Chemistry 210

Be sure to put your name on each page. This page can be removed from your exam so that you will have a Periodic Table handy throughout the exam, it does not need to be turned in. Show all your work for problems which require any sort of calculation, no credit will be given for answers without work shown. If you have shown a significant amount of work or multiple drawings for a problem, draw a box around what you consider your final answer.

 $k = Ae^{-Ea/RT}$

 $\ln(k) = \left(\frac{-E_a}{R}\right)\left(\frac{1}{T}\right) + \ln(A)$

 $\ln\left(\frac{k_1}{k_2}\right) = \frac{E_a}{R}\left(\frac{1}{T_2} - \frac{1}{T_1}\right)$

 $pH = pK_a + log\left(\frac{[conjugate base]}{[conjugate acid]}\right)$

Avogadro's Number =
$$6.022 \times 10^{23}$$
 ^{units}/_{mol}
32.00°F = $0.000°C = 273.15K$
Density of Water = $1.000^{g}/_{mL}$
 $R = 0.08206^{L*atm}/_{mol*K} = 8.314^{J}/_{mol*K}$
 $PV=nRT$
 $\Delta T_{fp/bp} = k_{fp/bp} \cdot m \cdot i$
For water, $k_{fp} = -1.86°C/_{m}$; $k_{bp} = 0.52°C/_{m}$
 $P_1 = X_1P_1^{o}$
 $P = cRTi$
 $C_1V_1 = C_2V_2$
Quadratic formula:
 $x = \frac{-b \pm \sqrt{b^2 - 4ac}}{2a}$

Integrated Rate Laws: $\begin{array}{l} ln[A]_t = -kt + ln[A]_o \\ 1/[A]_t = kt + 1/[A]_o \\ [A]_t = -kt + [A]_o \end{array}$

232.04

231.04

238.03

237.05

(244)

(243)

1																	2
Н																	Не
1.0079																	4.0026
3	4											5	6	7	8	9	10
Li	Be											В	С	Ν	0	F	Ne
6.941	9.0122											10.811	12.011	14.007	15.999	18.998	20.180
11	12											13	14	15	16	17	18
Na	Mσ											Al	Si	Р	S	CL	Ar
22.990	24.305											26.982	28.086	30.974	32.066	35.453	39.948
19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36
K	Ca	Sc	Ti	V	Cr	Mn	Fe	Со	Ni	Cu	Zn	Ga	Ge	As	Se	Br	Kr
39.098	40.078	44.956	47.88	50.942	51.996	54.938	55.847	58.933	58.69	63.546	65.39	69.723	72.61	74.922	78.96	79.904	83.80
37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54
Rb	Sr	Y	Zr	Nb	Mo	Tc	Ru	Rh	Pd	Ag	Cd	In	Sn	Sb	Те	Ι	Xe
85.468	87.62	88.906	91.224	92.906	95.94	(98)	101.07	102.91	106.42	107.87	112.41	114.82	118.71	121.76	127.60	126.90	131.29
55	56	57	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86
Cs	Ba	La	Hf	Ta	W	Re	Os	Ir	Pt	Au	Hg	Tl	Pb	Bi	Po	At	Rn
132.91	137.33	138.91	178.49	180.95	183.84	186.21	190.23	192.22	195.08	196.97	200.59	204.38	207.2	208.98	(209)	(210)	(222)
87	88	89	104	105	106	107	108	109	110	111	112		114		116		
Fr	Ra	Ac	Rf	Db	Sg	Bh	Hs	Mt									
(223)	226.03	227.03	(261)	(262)	(263)	(262)	(265)	(266)	(269)	(272)	(277)						
														_			
		58	59	60	61	62	63	64	65	66	67	68	69	70	71		
		C.	D	NJ	D	C	F	C.J	ть	D	II.	E-	Т	VL	Т		
		140.12	I I 140.01	144.24	F III (145)	5 11 150.26	EU 151.07	GU	1 D	Dy	П 0	EI 167.26	1 M	1 D	174.07		
		90	91	92	93	94	95	96	97	98	99	107.20	108.94	102	103		
		ТЬ	Da	TI	N	D	A	C	DI-	Cr.	E.	E	MJ	No	T		
		IN	ra	U	IND	ru	AM	UM	ĎК	U	ĽS	гm	IVIA	INO	Lr		

Exam 3

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1. Complete each row of the following tables for aqueous solutions at 25°C (4pts per box):

[H ₃ O ⁺]	[OH ⁻]	рН	рОН	Acidic, Basic or Neutral?		
3.19x10 ⁻⁴	3.13x10 ⁻¹¹	3.496	10.504	Acidic		
4.80x10 ⁻⁵	2.08×10^{-10}	4.319	9.681	Acidic		

Conjugate Acid	K _a @25°C	Conjugate Base	K _b @25°C
H ₂ CO ₃	4.2x10 ⁻⁷	HCO ₃ ⁻¹	2.4x10 ⁻⁸
HSO ₄ ⁻¹	1.2×10^{-2}	SO_4^{2-}	8.3x10 ⁻¹³

2. Does the combination listed result in an effective buffer solution? (4pts each)



3. How much 1.131 M NaOH(aq) must be added to 45.00mL of 0.984 M HCl(aq) to reach the equivalence point? What is the pH of this solution at the equivalence point? (12pts)

 $NaOH(aq) + HCl(aq) \rightarrow NaCl(aq) + H_2O(l)$ $\frac{(0.04500 \text{ L HCl(aq)})(0.984 \frac{\text{mol HCl}}{\text{L HCl(aq)}})(\frac{1 \text{ mol NaOH}}{1 \text{ mol HCl}})}{1.131 \frac{\text{mols NaOH}}{\text{L sol'n}}} = 0.0392 \text{ L NaOH(aq)} = 39.2 \text{ mL NaOH(aq)}$

Since this is a strong acid/strong base titration, the equivalence point should be at pH = 7.

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4. You have prepared a buffer solution by combining 0.164mols of nitrous acid (HNO₂, $K_a = 4.5 \times 10^{-4}$) and 0.186mols of sodium nitrite in enough water to make 500.0mL of solution. What is the pH of this buffer solution? (12pts)

Plugging into the Henderson-Hasselbalch equation:

 $pH = 3.35 + log \bigg(\frac{0.186 mols/0.5000 L}{0.164 mols/0.5000 L} \bigg) = 3.40$

NOTE: The H-H equation technically uses the *concentrations* of conjugate acid and conjugate base, but since they are both dissolved in the same volume of water, the volume cancels out.

5. Sketch the titration curve (pH vs. volume added) for the titration of 1 M potassium sulfite $(K_2SO_3, K_{b1}=1.6x10^{-7})$ and 1 M HClO₄(aq). Label all the axes (including approximately accurate numbers) and the major sulfite-based species present in solution at each point in the titration curve. Indicate the equivalence point(s) on the curve. (17pts)



Volume acid added \rightarrow

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6. What is the K_b of a base if 750.0mL of a solution containing 0.153 mol of the base and 0.141 mol of its conjugate acid has a pH of 4.165? Over what pH range would this conjugate acid/ conjugate base pair make an effective buffer? (16pts)

Plugging into the Henderson-Hasselbalch equation:

 $4.165 = pK_a + \log\left(\frac{0.153 \text{mols}/0.7500\text{L}}{0.141 \text{mols}/0.7500\text{L}}\right)$ $pK_a = 4.130$ $K_a = 7.42 \times 10^{-5}$ $K_b = 1.35 \times 10^{-10}$

This system would make an effective buffer over the range $3.130 \rightarrow 5.130$

7. You have titrated 25.00mL of a monoprotic acid ($K_a = 3.4 \times 10^{-5}$) with 0.613 M KOH(aq). If thymol blue (endpoint 7.8-9.5) is used as an indicator, you reach the endpoint when 31.37mL of base is added. Based on this data, what is the concentration of the acid? Would your result change if ethyl red (endpoint 4.0-5.8) was used as an indicator? If so, how? Which indicator (thymol blue or ethyl red) more correctly indicates the equivalence point in this titration? (Explain your answers!) (25pts)

$$\frac{\text{KOH}(\text{aq}) + \text{HA}(\text{aq}) \rightarrow \text{KA}(\text{aq}) + \text{H}_2\text{O}(1)}{\frac{(0.03137 \text{ L KOH}(\text{aq}))(0.613 \frac{\text{mol KOH}}{\text{L KOH}(\text{aq})})(\frac{1 \text{ mol HA}}{1 \text{ mol KOH}})}{0.02500 \text{ L HA}(\text{aq})} = 0.769 \text{ M HA}(\text{aq})$$

Since this is a weak acid/strong base titration, the equivalence point should be basic. Therefore, the proper indicator to use will have an endpoint that is basic. Thymol blue is an appropriate indicator.

If ethyl red is used as an indicator in this reaction, it will reach its endpoint *before* the titration reaches its equivalence point. This would result in a *calculated concentration* that is too low.

NOTE: A slight language note here. Some people said that if ethyl red was used, the concentration of the acid would be lower. The concentration of the acid is the same regardless of the indicator used, but the concentration that is calculated would be lower than the actual concentration.