase change $\Delta H, \Delta S$
+ , +
- , -

melting $\Delta S \equiv +, \Delta H \equiv +$

2 Theory: vapor pressure

know what the definition of v.p. is and you will be fine

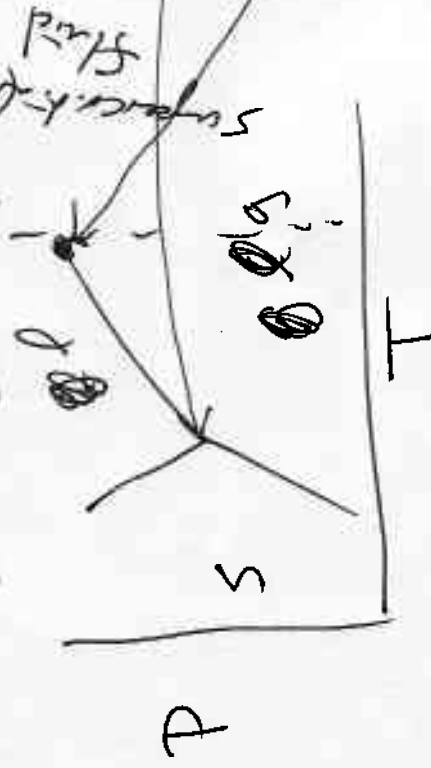


surface molecules pick up internal energy, overcome i.m.f. and vaporize.
The smaller the i.m.f. the larger the vapor

3 Theory: salt dissociation in water

A salt crystal held together by C.L. energy is attached by water molecules w/ a ΔH_{soln} . The ΔH_{soln} vs. C.L. energy determines ΔH_{soln} is + or - for NaCl. it is slightly positive so that according to LeChatelier as the $T \uparrow$ more salt dissolves.

4 Problem: phase diagram interpretation



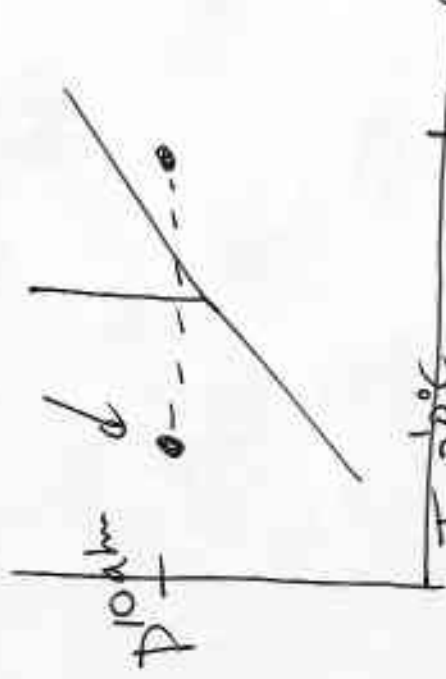
I will give a phase diagram.
Know all the famous points

There are equilibria for the phases changes. The triple point is where all phases exist simult. The critical point is the T beyond which no liquid can form as pressure increase

5 Problem: phase diagram navigation

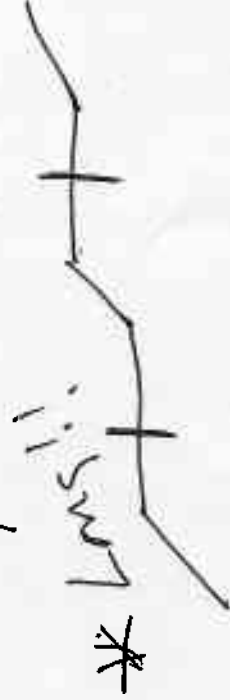
I give you a starting P, T. Then I tell you to walk around the diagram and then say "stop". you tell me where you are.

Example: (10 atm, 20°C) increase to 100°C, what is the phase?
Answer: gas (condensation occurs)



6 Calculation: ΔH for heating across phases

you have 5 possible calculations depending upon where T starts and stops.



3 calc.: ave $\Delta H = mCST$ (phase)
2 calc.: $\Delta H_{vap} \approx \Delta H_{fus}$ at b.p. or f.p.

14.4: This problem sucks. (Not water) So be ready for a non- H_2O problem.

7 Theory: gas solubility in liquids

Gases dissolve in l.s. v. d. s. become

- reactive like CO_2
- strong IMF like HCl
- wimpy little things that H_2O doesn't care about $\text{He}, \text{N}_2, \text{O}_2$

Typically for nonpolar gases

They have an exothermic

dissolution $\text{He(g)} \rightarrow \text{He(aq)}$ exo.

$\therefore$ As $T \uparrow$ [gas] $\downarrow$ and little fishes die

8 Ranking: miscibility of liquids

Employ "like dissolves like" You will be given a compound, either polar or nonpolar. You need to find out which in a list it is most "like". That is most misc. sol.

Least "like" most immiscible.

Example CHCl_3 H_2O CCl_4



9 Definition: colligative property

You have around 4 colligative properties.

- V.P. lowering $P = P^\circ X$ (Raoult's Law) why we mix antifreeze w/ H_2O to keep car from overheating
- $\Delta T_b = K_m$ says that as you add solute to solvent, the T_b goes up
- $\Delta T_f = K_m$ says that as you add solute $\rightarrow$ solvent f.p. $\downarrow$ (salt is added to ice)
- $\Pi = MRT$ (osmotic pressure) says that H_2O will cross a semipermeable membrane to equilibrate [Why cells explode]

10 Calculation: vapor pressure in binary system very similar to gas problem.
 Given 2 solvents, each w/ a mole fraction and their P^0 values
 (pure v.p. values) Perform Raoult's Law on each and
 give partial v.p. that you add together for P_{TOT} .

$$P_A = P_A^0 X_A \quad P_{TOT} = P_A + P_B$$

$$P_B = P_B^0 X_B$$

11 Calculation: Clausius Clapeyron

Plug + Chng. Given

ΔH_{vap} , R , P_1, T_1 , P_2, T_2 One of these is missing. Solve for it.

$$\ln \frac{P_2}{P_1} = \frac{\Delta H}{R} \left(\frac{1}{T_1} - \frac{1}{T_2} \right)$$

Hint. you don't get to solve for ΔH_{vap} this time.

Hint. Remember that b.p. $\equiv$ at 760 torr

Hint. save time.

12 Ranking: Van't Hoff and solution concentrations

This is the i in the coll. $\pm$ property calculations that corrects for # of particles found when a compd dissolves.

sugar 1 particle $\equiv i=1$

and 2 particles $\equiv i=2$

$CaCl_2$ 3 particles $\equiv i=3$

13 Problem: colligative property application

I have told some practical things about colligative properties. Know these. Explain to a friend.
How to deice icy roads How to cook pasta faster
How to keep radishes firm over winter How to kill your animals

14 Calculation: colligative property

$$P = P^0 X \quad \text{Raoult's Law}$$

$$\Delta T_b = K_b m$$

$$\Delta T_f = K_f m$$

$$\Pi = MRT$$

you get a pretty easy
plus + chng.
↑
easy.

Chapter 9

15 Problem: Setting up K from equilibrium expression

Given a chemical rxn. Simply write the stuff on right ~~side~~ in numerator, stuff on ~~left~~ in denominator.

• Put the coeff of rxn in the exponent
* • Ignore (s) (l), care about (g) (aq) *

easy problem.

16 Calculation: calculating K from RICE expression

Easy plug + chug. I give a chem rxn. you create K for it. I tell you the equl. amounts. you plug them into K + solve.

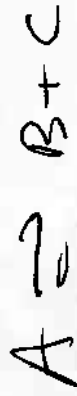


0	0	0

$$K = \frac{0 \cdot 0}{0} = \text{answer.}$$

17 Calculation: equilibrium concentrations from K

This is opposite of #16. I give you K and initial amounts. you tell me equl. amounts.



✓	✓	✓
?	?	?

given K.

Hint!!! This is the kind of problem where you set the answer equal to X and solve for everything in terms of X

18 Calculation: equilibrium concentrations from K (quadratic)

(a problem to do last)

I give you K + initial amounts. you assign a value for X and find all equl. [] in terms of X.

The answer will be a quadratic (X²)

Solve by back substit~~ution~~ or quad. form or fancy calc.

19 Problem: Reaction direction from Q and K

I will give you a K , and some initial $[]$ values.
you will solve for Q . Compare to K .
 $Q < K$ shift right $Q > K$ shift left.

Hint. TRICK TRICK TRICK.

make sure you convert your concentrations into molarity
by dividing by volume. first.

20 Problem: LeChatelier and reaction direction

Three steps.

1. Stress is applied to system ($[]$, P , T)
2. system does opposite
3. shift left or right to make step 2 happen.

Same.

21 Problem: LeChatelier and reaction direction

22 Calculation: Relationship of ΔG to K

Plug + Chng

$$\Delta G = -RT \ln K$$

I will K you find ΔG
or I will give ΔG you find K

But very routine.

H. at. K_p + K_c values

are solved exactly the same way.

Chapter 10

23 Theory of auto-prolysis of water

K is temp dependent. So as T varies for H_2O ,
 K varies. But K is what tells you the H^+ , OH^- amounts.
So $H^+ = OH^-$ are equal but not always 1×10^{-7} .

So real definition neutrality is $pH = pOH$ or $H^+ = OH^-$
not $pH = 7$. As $T \uparrow$ $K \uparrow$ $pH \downarrow$ $T \downarrow$ $K \downarrow$ $pH \uparrow$

24. Temperature dependence of K_w

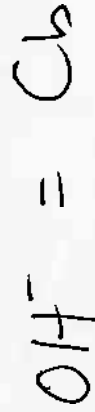
you will be given the K_w expression

$$K_w = [H^+][OH^-] \text{ with } [H_2O] \text{ a constant}$$

and will be asked to solve it. (P.S. this is a = bc)

H. at. know your exponent math + calc. or you will fail this test

25. Strong acid or base calculation



Pathetically simple calculation
You need to know S.A. + S.B.
Then the starting amount is the
answer. (May have to convert to pH)

26. Weak acid or base calculation

If you know you strong acids + bases, then the
guys + gals that aren't strong are weak. Then solve.

$$H^+ = (K_a C_a)^{1/2} \text{ for weak acids}$$

$$OH^- = (K_b C_b)^{1/2} \text{ for weak bases.}$$

You will do
This kind of
calculation
twice

Hint. When

you have a OH^- answer +
need to convert to pH, be
very to do so. Be careful!!

27 Weak acid or base calculation



any man
be correct



28. Identifying acid and base strength

Distinguishing S.A. + S.B. ~~S.A.~~

Strong acids are HCl , HBr , HI , HClO_4 , HClO_3 , H_2SO_4 , HNO_3

also know their names.

Strong bases are Group 1 hydroxides NaOH , LiOH , CsOH etc
and Group 2 hydroxides Ca(OH)_2 , Ba(OH)_2 etc.

29. Converting between pH, POH, H^+ and OH^-



$$1 \times 10^{-14} = K_w = (\text{H}^+)(\text{OH}^-) \Leftarrow \text{so if given } \text{H}^+ \text{ you can give } \text{OH}^-, \text{ etc}$$

also $\text{pH} = \text{pH} + \text{pOH} \Leftarrow \text{so if given pH you can give pOH.}$

30. Ranking acidity and basicity based on equilibrium constants

As K value increases, the rxn shifts more to the right.

So for acids $\text{HA} \rightleftharpoons \text{H}^+ + \text{A}^-$ as $K \uparrow$ $\text{H}^+ \uparrow$ $\text{pH} \downarrow$

$$K_1 = 1 \times 10^{-5}$$

$$K_2 = 1 \times 10^{-8} \leftarrow \text{weakest}$$

$$K_3 = 1 \times 10^{-12} \leftarrow \text{strongest}$$

∴ The larger K_a , the stronger the acid.

Similarly the larger K_b the stronger the base

I will save a list of K_s , find the strongest & weakest from it