

Name _____

TA's Name _____

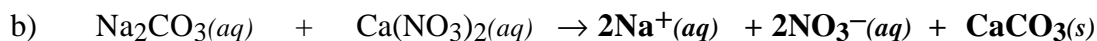
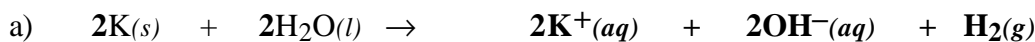
Section _____

INSTRUCTIONS:

1. This examination consists of a total of 8 different pages. The last three pages include a periodic table, a table of vapor pressures for water, a solubility table and a table of thermodynamic values. All work should be done in this booklet.
2. PRINT your name, TA's name and your lab section number now in the space at the top of this sheet. DO NOT SEPARATE THESE PAGES.
3. Answer all questions that you can and whenever called for show your work clearly. Your method of solving problems should pattern the approach used in lecture. You do not have to show your work for the multiple choice or short answer questions.
4. No credit will be awarded if your work is not shown in 4a, 4d and 8.
5. Point values are shown next to the problem number.
6. Budget your time for each of the questions. Some problems may have a low point value yet be very challenging. If you do not recognize the solution to a question quickly, skip it, and return to the question after completing the easier problems.
7. Look through the exam before beginning; plan your work; then begin.
8. ~~Relax~~ and do well.

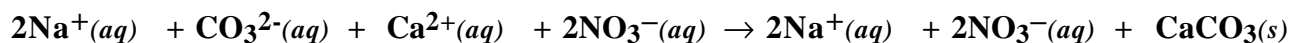
	Page 2	Page 3	Page 4	Page 5	TOTAL
SCORES	_____	_____	_____	_____	_____
	(23)	(35)	(18)	(24)	(100)

- (9) 1. Write the chemical formula(s) of the product(s) and balance the following reactions. Identify all products phases as either (g)as, (l)iquid, (s)olid or (aq)ueous. Soluble ionic compounds should be written in the form of their component ions.

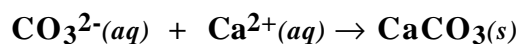


- (4) 2a. Write the ionic and net ionic chemical equation for 1a) or 1b).

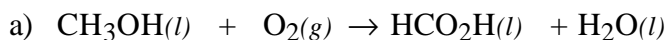
Ionic equation



Net Ionic equation



- (10) 3. Predict whether the entropy change in the system is positive or negative for each of the following processes.

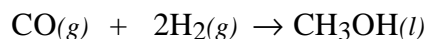


The phases are both liquid, while the reactants are a liquid and a gas. There are equal numbers of moles, and the liquid phase is more ordered compared to the gas phase, so the products are more ordered compared to the reactants and ΔS is $-$.



The reactants and products are all in the same phase, but there are more moles of products compared to reactants. So there are more possible arrangements in the products compared to the reactants and ΔS is $+$.

4. Methanol, CH₃OH, can be synthesized by reacting hydrogen gas with carbon monoxide, according to the chemical equation,



- (18) a) Calculate $\Delta H^\circ_{\text{rxn}}$, $\Delta S^\circ_{\text{rxn}}$ and $\Delta G^\circ_{\text{rxn}}$.

	$\text{CO}(g)$	+	$2\text{H}_2(g)$	→	$\text{CH}_3\text{OH}(l)$
$\Delta H_f^\circ \left(\frac{\text{kJ}}{\text{mol}} \right)$	-110.5		-0		-239
$S^\circ \left(\frac{\text{J}}{\text{K}\cdot\text{mol}} \right)$	198		131		127
$\Delta G_f^\circ \left(\frac{\text{kJ}}{\text{mol}} \right)$	-137		-0		-166

$$\begin{aligned} \Delta H_{\text{rxn}}^\circ &= \sum \Delta H_f^\circ (\text{products}) - \sum \Delta H_f^\circ (\text{reactants}) \\ &= [\Delta H_f^\circ (\text{CH}_3\text{OH}(l))] - [\Delta H_f^\circ (\text{CO}(g)) + 2\Delta H_f^\circ (\text{H}_2(g))] \\ &= (-239 \text{ kJ}) - [(-110.5 \text{ kJ}) + 2(0 \text{ kJ})] = -128 \text{ kJ} \end{aligned}$$

$$\begin{aligned} \Delta S_{\text{rxn}}^\circ &= \sum S^\circ (\text{products}) - \sum S^\circ (\text{reactants}) \\ &= [S^\circ (\text{CH}_3\text{OH}(l))] - [S^\circ (\text{CO}(g)) + 2S^\circ (\text{H}_2(g))] \\ &= \left[\left(127 \frac{\text{J}}{\text{K}} \right) \right] - \left[\left(198 \frac{\text{J}}{\text{K}} \right) + 2 \left(131 \frac{\text{J}}{\text{K}} \right) \right] = -333 \text{ J/K} \end{aligned}$$

$$\begin{aligned} \Delta G_{\text{rxn}}^\circ &= \sum \Delta G_f^\circ (\text{products}) - \sum \Delta G_f^\circ (\text{reactants}) \\ &= [\Delta G_f^\circ (\text{CH}_3\text{OH}(l))] - [\Delta G_f^\circ (\text{CO}(g)) + 2\Delta G_f^\circ (\text{H}_2(g))] \\ &= (-166 \text{ kJ}) - [(-137 \text{ kJ}) + 2(0 \text{ kJ})] = -29 \text{ kJ} \end{aligned}$$

- (4) b) Which factor, the change in enthalpy, ΔH° , or the change in entropy, ΔS° , provides the principal driving force for the reaction at 298 K? Explain.

ΔH is the principal driving force, because the sign of ΔH in the reaction is negative. A $-\Delta H$ favors spontaneity. ΔS is also negative. $-\Delta S$ does not favor spontaneity.

Since $\Delta G = \Delta H - T\Delta S$ a spontaneous reaction has a negative ΔG . In this equation spontaneity is favored by a $-\Delta H$ and a $+\Delta S$. In the reaction in part a only the sign of ΔH favors spontaneity.

- (5) c) For the reaction, how is the value of the standard free energy change, ΔG° , affected by an increase in temperature? Explain.

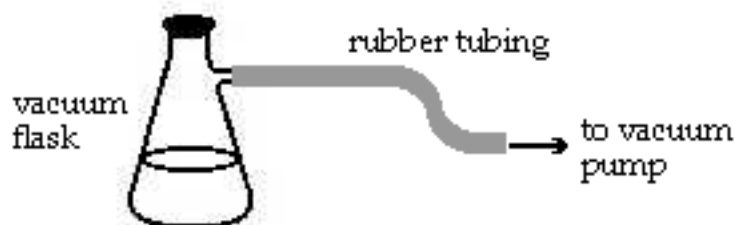
Using the equation $\Delta G = \Delta H - T\Delta S$, since ΔH is negative and ΔS is negative as the temperature increases the $T\Delta S$ term is becoming more negative, and $-T\Delta S$ is more positive, making ΔG° more positive and less spontaneous.

- (8) d) Calculate the ΔS°_f for CH₃OH(l).

	$\text{C}(s)$	+	$2\text{H}_2(g)$	+	$\frac{1}{2}\text{O}_2(g)$	→	$\text{CH}_3\text{OH}(l)$
$S^\circ \left(\frac{\text{J}}{\text{K}\cdot\text{mol}} \right)$	6		131		205		127

$$\begin{aligned} \Delta S_{\text{rxn}}^\circ &= \sum S^\circ (\text{products}) - \sum S^\circ (\text{reactants}) \\ &= [S^\circ (\text{CH}_3\text{OH}(l))] - [S^\circ (\text{C}(s)) + 2S^\circ (\text{H}_2(g)) + \frac{1}{2} S^\circ (\text{O}_2(g))] \\ &= \left[\left(127 \frac{\text{J}}{\text{K}} \right) \right] - \left[\left(6 \frac{\text{J}}{\text{K}} \right) + 2 \left(131 \frac{\text{J}}{\text{K}} \right) + \frac{1}{2} \left(205 \frac{\text{J}}{\text{K}} \right) \right] = -243 \text{ J/K} \end{aligned}$$

(12) 5. A sample of water is in the vacuum flask shown below.



The vacuum flask side arm has vacuum rubber tubing with one end attached to the flask and the other end attached to a vacuum pump (can not see in this diagram). When the vacuum pump is on any gas/vapor in the flask is removed.

In an experiment the water placed into the flask initially is at room temperature. The rubber tubing is attached to the flask and the vacuum pump, and the pump is turned on. After the pump has been on for several minutes the following two observations are made:

1. the volume of water in the flask has decreased;
2. what water remains has turned to ice.

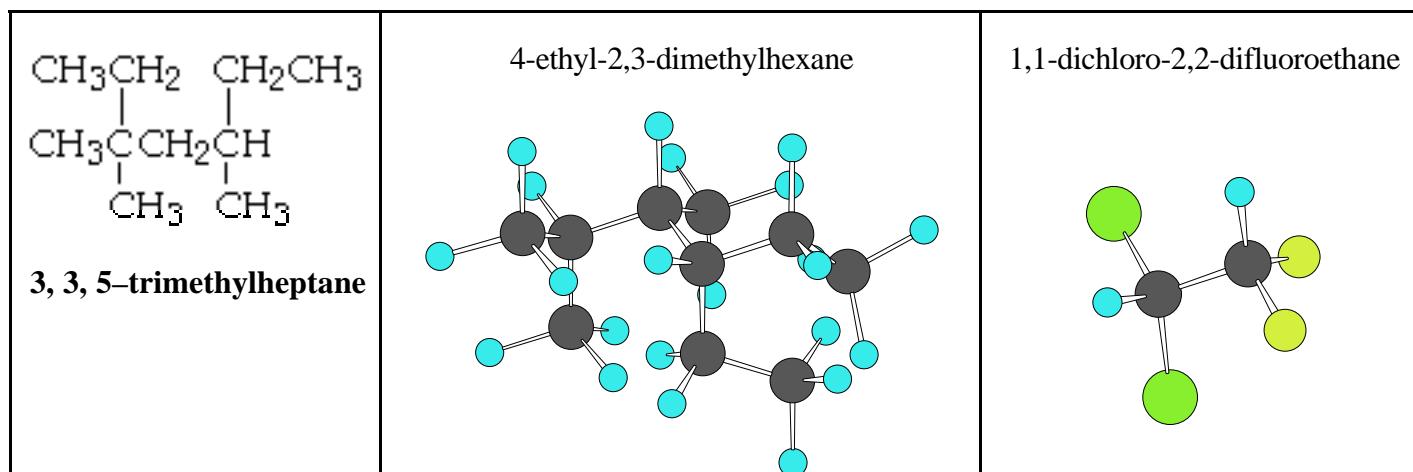
Explain both of these observations.

The volume of liquid water decreases because as the vacuum pump removes vapor from above the liquid, particles escape the liquid phase to maintain the equilibrium vapor pressure. Since particles are evaporating and few particles are condensing, the volume of the liquid will decrease.

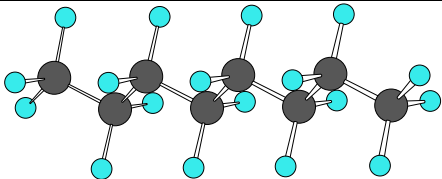
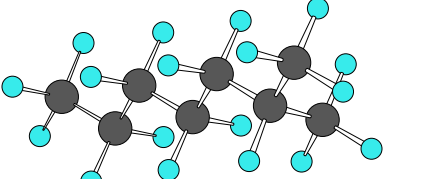
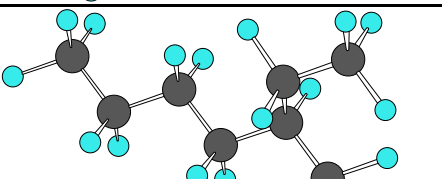
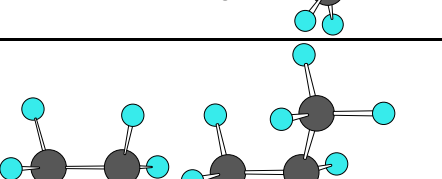
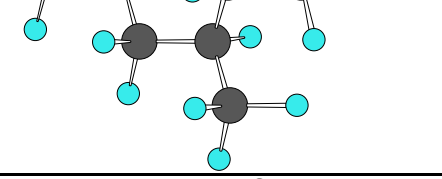
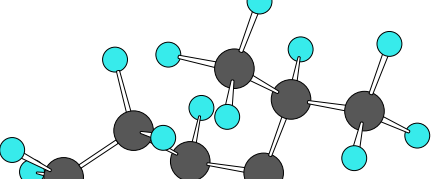
The reason the liquid eventually turns to solid is because evaporation is an endothermic process, and heat is removed from the system (the liquid water). When enough heat is removed the liquid freezes.

Evaporation is endothermic because the conversion of water in the liquid phase to water in the vapor phase requires breaking hydrogen-bonding forces. Breaking bonds is always an endothermic process.

(6) 6. Give the name or draw the complete Lewis structure (showing all C-H(X) bonds) for each of the following compounds.



(12) 7. Draw and name six different structural isomers for C_8H_{18} . (NOTE: You may use condensed formulas when representing the different structural isomers.)

	<i>n</i>-octane
	2-methylheptane
	3-methylheptane
	4-methylheptane
	2,3-dimethylhexane
	3,3-dimethylhexane

- (12) 8. CH_3I has a vapor pressure of 400 mm Hg at 25.3 °C. Calculate the temperature that CH_3I has a vapor pressure of 40.0 mm Hg. $\Delta H^\circ_{\text{vap}}$ for CH_3I is 29.2 kJ mol⁻¹.

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

$$\ln \frac{400}{40} = \frac{-29200 \frac{\text{J}}{\text{mol}}}{8.314 \frac{\text{J}}{\text{mol} \cdot \text{K}}} \left(\frac{1}{298.3 \text{ K}} - \frac{1}{T_1} \right)$$

$$\ln (10) = -3512 \text{ K} \left(3.35 \times 10^{-3} - \frac{1}{T_1} \right)$$

$$2.303 = -3512 \text{ K} \left(3.35 \times 10^{-3} - \frac{1}{T_1} \right)$$

$$-6.56 \times 10^{-4} = 3.35 \times 10^{-3} - \frac{1}{T_1}$$

$$\frac{1}{T_1} = 4.00 \times 10^{-3} \qquad T_1 = 250 \text{ K}$$

Periodic Table of the Elements

	IA																VIII A	
1	1 H 1.008																	2 He 4.00
2	3 Li 6.94	4 Be 9.01										5 B 10.81	6 C 12.01	7 N 14.01	8 O 16.00	9 F 19.00	10 Ne 20.18	
3	11 Na 22.99	12 Mg 24.30										13 Al 26.98	14 Si 28.09	15 P 30.97	16 S 32.06	17 Cl 35.45	18 Ar 39.95	
4	19 K 39.10	20 Ca 40.08	21 Sc 44.96	22 Ti 47.88	23 V 50.94	24 Cr 52.00	25 Mn 54.94	26 Fe 55.85	27 Co 58.93	28 Ni 58.69	29 Cu 63.55	30 Zn 65.38	31 Ga 69.72	32 Ge 72.59	33 As 74.92	34 Se 78.96	35 Br 79.90	36 Kr 83.80
5	37 Rb 85.47	38 Sr 87.62	39 Y 88.91	40 Zr 91.22	41 Nb 92.91	42 Mo 95.94	43 Tc (98)	44 Ru 101.1	45 Rh 102.9	46 Pd 106.4	47 Ag 107.9	48 Cd 112.4	49 In 114.8	50 Sn 118.7	51 Sb 121.8	52 Te 127.6	53 I 126.9	54 Xe 131.3
6	55 Cs 132.9	56 Ba 137.3	57 La 138.9	72 Hf 178.5	73 Ta 180.9	74 W 183.8	75 Re 186.2	76 Os 190.2	77 Ir 192.2	78 Pt 195.1	79 Au 197.0	80 Hg 200.6	81 Tl 204.4	82 Pb 207.2	83 Bi 209.0	84 Po (209)	85 At (210)	86 Rn (222)
7	87 Fr (223)	88 Ra 226.0	89 Ac 227.0	104 (261)	105 (262)	106 (263)												

Lanthanides	58 Ce 140.1	59 Pr 140.9	60 Nd 144.2	61 Pm (145)	62 Sm 150.4	63 Eu 152.0	64 Gd 157.2	65 Tb 158.9	66 Dy 162.5	67 Ho 164.9	68 Er 167.3	69 Tm 168.9	70 Yb 173.0	71 Lu 175.0
Actinides	90 Th 232.0	91 Pa 231.0	92 U 238.0	93 Np 237.0	94 Pu (244)	95 Am (243)	96 Cm (247)	97 Bk (247)	98 Cf (251)	99 Es (252)	100 Fm (257)	101 Md (258)	102 No (259)	103 Lr (260)

Useful Information

$$PV = nRT$$

$$\ln\left(\frac{vp_2}{vp_1}\right) = -\frac{\Delta H^\circ_{\text{vap}}}{R}\left(\frac{1}{T_2} - \frac{1}{T_1}\right)$$

$$\text{density of H}_2\text{O} = 1.00 \frac{\text{g}}{\text{cm}^3}$$

$$\Delta H^\circ_{\text{rxn}} = \sum n(\Delta H_f^\circ(\text{products})) - \sum m(\Delta H_f^\circ(\text{reactants}))$$

$$\Delta S^\circ_{\text{rxn}} = \sum n(S^\circ(\text{products})) - \sum m(S^\circ(\text{reactants}))$$

$$\Delta G^\circ_{\text{rxn}} = \sum n(\Delta G_f^\circ(\text{products})) - \sum m(\Delta G_f^\circ(\text{reactants}))$$

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

$$R = 0.0821 \frac{\text{L}\cdot\text{atm}}{\text{mol}\cdot\text{K}} = 8.314 \frac{\text{J}}{\text{mol}\cdot\text{K}}$$

$$\text{density of H}_2\text{O} = 1.00 \frac{\text{g}}{\text{cm}^3}$$

Temperature (°C)	Vapor Pressure(mmHg)	Temperature (°C)	Vapor Pressure(mmHg)
-5	3.2	50	92.5
0	4.6	55	118.0
5	6.52	60	149.4
10	9.20	65	187.5
15	12.8	70	233.7
20	17.5	75	289.1
25	23.8	80	355.1
30	31.8	85	433.6
35	42.1	90	525.8
40	55.3	95	633.9
45	71.9	100	760

Solubility Table

<u>Ion</u>	<u>Solubility</u>	<u>Exceptions</u>
NO ₃ ⁻	soluble	none
ClO ₄ ⁻	soluble	none
Cl ⁻	soluble	except Ag ⁺ , Hg ₂ ²⁺ , *Pb ²⁺
I ⁻	soluble	except Ag ⁺ , Hg ₂ ²⁺ , Pb ²⁺
SO ₄ ²⁻	soluble	except Ca ²⁺ , Ba ²⁺ , Sr ²⁺ , Hg ²⁺ , Pb ²⁺ , Ag ⁺
CO ₃ ²⁻	insoluble	except Group IA and NH ₄ ⁺
PO ₄ ³⁻	insoluble	except Group IA and NH ₄ ⁺
-OH	insoluble	except Group IA, *Ca ²⁺ , Ba ²⁺ , Sr ²⁺
S ²⁻	insoluble	except Group IA, IIA and NH ₄ ⁺
Na ⁺	soluble	none
NH ₄ ⁺	soluble	none
K ⁺	soluble	none

*slightly soluble

Thermodynamic Values (25 °C)

Substance and State	ΔH_f° ($\frac{\text{kJ}}{\text{mol}}$)	ΔG_f° ($\frac{\text{kJ}}{\text{mol}}$)	S° ($\frac{\text{J}}{\text{K}\cdot\text{mol}}$)	Substance and State	ΔH_f° ($\frac{\text{kJ}}{\text{mol}}$)	ΔG_f° ($\frac{\text{kJ}}{\text{mol}}$)	S° ($\frac{\text{J}}{\text{K}\cdot\text{mol}}$)
Carbon				Oxygen			
C(s) (graphite)	0	0	6	O ₂ (g)	0	0	205
C(s) (diamond)	2	3	2	O(g) 249	232	161	
CO(g)	-110.5	-137	198	O ₃ (g)	143	163	239
CO ₂ (g)	-393.5	-394	214	Nitrogen			
CH ₄ (g)	?	-51	186	N ₂ (g)	0	0	192
CH ₃ OH(g)	-201	-163	240	NCl ₃ (g)	230	271	-137
CH ₃ OH(l)	-239	-166	127	NF ₃ (g)	-125	-83.6	-139
CH ₃ Cl(g)	-80.8	-57.4	234	NH ₃ (g)	?	-17	193
CHCl ₃ (g)	-100.8			NH ₃ (aq)	?	-27	111
CHCl ₃ (l)	-131.8			NH ₂ CONH ₂ (aq)	?	?	174
H ₂ CO(g)	-116	-110	219	NO(g)	90	87	211
HCOOH(g)	-363	-351	249	NO ₂ (g)	32	52	240
HCN(g)	135.1	125	202	N ₂ O(g)	82	104	220
C ₂ H ₂ (g)	227	209	201	N ₂ O ₄ (g)	10	98	304
C ₂ H ₄ (g)	52	68	219	N ₂ O ₅ (g)	-42	134	178
CH ₃ CHO(g)	-166	-129	250	HNO ₃ (aq)	-207	-111	146
C ₂ H ₅ OH(l)	-278	-175	161	HNO ₃ (l)	-174	-81	156
C ₂ H ₆ (g)	-84.7	-32.9	229.5	NH ₄ Cl(s)	-314	-201	95
C ₃ H ₆ (g)	20.9	62.7	266.9	NH ₄ ClO ₄ (s)	-295	-89	186
C ₃ H ₈ (g)	-104	-24	270	Silver			
Bromine				Ag(s)	0	0	42.6
Br ₂ (l)	0	0	152.	Ag ⁺ (aq)	105.6	77.1	72.7
BrCl(g)	14.64	-0.96	240	Ag(S ₂ O ₃) ³⁻ (aq)	-1285.7	--	--
Chlorine				AgBr(s)	-100.4	-96.9	107.1
Cl ₂ (g)	0	0	223	AgCl(s)	-127.1	-109.8	96.2
Cl ₂ (aq)	-23	7	121	Sulfur			
Cl ⁻ (aq)	-167	-131	57	S(rhombic)	0	0	31.8
HCl(g)	-92	-95	187	SO ₂ (g)	-296.8	-300.2	248.8
Fluorine				SO ₃ (g)	-395.7	-371.1	256.3
F ₂ (g)	0	0	203	H ₂ S(g)	-20.17	-33.0	205.6
F ⁻ (aq)	-333	-279	-14	Titanium			
HF(g)	-271	-273	174	TiCl ₄ (g)	-763	-727	355
Hydrogen				TiO ₂ (s)	-945	-890	50
H ₂ (g)	0	0	131	Aluminum			
H(g) 217	203	115		AlCl ₃ (s)	-526	-505	184
H ⁺ (aq)	0	0	0	Barium			
OH ⁻ (aq)	-230	-157	-11	BaCl ₂ (aq)	-872	-823	123
H ₂ O(l)				Ba(OH) ₂ ·8H ₂ O(s)	-3342	-2793	427
H ₂ O(g)	-242	-229	189	Iodine			
Magnesium				I ₂ (s)	0	0	116.7
Mg(s)	0	0	33	HI(g)	25.94	1.30	206.3
Mg(aq)	-492	-456	-118				
MgO(s)	-601	-569	26.9				

