# Solution to Assignment questions <br> JIF 314 Thermodynamics 

## Based on the text book <br> Heat and thermodynamics by Zemansky and Dittman, $7^{\text {th }}$ edition, Mcgraw-Hill.

## Chapter 1

Problem 1.1. Solve using Excel.
First, calculate the value $\theta$ of the gas: $\theta=273.1 \mathrm{~K}\left(\frac{P}{P_{T P}}\right)$.

| $\mathrm{P}_{\text {TP }}(\mathrm{kPa})$ | $\mathrm{P}(\mathrm{kPa})$ | $\theta(\mathrm{K})$ |
| :---: | :---: | ---: |
| 33.331 | 51.19 | 419.5211785 |
| 66.661 | 102.37 | 419.4864944 |
| 99.992 | 153.54 | 419.4434195 |
| 133.32 | 204.69 | 419.390342 |


$\theta$ vs. $P_{\mathrm{TP}}$ is a straight line in the form of $y=m x+c$, where $y \equiv \theta, x \equiv P_{\mathrm{TP}}$. The value of $\theta$ when $P_{\mathrm{TP}}$ becomes zero is the value of the temperature of the gas. This value is simply the value of intersection, $c$, in the formula of the straight line in the form of $y=m x+c$.

From the formula of the straight line generated by Excell, the intersection of the straight line is $c=419.57$ in the graph of $\theta$ vs. $P_{\text {TP }}$.

Hence, the temperature of the gas in the bulb is $\theta=419.57 \mathrm{~K}$.

## Problem 1.3.

(a) The temperature with resistance measured to be $1000 \Omega$ can be calculated using the relationship between $R^{\prime}$ and $T$, as per
$\sqrt{\frac{\log R^{\prime}}{T}}=a+b \log R^{\prime}, a=-1.16, b=0.675$.

Setting $R^{\prime}=1000 \Omega$,
$\left(\sqrt{\frac{\log R^{\prime}}{T}}=a+b \log R^{\prime}\right)^{2}=\frac{\log R^{\prime}}{T}=a^{2}+b^{2}\left(\log R^{\prime}\right)^{2}+2 a b \log R^{\prime}$

$$
\begin{aligned}
& T=\frac{\log R^{\prime}}{a^{2}+b^{2}\left(\log R^{\prime}\right)^{2}+2 a b \log R^{\prime}} \\
& =\frac{\log (1000)}{(-1.16)^{2}+(0.675)^{2}[\log (1000)]^{2}+2(-1.16)(0.675) \log R^{\prime}} \\
& =\frac{0.6908}{(-1.16)^{2}+(0.675)^{2}[0.6908]^{2}+2(-1.16)(0.675)(0.6908)}=1.44
\end{aligned}
$$

Hence, the temperature of the helium cryostat is 1.44 K .
(b) Use Excell. Plot $\log R^{\prime}$ vs. $\log T$ graph by forming the following table:

| $R^{\prime}$ | $\log R^{\prime}$ | $T=\log R^{\prime} /\left(a+b \log R^{\prime}\right)^{\wedge} 2$ | $l o g T$ |
| ---: | :--- | ---: | ---: |
| 1000 | 6.907755 | 0.563018189 | 0.57444 |
| 5000 | 8.517193 | 0.404427271 | 0.90528 |
|  |  | - |  |
| 10000 | 9.21034 | 0.360158153 | 1.02121 |
| 15000 | 9.615805 |  | - |
|  |  | 0.338393713 | 1.08355 |
| 20000 | 9.903488 | 0.32444907 | 1.12563 |
|  |  |  | - |
| 25000 | 10.12663 | 0.31438398 | 1.15714 |
| 30000 | 10.30895 | 0.306603264 | -1.1822 |



Problem 1.9: $\theta\left({ }^{\circ} \mathrm{F}\right)=\frac{9}{5} \theta\left({ }^{\circ} \mathrm{C}\right)+32=\frac{9}{5}(99.974)+32=211.95^{\circ} \mathrm{F}(5$ significant figures $)$.

## Chapter 2

Problem 2.1
(a) Given the equation of state for a ideal gas $P V=n R T$, show that $\beta=\frac{1}{T}$.

## Solution:

Given equation of state for a ideal gas

$$
P V=n R T, \quad \text { Eq. (1) }
$$

and the definition of volume expansivity $\beta=\frac{1}{V}\left(\frac{\partial V}{\partial T}\right)$, it is easily verified that $\beta=1 / T$ by taking the partial derivate of Eq. (1) with respect to $T$ :

$$
\begin{equation*}
\frac{\partial}{\partial T}(P V=n R T) \rightarrow P \frac{\partial V}{\partial T}=n R \tag{2}
\end{equation*}
$$

Inserting $P V=n R T$ into Eq. (2), we arrive at

$$
\frac{\partial V}{\partial T}=\frac{n R}{P}=\frac{P V}{T} \frac{1}{P}=\frac{V}{T}
$$

Hence, $\beta=\frac{1}{V}\left(\frac{\partial V}{\partial T}\right)=\beta=\frac{1}{V}\left(\frac{V}{T}\right)=\frac{1}{T}$.
(b) Show that the isothermal compressivility $\kappa=1 / P$.

## Solution

Given equation of state for a ideal gas

$$
P V=n R T, \quad \text { Eq. (1) }
$$

and the definition of isothermal compressibility $\kappa=\frac{1}{B}=-\frac{1}{V}\left(\frac{\partial V}{\partial P}\right)$, it is easily verified that $\beta=1 / P$ by taking the partial derivate of Eq. (1) with respect to $P$ :

$$
\begin{equation*}
\frac{\partial}{\partial P}(P V=n R T) \rightarrow P \frac{\partial V}{\partial P}+V=\frac{\partial}{\partial P}(n R T)=0 \tag{2}
\end{equation*}
$$

Inserting $P V=n R T$ into Eq. (2), we arrive at

$$
\frac{\partial V}{\partial P}=-\frac{V}{P}
$$

Hence, $\kappa=-\frac{1}{V}\left(\frac{\partial V}{\partial P}\right)=-\frac{1}{V}\left(-\frac{V}{P}\right)=\frac{1}{P}$.

Problem 2.2: Given the equation of state of a van der Waals gas, $\left(P+\frac{a}{v^{2}}\right)(v-b)=R T$, calculate
(a) $\left(\frac{\partial P}{\partial v}\right)_{T}$, (b) $\left(\frac{\partial P}{\partial T}\right)_{v}$.

## Solution:

(a) Taking the partial derivative with respect to $v$, with constant $T$,
$\frac{\partial}{\partial v}\left[\left(P+\frac{a}{v^{2}}\right)(v-b)\right]_{T}=\left.\frac{\partial}{\partial v}(R T)\right|_{T}=0$
$\left.(v-b) \frac{\partial}{\partial v}\left(P+\frac{a}{v^{2}}\right)\right|_{T}+\left.\left(P+\frac{a}{v^{2}}\right) \frac{\partial}{\partial v}(v-b)\right|_{T}=0$
$(v-b)\left(\left.\frac{\partial P}{\partial v}\right|_{T}-\frac{2 a}{v^{3}}\right)+\left(P+\frac{a}{v^{2}}\right)=0$
$\left.\frac{\partial P}{\partial v}\right|_{T}=-\frac{P+\frac{a}{v^{2}}}{v-b}+\frac{2 a}{v^{3}}$
(b) Taking the partial derivative with respect to $T$, with constant $v$,

$$
\begin{aligned}
& \left.\frac{\partial}{\partial T}\left[\left(P+\frac{a}{v^{2}}\right)(v-b)\right]\right|_{v}=\left.\frac{\partial}{\partial T}(R T)\right|_{v} \\
& \left.(v-b) \frac{\partial}{\partial T}\left(P+\frac{a}{v^{2}}\right)\right|_{v}+\left.\left(P+\frac{a}{v^{2}}\right) \frac{\partial}{\partial T}(v-b)\right|_{v}=R \\
& (v-b)\left[\left.\frac{\partial P}{\partial T}\right|_{v}+\left.a \frac{\partial}{\partial T}\left(\frac{1}{v^{2}}\right)\right|_{v}\right]+\left.\left(P+\frac{a}{v^{2}}\right) \frac{\partial v}{\partial T}\right|_{v}=R \\
& (v-b)\left(\left.\frac{\partial P}{\partial T}\right|_{v}+0\right)+\left(P+\frac{a}{v^{2}}\right) \cdot 0=R \\
& \left.\frac{\partial P}{\partial T}\right|_{v}=\frac{R}{v-b}
\end{aligned}
$$

(c)

$$
\begin{aligned}
& \left(\frac{\partial P}{\partial v}\right)_{T}\left(\frac{\partial v}{\partial T}\right)_{P}=-\left(\frac{\partial P}{\partial T}\right)_{v} \\
& \rightarrow\left(\frac{\partial v}{\partial T}\right)_{P}=\frac{-\left(\frac{\partial P}{\partial T}\right)_{V}}{\left(\frac{\partial P}{\partial v}\right)_{T}}=-\frac{\frac{R}{v-b}}{-\frac{P+\frac{a}{v^{2}}}{v-b}+\frac{2 a}{v^{3}}}=\frac{R}{P}\left(\frac{1}{1+\frac{2 a b}{v^{3} P}-\frac{a}{v^{2} P}}\right)
\end{aligned}
$$

Problem 3.2
(a) Show that the work done by an ideal gas during the quasi-static, isothermal expansion from an initial pressure $P_{\mathrm{i}}$ to a final pressure $P_{\mathrm{f}}$, is given by $W=n R T \ln \left(P_{\mathrm{f}} / P_{\mathrm{i}}\right)$.

## Solution:

For isothermal process, $P_{i} V_{i}=P_{f} V_{f}$. Hence $V_{i} / V_{f}=P_{f} / P_{i}$. Substitute this into $W=-n R T \ln \left(V_{f} / V_{i}\right)$, we get $W=-n R T \ln \left(P_{i} / P_{f}\right)=n R T \ln \left(P_{f} / P_{i}\right)$.

Problem 3.3
An adiabatic chamber with rigid walls consists of two compartments, one containing a gas and the other evacuated; the partition between the two compartments is suddenly removed. Is the work done during an infinitesimal portion of this process (called an adiabatic expansion) equal $P \mathrm{~d} V$ ?

Answer: NO. Because there is no work done against the expansion of the gas-filled compartment by the evacuated compartment.

