## Name

$\qquad$

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

Please turn in:

1. Your Scantron with your name written in + bubbled answers. You don't have to bubble in your name.
2. This signature sheet. Please put your name on top. You are welcome to supply a 4 digit number of your choice if you would like your course results posted anonymously on the course web site.

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

Conversions: 1 atm. = 760 torr

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

$$
\begin{aligned}
& \mathrm{R}=8.31 \mathrm{~J} / \mathrm{mol} \cdot \mathrm{~K} \\
& \mathrm{R}=8.31 \times 10^{-3} \mathrm{~kJ} / \mathrm{mol} \cdot \mathrm{~K} \\
& \mathrm{~N}_{\mathrm{A}}=6.02 \times 10^{23} \mathrm{~mol}^{-1} \\
& \mathrm{~F}=96,500 \mathrm{Coul} / \mathrm{mol} \mathrm{e}^{-} \\
& \mathrm{C}=3.00 \times 10^{8} \mathrm{~m} / \mathrm{s} \text { (speed of light) }
\end{aligned}
$$

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



## ELECTROCHEMISTRY INFORMATION

Table 1: Standard Reduction Potentials
Reduction Half-Reactions $\mathrm{E}^{\circ}(\mathrm{V})$

| $\mathrm{F}_{2}+2 \mathrm{e}^{-} \rightarrow 2 \mathrm{~F}^{-}$ | +2.87 |
| :--- | ---: |
| $\mathrm{Au}^{3+}+3 \mathrm{e}^{-} \rightarrow \mathrm{Au}$ | +1.50 |
| $\mathrm{Cl}_{2}+2 \mathrm{e}^{-} \rightarrow 2 \mathrm{Cl}^{-}$ | +1.36 |
| $\mathrm{Br}_{2}+2 \mathrm{e}^{-} \rightarrow 2 \mathrm{Br}^{-}$ | +1.07 |
| $\mathrm{Hg}^{2+}+2 \mathrm{e}^{-} \rightarrow \mathrm{Hg}$ | +0.86 |
| $\mathrm{Ag}^{+}+1 \mathrm{e}^{-} \rightarrow \mathrm{Ag}$ | +0.80 |
| $\mathrm{I}_{2}+2 \mathrm{e}^{-} \rightarrow 2 \mathrm{I}^{-}$ | +0.54 |
| $\mathrm{Cu}^{2+}+2 \mathrm{e}^{-} \rightarrow \mathrm{Cu}$ | +0.34 |
| $\mathrm{Sn}^{2+}+2 \mathrm{e}^{-} \rightarrow \mathrm{Sn}$ | -0.14 |
| $\mathrm{Fe}^{3+}+3 \mathrm{e}^{-} \square \mathrm{Fe}$ | -0.04 |
| $\mathrm{Ni}^{2+}+2 \mathrm{e}^{-} \rightarrow \mathrm{Ni}$ | -0.25 |
| $\mathrm{Fe}^{2+}+2 \mathrm{e}^{-} \rightarrow \mathrm{Fe}$ | -0.44 |
| $\mathrm{Zn}^{2+}+2 \mathrm{e}^{-} \rightarrow \mathrm{Zn}$ | -0.76 |
| $\mathrm{Mn}^{2+}+2 \mathrm{e}^{--} \rightarrow \mathrm{Mn}$ | -1.18 |
| $\mathrm{Al}^{3+}+3 \mathrm{e}^{-} \rightarrow \mathrm{Al}$ | -1.66 |
| $\mathrm{Mg}^{2+}+2 \mathrm{e}^{-} \rightarrow \mathrm{Mg}$ | -2.37 |
| $\mathrm{~K}^{+}+1 \mathrm{e}^{-} \rightarrow \mathrm{K}$ | -2.93 |
| $\mathrm{Ki}^{+}+1 \mathrm{e}^{-} \rightarrow \mathrm{Li}$ | -3.05 |

Table 2: Some Reduction and Oxidation Potentials in Aqueous Solution

## Reduction Potentials

$2 \mathrm{H}_{2} \mathrm{O}+2 \mathrm{e}^{-} \rightarrow \mathrm{H}_{2}+2 \mathrm{OH}^{-}$

$$
\mathrm{Al}^{3+}+3 \mathrm{e}^{-} \rightarrow \mathrm{Al}
$$

$$
\mathrm{Zn}^{2+}+2 \mathrm{e}^{-} \rightarrow \mathrm{Zn}
$$

$$
\mathrm{Mg}^{2+}+2 \mathrm{e}^{-\quad} \quad \mathrm{Mg}
$$

$$
\mathrm{Fe}^{2+}+2 \mathrm{e}^{-}[\mathrm{Fe}
$$

$$
\begin{aligned}
& \mathrm{E}^{\circ}{ }_{\mathrm{red}}=-0.83 \mathrm{~V} \\
& \mathrm{E}^{\circ}{ }_{\mathrm{red}}=-1.66 \mathrm{~V} \\
& \mathrm{E}^{\circ}{ }_{\mathrm{red}}=-0.76 \mathrm{~V} \\
& \mathrm{E}^{\circ}{ }_{\mathrm{red}}=-2.37 \mathrm{~V} \\
& \mathrm{E}^{\mathrm{o}} \mathrm{red}=-2.71 \mathrm{~V} \\
& \mathrm{E}^{\circ}{ }_{\text {red }}=-0.44 \mathrm{~V}
\end{aligned}
$$

## Oxidation Potentials

| $2 \mathrm{H}_{2} \mathrm{O} \rightarrow \mathrm{O}_{2}+4 \mathrm{H}^{+}+4 \mathrm{e}^{-}$ | $\mathrm{E}^{\mathrm{o}}{ }_{\text {oxid }}=-1.23 \mathrm{~V}$ |
| :--- | :--- |
| $2 \mathrm{I}^{-} \rightarrow \mathrm{I}_{2}+2 \mathrm{e}^{-}$ | $\mathrm{E}^{\mathrm{o}}{ }_{\text {oxid }}=-0.54 \mathrm{~V}$ |
| $2 \mathrm{Br}^{-} \square \mathrm{Br}_{2}+2 \mathrm{e}^{-}$ | $\mathrm{E}^{\mathrm{o}}{ }_{\text {oxid }}=-1.07 \mathrm{~V}$ |
| $2 \mathrm{~F}^{-} \rightarrow \mathrm{F}_{2}+2 \mathrm{e}^{-}$ | $\mathrm{E}^{\mathrm{o}}{ }_{\text {oxid }}=-2.87 \mathrm{~V}$ |

## Some Electrochemical Equations

$F=96,500 \mathrm{C} / \mathrm{mol} \mathrm{e}^{-}$(Coulombs per mole of electrons)
$1 \mathrm{~J}=1 \mathrm{CxV}$ [i.e. 1 Joule $=1$ Coulomb $x$ Volt]
$\Delta G^{0}=-n F E^{\circ}$
$E=E^{\circ}-\frac{0.0592}{n} \cdot \log (Q)$
$\mathrm{Q}=\mathrm{ixt} \quad$ i.e. Charge (in Coul) $=$ Current (in Amps $=$ Coul/sec) $x$ time (in sec)]
$\mathrm{E}=\mathrm{Q} \times \mathrm{V} \quad$ i.e. Energy (in J) = Charge (in Coulombs) $\times$ Voltage (in Volts)
(note: 1 Coulomb-Volt = 1 Joule)
(60) 60 QUESTIONS (Mark the one correct answer to each question on your scantron)

## 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)

1. For the reaction, $\mathrm{A}+\mathrm{B} \rightarrow$ Products, the rate law is: Rate $=k \frac{[C]^{2}}{[B]}$ The units of the rate constant are:
(A) $\mathrm{s}^{-1}$
(B) $\mathrm{M}^{2} \mathrm{~s}^{-1}$
(C) $\mathrm{M}^{-2} \mathrm{~s}^{-1}$
(D) $\mathrm{M}^{-1} \mathrm{~s}^{-1}$
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) $\mathrm{k}[\mathrm{A}][\mathrm{B}]$
(C) $k[A]^{2}[B]^{3}$
(D) $\mathrm{k}[\mathrm{A}][\mathrm{B}]^{2}$
3. Consider a reaction which is zeroth order; i.e. $d[A] / d t=-k[A]^{0}=-k$ For this reaction, a plot of $\qquad$ vs. time is a straight line with a $\qquad$ slope.
(A) $\ln ([A] t)$, negative
(B) $[A] t$, negative
(C) (A]t, positive
(D) $1 /[A]_{t}$, positive
4. 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 0.20 M , the initial rate is $1.5 \mathrm{Ms}^{-1}$. When the initial concentration of $A$ is 0.60 M , the initial rate is $40.5 \mathrm{Ms}^{-1}$. 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 $0.02 \mathrm{M}^{-1} \mathrm{~s}^{-1}$. 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
7. For the reaction, $\mathrm{Sc}^{3+}(\mathrm{aq})+2 \mathrm{Ni}^{+}(\mathrm{aq}) \rightarrow \mathrm{Sc}^{+}(\mathrm{aq})+2 \mathrm{Ni}^{2+}(\mathrm{aq})$, the reaction mechanism is:

$$
\begin{aligned}
& 2 \mathrm{Ni}^{+} \stackrel{K}{\longleftrightarrow} \mathrm{Ni}+\mathrm{Ni}^{2+} \quad \text { Fast equilibrium } \\
& \mathrm{Ni}+\mathrm{Sc}^{3+} \xrightarrow{k} \mathrm{Ni}^{2+}+\mathrm{Sc}^{+} \quad \text { Slow step }
\end{aligned}
$$

The overall rate equation for this reaction is:
(A) Rate $=k^{\prime} \frac{\left[\mathrm{Ni}^{2+}\right]\left[\mathrm{Sc}^{3+}\right]}{\left[\mathrm{Ni}^{+}\right]^{2}}$
(B) Rate $=k^{\prime} \frac{\left[\mathrm{Ni}^{+}\right]\left[\mathrm{Sc}^{3+}\right]}{\left[\mathrm{Ni}^{2+}\right]}$
(C) Rate $=k^{\prime} \frac{\left[\mathrm{Ni}^{+}\right]^{2}\left[\mathrm{Sc}^{3+}\right]}{\left[\mathrm{Ni}^{2+}\right]}$
(D) Rate $=k^{\prime}[\mathrm{Ni}]\left[\mathrm{Sc}^{3+}\right]$
8. The gas phase equilibrium, $2 \mathrm{~A}(\mathrm{~g}) \leftrightharpoons 3 \mathrm{~B}(\mathrm{~g})+\mathrm{C}(\mathrm{g})$. The equilibrium constant is $\mathrm{K}_{\mathrm{c}}=1 \times 10^{-4}$. 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, $2 \mathrm{~A}(\mathrm{~g}) \leftrightharpoons \mathrm{B}(\mathrm{g})$. 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, $\mathrm{K}_{\mathrm{c}}$ is approximately:
(A) 3.3
(B) 2.0
(C) 0.67
(D) 0.37
10. The equilibrium constant for the reaction, $\mathrm{N}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \leftrightharpoons 2 \mathrm{NO}(\mathrm{g})$ is $1.7 \times 10^{-3}$ (at $2300 \mathrm{~K})$. What is the equilibrium constant for the reaction, $6 \mathrm{NO}(\mathrm{g}) \leftrightharpoons 3 \mathrm{~N}_{2}(\mathrm{~g})+3 \mathrm{O}_{2}(\mathrm{~g})$ ?
(A) $4.9 \times 10^{-9}$
(B) 8.4
(C) $1.7 \times 10^{3}$
(D) $2.0 \times 10^{8}$

For \#11-\#13: Consider the gas phase equilibrium, $4 \mathrm{NO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \leftrightharpoons 2 \mathrm{~N}_{2} \mathrm{O}_{5}(\mathrm{~g})$. This is an Exothermic reaction.
11. For the above reaction, if $\mathrm{O}_{2}(\mathrm{~g})$ is added to the mixture, then the ratio, $\left[\mathrm{NO}_{2}\right] /\left[\mathrm{N}_{2} \mathrm{O}_{5}\right]$ will $\qquad$ and $\mathrm{K}_{\mathrm{c}}$ will $\qquad$
(A) decrease, Increase
(B) increase, decrease
(C) decrease, remain constant
(D) increase, remain constant
12. For the above reaction, if $\mathrm{Ne}(\mathrm{g})$ is added to the mixture in a container at fixed total pressure, the ratio, $\left[\mathrm{NO}_{2}\right] /\left[\mathrm{N}_{2} \mathrm{O}_{5}\right]$ will $\qquad$ and
$K_{c}$ will $\qquad$
(A) decrease, Increase
(B) increase, decrease
(C) decrease, remain constant
(D) increase, remain constant
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 $3 \times 10^{-11} \%$. Therefore, the concentration of Thalium, in parts per trillion (ppt) is:
(A) 0.30 ppt
(B) $3 \times 10^{-2} \mathrm{ppt}$
(C) $3 \times 10^{-4} \mathrm{ppt}$
(D) 3.0 ppt
15. What is the molality of a solution prepared by adding 138 grams of Glycerol ( $\mathrm{C}_{3} \mathrm{H}_{8} \mathrm{O}_{3}, \mathrm{M}=92$.) 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 $\left(\mathrm{C}_{3} \mathrm{H}_{6} \mathrm{O}_{2}, \mathrm{M}=62\right)$ to 400 grams of water is $1.30 \mathrm{~g} / \mathrm{mL}$. 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) $0.20 \mathrm{~m} \mathrm{~K}_{3} \mathrm{PO}_{4}$
(B) $0.30 \mathrm{~m} \mathrm{Ca}\left(\mathrm{NO}_{3}\right)_{2}$
(C) $0.70 \mathrm{~m} \mathrm{C}_{6} \mathrm{H}_{12} \mathrm{O}_{6}$
(D) 0.32 m NaBr
18. When a sample of the strong electrolyte, aluminum nitrate, $\mathrm{Al}\left(\mathrm{NO}_{3}\right)_{3}$, is dissolved in 600 mL of aqueous solution, the osmotic pressure at $25^{\circ} \mathrm{C}$ is 350 torr. Approximately how many moles of $\mathrm{Al}\left(\mathrm{NO}_{3}\right)_{3}$ are dissolved in the 600 mL ?
(A) $1.1 \times 10^{-2} \mathrm{~mol}$
(B) $2.8 \times 10^{-3} \mathrm{~mol}$
(C) $7.9 \times 10^{-3} \mathrm{~mol}$
(D) $3.9 \times 10^{-3} \mathrm{~mol}$
19. The normal boiling point of pure $\mathrm{CCl}_{4}(\mathrm{I})$ is $77.0^{\circ} \mathrm{C}$ and the boiling point elevation constant is $5.0^{\circ} \mathrm{C} / \mathrm{m}$. When 80. grams of an unknown compound is placed in 750 grams of $\mathrm{CCl}_{4}$, the boiling point of the solution is $80.5^{\circ} \mathrm{C}$. The Molar Mass of the unknown compound is approximately:
(A) $86 \mathrm{~g} / \mathrm{mol}$
(B) $69 \mathrm{~g} / \mathrm{mol}$
(C) $152 \mathrm{~g} / \mathrm{mol}$
(D) $114 \mathrm{~g} / \mathrm{mol}$
20. A 300. L sample of nitric acid $\left(\mathrm{HNO}_{3}, \mathrm{M}=63\right)$ 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 $(\mathrm{pH}>7)$ ?
(i) Potassium Lactate ( $\left.\mathrm{KLac}^{( }\right)$
(ii) Sodium Nitrate $\left(\mathrm{NaNO}_{3}\right)$
(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, $\mathrm{K}_{\mathrm{a}}$, 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 $2.0 \times 10^{-9}$. 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, $\mathrm{K}_{\mathrm{b}}=4.3 \times 10^{-10}$. 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 $0.40 \mathrm{M} \mathrm{HNO}_{3}$, which of the following would form a buffer?
(1) 0.6 mol of potassium lactate (KLac)
(2) 0.6 mol of sodium carbonate $\left(\mathrm{Na}_{2} \mathrm{CO}_{3}\right)$
(3) 1.2 mol of sodium acetate ( NaAc )
(4) 1.2 mol of ammonium bromide $\left(\mathrm{NH}_{4} \mathrm{Br}\right)$
(A) $1 \& 2 \& 3$
(B) $2 \& 3 \& 4$
(C) 3 only
(D) $2 \& 3$

For \#26-\#30: Consider Carbonic acid, $\mathrm{H}_{2} \mathrm{CO}_{3}$, which is a diprotic acid with acid dissociation constants, $\mathrm{Ka}=4.2 \times 10^{-7}$ and $\mathrm{Ka}^{\prime \prime}=4.8 \times 10^{-11}$.
26. What is the approximate pH of a 0.05 M solution of sodium carbonate, $\mathrm{Na}_{2} \mathrm{CO}_{3}$ ?
(A) 5.8
(B) 11.5
(C) 10.2
(D) 8.2
27. What is the approximate pH of a solution containing $0.25 \mathrm{M} \mathrm{K}_{2} \mathrm{CO}_{3}$ and $0.40 \mathrm{M} \mathrm{NaHCO}_{3}$ ?
(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 $\mathrm{HNO}_{3}$ to 4 L of $0.40 \mathrm{M} \mathrm{Na}_{2} \mathrm{CO}_{3}$ ?
(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 $0.40 \mathrm{M} \mathrm{H}_{2} \mathrm{CO}_{3}$ ?
(A) 5.9
(B) 10.9
(C) 9.0
(D) 9.7
30. What is the approximate ratio of concentrations, $\left[\mathrm{H}_{2} \mathrm{CO}_{3}\right] /\left[\mathrm{HCO}_{3}^{-}\right]$, of a buffer solution with $\mathrm{pH}=6.80$ ?
(A) 2.7
(B) 1.5
(C) 0.4
(D) 0.7
31. 600 mL of $0.40 \mathrm{M} \mathrm{NaOH}(\mathrm{aq})$ is needed to completely neutralize a sample of aqueous $\mathrm{H}_{3} \mathrm{PO}_{4}(\mathrm{aq})$. Approximately how many grams of $\mathrm{H}_{3} \mathrm{PO}_{4}(\mathrm{M}=98 \mathrm{~g} / \mathrm{mol})$ 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 $\mathrm{H}_{3} \mathrm{PO}_{4}$ sample
32. Consider the slightly soluble salt, $A B_{3}$. The solubility product constant for $A B_{3}$ is $\mathrm{K}_{\text {sp }}=6.0 \times 10^{-23}$. What is the solubility of $\mathrm{AB}_{3}$ in a solution containing 0.05 M $\mathrm{NaB}(\mathrm{aq})$ (which is a strong electrolyte)?
(A) $1.8 \times 10^{-20} \mathrm{M}$
(B) $7.4 \times 10^{-12} \mathrm{M}$
(C) $4.8 \times 10^{-19} \mathrm{M}$
(D) $1.2 \times 10^{-21} \mathrm{M}$
33. The enthalpy of vaporization of toluene is $39.2 \mathrm{~kJ} / \mathrm{mol}$ and the normal boiling point is $111^{\circ} \mathrm{C}$.. What is the entropy change of the system when 2 (two) moles of toluene gas condenses at $111^{\circ} \mathrm{C}$ ?
(A) $-204 \mathrm{~J} / \mathrm{K}$
(B) $-102 \mathrm{~J} / \mathrm{K}$
(C) $-706 \mathrm{~J} / \mathrm{K}$
(D) $+102 \mathrm{~J} / \mathrm{K}$
34. The enthalpy of vaporization of liquid benzene, $\mathrm{C}_{6} \mathrm{H}_{6}(\mathrm{I})$, is $+30.7 \mathrm{~kJ} / \mathrm{mol}$. What is the entropy change of the surroundings for the condensation of 0.50 mol of benzene gas at the boiling point, $80^{\circ} \mathrm{C}$ ?
(A) $-43.5 . \mathrm{J} / \mathrm{K}$
(B) $-87 \mathrm{~J} / \mathrm{K}$
(C) $+87 . \mathrm{J} / \mathrm{K}$
(D) $+43.5 \mathrm{~J} / \mathrm{K}$
35. For the reaction, $\mathrm{CH}_{3} \mathrm{OH}(\mathrm{g}) \rightleftarrows \mathrm{CO}(\mathrm{g})+2 \mathrm{H}_{2}(\mathrm{~g}), \Delta \mathrm{H}^{\circ}=+91 \mathrm{~kJ}$ and $\Delta \mathrm{S}^{\circ}=+221 \mathrm{~J} / \mathrm{K}$. What is the entropy change of the surroundings, $\Delta \mathrm{S}_{\text {surr, }}$ for the related reaction, $2 \mathrm{CO}(\mathrm{g})+4 \mathrm{H}_{2}(\mathrm{~g}) \rightleftarrows 2 \mathrm{CH}_{3} \mathrm{OH}(\mathrm{g})$, at $25^{\circ} \mathrm{C}$ ?
(A) $+611 \mathrm{~J} / \mathrm{K}$
(B) $-442 \mathrm{~J} / \mathrm{K}$
(C) $-611 \mathrm{~J} / \mathrm{K}$
(D) $+442 \mathrm{~J} / \mathrm{K}$
36. For the hypothetical reaction, $\mathrm{A} \leftrightharpoons \mathrm{B}, \Delta \mathrm{S}^{\circ}=-70 \mathrm{~J} / \mathrm{K}$ (independent of temperature). The equilibrium constant for the reaction at $150^{\circ} \mathrm{C}$ is $2.0 \times 10^{-3}$. What is the enthalpy change, $\Delta \mathrm{H}^{0}$ 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 $25^{\circ} \mathrm{C}$ and the entropy change is $-20 \mathrm{~J} / \mathrm{K}$. What can be concluded about the enthalpy change for this reaction?
(A) $\Delta \mathrm{H}<-18 \mathrm{~kJ}$
(B) $\Delta \mathrm{H}>+6 \mathrm{~kJ}$
(C) $\Delta \mathrm{H}<-6 \mathrm{~kJ}$
(D) No conclusion can be made about $\Delta \mathrm{H}$
38. The equilibrium constant for the reaction, $4 \mathrm{~N}_{2} \mathrm{O}_{5} \leftrightharpoons 4 \mathrm{~N}_{2}(\mathrm{~g})+10 \mathrm{O}_{2}(\mathrm{~g})$, is $5.8 \times 10^{82}$ at $25^{\circ} \mathrm{C}$. What is the approximate value of the Gibbs Free Energy of Formation of $\mathrm{N}_{2} \mathrm{O}_{5}(\mathrm{~g})$ ?
(A) $+120 . \mathrm{kJ} / \mathrm{mol}$
(B) $+240 . \mathrm{kJ} / \mathrm{mol}$
(C) $-120 \mathrm{~kJ} / \mathrm{mol}$
(D) Insufficient data is given to determine $\Delta \mathrm{G}^{0}\left(\mathrm{~N}_{2} \mathrm{O}_{5}\right)$

For \#39-\#40: The Enthalpy of Vaporization of $\mathrm{Cl}_{2}(\mathrm{liq})$ is $10.2 \mathrm{~kJ} / \mathrm{mol}$. The Entropy of Vaporization of $\mathrm{Cl}_{2}(\mathrm{liq})$ is $42.9 \mathrm{~J} / \mathrm{mol}-\mathrm{K}$.
39. The Entropy change of the universe, $\Delta$ Suniv, when 3 (three) moles of $\mathrm{Cl}_{2}$ (liq) vaporizes to $\mathrm{Cl}_{2}$ (gas) at $-60^{\circ} \mathrm{C}$ is approximately:
(A) $+5.0 \mathrm{~J} / \mathrm{mol}-\mathrm{K}$
(B) $-15.0 \mathrm{~J} / \mathrm{mol}-\mathrm{K}$
(C) $-5.0 \mathrm{~J} / \mathrm{mol}-\mathrm{K}$
(D) None of the above
40. The Gibbs Energy Change, $\Delta \mathrm{G}^{\mathrm{o}}$, when 3 (three) moles of $\mathrm{Cl}_{2}$ (gas) condenses to $\mathrm{Cl}_{2}$ (liq) at $-60^{\circ} \mathrm{C}$ is approximately:
(A) $+1.8 \mathrm{~kJ} / \mathrm{mol}$
(B) $+3.2 \mathrm{~kJ} / \mathrm{mol}$
(C) $-1.0 \mathrm{~kJ} / \mathrm{mol}$
(D) $-3.2 \mathrm{~kJ} / \mathrm{mol}$
41. Regarding the reaction, $\mathrm{Br}_{2} \mathrm{O}_{5}(\mathrm{~s})+5 \mathrm{CO}(\mathrm{g}) \rightarrow \mathrm{Br}_{2}(\mathrm{I})+5 \mathrm{CO}_{2}(\mathrm{~g})$, which of the following statements is/are true?
(1) $\mathrm{Br}_{2} \mathrm{O}_{5}$ is the reducing agent
(2) $\mathrm{Br}_{2}$ is oxidized
(3) CO is oxidized
(4) $\mathrm{Br}_{2} \mathrm{O}_{5}$ is reduced
(A) $1 \& 2$
(B) $3 \& 4$
(C) $2 \& 3 \& 4$
(D) $1 \& 2 \& 4$
42. For the electrochemical cell given by $\mathrm{Al}^{2} \mathrm{Al}^{3+}| | \mathrm{Cu}^{2+} \mid \mathrm{Cu}$, which of the following statements is/are correct?
(1) The cathode reaction is $\mathrm{Cu} \rightarrow \mathrm{Cu}^{2+}+2 \mathrm{e}^{-}$
(2) Electrons flow from the Al electrode to the Cu electrode through an external circuit
(3) The anode reaction is $\mathrm{Al} \rightarrow \mathrm{Al}^{3+}+3 e^{-}$
(4) If the half-cells are separated by a $\mathrm{K}_{2} \mathrm{SO}_{4}$ salt bridge, $\mathrm{SO}_{4}{ }^{2-}$ 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) $\mathrm{Cl}^{-}\left|\mathrm{Cl}_{2}\right|\left|\mathrm{Hg}^{2+}\right| \mathrm{Hg}$
(2) $\mathrm{I}^{-}\left|\mathrm{I}_{2}\right|\left|\mathrm{Ag}^{+}\right| \mathrm{Ag}$
(3) $\mathrm{Fe}\left|\mathrm{Fe}^{2+} \| \mathrm{Ni}^{2+}\right| \mathrm{Ni}$
(4) $\mathrm{Ag}\left|\mathrm{Ag}^{+} \| \mathrm{Cu}^{2+}\right| \mathrm{Cu}$
(A) $2 \& 3$
(B) $1 \& \& 3 \& 4$
(C) 3 only
(D) $1 \& 4$
44. For the electrochemical cell given by $\mathrm{Hg}\left|\mathrm{Hg}^{2+}\right|\left|\mathrm{Be}^{2+}\right| \mathrm{Be}$, the cell potential is $\mathrm{E}^{\circ}$ cell $=-2.71 \mathrm{~V}$. What is the reduction potential of $\mathrm{Be}^{2+}$ ?
(A) -1.85 V
(B) +3.57 V
(C) -3.57 V
(D) +1.85 V
.45. For the redox reaction given by $\mathrm{Fe}_{\mathrm{F}} \mathrm{Fe}^{3+}| | \mathrm{Cu}^{2+} \mid \mathrm{Cu}$, the cell potential is: $\mathrm{E}^{\mathrm{o}}$ cell $=+0.38 \mathrm{~V}$. 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) $3.1 \times 10^{-39}$
45. What is the standard Gibbs Free Energy change for the electrochemical reaction, $\mathrm{Ni}^{2+}+2 \mathrm{I}^{-} \rightarrow \mathrm{Ni}+\mathrm{I}_{2}$ ?
(A) -56 kJ
(B) +56 kJ
(C) +152 kJ
(D) +76 kJ
46. Consider the following electrochemical cell reaction (values in parentheses indicate Molar Concentrations): $\mathrm{Al}(\mathrm{s})\left|\mathrm{Al}^{3+}(1.50 \mathrm{M})\right|\left|\mathrm{Mn}^{2+}(0.002 \mathrm{M})\right| \mathrm{Mn}(\mathrm{s})$. Under standard conditions, the cell potential is $\mathrm{E}^{\circ}$ cell $=+0.48 \mathrm{~V}$.
What is the approximate cell potential, Ecell, at the concentrations shown in the reaction?
(A) -0.40 V
(B) +0.56 V
(C) +0.73 V
(D) +0.40 V
47. Consider a concentration cell, containing $0.10 \mathrm{M} \mathrm{H}^{+}$in the reference compartment (the cathode) and a solution containing $\mathrm{H}^{+}$in the sample cell (the anode) with $\mathrm{pH}=3.50$. (Note: $\mathbf{n}=\mathbf{1}$ for this cell)
In cell notation, this can be written as: $\mathrm{H}_{2}(\mathrm{~g})\left|\mathrm{H}^{+}(\mathrm{pH}=3.50)\right|\left|\mathrm{H}^{+}(0.10 \mathrm{M})\right| \mathrm{H}_{2}(\mathrm{~g})$. The cell potential, Ecell is approximately:
(A) +0.15 V
(B) +0.34 V
(C) -0.15 V
(D) -0.21 V
48. The concentration of Arsenic [As, $M=74.9$ ] in drinking water was determined using a concentration cell with 0.40 M Arsenic(III) nitrate, $\mathrm{As}\left(\mathrm{NO}_{3}\right)_{3}$, in the reference cell (the cathode) and a sample of water with an unknown concentration of $\mathrm{As}^{3+}(\mathrm{xx})$ in the sample cell (the anode). In cell notation, this can be written as: As(s)|As ${ }^{3+}(x x)| | A s^{3+}(0.40 \mathrm{M}) \mid A s(s)$.

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 ( $\mu \mathrm{g} / \mathrm{L}$ ) is approximately:
(A) $280 \mu \mathrm{~g} / \mathrm{L}$
(B) $3,900 \mu \mathrm{~g} / \mathrm{L}$
(C) $45 \mu \mathrm{~g} / \mathrm{L}$
(D) $111 \mu \mathrm{~g} / \mathrm{L}$
50. The solubility product of $\mathrm{Li}_{3} \mathrm{PO}_{4}$ is $\mathrm{K}_{s p}=3.2 \times 10^{-9}$. A Saturated solution of $\mathrm{Li}_{3} \mathrm{PO}_{4}$ 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+(Saturated Solution) || $\mathrm{Li}^{+}(2.00 \mathrm{M}) \mid \mathrm{Li}(\mathrm{s})$.
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 $\mathrm{Zn}(\mathrm{II})$ Fluoride, $\mathrm{ZnF}_{2}(\mathrm{aq})$ is placed in an electrolysis cell, and a voltage is applied, what will be the principal products of the electrolysis?
(A) $\mathrm{Zn}, \mathrm{F}_{2}$
(B) $\mathrm{F}_{2}, \mathrm{H}_{2}, \mathrm{OH}^{-}$
(C) $\mathrm{H}_{2}, \mathrm{OH}^{-}, \mathrm{O}_{2}, \mathrm{H}^{+}$
(D) $\mathrm{Zn}, \mathrm{O}_{2}, \mathrm{H}^{+}$
52. If aqueous Aluminum Iodide, $\mathrm{AlI}_{3}(\mathrm{aq})$ is placed in an electrolysis cell, and a voltage is applied, what will be the principal products of the electrolysis?
(A) $\mathrm{Al}, \mathrm{I}_{2}$
(B) $\mathrm{I}_{2}, \mathrm{H}_{2}, \mathrm{OH}^{-}$
(C) $\mathrm{H}_{2}, \mathrm{OH}^{-}, \mathrm{O}_{2}, \mathrm{H}^{+}$
(D) $\mathrm{Al}, \mathrm{O}_{2}, \mathrm{H}$
53. A current of 0.80 Amps (Coul/sec) is passed through a solution of Aluminum Nitrate, $\mathrm{Al}\left(\mathrm{NO}_{3}\right)_{3}(\mathrm{aq})$, for 7 . hours. Approximately what mass of solid Aluminum, $\mathrm{Al}(\mathrm{s})[\mathrm{M}=27]$, 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 $\mathrm{Sn}(\mathrm{s})$ [ $\mathrm{M}=118.7$ ] by electrolysis from a solution containing $\mathrm{Sn}\left(\mathrm{NO}_{3}\right)_{2}(\mathrm{aq})$ ? Assume that the voltage is 9 . Volts.
(A) 660 kJ
(B) $2,050 \mathrm{~kJ}$
(C) 330 kJ
(D) $1,320 \mathrm{~kJ}$
55. The type of nuclides with the highest number of stable isotopes have a(n)
$\qquad$ number of protons and $\qquad$ number of neutrons
(A) even , odd
(B) odd, even
(C) even, even
(D) odd, odd
56. What nuclide will undergo positron emission to form $\mathrm{Ar}-38$ ?
(A) $\mathrm{Ca}-42$
(B) $\mathrm{Cl}-38$
(C) $\mathrm{Cl}-40$
(D) $\mathrm{K}-38$
57. Which of the following is a likely decay path for $\mathrm{Ne}-19$. Stable isotopes in this range typically have $\mathrm{N} / \mathrm{Z}=1.0$.
(A) ${ }_{10}^{19} \mathrm{Ne} \rightarrow{ }_{-1}^{0} e+{ }_{11}^{19} \mathrm{Na}$
(B) ${ }_{10}^{19} \mathrm{Ne} \rightarrow{ }_{2}^{4} \mathrm{He}+{ }_{8}^{15} \mathrm{O}$
(C) ${ }_{10}^{19} \mathrm{Ne} \rightarrow{ }_{+1}^{0} e+{ }_{9}^{19} \mathrm{~F}$
(D) ${ }_{10}^{19} \mathrm{Ne} \rightarrow{ }_{1}^{1} \mathrm{H}+{ }_{9}^{18} \mathrm{~F}$
58. 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) $\mathrm{Cm}-249$
(C) $\mathrm{Md}-256$
(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.
$\mathrm{m}\left(1^{1} \mathrm{H}\right)=1.008 \mathrm{~g} / \mathrm{mol}, \mathrm{m}\left(0^{1} \mathrm{n}\right)=1.009 \mathrm{~g} / \mathrm{mol}, \mathrm{m}\left(36^{92} \mathrm{Kr}\right)=91.926 \mathrm{~g} / \mathrm{mol}$
(A) $4.7 \times 10^{8} \mathrm{~kJ} / \mathrm{mol}$
(B) $8.5 \times 10^{8} \mathrm{~kJ} / \mathrm{mol}$
(C) $8.5 \times 10^{11} \mathrm{~kJ} / \mathrm{mol}$
(D) $7.8 \times 10^{10} \mathrm{~kJ} / \mathrm{mol}$

