# leyChemistry 1A, Spring 2011 

Midterm 3<br>April 11, 2011<br>(90 min, closed book)

Name: $\qquad$

SID: $\qquad$

TA Name: $\qquad$

- There are 40 multiple choice questions worth 3 points each.
- There is only one correct answer for each question unless otherwise specified.
- Only answers on the Scantron form will be graded.
- Scantron must be properly filled in and cannot contain any smudges or other marks. Scantrons will not be rescanned!
- You can tear off the equation sheet and the periodic table for your convenience.
- You can use the page margin or the back of the pages as scratch paper.
- You can take the exam booklet with you after the exam.


## Quantum:

$$
\begin{aligned}
& \mathrm{E}=\mathrm{h} \nu \\
& \lambda \nu=\mathrm{c} \\
& \lambda_{\text {deBroglie }}=\mathrm{h} / \mathrm{p}=\mathrm{h} / \mathrm{mv} \\
& \mathrm{E}_{\text {kin }}(\mathrm{e}-)=\mathrm{h} \nu-\Phi=\mathrm{h} \nu-\mathrm{h} v_{0} \\
& E_{n}=-\frac{\mathrm{Z}^{2}}{n^{2}} R_{\infty} \\
& \Delta \mathrm{x} \Delta \mathrm{p} \sim \mathrm{~h} \\
& \mathrm{p}=\mathrm{mv} \\
& \mathrm{E}_{\mathrm{n}}=\mathrm{h}^{2} \mathrm{n}^{2} / 8 \mathrm{~mL}^{2} ; \mathrm{n}=1,2,3 . \ldots \\
& \mathrm{E}_{\mathrm{v}}=(\mathrm{v}+1 / 2) \mathrm{hA} / 2 \pi ; \mathrm{A}=(\mathrm{k} / \mathrm{m})^{1 / 2} \\
& \mathrm{E}_{\mathrm{n}}=\mathrm{n}(\mathrm{n}+1) \mathrm{hB} ; \mathrm{B}=\mathrm{h} / 8 \pi^{2} \mathrm{I} ; \mathrm{I}=2 \mathrm{mr}^{2} \\
& \mathrm{~m}=\mathrm{m}_{\mathrm{A}} \mathrm{~m}_{\mathrm{B}} /\left(\mathrm{m}_{\mathrm{A}}+\mathrm{m}_{\mathrm{B}}\right)
\end{aligned}
$$

## Ideal Gas:

$$
\begin{aligned}
& \mathrm{PV}=\mathrm{nRT} \\
& E_{k i n}=\frac{3}{2} R T \\
& \mathrm{v}_{\mathrm{rms}}=\sqrt{\frac{3 \mathrm{RT}}{\mathrm{M}}}
\end{aligned}
$$

## Constants:

$\mathrm{N}_{0}=6.02214 \times 10^{23} \mathrm{~mol}^{-1}$
$\mathrm{R}_{\infty}=2.179874 \times 10^{-18} \mathrm{~J}$
$\mathrm{R}_{\infty}=3.28984 \times 10^{15} \mathrm{~Hz}$
$\mathrm{k}=1.38066 \times 10^{-23} \mathrm{~J} \mathrm{~K}^{-1}$
$\mathrm{h}=6.62608 \times 10^{-34} \mathrm{~J} \mathrm{~s}$
$\mathrm{m}_{\mathrm{e}}=9.101939 \times 10^{-31} \mathrm{~kg}$
$\mathrm{c}=2.99792 \times 10^{8} \mathrm{~m} \mathrm{~s}^{-1}$
$\mathrm{T}(\mathrm{K})=\mathrm{T}(\mathrm{C})+273.15$
$\mathrm{F}=96,485 \mathrm{C} / \mathrm{mol}$
$1 \mathrm{~V}=1 \mathrm{~J} / \mathrm{C}$
Gas Constant:
$\mathrm{R}=8.31451 \mathrm{~J} \mathrm{~K}^{-1} \mathrm{~mol}^{-1}$
$\mathrm{R}=8.20578 \times 10^{-2} \mathrm{~L} \mathrm{~atm} \mathrm{~K}^{-1} \mathrm{~mol}^{-1}$
$1 \mathrm{~nm}=10^{-9} \mathrm{~m}$
$1 \mathrm{~kJ}=1000 \mathrm{~J}$
$1 \mathrm{~atm}=760 \mathrm{~mm} \mathrm{Hg}=760$ torr $\approx 1$ bar $1 \mathrm{~L} \mathrm{~atm} \approx 100 \mathrm{~J}$

## Thermodynamics:

$$
\Delta \mathrm{G}^{\circ}=\Delta \mathrm{H}^{\circ}-\mathrm{T} \Delta \mathrm{~S}^{\circ}
$$

$$
\Delta \mathrm{H}^{\circ}=\sum \Delta \mathrm{H}_{\mathrm{f}}^{\circ} \text { (products) }-\sum \Delta \mathrm{H}_{\mathrm{f}}^{\circ} \text { (reactants) }
$$

$$
\Delta \mathrm{S}^{\circ}=\sum \mathrm{S}^{\circ} \text { (products) }-\Sigma \mathrm{S}^{\circ} \text { (reactants) }
$$

$$
\Delta \mathrm{G}^{\circ}=\sum \Delta \mathrm{G}_{\mathrm{f}}^{\circ} \text { (products) }-\sum \Delta \mathrm{G}_{\mathrm{f}}^{\circ} \text { (reactants) }
$$

$$
\mathrm{S}=\mathrm{k}_{\mathrm{B}} \ln \mathrm{~W}
$$

$$
\Delta \mathrm{S}=\mathrm{q}_{\mathrm{rev}} / \mathrm{T}
$$

$$
\Delta \mathrm{E}=\mathrm{q}+\mathrm{w}
$$

$$
\mathrm{w}=-\mathrm{P}_{\mathrm{ext}} \Delta \mathrm{~V}
$$

$$
\text { for } \mathrm{aA}+\mathrm{bB} \rightleftarrows \mathrm{cC}+\mathrm{dD}
$$

$$
Q=\frac{[C]^{c}[D]^{d}}{[A]^{a}[B]^{b}} \quad \text { At equilibrium, } \mathrm{Q}=\mathrm{K}
$$

$$
\Delta \mathrm{G}=\Delta \mathrm{G}^{\circ}+\mathrm{RT} \ln \mathrm{Q}
$$

$$
\mathrm{G}=\mathrm{G}^{\circ}+\mathrm{RT} \ln (\mathrm{a}) ; \mathrm{a}=\text { activity }=\gamma \mathrm{P} / \mathrm{P}^{\circ} \text { or } \gamma[\mathrm{A}] /[\mathrm{A}]^{\circ}
$$

$$
\Delta \mathrm{G}^{\circ}=-\mathrm{RT} \ln \mathrm{~K}
$$

$$
\Delta G^{\circ}=-n F \Delta \epsilon^{\circ}
$$

$$
\Delta \mathrm{C}=\Delta \mathrm{E}^{\mathrm{o}}-(\mathrm{RT} / \mathrm{nF}) \ln \mathrm{Q}
$$

$$
\ln K=-\frac{\Delta H^{\circ}}{R} \frac{1}{T}+\frac{\Delta S^{\circ}}{R}
$$

$$
\Delta \mathrm{T}=\mathrm{i} \mathrm{k}_{\mathrm{b}, \mathrm{f}} \mathrm{~m}
$$

$$
\Pi=\mathrm{iMRT}
$$

$$
\mathrm{P}_{\text {total }}=\mathrm{P}_{\mathrm{A}}+\mathrm{P}_{\mathrm{B}}=\mathrm{X}_{\mathrm{A}} \mathrm{P}_{\mathrm{A}}{ }^{\circ}+\mathrm{X}_{\mathrm{B}} \mathrm{P}_{\mathrm{B}}^{\circ}
$$

$$
\mathrm{q}=\mathrm{cm}\left(\mathrm{~T}_{2}-\mathrm{T}_{1}\right)
$$

## Kinetics:

$$
\begin{aligned}
& {[\mathrm{A}]_{\mathrm{t}}=[\mathrm{A}]_{0} \mathrm{e}^{-\mathrm{kt}}} \\
& \ln [\mathrm{~A}]_{\mathrm{t}}=\ln [\mathrm{A}]_{0}-\mathrm{kt} \\
& \mathrm{t}_{1 / 2}=\ln 2 / \mathrm{k} \\
& 1 /[\mathrm{A}]_{\mathrm{t}}=1 /[\mathrm{A}]_{0}+\mathrm{kt} \\
& \mathrm{k}=\mathrm{A} \mathrm{e}^{(-\mathrm{Ea} / \mathrm{RT})} \\
& \ln \left(\mathrm{k}_{1} / \mathrm{k}_{2}\right)=\mathrm{E}_{\mathrm{a}} / \mathrm{R}\left(1 / \mathrm{T}_{2}-1 / \mathrm{T}_{1}\right) \\
& \mathrm{t}_{1 / 2}=1 /[\mathrm{A}]_{0} \mathrm{k} \\
& \mathrm{t}_{1 / 2}=[\mathrm{A}]_{0} / \mathrm{kt}
\end{aligned}
$$

1. One of the underlying assumptions of the kinetic theory of gases is
A. Gas molecules move randomly at fixed velocities.
B. Gas molecules only exert forces when they move close to one another.
C. Energy is lost only during collisions with the container.
D. All of these are assumptions of the kinetic theory of gases.
E. None of these is an assumption of the kinetic theory of gases.

ANS: E
2. At $25^{\circ} \mathrm{C}$ we have the following root-mean-square speeds (in m/s): $\mathrm{Xe}-238, \mathrm{Ar}-431, \mathrm{He}-$ 1360 . Thus sulfur dioxide gas molecules at $25^{\circ} \mathrm{C}$ should have a root-mean-square speed
A. $<238 \mathrm{~m} / \mathrm{s}$.
B. $>238 \mathrm{~m} / \mathrm{s}$ but $<431 \mathrm{~m} / \mathrm{s}$.
C. $>431 \mathrm{~m} / \mathrm{s}$ but $<1360 \mathrm{~m} / \mathrm{s}$.
D. $>1360 \mathrm{~m} / \mathrm{s}$.
E. This cannot be predicted.

ANS: B
3. At a certain absolute temperature the average speed of $\underline{H}_{2}$ molecules is $1780 \mathrm{~m} / \mathrm{s}$. Thus at an absolute temperature twice as great, the average speed of He atoms would be
A. $890 \mathrm{~m} / \mathrm{s}$
B. $1260 \mathrm{~m} / \mathrm{s}$
C. $1780 \mathrm{~m} / \mathrm{s}$
D. $3560 \mathrm{~m} / \mathrm{s}$
E. none of these

ANS: C
4. What happens to bicycle tires on hot days versus cold days?
A. The tires expand because an increase in temperature corresponds to an increase in molecular speed which causes an increase in the force of gas collisions on the inside of the tire.
B. The tires shrink because an increase in temperature corresponds to a decrease in molecular speed which causes a decrease in the force of gas collisions on the inside of the tire.
C. The tires expand because an increase in temperature causes the gas molecules to be less attracted to each other and therefore they take up more space.
D. The tires shrink because an increase in temperature causes the gas molecules to be more attracted to each other and therefore they take up less space.
E. The tires stay the same volume because the amount of gas contained within the tire does not change.
ANS: A
5. Which of the following conditions should minimize deviations from ideal gas behavior in a real gas?
A. high $T$ and high $P$
B. low $T$ and low $P$
C. low $T$ and high $P$
D. high $T$ and low $P$
E. This requires specific knowledge of the gas to predict.

ANS: D
6. What is the pressure of 2.50 mol of $\mathrm{CO}_{2}$ in a $1.00-\mathrm{L}$ flask at 298 K when calculated using the van der Waals equation? For $\mathrm{CO}_{2}$, the values of the van der Waals constants are $\mathrm{a}=3.59 \mathrm{~L}^{2}$ $\mathrm{atm} / \mathrm{mol}^{2}$ and $\mathrm{b}=0.0427 \mathrm{~L} / \mathrm{mol}$. The van der Waal's equation is given below.
$\left(P+a \frac{n^{2}}{V^{2}}\right)(V-n b)=n R T$
A. 5.74
B. 46.0
C. 59.5
D. 61.1
E. 68.5

ANS: B
7. The temperature at point $a$ is the

A. critical point.
B. triple point.
C. absolute freezing point.
D. normal freezing point.
E. normal boiling point.

ANS: D
8. The transition from Phase I to Phase II is called

A. melting.
B. freezing.
C. sublimation.
D. evaporation.
E. condensation.

ANS: A
9. For a balloon expanding against atmospheric pressure, the work $w$ done on the balloon is
A. positive
B. negative
C. of indeterminate sign
D. zero
E. This cannot be answered without additional information. ANS: B

10 . A gas is compressed from 39.92 L to 12.97 L at a constant pressure of 5.00 atm . In the course of this compression 9.82 kJ of energy is released as heat. The heat $q$ for this process is
A. 135 kJ
B. -135 kJ
C. -9.82 kJ
D. 9.82 kJ
E. This cannot be determined without additional information.

ANS: C
11. Which of the following processes is expected to be endothermic?
A. combustion of a hydrocarbon
B. dilution of concentrated sulfuric acid with water
C. freezing of liquid $\mathrm{N}_{2}$ at its normal melting point ( 53 K )
D. vaporization of zinc at its normal boiling point (1179 K)
E. All of these are endothermic.

ANS: D
12. The First Law of Thermodynamics is the law of
A. conservation of energy
B. conservation of matter
C. conservation of enthalpy
D. All of these are involved.
E. none of these

ANS: A
13. Propanol (pro) burns cooler than isopropanol (iso). What can you conclude about the isomerization of propanol to isopropanol?

isopropanol
propanol
A. $\Delta \mathrm{H}^{\circ}{ }_{\mathrm{rxn}}<0$
B. $\Delta \mathrm{H}^{\circ}{ }_{\mathrm{rxn}}>0$
C. $\Delta \mathrm{H}_{\mathrm{rxn}}^{\circ}=0$
D. $\Delta \mathrm{H}^{\circ}{ }_{\mathrm{f}}($ propanol $)=\Delta \mathrm{H}^{\circ}{ }_{\mathrm{f}}($ isopropanol $)$

E . none of these
B
14. Gaseous methane $\left(\mathrm{CH}_{4}\right)$ burns completely in air with a heat of combustion of $-890 \mathrm{~kJ} \mathrm{~mol}^{-1}$. Calculate the heat liberated by burning 4.00 L of $\mathrm{CH}_{4}(\mathrm{~g})$ measured at $101^{\circ} \mathrm{C}$ and 723 torr.
Assume methane is an ideal gas at this condition.
A. 110 kJ
B. 159 kJ
C. 409 kJ
D. 890 kJ
E. none of these

ANS: A
15. Determine the heat of reaction for the process
$\mathrm{Fe}_{2} \mathrm{O}_{3(\mathrm{~s})}+\mathrm{FeO}_{(\mathrm{s})} \rightarrow \mathrm{Fe}_{3} \mathrm{O}_{4(\mathrm{~s})}$
using the information given below:
$2 \mathrm{Fe}_{(\mathrm{s})}+\mathrm{O}_{2(\mathrm{~g})} \rightarrow 2 \mathrm{FeO}_{(\mathrm{s})}$

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

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

$$
\begin{aligned}
& \Delta \mathrm{H}^{\circ}=-544.0 \mathrm{~kJ} \\
& \Delta \mathrm{H}^{\circ}=-1648.8 \mathrm{~kJ} \\
& \Delta \mathrm{H}^{\circ}=-1118.4 \mathrm{~kJ}
\end{aligned}
$$

A. -1074.0 kJ
B. -22.0 kJ
C. 22.2 kJ
D. 249.8 kJ
E. 1074.0 kJ

ANS: B
16. Which of the following is not a state function?
A. $q$
B. $E$
C. $H$
D. $P V$
E. All of these are state functions.

ANS: A
17. A 43.9-g piece of copper ( $c_{\mathrm{s}}=0.385 \mathrm{~J} \mathrm{~g}^{-1} \mathrm{~K}^{-1}$ ) at $135.0^{\circ} \mathrm{C}$ is plunged into 254 g of water at $39.0^{\circ} \mathrm{C}$. Assuming that no heat is lost to the surroundings, what will the final temperature of the system be? The heat capacity of water is $4.184 \mathrm{~J} \mathrm{~g}^{-1} \mathrm{~K}^{-1}$.
A. $100.0^{\circ} \mathrm{C}$
B. $87.0^{\circ} \mathrm{C}$
C. $53.1^{\circ} \mathrm{C}$
D. $40.5^{\circ} \mathrm{C}$
E. none of these

ANS: D
18. The following process of bond-breaking $\rightarrow$ bond formation corresponds to the combustion of which compound?

$$
3 \mathrm{C}-\mathrm{H}+\mathrm{O}-\mathrm{H}+\mathrm{C}-\mathrm{O}+3 / 2 \mathrm{O}=\mathrm{O} \rightarrow 2 \mathrm{C}=\mathrm{O}+4 \mathrm{O}-\mathrm{H}
$$

A.Ethanol B.Ethane $\quad$ C.Methanol
19. Methane can react with a fluorine atom to produce a methyl radical and hydrogen fluoride.


$$
\Delta \mathrm{H}^{\circ}{ }_{\mathrm{rxn}}=-153 \mathrm{~kJ} / \mathrm{mol}
$$

Based on the $\Delta \mathrm{H}^{\circ}{ }_{\mathrm{rxn}}$, which of the following is true?
A. Breaking chemical bonds releases energy.
B. All reactions involving bond-breaking and bond formation are exothermic.
C. The H-F bond is stronger than the C-H bond.
D. The H-F bond is weaker than the C-H bond.
E. The heat of the reactants is absorbed by the products.

C
20. The decomposition:
$\mathrm{H}_{2} \mathrm{O}_{2} \rightarrow \mathrm{H}_{2} \mathrm{O}+1 / 2 \mathrm{O}_{2}$
is exothermic. What is the molecular structure of $\mathrm{H}_{2} \mathrm{O}_{2}$ ?

ANS: A
21. As a demonstration, three balloons were filled with the following mixtures. Which balloons produces the most heat when the contents undergo combustion reactions?
A. 2 moles $\mathrm{C}_{8} \mathrm{H}_{18}$
B. 3 moles $\mathrm{C}_{8} \mathrm{H}_{18}$
C. 2 moles $\mathrm{C}_{8} \mathrm{H}_{18}$ and 2 moles $\mathrm{O}_{2}$

B
22. Which process is accompanied by the largest increase in entropy?

A. I $\rightarrow$ II
B. I $\rightarrow$ III
C. II $\rightarrow$ III
D. III $\rightarrow$ IV
E. V $\rightarrow$ VI

B
23. Which of the following processes should show the greatest increase in entropy?
A. $\mathrm{C}_{6} \mathrm{H}_{6}(\mathrm{l})+\frac{15}{2} \mathrm{O}_{2}(g) \rightarrow 6 \mathrm{CO}_{2}(g)+3 \mathrm{H}_{2} \mathrm{O}(g)$
B. $2 \mathrm{NO}_{2}(g) \rightarrow \mathrm{N}_{2} \mathrm{O}_{4}(g)$
C. $\mathrm{C}_{2} \mathrm{H}_{4}(\mathrm{~g})+\mathrm{H}_{2}(\mathrm{~g}) \rightarrow \mathrm{C}_{2} \mathrm{H}_{6}(\mathrm{~g})$
D. $\mathrm{BaS}(s)+2 \mathrm{NaNO}_{3}(s) \rightarrow \mathrm{Ba}\left(\mathrm{NO}_{3}\right)_{2}(s)+\mathrm{Na}_{2} \mathrm{~S}(s)$
E. This cannot be predicted without additional information.

ANS: A
24. Calculate the value of $\Delta S^{\circ}$ for the reaction shown:

$$
\mathrm{N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{NH}_{3}(\mathrm{~g})
$$

At $25^{\circ} \mathrm{C}$ the values of entropy in $\mathrm{J} \mathrm{K}^{-1} \mathrm{~mol}^{-1}$ are nitrogen, 191.61: hydrogen, 130.68 ; and ammonia, 192.77.
A. $-198.11 \mathrm{~J} / \mathrm{K}$
B. $-259.03 \mathrm{~J} / \mathrm{K}$
C. $-390.88 \mathrm{~J} / \mathrm{K}$
D. $-393.20 \mathrm{~J} / \mathrm{K}$
E. $-969.19 \mathrm{~J} / \mathrm{K}$

ANS: A
25. Consider the process whereby 1.00 mol of $\mathrm{H}_{2} \mathrm{O}(s)$ is melted to $\mathrm{H}_{2} \mathrm{O}(\mathrm{l})$ at the normal melting point of ice, 273.15 K . Which of the following is true for this process?
A. $\Delta H>0$
B. $\Delta G=0$
C. $\Delta H=T \Delta S$
D. All of these are true.
E. None of these is true.

ANS: D
26. Calculate $\Delta \mathrm{G}_{\mathrm{rxn}}(\mathrm{kJ})$ at 298 K under the conditions shown below for the following reaction.
$\mathrm{Fe}_{2} \mathrm{O}_{3}(\mathrm{~s})+3 \mathrm{CO}(\mathrm{g}) \rightarrow 2 \mathrm{Fe}(\mathrm{s})+3 \mathrm{CO}_{2}(\mathrm{~g})$

$$
\Delta \mathrm{G}^{0}=-28.0 \mathrm{~kJ} \text { at } 298 \mathrm{~K}
$$

$\mathrm{P}(\mathrm{CO})=1.4 \mathrm{~atm}, \mathrm{P}\left(\mathrm{CO}_{2}\right)=2.1 \mathrm{~atm}$
A. 31.0 kJ
B. 2.99 kJ
C. -30.7 kJ
D. 17.5 kJ
E. -25.0 kJ

ANS: E
27. Which is the correct plot of $\Delta \mathrm{G}^{\circ}$ vs. T for the reaction $\mathrm{H}_{2} \mathrm{O}(\mathrm{l}) \leftrightarrow \mathrm{H}_{2} \mathrm{O}$ (g)?


ANS: C
28. For the reaction, $\mathrm{X}+\mathrm{Y} \rightarrow \mathrm{A}+\mathrm{B}, \Delta G_{\mathrm{rxn}}$ is $-1324 \mathrm{~kJ} \mathrm{~mol}^{-1}$. Which one of the following statements CANNOT be concluded concerning this reaction?
A. The reaction is thermodynamically favorable.
B. The reaction is spontaneous as written.
C. The products are more stable than the reactants.
D. The reaction will proceed rapidly from left to right.
E. All of these are valid.

ANS: D
29. Which is true for the following reaction under standard conditions?
$\mathrm{C}_{2} \mathrm{H}_{6}(\mathrm{~g}) \rightarrow \mathrm{C}_{2} \mathrm{H}_{4}(\mathrm{~g})+\mathrm{H}_{2}(\mathrm{~g})$
$\Delta \mathrm{H}^{\circ}$ is 137 kJ and $\Delta \mathrm{S}^{\circ}$ is $120 \mathrm{~J} / \mathrm{K}$.
A. spontaneous at all temperatures
B. spontaneous only at high temperature
C. spontaneous only at low temperature
D. not spontaneous at all temperatures
E. cannot be determined

ANS: B
30. If a catalyst is added to a chemical reaction, the equilibrium yield of a product will be
$\qquad$ , and the time taken to come to equilibrium will be $\qquad$ than before.
A. higher; less
B. lower; the same
C. higher; the same
D. the same; less
E. lower; less

ANS: D
31. Consider the endothermic reaction
$\mathrm{C}(\mathrm{s})+\mathrm{CO}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{CO}(\mathrm{g})$
If such a system at equilibrium is heated, equilibrium will $\qquad$ because $\qquad$ .
A. be unchanged; temperature has no effect on equilibrium
B. shift to the left; increased temperature favors an exothermic reaction
C. shift to the right; increased temperature favors an exothermic reaction
D. shift to the right; increased temperature favors an endothermic reaction
E. shift to the left; increased temperature favors an endothermic reaction

ANS: D
32. Consider the reaction
$\mathrm{NH}_{4} \mathrm{Cl}(\mathrm{s}) \rightleftharpoons \mathrm{NH}_{3}(\mathrm{~g})+\mathrm{HCl}(\mathrm{g})$.
If the pressure is decreased on an equilibrium mixture of these three substances, equilibrium will
$\qquad$ ,because $\qquad$ .
A. shift to the right; lower pressure favors fewer moles of gas
B. shift to the right; lower pressure favors more moles of gas
C. shift to the left; lower pressure favors fewer moles of gas
D. shift to the left; lower pressure favors more moles of gas
E. be unchanged; solid $\mathrm{NH}_{4} \mathrm{Cl}$ does not appear in the equilibrium constant expression.

## ANS: B

33. Consider the equilibrium system $\mathrm{C}(\mathrm{s})+\mathrm{CO}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{CO}(\mathrm{g})$.
If more $\mathrm{C}(\mathrm{s})$ is added, the equilibrium will $\qquad$ ; if CO is removed the equilibrium will $\qquad$ .
A. shift to the left; shift to the left
B. shift to the right; shift to the right
C. shift to the right; shift to the left
D. be unchanged; shift to the left
E. be unchanged; shift to the right

ANS: E
34. Which set of conditions describes a reaction that is most likely to proceed?
A. endothermic, decreasing entropy, high activation energy
B. exothermic, decreasing entropy, high activation energy
C. exothermic, increasing entropy, high activation energy
D. exothermic, increasing entropy, low activation energy
E. endothermic, decreasing entropy, low activation energy

ANS: D
35. What is the value of the equilibrium constant at $25^{\circ} \mathrm{C}$ for a reaction, if the value of $\Delta \mathrm{G}^{\circ}{ }_{\mathrm{rxn}}$ is 47.8 kJ at $25^{\circ} \mathrm{C}$ ?
A. 1.70
B. 6.88
C. $2.30 \times 10^{2}$
D. $2.74 \times 10^{5}$
E. $2.37 \times 10^{8}$

ANS: E
36. If the equilibrium constants for the two reactions

$$
2 \mathrm{HCl}(\mathrm{~g}) \rightleftharpoons \mathrm{H}_{2}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g})
$$

and

$$
\mathrm{I}_{2}(\mathrm{~g})+\mathrm{Cl}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{ICl}(\mathrm{~g})
$$

are denoted $\mathrm{K}_{1}$ and $\mathrm{K}_{2}$ respectively, then the equilibrium constant for the reaction
$2 \mathrm{HCl}(\mathrm{g})+\mathrm{I}_{2}(\mathrm{~g}) \rightleftharpoons 2 \mathrm{ICl}(\mathrm{g})+\mathrm{H}_{2}(\mathrm{~g})$ equals
A. $\left(\mathrm{K}_{1} / \mathrm{K}_{2}\right)^{2}$.
B. $\left(K_{1} K_{2}\right)^{2}$.
C. $\mathrm{K}_{1} \mathrm{~K}_{2}$.
D. $\mathrm{K}_{1}+\mathrm{K}_{2}$.
E. $\mathrm{K}_{1} \mathrm{~K}_{2} / 2$.

ANS: C
37. Consider the equilibrium reaction
$2 \mathrm{NO}_{2}(\mathrm{~g}) \rightleftharpoons \mathrm{N}_{2} \mathrm{O}_{4}(\mathrm{~g})$.
A sample of pure $\mathrm{NO}_{2}(\mathrm{~g})$ at 0.140 atm is allowed to come to equilibrium. It is then found that $57.0 \%$ of the $\mathrm{NO}_{2}(\mathrm{~g})$ has reacted to form $\mathrm{N}_{2} \mathrm{O}_{4}(\mathrm{~g})$. What is the value of K ?
A. 0.211
B. 0.377
C. 0.754
D. 4.73
E. 11.0

ANS: E
38. In the experiment "How Hot is that Flame", A bomb calorimeter is calibrated with benzoic acid, for which the specific heat of combustion is $-26.41 \mathrm{~kJ} / \mathrm{g}$. The combustion of 1.327 g of benzoic acid causes the temperature of the calorimeter to rise from $25.998^{\circ} \mathrm{C}$ to $27.918^{\circ} \mathrm{C}$. Calculate the calorimeter constant $\mathrm{K}_{\text {cal }}$.
A. $10.37 \mathrm{~kJ} \mathrm{~K}^{-1}$
B. $18.25 \mathrm{~kJ} \mathrm{~K}^{-1}$
C. $38.21 \mathrm{~kJ} \mathrm{~K}^{-1}$
D. $67.29 \mathrm{~kJ} \mathrm{~K}^{-1}$
E. none of these

ANS: B
39. A student explored the thermodynamic properties of the dissolution of borax using titrations at different temperatures. The following plot was obtained. What is $\Delta \mathrm{S}^{\circ}$ in $\mathrm{J} / \mathrm{mol} \cdot \mathrm{K}$ for the dissolution of borax?

F. 53.727
G. -16780
H. 446.69
I. $\quad 140.26$
J. -140.26

C
40. A student has a solution of potassium chromate $\left(\mathrm{K}_{2} \mathrm{CrO}_{4}\right)$ in a test tube. The solution is yellow. To this test tube, several drops of a solution of sulfuric acid are added. Given the reactions below, predict what changes will occur after the addition of acid and why the changes occur.

A. The solution remains yellow because sulfuric acid is colorless
B. The solution remains yellow because no additional reactions take place
C. The solution turns a lighter yellow because it is more dilute after adding acid
D. The solution turns orange because the acid reacts with the chromate to convert it to dichromate which is orange.
D

Page 14 of 15


