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<b>CHEM 142</b>	3 - Final	Exam –	May 1	12, 2015
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Name\_Salabam

If you wish to have your final exam and course grade posted on the Web site, please provide me with a four (4) digit number which will be the ID number for your grade.

Four (4) digit number for posting.

**Problem (4 pts)**: A concentration cell is prepared with 0.60 M Calcium Nitrate, Ca(NO<sub>3</sub>)<sub>2</sub>, in the reference compartment (cathode) and a saturated solution of Calcium Phosphate, Ca<sub>3</sub>(PO<sub>4</sub>)<sub>2</sub>, in the sample compartment (anode).

The cell reaction can be written as: Ca(s)|Ca<sup>2+</sup>(xx M)||Ca<sup>2+</sup>(0.60 M)|Ca(s)

The measured cell voltage is +0.161 V. Calculate the Solubility Product, K<sub>sp</sub>, of Ca<sub>3</sub>(PO<sub>4</sub>)<sub>2</sub>

 $\frac{1}{\sqrt{\frac{xx}{0.60}}} = \frac{1}{\sqrt{\frac{xx}{0.60}}} = \frac{1}{\sqrt{\frac{xx}{0.60}}$ 

(2) Calentato  $K_{SP}$   $C_{a3}(PO_{4})_{2} \rightarrow 3 \frac{20}{6} \times PO_{4}^{3}$   $5 = 2.18 \times 10^{-7}$   $5 = 2.28 \times 10^{-7}$   $5 = 2.28 \times 10^{-7}$   $K_{ap} = (2.18 \times 10^{-7})^{2} = (2.18 \times 10^{-7})^{3} (148 \times 10^{-7})$   $= 2.28 \times 10^{-79} \approx 2.8 \times 10^{-79}$ 



Conversions: 1 atm. = 760 torr

Constants:  $R = 0.082 \text{ L-atm/mol} \cdot \text{K}$ 

R = 8.31 J/mol·K

 $R = 8.31x10^{-3} \text{ kJ/mol} \cdot \text{K}$ 

 $N_A = 6.02 \times 10^{23} \text{ mol}^{-1}$ 

F = 96,500 Coul/mol e<sup>-</sup>

 $c = 3.00x10^8$  m/s (speed of light)

Molar Masses: Given with each question: [M=xx]

18    A	4.003 10 <b>Ne</b>	0.18 18 9.95	3.80 3.80	<b>6</b> 2	<b>3</b> 88 53	(222				
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<b>1</b> √		∞└	6 <b>T</b> 8	4₽	101.1 8,0	190.2	708 (265)			
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_		S VB	8> g	4 <b>Z</b>	92.91 73	180.9	105 <b>Db</b> (262)	anlde	O 4	2 0 0 B
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	<u> </u>						(% <b>C</b> 3	/_		
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-≤	- <b>エ</b> 00 - <b>コ</b>	6.941 11 23.00	<b>5×</b> 5	33 <b>8</b>	85.47 55	132.9	87 <b>F</b> (223)			
	- 0	69	4	ın	9		٠ ;			

#### **ELECTROCHEMISTRY INFORMATION**



## **Table 1: Standard Reduction Potentials**

# Reduction Half-Reactions E° (V)

	_ ( - ,
$F_2 + 2 e^- \rightarrow 2 F^-$	+2.87
$Au^{3+} + 3e^- \rightarrow Au$	+1.50
$Cl_2 + 2 e^- \rightarrow 2 Cl^-$	+1.36
$Br_2 + 2 e^- \rightarrow 2 Br$	+1.07
$Hg^{2+} + 2 e^{-} \rightarrow Hg$	+0.86
$Ag^+ + 1 e^- \rightarrow Ag$	+0.80
$l_2 + 2 e^- \rightarrow 2 l^-$	+0.54
$Cu^{2+} + 2 e^- \rightarrow Cu$	+0.34
$Fe^{3+} + 3e^{-} \rightarrow Fe$	-0.04
$Sn^{2+} + 2 e^- \rightarrow Sn$	-0.14
$Ni^{2+} + 2 e^- \rightarrow Ni$	-0.25
$Zn^{2+} + 2 e^- \rightarrow Zn$	-0.76
$Mn^{2+} + 2 e^- \rightarrow Mn$	-1.18
$Al^{3+} + 3 e^{-} \rightarrow Al$	-1.66
$Mg^{2+} + 2 e^{-} \rightarrow Mg$	-2.37
$K^+ + 1 e^- \rightarrow K$	-2.93
$Li^+ + 1 e^- \rightarrow Li$	-3.05



#### Table 2: Some Reduction and Oxidation Potentials in Aqueous Solution

#### **Reduction Potentials**

#### **Oxidation Potentials**

$$2 H_2O \rightarrow O_2 + 4 H^+ + 4 e^ E^o_{oxid} = -1.23 V$$
  
 $2 I^- \rightarrow I_2 + 2 e^ E^o_{oxid} = -0.54 V$   
 $2 Br \rightarrow Br_2 + 2 e^ E^o_{oxid} = -1.07 V$   
 $2 F^- \rightarrow F_2 + 2 e^ E^o_{oxid} = -2.87 V$ 

### Some Electrochemical Equations

F = 96,500 C/mol e<sup>-</sup> (Coulombs per mole of electrons)

$$E = E^{\circ} - \frac{0.0592}{n} \cdot \log(Q)$$



(56) MULTIPLE CHOICE (Mark the one correct answer to each question on your scantron)

> Turn in: (a) Your scantron with your name and answers (there is no need to bubble in your ID.

(b) The cover sheet with your Electrochemistry Problem + four (4) digit number if you would like your results posted on the course web site

nu	mper ir you wou	ia like your results	s postea on the cour	se web site.
4 [	ooints, yielding a	total of 60 points	rth 1 point. The probon the test. Your sony further analysis.	
1.	7.1		$3A + B \rightarrow 2C$ . If the rarate" of the reaction?	ite of change of [A] is
(	(A) +0.20 M hr <sup>1</sup>		(B) -0.40 M hr <sup>1</sup>	
	(C) -0.20 M hr <sup>r</sup>		(D) +0.40 M hr <sup>-1</sup>	
2.	measured. It is for the B concentrate to the first exper rate increases by	ound that if the initiation the same, the raiment). If the conce	ate increases by a fac entrations of both A an	sed is tripled, keeping tor of nine (9) (relative
	(A) $k[A]^2[B]^2$	(B) k[A][B] <sup>3</sup>	(C) k[A] <sup>2</sup> [B] <sup>3</sup>	(D) k[A] <sup>3</sup> [B] <sup>2</sup>
3.	When the initial	concentration of A is ntration of A is 0.20	s 0.60 M, the initial rat	o [A]; i.e. Rate = k[A] <sup>n</sup> . te is 0.50 Ms <sup>-1</sup> . When 50 Ms <sup>-1</sup> . The order of
(	(A) -2	(B) -1	(C) +1	(D) +2
4.	[A]? When the i		of A is 0.80 M, the init	order with respect to tial rate is 0.45 Ms <sup>-1</sup> .
	(A) 1.4 M <sup>-3</sup> s <sup>-1</sup>	(B) 0.73 M <sup>-3</sup> s <sup>-1</sup>	(C) $0.91 \text{ M}^{-3}\text{s}^{-1}$ ((I	D) 1.1 M <sup>-3</sup> s <sup>-1</sup>
5.			oducts, the rate const what is the approxima	ant is 0.025 s <sup>-1</sup> . If the te concentration of A
	(A) 0.40 M	(B) 0.30 M	(C) 0.26 M	(D) 0.61 M



6.		Products, when the initial concentration of A entration to decrease to 0.30 M. The rate nately:
	(A) 0.027 M <sup>-1</sup> s <sup>-1</sup> (B) 0.056 M <sup>-1</sup>	s <sup>-1</sup> (C) 0.015 M <sup>-1</sup> s <sup>-1</sup> (D) 0.082 M <sup>-1</sup> s <sup>-1</sup>
7.	For the reaction, 2 NO(g) + O <sub>2</sub> (g) $\rightarrow$	2 NO <sub>2</sub> (g), the reaction mechanism is:
	$2 \text{ NO} \rightleftarrows \text{N}_2\text{O}_2$	ast Equilibrium (N <sub>2</sub> O <sub>2</sub> ) is an intermediate
	$N_2O_2 + O_2 \rightarrow 2 NO_2$ S	low rate determining step
	The overall rate equation for this rea	action is:
	(A) Rate = $k'[NO]^2/[O_2]$	(B) Rate = $k'[O_2][2NO]$
	(C) Rate = $k'[O_2][NO]$	(D) Rate = $k'[O_2][NO]^2$
For	or #8-#9: Consider the gas phase equ	uilibrium, 2 POBr <sub>3</sub> (g) $\rightarrow$ 2 PBr <sub>3</sub> (g) + O <sub>2</sub> (g).
8.	For the above reaction, if the volum and K <sub>c</sub> will	e is <b>decreased</b> , the ratio [PBr <sub>3</sub> ]/[POBr <sub>3</sub> ] will
	(A) increase , increase	(B) increase , remain constant
	(C) decrease, remain constant	(D) decrease, decrease
9.		added to the mixture at constant pressure
	(A) increase , increase	(B) increase , remain constant
	(C) decrease , remain constant	(D) remain constant , remain constant
10.	container with A at a concentration	reaction, A(g) $\rightleftharpoons$ 2 B(g). If one initially fills a of 2.0 M, and then allows it to come to ibrium concentration of A is 1.6 M. Therefore it, $K_c$ is approximately:
1	(A) 0.40 (B) 0.50	(C) 0.10 (D) 0.67
11.	1. Consider the reaction: $2HBr(g) \stackrel{k}{\longleftarrow}$ is $K_c = 15.0$ at $100$ °C. The Enthalp $\Delta H^o = +70$ . kJ/mol. What is the app	
	(A) 33. (B) 0.030	(C) 490 (D) 0.45



12.	The gas phase	e molecule	e, A, dissocia	ates acc	cording to the	e equil	ibrium,	
	A(g)   ⇒ 3 B(g) initial concentration dissociates]	ation of 2	.0 M of A in	to a flas	k, what is th	ie appr	oximate	
	(A) 0.15 M	(B) 0.2	28 M	(C) 0.	09 M	(D)	0.15 M	
13.	The concentra Approximately the solution?:							0 mL of
(	(A) 6.8x10 <sup>4</sup> ng		(B) 4.5	x10⁵ ng	((	C) 68 r	ng	
	(D) None of the	ne above	, ,		·			
14.	When 16 gram [M=18], the description is:		•	-	-	•	-	
	(A) 3.63 M	) (B)	4.17 M	(	C) 4.48 M		(D) 4.63	М
15.	A sample of el (K <sub>f</sub> = 1.86 °C/r how many mo	n). The fr les of ethy	eezing point lene glycol	of the s	solution is -3	3.6 °C.	Approxima	
	(A) 0.42 mol	(B)	1.35 mol	(C	) 1.94 mol	(	(D) 2.76 m	nol
16.	Which of the fo	ollowing s	olutions has	the low	<b>rest</b> freezing	g point'	?	
	(A) 0.32 m C <sub>6</sub>	H <sub>12</sub> O <sub>6a</sub>		(B) 0.1	0 m Ca(NO	3)2		
	(C) 0.09 m K <sub>3</sub>	PO <sub>4</sub>		(D) 0.2	0 m NaBr			
17.	What is the ap strong electrol aqueous solut	yte, Calcii	um Phospha					
	(A) 15 torr	(B)	77 torr	(	C) 0.10 torr		(D) 46 t	orr
18.	The normal bo constant is 5.0 grams of CCl <sub>4</sub> unknown com	) °C/m. W , the boilir	hen 60. grar ig point of th	ns of ar ie soluti	unknown c	ompou	ınd is place	ed in 750
	(A) 64 g/mol	(B)	86 g/mol	(c	) 114 g/mol		(D) 153 g/	mol
19.	Approximately aqueous solut			-	-	be dis	solved in 1	5. L of
	(A) 7.5x10 <sup>-2</sup> g	(B)	7.5x10 <sup>-4</sup> g	(C)	2.0x10 <sup>-3</sup> g		D) 3.0x10	<sup>2</sup> g)



(A) 1.6x10 <sup>-5</sup> (B) 6.3x10 <sup>-10</sup> (C) 1.6x10 <sup>-17</sup> (D) 4.8 21. Benzoic Acid (HBenz) has an acid dissociation constant of 1.6x10 <sup>-9</sup> . W	₹∨10-7
21 Benzoic Acid (HBenz) has an acid dissociation constant of 1 6v10-9 W	W10.
approximate pH a 0.20 M solution of aqueous sodium benzoate (NaBer	
(A) 11.0 (B) 9.3 (C) 4.7 (D) 3.0	
22. The weak base, aniline (Anil), has a base equilibrium constant, K <sub>b</sub> = 4.3 What is the pH of a 0.05 M aqueous solution of anilinium chloride (Anilinium chloride)	
(A) 11.0 (B) 8.7 (C) 5.3 (D) 3.0	>
23. If added to 2 L of 0.80 M NaOH, which one of the following would form	a buffer?
(A) 2. L of 0.50 M Nitric Acid (HNO <sub>3</sub> )	
(B) 2. L of 0.50 M Acetic Acid (HAc)	
(C) 2. L of 1.0 M Lactic Acid (HLac)	
(D) 2. L of 1.0 M Potassium Acetate (KAc)	
For #24 - #28: Tellurous acid, $H_2TeO_3$ , is a diprotic acid with acid dissociation constants, $K_a' = 3.0 \times 10^{-3}$ and $K_a'' = 2.0 \times 10^{-8}$	
24. What is the pH of a 0.04 M solution of sodium tellurite, Na <sub>2</sub> TeO <sub>3</sub> ?	
(A) 9.45 (B) 10.15 (C) 11.25 (D) 3.85	
25. What is the pH of a solution containing 0.20 M KHTeO <sub>3</sub> and 0.50 M Na <sub>2</sub>	₂TeO₃?
(A) 2.92 (B) 7.30 (C) 8.10 (D) 2.12	
26. What is the pH of a solution prepared by adding 0.35 mol of KOH to 2.0 L of 0.50 M H <sub>2</sub> TeO <sub>3</sub> ?	
(A) 7.43 (B) 2.79 (C) 2.06 (D) 2.25	>
27. What is the pH of a solution prepared by adding 2 L of 0.70 M HCl to 2 0.45 M Na <sub>2</sub> TeO <sub>3</sub> ?	L of
(A) 7.60 (B) 2.42 (C) 7.80 (D) 2.62	``
28. What ratio of [HTeO <sub>3</sub> -]/[TeO <sub>3</sub> <sup>2</sup> -] will give a pH of 7.00	



29.		/I KOH(aq) are requ aq) solution.  What		neutralize 400 mL of a he acid solution?	an
(	(A) 0.23 M	(B) 0.36 M	(C) 0.45 M	(D) 0.90 M	
30.	-	•	und, silver carbona	ate, Ag <sub>2</sub> CO <sub>3</sub> . The solu	bility
	product constant	is $K_{sp}$ = 6.2x10 <sup>-12</sup> .			
	What is the conc and 0.1 M K <sub>2</sub> CO:		ns, [Ag+], in a solu	ution containing Ag₂CC	Эз
	(A) 1.6x10 <sup>-5</sup> M	(B) 7.9x10 <sup>-6</sup> M	(C) 3.9x10 <sup>-6</sup> M	(D) 2.5x10 <sup>-6</sup> M	
31.	Consider the rea	ction: 2 NO <sub>2</sub> (g) $\rightarrow$ N	$N_2(g) + 2 O_2(g)$ , $\Delta$	.H° < 0. This reaction	is:
	(A) Reactant Fa	vored at all tempera	atures		
		ored at all temperat			
	` '	ored at low tempera			
	(D) Product Fav	ored at high tempe	ature		
32.		al reaction, $A \rightleftharpoons B$ , $\Delta$ 0x10 <sup>-11</sup> at 25 °C. $\Delta$		e equilibrium constant is approximately:	for
	(A) -470 J/K	(B) -67 J/K	(C) +470 J/K	(D) +67 J/K	
	he entropy char	ge of the <b>surround</b>	lings, $\Delta S_{ ext{surr}}$ , for th	), is +30.7 kJ/mol. Whe condensation of 0.5	
	•	at the boiling point,  (B) -192 J/K	(C) -43.5 J/K	) (D) +87 J/K	
	(A) -87. J/K	(D) -192 J/K	(C) -43.3 J/K	(b) 101 3/K	
34.	is the approxima		llibrium Constant f	Go = -236 kJ at 25°C. \ For the <b>related reactio</b>	
	(A) 2.0x10 <sup>-21</sup>	(B) 2.4x	10+41 (	C) 2.0x10 <sup>-42</sup>	
	(D) None of the	above			
35.				8 kJ and ∆S° = +85 J/k <b>⊵low</b> °C (Cels	
	(A) Product, 29	<u>2 °</u> C	(B) Product, 565	5 °C	
(	(C) Reactant, 2	.92 °C	(D) Reactant, 56	85 °C	



at 2	.40 °C. What i			ium constant, $K = 220$ , $C$ when $P(N_2) = P(H_2) =$
(A)	-33.4 kJ	(B) -7.4 kJ	(C) + 10.4  kJ	(D) -56.4 kJ
	· ·	B, is <b>exergonic</b> at 2 cluded about the Entl		opy change is -95 J/K. is reaction?
` '	$\Delta H < 45.6 \text{ kJ}$ $\Delta H \text{ cannot be}$	(B) ΔH <		(C) ∆H > +28.3 kJ of ∆G
38. Reg	garding the foll	owing reaction, whic	h of the statements	s below is/are correct?
		O <sub>3</sub> (s) + 3 CO(g) → exidizing agent e transfered (B) 1 & 3 & 4		ed ed (D) 2 & 3
	the electroche ements is/are	emical cell given by C correct?	Cu Cu²+  Au³+ Au, w	which of the following
<ul><li></li></ul>	Electrons flow circuit The anode rea	action is $Cu \rightarrow Cu^{2+}$ s are separated by a	de to the Cu electro + 2 e <sup>-</sup>	ode through an external NO <sub>3</sub> - ions flow towards
(A)	1 & 2 & 3	(B) 1 & 2 & 4	(C) 1 & 3	(D) 2 & 3 & 4
	- #46: Use T of the test.	able 1 (Standard Re	eduction Potentia	ls), as necessary, near
40. Wh	ich of the follo	wing reactions are <b>re</b>	actant favored?	
Red (1)	Hg <sup>2+</sup> + 2 Cl <sup>-</sup> -	$\rightarrow$ Hg + Cl <sub>2</sub> (2)	$\int 2 Ag^+ + 2 I^- \rightarrow 2 I$	Ag + 2 l <sub>2</sub>
SKIP this Question (A)	Hg <sup>2+</sup> + 2 Fe <sup>2+</sup>	$\rightarrow$ Hg + 2 Fe <sup>3+</sup> (4)	$\chi$ Cu <sup>2+</sup> + 2 Ag $\rightarrow$ 2	Ag+ + Cu
(A)	2 & 3	(B) 1 & 3 & 4	(C) 3 only	(D) 1 & 4
		ction, 2 K $^+$ + Cd $\rightarrow$ 2 What is the reduction	· · · · · · · · · · · · · · · · · · ·	•
(A)	-0.40	(B) -5.46 V	(C) -3.33 V	(D) +0.40 V
	at is the stand $^{+} + 2  ^{-} \rightarrow Ni^{-}$		gy change for the	electrochemical reaction
(A)	-56 kJ	(B) +76 kJ	(C) +56 kJ	(D) +152 kJ



43. For the redox rea E° <sub>cell</sub> = -0.48 V. V	action, 2 Al <sup>3+</sup> + 3 Mn What is the equilibri		
(A) 4.7x10 <sup>-25</sup>	(B) 2.2x10 <sup>-49</sup>	(C) 4.5x10 <sup>+48</sup>	(D) 7.8x10 <sup>-9</sup>
this reaction, E°c	•	s) Ag⁺( 0.002 M)Ì A is the cell potential	\u <sup>3+</sup> (3.00 M) Au(s) . For
(A) +0.36 V	(B) +1.21 V	(C) +0.53 V	(D) +0.87 V
a concentration of (the cathode) and in the sample ce	cell with 0.30 M lead d a sample of water	I(II) nitrate, Pb(NO: with an unknown	ter was determined using (3)2, in the reference cell concentration of Pb <sup>2+</sup> (xx) Pb <sup>2+</sup> (0.20 M) Pb(s).
•	oncentration of lead		neasured to be +0.140 V. nilligrams per Liter
(A) 1.2 mg/L	(B) 550 mg/L	(C) 3.9 mg/L	(D) 3.9x10 <sup>-3</sup> mg/L
for this reaction i Based upon elec	s given by: O <sub>2</sub> + 2 I ctrochemical conside corrode (i.e. undergo	$H_2O + 4 e^- \rightarrow 4 OH$ erations, which of t	and the reduction potetial $F_{red}^{o}(O_2) = +0.40 \text{ V}$ he following metals would presence of $O_2$ and
(A) Cu only	(B) Hg & Cu	(C) Sn & Cu	(D) Ag & Hg
For #47 - #48: Use Aqueous Solution)			ation Potentials in
	Fluoride, ZnF₂(aq) will be the principal		ctrolysis cell, and a voltage

ge

(A) Zn,  $F_2$ 

- (B) Zn, O<sub>2</sub>, H<sup>+</sup>
- (C)  $F_2$ ,  $H_2$ ,  $OH^-$

(C)  $H_2$ ,  $OH^-$ ,  $O_2$ ,  $H^+$ 

48. If aqueous Aluminum Iodide, AlI<sub>3</sub>(aq) is placed in an electrolysis cell, and a voltage is applied, what will be the principal products of the electrolysis?

(C) I<sub>2</sub> , H<sub>2</sub> , OH

- (B) Al,  $O_2$ ,  $H^{\dagger}$
- (C)  $H_2$ ,  $OH^-$ ,  $O_2$ ,  $H^+$



49.	. Approximately how long would it take to electroplate a metal surface with 0.15 g
•	of Nickel [M=58.7] metal from a Ni(NO <sub>3</sub> ) <sub>2</sub> (aq) solution with a current of 150 mA
	(milliAmps)?

54.8 min )

(B) 157 min

(C) 27.5 min

(D) 32.9 min

50. A total of 850 kJ of energy was required to plate out Al(s) [M=27.] by electrolysis of a Al(NO<sub>3</sub>)<sub>3</sub>(aq) solution. The voltage was 8. Volts. Approximately how many grams of Al(s) were plated out by electrolysis?

(A) 89.2 g

(B) 9.9 a

(C) 5.4 q

(D) 29.7 g

51. Consider the nuclear reaction,  ${}^{252}_{98}Cf + X \rightarrow 3\,{}^1_0n + {}^{259}_{103}Lr$ . What is X in this equation?

(A) <sup>4</sup><sub>2</sub>He

(B) <sup>9</sup><sub>5</sub>B

(C) 16<sub>8</sub>O

52. What nuclide will undergo electron capture to form Pt-195?

(A) Ir-196

(B) Ir-195

(D) Pt-196

53. Which of the following decay paths is the most likely one for Rn-222?

### SKIP this Question

(A) 
$$_{86}^{222} Ra \rightarrow_{+1}^{0} e +_{85}^{222} At$$
  
(C)  $_{86}^{222} Ra \rightarrow_{-1}^{0} e +_{87}^{222} Fr$ 

$$(B)$$
  $_{86}^{222} Ra \rightarrow_{2}^{4} He +_{84}^{218} Po$ 

(C) 
$${}^{222}_{86}Ra \rightarrow {}^{0}_{-1}e + {}^{222}_{87}Fr$$

(D) 
$$_{86}^{222} Ra +_{-1}^{0} e \rightarrow_{85}^{222} At$$

54. Which of the following is/are likely decay paths for Mg-22. Stable isotopes in this range typically have N/Z = 1.05.



$$(2)^{22}_{12}Mg \rightarrow {}^{0}_{+1}e + {}^{22}_{11}Na$$

Two more MC questions on next page.



- 55. One nuclear fusion reaction involves the reaction of a deuterium and tritium nucleus to form helium:  ${}_1^2H + {}_1^3H \rightarrow {}_2^4He + {}_0^1n$ . This reaction is highly exothermic because:
  - (A) The n-n repulsions in deuterium and tritium are higher than in helium
  - (B) Helium has a lower Binding Energy per nucleon than deuterium or tritium
  - (C) The p-p attractions are greater in helium than in deuterium or tritoi,
  - (D) Helium has a higher Binding Energy per nucleon than deuterium or tritium
- 56. Use the Molar Masses below to calculate the approximate Binding Energy per Nucleon (Eb/N) of Pb-208.

 $m(_1^1H) = 1.008 \text{ g/mol}$ ,  $m(_0^1n) = 1.009 \text{ g/mol}$ ,  $m(_{82}^{208}Pb) = 207.977 \text{ g/mol}$ 

(A) 1.6x10<sup>11</sup> kJ/mol

(B) 7.8x10<sup>11</sup> kJ/mol

(C) 7.8x108 kJ/mol

(D) 1.6x10<sup>14</sup> kJ/mol

Version B

CHEM 1423 - Final Exam - May 12, 2015 ,

Name Saladun

If you wish to have your final exam and course grade posted on the Web site, please provide me with a four (4) digit number which will be the ID number for your grade.

Four (4) digit number for posting.

**Problem (4 pts)**: A concentration cell is prepared with 0.20 M Calcium Nitrate, Ca(NO<sub>3</sub>)<sub>2</sub>, in the reference compartment (cathode) and a saturated solution of Calcium Phosphate, Ca<sub>3</sub>(PO<sub>4</sub>)<sub>2</sub>, in the sample compartment (anode).

The cell reaction can be written as: Ca(s)|Ca<sup>2+</sup>(xx M)||Ca<sup>2+</sup>(0.20 M)|Ca(s)

The measured cell voltage is +0.146 V. Calculate the Solubility Product, K<sub>sp</sub>, of Ca<sub>3</sub>(PO<sub>4</sub>)<sub>2</sub>

1) Calculati [ 6 1 ] = XX ] = 0 - 0,0192 / 07 4X 12(xx) = -NF = -2(0,146) = -4.93 2x = 10 - 0.493 = 1.07×10 - 5617 = xx = 2.34×10 } 6) Calubte Kip/ Ca3(Aby) = 36 to 2 Pay [aV]= 35= 234×106 5 = 2 79 x10 -7  $R_{\text{sp}}^{2} = \frac{5a^{3}}{3.40} = \frac{2.34 \times 10^{2}}{3.40}$   $= \frac{3.1 \times 10^{-29}}{3.40} = \frac{3.40}{3.40}$ Spa; ]=252 1.56×10

(56)	MULTIPLE CHOICE (Mark the one correct answer to each question on
	your scantron)

Turn in: (a) Your scantron with your name and answers (there is no need to bubble in your ID.

(b) The cover sheet with your Electrochemistry Problem + four (4) digit number if you would like your results posted on the course web site.

Each Multiple Choice question is worth 1 point. The problem is worth 4 points, yielding a total of 60 points on the test. Your score will be converted to a percentage prior to any further analysis.

1.	The reaction, $A \rightarrow Products$ , is of order "n" with respect to [A]; i.e.	Rate = $k[A]^n$
	When the initial concentration of A is 0.60 M, the initial rate is 0.50	Ms <sup>-1</sup> . When
	the initial concentration of A is 0.20 M, the initial rate is 4.50 Ms <sup>-1</sup> .	The order of
	this reaction, n, is:	_

(A) +2

(B) -1

(C) +1

(D) -2

The rate law for a given reaction, A → Products, is **fourth** order with respect to [A]? When the initial concentration of A is 0.80 M, the initial rate is 0.45 Ms<sup>-1</sup>. The rate constant for this reaction is approximately:

(A) 1.4 M<sup>-3</sup>s<sup>-1</sup>

(B) 1.1 M<sup>-3</sup>s<sup>-1</sup>

(C) 0.91 M<sup>-3</sup>s<sup>-1</sup>

(D) 0.73 M<sup>-3</sup>s<sup>-1</sup>

3. For the **first** order reaction, A → Products, the rate constant is 0.025 s<sup>-1</sup>. If the initial concentration of A is 0.50 M, what is the approximate concentration of A after 20 s?

(A) 0.40 M

(B) 0.26 M.

(C) 0.30 M

(D) 0.61 M

4. The rate of the chemical reaction involving two substances, A and B, is measured. It is found that if the initial concentration of A used is tripled, keeping the B concentration the same, the rate increases by a factor of nine (9) (relative to the first experiment). If the concentrations of both A and B are doubled, the rate increases by a factor of thirty-two (32) (relative to the first experiment). The rate law for this reaction is: Rate =

(A)  $k[A]^2[B]^2$ 

(B)  $k[A][B]^3$ 

(C)  $k[A]^3[B]^2$ 

(D) k[A]<sup>2</sup>[B]<sup>3</sup>

5. Consider the hypothetical reaction,  $3A + B \rightarrow 2C$ . If the rate of change of [A] is  $\Delta[A]/dt = -0.60 \text{ M hr}^{-1}$ . What is the "rate" of the reaction?

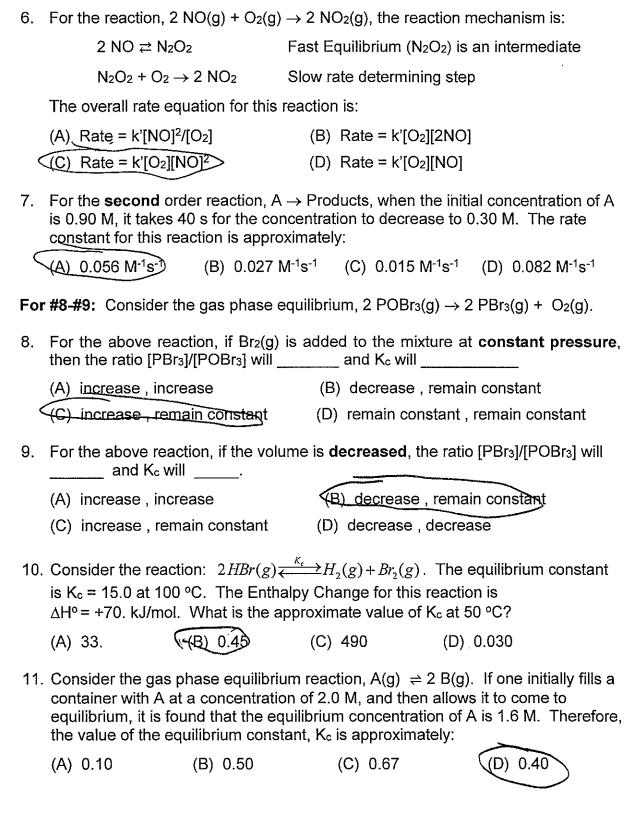
(A)  $-0.40 \text{ M hr}^{-1}$ 

(B)  $+0.40 \text{ M hr}^{-1}$ 

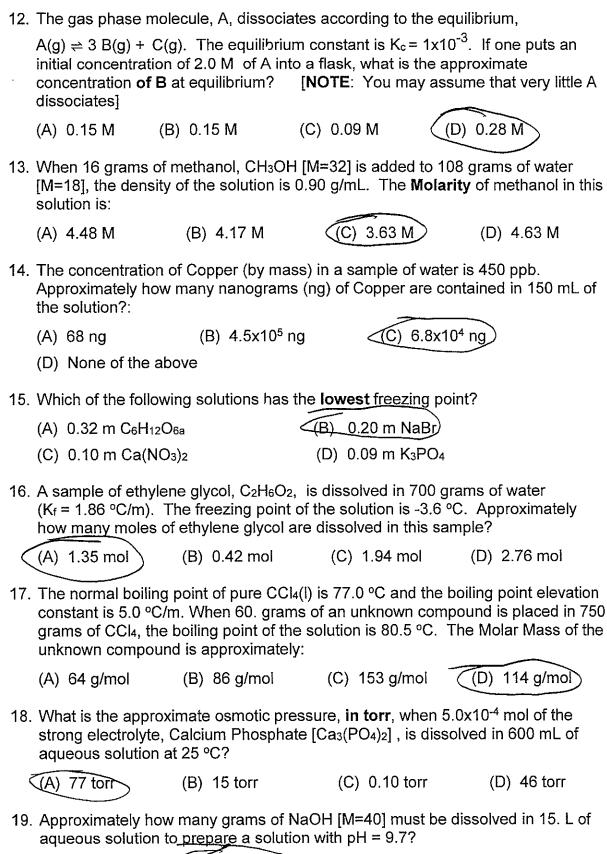
(C) -0.20 M hrr

(D) +0.20 M hr<sup>-1</sup>









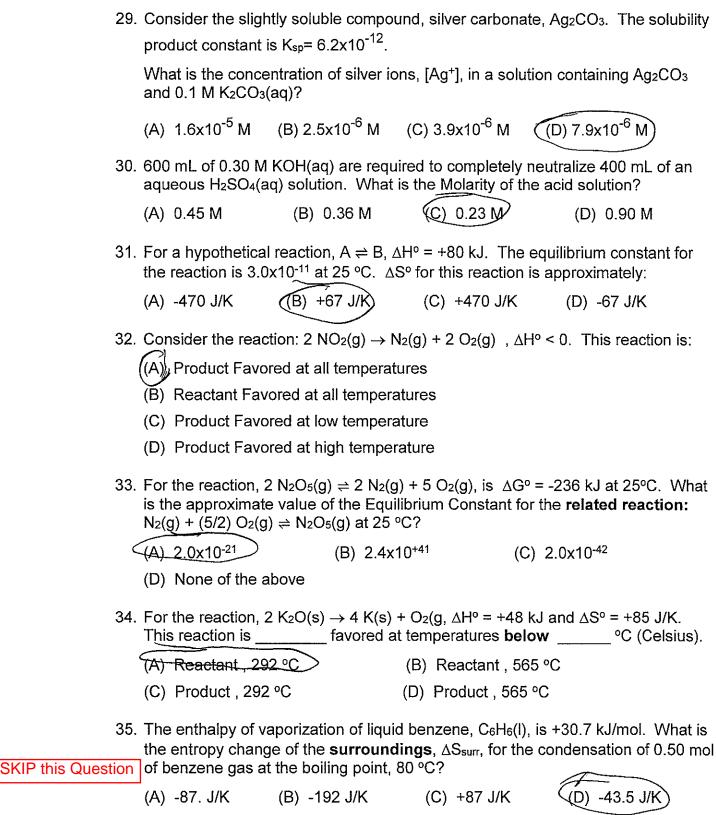
(B)  $3.0 \times 10^{-2}$  a)

(A)  $7.5 \times 10^{-2}$  g

(C)  $2.0x10^{-3}$  g

(D)  $7.5 \times 10^{-4}$  g

20.			-	onstant, $K_b = 4.3 \times 10^{-10}$ . in chloride (AnilHCl)?			
	(A) 11.0	(B) 3.0	(C) 5.3	(D) 8.7			
21.	,	•		t of 1.6x10 <sup>-9</sup> . What is the enzoate (NaBenz)?			
	(A) 9.3	(B) 3.0	(C) 11.0	(D) 4.7			
22.	2. The pH of a 0.10 M acetate (KAc) is 8.90 . The base equilibrium constnat, $K_b$ of the acetate ion (Ac <sup>-</sup> ) is approximately:						
	(A) 1.6x10 <sup>-5</sup>	(B) 6.3x10 <sup>-10</sup>	(C) 1.6x10 <sup>-17</sup>	(D) 4.8x10 <sup>-7</sup>			
23.	If added to 2 L of	0.80 M NaOH, whi	ch one of the follow	ring would form a buffer?			
	(A) 2. L of 0.50 M	¹ Nitric Acid (HNO₃	)				
(	(B) 2). L of 1.0 M	Lactic Acid (HLac)					
	(C) 2. L of 0.50 M	Acetic Acid (HAc	)				
	(D) 2. L of 1.0 M	Potassium Acetate	e (KAc)				
For #24 - #28: Tellurous acid, $H_2TeO_3$ , is a diprotic acid with acid dissociation constants, $K_a$ ' = $3.0x10^{-3}$ and $K_a$ " = $2.0x10^{-8}$							
24.	What is the pH of	a solution containi	ng 0.20 M KHTeO₃	and 0.50 M Na₂TeO₃?			
<	(A) 8.10	(B) 7.30	(C) 2.92	(D) 2.12			
25.	What is the pH of 2.0 L of 0.50 M H	• •	d by adding 0.35 m	ol of KOH to			
	(A) 7.43	(B) 2.79	(C) 2.25	(D) 2.06			
26.	26. What is the pH of a solution prepared by adding 2 L of 0.70 M HCl to 2 L of 0.45 M Na₂TeO₃ ?						
	(A) 7.60	(B) 2.62	(C) 7.80	(D) 2.42			
27. What is the pH of a 0.04 M solution of sodium tellurite, Na <sub>2</sub> TeO <sub>3</sub> ?							
£ 1 .	What is the pH of	a 0.04 M solution	of sodium tellurite,	Na₂TeO₃ ?			
<	What is the pH of (A) 10.15	a 0.04 M solution (B) 9.45	of sodium tellurite, (C) 11.25	Na₂TeO₃ ? (D) 3.85			
<	(A) 10.15		(C) 11.25				



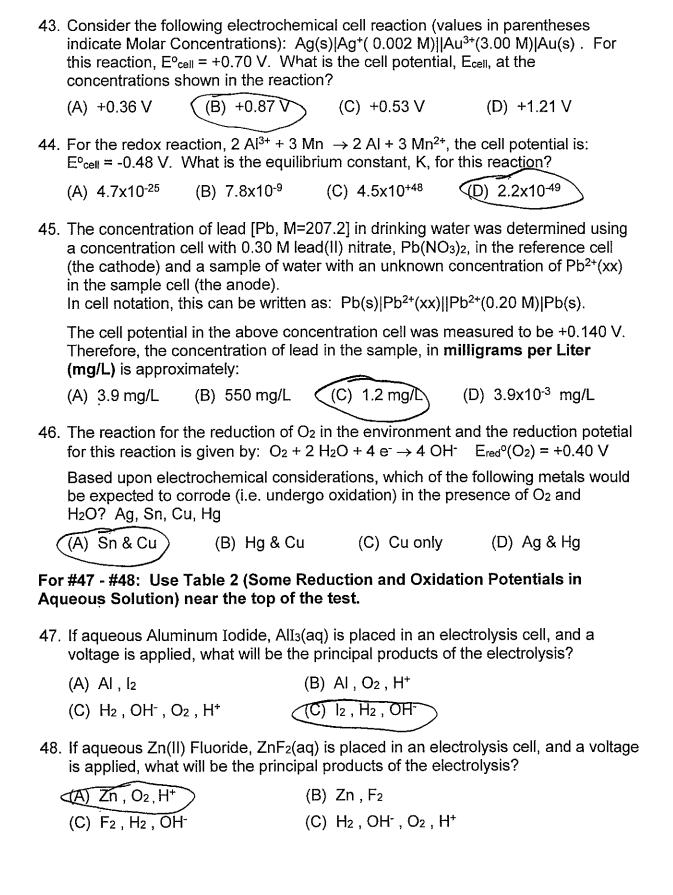


(D) +152 kJ

			t 25 °C and the Ent thalpy change for t	ropy change is -95 J/K. his reaction?
(A)	ΔH < 45.6 kJ	(B) ∆H :	> +28.3 J/K	(C) $\Delta H < -28.3 \text{ kJ}$
(D)	ΔH cannot be	determined withou	t knowing the sign	of ΔG
at 2		s the approximate v		orium constant, $K = 220$ , ${}^{\circ}C$ when $P(N_2) = P(H_2) =$
(A)	+10.4 kJ	(B) -7.4 kJ	(C) -33.4 kJ	(D) -56.4 kJ
	the electroche tements is/are		Cu Cu²+  Au³+ Au,	which of the following
₹ (2)	Electrons flow circuit		ode to the Cu elect	rode through an external
$\lambda(4)$				, NO₃⁻ ions flow towards
(A)	1 & 3	(B) 1 & 2 & 4	(C) 1 & 2 & 3	(D) 2 & 3 & 4
39. Re	garding the folk	owing reaction, whi	ich of the statemen	ts below is/are correct?
_	Fe <sub>2</sub> C	$O_3(s) + 3 CO(g) \rightarrow$	2 Fe + 3 CO <sub>2</sub> (g)	
		xidizing agent e transfered	✓(2) Fe is reduction ✓(4) CO is oxidi	
	3 & 4	(B) 1 & 4	(C) 1&3&4	(D) 2 & 3
	- #46: Use Ta of the test.	able 1 (Standard F	Reduction Potenti	als), as necessary, near
			2 K + Cd <sup>2+</sup> , the cell on potential for Cd <sup>2</sup>	
(A)	+0.40 V	(B) -5.46 V	(C) -3.33 V	(D) -0.40 V
41. Wh	ich of the follov	ving reactions are	reactant favored?	
SKIP this Question (3)	Hg <sup>2+</sup> + 2 Cl <sup>-</sup>	Hg + Cl <sub>2</sub> Pol (	2) $2 \text{ Ag}^+ + 2 \text{ I}^- \rightarrow 2$	. Ag + 2 l <sub>2</sub>
Prof (3)	$Hg^{2+} + 2 Fe^{2+}$	$\rightarrow$ Hg + 2 Fe <sup>3+</sup>	4) $Cu^{2+} + 2 Ag \rightarrow 2$	2 Ag+ + Cu
(A)	2 & 3	(B) 1 & 4	(C) 3 only	(D) 1 & 3 & 4
	at is the standath $^{+}$ + 2 l $^{-}$ $\rightarrow$ Ni +		ergy change for the	electrochemical reaction,

(A) -56 kJ (B) +76 kJ (C) +56 kJ







49. A total of 850 kJ of energy was required to plate out Al(s) [M=27.] by electr	olysis
of a Al(NO <sub>3</sub> ) <sub>3</sub> (aq) solution. The voltage was 8. Volts. Approximately how r	nany
grams of Al(s) were plated out by electrolysis?	·

- (A) 89.2 g
- (B) 5.4 g
- (C) 9.9 g
- (D) 29.7 g
- 50. Approximately how long would it take to electroplate a metal surface with 0.15 g of Nickel [M=58.7] metal from a Ni(NO<sub>3</sub>)<sub>2</sub>(aq) solution with a current of 150 mA (milliAmps)?
  - (A) 32.9 min
- (B) 157 min
- (C) 27.5 min
- 51. What nuclide will undergo electron capture to form Pt-195?
  - (A) Ir-196
- (B) Ir-195
- (C) Pt-196
- (D) Au-195
- 52. Which of the following decay paths is the most likely one for Rn-222?

## SKIP this Question

(A) 
$$_{86}^{222}Ra \rightarrow_{+1}^{0} e +_{85}^{222} At$$

(B) 
$$_{86}^{222} Ra \rightarrow_{2}^{4} He +_{84}^{218} Po$$

(C) 
$$^{222}_{86}Ra \rightarrow^{0}_{-1} e +^{222}_{87} Fr$$

(D) 
$$_{86}^{222} Ra +_{-1}^{0} e \rightarrow_{85}^{222} At$$

- 53. Consider the nuclear reaction,  ${}^{252}_{98}Cf + X \rightarrow 3 {}^{1}_{0}n + {}^{259}_{103}Lr$ . What is X in this equation?
  - (A) 42He
- (B)  $^{10}5B$
- $(C)^{-16}8O$
- (D) 8<sub>5</sub>B
- 54. Which of the following is/are likely decay paths for Mg-22. Stable isotopes in this range typically have N/Z = 1.05.
  - (1)  $^{22}_{12}Mg \rightarrow ^{0}_{-1}e + ^{22}_{13}Al$
- (2)  $^{22}_{12}Mg \rightarrow ^{0}_{+1}e + ^{22}_{11}Na$
- (3)  $^{22}_{12}Mg + ^{0}_{-1}e \rightarrow ^{22}_{11}Na$
- (4)  ${}_{12}^{22}Mg \rightarrow {}_{2}^{4}He + {}_{10}^{18}Ne$

- (A) 1 & 4 (B) 4 only
- (D) 1&2

Two more MC questions on next page.

55. Use the Molar Masses below to calculate the approximate Binding Energy per Nucleon (Eb/N) of Pb-208.

 $m(_1^1H) = 1.008 \text{ g/mol}$ ,  $m(_0^1n) = 1.009 \text{ g/mol}$ ,  $m(_{82}^{208}Pb) = 207.977 \text{ g/mol}$ 

(A) 1.6x10<sup>14</sup> kJ/mol

(B) 7.8x10<sup>11</sup> kJ/mol

(C) 7.8x108 kJ/mol

- (D) 1.6x10<sup>11</sup> kJ/mol
- 56. One nuclear fusion reaction involves the reaction of a deuterium and tritium nucleus to form helium:  ${}_1^2H + {}_1^3H \rightarrow {}_2^4He + {}_0^1n$ . This reaction is highly exothermic because:
  - (A) The n-n repulsions in deuter um and tritium are higher than in helium
  - (B) Helium has a higher Binding Energy per nucleon than deuterium or tritium
  - (C) The p-p attractions are greater in helium than in deuterium or tritoi,
  - (D) Helium has a lower Binding Energy per nucleon than deuterium or tritium