INFORMATION PAGE (Use for reference and for scratch paper) Constants and Conversion Factors:
$\mathrm{R}=0.082 \mathrm{~L}-\mathrm{atm} / \mathrm{mol}-\mathrm{K}=8.31 \mathrm{~J} / \mathrm{mol}-\mathrm{K}=8.31 \mathrm{kPa}-\mathrm{L} / \mathrm{mol}-\mathrm{K}$
$1 \mathrm{~L}-\mathrm{atm}=101 \mathrm{~J}$
$1 \mathrm{~L}-\mathrm{bar}=100 \mathrm{~J}$
$1 \mathrm{kPa}-\mathrm{L}=1 \mathrm{~J}$
1 bar $=100 \mathrm{kPa}$
1 bar = 750 torr
$1 \mathrm{~atm}=760$ torr

Trouton's Rule: $\Delta_{\text {vap }} S^{\circ}=85 . \mathrm{J} / \mathrm{mol}-\mathrm{K}$

The relation between the Molar Mass (M), density ( $\rho$ ) and Molar Volume ( $\mathrm{V}_{\mathrm{m}}$ ) of a material is: $\rho=\frac{M}{V_{m}}$

## CHEM 5200-Exam 2 - October 17, 2017

## Name

$\qquad$

## (45) MULTIPLE CHOICE [3 points per question] (Circle the ONE correct answer)

For \#1-\#2: Consider the reaction, $2 \mathrm{~N}_{2} \mathrm{O}_{5}(\mathrm{~g}) \rightarrow 4 \mathrm{NO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g})$. Relevant thermodynamic data is given in the table below:

| Compound | $\mathbf{S}_{\mathbf{m}}{ }^{\circ}$ | $\Delta \mathbf{G}^{0}$ |
| :--- | :--- | :--- |
| $\mathrm{NO}_{2}(\mathrm{~g})$ | $240 . \mathrm{J} / \mathrm{mol}-\mathrm{K}$ | $+51 . \mathrm{kJ} / \mathrm{mol}$ |
| $\mathrm{O}_{2}(\mathrm{~g})$ | 205. |  |
| $\mathrm{~N}_{2} \mathrm{O}_{5}(\mathrm{~g})$ | 356. | +115. |

1. The Gibbs Energy Change $\left(\Delta_{\mathrm{r}} \mathrm{G}^{\circ}\right)$ for the above reaction at $25^{\circ} \mathrm{C}$ (in kJ) is approximately:
(A) -128. kJ
(B) $-26 . \mathrm{kJ}$
(C) +26. kJ
(D) Insufficient data is given
2. The standard Enthalpy Change ( $\Delta r^{H} \mathrm{H}^{\circ}$ ) for the above reaction at $25^{\circ} \mathrm{C}$ (in kJ$)$ is approximately:
(A) -161 kJ
(B) +109 kJ
(C) $-109 . \mathrm{kJ}$
(D) Insufficient data is given
3. When two (2) moles of $\mathrm{N}_{2}(\mathrm{~g})$ at $25^{\circ} \mathrm{C}$ and 50 L are compressed reversibly and isothermally to a final volume of 20 L , the entropy change is
(A) $-7.6 \mathrm{~J} / \mathrm{K}$
(B) $+15.2 \mathrm{~J} / \mathrm{K}$
(C) $-4.5 \mathrm{~kJ} / \mathrm{K}$
(D) $-15.2 \mathrm{~J} / \mathrm{K}$
4. The constant pressure molar heat capacity of $\mathrm{CO}_{2}(\mathrm{~g})$ is $37.1 \mathrm{~J} / \mathrm{mol}-\mathrm{K}$. What is $\Delta \mathrm{S}$ when 5 moles of $\mathrm{CO}_{2}(\mathrm{~g})$ is heated at constant volume from $100^{\circ} \mathrm{C}$ to $400^{\circ} \mathrm{C}$ ?
(A) $+25 \mathrm{~J} / \mathrm{K}$
(B) $+109 \mathrm{~J} / \mathrm{K}$
(C) $+85 \mathrm{~J} / \mathrm{K}$
(D) $+300 \mathrm{~J} / \mathrm{K}$
5. A sample of 2 moles of $\mathrm{N}_{2}(\mathrm{~g})$ at 50 kPa and 20 L is compressed reversibly and adiabatically to a final pressure of 400 kPa . What is $\Delta \mathrm{S}$ for this process?
(A) $+34.6 \mathrm{~J} / \mathrm{K}$
(B) $0 \mathrm{~J} / \mathrm{K}$
(C) $-34.6 \mathrm{~J} / \mathrm{K}$
(D) $-97.4 \mathrm{~J} / \mathrm{K}$
6. The normal boiling point of napthalene, $\mathrm{C}_{10} \mathrm{H}_{8}$, is $218{ }^{\circ} \mathrm{C}$. An estimate of the Enthalpy of Vaporization of Napthalene using Trouton's Rule is:
(A) $41.7 \mathrm{~kJ} / \mathrm{mol}$
(B) $52.1 \mathrm{~kJ} / \mathrm{mol}$
(C) $18.5 \mathrm{~kJ} / \mathrm{mol}$
(D) $28.9 \mathrm{~kJ} / \mathrm{mol}$
7. The normal boiling point of methanol is $64^{\circ} \mathrm{C}$. The Enthalpy of Vaporization of methanol is $35.3 \mathrm{~kJ} / \mathrm{mol}$. What is the entropy change of the system when two (2) moles of liquid methanol are vaporized reversibly to the gas at $64^{\circ} \mathrm{C}$ ?
(A) $+210 \mathrm{~J} / \mathrm{K}$
(B) $+105 \mathrm{~J} / \mathrm{K}$
(C) $-105 \mathrm{~J} / \mathrm{K}$
(D) $-210 \mathrm{~J} / \mathrm{K}$
8. The normal melting point of mercury is $-39^{\circ} \mathrm{C}$. The enthalpy of fusion of mercury is $2.3 \mathrm{~kJ} / \mathrm{mol}$. What is the entropy change of the surroundings when 4 (four) moles of liquid mercury are crystallized to the solid reversibly at $-39^{\circ} \mathrm{C}$ ?
(A) $+9.8 \mathrm{~J} / \mathrm{K}$
(B) $-39.3 \mathrm{~J} / \mathrm{K}$
(C) $-9.8 \mathrm{~J} / \mathrm{K}$
(D) $+39.3 \mathrm{~J} / \mathrm{K}$
9. The change in the Gibbs energy (in $\mathbf{k J}$ ) when the volume of $\mathbf{2 . 5}$ moles of ethane gas $\left[\mathrm{C}_{2} \mathrm{H}_{6}(\mathrm{~g})\right]$ is increased from isothermally 0.50 Liters to 10 Liters at $300^{\circ} \mathrm{C}$ is:
(A) +35.7 kJ
(B) +14.3 kJ
(C) -35.7 kJ
(D) -18.7 kJ
10. The density of liquid toluene, $\mathrm{C}_{7} \mathrm{H}_{8}(\mathrm{l})[\mathrm{M}=92]$, is $0.90 \mathrm{~g} / \mathrm{mL}$ at $50^{\circ} \mathrm{C}$. Therefore, the change in the Gibbs energy [in J] of one(1) mole of liquid toluene when the pressure is increased isothermally from 100 kPa to $5,000 \mathrm{kPa}$ at $50^{\circ} \mathrm{C}$ is approximately
(A) $1.6 \times 10^{3} \mathrm{~J}$
(B) $5.0 \times 10^{2} \mathrm{~J}$
(C) $1.1 \times 10^{4} \mathrm{~J}$
(D) $5.0 \times 10^{5} \mathrm{~J}$
11. A solid has two crystalline forms, $A(s)$ and $B(s)$. For the transition $A(s) \rightarrow B(s)$, $\Delta \mathrm{G}^{\circ}=-9.0 \mathrm{~kJ} / \mathrm{mol}$ (i.e. at 1 bar pressure). The difference in molar volumes of the two forms is $V_{m}(B)-V_{m}(A)=\Delta V_{m}=+2.0 \times 10^{-2} \mathrm{~L} / \mathrm{mol}$. This transition will be spontaneous at pressures $\qquad$ a pressure of $\qquad$ bar.
(A) above, $4.5 \times 10^{5}$ bar
(B) above , 4500 bar
(C) below , 4500 bar
(D) Spontaneous at all pressures

There are Four more MC questions on the following page

## MULTIPLE CHOICE QUESTIONS (Continued)

For \#12-\#15, consider the phase diagram to the right

12. The Critical Point of the substance is represented by:
(A) Point A
(B) Point B
(C) Point C
(D) Point D
13. The slope of curve $A-D$ is greater than the slope of curve $A-C$ because
(A) $\Delta_{\text {sub }} \mathrm{V}>\Delta_{\text {vap }} \mathrm{V}$
(B) $\Delta_{\text {vap }} S>\Delta_{\text {sub }} S$
(C) $\Delta_{\text {sub }} S>\Delta_{\mathrm{vap}} S$
(D) $\Delta_{\text {sub }} V<\Delta_{\text {vap }} V$
14. The slope of curve $A-B$ is much greater than the slope of curve $A-C$ because
(A) $\Delta$ fus $V<0$
(B) $\Delta$ fus $V \ll \Delta$ vap $V$
(C) $\Delta_{\text {fus }} S \ll \Delta_{\text {vap }} S$
(D) $\Delta$ fus $S \gg \Delta$ vap $S$
15. If the pressure on this substance is decreased from 2000. bar to 1. bar, the melting point temperature will $\qquad$ and the boiling point temperature will $\qquad$
(A) Decrease, Decrease
(B) Increase, Decrease
(C) Decrease, Increase
(D) Increase, Increase
(12) 1. A Perfect Gas has a temperature dependent molar constant pressure heat capacity, $C_{p, m}=a+b T^{3}$ with $\mathrm{a}=30 \mathrm{~J} / \mathrm{mol}-\mathrm{K}$ and $\mathrm{b}=8 \times 10^{-8} \mathrm{~J} /\left(\mathrm{mol}-\mathrm{K}^{4}\right)$.
Two (2) moles of this gas, originally at a temperature, of $250^{\circ} \mathrm{C}$ and volume of 30 L is heated reversibly at constant pressure to a temperature of $600^{\circ} \mathrm{C}$.

Calculate the Entropy change, $\Delta \mathrm{S}$, for this process (in $\mathrm{J} / \mathrm{K}$ )
(15) 2. Consider a hypothetical gas that obeys the equation of state:
$\mathrm{p}(\mathrm{V}-\mathrm{Ap})=\mathrm{nRT} \quad$ Note: $\mathbf{A}$ is an arbitrary constant, and NOT the Helmholtz Energy.
If this gas undergoes an isothermal compression from $p_{1}$ to $p_{2}$, develop INTEGRATED expressions for $\Delta \mathrm{U}, \Delta \mathrm{H}$ and $\Delta \mathrm{A}$ in terms of $\mathrm{n}, \mathrm{R}, \mathrm{T}, \mathrm{A}, \mathrm{p}_{1}$ and $\mathrm{p}_{2}$.
(16) 3. The normal boiling point of Bromine liquid, $\mathrm{Br}_{2}\left(\right.$ liq) is $59{ }^{\circ} \mathrm{C}$.

The Enthalpy of Vaporization is $29.5 \mathrm{~kJ} / \mathrm{mol}$ at $59^{\circ} \mathrm{C}$.
The constant pressure molar heat capacity of bromine liquid, $\mathrm{Br}_{2}(\mathrm{liq})$ is $76 . \mathrm{J} / \mathrm{mol}-\mathrm{K}$ The constant pressure molar heat capacity of bromine gas, $\mathrm{Br}_{2}$ (gas), is $36 . \mathrm{J} / \mathrm{mol}-\mathrm{K}$.

Calculate the entropy change of the Surroundings, $\Delta \mathbf{S}_{\text {surr }}$, (in $\mathrm{J} / \mathrm{mol}-\mathrm{K}$ ) when one (1) mole of supercooled $\mathrm{Br}_{2}$ (gas) condenses to $\mathrm{Br}_{2}\left(\right.$ liq) at $35^{\circ} \mathrm{C}$.
(12) 4 The densities of solid and liquid copper [Cu, $\mathrm{M}=63.5$ ] are $8.9 \mathrm{~g} / \mathrm{mL}$ and $8.0 \mathrm{~g} / \mathrm{mL}$, respectively. The normal melting point of copper is $1085^{\circ} \mathrm{C}$. The enthalpy of fusion of copper is $13.3 \mathrm{~kJ} / \mathrm{mol}$.

What pressure must be applied (in bar) to increase melting point of Cu to $1100^{\circ} \mathrm{C}$ ?

