

CHEM 1515.001 - 006
Exam V
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May 6, 2002

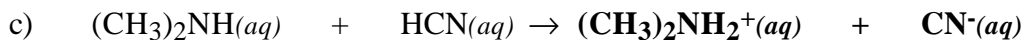
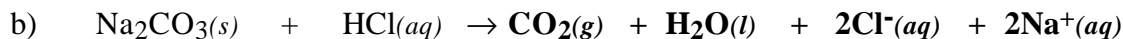
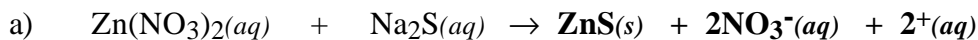
Name _____
TA's Name _____
Section _____

INSTRUCTIONS:

1. This examination consists of a total of 11 different pages. The last four pages include a periodic table; many useful equations and constants; a table of vapor pressures for water; a solubility table; a table of equilibrium constants for acids and bases; 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 6.
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.

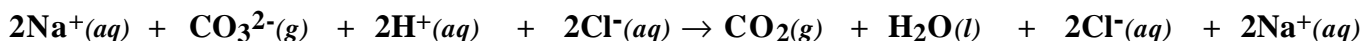
	Page 2	Page 3	Page 4	Page 5	Page 6	Page 7	TOTAL
SCORES	<u> </u> (18)	<u> </u> (16)	<u> </u> (16)	<u> </u> (18)	<u> </u> (20)	<u> </u> (12)	<u> </u> (100)

- (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.

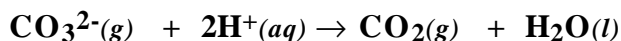


- (4) 2a. Write the ionic and net ionic chemical equation for 1a), 1b) or 1c).

Ionic equation



Net Ionic equation

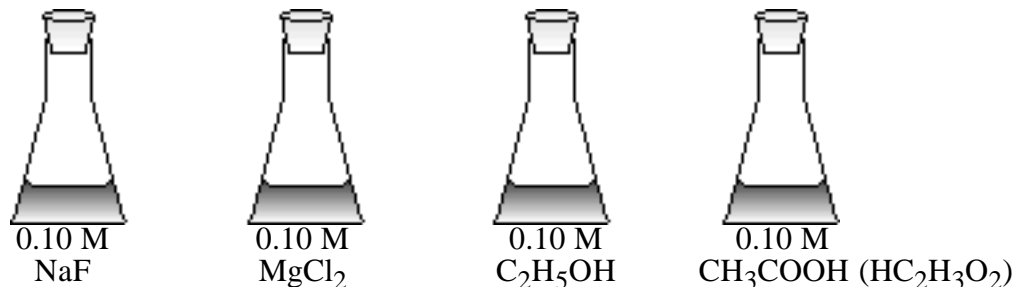


- (5) 3. Account for the following observations about NH_3 and NF_3 . In your answer, use appropriate principles of intermolecular forces. Your answer must include reference to both substances.

- a) NH_3 has a normal boiling point of -33.4°C where as NF_3 has a boiling point of -128.8°C .

NH_3 is polar and can hydrogen bond. Hydrogen-bonding is a strong intermolecular attractive force and causes the boiling point of NH_3 to be high. NF_3 is polar but can not hydrogen bond. The dipole-dipole forces are not as strong an intermolecular attractive force so the boiling point of NF_3 is much lower compared to NH_3 .

(16) 4.



Answer the following questions, which refer to the 100 mL samples of aqueous solutions at 25 °C in the stoppered flasks shown above

(a) Which solution has the lowest electrical conductivity? Explain.

The lowest electrical conductivity has the fewest ions... C₂H₅OH. All other substances above form ions in solution.

(b) Which solution has the lowest freezing point? Explain.

MgCl₂ has the lowest freezing point. Since MgCl₂ produces the most ions in solution with an *i* value of 3, ΔT_f will be the largest.

(c) Above which solution is the pressure of water vapor the greatest? Explain.

C₂H₅OH does not ionize, so there are the fewest particles in solution so it will have the higher vapor pressure.

(d) Which solution has the highest pH? Explain.

NaF will have the highest pH. F⁻ is a weak base.

(16) 5a. Define the term 'equilibrium vapor pressure'.

- (b) If a sample of a liquid is injected into an evacuated container, describe what phase change occurs. (Note the amount of the liquid injected is small compared to the volume of the container.)

Since the container is evacuated when some liquid is introduced, some of the liquid will immediately vaporize until the equilibrium vapor pressure of the liquid is reached, or until all of the liquid evaporates. So if the equilibrium vapor pressure is reached, both liquid and vapor are present. Otherwise only vapor.

- (c) Is the phase change exothermic or endothermic? Explain?

Evaporation is endothermic, because to escape from the liquid phase to the vapor phase intermolecular attractive forces must be overcome. Although weak these forces must be broken for particles to enter the vapor phase.

- (d) In terms of the 'pressure exerted by the vapor' and the 'equilibrium vapor pressure' how do we determine what phase(s) are present in the container after adding the sample?

If the pressure due to the vapor is equal to the equilibrium vapor pressure both liquid and vapor must be present. If the pressure due to the vapor is less than the equilibrium vapor pressure only vapor is present. The pressure due to the vapor can never exceed the equilibrium vapor pressure.

- (18) 6. A commercial aqueous solution of ammonia is 28% NH_3 by mass and has a density of 0.900 g mL^{-1} .
(a) Calculate the molarity of the solution.

Assume 100 gram of solution

28 g of NH_3 and 72 grams of water. Convert the mass of NH_3 to moles, then divide the 100 gram of solution by the density to get mLs of solution. Convert mLs to liters and divide into the moles for the molarity

Answer is 14.8 M

- (b) Calculate the molality of the solution.

Use the moles determined in part a and divide by the mass of water in kilograms.

Answer is 22.9 molal

- (c) Describe how you would prepare 1.00 L of a 0.400 M NH_3 solution from the more concentrated solution in part a).

Using the dilution equation $M_c \cdot V_c = M_d \cdot V_d$

The volume of concentrated ammonia needed is 27.0 mL.

Add 27 mL of NH_3 to about 900 mL of water, be sure all the NH_3 is dissolved then add enough water to reach a final volume of 1000 mLs.

- (10) (d) Calculate the pH of the solution prepared in part c)

The pH of the solution is 11.43. This is a simple weak base calculation where the initial concentration of NH_3 is 0.400 M

- (10) (e) Calculate the pH of the solution prepared by adding 0.250 mol of solid NH_4Cl to the solution prepared in part c).

This is a common ion/buffer solution. The pH is 9.45.

- (12) 7. For the gaseous equilibrium represented below, it is observed that greater amounts of PCl_3 and Cl_2 are produced as the temperature is increased.



- (a) What is the sign of ΔS° for the reaction? Explain.

ΔS° is positive. In the balanced equation there are more moles of gas in the products compared to the reactants. So the products are more disordered.

- (b) What change, if any, will occur in ΔG° for the reaction as the temperature is increased? Explain your reasoning in terms of thermodynamic principles.

The reaction is endothermic because , ", it is observed that greater amounts of PCl_3 and Cl_2 are produced as the temperature is increased ". So if ΔH° is positive and ΔS° is positive ΔG° will become more negative with increasing temperature. The equation $\Delta G^\circ = \Delta H^\circ - T\Delta S^\circ$ means the reaction will become more spontaneous as T increases. The $T\Delta S^\circ$ term will eventually be greater than ΔH° and the reaction will be spontaneous.

- (c) If the volume of the reaction mixture is decreased at constant temperature to half the original volume, what will happen to the number of moles of Cl_2 in the reaction vessel? Explain.

If the volume is decreased the pressure will increase in the container and the reaction will proceed in a direction to relieve the pressure, so it will go from right to left and the amount of Cl_2 will decrease.

Q is inversely related to the volume, so as volume decreases Q increases. To re-establish equilibrium the reaction must go from R to L and amount of Cl_2 will decrease.

Periodic Table of the Elements

	IA															VIIIA		
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
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)

Actinides

Useful Information

$$PV = nRT \quad R = 0.0821 \frac{\text{L}\cdot\text{atm}}{\text{mol}\cdot\text{K}} = 8.314 \frac{\text{J}}{\text{mol}\cdot\text{K}} \quad 6.02 \times 10^{23}$$

$$\ln\left(\frac{vp_2}{vp_1}\right) = -\frac{\Delta H^\circ_{\text{vap}}}{R} \left(\frac{1}{T_2} - \frac{1}{T_1}\right) \quad \text{density of H}_2\text{O} = 1.00 \frac{\text{g}}{\text{cm}^3}$$

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

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

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

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

$$\text{pH} = -\log[\text{H}^+]$$

$$\text{pH} + \text{pOH} = 14$$

$$\text{pOH} = -\log[\text{OH}^-]$$

$$K_w = 1.00 \times 10^{-14}$$

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

$$K_p = K_c(RT)^{\Delta n}$$

$$x_{1,2} = \frac{-b \pm \sqrt{b^2 - 4ac}}{2a} \quad \text{for } ax^2 + bx + c = 0$$

$$1 \text{ pm} = 10^{-12} \text{ m}$$

$$P_{\text{solution}} = \chi_{\text{solvent}} P^{\circ}_{\text{solvent}}$$

$$\Delta T = i k m \quad k_f(\text{H}_2\text{O}) = 1.86 \frac{^{\circ}\text{C}}{\text{m}} \quad k_b(\text{H}_2\text{O}) = 0.512 \frac{^{\circ}\text{C}}{\text{m}}$$

$$\text{edge length (l)} = 2r$$

$$\text{edge length (l)} = 2\sqrt{2} \cdot r$$

$$\text{edge length (l)} = \frac{4r}{\sqrt{3}}$$

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

Ion	Solubility	Exceptions
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

Name	Formula	K_{a1}	K_{a2}	K_{a3}
Acetic	$\text{HC}_2\text{H}_3\text{O}_2$	1.8×10^{-5}		
Ascorbic	$\text{HC}_6\text{H}_7\text{O}_6$	8.0×10^{-3}		
Arsenic	H_3AsO_4	5.6×10^{-3}	1.0×10^{-7}	3.0×10^{-12}
Arsenous	H_3AsO_3	6.0×10^{-10}		
Benzoic	$\text{HC}_7\text{H}_5\text{O}_2$	6.5×10^{-5}		
Boric	H_3BO_3	5.8×10^{-10}		
Butyric acid	$\text{HC}_4\text{H}_7\text{O}_2$	1.5×10^{-5}		
Carbonic	H_2CO_3	4.3×10^{-7}	5.6×10^{-11}	
Cyanic	HCNO	3.5×10^{-4}		
Citric	$\text{H}_3\text{C}_6\text{H}_5\text{O}_7$	7.4×10^{-4}	1.7×10^{-5}	4.0×10^{-7}
Formic	HCHO_2	1.8×10^{-4}		
Hydroazoic	HN_3	1.9×10^{-5}		
Hydrocyanic	HCN	4.9×10^{-10}		
Hydrofluoric	HF	7.2×10^{-4}		
Hydrogen chromate ion	HCrO_4^-	3.0×10^{-7}		
Hydrogen peroxide	H_2O_2	2.4×10^{-12}		
Hydrogen selenate ion	HSeO_4^-	2.2×10^{-2}		
Hydrogen sulfate ion	HSO_4^-	1.2×10^{-2}		
Hydrogen sulfide	H_2S	5.7×10^{-8}	1.3×10^{-13}	
Hypobromous	HBrO	2.0×10^{-9}		
Hypochlorous	HClO	3.0×10^{-8}		
Hypoiodous	HIO	2.0×10^{-11}		
Iodic	HIO_3	1.7×10^{-1}		
Lactic	$\text{HC}_3\text{H}_5\text{O}_3$	1.4×10^{-4}		
Malonic	$\text{H}_2\text{C}_3\text{H}_2\text{O}_4$	1.5×10^{-3}	2.0×10^{-6}	
Oxalic	$\text{H}_2\text{C}_2\text{O}_4$	5.9×10^{-2}	6.4×10^{-5}	
Nitrous	HNO_2	4.5×10^{-4}		
Phenol	$\text{HC}_6\text{H}_5\text{O}$	1.3×10^{-10}		
Phosphoric	H_3PO_4	7.5×10^{-3}	6.2×10^{-8}	4.2×10^{-13}
Paraperiodic	H_5IO_6	2.8×10^{-2}	5.3×10^{-9}	
Propionic	$\text{HC}_3\text{H}_5\text{O}_2$	1.3×10^{-5}		
Pyrophosphoric	$\text{H}_4\text{P}_2\text{O}_7$	3.0×10^{-2}	4.4×10^{-3}	
Selenous	H_2SeO_3	2.3×10^{-3}	5.3×10^{-9}	
Sulfuric	H_2SO_4	strong acid	1.2×10^{-2}	
Sulfurous	H_2SO_3	1.7×10^{-2}	6.4×10^{-8}	
Tartaric	$\text{H}_2\text{C}_4\text{H}_4\text{O}_6$	1.0×10^{-3}	4.6×10^{-5}	

E.2 DISSOCIATION CONSTANTS FOR BASES AT 25°C

Name	Formula	K_b	Name	Formula	K_b
Ammonia	NH_3	1.8×10^{-5}	Hydroxylamine	HONH_2	1.1×10^{-8}
Aniline	$\text{C}_6\text{H}_5\text{NH}_2$	4.3×10^{-10}	Methylamine	CH_3NH_2	4.4×10^{-4}
Dimethylamine	$(\text{CH}_3)_2\text{NH}$	5.4×10^{-4}	Pyridine	$\text{C}_5\text{H}_5\text{N}$	1.7×10^{-9}
Ethylamine	$\text{C}_2\text{H}_5\text{NH}_2$	6.4×10^{-4}	Trimethylamine	$(\text{CH}_3)_3\text{N}$	6.4×10^{-5}
Hydrazine	H_2NNH_2	1.3×10^{-6}			

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				