Exam 3 Chm 205 (Dr Mattson) 28 March 2018

Academic Integrity Pledge: In keeping with Creighton University's ideals and with the Academic Integrity Code, I pledge that this work is my own and that I have neither given nor received inappropriate assistance in preparing it.

Name:

Chemistry Student Number:

Signature:

Instructions: Show all work whenever a calculation box is provided! Write legibly. Include units whenever appropriate. You will receive credit for how you worked each problem as well as for the correct answer. If you need more space, you may use the back of the data sheet provided — Write: "See data sheet" in the answer box – then write your name on the data sheet. On your desk you are allowed only pencils (but no pencil pouch), an eraser, and a non-programmable calculator without a slipcover. Backpacks, bags, and purse-like items must be stored on the tables in the back of the room. Cell phones must be silent and placed in your backpack/bag/purse – not in your pocket.

- (4 pts) Write the net ionic neutralization reaction for a strong acid with a strong base. Include appropriate arrows and a numerical value for K_n.
- 2. (4 pts) Write the **net ionic** neutralization reaction for a weak acid, HA ($K_a = 7.0 \times 10^{-6}$), with a strong base. Include long/short arrows and a **numerical value** for K_n .
- 3. (4 pts) Write the **net ionic** neutralization reaction for a weak base, A⁻, with a strong acid. Include long/short arrows and **numerical value** for K_n (K_a^{HA} = 6.4 x 10⁻⁵)
- 4. (5 pts) Which of these are buffers and if so, what recipe produced the buffer: Circle: Recipe 1 (R1): wa + wb mixture, R2: wa + OH⁻, R3: wb + H₃O⁺ or Not a buffer.

5.5 mmol KF + 3.0 mmol HNO ₃	R1 R2 R3 Not
7.0 mmol HF + 7.0 mmol NaOH	R1 R2 R3 Not
40 mL 1 M KCN + 15 mL 1 M HBr	R1 R2 R3 Not
5.0 g NaC $_2$ H $_3$ O $_2$ + 3.0 g HC $_2$ H $_3$ O $_2$	R1 R2 R3 Not
2.0 mmol HClO ₂ + 1.2 mmol KOH	R1 R2 R3 Not

5a. (5 pts) What is the pH of a buffer prepared by combining 1.290 g NaNO₂ (MM = 69.00 g/mol) with

500.0 mL 0.100 M HNO₂(aq)? K_a^{HNO₂} = 3.7 x 10⁻⁴

Answer with correct significant figures:

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5b. (4 pts) What is the pH of the buffer from Question 5a if 0.0030 mol NaOH is added?



mL to reach the equivalence point.

6a. (4 pts) What was the original concentration of the acid? This is a calculation – do not estimate from graph.



6b. (4 pts) What is K_a for the acid?

Answer with correct significant figures: _

6c. (5 pts) What was the pH after 13.00 mL NaOH has been added? This is a calculation – do not estimate.



- 6d. (2 pts) From the information provided above, directly or indirectly, including from the graph, do you have everything needed to calculate the pH at the equivalence point? Circle: Yes or No
- (5 pts) Suppose 25.00 mL of a 0.160 M HNO₃(aq) is titrated with 0.205 M KOH(aq). Calculate the pH after 18.00 mL KOH has been added.

Answer to the hundredths place:

8a. (4 pt) Calcium fluoride, CaF_2 is a sparingly soluble salt. Write its equilibrium expression and define its K_{sp} in terms the ions produced.

Answer with units:

8a. (5 pt) What is the molar solubility of calcium fluoride if $K_{sp} = 3.5 \times 10^{-11}$? Hint: Set up MICE table.

Answer with units:

8b. (4 pts) What is the molar solubility of calcium fluoride in a solution that is 0.040 M KF?

Answer with units:

8c. (4 pts) Will a precipitate form if equal volumes of 0.0040 M Ca(NO₃)₂ and NaF are mixed? Show Q_{sp} calculation! Note: The two solutions dilute each other.



- 9. (5 pts) Which of these processes is entropy-favored?
 - a. dissolving sugar in hot coffee
 - b. mixing of nitrogen gas and bromine vapor
 - c. ethanol boiling
 - d. $N_2(g) + 2 H_2(g) \rightarrow N_2H_4(g)$
 - e. HCl(aq) + Zn(s) \rightarrow ZnCl₂(aq) + H₂(g)

10. (5 pts) Which of these processes are spontaneous?

- a. dissolving sugar in hot coffee
- b. mixing of nitrogen gas and bromine vapor
- c. ethanol boiling at 25 °C and 1 atm
- d. $CH_4(g) + 2 O_2(g) \rightarrow CO_2(g) + 2 H_2O(g)$
- e. HCl(aq) + NaOH(aq) \rightarrow H₂O(aq) + NaCl(aq)

11a. (5 pts) Calculate ∆S ^o	S ^o (J mol⁻¹ K⁻¹)	
for the reaction in the	H ₂ O ₂ (aq)	144
standard entropies.	H ₂ O(I)	70
Show units in your calculation.	O ₂ (g):	205

 $2 H_2O_2(aq) \rightarrow 2 H_2O(l) + O_2(g)$

Answer with units:

11b. (4 pts) Given ΔH_f^o = -188 kJ for the reaction above, calculate ΔG^o for the reaction.



A. all temperatures B. never

E lievel

- C. high temperatures only D. low temperatures only
- 12a. (4 pts) Estimate the boiling point of benzene, given: $\Delta H_{vap} = 30.7 \text{ kJ/mol} \text{ and } \Delta S_{vap} = 87.0 \text{ J mol}^{-1} \text{ K}^{-1}.$

Answer with units:

12b. (4 pts) Estimate ΔG_{vap} at 85 °C for benzene. Is vaporization spontaneous at 85 °C?

13a-d (4 pts). Consider the all-gas reaction at 25 °C.

 $P_4 + 6 H_2 \rightarrow 4 PH_3 \quad \Delta G^0 = +29.5 \text{ kJ} \quad \Delta H^0 = -37.3 \text{ kJ}$

13a. Is the reaction enthalpy-favored? Yes or No

13b. Is the reaction spontaneous at 25 °C? Yes or No

- 13c. Is the reaction entropy-favored? Yes or No
- 13d. Is the reaction ever spontaneous? Use the same answer choices as in Question 11c: Circle: A B C D
- 13e. (4 pts) Calculate ΔG at 25 °C starting with 1.0 atm P₄(g), 1.0 atm H₂(g) and 5.0 atm PH₃(g).



Answer: _____ Total score (out of 100): _____

 $\mathsf{A}+ \geq 95\% \hspace{0.2cm} \mathsf{A} \geq 90\% \hspace{0.2cm} \mathsf{B}+ \geq 85\% \hspace{0.2cm} \mathsf{B} \geq 80\% \hspace{0.2cm} \mathsf{C}+ \geq 75\% \hspace{0.2cm} \mathsf{C} \geq 70\% \hspace{0.2cm} \mathsf{D} \geq 60\%$

Answers

1. $H_3O^+ + OH^- \implies H_2O K_n = 1 \times 10^{+14}$ 2. HA + OH⁻ \iff H₂O + A⁻ K_n = $^{1}/_{K_n}$ = 7.0 x 10⁺⁸ 3. $A^- + H_3O^+ - H_2O + HA K_n = \frac{1}{K_a} = 1.6 \times 10^{+4}$ 4. R3, Not, R3, R1, R2 5a. 3.00 5b. 3.09 6a. 0.164 M 6b. $K_a = 1.82 \times 10^{-6}$ 6c. 5.55 6d. Yes 7.2.14 8a. $BaF_2 \longleftarrow Ba^{+2}(aq) + 2 F^{-}(aq) K_{sp} = [Ba^{+2}][F^{-}]^2$ 8b. x = $2.06 \times 10^{-4} M$ 8c. x = 2.19 x 10⁻⁸ M 8d. Q_{sp} = 8.0 x 10⁻⁹ YES 9. a, b, c, e 10. a, b, d, e 11a. ∆S^o = 57 J/deg 11b. ∆G⁰ = -205 kJ 11c. A 12a. 353 K 12b. ΔG_{vap} = -0.446 kJ (kJ/mol is also ok this time) 13a-d (4 pts). Consider the all-gas reaction at 25 °C. $P_4 + 6 H_2 \rightarrow 4 PH_3 \quad \Delta G^0 = +29.5 \text{ kJ} \quad \Delta H^0 = -37.3 \text{ kJ}$ 13a. Yes 13b. No 13c. No 13d. D 13e. ∆G = 45.4 kJ 13f. $K_p = 6.74 \times 10^{-6}$