## Part 1: Multiple Choice

1. Calculate the solubility product constant for lead(II) iodide if 0.0024 mole of I ion is present in 2.0 L of a saturated lead(II) iodide solution
a. $1.4 \times 10^{-5}$
b. $8.6 \times 10^{-10}$
c. $5.2 \times 10^{-8}$
d. $3.5 \times 10^{-6}$
e. $4.6 \times 10^{-9}$
2. Calculate the number of moles of $\mathrm{Ag}^{+}$ion present in 2.0 L of a saturated solution of silver chromate. For silver chromate, $\mathrm{K}_{\mathrm{sp}}=1.1 \times 10^{-12}$.
a. $2.6 \times 10^{-4}$
b. $1.3 \times 10^{-4}$
c. $2.1 \times 10^{-4}$
d. $1.1 \times 10^{-4}$
e. $4.1 \times 10^{-4}$
3. Calculate the molar solubility of silver carbonate in 1.0 M sodium carbonate solution. For silver carbonate, $\mathrm{K}_{\mathrm{sp}}=8.1 \times 10^{-12}$.
a. $8.1 \times 10^{-12}$
b. $2.8 \times 10^{-6}$
c. $1.4 \times 10^{-6}$
d. $1.4 \times 10^{-8}$
e. $2.0 \times 10^{-4}$
4. Calculate the pH of a solution necessary to just begin the precipitation of magnesium hydroxide when the concentration of magnesium ion $=0.001 \mathrm{M}$. For magnesium hydroxide $K_{\text {sp }}=1.2 \times 10^{-11}$.
a. 11
b. 10
c. 9
d. 8
e. 4
5. The line notation, $\mathrm{Mg}(\mathrm{s})\left|\mathrm{Mg}^{2+}(\mathrm{aq})\right|\left|\mathrm{Fe}^{2+}(\mathrm{aq})\right| \mathrm{Fe}(\mathrm{s})$, indicates that:
a. iron metal is the reducing agent
b. magnesium metal is the cathode
c. $\mathrm{Fe}^{2+}$ ions are oxidized
d. magnesium metal is the reducing agent
e. $\mathrm{Mg}^{2+}$ ion is the reducing agent
6. Consider the following two electrode reactions and their standard electrode potentials:

$$
\begin{aligned}
& \mathrm{Al}^{+3}(\mathrm{aq})+3 \mathrm{e}^{-} \rightarrow \mathrm{Al}(\mathrm{~s}) \mathrm{E}^{\circ}=-1.66 \mathrm{~V} \\
& \mathrm{Cd}^{+2}(\mathrm{aq})+2 \mathrm{e}^{-} \rightarrow \mathrm{Cd}(\mathrm{~s}) \mathrm{E}^{\circ}=-0.40 \mathrm{~V}
\end{aligned}
$$

Write the cell reaction for a voltaic cell based on these two electrodes, and calculate the standard cell potential
a. $2 \mathrm{Al}^{+3}(\mathrm{aq})+3 \mathrm{Cd}^{+2}(\mathrm{aq}) \rightarrow 2 \mathrm{Al}(\mathrm{s})+3 \mathrm{Cd}(\mathrm{s}) \mathrm{E}^{\circ}$ cell $=2.10 \mathrm{~V}$
b. $2 \mathrm{Al}(\mathrm{s})+3 \mathrm{Cd}^{+2}(\mathrm{aq}) \rightarrow 2 \mathrm{Al}^{+3}(\mathrm{aq})+3 \mathrm{Cd}(\mathrm{s}) \mathrm{E}^{\circ}$ cell $=1.26 \mathrm{~V}$
c. $2 \mathrm{Al}(\mathrm{s})+3 \mathrm{Cd}^{+2}(\mathrm{aq}) \rightarrow 2 \mathrm{Al}^{+3}(\mathrm{aq})+3 \mathrm{Cd}(\mathrm{s}) \mathrm{E}^{\circ}$ cell $=3.78 \mathrm{~V}$
d. $2 \mathrm{Al}^{+3}(\mathrm{aq})+3 \mathrm{Cd}(\mathrm{s}) \rightarrow 2 \mathrm{Al}(\mathrm{s})+3 \mathrm{Cd}^{+2}(\mathrm{aq}) \mathrm{E}^{\circ}{ }_{\text {cell }}=1.26 \mathrm{~V}$
e. $2 \mathrm{Al}^{+3}(\mathrm{aq})+3 \mathrm{Cd}(\mathrm{s}) \rightarrow 2 \mathrm{Al}(\mathrm{s})+3 \mathrm{Cd}^{+2}(\mathrm{aq}) \mathrm{E}^{\circ}{ }_{\text {cell }}=2.10 \mathrm{~V}$
7. A voltaic cell consists of $\mathrm{Mn} / \mathrm{Mn}^{2+}$ and $\mathrm{Cd} / \mathrm{Cd}^{2+}$ half-cells with concentrations $\left[\mathrm{Mn}^{2+}\right]=0.75 \mathrm{M}$ and $\left[\mathrm{Cd}^{2+}\right]=0.15 \mathrm{M}$. Calculate the cell potential at $25^{\circ} \mathrm{C}$.

$$
\begin{aligned}
& \mathrm{Cd}^{+2}(\mathrm{aq})+2 \mathrm{e}^{-} \rightarrow \mathrm{Cd}(\mathrm{~s}) \mathrm{E}^{\circ}=-0.40 \mathrm{~V} \\
& \mathrm{Mn}^{+2}(\mathrm{aq})+2 \mathrm{e}^{-} \rightarrow \mathrm{Mn}(\mathrm{~s}) \mathrm{E}^{\circ}=-1.18 \mathrm{~V}
\end{aligned}
$$

a. 1.60 V
b. 1.56 V
c. 1.54 V
d. 0.80 V
e. 0.76 V
8. The standard reference electrode that is used to measure all other standard electrode potentials is called the "standard $\qquad$ electrode."
a. sulfur
b. oxygen
c. hydrogen
d. iron
e. platinum
9. A constant current was passed through a solution of $\mathrm{KAuCl}_{4}$ between gold electrodes. Over a period of 20.00 min, the cathode increased in mass by 2.664 g. What was the current in amperes?
( $\mathrm{F}=96500 \mathrm{C} / \mathrm{mol}$ ) Cathode half-reaction: $\mathrm{AuCl}_{4}^{-}(\mathrm{aq})+3 \mathrm{e}^{-} \rightarrow \mathrm{Au}(\mathrm{s})+4 \mathrm{Cl}^{-}(\mathrm{aq})$
a. 1.08 A
b. 3.26 A
c. 2.17 A
d. 6.52 A
e. 3.48 A
10. Calculate the equilibrium constant $\mathrm{K}_{\mathrm{c}}$ for this reaction at $25^{\circ} \mathrm{C}$ :
$2 \mathrm{Cr}(\mathrm{s})+3 \mathrm{~Pb}^{2+}(\mathrm{aq}) \rightarrow 3 \mathrm{~Pb}(\mathrm{~s})+2 \mathrm{Cr}^{3+}(\mathrm{aq}) \mathrm{E}^{\circ}$ cell $=0.61 \mathrm{~V}$
a. $6.7 \times 10^{61}$
b. $8.1 \times 10^{30}$
c. $9.2 \times 10^{45}$
d. $3.2 \times 10^{51}$
e. $4.6 \times 10^{22}$
11. Consider the following half-reactions and select the strongest oxidizing agent present:
$\mathrm{Sr}^{2+}(\mathrm{aq})+2 \mathrm{e}^{-} \rightarrow \mathrm{Sr}(\mathrm{s}) \mathrm{E}^{\circ}=-2.89 \mathrm{~V}$
$\mathrm{Cr}^{2+}(\mathrm{aq})+2 \mathrm{e}^{-} \rightarrow \mathrm{Cr}(\mathrm{s}) \mathrm{E}^{\circ}=-0.913 \mathrm{~V}$
$\mathrm{Co}^{2+}(\mathrm{aq})+2 \mathrm{e}^{-} \rightarrow \mathrm{Co}(\mathrm{s}) \mathrm{E}^{\circ}=-0.28 \mathrm{~V}$
a. $\mathrm{Cr}^{2+}(\mathrm{aq})$
b. $\mathrm{Sr}^{2+}(\mathrm{aq})$
c. $\mathrm{Co}^{2+}(\mathrm{aq})$
d. $\mathrm{Sr}(\mathrm{s})$
e. $\mathrm{Co}(\mathrm{s})$
12. The standard free energy change for the following reaction is -210 kJ . What is the cell potential?

$$
2 \mathrm{H}_{2} \mathrm{O}_{2}(\mathrm{aq}) \leftrightarrow 2 \mathrm{H}_{2} \mathrm{O}(\mathrm{I})+\mathrm{O}_{2}(\mathrm{~g})
$$

a. +0.640 V
b. +1.09 V
c. +0.420 V
d. +0.547 V
e. +0.752 V
13. In the electrolysis of aqueous sodium sulfate, which one of the following species is oxidized?
$\mathrm{S}_{2} \mathrm{O}_{8}{ }^{2-}(\mathrm{aq})+2 \mathrm{e}^{-} \rightarrow 2 \mathrm{SO}_{4}{ }^{2-}(\mathrm{aq}) \quad \mathrm{E}^{\circ}=2.10 \mathrm{~V}$
$\mathrm{Na}^{+}(\mathrm{aq})+\mathrm{e}^{-} \rightarrow \mathrm{Na}(\mathrm{s}) \quad \mathrm{E}^{\circ}=-2.71 \mathrm{~V}$
$2 \mathrm{H}_{2} \mathrm{O}(\mathrm{I})+2 \mathrm{e}^{-} \rightarrow \mathrm{H}_{2}(\mathrm{~g})+2 \mathrm{OH}^{-}(\mathrm{aq}) \quad \mathrm{E}^{\circ}=-0.38 \mathrm{~V}$
$\mathrm{O}_{2}(\mathrm{~g})+4 \mathrm{H}^{+}(\mathrm{aq})+4 \mathrm{e}^{-} \rightarrow \mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \quad \mathrm{E}^{\circ}=1.23 \mathrm{~V}$
a. sodium ion
b. oxygen
c. sulfate ion
d. water
e. hydronium ion
14. Calculate $\Delta S^{\circ}$ for the following reaction:

Standard molar entropies, $\mathrm{S}^{\circ}$ (J/mol $\cdot \mathrm{K}$ ): $\mathrm{Cr}(\mathrm{s}), 23.8 ; \mathrm{O}_{2}(\mathrm{~g}), 205.1 ; \mathrm{Cr}_{2} \mathrm{O}_{3}(\mathrm{~s}), 81.2$

$$
4 \mathrm{Cr}(\mathrm{~s})+3 \mathrm{O}_{2}(\mathrm{~g}) \rightarrow 2 \mathrm{Cr}_{2} \mathrm{O}_{3}(\mathrm{~s})
$$

a. $548.1 \mathrm{~J} / \mathrm{K}$
b. $147.7 \mathrm{~J} / \mathrm{K}$
c. $-147.7 \mathrm{~J} / \mathrm{K}$
d. $-548.1 \mathrm{~J} / \mathrm{K}$
e. $-66.5 \mathrm{~J} / \mathrm{K}$
15. When crystalline solid barium hydroxide octahydrate and crystalline solid ammonium nitrate are mixed in a beaker at room temperature, a spontaneous reaction occurs. The temperature of the beaker contents rapidly falls to below $0^{\circ} \mathrm{C}$. Use this information to decide whether the reaction is exothermic or endothermic and what the signs of $\Delta \mathrm{H}$ and $\Delta \mathrm{S}$ are.
a. endothermic; $\Delta \mathrm{H}>0 ; \Delta \mathrm{S}>0$
b. exothermic; $\Delta \mathrm{H}<0 ; \Delta \mathrm{S}>0$
c. endothermic; $\Delta \mathrm{H}<0 ; \Delta \mathrm{S}<0$
d. endothermic; $\Delta \mathrm{H}<0 ; \Delta \mathrm{S}>0$
e. exothermic; $\Delta \mathrm{H}>0 ; \Delta \mathrm{S}<0$
16. A certain reaction has $\Delta H^{\circ}=-14.2 \mathrm{~kJ}$ and $\Delta \mathrm{S}^{\circ}=+87.9 \mathrm{~J} / \mathrm{K}$. What is the value of $\Delta \mathrm{G}^{\circ}$ for this reaction? (Temperature is $25^{\circ} \mathrm{C}$ )
a. +40.4 kJ
b. -16.4 kJ
c. -26200 kJ
d. -40.4 kJ
e. -7820 kJ
17. A reaction has an equilibrium constant $\mathrm{K}_{\mathrm{c}}=7.0$ at $35^{\circ} \mathrm{C}$. Calculate the value of $\Delta G^{\circ}$ for the reaction
a. -4.98 kJ
b. -2.46 kJ
c. -5.66 kJ
d. -2.16 kJ
e. none of the above
18. The compound 1-pentanol has an enthalpy of vaporization of $55.5 \mathrm{~kJ} / \mathrm{mol}$ and an entropy of vaporization of $148 \mathrm{~J} / \mathrm{K} \cdot \mathrm{mol}$. Calculate its approximate boiling point.
a. $45^{\circ} \mathrm{C}$
b. $102^{\circ} \mathrm{C}$
C. $93^{\circ} \mathrm{C}$
d. $210^{\circ} \mathrm{C}$
e. $375^{\circ} \mathrm{C}$
19. Which of the following statements about entropy and enthalpy of a system is correct?
a. The absolute entropy of pure oxygen at $25^{\circ} \mathrm{C}$ and 1 atm is zero.
b. When ice melts, $\Delta \mathrm{S}$ is positive and $\Delta \mathrm{H}$ is negative.
c. When a candle burns, $\Delta \mathrm{S}$ is positive and $\Delta \mathrm{H}$ is negative.
d. The entropy of a system must increase for the reaction to be spontaneous.
e. None of the above statements are correct.
20. Sodium carbonate can be made by heating sodium bicarbonate carbonate:

$$
2 \mathrm{NaHCO}_{3}(\mathrm{~s}) \rightarrow \mathrm{Na}_{2} \mathrm{CO}_{3}(\mathrm{~s})+\mathrm{CO}_{2}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{~g})
$$

For this reaction, $\Delta \mathrm{H}^{\circ}=128.9 \mathrm{~kJ}$ and $\Delta \mathrm{S}^{\circ}=321 \mathrm{~J} / \mathrm{K}$. At approximately what temperature will $\mathrm{K}=1$ ?
a. $401.6^{\circ} \mathrm{C}$
b. 401.6 K
c. $33.1^{\circ} \mathrm{C}$
d. 33.1 K
e. None of the above

Part 2: Free Response. Please show all work.

1. Will a precipitate form when 125 ml of 0.0250 M aluminum nitrate and 25.0 ml of 0.000100 M calcium hydroxide are mixed together? Why? $\mathrm{K}_{\mathrm{sp}}$ of aluminum hydroxide $=3.7 \times 10^{-15}$
2. In the electrolysis of a molten mixture of potassium iodide and magnesium fluoride, identify which product forms at the positive electrode, and what product forms at the negative electrode.
$\mathrm{K}^{+}(\mathrm{aq})+\mathrm{e}^{-} \rightarrow \mathrm{K}(\mathrm{s}) \quad \mathrm{E}^{\circ}=-2.93 \mathrm{~V}$
$\mathrm{I}_{2}(\mathrm{~s})+2 \mathrm{e}^{-} \rightarrow 2 \mathrm{I}^{-} \quad \mathrm{E}^{\circ}=-0.53 \mathrm{~V}$
$\mathrm{Mg}^{2+}(\mathrm{aq})+2 \mathrm{e}^{-} \rightarrow \mathrm{Mg}(\mathrm{s}) \quad \mathrm{E}^{\circ}=-2.37 \mathrm{~V}$
$\mathrm{F}_{2}(\mathrm{~g})+2 \mathrm{e}^{-} \rightarrow 2 \mathrm{~F}^{-} \quad \mathrm{E}^{\circ}=2.87 \mathrm{~V}$
3. In a $\mathrm{Cd}^{2+} / \mathrm{Cd}(\mathrm{s})$ and $\mathrm{Au}^{3+} / \mathrm{Au}(\mathrm{s})$ voltaic cell the electronic voltmeter measures to be 1.92 -Volts. What concentration of cadmium (II) ion must be present in the cell if the gold (III) ion concentration is known to be 0.10 M .

$$
\begin{array}{ll}
\mathrm{Au}^{3+}(\mathrm{aq})+3 \mathrm{e}^{-} \rightarrow \mathrm{Au}(\mathrm{~s}) & \mathrm{E}^{\circ}=1.50 \\
\mathrm{Cd}^{2+}(\mathrm{aq})+2 \mathrm{e}^{-} \rightarrow \mathrm{Cd}(\mathrm{~s}) & \mathrm{E}^{\circ}=-0.40
\end{array}
$$

