$$
\begin{gathered}
\text { ipt Instituto Politécnico de Tomar } \\
\text { Frequency of CHEMICAL THERMODYNAMICS }- \text { January } 8,2014 \\
\mathrm{R}=8.314 \mathrm{~J} \cdot \mathrm{~K}^{-1} \cdot \mathrm{~mol}^{-1}=0.082 \mathrm{~atm} \text { L. } \mathrm{mol}^{-1} \cdot \mathrm{~K}^{-1} ; 1 \mathrm{~atm}=101325 \mathrm{~Pa}=760 \mathrm{mmHg} \\
\text { Maximum time: } 2 \mathrm{H} 30 \mathrm{~m}
\end{gathered}
$$

1. Two moles of oxygen (considered as an ideal gas) are compressed adiabatically with application of constant exterior pressure of 10 atm , since the initial temperature of $26.85^{\circ} \mathrm{C}$ and 1 atm , until a final equilibrium state. Considering $\mathrm{Cp}=7 / 2 \mathrm{R}$ :
1.1. Calculate the final temperature of the gas.
1.2. Calculate the work, $w$, heat, $Q$, internal energy change, $\Delta \mathrm{U}$ and enthalpy change, $\Delta \mathrm{H}$ (If you do not solved the preceding paragraph consider the final temperature equal to 1071 K ).
2. The reaction of nitrogen monoxide oxidation is important in the process of formation of smoke pollutants

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\mathrm{NO}(\mathrm{~g})+1 / 2 \mathrm{O}_{2}(\mathrm{~g}) \rightarrow \mathrm{NO}_{2}(\mathrm{~g})
$$

Calculate for this reaction:
2.1. The standard enthalpy of reaction, $\Delta H^{o}{ }_{r}$ at 298 K .
2.2. The enthalpy of reaction at 200 K .

| Compound | $\Delta H_{f}^{o}$ at $298 \mathrm{~K} / \mathrm{kJ} . \mathrm{mol}^{-1}$ | $C_{p} / \mathrm{J} \cdot \mathrm{mol}^{-1} \cdot \mathrm{~K}^{-1}$ |
| :--- | :---: | :---: |
| $\mathrm{NO}(\mathrm{g})$ | 90.29 | 30.3 |
| $\mathrm{O}_{2}(\mathrm{~g})$ | ---- | 30.0 |
| $\mathrm{NO}_{2}(\mathrm{~g})$ | 33.10 | 39.4 |

3. Entropy is the fundamental property associated with the $2^{\text {nd }}$ law of Thermodynamics 3.1. State the second Law.
3.2. Calculate the entropy change when 1 mol of water vapor is heated from $200{ }^{\circ} \mathrm{C}$ until $400{ }^{\circ} \mathrm{C}$ at 1 atm . The molar heat capacity of water is given by the equation: $C_{p} / \mathrm{J} \cdot \mathrm{K}^{-1} \cdot \mathrm{~mol}^{-1}=30.51+1.03 \times 10^{-2} \mathrm{~T}$
4. Using the Trouton rule $\left(\Delta S_{\text {vap }}=85 \mathrm{~J} \cdot \mathrm{~K}^{-1} \cdot \mathrm{~mol}^{-1}\right.$ at the normal boiling temperature) calculate the vapor pressure of dipropylic ether at $33^{\circ} \mathrm{C}$, knowing that the normal boiling temperature is $89.5^{\circ} \mathrm{C}$.
5. It was found that a solution of benzene and toluene at 293 K showed a total vapor pressure of 46 mmHg . At that temperature, vapor pressures of pure benzene and toluene are equal to 74.7 mmHg and 22.3 mmHg .
5.1. Determine the mole fraction of benzene in solution and in the vapor phase in equilibrium with it.
5.2. Calculate the $\Delta \mathrm{G}_{\text {mist }}$ assuming the ideality in liquid phase.
