CHEM 1423 - Final Exam – May 9, 2017

Name\_\_\_\_\_

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Please turn in:

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You can keep the test (below) and use it to compare results with the answer key.

**Conversions:** 1 atm. = 760 torr

Constants: R = 0.082 L-atm/mol·K R = 8.31 J/mol·K R = 8.31x10<sup>-3</sup> kJ/mol·K N<sub>A</sub> =  $6.02x10^{23}$  mol<sup>-1</sup> 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]



### **ELECTROCHEMISTRY INFORMATION**

### Table 1: Standard Reduction Potentials

## Reduction Half-Reactions E<sup>o</sup>(V)

$F_2 + 2 e^- \rightarrow 2 F^-$	+2.87
$Au^{3+} + 3 e^- \rightarrow Au$	+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
$I_2 + 2 e^- \rightarrow 2 I^-$	+0.54
$Cu^{2+} + 2 e^{-} \rightarrow Cu$	+0.34
$Sn^{2+} + 2 e^{-} \rightarrow Sn$	-0.14
Fe <sup>3+</sup> + 3 e <sup>-</sup> 🛛 Fe	-0.04
$Ni^{2+} + 2 e^{-} \rightarrow Ni$	-0.25
$Fe^{2+} + 2e^{-} \rightarrow Fe$	-0.44
$Zn^{2+} + 2 e^{-} \rightarrow Zn$	-0.76
$Mn^{2+} + 2 e^{} \rightarrow Mn$	-1.18
$AI^{3+} + 3 e^{-} \rightarrow AI$	-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**

$2 \text{ H}_2\text{O} + 2 \text{ e}^- \rightarrow \text{H}_2 + 2 \text{ OH}^-$	$E^{o}_{red}$ = -0.83 V
$AI^{3+} + 3 e^- \rightarrow AI$	$E^{o}_{red}$ = -1.66 V
$Zn^{2+} + 2 e^{-} \rightarrow Zn$	$E^{o}_{red}$ = -0.76 V
Mg <sup>2+</sup> + 2 e <sup>-</sup> [] Mg	$E^{o}_{red}$ = -2.37 V
Na⁺ + 3 e⁻ 🛛 Na	$E^{o}_{red}$ = -2.71 V
Fe <sup>2+</sup> + 2 e <sup>-</sup> [] Fe	$E^{o}_{red}$ = -0.44 V

#### **Oxidation Potentials**

$2 \text{ H}_2\text{O} \rightarrow \text{O}_2 + 4 \text{ H}^+ + 4 \text{ e}^-$	$E^{o}_{oxid}$ = -1.23 V
$2 \text{ I}^{-} \rightarrow \text{I}_2 + 2 \text{ e}^{-}$	$E^{o}_{oxid}$ = -0.54 V
2 Br <sup>-</sup> [] Br <sub>2</sub> + 2 e <sup>-</sup>	$E^{o}_{oxid}$ = -1.07 V
$2 F^- \rightarrow F_2 + 2 e^-$	$E^{o}_{oxid}$ = -2.87 V

#### **Some Electrochemical Equations**

- $F = 96,500 \text{ C/mol e}^{-}$  (Coulombs per mole of electrons)
- 1 J = 1 CxV [i.e. 1 Joule = 1 Coulomb x Volt]

 $\Delta G^{\circ} = -nFE^{\circ}$ 

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

Q = i x t i.e. Charge (in Coul) = Current (in Amps = Coul/sec) x time (in sec)]

E = Q x V i.e. Energy (in J) = Charge (in Coulombs) x Voltage (in Volts) (note: 1 Coulomb-Volt = 1 Joule)

# (60) 60 QUESTIONS (Mark the one correct answer to each question on your scantron)

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Each Question is worth 1 point. Your score will be converted to percent by:
Score = XX/60 x 100 (where XX is the number of correct answers)
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1. For the reaction, A + B  $\rightarrow$  Products, the rate law is:  $Rate = k \frac{[C]^2}{[B]}$  The units of the rate constant are:

(A) s<sup>-1</sup> (B) M<sup>2</sup>s<sup>-1</sup> (C) M<sup>-2</sup>s<sup>-1</sup> (D) M<sup>-1</sup>s<sup>-1</sup>

2. 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 9 (relative to the first experiment). If the concentrations of both A and B are doubled, the rate increases by a factor of 32 (relative to the first experiment). The rate law for this reaction is: Rate =

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

Consider a reaction which is zeroth order; i.e. d[A]/dt = -k[A]<sup>0</sup> = -k For this reaction, a plot of \_\_\_\_\_ vs. time is a straight line with a \_\_\_\_\_ slope.

(A)	In([A]t) , negative	(B)	[A]t , negative
(C)	(A] <sub>t</sub> , positive	(D)	1/[A] <sub>t</sub> , positive

- 4. The reaction,  $A \rightarrow Products$ , is of order "n" with respect to [A]; i.e. Rate = k[A]<sup>n</sup>. When the initial concentration of A is 0.20 M, the initial rate is 1.5 Ms<sup>-1</sup>. When the initial concentration of A is 0.60 M, the initial rate is 40.5 Ms<sup>-1</sup>. The order of this reaction, n, is:
  - (A) -2 (B) +1 (C) +2 (D) +3

For #5 - #6: Consider a **second** order reaction,  $A \rightarrow$  Products. The rate constant for this reaction is 0.02 M<sup>-1</sup>s<sup>-1</sup>. The initial concentration of A is 0.75 M

- 5. Approximately how long will it take for the reactant concentration to decrease to 0.20 M?
  - (A) 45 s (B) 100 s (C) 180 s (D) 80 s
- 6. What will be the concentration of A 60 s after the start of the reaction?
  - (A) 0.39 M (B) 0.23 M (C) 0.47 M (D) 0.56 M

For the reaction, Sc<sup>3+</sup>(aq) + 2 Ni<sup>+</sup>(aq) → Sc<sup>+</sup>(aq) + 2 Ni<sup>2+</sup>(aq), the reaction mechanism is:

$$2Ni^+ \xleftarrow{K} Ni + Ni^{2+}$$
 Fast equilibrium

$$Ni + Sc^{3+} \xrightarrow{k} Ni^{2+} + Sc^{+}$$
 Slow step

The overall rate equation for this reaction is:

(A) 
$$Rate = k' \frac{[Ni^{2+}][Sc^{3+}]}{[Ni^{+}]^{2}}$$
 (B)  $Rate = k' \frac{[Ni^{+}][Sc^{3+}]}{[Ni^{2+}]}$   
(C)  $Rate = k' \frac{[Ni^{+}]^{2}[Sc^{3+}]}{[Ni^{2+}]}$  (D)  $Rate = k'[Ni][Sc^{3+}]$ 

8. The gas phase equilibrium, 2 A(g) = 3 B(g) + C(g). The equilibrium constant is

 $K_c = 1 \times 10^{-4}$ . If one puts an initial concentration of 2.0 M of A into a flask, what is the approximate concentration of B at equilibrium? [NOTE: You may assume that very little A reacts]

- (A) 0.16 M (B) 0.19 M (C) 0.28 M (D) 0.06 M
- 9. Consider the gas phase equilibrium reaction,  $2 A(g) \rightleftharpoons B(g)$ . If one initially fills a container with A at a concentration of 3.0 M, and then allows it to come to equilibrium, it is found that the equilibrium concentration of B is 1.2 M. Therefore, the value of the equilibrium constant, K<sub>c</sub> is approximately:
  - (A) 3.3 (B) 2.0 (C) 0.67 (D) 0.37
- 10. The equilibrium constant for the reaction, N<sub>2</sub>(g) + O<sub>2</sub>(g) = 2 NO(g) is 1.7x10<sup>-3</sup> (at 2300 K). What is the equilibrium constant for the reaction, 6 NO(g) = 3 N<sub>2</sub>(g) + 3 O<sub>2</sub>(g) ?
  - (A) 4.9x10<sup>-9</sup> (B) 8.4 (C) 1.7x10<sup>3</sup> (D) 2.0x10<sup>8</sup>

**For #11 - #13:** Consider the gas phase equilibrium,  $4NO_2(g) + O_2(g) \rightleftharpoons 2N_2O_5(g)$ . This is an Exothermic reaction.

- 11. For the above reaction, if O<sub>2</sub>(g) is added to the mixture, then the ratio, [NO<sub>2</sub>]/[N<sub>2</sub>O<sub>5</sub>] will \_\_\_\_\_ and K<sub>c</sub> will \_\_\_\_\_
  - (A) decrease, Increase (B) increase, decrease
  - (C) decrease, remain constant (D) increase, remain constant

12. For the above reaction, if Ne(g) is added to the mixture in a container at fixed **total pressure**, the ratio, [NO<sub>2</sub>]/[N<sub>2</sub>O<sub>5</sub>] will \_\_\_\_\_ and

Kc will \_\_\_\_\_

(A) decrease, Increase

- (B) increase, decrease
- (C) decrease, remain constant
- (D) increase, remain constant
- decrease, remain constant (i
- 13. For the above reaction, if the temperature is decreased,
  - (A) the equilibrium will move to the left and K will decrease
  - (B) the equilibrium will move to the right and K will increase
  - (C) the equilibrium will move to the left and K will remain constant
  - (D) the equilibrium will move to the right and K will remain constant
- 14. The weight percent of Thalium in a sample of water is 3x10<sup>-11</sup> %. Therefore, the concentration of Thalium, in parts per trillion (ppt) is:
  - (A) 0.30 ppt (B) 3x10<sup>-2</sup> ppt (C) 3x10<sup>-4</sup> ppt (D) 3.0 ppt
- 15. What is the molality of a solution prepared by adding 138 grams of Glycerol (C<sub>3</sub>H<sub>8</sub>O<sub>3</sub>, M=92.) to 600 grams of water?
  - (A) 2.033 molal (B) 3.75 molal (C) 2.5 molal
  - (D) Determination of the molality requires the density of the soluton
- 16. The density of a solution prepared by adding 93 grams of Ethylene Glycol (C<sub>3</sub>H<sub>6</sub>O<sub>2</sub>, M=62) to 400 grams of water is 1.30 g/mL. What is the approximate Molarity of this solution?
  - (A) 3.95 Molar (B) 3.75 Molar (C) 4.95 Molar (D) 3.45 Molar
- 17. Which of the following aqueous solutions has the lowest boiling point?
  - (A) 0.20 m K<sub>3</sub>PO<sub>4</sub> (B) 0.30 m Ca(NO<sub>3</sub>)<sub>2</sub>
  - (C)  $0.70 \text{ m } \text{C}_6\text{H}_{12}\text{O}_6$  (D) 0.32 m NaBr
- 18. When a sample of the strong electrolyte, aluminum nitrate, Al(NO<sub>3</sub>)<sub>3</sub>, is dissolved in 600 mL of aqueous solution, the osmotic pressure at 25 °C is 350 torr. Approximately how many moles of Al(NO<sub>3</sub>)<sub>3</sub> are dissolved in the 600 mL?
  - (A) 1.1x10<sup>-2</sup> mol (B) 2.8x10<sup>-3</sup> mol (C) 7.9x10<sup>-3</sup> mol (D) 3.9x10<sup>-3</sup> mol
- 19. The normal boiling point of pure CCl<sub>4</sub>(I) is 77.0 °C and the boiling point elevation constant is 5.0 °C/m. When 80. grams of an unknown compound is placed in 750 grams of CCl<sub>4</sub>, the boiling point of the solution is 80.5 °C. The Molar Mass of the unknown compound is approximately:
  - (A) 86 g/mol (B) 69 g/mol (C) 152 g/mol (D) 114 g/mol

- 20. A 300. L sample of nitric acid (HNO<sub>3</sub>, M=63) has a **pOH** of 11.5. Approximately how many grams of nitric acid are contained in the sample?
  - (A) 60. grams (B) 0.20 grams (C) 0.62 grams (D) 23. grams
- 21. Which of the following aqueous solutions is/are basic (pH > 7)?
  - (i) Potassium Lactate (KLac)
  - (ii) Sodium Nitrate (NaNO<sub>3</sub>)
  - (iii) Pyridinium Bromide (PyrHBr)
  - (iv) Sodium Propanoate (NaProp)
  - (A) i & iv (B) ii & iii (C) i & ii & iv (D) iv only
- 22. The pH of a 0.10 M solution of hydrocyanic acid, HCN, is 5.10. Therefore, the acid dissociation constant, K<sub>a</sub>, for this acid is:
  - (A)  $6.8 \times 10^{-12}$  (B)  $7.9 \times 10^{-5}$  (C)  $6.3 \times 10^{-10}$  (D)  $4.8 \times 10^{-7}$
- 23. For the weak acid, Hypobromous acid, HBrO, the acid ionization constant is 2.0x10<sup>-9</sup>. What is the approximate pH of a 0.05 M solution of this acid?
  - (A) 9.0 (B) 5.0 (C) 3.4 (D) 4.6
- 24. The weak base, aniline (Anil), has a base equilibrium constant,  $K_b = 4.3 \times 10^{-10}$ . What is the pH of a 0.05 M aqueous solution of anilinium chloride (AnilHCl)?
  - (A) 11.0 (B) 8.7 (C) 5.3 (D) 3.0
- 25. If added to 2 L of 0.40 M HNO<sub>3</sub>, which of the following would form a buffer?
  - (1) 0.6 mol of potassium lactate (KLac)
  - (2) 0.6 mol of sodium carbonate (Na<sub>2</sub>CO<sub>3</sub>)
  - (3) 1.2 mol of sodium acetate (NaAc)
  - (4) 1.2 mol of ammonium bromide (NH<sub>4</sub>Br)
  - (A) 1 & 2 & 3 (B) 2 & 3 & 4 (C) 3 only (D) 2 & 3

**For #26 - #30:** Consider Carbonic acid,  $H_2CO_3$ , which is a diprotic acid with acid dissociation constants, Ka' =  $4.2 \times 10^{-7}$  and Ka'' =  $4.8 \times 10^{-11}$ .

- 26. What is the approximate pH of a 0.05 M solution of sodium carbonate, Na<sub>2</sub>CO<sub>3</sub>?
  - (A) 5.8 (B) 11.5 (C) 10.2 (D) 8.2
- 27. What is the approximate pH of a solution containing 0.25 M K<sub>2</sub>CO<sub>3</sub> and 0.40 M NaHCO<sub>3</sub> ?
  - (A) 10.5 (B) 9.4 (C) 10.1 (D) 6.6

- 28. What is the approximate pH of a solution prepared by adding 0.70 mol of HNO<sub>3</sub> to 4 L of 0.40 M Na<sub>2</sub>CO<sub>3</sub>?
  - (A) 10.4 (B) 10.2 (C) 5.5 (D) 10.7
- 29. What is the approximate pH of a solution prepared by adding 2.40 mol of NaOH to 5 L of 0.40 M H<sub>2</sub>CO<sub>3</sub> ?
  - (A) 5.9 (B) 10.9 (C) 9.0 (D) 9.7
- 30. What is the approximate ratio of concentrations, [H<sub>2</sub>CO<sub>3</sub>]/[HCO<sub>3</sub><sup>-</sup>], of a buffer solution with pH = 6.80?
  - (A) 2.7 (B) 1.5 (C) 0.4 (D) 0.7
- 31. 600 mL of 0.40 M NaOH(aq) is needed to completely neutralize a sample of aqueous H<sub>3</sub>PO<sub>4</sub>(aq). Approximately how many grams of H<sub>3</sub>PO<sub>4</sub> (M=98 g/mol) were in the sample?
  - (A) 70.6 g (B) 7.8 g (C) 23.5 g
  - (D) Cannot be determined without the volume of the H<sub>3</sub>PO<sub>4</sub> sample
- 32. Consider the slightly soluble salt, AB<sub>3</sub>. The solubility product constant for AB<sub>3</sub> is  $K_{sp} = 6.0 \times 10^{-23}$ . What is the solubility of AB<sub>3</sub> in a solution containing 0.05 M NaB(aq) (which is a strong electrolyte)?
  - (A) 1.8x10<sup>-20</sup> M (B) 7.4x10<sup>-12</sup> M (C) 4.8x10<sup>-19</sup> M (D) 1.2x10<sup>-21</sup> M
- 33. The enthalpy of vaporization of toluene is 39.2 kJ/mol and the normal boiling point is 111 °C.. What is the entropy change of the **system** when **2 (two) moles** of toluene gas condenses at 111 °C?
  - (A) -204 J/K (B) -102 J/K (C) -706 J/K (D) +102 J/K
- 34. The enthalpy of vaporization of liquid benzene, C<sub>6</sub>H<sub>6</sub>(I), is +30.7 kJ/mol. What is the entropy change of the **surroundings** for the condensation of 0.50 mol of benzene gas at the boiling point, 80 °C?
  - (A) -43.5. J/K (B) -87 J/K (C) +87. J/K (D) +43.5 J/K

35. For the reaction,  $CH_3OH(g) \longrightarrow CO(g) + 2H_2(g)$ ,  $\Delta H^{\circ} = +91$  kJ and

 $\Delta S^{\circ} = +221 \text{ J/K}$ . What is the entropy change of the **surroundings**,  $\Delta S_{\text{surr}}$ , for the related reaction,  $2CO(g) + 4H_2(g) \rightleftharpoons 2CH_3OH(g)$ , at 25 °C?

(A) +611 J/K (B) -442 J/K (C) -611 J/K (D) +442 J/K

- 36. For the hypothetical reaction, A = B,  $\Delta S^{\circ} = -70 \text{ J/K}$  (independent of temperature). The equilibrium constant for the reaction at 150 °C is 2.0x10<sup>-3</sup>. What is the enthalpy change,  $\Delta H^{\circ}$  for this reaction?
  - (A) +51.5 kJ (B) -7.8 kJ (C) -37.3 kJ (D) -51.5 kJ
- 37. The reaction,  $A \rightarrow B$ , is **exergonic** at 25 °C and the entropy change is -20 J/K. What can be concluded about the enthalpy change for this reaction?
  - (B)  $\Delta H > +6 \text{ kJ}$ (C)  $\Delta H < -6 \text{ kJ}$ (A)  $\Delta H < -18 \text{ kJ}$
  - (D) No conclusion can be made about  $\Delta H$
- 38. The equilibrium constant for the reaction,  $4 N_2 O_5 = 4 N_2(q) + 10 O_2(q)$ , is 5.8x10<sup>82</sup> at 25 °C. What is the approximate value of the Gibbs Free Energy of Formation of  $N_2O_5(q)$ ?
  - (A) +120. kJ/mol (B) +240. kJ/mol (C) -120 kJ/mol

(D) Insufficient data is given to determine  $\Delta G_{f^0}(N_2O_5)$ 

For #39 - #40: The Enthalpy of Vaporization of Cl<sub>2</sub>(liq) is 10.2 kJ/mol. The Entropy of Vaporization of Cl<sub>2</sub>(liq) is 42.9 J/mol-K.

- 39. The Entropy change of the universe,  $\Delta S_{univ}$ , when **3 (three)** moles of Cl<sub>2</sub>(liq) vaporizes to Cl<sub>2</sub>(gas) at -60 °C is approximately:
  - (A) +5.0 J/mol-K (B) -15.0 J/mol-K (C) -5.0 J/mol-K
  - (D) None of the above
- 40. The Gibbs Energy Change,  $\Delta G^{\circ}$ , when **3** (three) moles of Cl<sub>2</sub>(gas) condenses to Cl<sub>2</sub>(liq) at -60 °C is approximately:
  - (A) +1.8 kJ/mol (B) +3.2 kJ/mol (C) -1.0 kJ/mol (D) -3.2 kJ/mol
- 41. Regarding the reaction,  $Br_2O_5(s) + 5 CO(g) \rightarrow Br_2(l) + 5 CO_2(g)$ , which of the following statements is/are true?
  - (1)  $Br_2O_5$  is the reducing agent (2) Br<sub>2</sub> is oxidized (3) CO is oxidized
    - (4) Br<sub>2</sub>O<sub>5</sub> is reduced
  - (D) 1 & 2 & 4 (A) 1 & 2 (C) 2 & 3 & 4 (B) 3 & 4

- 42. For the electrochemical cell given by Al|Al<sup>3+</sup>||Cu<sup>2+</sup>|Cu, which of the following statements is/are correct?
  - (1) The cathode reaction is  $Cu \rightarrow Cu^{2+} + 2e^{-}$
  - (2) Electrons flow from the AI electrode to the Cu electrode through an external circuit
  - (3) The anode reaction is  $AI \rightarrow AI^{3+} + 3e^{-}$
  - (4) If the half-cells are separated by a K<sub>2</sub>SO<sub>4</sub> salt bridge, SO<sub>4</sub><sup>2-</sup> ions flow towards the Cu electrode
  - (A) 2 & 3 (B) 1 & 3 & 4 (C) 1 & 2 & 3 (D) 1 & 4

#### For #43 - #50: Use Table 1 (Standard Reduction Potentials) as needed.

43. Which of the following reactions are **reactant** favored?

(1)	$CI^{-} CI_{2}   Hg^{2+}$	Hg	(2) I⁻   I₂    Ag⁺  Ag	
(3)	Fe   Fe <sup>2+</sup>    Ni <sup>2</sup>	+  Ni	(4) Ag   Ag+    Cu <sup>2+</sup>   Cu	
(A)	2&3	(B) 1 & & 3 & 4	(C) 3 only	(D) 1 & 4

44. For the electrochemical cell given by Hg|Hg<sup>2+</sup>||Be<sup>2+</sup>|Be , the cell potential is  $E^{o}_{cell} = -2.71 \text{ V}$ . What is the **reduction potential** of Be<sup>2+</sup>?

(A) -1.85 V (B) +3.57 V (C) -3.57 V (D) +1.85 V

.45. For the redox reaction given by  $Fe|Fe^{3+}||Cu^{2+}|Cu$ , the cell potential is:  $E^{o}_{cell} = +0.38 \text{ V}$ . What is the equilibrium constant, K, for this reaction?

(A) 
$$6.9 \times 10^{+12}$$
 (B)  $3.3 \times 10^{+38}$  (C)  $5.3 \times 10^{+16}$  (D)  $3.1 \times 10^{-39}$ 

- 46. What is the standard Gibbs Free Energy change for the electrochemical reaction, Ni<sup>2+</sup> + 2 l<sup>-</sup>  $\rightarrow$  Ni + l<sub>2</sub> ?
  - (A) -56 kJ (B) +56 kJ (C) +152 kJ (D) +76 kJ
- 47. Consider the following electrochemical cell reaction (values in parentheses indicate Molar Concentrations): Al(s)|Al<sup>3+</sup>(1.50 M)||Mn<sup>2+</sup>(0.002 M)|Mn(s). Under standard conditions, the cell potential is E<sup>o</sup><sub>cell</sub> = +0.48 V.

What is the approximate cell potential, E<sub>cell</sub>, at the concentrations shown in the reaction?

(A) -0.40 V (B) +0.56 V (C) +0.73 V (D) +0.40 V

48. Consider a concentration cell, containing 0.10 M H<sup>+</sup> in the reference compartment (the cathode) and a solution containing H<sup>+</sup> in the sample cell (the anode) with pH = 3.50. (**Note: n = 1 for this cell**)

In cell notation, this can be written as:  $H_2(g)|H^+(pH=3.50)||H^+(0.10 \text{ M})|H_2(g)$ . The cell potential,  $E_{cell}$  is approximately:

(A) +0.15 V (B) +0.34 V (C) -0.15 V (D) -0.21 V

49. The concentration of Arsenic [As, M=74.9] in drinking water was determined using a concentration cell with 0.40 M Arsenic(III) nitrate, As(NO<sub>3</sub>)<sub>3</sub>, in the reference cell (the cathode) and a sample of water with an unknown concentration of As<sup>3+</sup>(xx) in the sample cell (the anode). In cell notation, this can be written as: As(s)|As<sup>3+</sup>(xx)||As<sup>3+</sup>(0.40 M)|As(s).

The cell potential in the above concentration cell was measured to be +0.115 V. Therefore, the concentration of Arsenic in the sample, in **micrograms per Liter** ( $\mu$ g/L) is approximately:

- (A)  $280 \ \mu g/L$  (B)  $3,900 \ \mu g/L$  (C)  $45 \ \mu g/L$  (D)  $111 \ \mu g/L$
- 50. The solubility product of Li<sub>3</sub>PO<sub>4</sub> is K<sub>sp</sub> = 3.2x10<sup>-9</sup>. A Saturated solution of Li<sub>3</sub>PO<sub>4</sub> is placed in the sample compartment (anode) of an electrochemical cell and a standard 2.00 M solution of LiCl (a strong electrolyte) is placed in the reference compartment (cathode). In cell notation, this can be written as: Li(s) | Li<sup>+</sup>(Saturated Solution) || Li<sup>+</sup>(2.00 M) | Li(s).

What will be the cell voltage, Ecell ?

(A) -0.052 V (B) +0.136 V (C) +0.052 V (D) +0.068 V

# For #51 - #52: Use Table 2 (Some Reduction and Oxidation Potentials in Aqueous Solution) near the top of the test.

51. If aqueous Zn(II) Fluoride, ZnF<sub>2</sub>(aq) is placed in an electrolysis cell, and a voltage is applied, what will be the principal products of the electrolysis?

(A) Zn , F <sub>2</sub>	(B) F <sub>2</sub> , H <sub>2</sub> , OH <sup>-</sup>
(C) H <sub>2</sub> , OH <sup>-</sup> , O <sub>2</sub> , H <sup>+</sup>	(D) Zn , O <sub>2</sub> , H <sup>+</sup>

- 52. 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?
  - (A) AI,  $I_2$  (B)  $I_2$ ,  $H_2$ ,  $OH^-$
  - $(C) \ H_2 \ , \ OH^- \ , \ O_2 \ , \ H^+ \ (D) \ AI \ , \ O_2 \ , \ H \ (C)$

- 53. A current of 0.80 Amps (Coul/sec) is passed through a solution of Aluminum Nitrate, Al(NO<sub>3</sub>)<sub>3</sub>(aq), for 7. hours. Approximately what mass of solid Aluminum, Al(s) [M=27], will be deposited at the cathode?
  - (A) 1.9 g (B) 1.3 g (C) 16.9 g (D) 5.6 g
- 54. Approximately how much energy (in kJ) is needed to plate out 90. grams of Sn(s) [M=118.7] by electrolysis from a solution containing Sn(NO<sub>3</sub>)<sub>2</sub>(aq)? Assume that the voltage is 9. Volts.
  - (A) 660 kJ (B) 2,050 kJ (C) 330 kJ (D) 1,320 kJ
- 55. The type of nuclides with the highest number of stable isotopes have a(n) \_\_\_\_\_ number of protons and \_\_\_\_\_\_number of neutrons
  - (A) even , odd (B) odd , even (C) even , even (D) odd , odd
- 56. What nuclide will undergo positron emission to form Ar-38?
  - (A) Ca-42 (B) Cl-38 (C) Cl-40 (D) K-38
- 57. Which of the following is a likely decay path for Ne-19. Stable isotopes in this range typically have N/Z = 1.0.
  - (A)  ${}^{19}_{10}Ne \rightarrow {}^{0}_{-1}e + {}^{19}_{11}Na$ (B)  ${}^{19}_{10}Ne \rightarrow {}^{4}_{2}He + {}^{15}_{8}O$ (C)  ${}^{19}_{10}Ne \rightarrow {}^{0}_{+1}e + {}^{19}_{9}F$ (D)  ${}^{19}_{10}Ne \rightarrow {}^{1}_{1}H + {}^{18}_{9}F$
- Californium-252 (Cf-252) combines with an alpha particle to form a neutron and a new element. The new element is:
  - (A) Md-255 (B) Cm-249 (C) Md-256 (D) Fm-255

The last 2 questions are on the following page

- 59. Typical values of N/Z for stable nuclei rise with increasing atomic numbers because:
  - (A) More neutrons are required to form attractive interactions with the protons
  - (C) More long-range neutron-neutron attractions are required to counter the increasing short-range proton-proton repulsions
  - (B) Increased numbers of neutrons shield the short range proton-proton repulsions
  - (D) More short-range neutron-neutron attractions are required to counter the increasing long-range proton-proton repulsions.
- 60. Use the Molar Masses below to calculate the approximate Binding Energy per Nucleon (Eb/N) of Kr-92.
  - $m(_{1}H) = 1.008 \text{ g/mol}$ ,  $m(_{0}n) = 1.009 \text{ g/mol}$ ,  $m(_{36}^{92}Kr) = 91.926 \text{ g/mol}$
  - (A) 4.7x10<sup>8</sup> kJ/mol (B) 8.5x10<sup>8</sup> kJ/mol
  - (C) 8.5x10<sup>11</sup> kJ/mol (D) 7.8x10<sup>10</sup> kJ/mol