This print-out should have 25 questions. Multiple-choice questions may continue on the next column or page - find all choices before answering.

Please refer to the cover page for needed thermodynamic values and formulas.

## $001 \quad 10.0$ points

An isolated system allows for the flow of...?

1. none of these
2. sound waves
3. kinetic energy
4. matter
5. heat

## 00210.0 points

Which of the following is true of a general thermodynamic state function?

1. The change in the value of a state function is always negative for a spontaneous reaction.
2. The value of the state function remains constant.
3. The value of a state function does NOT change with a change in temperature of a process.
4. The change of the value of a state function is independent of the path of a process.
5. The change in the value of the state function is always positive for endothermic processes.

## 00310.0 points

Consider the reaction
$\mathrm{C}_{6} \mathrm{H}_{14}(\ell)+9.5 \mathrm{O}_{2}(\mathrm{~g}) \rightarrow 6 \mathrm{CO}_{2}(\mathrm{~g})+7 \mathrm{H}_{2} \mathrm{O}(\ell)$
at constant pressure. Which response is true?
2. Work is done by the system as the reaction occurs.
3. Work is done on the system as the reaction occurs.
4. Work may be done on or by the system as the reaction occurs, depending upon the temperature.

## $004 \quad 10.0$ points

Consider the following specific heats: copper, $0.384 \mathrm{~J} / \mathrm{g} \cdot{ }^{\circ} \mathrm{C}$; lead, $0.159 \mathrm{~J} / \mathrm{g} \cdot{ }^{\circ} \mathrm{C}$; water, $4.18 \mathrm{~J} / \mathrm{g} \cdot{ }^{\circ} \mathrm{C}$; glass, $0.502 \mathrm{~J} / \mathrm{g} \cdot{ }^{\circ} \mathrm{C}$. If the same amount of heat is added to identical masses of each of these substances, which substance attains the highest temperature? (Assume that they all have the same initial temperature.)

## 1. copper

2. water
3. lead
4. glass

## $005 \quad 10.0$ points

Which of the following statements is/are true concerning the first law of thermodynamics?
I) The internal energy $(U)$ of the universe is conserved.
II) The internal energy of a system plus that of its surroundings is conserved.
III) The change in internal energy $(\Delta U)$ of a system and its surroundings can have the same sign.

1. I, II
2. III only
3. II, III
4. I only
5. I, III
6. No work is done as the reaction occurs.
7. II only
8. I, II, III

## $006 \quad 10.0$ points

1.95 mol of an ideal gas at 300 K and 3.00 atm expands from 16 L to 28 L and a final pressure of 1.20 atm in two steps:
(1) the gas is cooled at constant volume until its pressure has fallen to 1.20 atm , and
(2) it is heated and allowed to expand against a constant pressure of 1.20 atm until its volume reaches 28 L .

Which of the following is CORRECT?

1. $w=0$ for the overall process
2. $w=-6.03 \mathrm{~kJ}$ for the overall process
3. $w=-4.57 \mathrm{~kJ}$ for (1) and $w=-1.46 \mathrm{~kJ}$ for (2)
4. $w=0$ for (1) and $w=-1.46 \mathrm{~kJ}$ for (2)
5. $w=-4.57 \mathrm{~kJ}$ for the overall process

## $007 \quad 10.0$ points

A CD player and its battery together do 500 kJ of work, and the battery also releases 250 kJ of energy as heat and the CD player releases 50 kJ as heat due to friction from spinning. What is the change in internal energy of the system, with the system regarded as the battery and CD player together?

1. -800 kJ
2. +200 kJ
3. -200 kJ
4. -700 kJ
5. -750 kJ

## $008 \quad 10.0$ points

The specific heat of liquid water is 4.184 $\mathrm{J} / \mathrm{g} \cdot{ }^{\circ} \mathrm{C}$, and of steam $2.03 \mathrm{~J} / \mathrm{g} \cdot{ }^{\circ} \mathrm{C}$. The heat of vaporization of water $(\ell)$ is $2.26 \mathrm{~kJ} / \mathrm{g}$ and
its boiling point is $100^{\circ} \mathrm{C}$. What is the total heat flow when 18 grams of water at $12^{\circ} \mathrm{C}$ are heated to become steam at $109^{\circ} \mathrm{C}$ ?

1. 44.4 kJ
2. 47.6 kJ
3. under 28 kJ
4. 48.9 kJ
5. over 55 kJ
6. 31.7 kJ
7. 40.7 kJ

## $009 \quad 10.0$ points

Refer to the potential energy diagram shown below.


What is the change in enthalpy $(\Delta H)$ for the reaction

$$
\mathrm{A} \rightarrow \mathrm{~B} ?
$$

1. +50 kJ , endothermic
2. -50 kJ , exothermic
3. +50 kJ , exothermic
4. -250 kJ , endothermic
5. -150 kJ , endothermic
6. -50 kJ , endothermic
7. +300 kJ , exothermic
8. -150 kJ , exothermic
9. +300 kJ , endothermic
10. -250 kJ , exothermic

## $010 \quad 10.0$ points

Which of the following is NOT a feature of the bomb calorimetry apparatus used to measure the internal energy of a reaction?

1. The heat capacity of the calorimeter should be known to accurately correct for any heat lost to it.
2. The thermometer is inserted directly into the reaction vessel to measure $\Delta T$ of the reaction.
3. Large quantities of water surrounding the reaction vessel absorb the majority of the heat loss.
4. The volume of the reaction vessel is held constant to eliminate energy released as work.
5. The large heat capacity of water is beneficial in measuring heat released by combustion reactions.

## $011 \quad 10.0$ points

A 1.00 g sample of $n$-hexane $\left(\mathrm{C}_{6} \mathrm{H}_{14}\right)$ undergoes complete combustion with excess $\mathrm{O}_{2}$ in a bomb calorimeter. The temperature of the 1502 g of water surrounding the bomb rises from $22.64^{\circ} \mathrm{C}$ to $29.30^{\circ} \mathrm{C}$. The heat capacity of the hardware component of the calorimeter (everything that is not water) is $4042 \mathrm{~J} /{ }^{\circ} \mathrm{C}$. What is $\Delta U$ for the combustion of $n-\mathrm{C}_{6} \mathrm{H}_{14}$ ? One mole of $n-\mathrm{C}_{6} \mathrm{H}_{14}$ is 86.1 g . The specific heat of water is $4.184 \mathrm{~J} / \mathrm{g} \cdot{ }^{\circ} \mathrm{C}$.

1. $-1.15 \times 10^{4} \mathrm{~kJ} / \mathrm{mol}$
2. $-4.52 \times 10^{3} \mathrm{~kJ} / \mathrm{mol}$
3. $-7.40 \times 10^{4} \mathrm{~kJ} / \mathrm{mol}$
4. $-5.92 \times 10^{3} \mathrm{~kJ} / \mathrm{mol}$
5. $-9.96 \times 10^{3} \mathrm{~kJ} / \mathrm{mol}$

## $012 \quad 10.0$ points

Which of the reactions below is a formation reaction?

$$
\begin{aligned}
& \text { 1. } \mathrm{H}_{2}(\mathrm{~g})+\mathrm{C}_{\text {graphite }}(\mathrm{s})+\frac{3}{2} \mathrm{O}_{2}(\mathrm{~g}) \\
& \quad \rightarrow \mathrm{CO}_{2}(\mathrm{~g})+\mathrm{H}_{2} \mathrm{O}(\mathrm{~g})
\end{aligned}
$$

2. $2 \mathrm{H}_{2}(\mathrm{~g})+2 \mathrm{C}_{\text {graphite }}(\mathrm{s})+\mathrm{O}_{2}(\mathrm{~g})$
$\rightarrow 2 \mathrm{CH}_{2} \mathrm{O}(\ell)$
3. $2 \mathrm{Fe}(\mathrm{s})+\frac{3}{2} \mathrm{O}_{2}(\mathrm{~g}) \rightarrow \mathrm{Fe}_{2} \mathrm{O}_{3}(\mathrm{~s})$
4. $\mathrm{N}_{2}(\ell)+2 \mathrm{H}_{2}(\mathrm{~g}) \rightarrow \mathrm{N}_{2} \mathrm{H}_{4}(\ell)$

## $013 \quad 10.0$ points

Calculate the standard reaction enthalpy for the reaction

$$
\mathrm{N}_{2} \mathrm{H}_{4}(\ell)+\mathrm{H}_{2}(\mathrm{~g}) \rightarrow 2 \mathrm{NH}_{3}(\mathrm{~g})
$$

given

$$
\begin{aligned}
\mathrm{N}_{2} \mathrm{H}_{4}(\ell)+\mathrm{O}_{2}(\mathrm{~g}) \rightarrow & \mathrm{N}_{2}(\mathrm{~g})+2 \mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \\
& \Delta H^{\circ}=-543 \mathrm{~kJ} \cdot \mathrm{~mol}^{-1} \\
2 \mathrm{H}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightarrow & 2 \mathrm{H}_{2} \mathrm{O}(\mathrm{~g}) \\
& \Delta H^{\circ}=-484 \mathrm{~kJ} \cdot \mathrm{~mol}^{-1} \\
\mathrm{~N}_{2}(\mathrm{~g})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightarrow & 2 \mathrm{NH}_{3}(\mathrm{~g}) \\
& \Delta H^{\circ}=-92.2 \mathrm{~kJ} \cdot \mathrm{~mol}^{-1}
\end{aligned}
$$

1. $-1119 \mathrm{~kJ} \cdot \mathrm{~mol}^{-1}$
2. $-151 \mathrm{~kJ} \cdot \mathrm{~mol}^{-1}$
3. $-935 \mathrm{~kJ} \cdot \mathrm{~mol}^{-1}$
4. $-243 \mathrm{~kJ} \cdot \mathrm{~mol}^{-1}$
5. $-59 \mathrm{~kJ} \cdot \mathrm{~mol}^{-1}$

## $014 \quad 10.0$ points

Estimate the heat released when ethene $\left(\mathrm{CH}_{2}=\mathrm{CH}_{2}\right)$ reacts with HBr to give $\mathrm{CH}_{3} \mathrm{CH}_{2} \mathrm{Br}$.

1. $200 \mathrm{~kJ} / \mathrm{mol}$
2. $76 \mathrm{~kJ} / \mathrm{mol}$
3. $1036 \mathrm{~kJ} / \mathrm{mol}$
4. $424 \mathrm{~kJ} / \mathrm{mol}$
5. $470 \mathrm{~kJ} / \mathrm{mol}$

## $015 \quad 10.0$ points

Consider a reaction that is non-spontaneous at all temperatures. What would be the signs of $\Delta G_{\text {sys }}, \Delta H_{\text {surr }}$, and $\Delta S_{\text {univ }}$ respectively for such a reaction?
1.,,+-+
2.,,-+-
3.,,+--
4.,,-++
5.,,+++

## $016 \quad 10.0$ points

Which one shows the substances in the decreasing order of their molar entropy?

1. $\mathrm{C}(\mathrm{s}), \mathrm{H}_{2} \mathrm{O}(\ell), \mathrm{H}_{2} \mathrm{O}(\mathrm{g}), \mathrm{H}_{2} \mathrm{O}(\mathrm{s})$
2. $\mathrm{H}_{2} \mathrm{O}(\mathrm{g}), \mathrm{H}_{2} \mathrm{O}(\ell), \mathrm{H}_{2} \mathrm{O}(\mathrm{s}), \mathrm{C}(\mathrm{s})$
3. None of these
4. $\mathrm{H}_{2} \mathrm{O}(\mathrm{s}), \mathrm{H}_{2} \mathrm{O}(\ell), \mathrm{H}_{2} \mathrm{O}(\mathrm{g}), \mathrm{C}(\mathrm{s})$
5. $\mathrm{C}(\mathrm{s}), \mathrm{H}_{2} \mathrm{O}(\mathrm{g}), \mathrm{H}_{2} \mathrm{O}(\ell), \mathrm{H}_{2} \mathrm{O}(\mathrm{s})$
6. $\mathrm{C}(\mathrm{s}), \mathrm{H}_{2} \mathrm{O}(\mathrm{s}), \mathrm{H}_{2} \mathrm{O}(\ell), \mathrm{H}_{2} \mathrm{O}(\mathrm{g})$
$017 \quad 10.0$ points
Consider the following processes. (Treat all gases as ideal.)
I) The pressure of one mole of oxygen gas is allowed to double isothermally.
II) Carbon dioxide is allowed to expand isothermally to 10 times its original volume.
III) The temperature of one mole of helium
is increased $25^{\circ} \mathrm{C}$ at constant pressure.
IV) Nitrogen gas is compressed isothermally to one half its original volume.
V) A glass of water loses 100 J of energy reversibly at $30^{\circ} \mathrm{C}$.
Which of these processes leads to an increase in entropy?

## 1. III and V

2. I and II
3. I and IV
4. V
5. II and III

## $018 \quad 10.0$ points

What is the entropy at $T=0 \mathrm{~K}$ for one mole of chloroform $\left(\mathrm{CHCl}_{3}\right)$ ?

1. $1.9 \times 10^{-23} \mathrm{~J} \mathrm{~K}^{-1}$
2. $1.38 \mathrm{~J} \mathrm{~K}^{-1}$
3. $11.5 \mathrm{~J} \mathrm{~K}^{-1}$
4. $-11.5 \mathrm{~J} \mathrm{~K}^{-1}$
5. $0 \mathrm{~J} \mathrm{~K}^{-1}$

## $019 \quad 10.0$ points

What is the total non-vibrational internal energy of 10 nitrous oxide $\left(\mathrm{N}_{2} \mathrm{O}\right)$ molecules?

1. 15 k T
2. 10 k T
3. 25 kT
4. 10 R T
5. 25 R T
6. 15 R T

Advertising claims sometimes state that adding something mechanical to a car's engine will allow it to recover $100 \%$ of the energy that comes from burning gasoline. You should be skeptical of such claims because they violate the

1. first law of thermodynamics.
2. activation energy requirements of all chemical reactions.
3. law of conservation of matter.
4. second law of thermodynamics.

## $021 \quad 10.0$ points

Calculate $\Delta S_{\text {surr }}^{\circ}$ at 298 K for the reaction

$$
\begin{aligned}
& \quad 6 \mathrm{C}(\mathrm{~s})+3 \mathrm{H}_{2}(\mathrm{~g}) \rightarrow \mathrm{C}_{6} \mathrm{H}_{6}(\ell) \\
& \Delta H_{\mathrm{r}}^{\circ}=+49.0 \mathrm{~kJ} \cdot \mathrm{~mol}^{-1} \text { and } \Delta S_{\mathrm{r}}^{\circ}=-253 \\
& \mathrm{~J} \cdot \mathrm{~K}^{-1} \cdot \mathrm{~mol}^{-1} .
\end{aligned}
$$

1. $-417 \mathrm{~J} \cdot \mathrm{~K}^{-1} \cdot \mathrm{~mol}^{-1}$
2. $-164 \mathrm{~J} \cdot \mathrm{~K}^{-1} \cdot \mathrm{~mol}^{-1}$
3. $+253 \mathrm{~J} \cdot \mathrm{~K}^{-1} \cdot \mathrm{~mol}^{-1}$
4. $-253 \mathrm{~J} \cdot \mathrm{~K}^{-1} \cdot \mathrm{~mol}^{-1}$
5. $+164 \mathrm{~J} \cdot \mathrm{~K}^{-1} \cdot \mathrm{~mol}^{-1}$
$022 \quad 10.0$ points
For the evaporation of water from an open pan at $25^{\circ} \mathrm{C}$, the values of $\Delta S$ for the water, the surroundings, and the universe must be, respectively,
6. positive, positive, positive.
7. positive, negative, zero.
8. None of these is correct.
9. negative, negative, negative.
10. positive, negative, positive.

## $023 \quad 10.0$ points

For the reaction

$$
2 \mathrm{SO}_{3}(\mathrm{~g}) \rightarrow 2 \mathrm{SO}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g})
$$

$\Delta H_{\mathrm{r}}^{\circ}=+198 \mathrm{~kJ} \cdot \mathrm{~mol}^{-1}$ at 298 K . Which statement is true for this reaction?

1. The reaction is driven by the enthalpy.
2. The reaction will not be spontaneous at any temperature.
3. The reaction will not be spontaneous at high temperatures.
4. $\Delta G_{\mathrm{r}}^{\circ}$ will be positive at high temperatures.
5. $\Delta G_{\mathrm{r}}^{\circ}$ will be negative at high temperatures.

## $024 \quad 10.0$ points

What is $\Delta G_{\mathrm{r}}^{\circ}$ for the combustion of liquid $n$-pentane?

1. $3389 \mathrm{~kJ} / \mathrm{mol}$
2. $-383 \mathrm{~kJ} / \mathrm{mol}$
3. $383 \mathrm{~kJ} / \mathrm{mol}$
4. $-3389 \mathrm{~kJ} / \mathrm{mol}$
5. $-451 \mathrm{~kJ} / \mathrm{mol}$
6. $451 \mathrm{~kJ} / \mathrm{mol}$

## $025 \quad 10.0$ points

Consider the date below regarding different allotropes of carbon.

|  | $\Delta G_{\mathrm{f}}^{\circ}$ <br> carbon allotrope |
| :--- | ---: |
| $\mathrm{CJJ} \cdot \mathrm{mol}^{-1}$ |  |

C(s, diamond)
2.9
$\mathrm{C}_{60}$ (s, buckminsterfullerene) 24
Which of the following statements is supported by these data?

1. $\mathrm{C}_{60}$ is thermodynamically more stable than graphite under standard conditions.
2. Graphite could spontaneously form $\mathrm{C}_{60}$ under standard conditions.
3. Formation of graphite from $\mathrm{C}_{60}$ would be exergonic under standard conditions.
4. Diamond is the least thermodynamically stable allotrope of carbon under standard conditions.
