# EXAM INSTRUCTIONS/SUGGESTIONS. READ THIS CAREFULLY!

YOU MAY OMIT TWO (2) OF THE 16 QUESTIONS ON THIS EXAM. EACH QUESTION IS WORTH 10 POINTS. IF YOU ANSWER MORE THAN 14 QUESTIONS, ONLY THE FIRST 14 ANSWERED WILL BE COUNTED.

INDICATE ON THE NEXT PAGE WHICH QUESTIONS ARE NOT TO BE GRADED BY WRITING <u>DNG</u> (DO NOT GRADE) NEXT TO PROBLEM NUMBER.

YOU CAN MAKE ALL REASONABLE APPROXIMATIONS IN DOING THE pH AND SOLUBILITY PROBLEMS UNLESS AN APPROXIMATION IS NOT NECESSARY.

NO PARTIAL CREDIT on any question except where indicated by the statement SHOW WORK. When work is requested, set up equations (with numbers substituted in appropriate units) in space provided, but do the calculations on scrap sheet.

IF QUESTION STATES "SHOW WORK" AND YOU <u>GUESS</u> CORRECTLY WITHOUT SHOWING WORK - YOU GET 1/2 CREDIT.

CHECK FRONT BLACKBOARD FOR CORRECTIONS/CHANGES.

DO THE SIMPLER QUESTIONS FIRST.

IF ANY QUESTION IS NOT CLEAR - ASK TANUJA OR ME ABOUT IT!

CONSTANTS AND CONVERSION FACTORS ARE ON THE PAGE AT THE END OF THE EXAM. PLEASE LOOK THERE!

LAST THREE PAGES ARE FOR SCRAP WORK.

FEEL FREE TO TEAR THESE PAGES OFF.

PLEASE REMOVE THIS SHEET PRIOR TO STARTING EXAM.

CHEMISTRY S14O4 PROFESSOR J. MORROW	FIRST EXAM	7/15/99
PRINT NAME, LAST:		
FIRST:		
	ESTION IS WORTH 10	
1	7	12
2	8	13
3	9	14
4	10	15
5	11	16
6		
COLUMN TOTALS: (MAXI	MUM)	
(60)	(50)	(50)
EXAM TOTAL (140 pts)		
	JO	JT OF 100

#### THIS INFORMATION IS FOR PROBLEMS 1 AND 2.

You are given a triprotic acid,  $H_3X$ , with  $K_{a1} = 1.00 \times 10^{-7}$ ,  $K_{a2} = 1.00 \times 10^{-12}$ 

$$\text{and } K_{a3} = 1.00 \times 10^{\text{-}16} \; . \; \; \kappa_{a1} = \; \frac{[\text{H}^+][\text{H}_2 \text{X}^-]}{[\text{H}_3 \text{X}]} \; \; , \quad \; \kappa_{a2} = \; \frac{[\text{H}^+][\text{HX}^{2^-}]}{[\text{H}_2 \text{X}^-]} \; \; , \quad \; \kappa_{a3} = \; \frac{[\text{H}^+][\text{X}^{3^-}]}{[\text{HX}^{2^-}]} \; .$$

1) Calculate the ratio  $\frac{[HX^{2-}]}{[H_2X^-]}$  present in a solution whose pH = 4.00 if

 $H_3X$  is the only solute.

- i)  $1.0x10^{-3}$  ii)  $1.0x10^{-12}$  iii)  $1.0x10^{-8}$  iv)  $1.0x10^{-10}$

ANSWER IS: \_\_\_\_\_

- 2) Calculate the  $[H_3X]$  in the above solution from problem 1.
  - i) 0.30
- ii) 0.25
- iii) 0.15
- iv) 0.10
- v) 0.050

ANSWER IS: \_\_\_\_\_

#### THE FOLLOWING INFORMATION IS FOR PROBLEMS 3 AND 4.

Given a sparingly soluble salt, M<sub>3</sub>X<sub>2</sub>, where X is the anion from the weak triprotic acid given in problem 1.  $(K_{a1} = 1.00x10^{-7}, K_{a2} = 1.00x10^{-12}, and$ and  $K_{a3} = 1.00 \times 10^{-16}$ ).

For this sparingly soluble salt,  $M_3X_2$ ,  $K_{sp} = 1.00x10^{-44}$ .

EACH QUESTION IS INDEPENDENT OF THE OTHER. **SHOW WORK** 

				AN	NSWER IS:
			in a solution co	ntaining O.C	010 M
Na <sub>3</sub> X	? Neglect the hy	ydrolysis	of $X^{3-}$ .		
i) 1.55	5x10 <sup>-14</sup>	ii) 4.6	54x10 <sup>-14</sup> iii) 2.3	1x10 <sup>-14</sup>	iv) 0.77x10 <sup>-14</sup>
					ANSWER IS:
~			~		
Consid	er the weak acid	ls, A, B, <b>(</b>	C, and D;		
	ACID	A	В	С	D
	ACID pK <sub>a</sub>	A 8.3	B 2.7	<u>C</u> 12.9	D 5.6
List the			OK		
	se bases in orde	er of incr	OK easing acidity (	decreasing ba	asicity);
	se bases in orde	er of incr	OK easing acidity (	decreasing ba	
) A O.1	se bases in orde LEAST ACIDIO OO M solution	er of incre	OK easing acidity (	decreasing ba	asicity);  MOST ACIDIC s a pH = 9.00.
) A O.1 Calcul	se bases in orden LEAST ACIDIO OO M solution ate the value of	of a salt, $K_a$ for th	OK easing acidity (  < <  KX (which hyd	decreasing backers <pre> crolyzes), has County </pre>	asicity);  MOST ACIDIC s a pH = 9.00.
A O.1 Calcul	ese bases in order LEAST ACIDIC OO M solution at the value of on is, $X^2 + H_2$	of a salt, $K_{a} \text{ for th}$	OK easing acidity (	decreasing bands  < rolyzes), has  X. The hydro	asicity);  MOST ACIDIC s a pH = 9.00. olysis
) A O.1 Calcul reactio	ese bases in order LEAST ACIDIC OO M solution at the value of on is, $X^2 + H_2$	of a salt, $K_{a} \text{ for th}$	OK easing acidity (	decreasing bands  < rolyzes), has  X. The hydro	asicity);  MOST ACIDIC s a pH = 9.00. olysis
) A O.1 Calcul reactio	ese bases in order LEAST ACIDIC OO M solution at the value of on is, $X^2 + H_2$	of a salt, $K_{a} \text{ for th}$	OK easing acidity (	decreasing bands  < rolyzes), has  X. The hydro	asicity);  MOST ACIDIC s a pH = 9.00. olysis
) A O.1 Calcul reactio	ese bases in order LEAST ACIDIC OO M solution at the value of on is, $X^2 + H_2$	of a salt, $K_{a} \text{ for th}$	OK easing acidity (	decreasing bands  < rolyzes), has  X. The hydro	asicity);  MOST ACIDIC s a pH = 9.00. olysis

i) 1.58x1O<sup>-10</sup> ii) 6.21x1O<sup>-10</sup> iii) 18.6x1O<sup>-10</sup> iv) 12.4x1O<sup>-10</sup>

3) What is the solubility of  $M_3X_2$  in pure water. Neglect the

hydrolysis of  $X^{3-}$ .

### THE FOLLOWING INFORMATION IS FOR PROBLEMS 7, 8, AND 9.

- 25.00 mL of 0.100 M methylamine, CH<sub>3</sub>NH<sub>2</sub>, is titrated with 0.100 M HCl. Calculate the pH;  $(K_b = 4.4 \times 10^{-5})$
- 7) before any HCl is added.  $(CH_3NH_2 + H_2O \rightarrow CH_3NH_3^+ + OH^-)$ (SHOW WORK)

pH = \_\_\_\_\_

8) halfway to the equivalence point.

pH = \_\_\_\_\_

- 9) at the equivalence point. (HINT: First solve for  $K_h$ ) SHOW WORK
  - i) 2.68
- ii) 6.36 iii) 5.47
- iv) 10.95 v) 8.24

pH = \_\_\_\_\_

- 10) How many milliliters of a 0.100 M HX (a weak acid) solution should be added to 500 mL of O.100 M NaX solution, to produce a buffer of pH = 5.00 ? For HX,  $K_a = 4.00 \times 10^{-5}$  (SHOW WORK)
  - i) 125 ii) 250
- iii) 500
- iv) 750
- v) 1000

ANSWER IS: \_\_\_\_\_

11) Complete the following table. (for  $M(OH)_3$ ,  $K_{sp} = 6.Ox1O^{-2O}$ )

$[M^{3+}]$	[OH <sup>-</sup> ]
	1.Ox1O <sup>-4</sup> M
6.Ox1O <sup>-3</sup> M	

12) Given the following  $K_{sp}$  values: for  $M(OH)_3$ ,  $K_{sp(M)}=4.00 \times 10^{-14}$ ; for  $N(OH)_3$ ,  $K_{sp(N)}=1.00 \times 10^{-13}$ .

One mole of each of the above solids is placed in a beaker containing 1 (one) liter of pure water. These solids go into equilibrium with their ions. Calculate the pH of this solution. Do this calculation as accurately as possible. Use of the approximate method is worth 7 points. (SHOW WORK)

- i) 10.91
- ii) 10.87
- iii) 1O.29
- iv) 10.80

ANSWER IS: \_\_\_\_\_

13) Given the following reactions and their equilibrium constants:

1. 
$$HS^{-}$$
  $H^{+} + S^{2-}$   $K_{1}$ 
2.  $HCO_{3}^{-}$   $H^{+} + CO_{3}^{2-}$   $K_{2}$ 
3.  $PbS$   $Pb^{2+} + S^{2-}$   $K_{sp3}$ 
4.  $PbCO_{3}$   $Pb^{2+} + CO_{3}^{2-}$   $K_{sp4}$ 
5.  $H_{2}O$   $H^{+} + OH^{-}$   $K_{w}$ 

Calculate the value of K (in terms of any or all of the above K values) for the following reaction;

ANSWER IS: \_\_\_\_\_

14) A certain metal has a work function,  $\Phi$ , of 1.50 eV (= 2.403x10<sup>-19</sup> J). When monochromatic light of wavelength,  $\lambda$ , falls on this metal, photoelectrons are expelled. The kinetic energy of these photoelectrons is 2.64 eV? What is the wavelength,  $\lambda$  (in nm), of this light? SHOW WORK

- i) 255
- ii) 300
- iii) 334
- iv) 319

ANSWER IS: \_\_\_\_\_

15) An atom has a velocity of 50.0 $\frac{m}{sec}$ and a deBroglie wavelength of
1.991 Å. Which of the following is the element? (Hint:The atomic
weights in the Periodic Table are averages over the weights
of isotopes. To answer this question, just round off the numbers in
the Periodic Table to the nearest integer.) BE CAREFUL WITH UNITS!
(SHOW WORK)

i) H

ii) C

iii) Mg

iv) Ti v) Ca

ANSWER IS: \_\_\_\_\_

16) The linear velocity, v, of an electron in the hydrogen atom (according to the Bohr Theory) is,

$$v = \frac{2\delta e^2}{nh}$$

where: n = principle quantum number,

h = Plancks constant

 $m_e = electron mass,$ 

 $e = q = electron charge, \quad \eth = 3.14$ 

What is the de Broglie wavelength,  $\lambda$  (in Å), of an electron in the first shell of the H atom? (HINT: Use the cgs system) SHOW WORK

i) O.O332

ii) 384

iii) 3.32

iv) 6.64

v) 332

ANSWER IS: \_\_\_\_\_

# CONVERSION FACTORS and POTENTIALLY (BUT NOT NECESSARILY) USEFUL EQUATIONS

$$C = 3.00 \text{ x } 10^8 \text{ m/sec} = 3.00 \text{ x } 10^{10} \text{ cm/sec}$$

$$h = 6.626 \times 10^{-34} \text{ J·sec} = 6.626 \times 10^{-27} \text{ erg sec}$$

$$m_e = 9.108 \times 10^{-28} g = 9.108 \times 10^{-31} kg$$

$$\varepsilon = hv$$

$$N_A = 6.022 \times 10^{23}$$

$$R = O.O821 \quad \frac{L \cdot atm}{mol \cdot degK} \quad = \quad 82.1 \quad \frac{cm^3 \cdot atm}{mol \cdot degK}$$

$$\lambda \nu = c$$

$$1 \text{ Å} = 0.1 \text{ nm} = 10^{-8} \text{ cm} = 10^{-10} \text{ m}$$

$$\lambda = \frac{h}{p}$$
 (deBroglie equation) where  $p = mv$ 

Heisenberg Uncertainty principle:  $P = X = \frac{h}{4\delta}$  where p = mv

$$q = e = 1.6O22 \ x \ 10^{-19} \ coulombs = 4.8O6 \ x \ 10^{-1O} \ esu$$

$$1 \text{ eV} = 1.6022 \text{ x } 10^{-19} \text{ J}$$

$$\mathcal{E}_{n} = -\frac{2.179 \times 10^{-18}}{n^{2}} J$$

$$\epsilon = \mathsf{q} v = \mathsf{h} \nu$$
 -  $\Phi$ 

## **UNIT RELATIONSHIPS**

$$1 \text{ amp} = 1 \frac{\text{coulomb}}{\text{sec}} \qquad \qquad 1 \text{ volt·coulomb} = 1 \text{ Joule}$$

$$1 \text{ esu}^2 = 1 \text{ erg cm}$$
  $1 \text{ Watt} = 1 \frac{\text{joule}}{\text{sec}}$ 

1 Joule = 
$$1 \times 10^7$$
 erg 1 Joule =  $1 \frac{\text{kg m}^2}{\text{sec}^2}$  1 erg =  $1 \frac{\text{g cm}^2}{\text{sec}^2}$ 

# SCRAP WORK SHEET

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