Chemistry 277, Spring 2019	Your Name:
Prof. Jason Kahn	
University of Maryland, College Park	Your SID #:
General Chemistry and Energetics	
<u>Final Exam (100 points)</u>	Your Section # or time:

May 16, 2019

You have 50 minutes for this exam.

Explanations should be <u>concise</u> and <u>clear</u>. There is extra space on the last page if you need it. You will not need a calculator for this exam. No other study aids or materials are permitted. Generous partial credit will be given, *i.e.*, if you don't know, guess.

## **Useful Equations:**

4 Г 21		(2) $(2)$
$\sigma_{Y}^{2} = \lim_{N \to \infty} \frac{1}{N} \left[ \sum_{i} \left( Y_{i} - \overline{Y} \right)^{2} \right]$	$pH = -\log([H^+])$	$\sigma_Y^2 = \left(\frac{\partial Y}{\partial u}\right)^- \sigma_u^2 + \left(\frac{\partial Y}{\partial v}\right)^- \sigma_v^2 + \cdots$
$R = 0.08206 L \cdot atm/mole K$	$T^2 = 4\pi^2 a^3/GM$	$\ln K_{eq} = -\Delta H^{\circ}/(\mathbf{R}T) + \Delta S^{\circ}/\mathbf{R}$
R = 8.314  J/mole K = 1.987  c	cal/mole $K = N_A k_B$	$SEM = \frac{\sigma}{\sqrt{n}}$
$^{\circ}C = ^{\circ}K - 273.15$	$P(v)dv = Cv^2 exp(-mv^2/2kT)$	$\ln k = (-E_a/RT) + \ln A$
$pH = pK_a + \log([A^-]/[HA])$	$K_p = K_c(\mathbf{R}T)^{\Delta n}$	$K_w = [\mathrm{H}^+][\mathrm{OH}^-] = 10^{-14}$
Absorbance = $\varepsilon c \ell$	$PV = n\mathbf{R}T$	$N(E) = N_0 \omega \exp(-E/k_B T)$
$\mathbf{p}K_a = -\log(K_a)$	$pH(e.p.) = \frac{1}{2} (pK_{a1} + pK_{a2})$	$S = k_B \ln W$
	• ·• ·• •	

# Honor Pledge: At the end of the examination time, please write out the following sentence and sign it, or talk to me about it:

"I pledge on my honor that I have not given or received any unauthorized assistance on this examination."

#### **<u>1. Redox titration (25 pts)</u>**

In Lab 7B, we used iodate  $IO_3^-$  + excess iodide to generate triiodide according to:

 $IO_3^- + 8 I^- + 6 H^+ \rightarrow 3 I_3^- + 3 H_2O_2$ 

then used the triiodide in excess to oxidize ascorbic acid, as in:

ascorbic acid +  $I_3^- \rightarrow$  dehydroascorbic acid +  $3I^- + 2H^+$ ,

(a; 6 pts) Fill in the blanks:

...and then back-titrated with (word): \_\_\_\_\_\_\_to find out how much triiodide was still left

according to this reaction:  $I_3^- + 2 \_ \longrightarrow 3 I^- + S_4 O_6^{2^-}$ .

(b; 4 pts) If you Google for ascorbic acid titration, you can find a few undergraduate labs that do an acidbase titration to determine the purity of commercial vitamin C tablets. Why is this inferior to the redox titration that we and most others use?

(b; 3 pts) Why is periodate used as the primary concentration standard in this lab, as opposed to triiodide or thiosulfate?

(d; 3 pts) What result would you have obtained if the supplier had a QC problem and supplied KIO<sub>3</sub> at 90% purity, with the remainder being entirely soluble and inert? Circle the best answer:

- (*i*) There would have been no change in the experiment.
- (ii) The calculated ascorbic acid concentration would be 10 % too low.
- (iii) The calculated ascorbic acid concentration would be 10 % too high.
- (*iv*) The experiment would have failed completely.

(e; 3 pts) Briefly explain your answer to (d).

(f; 6 pts) Briefly discuss how/why a truly terrible KIO<sub>3</sub> purity of  $\sim 15$  % would have made (*iv*) the correct answer to part (d), and state what you could have done to recover.

#### 2. Acid-base and methods development (16 pts)

Ammonia gas (NH<sub>3</sub>) dissolves in water and is in equilibrium with ammonium hydroxide, but the pKb of NH<sub>3</sub> is about 4.75 so the concentration of ammonium cation is low:

$$NH_3(g) + H_2O(l) \rightleftharpoons NH_3(aq) + H_2O(l) \rightleftharpoons NH_4^+(aq) + HO^-(aq)$$

In the Kjeldahl assay for determining protein concentration, a sample is reacted completely with strong acid to convert all organic nitrogen to  $NH_4^+$  (*aq*). The solution is then made basic, and whatever is volatile (i.e. not ions) is distilled into a receiving flask containing an acid, often the weak acid boric acid B(OH)<sub>3</sub>.

$$B(OH)_3(aq) + H_2O \rightleftharpoons B(OH)_4^-(aq) + H^+(aq)$$

Then the ammonia and/or ammonium content in the receiving flask is measured.

(Not fun at all fact: The Kjeldahl assay can be fooled by the nitrogen-rich but non-nutritious and poisonous compound melamine, which was used in 2007-2008 by unscrupulous pet food and baby formula manufacturers to simulate protein in their products, killing dogs and babies.)

(a; 4 pts) Why is it necessary to raise the pH

before doing the distillation?



Picture credit: By Roshan220195 - Own work, CC BY-SA 3.0, https://commons.wikimedia.org/w/index.php?curid=18937792

(b; 6 pts) Why is it necessary to collect the distillate in an acidic receiving solution? What are the products of bubbling ammonia into boric acid?

(c; 6 pts) Experimentally, how could you determine the amount of ammonia that ended up in the collection flask?

# 3. SDL (12 pts)

Describe an unexpected finding from your SDL, why it was a surprise or disappointment, and how you would either make the issue go away by improving your methods or else investigate it further.

## 4. Nanoparticles and Chelators (22 pts)

(a; 6 pts) Write the name of the physical phenomenon that causes nanoparticles of different sizes to be

different colors: \_\_\_\_\_\_. What technique provided the

calibration curve we used to estimate size from  $\lambda_{max}$ ?

(b; 3 pts) What color do you get if all wavelengths are absorbed equally?

- In Lab 6, we used EDTA to rip  $Zn^{+2}$  away from a xylenol orange indicator.
- (c; 4 pts) Explain why we used EDTA in this lab instead of just titrating directly with xylenol orange until no further color change was observed.



https://pediaa.com/difference-between-disodium-edta-and-tetrasodium-edta/

(d; 3 pts) Explain why EDTA has such a strong binding affinity for  $Zn^{+2}$  (and many other metal ions).

(e; 6 pts) Explain why EDTA is used in the treatment of lead and mercury poisoning, but why large doses should be avoided unless one has actually been poisoned.

## 5. Using the computer as if it were a piece of apparatus (25 pts)

(a; 12 pts) Sketch a curve showing pH vs. volume of base added with two titration curves, one curve for an acid with a pKa of 3.5 with strong base, and the other curve for the same acid at the same concentration with the same concentration of a weak base. Indicate how you would measure the pKa of the acid.



(d; 6 pts) How would you exercise the micro\_movr3 program to see whether its toy model for entropy is an extensive state function (which

is the idea that two moles of a pure substance at a given temperature has twice the entropy of one mole)?