Chem. 1B Midterm 2
Version B
March 3, 2017

Name:
Print Neatly. You will lose 1 point if I cannot read your name or perm number.

Perm Number: $\qquad$

All work must be shown on the exam for partial credit. Points will be taken off for incorrect or missing units. Calculators are allowed. Cell phones may not be used as calculators. On fundamental and challenge problems you must show your work in order to receive credit for the problem. If your cell phone goes off during the exam, you will have your exam removed from you.

| Fundamentals <br> (of 36 possible) |  |
| :---: | :--- |
| Problem 1 <br> (of 20 possible) |  |
| Problem 2 <br> (of 14 possible) |  |
| Multiple Choice <br> (of 30 possible) |  |
| Midterm Total <br> (of 100 possible) |  |

## Fundamental Questions

Each of these fundamental chemistry questions is worth 6 points. You must show work to get credit. Little to no partial credit will be awarded. Make sure to include the correct units on your answers.

1) 6 pts Use the information on the constant sheet to sort the following chemical species by oxidizing power. 1 will be the species that has the largest oxidizing power and 4 will be the species with the lowest oxidizing power.

| Species | Oxidizing Power |
| :--- | :--- |
| $\mathrm{Zn}^{2+}(\mathrm{aq})$ | 3 |
| $\mathrm{Ag}^{+}(\mathrm{aq})$ | 1 (highest) |
| $\mathrm{Ni}^{2+}(\mathrm{aq})$ | 2 |
| $\mathrm{Al}^{3+}(\mathrm{aq})$ | 4 (lowest) |

2) 6 pts

The rate constant k for a certain reaction is measured at two different temperatures:

| temperature | k |
| :---: | :---: |
| $104.0^{\circ} \mathrm{C}$ | $4.8 \times 10^{3}$ |
| $-36.0^{\circ} \mathrm{C}$ | 0.023 |

Assuming the rate constant obeys the Arrhenius equation, calculate the activation energy $\mathrm{E}_{\mathrm{a}}$ for this reaction.
Round your answer to 2 significant digits.

$$
\begin{aligned}
& \ln \left(\frac{k_{1}}{k_{2}}\right)=\frac{E_{a}}{R}\left(\frac{1}{T_{2}}-\frac{1}{T_{1}}\right) \\
& \ln \left(\frac{4.8 \times 10^{3}}{0.023}\right)=\frac{E_{a}}{8.3145 \frac{\mathrm{~J}}{\mathrm{~mol} \cdot \mathrm{~K}}}\left(\frac{1}{237 \mathrm{~K}}-\frac{1}{377 \mathrm{~K}}\right) \\
& \mathrm{E}_{\mathrm{a}}=65,000 \frac{\mathrm{~J}}{\mathrm{~mol}}=65 \frac{\mathrm{~kJ}}{\mathrm{~mol}}
\end{aligned}
$$

3) 6 pts Determine the rate law for $2 \mathrm{~A}+\mathrm{B} \rightarrow 2 \mathrm{C}$ (you do not need to find the numerical value of k ):

| Experiment | $[\mathrm{A}]_{\mathrm{o}}(\mathrm{M})$ | $[\mathrm{B}]_{\mathrm{o}}(\mathrm{M})$ | Initial Rate $\left(\frac{\mathrm{mol}}{\mathrm{L} \cdot \mathrm{s}}\right)$ |
| :---: | :---: | :---: | :---: |
| 1 | 0.050 | 0.10 | 0.074 |
| 2 | 0.10 | 0.20 | 0.888 |
| 3 | 0.050 | 0.20 | 0.222 |

General rate law: Rate $=k[A]^{x}[B]^{y}$
Determine Reaction order for $A$ (Experiment 2\&3)
Double [A] Quadruple rate $\quad$ Rate $=k[A]^{2}[B]^{y}$
Determine reaction order for B (Experiment 1\&3)

$$
\begin{aligned}
& \frac{0.074 \frac{\text { mol }}{L . s}=k(0.050 M)^{2}(0.10 M)^{y}}{0.222 \frac{m o l}{L \cdot s}=k(0.050 M)^{2}(0.20 M)^{y}} \\
& 0.333=\frac{(0.10 M)^{y}}{(0.20 M)^{y}}=(0.50)^{y} \\
& \log (0.333)=y \log (0.50) \\
& y=1.6 \\
& \text { Rate }=k[A]^{2}[B]^{1.6}
\end{aligned}
$$

4) 6 pts $\quad$ What is the $\mathrm{E}_{\text {cell }}$ for the following at $25^{\circ} \mathrm{C}$ ?

$$
\mathrm{Al}(\mathrm{~s})\left|\mathrm{Al}^{3+}(0.18 \mathrm{M})\right|\left|\mathrm{Fe}^{2+}(0.85 \mathrm{M})\right| \mathrm{Fe}(\mathrm{~s})
$$

$$
\begin{array}{ll}
\text { The reaction occurring in cell } & \mathrm{E}^{\circ}=1.66 \mathrm{~V} \\
\qquad \begin{array}{cl}
\text { Anode: } 2\left(\mathrm{Al}(\mathrm{~s}) \rightarrow \mathrm{Al}^{3+}(\mathrm{aq})+3 \mathrm{e}^{-}\right) & \mathrm{E}^{\circ}=-0.44 \mathrm{~V} \\
\text { Cathode: } 3\left(\mathrm{Fe}^{2+}(\mathrm{aq})+2 \mathrm{e}^{-} \rightarrow \mathrm{Fe}(\mathrm{~s})\right) & 2 \mathrm{Al}(\mathrm{~s})+3 \mathrm{Fe}^{2+}(\mathrm{aq}) \rightarrow 2 \mathrm{Al}^{3+}(\mathrm{aq})+3 \mathrm{Fe}(\mathrm{~s}) \\
\mathrm{E}^{\circ}=1.22 \mathrm{~V} \\
E=E^{\circ}-\frac{R T}{n F} \ln (Q)=E^{\circ}-\frac{R T}{n F} \ln \left(\frac{\left[\mathrm{Al}^{3+}\right]^{2}}{\left[F e^{2+}\right]^{3}}\right) & \\
E=1.22 \mathrm{~V}-\frac{\left(8.3145 \frac{J}{m o l \cdot K}\right)(298 K)}{(6)\left(96,485 \frac{C}{m o l}\right)} \ln \left(\frac{0.18^{2}}{0.85^{3}}\right)=1.23 \mathrm{~V} &
\end{array}
\end{array}
$$

5) 6 pts Write the overall reaction and rate law that corresponds to the following reaction mechanism. Be sure to eliminate intermediates from the answer.

| $\stackrel{k_{1}}{\rightleftharpoons} \mathrm{~B}+\mathrm{C}$ | fast equilibrium |
| :---: | :--- |
| $k_{-1}$ |  |
| $\mathrm{C}+\mathrm{D} \underset{2}{\rightleftharpoons} \mathrm{E}$ | fast equilibrium |
| $k_{-2}$ |  |
| $\mathrm{E} \xrightarrow{k_{3}} \mathrm{~F}$ | slow |

Overall Reaction: $\mathrm{A}+\mathrm{D} \rightarrow \mathrm{B}+\mathrm{F}$
Rate $=k_{3}[E]$
E is an intermediate use equation 2 to eliminate E

$$
\begin{aligned}
& \quad k_{2}[C][D]=k_{-2}[E] \\
& \text { Rate }=\frac{\left.k_{2} k_{3}\right]}{k_{-2}}[C][D]
\end{aligned}
$$

C is an intermediate use equation 1 to eliminate C

$$
\begin{aligned}
& k_{1}[A]=k_{-1}[B][C] \\
\text { Rate }= & =\frac{k_{1} k_{2} k_{3}}{k_{-1} k_{-2}} \frac{[A][D]}{[B]}
\end{aligned}
$$

6) $6 p t s$

Draw the following galvanic cell:

$$
\mathrm{Pb}(\mathrm{~s})+\mathrm{Cl}_{2}(\mathrm{~g}) \rightarrow \mathrm{Pb}^{2+}(\mathrm{aq})+2 \mathrm{Cl}^{-}(\mathrm{aq})
$$

Calculate $\mathrm{E}^{\circ}$, show the direction of electron flow and the direction of ion migration through the salt bridge, and identify the cathode and anode.


## Challenge Problems

Each of the following short answer questions are worth the noted points. Partial credit will be given. You must show your work to get credit. Make sure to include proper units on your answer.

1) The decomposition of hydrogen peroxide was studied at a particular temperature. The following data were obtained, where

| rate $=-\frac{d\left[\mathrm{H}_{2} \mathrm{O}_{2}\right]}{d t}$ |  |
| :---: | :---: |
| Time $(\mathrm{s})$ | $\left[\mathrm{H}_{2} \mathrm{O}_{2}\right](\mathrm{mol} / \mathrm{L})$ |
| 0 | 1.00 |
| $120 \pm 1$ | 0.91 |
| $300 \pm 1$ | 0.78 |
| $600 \pm 1$ | 0.59 |
| $1200 \pm 1$ | 0.37 |
| $1800 \pm 1$ | 0.22 |
| $2400 \pm 1$ | 0.13 |
| $3000 \pm 1$ | 0.082 |
| $3600 \pm 1$ | 0.050 |

1a) $8 p t s$
Determine the integrated rate law
Find time points that are evenly spaced ( $0,600,1200,1800$ )
Determine what give a straight $\left[\mathrm{H}_{2} \mathrm{O}_{2}\right]$ vs. $\mathrm{t}, \ln \left[\mathrm{H}_{2} \mathrm{O}_{2}\right]$ vs. t , or $\left[\mathrm{H}_{2} \mathrm{O}_{2}\right]^{-1}$ vs. t

| Time (s) | $\left[\mathrm{H}_{2} \mathrm{O}_{2}\right]$ | $\ln \left[\mathrm{H}_{2} \mathrm{O}_{2}\right]$ | $\frac{1}{\left[\mathrm{H}_{2} \mathrm{O}_{2}\right]}$ |
| :---: | :---: | :---: | :---: |
| 0 | $1.00(0.41)$ | $0.00(0.53)$ | $1.00(-0.69)$ |
| 600 | $0.59(0.22)$ | $-0.53(0.46)$ | $1.69(-1.01)$ |
| 1200 | $0.37(0.15)$ | $-0.99(0.52)$ | $2.70(-1.85)$ |
| 1800 | 0.22 | -1.51 | 4.55 |

Therefore, the only number that are evenly spaced are the $\ln \left[\mathrm{H}_{2} \mathrm{O}_{2}\right]$ so the reaction is $1^{\text {st }}$ order and the integrated rate law is: $\ln \left[\mathrm{H}_{2} \mathrm{O}_{2}\right]=-k t+\ln \left[\mathrm{H}_{2} \mathrm{O}_{2}\right]$ 。

1b) $2 p t s$
Determine the differential rate law.
Rate $=k\left[H_{2} O_{2}\right]$
1c) 4 pts Determine the value of the rate constant.
Since the reaction was first order with respect to $\left[\mathrm{O}_{3}\right]$ the slope of the line equals -k

$$
\mathrm{k}=- \text { slope }=-\frac{\text { rise }}{\text { run }}=-\frac{\ln \left(0.22 \frac{\mathrm{~mol}}{\mathrm{~L}}\right)-\ln \left(1.00 \frac{\mathrm{~mol}}{\mathrm{~cm}^{3}}\right)}{1,800 \cdot s-0 \mathrm{~s}}=8.4 \times 10^{-4} \frac{1}{\mathrm{~s}}
$$

1d) 6 pts Calculate the $\left[\mathrm{H}_{2} \mathrm{O}_{2}\right]$ at 4000 . s after the start of the reaction.
$\ln \left[\mathrm{H}_{2} \mathrm{O}_{2}\right]=-k t+\ln \left[\mathrm{H}_{2} \mathrm{O}_{2}\right]_{\circ}=-\left(8.4 \times 10^{-4} \frac{1}{s}\right)(4000 . \mathrm{s})+\ln (1.00 \mathrm{M})$
$\left[\mathrm{H}_{2} \mathrm{O}_{2}\right]=0.035 \mathrm{M}$

2a) 8 pts Suppose that 2.69 g of a silver salt $(\mathrm{AgX})$ is dissolved in 550 . mL of water. With a current of $3.50 \mathrm{~A}, 395.0 \mathrm{~s}$ was needed to plate out all the silver. What is the mass percentage of silver in the salt? (Hint: Silver forms the following ion: $\mathrm{Ag}^{+}$)

Determine the equation of interest

$$
\mathrm{Ag}^{+}+\mathrm{e}^{-} \rightarrow \mathrm{Ag}
$$

Determine the moles of $\mathrm{e}^{-}$and moles of Ag
$n=\frac{I t}{F}=\frac{(3.50 \mathrm{~A})(395.0 \mathrm{~s})}{\left(96,485 \frac{C}{\mathrm{~mol}}\right)}=0.0143 \mathrm{~mol} \mathrm{e}^{-}$
$0.0143 \mathrm{~mol} e^{-\left(\frac{1 \mathrm{~mol} \mathrm{Ag}}{1 \mathrm{~mol}^{-}}\right)}=0.0143 \mathrm{~mol} \mathrm{Ag}$
Determine mass of Ag

$$
0.0143 \mathrm{~mol} \mathrm{Ag}\left(\frac{107.87 \mathrm{~g}}{1 \mathrm{~mol} \mathrm{Ag}}\right)=1.54 \mathrm{~g}
$$

Determine mass Percent
mass $\% A g=\frac{m_{A g}}{m_{t o t}}(100 \%)=\frac{1.54 g}{2.69 g}(100 \%)=57.2 \%$
2b) 6 pts What is the formula of the salt?

Determine the identity of $X$
For every one mol of Ag there is one mol of $X$
Determine the molar mass of $X$

$$
M=\frac{m_{x}}{n_{x}}=\frac{2.69 \mathrm{~g}-1.54 \mathrm{~g}}{0.0143 \mathrm{~mol}}=80.4 \frac{\mathrm{~g}}{\mathrm{~mol}}
$$

Therefore, X is bromine and the formula is AgBr

## Multiple Choice Questions

On the ParScore form you need to fill in your answers, perm number, test version, and name. Failure to do any of these things will result in the loss of 1 point. Your perm number is placed and bubbled in under the "ID number." Do not skip boxes or put in a hyphen; unused boxes should be left blank. Bubble in your test version (B) under the "test form." Note: Your ParScore form will not be returned to you, therefore, for your records, you may want to mark your answers on this sheet. Each multiple choice question is worth 5 points.

1. When the equation $\mathrm{Cl}_{2} \rightarrow \mathrm{Cl}^{-}+\mathrm{ClO}_{3}^{-}$(basic solution) is balanced using the smallest wholenumber coefficients, what is the coefficient of $\mathrm{OH}^{-}$?
a. 1
b. 2
c. 3
d. 4
e. None of the above
2. For a reaction in a voltaic cell, both $\Delta H^{\circ}$ and $\Delta S^{\circ}$ are positive. Which of the following statements is true?
a. $E^{\circ}{ }_{c e l l}$ will decrease with an increase in temperature.
b. $E^{\circ}$ cell will not change when the temperature increases.
c. $E^{\circ}$ cell will increase with an increase in temperature.
d. $\Delta G^{\circ}>0$ for all temperatures.
e. None of the above statements is true.

An antique automobile bumper is to be chrome plated. The bumper, which is dipped into an acidic $\mathrm{Cr}_{2} \mathrm{O}_{7}{ }^{2-}$ solution, serves as a cathode of an electrolytic cell. The atomic mass of Cr is $51.996 ; 1$ faraday $=96,485$ coulombs.
3. If the current is 34.4 amperes, how long will it take to deposit $1.58 \times 10^{2} \mathrm{~g}$ of $\mathrm{Cr}(s)$ onto the bumper?
a. 9.5 h
b. 14.2 h
c. 23.4 h
d. 28.4 h
e. None of the above
4. What reaction occurs at the anode and at the cathode when electrolysis of aqueous $\mathrm{Mgl}_{2}$ occurs? Assume standard conditions.
Helpful information: $I_{2}+2 e^{-} \rightarrow 21-\quad E^{\circ}=0.54 \mathrm{~V}$

$$
\mathrm{Mg}^{2+}+2 e^{-} \rightarrow \mathrm{Mg} \quad E^{\circ}=-2.37 \mathrm{~V}
$$

Anode/Cathode
a. Magnesium/Water
b. Iodine/Magnesium
c. Water/Water
d. Water/Magnesium
e. None of the above
5. The rate law for the following reaction

$$
2 \mathrm{NO}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightarrow 2 \mathrm{NO}_{2}(\mathrm{~g})
$$

was experimentally found to be in the form
rate $=\mathrm{k}[\mathrm{NO}]^{\times}\left[\mathrm{O}_{2}\right]^{\mathrm{y}}$
It was also found that when the NO concentration was doubled, the rate of the reaction increases by a factor of 4 . In addition, when both the $\mathrm{O}_{2}$ and the NO concentration were doubled, the rate increases by a factor of 8 . What is the reaction order of $\mathrm{O}_{2}$ ?
a. $2^{\text {nd }}$
b. $1^{\text {st }}$
c. $3^{\text {rd }}$
d. $0^{\text {th }}$
e. None of the above
6. Calculate the solubility product of silver iodide at $25^{\circ} \mathrm{C}$, given the following data:
$E^{\circ}(\mathrm{V})$
$\mathrm{Ag}(\mathrm{s})+\mathrm{e}^{-} \rightarrow \mathrm{Ag}(\mathrm{s})+\mathrm{I}^{-}$
-0.15
$\mathrm{I}_{2}(\mathrm{~s})+2 \mathrm{e}^{-} \rightarrow 21^{-}$
+0.54
$\mathrm{Ag}^{+}+\mathrm{e}^{-} \rightarrow \mathrm{Ag}(s) \quad+0.80$
a. $2.9 \times 10^{-3}$
b. $8.4 \times 10^{-17}$
c. $3.5 \times 10^{-20}$
d. $1.9 \times 10^{-4}$
e. None of the above

E, C, B, E, B, B

