Name of Paper
Name of the Course
Semester
Duration
Maximum Marks
: Thermal Physics
: B.Sc. (Hons.) Physics
: III
: 3 Hours
: 75

Attempt five questions in all.
Question No. 1 is compulsory.
Answer any four of the remaining six, attempting any two parts from each question.
Q. 1. Attempt all parts.
(a) Which of the two, an isothermal or an adiabatic, has greater slope? Prove mathematically.

Ans. For isothermal process
PV = constant

Differentiating this we get

$$
\begin{aligned}
\mathrm{P} d \mathrm{~V}+\mathrm{V} d \mathrm{P} & =\mathrm{Q} \\
\frac{d \mathrm{P}}{d \mathrm{~V}} & =\frac{-\mathrm{P}}{\mathrm{~V}}
\end{aligned}
$$

So, slope isothermal $=\frac{-\mathrm{P}}{\mathrm{V}}$
For adiabatic process

$$
P V^{r}=\text { constant }
$$

Differentiating this we get:

$$
(d \mathrm{P}) \mathrm{V}^{\mathrm{r}}+\mathrm{P} r \mathrm{~V}^{r-1} d \mathrm{~V}=0
$$

$$
\frac{d \mathrm{P}}{d \mathrm{~V}}=-r \frac{\mathrm{PV}^{r-1}}{\mathrm{~V}^{r}}=-r\left(\frac{\mathrm{P}}{\mathrm{~V}}\right)
$$

So, slope of adisbatic $=-r \frac{\mathrm{P}}{\mathrm{V}}$
Since, $r>1$ the slope of adisbatic curve is greater.
(b) A Carnot's engine whose sink is at $27^{\circ} \mathrm{C}$ has an efficiency of $50 \%$. By how much the temperature of the source be changed to decrease its efficiency to $40 \%$ ?

Ans. $\quad$ Efficiency $=n=1-\frac{\mathrm{T}_{2}}{\mathrm{~T}_{1}}$

$$
\begin{aligned}
\mathrm{T}_{1} & =\text { Temperature of source } \\
\mathrm{T}_{2} & =\text { Temperature of sink } \\
\mathrm{T}_{2} & =27^{\circ} \mathrm{C}=300 \mathrm{~K} \text { (given) } \\
n & =50 \%=\frac{50}{100}=\frac{1}{2}
\end{aligned}
$$

So,

So,

$$
\begin{aligned}
\frac{2}{5} & =1-\frac{300}{\mathrm{~T}_{1}} \\
\frac{300}{\mathrm{~T}_{1}} & =1-\frac{2}{5}=\frac{3}{5} \\
\mathrm{~T}_{1} & =\frac{300 \times 5}{3}=500
\end{aligned}
$$

$\mathrm{T}_{1}=500 \mathrm{~K} \rightarrow$ New temperature of source
So, Difference $=600-500=100 \mathrm{~K}$
(c) One kilogram of water is heated from $0^{\circ} \mathrm{C}$ to $100^{\circ} \mathrm{C}$ and converted into steam at the same temperature. Calculate the increase in entropy. Given that specific heat of water is $4.18 \times 103 \mathrm{Jkg}^{-1} \mathrm{~K}^{-1}$ and latent heat of vaporisation is $2.24 \times 106 \mathrm{Jkg}^{1}$.

Ans. Heating is carried out then entrops increase is given by

$$
\begin{aligned}
\mathrm{T}_{1} & =0^{\circ} \mathrm{C}=273 \mathrm{~K} \\
\mathrm{~T}_{2} & =100^{\circ} \mathrm{C}=373 \mathrm{~K} \\
\Delta \mathrm{~S}_{1} & =\mathrm{MC} \ln \left(\frac{\mathrm{~T}_{2}}{\mathrm{~T}_{1}}\right) \\
& =1 \times 4.18 \times 10^{3} \times 2.303 \times \log _{10}\left(\frac{373}{273}\right) \\
\Delta \mathrm{S}_{1} & =1305 \mathrm{JK}^{-1}
\end{aligned}
$$

Entrapy when water is converted into

$$
\begin{aligned}
& \text { Steam }=\Delta \mathrm{S}_{2} \\
& \Delta \mathrm{~S}_{2}=\frac{\delta \mathrm{Q}}{\mathrm{~T}}=\frac{m \mathrm{~L}}{\mathrm{~T}}
\end{aligned}
$$

$$
\Delta \mathrm{S}_{2}=\frac{1 \times 2.24 \times 10^{6}}{373}=6 \times 10^{3}
$$

$$
\Delta \mathrm{S}_{2}=6 \times 10^{3} \mathrm{JK}^{-1}
$$

Total at entropy $=\Delta \mathrm{S}_{1}+\Delta \mathrm{S}_{2}$

$$
\begin{align*}
& S=1305+6000 \\
& S=73.05 \mathrm{JK}^{-1} . \tag{4}
\end{align*}
$$

(d) Using Carnot's cycle derive Clausius-Clapeyron latent heat equation.

Ans. The solution to this questions is in this textbook.
(e) A substance has volume expansivity $=2 b \mathrm{~T} / \mathrm{V}$ and isothermal compressibility $=a / \mathrm{V}$, where ' $a$ ' and ' $b$ ' are constant. Find the equation of state.

Ans.

$$
\begin{equation*}
\text { Volume expansivity }=\frac{2 b \mathrm{~T}}{\mathrm{~V}}=\alpha \tag{3}
\end{equation*}
$$

$$
\begin{aligned}
\text { Isothermal compressibility } & =B_{T}=\frac{a}{V} \\
V & =V(P, T)
\end{aligned}
$$

$$
d \mathrm{~V}=\left(\frac{\partial \mathrm{V}}{\partial \mathrm{P}}\right)_{\mathrm{T}} d \mathrm{P}+\left(\frac{\partial \mathrm{V}}{\partial \mathrm{~T}}\right)_{\mathrm{P}} d \mathrm{~T}
$$

In terms of $\alpha$ and $B_{T}$

$$
\begin{aligned}
& \frac{d \mathrm{~V}}{\mathrm{~V}}=\mathrm{B}_{\mathrm{T}} d \mathrm{P}+\alpha d \mathrm{~T} \\
& \frac{d \mathrm{~V}}{\mathrm{~V}}=\frac{-a}{\mathrm{~V}} d \mathrm{P}+\frac{2 b \mathrm{~T}}{\mathrm{~V}} d \mathrm{~T}
\end{aligned}
$$

$$
d \mathrm{~V}+a d \mathrm{P}-2 b \mathrm{~T} d \mathrm{~T}=0
$$

Integrate both sides

$$
\begin{equation*}
\mathrm{V}+a \mathrm{P}-b \mathrm{~T}^{2}=\text { constant } \rightarrow \text { equation of state } \tag{2}
\end{equation*}
$$

(f) Define Boyle Temperature. Give relation between Boyle temperature, Temperature of inversion and Critical temperature.

Ans. The solution to this questions is in this textbook.
$(g)$ What is Brownian motion? Give its characteristics.
Ans. The solution to this questions is in this textbook.
Q. 2. (a) (i) State first law of thermodynamics. What are its physical significance and limitations? Write first law of thermodynamics for an adiabatic, isobaric and isochoric processes.
(ii) Derive the work done by an ideal gas in expanding adiabatically from initial state ( $\mathrm{P}_{i^{\prime}}, \mathrm{V}_{i}, \mathrm{~T}_{i}$ ) to the final state ( $\mathrm{P}_{f} \mathrm{~V}_{f}, \mathrm{~T}_{f}$ ) .

Ans. (i) The solution to this questions is in this textbook.
(ii) The solution to this questions is in this textbook.
(b) Using first law of thermodynamics, prove that
(i) $\left(\frac{\partial \mathrm{U}}{\partial \mathrm{P}}\right)_{\mathrm{V}}=\frac{\mathrm{C}_{\mathrm{V}} \mathrm{K}_{\mathrm{T}}}{\beta}$
(ii) $\left(\frac{\partial U}{\partial V}\right)_{P}=\frac{C_{P}}{\beta V}-P$
where $\beta$ and $K_{T}$ are volume expansion coefficient and isothermal compressibility respectively.

Ans. Write U in terms of V and P

$$
\begin{aligned}
\mathrm{U} & =\mathrm{U}(\mathrm{~V}, \mathrm{P}) \\
d \mathrm{U} & =\left(\frac{\partial \mathrm{U}}{\partial \mathrm{~V}}\right)_{\mathrm{P}} d \mathrm{~V}+\left(\frac{\partial \mathrm{U}}{\partial \mathrm{P}}\right)_{\mathrm{V}} d \mathrm{P}
\end{aligned}
$$

$$
\left(\frac{\partial \mathrm{U}}{\partial \mathrm{P}}\right)_{V}=\left(\frac{\partial \mathrm{U}}{\partial \mathrm{~V}}\right)_{\mathrm{P}}\left(\frac{\partial \mathrm{~V}}{\partial \mathrm{P}}\right)_{\mathrm{V}}+\left(\frac{\partial \mathrm{V}}{\partial \mathrm{P}}\right)_{\mathrm{V}} \frac{\partial \mathrm{P}}{\partial \mathrm{P}}
$$

So,

$$
\begin{align*}
\left(\frac{\partial U}{\partial P}\right)_{V} & =\left(\frac{\partial V}{\partial P}\right)_{V} \times \frac{\partial T}{\partial T} \\
\left(\frac{\partial U}{\partial P}\right)_{V} & =\left(\frac{\partial U}{\partial T}\right)_{V} \times\left(\frac{\partial T}{\partial P}\right)_{V}  \tag{a}\\
C_{V} & =\left(\frac{\partial U}{\partial T}\right)_{V} \text { and } \beta=\frac{1}{V}\left(\frac{\partial V}{\partial T}\right) \tag{1}
\end{align*}
$$

$$
\mathrm{K}_{\mathrm{T}}=+\frac{1}{\mathrm{~V}}\left(\frac{\partial \mathrm{~V}}{\partial \mathrm{P}}\right)
$$

$$
\begin{equation*}
\frac{\mathrm{K}_{\mathrm{T}}}{\beta}=\left(\frac{\partial \mathrm{T}}{\partial \mathrm{P}}\right)_{\mathrm{V}} \tag{3}
\end{equation*}
$$

Put (1) and (2) in (a)

$$
\left(\frac{\partial \mathrm{U}}{\partial \mathrm{P}}\right)_{\mathrm{V}}=\frac{\mathrm{C}_{\mathrm{V}} \mathrm{~K}_{\mathrm{T}}}{\beta}
$$

(ii) $\quad\left(\frac{\partial \mathrm{U}}{\partial \mathrm{V}}\right)_{\mathrm{P}}=\left(\frac{\partial \mathrm{U}}{\partial \mathrm{T}}\right)_{\mathrm{P}} \times\left(\frac{\partial \mathrm{T}}{\partial \mathrm{V}}\right)$

Now, from first law

$$
\partial \mathrm{U}=\partial \theta-\partial w
$$

So, $\quad\left(\frac{\partial \mathrm{U}}{\partial \mathrm{V}}\right)_{\mathrm{P}}=\left(\frac{\partial \theta-\partial w}{\partial \mathrm{~T}}\right)_{\mathrm{P}} \mathrm{V} \frac{\partial \mathrm{T}}{\partial \mathrm{V}}$

$$
\begin{aligned}
& =\left(\frac{\partial \theta}{\partial \mathrm{T}}\right)_{\mathrm{P}}\left(\frac{\partial \mathrm{~T}}{\partial \mathrm{~V}}\right)_{\mathrm{P}}-\left(\frac{\partial w}{\partial \mathrm{~T}}\right) \times\left(\frac{\partial \mathrm{T}}{\partial \mathrm{~V}}\right) \\
\mathrm{C}_{\mathrm{P}} & =\left(\frac{\partial \theta}{\partial \mathrm{T}}\right)_{\mathrm{P}} \text { and } \beta=\frac{1}{\mathrm{~V}} \frac{\partial \mathrm{~V}}{\partial \mathrm{~T}} \\
8 \partial w & =\mathrm{P} \partial \mathrm{~V}
\end{aligned}
$$

Using this

$$
\left(\frac{\partial \mathrm{U}}{\partial \mathrm{~V}}\right)_{\mathrm{P}}=\frac{\mathrm{C}_{\mathrm{P}}}{\beta \mathrm{~V}}-\mathrm{P}
$$

(c) Find $\Delta W$ and $\Delta U$ for an iron cube of side 6 cm as it is heated from $20^{\circ} \mathrm{C}$ to $300^{\circ} \mathrm{C}$. For iron $\mathrm{C}=0.11 \mathrm{cal} / \mathrm{g}^{\circ} \mathrm{C}$ and volume coefficient of expansion is $\beta=3.6 \times 10^{-5 \circ} \mathrm{C}^{-1}$. Given, Mass of the cube is 1700 gm .

Ans.

$$
\begin{align*}
\mathrm{T}_{1} & =20^{\circ} \mathrm{C}=293 \mathrm{~K}  \tag{7}\\
\mathrm{~T}_{2} & =300^{\circ} \mathrm{C}=573 \mathrm{~K} \\
\mathrm{C} & =0.11 \mathrm{cal} / \mathrm{g}^{\circ} \mathrm{C} . \beta=3.6 \times 10^{-5}{ }^{\circ} \mathrm{C}^{-1} \\
\mathrm{M} & =1700 \mathrm{gm} \\
\Delta \mathrm{Q} & =\text { Heat }=m \mathrm{C} \Delta \mathrm{~T} \\
& =1700 \times 0.11 \times[573-293] \\
& =52360 \mathrm{Cal} \\
\mathrm{P} & =1 \mathrm{~atm}=10^{-5} \mathrm{~N} / \mathrm{m}^{2} \\
w & =\mathrm{P} d \mathrm{~V} \\
\beta & =\frac{1}{\mathrm{~V}}\left(\frac{\partial \mathrm{~V}}{\partial \mathrm{~T}}\right)
\end{align*}
$$

So, $d \mathrm{~V}=\beta \mathrm{V} d \mathrm{~T}$

$$
\begin{aligned}
& =3.6 \times 10^{-5} \times(6)^{3} \times 280 \\
& =2.177 \mathrm{~cm}^{3}=2.177 \times 10^{-6} \mathrm{~m}^{3} \\
w & =\mathrm{P} d \mathrm{~V}=105 \times 2.177 \times 10-6 \\
w & =0.21775 \\
& =\frac{0.2177}{4184} \mathrm{col}=5 \times 10^{-5} \mathrm{cal} \\
\Delta \mathrm{U} & =\Delta \mathrm{Q}-\Delta w \\
\Delta \mathrm{~V} & =52360 \mathrm{cal}
\end{aligned}
$$

Q. 3. (a) What are reversible and irreversible processes? Give one example of each. Prove that if Kelvin-Planck statement of second law is violated then Clausius statement is also violated.

Ans. The solution to this questions is in this textbook.
(b) If, two Carnot engines $R$ and $S$ are operated in series such as engine $T$ absorbs heat at temperature $T_{1}$ and rejects heat to the sink at temperature $T_{2^{\prime}}$ while Engine $S$ absorbs half of the heat rejected by engine $R$ and rejects heat to the sink at temperature $\mathrm{T}_{3}$. If the work done in both the cases is equal, show that $\mathrm{T}_{2}=\left(\mathrm{T}_{3}+2 \mathrm{~T}_{1}\right) / 3$.

Ans. The solution to this questions is in this textbook.
(c) (i) A refrigerator freezes 6 kg of water at $0^{\circ} \mathrm{C}$ into ice in a time interval of 20 min . Assume that room temp, is $25^{\circ} \mathrm{C}$, calculate the power needed to accomplish it.
(ii) If coefficient of performance of a refrigerator is 5 and operates at the room temperature $27^{\circ} \mathrm{C}$, find the temperature inside the refrigerator.

Ans. $\quad m=6 \mathrm{~kg} \quad$ time $=20 \mathrm{~min}$
Heat required to convert $0^{\circ} \mathrm{C}$ water to $0^{\circ} \mathrm{C}$ ice

$$
\mathrm{Q}=6 \times 80=480 \mathrm{~K} \mathrm{cal}
$$

Room temperature $\mathrm{T}=25^{\circ} \mathrm{C}=298 \mathrm{~K}$
convert it to $0^{\circ} \mathrm{C}\left(\mathrm{T}^{1}\right)=273 \mathrm{~K}$
work done $=w$

$$
\begin{aligned}
& \frac{\theta}{w}=\frac{\mathrm{T}^{1}}{\mathrm{~T}-\mathrm{T}^{1}} \\
& w=\theta \frac{\left(\mathrm{T}-\mathrm{T}^{1}\right)}{\mathrm{T}^{1}} \\
& w=480\left(\frac{298-273}{273}\right)
\end{aligned}
$$

$$
\begin{align*}
& =43.95 \mathrm{~K} \mathrm{cal} \\
w & =43.95 \times 10^{3} \times 4.25  \tag{7}\\
\mathrm{P} & =\frac{w}{t}=\frac{43.95 \times 10^{3} \times 4.2}{20 \times 60} \\
\mathrm{P} & =153.825 \text { watt. } \\
\beta & =5, \text { Room temperature }=27^{\circ} \mathrm{C} \\
\mathrm{~T} & =300 \mathrm{~K} \\
\mathrm{~T}^{1} & =\text { temperature inside } \\
\mathrm{S} & =\frac{\mathrm{T}^{1}}{200-\mathrm{T}^{1}} \\
1500-5 \mathrm{~T}^{1} & =\mathrm{T}^{1} \Rightarrow 6 \mathrm{~T}^{1}=1500 \\
\mathrm{~T}^{1} & =250 \mathrm{~K}
\end{align*}
$$

(ii)
Q.4. (a) Define entropy. What is principle of increase of entropy? Find increase in entropy for reversible and irreversible processes.

Ans. The solution to this questions is in this textbook.
(b) If two bodies have equal mass $m$ and heat capacity $c$, are kept at different temperature $T_{1}$ and $T_{2}$ respectively, taking $T_{1}>T_{2}$ and the first body as source of heat for reversible engine and the second as sink, find out the maximum work done.

Ans. The solution to this questions is in this textbook.
(c) (i) The temperature variation of $C_{P}$ is given by the relation $\mathrm{C}_{\mathrm{P}}=0.4 \mathrm{~T}-0.05 \mathrm{~T}^{2}-0.25$, in the temperature range 50 K to 100 K in cal $/ \mathrm{K}$. If 4 moles of the substance is heated from 50 K to 100 K , calculate the change in entropy.
(ii) An ideal gas is confined to a cylinder by a piston. The piston is slowlyn pushed such that the gas temperature remains at $20^{\circ} \mathrm{C}$. During compression, 730 J of work is done on the gas. Find the entropy change of the gas.

Ans. (i) Change in intrapy $=\Delta S=\int \frac{\partial \theta}{\mathrm{T}}$

$$
\begin{aligned}
& \mathrm{C}_{\mathrm{P}}=\frac{d \theta}{d \mathrm{~T}} \quad \Rightarrow \quad \partial \theta=\mathrm{C}_{\mathrm{P}} d \mathrm{~T} \\
& \Delta \mathrm{~S}=n \int \frac{\mathrm{C}_{\mathrm{P}} d \mathrm{~T}}{\mathrm{~T}} \quad n=4 \mathrm{~mol} \\
& \Delta \mathrm{~S}=4 \int_{\mathrm{T}_{1}}^{\mathrm{T}_{2}} \frac{0.4 \mathrm{~T}-0.05 \mathrm{~T}^{2}-0.25 d \mathrm{~T}}{\mathrm{~T}}
\end{aligned}
$$

$$
\begin{aligned}
& =4 \int_{\mathrm{T}_{1}}^{\mathrm{T}_{2}}\left(0.4-0.05 \mathrm{~T}-\frac{0.25}{\mathrm{~T}}\right) d \mathrm{~T} \\
& =4\left[0.4\left(\mathrm{~T}_{2}-\mathrm{T}_{1}\right)-\frac{0.05}{2}\left(\mathrm{~T}_{2}^{2}-\mathrm{T}_{1}^{2}\right)-0.25 \ln \mathrm{~T} \mathrm{~T}\right]_{\mathrm{T}_{1}}^{\mathrm{T}_{2}} \\
\Delta \mathrm{~S} & \left.=4\left[0.4(100-50)-\frac{0.05}{2}[100]^{2}-(50)^{2}\right]-0.25 \ln \frac{100}{50}\right] \\
& =4\left[20-\frac{0.05}{2} \times 7500-0.25 \ln 2\right] \\
\Delta \mathrm{S} & =4[20-187.5-0.25 \ln 2]
\end{aligned}
$$

(ii)

$$
\begin{aligned}
\mathrm{T} & =20^{\circ} \mathrm{C}=293 \mathrm{~K} \\
w & =730 \mathrm{~S}
\end{aligned}
$$

Temperature is fixed. So, $\Delta \mathrm{U}=0$

$$
\Delta \theta=\Delta w
$$

$$
\lambda S=\frac{d \theta}{\mathrm{~T}}=\frac{d w}{\mathrm{~T}}
$$

$$
\text { Change in entrapy }=\frac{d w}{\mathrm{~T}}=\frac{730}{293}
$$

$$
\Delta S=2.49 \mathrm{cal} \mathrm{~K}^{-1} .
$$

Q. 5. (a) What are thermodynamic potentials? Why are they so called? Give relations for them. Write physical significance of Gibb's free energy.

Ans. The solution to this questions is in this textbook.
(b) Apply Maxwell's relation to prove that the difference of isothermal compressibility and adiabatic compressibility is equal to $T V \beta^{2} / \mathrm{C}_{\mathrm{P}}$.

Ans. The solution to this questions is in this textbook.
(c) Minute droplets of water are slowly pushed out of an atomizer into air. The average radius of the droplets is $10^{-4} \mathrm{~cm}$. If 1 kg of water is atomized isothermally at $25^{\circ} \mathrm{C}$, calculate the amount of heat transferred. The specific volume of water at $25^{\circ} \mathrm{C}$ is $1.00187 \times 10^{-3} \mathrm{~m}^{3} \mathrm{~kg}^{-1}$ and the rate of change of surface tension of water


Ans.

$$
\begin{equation*}
r=10^{-4} \mathrm{~cm}, \quad m=1 \mathrm{~kg}, \quad \text { temperature }=25^{\circ} \mathrm{C} \tag{7}
\end{equation*}
$$

Specific volume $=1.00187 \times 10^{-3} \mathrm{~m}^{3} \mathrm{~kg}^{-1}$

$$
\frac{\Delta \mathrm{S}}{\mathrm{~T}}=-0.152 \times 10^{-3} \mathrm{Nm}^{-1} \mathrm{k}^{-1}
$$

$$
\begin{aligned}
\text { Excess pressure } & =\mathrm{P}=\frac{2 \times \Delta \mathrm{S}}{r} \\
\mathrm{P} & =\frac{2 \times\left(-0.152 \times 10^{-3}\right) \times 298}{10^{-4}} \\
& =905.9 \\
\text { Volume } & =\text { Specific Volume } \times \text { mass } \\
& =1.00187 \times 10^{-3} \times 1=1.00187 \times 10^{-3} \mathrm{~m}^{3} \\
\text { Work done } & =\mathrm{P} \Delta \mathrm{~V}=-905.9 \times 1.00187 \times 10^{-3} \\
w & =-0.90761 \\
\text { Energy } & =\text { Surface tension } \times \text { Area } \\
& =0.152 \times 10^{-3} \times 298 \times 4 \times 3.14 \times 10^{-8} \times 10^{-8} \\
& =568.91 \times 10-19 \mathrm{~S} \\
\Delta \mathrm{~S} & =\Delta \mathrm{U}-\Delta w \\
\Delta \theta & =0.90761 .
\end{aligned}
$$

Q.6. (a) Define mean free path $(\lambda)$ of molecules of a gas. Derive the expression $\lambda=\frac{3}{4 \pi \sigma^{2} n}$. Where $\sigma$ is the diameter of the gas molecules and $n$ is the no. of molecules per unit volume. (Assuming that all molecules move with the same velocity i.e., that the average velocity of the gas.

Ans. The solution to this questions is in this textbook.
(b) (i) Plot Maxwell distribution function for molecular speeds temperatures $T_{1}, T_{2}$ and $T_{3}$ such as $T_{1}<T_{2}<T_{3}$. Write the necessary inference from these curves. Write the necessary inference from these curves.
(ii) Calculate the value of $v_{x}$ for which the probability of a molecule having $x$-velocity falls to half of its maximum value.

Ans. (i) The solution to this questions is in this textbook.
(ii) The solution to this questions is in this textbook.
(c) (i) Calculate the probability that the speed of oxygen molecule lies between 109.5 and 110.5 metre/sec. at 300 K .
(ii) Hydrogen and Nitrogen are maintained under identical conditions of temperature and pressure. Calculate the ratio of their coefficients of viscosity if the diameters of these molecules are $2.5 \times 10^{-10} \mathrm{~m}$ and $3.5 \times 10^{-10} \mathrm{~m}$ respectively.

Ans. (i) The probability that a molecule will have speed between $\mathrm{VS} \mathrm{V}+d \mathrm{~V}$ is

$$
\frac{d \mathrm{~N}_{\mathrm{V}}}{\mathrm{~N}}=4 \pi\left(\frac{m}{2 \pi k_{\mathrm{B}} \mathrm{~T}}\right)^{3 / 2} \mathrm{~V}^{2} e-\left(\frac{m \mathrm{~V}^{2}}{2 k_{\mathrm{B}} \mathrm{~T}}\right)
$$

$$
\begin{aligned}
& m=32 a=\frac{32}{6 \times 10^{26}} \mathrm{~kg} \\
& \mathrm{~V}=109.5 \\
& d \mathrm{~V}=110.5-109.5=1 \mathrm{~m} / \mathrm{s} \\
& k_{\mathrm{B}}=1.38 \times 10^{-23} \\
& \mathrm{~T}=300 k \\
& \frac{d^{\mathrm{N}} \mathrm{~V}}{\mathrm{~N}}=4 \pi\left(\frac{\frac{32}{6} \times 10^{26}}{2 \times 3.14 \times 1.38 \times 10^{-23} \times 300}\right) \times(109.5)^{2} \\
& \times \exp \left[\frac{-\frac{32}{6} \times 10^{-26} \times 109 \times 109}{2 \times 1.38 \times 10^{-23} \times 300}\right] \times 1 \\
& \frac{d^{\mathrm{N}} \mathrm{~V}}{\mathrm{~N}}=8.11 \times 10^{-4} . \\
& n=\text { viscosity }=\frac{1}{3 \sqrt{2}} \frac{1}{\pi d^{2}} m \overline{\mathrm{~V}}
\end{aligned}
$$

Since both have at same temperature
So, $\overline{\mathrm{V}}$ same.

So,

$$
\begin{aligned}
& \frac{n_{1}}{n_{2}}=\frac{d_{2}^{2}}{d_{1}^{2}} \\
& \frac{n_{\mathrm{H}}}{n_{\mathrm{N}}}=\frac{d_{\mathrm{N}}^{2}}{d_{\mathrm{H}}^{2}}=\frac{(3.5)^{2} \times\left(10^{-10}\right)^{2}}{(2.5)^{2} \times\left(10^{-10}\right)^{2}} \\
& \frac{n_{\mathrm{H}}}{n_{\mathrm{N}}}=1.96
\end{aligned}
$$

Q. 7. (a) Discuss Joule-Thomson porous plug experiment. Obtain equation for Joule-Thomson co-efficient.

Ans. The solution to this questions is in this textbook.
(b) What are the limitations of Van der waal's equation of state. Draw and discuss similarities and dis-similarities of theoretical experimental curves for $\mathrm{CO}_{2}$ gas.

Ans. The solution to this questions is in this textbook.
(c) The Van der Waal's constant for Hydogen are $a=0.247 \mathrm{~atm}$. litre ${ }^{2} \mathrm{~mol}^{-2}$ and $b=2.65 \times 10^{-2}$ litre $/ \mathrm{mol}$. Calculate :
(i) The temperature of inversion
(ii) Joule Thomson coefficient for 2 atm fall of pressure, initial temp, being 100 K . Given $\mathrm{R}={ }^{224} / 273$ atoms litre $/ \mathrm{mol} / \mathrm{K}$.

Ans. (i) $\quad a=0.247 \mathrm{~atm}, \quad b=2.65 \times 10^{-2}$ litre $^{2} \mathrm{~mol}^{2}$
Inversion temperature $=\mathrm{T}_{i}=\frac{2 a}{12 b}$

$$
\mathrm{T}_{i}=\frac{2 \times 0.247 \times 273}{224 \times 2.65 \times 10^{-2}}=22.7 \mathrm{k}
$$

(ii)

$$
\mathrm{T}_{i}=22.7 \mathrm{k}
$$

$$
\Delta \mathrm{P}=2 \mathrm{~atm}
$$

Initial temperature $=100 k$
Change in temperature $=100-22.7$

$$
\begin{aligned}
\Delta \mathrm{T} & =77.3 \mathrm{k} \\
\Delta \mathrm{~T} & =\mu \Delta \mathrm{P}
\end{aligned}
$$

$$
\begin{aligned}
& \mu=\frac{\Delta T}{\Delta \mathrm{P}}=\frac{77.3}{2} \\
& \mu=38.65 \mathrm{katm}^{-1} .
\end{aligned}
$$

## All Lab Experiments

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