## SAMPPLz OONHFINT

## Precise

## PiYYICS

## BASED ON NEW PAPER PATTERN



## Written as per the latest textbook prescribed by the Maharashtra State Bureau of Textbook Production and Curriculum Research, Pune.

## Precise <br> PHYSICS (vol.1) Std. XII Sci.

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[^0]Precise Physics Vol. I, Std. XII Sci. is intended for every Maharashtra State Board aspirant of Std. XII, Science. The scope, sequence, and level of the book are designed to match the new textbook of Maharashtra State board.

We believe that the study of Physics needs driving desire to learn and willingness to work hard. The earnestness to dive into the subject eventuates naturally when students are presented with meaningful content that is easy to read and understand rather than being mired down with facts and information. Students do much better when they grasp the nub of the subject.

While beginning with a chapter in Physics, students should study to understand the content and not merely read point blankly. They can go slowly, knowing Physics can't be read like a novel, choosing their own pace. But it is essential for students to comprehend the concepts involved, ruminate and reproduce their own versions of the same.

## To quote Albert Einstein, "If you can't explain it simply, you don't understand it yourself."

Students should then attempt theoretical questions based on these concepts to gauge the level of understanding achieved.

Next advance after gaining command over theory would be numericals. Though Physics is communicated in English, it is expressed in Mathematics. Hence, it is essential to befriend formulae and derivations. These should be learnt and memorized. Once physical mathematics of concept is ingrained, solved numericals should be studied, starting from simple problems to difficult by escalating level of complexity gradually. Students are required to practise numericals and ascertain their command on problem solving. Calculations at this stage must be done using log table keeping in mind that calculators are not allowed in Board Exams. When it comes to problems in Physics nothing makes students perfect like practice!

Amongst building concepts, advancing into numbers and equations, it is essential to ponder underlying implications of subject. Students should read from references, visit authentic websites and watch relevant fascinating links.

Such a holistic preparation of subject is the key to succeed in the board examination.
After all, "Success is no accident. It is hard work, perseverance, learning, studying, sacrifice and most of all, love of what you are doing or learning to do," as said by legendary football player Pele and students should bear it in mind!

Our Precise Physics Vol. I, Std. XII Sci. adheres to our vision and achieves several goals: building concepts, developing competence to solve numericals, recapitulation and self-study -all while encouraging students toward cognitive thinking.

Features of the book presented below will explicate more about the same!
We hope the book benefits the learner as we have envisioned.
Publisher
Edition: Fifth

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- There will be single question paper of 70 Marks and practical examination of 30 Marks in Physics.
- Duration of the paper will be 3 hours.


## Section A:

This section will contain Multiple Choice Questions and Very Short Answer (VSA) type of questions. There will be 10 MCQs and 8 VSA type of questions, each carrying one mark.
Students will have to attempt all these questions.

## Section B:

This section will contain 12 Short Answer (SA-I) type of questions, each carrying 2 marks.
Students will have to attempt any 8 questions.

## Section C:

(24 Marks)
This section will contain 12 Short Answer (SA-II) type of questions, each carrying 3 marks. Students will have to attempt any 8 questions.

## Section D:

This section will contain 5 Long Answer (LA) type of questions, each carrying 4 marks.
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Distribution of Marks According to the Type of Questions

| Type of Questions |  |  |
| :--- | :---: | :---: |
| MCQ | 1 Mark each | 10 Marks |
| VSA | 1 Mark each | 8 Marks |
| SA - I | 2 Marks each | 16 Marks |
| SA - II | 3 Marks each | 24 Marks |
| LA | 4 Marks each | 12 Marks |


| Percentage wise distribution of marks |  |
| :---: | :---: |
| Theory | $63 \%$ |
| Numerical | $37 \%$ |

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[Reference: Maharashtra State Board of Secondary and Higher Secondary Education, Pune - 04]

Note: 1. * mark represents Textual question.
2. \# mark represents Intext question.
3. + mark represents Textual examples.
4. 㧚谷 symbol represents textual questions that need external reference for an answer.
5. Chapters 8 to 16 are a part of Std. XII: Precise Physics (Vol. II).

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## 3 Kinetic Theory of Gases and Radiation

## Contents and Concepts

3.1 Introduction
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### 3.1 Introduction

Q.1. Using the gas laws, prove that $\mathbf{P V}=\mathbf{N k}_{\mathrm{B}} \mathbf{T}$. Also state the significance of terms involved. OR
Can you recall? (Textbook page no. 56)
How do you get ideal gas equation from the gas laws?
[3 Marks]
Ans:
i. The three gas laws applied to a fixed mass $m$ of an enclosed gas are,
a. Boyle's law: $\mathrm{V} \propto \frac{1}{\mathrm{P}}$ at constant T .
b. Charles' law: $\mathrm{V} \propto \mathrm{T}$ at constant P .
c. Gay-Lussac's law: $\mathrm{P} \propto \mathrm{T}$ at constant V .
ii. Combining the three laws,
$P V \propto T$ i.e., $\frac{P_{1} V_{1}}{T_{1}}=\frac{P_{2} V_{2}}{T_{2}}$
iii. In terms of number of moles ( n ) of the gas,
$\mathrm{PV} \propto \mathrm{nT}$
$\therefore \quad \mathrm{PV}=\mathrm{nRT}$
Where, $\mathrm{R}=$ proportionality constant $=$ Universal gas constant.
iv. But, number of moles ( n )
$=\frac{\text { mass of the gas }(\mathrm{m})}{\text { molar mass }\left(\mathrm{M}_{0}\right)}=\frac{\mathrm{N}}{\mathrm{N}_{\mathrm{A}}}$
Where, $\mathrm{N}=$ number of molecules,

$$
\mathrm{M}_{0}=\text { mass of } 1 \text { mole of gas }
$$

$\mathrm{N}_{\mathrm{A}}=$ Avogadro's number
v. The universal gas constant can also be expressed in terms of Boltzmann constant $\left(\mathrm{k}_{\mathrm{B}}\right)$ as, $\mathrm{R}=\mathrm{N}_{\mathrm{A}} \mathrm{k}_{\mathrm{B}}$
Substituting equations (2) and (3) in equation (1), we get,
$\therefore \quad \mathrm{PV}=\frac{\mathrm{N}}{\mathrm{N}_{\mathrm{A}}} \times \mathrm{N}_{\mathrm{A}} \mathrm{k}_{\mathrm{B}} \mathrm{T}=\mathrm{Nk}_{\mathrm{B}} \mathrm{T}$
Q.2. Why is an ideal gas equation known as equation of state?
[1 Mark]
Ans: For a gas, its state is specified by a number of physical quantities such as pressure P , temperature T, volume V, internal energy E, etc. Hence, the equation relating these quantities i.e., ideal gas equation is known as the equation of state.
*Q.3. Write ideal gas equation for a mass of 7 g of nitrogen gas.
[1 Mark]
Ans: Ideal gas equation for 7 g of nitrogen gas is, $\mathrm{PV}=\frac{\mathrm{RT}}{4}$.

## Reading between the lines

Ideal gas equation, $P V=n R T$
Here, $n=\frac{\mathrm{m}}{\mathrm{M}_{0}}$
$m=7 \mathrm{~g},\left(\mathrm{M}_{0}\right)_{\mathrm{N}_{2}}=28 \mathrm{~g} / \mathrm{mol}$
$\therefore \quad n=\frac{7}{28}=\frac{1}{4}$
$\therefore \quad P V=\frac{1}{4} \times \mathrm{RT}=\frac{\mathrm{RT}}{4}$


## Solved Examples

Q.4. Using the ideal gas equation, determine the value of $R$. Given that one gram molecule of a gas at S.T.P. occupies 22.4 litre.
[2 Marks]

## Solution:

Given:

$$
\text { At S.T.P., } \mathrm{P}=1 \times 10^{5} \mathrm{~N} / \mathrm{m}^{2},
$$

$\mathrm{V}=22.4$ litre

$$
=22.4 \times 10^{-3} \mathrm{~m}^{3}
$$

$\mathrm{T}=273 \mathrm{~K}$
To find
Universal gas constant (R)
Formula: $\quad \mathrm{PV}=\mathrm{RT}$
Calculation: From formula,

$$
\begin{aligned}
& \mathrm{R}=\frac{\mathrm{PV}}{\mathrm{~T}} \\
\therefore & \mathrm{R}=\frac{1 \times 10^{5} \times 22.4 \times 10^{-3}}{273} \\
\therefore & \mathrm{R}=\frac{320}{39} \\
\therefore & \mathrm{R}=\mathbf{8 . 2 0 5} \mathbf{~ J ~ m o l}^{\mathbf{- 1}} \mathbf{K}^{\mathbf{- 1}}
\end{aligned}
$$

Ans: The value of universal gas constant $R$ is $8.205 \mathrm{~J} \mathrm{~mol}^{-1} \mathrm{~K}^{-1}$.

## Reading between the lines

There is difference between values of pressure and temperature at S.T.P. (standard temperature pressure) and N.T.P. (normal temperature pressure) conditions.
At S.T.P., $P=10^{5} P a=1 \mathrm{bar}=1 \mathrm{~atm}$ and $T=0{ }^{\circ} \mathrm{C}=273 \mathrm{~K}$
At N.T.P., $P=101.32 \mathrm{kPa}=1.013 \times 10^{5} \mathrm{~Pa}$ and $T=20^{\circ} \mathrm{C}=293 \mathrm{~K}$ (Room temperature)
Q.5. 16 g of oxygen occupy $0.025 \mathrm{~m}^{3}$ at $27^{\circ} \mathrm{C}$. If the universal gas constant is $8.4 \mathrm{~J} / \mathrm{mol} \mathrm{K}$. Find the pressure exerted by it. [Molecular weight of oxygen $=32$ ]
[2 Marks]

## Solution:

Given:
$\mathrm{m}=16 \mathrm{~g}, \mathrm{~V}=0.025 \mathrm{~m}^{3}$,
$\mathrm{T}=27^{\circ} \mathrm{C}=273+27=300 \mathrm{~K}$,
$\mathrm{R}=8.4 \mathrm{~J} / \mathrm{mol} \mathrm{K}, \mathrm{M}=32 \mathrm{~g} / \mathrm{mol}$
To find: $\quad$ Pressure (P)
Formula: $\quad \mathrm{PV}=\mathrm{nRT}$
Calculation: Since, $\mathrm{n}=\frac{\mathrm{m}}{\mathrm{M}}=\frac{16}{32}=\frac{1}{2}=0.5$
From formula,

$$
\begin{aligned}
\mathrm{P} & =\frac{\mathrm{nRT}}{\mathrm{~V}} \\
& =\frac{0.5 \times 8.4 \times 300}{0.025} \\
\therefore \quad \mathrm{P} & =\mathbf{5 0 . 4} \times \mathbf{1 0}^{\mathbf{3}} \mathbf{N} / \mathbf{m}^{\mathbf{2}}
\end{aligned}
$$

Ans: The pressure exerted by 16 g of oxygen is $50.4 \times 10^{3} \mathrm{~N} / \mathrm{m}^{2}$.
*Q.6. Two vessels $A$ and $B$ are filled with same gas where volume, temperature and pressure in vessel $A$ is twice the volume, temperature and pressure in vessel $B$. Calculate the ratio of number of molecules of gas in vessel $A$ to that in vessel $B$.
[2 Marks]

## Solution:

Given: $\quad \mathrm{P}_{\mathrm{A}}=2 \mathrm{P}_{\mathrm{B}}, \mathrm{T}_{\mathrm{A}}=2 \mathrm{~T}_{\mathrm{B}}, \mathrm{V}_{\mathrm{A}}=2 \mathrm{~V}_{\mathrm{B}}$
To find: $\quad$ Ratio of number of molecules $\left(\mathrm{N}_{1}: \mathrm{N}_{2}\right)$
Formulae: i

$$
\text { i. } \quad \mathrm{n}=\frac{\mathrm{PV}}{\mathrm{RT}} \quad \text { ii. } \quad \mathrm{n}=\frac{\mathrm{N}}{\mathrm{~N}_{\mathrm{A}}}
$$

## Calculation:

From formula (i)
$\mathrm{n}_{\mathrm{A}}=\frac{\mathrm{P}_{\mathrm{A}} \mathrm{V}_{\mathrm{A}}}{\mathrm{R} \mathrm{T}_{\mathrm{A}}}$
$\mathrm{n}_{\mathrm{B}}=\frac{\mathrm{P}_{\mathrm{B}} \mathrm{V}_{\mathrm{B}}}{\mathrm{RT} \mathrm{T}_{\mathrm{B}}}$
Dividing equation (1) by equation (2),

$$
\begin{array}{ll}
\therefore & \frac{\mathrm{n}_{\mathrm{A}}}{\mathrm{n}_{\mathrm{B}}}=\frac{2 \mathrm{P}_{\mathrm{B}} \times 2 \mathrm{~V}_{\mathrm{B}} \times \mathrm{T}_{\mathrm{B}}}{2 \mathrm{~T}_{\mathrm{B}} \times \mathrm{P}_{\mathrm{B}} \times \mathrm{V}_{\mathrm{B}}} \\
\therefore & \frac{\mathrm{n}_{\mathrm{A}}}{\mathrm{n}_{\mathrm{B}}}=\frac{2}{1}
\end{array}
$$

From formula (ii),
$\frac{\left(\mathrm{N}_{1} / \mathrm{N}_{\mathrm{A}}\right)}{\left(\mathrm{N}_{2} / \mathrm{N}_{\mathrm{A}}\right)}=\frac{2}{1}$
$\therefore \quad \frac{\mathrm{N}_{1}}{\mathrm{~N}_{2}}=\frac{\mathbf{2}}{\mathbf{1}}$
Ans: The ratio of number of molecules is $\mathbf{2 : 1}$.

### 3.2 Behaviour of a Gas

Q.7. Why is it necessary to make assumptions while studying behaviour of a gas? [2 Marks]
Ans:
i. For any solid object, its motion can be described well with the help of Newton's laws of motion.
ii. Similarly, a gas enclosed in a container can be characterised by macroscopic state variables like pressure, volume and temperature.
iii. However, as gas molecules are always in random motion, it is difficult to understand behaviour of a gas in terms of motion of a single molecule.
iv. The number of molecules in the gas is so large $\left(\approx 10^{23}\right.$ per $\left.\mathrm{m}^{3}\right)$ that motion of individual molecule cannot be related with macroscopic parameters $\mathrm{P}, \mathrm{V}, \mathrm{T}$ and energy E .
Hence, it is necessary to make certain assumptions while studying behaviour of a gas.

### 3.3 Ideal Gas and Real Gas

*Q.8. What is an ideal gas? Does an ideal gas exist in practice?
[2 Marks]
Ans: A gas which obeys ideal gas equation at all pressures and temperatures is an ideal gas. No, such gas does not exist in reality.
Q.9. Mention the conditions under which a real gas obeys ideal gas equation.
[1 Mark]
Ans: Conditions under which a real gas obeys ideal gas equation are:
Low density, low pressure or high temperature. In other words, at condition where gas molecules are far apart so that molecular interactions are negligible.

## Reading between the lines

i. In an ideal gas, there are no intermolecular interactions whereas, real gases are composed of atoms or molecules which do interact with each other.
ii. When the atoms or molecules of a real gas are so far apart that there are no interatomic or intermolecular interactions possible, the real gas is said to be in ideal state.

### 3.4 Mean Free Path

Q.10. Explain the concept of mean free path of a gas molecule.
[3 Marks]

## Ans:

i. The molecules of a gas are uniformly dispersed throughout the volume of the gas, but are executing random motion. The typical path followed by a particle during Brownian motion is shown in figure.


A typical molecule in a gas executing random motion
ii. When a molecule approaches another molecule, there is a repulsive force between them, due to which the molecules behave as small hard spherical particles. This leads to elastic collisions between the molecules.
iii. Therefore, both the speed and the direction of motion of the molecules change abruptly. The molecules also collide with the walls of the container.
iv. Molecules exert force on each other only during collisions. Thus, in between two successive collisions the molecules move along straight paths with constant velocity.
v. The average distance traversed by a molecule with constant velocity between two successive collisions is called mean free path.
vi. The mean free path $(\lambda)$ varies
a. inversely with density $\rho=\frac{\mathrm{N}}{\mathrm{V}}$ of the gas. Where, $\mathrm{N}=$ number of molecules,

$$
\mathrm{V}=\text { volume of the gas }
$$

b. inversely with square of the diameter of molecule $d^{2}$ (because it depends on the cross-section of a molecule).
vii. Thus, from above proportionalities, it can be shown that,

$$
\lambda=\frac{1}{\sqrt{2} \pi \mathrm{~d}^{2}(\mathrm{~N} / \mathrm{V})}
$$

Q.11. How does mean free path varies when pressure is reduced?
[1 Mark]
Ans: If the pressure of a gas in an enclosure is reduced by evacuating it, the density of the gas decreases and the mean free path increases.

## Solved Examples

+Q.12.Obtain the mean free path of nitrogen molecule at $0{ }^{\circ} \mathrm{C}$ and 1.0 atm pressure. The molecular diameter of nitrogen is 324 pm (assume that the gas is ideal).
(Example 3.1 of Textbook page no. 58)
[3 Marks]

## Solution:

Given:

$$
\begin{array}{ll}
\text { Given: } & \mathrm{T}=0^{\circ} \mathrm{C}=273 \mathrm{~K}, \\
& \mathrm{P}=1.0 \mathrm{~atm}=1.01 \times 10^{5} \mathrm{~Pa}, \\
& \mathrm{~d}=324 \mathrm{pm}=324 \times 10^{-12} \mathrm{~m}, \\
& \mathrm{k}_{\mathrm{B}}=1.38 \times 10^{-23} \mathrm{~J} / \mathrm{K} \\
\text { To find: } & \text { Mean free path of } \mathrm{N}_{2}(\lambda) \\
\text { Formulae: } & \text { i. } \quad \mathrm{PV}=\mathrm{Nk}_{\mathrm{B}} \mathrm{~T} \\
& \text { ii. } \quad \lambda=\frac{1}{\sqrt{2} \pi \mathrm{~d}^{2}\left(\frac{\mathrm{~N}}{\mathrm{~V}}\right)}
\end{array}
$$

Formulae: i. $\quad \mathrm{PV}=\mathrm{Nk}_{\mathrm{B}} \mathrm{T}$

Calculation:
From formula (i),
$\frac{N}{V}=\frac{P}{k_{B} T}$
Using this in formula (ii),

$$
\begin{aligned}
\lambda & =\frac{\mathrm{k}_{\mathrm{B}} \mathrm{~T}}{\sqrt{2} \pi \mathrm{~d}^{2} \mathrm{P}} \\
& =\frac{1.38 \times 10^{-23} \times 273}{\sqrt{2} \pi\left(324 \times 10^{-12}\right)^{2} \times 1.01 \times 10^{5}}
\end{aligned}
$$

$$
=\operatorname{antilog}\{\log (1.38)+\log (273)-\log (1.414)
$$

$$
-\log (3.142)-2 \log (324)-\log (1.01)\}
$$

$$
\times 10^{-4}
$$

$\ldots$... using $\sqrt{2}=1.414$ )

$$
\begin{aligned}
& =\operatorname{antilog}\{0.1399+2.4362-0.1504 \\
& \quad-0.4972-5.0210-0.0043\} \times 10^{-4} \\
& =\operatorname{antilog}\{\overline{4} .9032\} \times 10^{-4} \\
& =8.002 \times 10^{-4} \times 10^{-4} \approx 8 \times 10^{-8} \\
& =\mathbf{0 . 8} \times \mathbf{1 0}^{-7} \mathbf{~ m}
\end{aligned}
$$

Ans: Mean free path of $\mathrm{N}_{2}$ molecule is $\mathbf{0 . 8} \times \mathbf{1 0}^{\mathbf{- 7}} \mathbf{~ m}$.
Q.13. Mean free path of oxygen molecule is 70 nm at S.T.P. Find the molecular diameter of the oxygen (in pm) assuming it to be an ideal gas. (Take value of Boltzmann constant as $1.38 \times 10^{-23} \mathrm{~J} / \mathrm{K}$.)
[2 Marks]

## Solution:

Given:

$$
\begin{aligned}
& \text { At S.T.P., } \mathrm{T}=0^{\circ} \mathrm{C}=273 \mathrm{~K}, \\
& \mathrm{P}=1 \times 10^{5} \mathrm{~Pa}, \\
& \lambda=70 \mathrm{~nm}=7 \times 10^{-8} \mathrm{~m}, \\
& \mathrm{k}_{\mathrm{B}}=1.38 \times 10^{-23} \mathrm{~J} / \mathrm{K}
\end{aligned}
$$

To find: Molecular diameter of oxygen (d)
Formula: $\quad \lambda=\frac{\mathrm{k}_{\mathrm{B}} \mathrm{T}}{\sqrt{2} \pi \mathrm{~d}^{2} \mathrm{P}}$
Calculation: From formula,

$$
\begin{aligned}
\mathrm{d}^{2}= & \frac{\mathrm{k}_{\mathrm{B}} \mathrm{~T}}{\sqrt{2} \pi \lambda \mathrm{P}} \\
= & \frac{1.38 \times 10^{-23} \times 273}{\sqrt{2} \times 3.142 \times 7 \times 10^{-8} \times 10^{5}} \\
= & \frac{1.38 \times 273}{1.414 \times 3.142 \times 7} \times 10^{-20} \\
= & \frac{1.38 \times 39}{1.414 \times 3.142} \times 10^{-20} \\
= & \operatorname{antilog}\{\log (1.38)+\log (39) \\
& \quad-\log (1.414)-\log (3.142)\} \times 10^{-20} \\
= & \operatorname{antilog}\{0.1399+1.5911-0.1504 \\
= & \operatorname{antilog}\{1.0834\} \times 10^{-20} \\
\mathrm{~d}^{2} & =1.212 \times 10^{-20} \times 10 \\
\mathrm{~d}= & 3.481 \times 10^{-10} \mathrm{~m} \\
= & \mathbf{3 4 8 . 1} \mathbf{~ p m}
\end{aligned}
$$

Ans: Molecular diameter of oxygen is $348.1 \mathbf{~ p m}$.

### 3.5 Pressure of Ideal Gas

Q.14. Derive an expression for a pressure exerted by a gas on the basis of kinetic theory of gases.
[4 Marks] [Mar 22]

## Ans:

i. Let there be $n$ moles of an ideal gas enclosed in a cubical box of volume $V\left(=L^{3}\right)$ with sides of the box parallel to the coordinate axes, as shown in figure. The walls of the box are kept at a constant temperature T .
ii. The gas molecules are in continuous random motion, colliding with each other and hitting the walls of the box and bouncing back.
iii. As per one of the assumptions, we neglect intermolecular collisions and consider only elastic collisions with the walls.

iv. A typical molecule moving with the velocity $\overrightarrow{\mathrm{v}}$, about to collide elastically with the shaded wall of the cube parallel to yz-plane.
v. During elastic collision, the component $\mathrm{v}_{\mathrm{x}}$ of the velocity will get reversed, keeping $\mathrm{v}_{\mathrm{y}}$ and $\mathrm{v}_{\mathrm{z}}$ components unaltered.
vi. Hence the change in momentum of the particle is only in the $x$ component of the momentum, $\Delta p_{x}$ is given by,
$\Delta p_{x}=$ final momentum - initial momentum

$$
\begin{equation*}
=\left(-\mathrm{mv}_{\mathrm{x}}\right)-\left(\mathrm{mv}_{\mathrm{x}}\right)=-2 \mathrm{mv}_{\mathrm{x}} \tag{1}
\end{equation*}
$$

vii. Thus, the momentum transferred to the wall during collision is $+2 \mathrm{mv}_{\mathrm{x}}$. The re-bounced molecule then goes to the opposite wall and collides with it.
viii. After colliding with the shaded wall, the molecule travels to the opposite wall and travels back towards the shaded wall again.
ix. This means that the molecule travels a distance of 2 L in between two collisions.
x. As $L$ is the length of the cubical box, the time for the molecule to travel back and forth to the shaded wall is $\Delta t=\frac{2 \mathrm{~L}}{\mathrm{v}_{\mathrm{x}}}$.
xi. Average force exerted on the shaded wall by molecule 1 is given as,
Average force $=$ Average rate of change of
momentum
$\therefore \quad F_{a v g}=\frac{2 m v_{x_{1}}}{2 L / v_{x_{1}}}=\frac{m v_{x_{1}}^{2}}{L}$
where $v_{x_{1}}$ is the $x$ component of the velocity of molecule 1 .
xii. Considering other molecules $2,3,4 \ldots$ with the respective $x$ components of velocities $\mathrm{v}_{\mathrm{x}_{2}}, \mathrm{v}_{\mathrm{x}_{3}}, \mathrm{v}_{\mathrm{x}_{4}} \ldots$, the total average force on the wall is,
$F_{\text {avg }}=\frac{m}{L}\left(v_{x_{1}}^{2}+v_{x_{2}}^{2}+v_{x_{3}}^{2}+\ldots.\right) \quad \ldots .[$ From (2)]
$\therefore \quad$ The average pressure
$\mathrm{P}=\frac{\text { Average force }}{\text { Area of shaded wall }}$

$$
=\frac{\mathrm{m}\left(\mathrm{v}_{\mathrm{x}_{1}}^{2}+\mathrm{v}_{\mathrm{x}_{2}}^{2}+\ldots .\right)}{\mathrm{L} \times \mathrm{L}^{2}}
$$

xiii. The average of the square of the $x$ component of the velocities is given by,
$\overline{v_{x}^{2}}=\frac{v_{x_{1}}^{2}+v_{x_{2}}^{2}+v_{x_{3}}^{2}+\ldots .+v_{N}^{2}}{N}$
$\therefore \quad \mathrm{P}=\frac{\mathrm{mN} \overline{\mathrm{v}_{\mathrm{x}}^{2}}}{\mathrm{~V}}$
where $\overline{v_{x}^{2}}$ is the average over all possible values of $\mathrm{v}_{\mathrm{x}}$.

## Quick Review



## Law of equipartition of energy

- For a gas in thermal equilibrium at temperature T , the average energy for molecule associated with each quadratic term is $\frac{1}{2} k_{B} T$.
- Translational K.E.:
$\frac{1}{2} m v_{x}^{2}+\frac{1}{2} m v_{y}^{2}+\frac{1}{2} m v_{z}^{2}$
- Rotational K.E.:
$\frac{1}{2} \mathrm{I} \omega_{\mathrm{x}}^{2}+\frac{1}{2} \mathrm{I} \omega_{\mathrm{y}}^{2}+\frac{1}{2} \mathrm{I} \omega_{z}^{2}$
- Vibrational K.E.: $\frac{1}{2} \mathrm{mu}^{2}+\frac{1}{2} \mathrm{kr}^{2}$


## Degrees of Freedom

- The total number of coordinates or independent quantities required to describe the position and configuration of the system completely
- Monatomic Gas:

Translation dof : 3

- Diatomic Gas:

Translation dof : 3
Rotational dof : 2 or 3
Vibrational dof : 0 or 2



## Impotant formulae

## 1. Ideal gas equation:

i. $\quad \mathrm{PV}=\mathrm{nRT}$
ii. $\quad \mathrm{PV}=\mathrm{Nk}_{\mathrm{B}} \mathrm{T}$
2. Mean free path:
i. $\lambda=\frac{1}{\sqrt{2} \pi \mathrm{~d}^{2}\left(\frac{\mathrm{~N}}{\mathrm{~V}}\right)}$
ii. $\quad \lambda=\frac{\mathrm{k}_{\mathrm{B}} \mathrm{T}}{\sqrt{2} \pi \mathrm{~d}^{2} \mathrm{P}}$
3. Pressure exerted by gas:
$P=\frac{1}{3} \frac{\mathrm{~N}}{\mathrm{~V}} \mathrm{~m} \overline{\mathrm{v}^{2}}$
4. Root mean square speed:
i. $\quad \mathrm{V}_{\mathrm{rms}}=\sqrt{\frac{3 \mathrm{RT}}{\mathrm{M}_{0}}}$
ii. $\quad V_{\text {rms }}=\sqrt{\frac{3 P}{\rho}}$
iii. $\quad \mathrm{V}_{\mathrm{rms}}=\sqrt{\frac{3}{2} \mathrm{k}_{\mathrm{B}} \mathrm{T}}$
5. Kinetic energy of gas molecule:
i. K.E of gas molecules $=\frac{3}{2} \mathrm{PV}$
ii. K.E per unit mole $=\frac{3}{2}$ RT
iii. K.E per unit mass $=\frac{3}{2} \frac{\mathrm{RT}}{\mathrm{M}_{0}}$
iv. K.E per molecule $=\frac{3}{2} \mathrm{k}_{\mathrm{B}} \mathrm{T}$
6. Relation between $\mathbf{C}_{\mathbf{p}}$ and $\mathrm{C}_{\mathbf{v}}$ :
i. $\quad C_{P}-C_{V}=R$
....(When all quantities are expressed in same unit.)
$C_{P}-C_{V}=\frac{R}{J}$
$\ldots$.... When $\mathrm{C}_{\mathrm{P}}, \mathrm{C}_{\mathrm{V}}$ are in heat units and R is in work unit.)
ii. $\quad \frac{C_{P}}{C_{V}}=\gamma$
iii. $\quad S_{p}-S_{v}=\frac{R}{M_{0} J}$
7. Radiant energy incident on a surface: $\mathrm{Q}=\mathrm{Q}_{\mathrm{a}}+\mathrm{Q}_{\mathrm{r}}+\mathrm{Q}_{\mathrm{t}}$
8. Coefficient of radiation:
i. Coefficient of absorption, $a=\frac{Q_{a}}{Q}$
ii. Coefficient of reflection, $r=\frac{Q_{r}}{Q}$
iii. Coefficient of transmission, $t_{r}=\frac{Q_{t}}{Q}$
9. Relation between $\mathbf{a}, \mathbf{r}$, and $\mathbf{t}: \mathrm{a}+\mathrm{r}+\mathrm{t}_{\mathrm{r}}=1$
10. Coefficient of emission (Emissivity): $e=\frac{R}{R_{b}}=a$
11. Emissive power: $\mathrm{R}=\frac{\mathrm{Q}}{\mathrm{At}}$
12. Quantity of radiant heat emitted by a blackbody:
i. $\quad \mathrm{Q}=\sigma \mathrm{AT}^{4} \mathrm{t}$
(When temperature of surrounding is not given)
ii. $\quad \mathrm{Q}=\sigma \mathrm{A}\left(\mathrm{T}^{4}-\mathrm{T}_{0}^{4}\right) \mathrm{t}$
(When temperature of the surrounding is given)
13. Radiant energy emitted by ordinary body:
i. $\quad Q=e A \sigma T^{4} t \quad$ ii. $\quad Q=e A \sigma\left(T^{4}-T_{0}^{4}\right) t$
14. Wien's law: $\lambda_{\max }=\frac{\mathrm{b}}{\mathrm{T}}$
15. Rate of heat radiation: $\frac{d Q}{d t}=e \mathrm{~A} \sigma\left(\mathrm{~T}^{4}-\mathrm{T}_{0}^{4}\right)$
16. Total radiant energy emitted from a body: $\mathrm{Q}=\mathrm{e} \operatorname{At\sigma }\left(\mathrm{T}^{4}-\mathrm{T}_{0}^{4}\right)$

## Exercise

### 3.1 Introduction

1. Prove the relation, $\mathrm{PV}=\mathrm{Nk}_{\mathrm{B}} \mathrm{T}$. [3 Marks]

Ans: Refer Q. 1
2. $\quad 14 \mathrm{~g}$ of nitrogen occupy $0.028 \mathrm{~m}^{3}$ at $27^{\circ} \mathrm{C}$. If the universal gas constant is $8.4 \mathrm{~J} / \mathrm{mol} \mathrm{K}$. Find the pressure exerted by it. [Molecular weight of nitrogen $=28$ ]
[2 Marks]
Ans: $45 \times 10^{3} \mathrm{~N} / \mathrm{m}^{2}$

### 3.2 Behaviour of a Gas

3. Why cannot the behaviour of a gas molecule be studied using Newtonian mechanics? [2 Marks]
Ans: Refer $Q .7$

### 3.3 Ideal Gas and Real Gas

4. Define ideal gas.
[1 Mark]
Ans: Refer $Q .8$
5. When can a real gas be treated as ideal gas?
[1 Mark]
Ans: Refer $Q .9$

### 3.4 Mean Free Path

6. What is a mean free path?
[1 Mark]
Ans: Refer Q. 10 (v)
7. How does a mean free path of a gas molecule vary with number density and size of the molecule?
[1 Mark]
Ans: Refer Q. 10 (vi)
8. Obtain the mean free path of nitrogen molecule at $0^{\circ} \mathrm{C}$ and 1.0 atm pressure. The molecular diameter of oxygen is 335 pm (assume that the gas is ideal). (Take $\mathrm{k}_{\mathrm{B}}=1.38 \times 10^{-23} \mathrm{~J} / \mathrm{K}$ )
[3 Marks]
Ans: 74.65 nm

### 3.5 Pressure of Ideal Gas

9. Prove the relation between pressure of the gas and speed of its molecules.
[4 Marks]
Ans: Refer Q. 14
10. A gas in a cylinder is at pressure $500 \mathrm{~N} / \mathrm{m}^{2}$. If the masses of all the molecules are made one fifth of their original value and their speeds are doubled, then find the resultant pressure.
[2 Marks]
Ans: $400 \mathrm{~N} / \mathrm{m}^{2}$

### 3.6 Root Mean Square (rms) Speed

11. Show that r.m.s. velocity of a gas molecule is directly proportional to the square root of the absolute temperature of the gas.
[2 Marks] [Mar 14]
Ans: Refer Q. 17
12. State the relation between mean square velocity of a gas molecule and its absolute temperature.
[1 Marks]
Ans: Refer Q. 18
13. Compare the speed of sound in a gas and r.m.s. speed of that gas molecule.
[2 Marks]
Ans: Refer Q. 20
14. Calculate the temperature (in ${ }^{\circ} \mathrm{C}$ ) at which the r.m.s velocity of $\mathrm{O}_{2}$ gas molecule will be $0.5 \mathrm{~km} \mathrm{~s}^{-1}$. [Given $\mathrm{R}=8.311 \mathrm{~J} \mathrm{~mol}^{-1} \mathrm{~K}^{-1}$ ]
[2 Marks]
Ans: $47.86{ }^{\circ} \mathrm{C}$
15. Determine the pressure of oxygen at $0^{\circ} \mathrm{C}$, if the density of oxygen at N.T.P is $1.44 \mathrm{~kg} / \mathrm{m}^{3}$ and r.m.s. speed of the molecules at N.T.P is $456.4 \mathrm{~m} / \mathrm{s}$.
[2 Marks]
Ans: $10^{5} \mathrm{~N} / \mathrm{m}^{2}$
16. Calculate the ratio of mean square speeds of molecules of a gas at 30 K and 150 K . [1 Mark]
Ans: 1:5
17. The r.m.s speed of oxygen molecules at N.T.P is $461.2 \mathrm{~m} / \mathrm{s}$. Find the temperature at which the r.m.s speed becomes $90 \%$ of the speed at N.T.P.
[2 Marks]
Ans: 221.13 K

### 3.7 Interpretation of Temperature in Kinetic Theory

18. Explain how kinetic energy of gas molecules varies with the temperature using kinetic theory of gases.
[3 Marks]
Ans: Refer Q. 29
19. Deduce Boyle's law using the expression for pressure exerted by the gas. [2 Marks][Feb 20]
Ans: Refer Q. 30
20. Find kinetic energy of 2.5 litre of a gas at S.T.P. Given standard pressure is $1 \times 10^{5} \mathrm{~N} / \mathrm{m}^{2}$.
[2 Marks]
Ans: 375 J
21. The kinetic energy of 1 kg of oxygen at 300 K is $1.356 \times 10^{6} \mathrm{~J}$. Find the kinetic energy of 4 kg of oxygen at 400 K .
[2 Marks]
Ans: $7.232 \times 10^{6} \mathrm{~J}$
22. Find the average kinetic energy of a molecule of nitrogen at $27{ }^{\circ} \mathrm{C}$. [Boltzmann constant, $\mathrm{k}_{\mathrm{B}}=1.381 \times 10^{-23} \mathrm{~J} /$ molecule K ] [2 Marks]
Ans: $6.215 \times 10^{-21} \mathrm{~J}$
23. Calculate the molecular K.E per k mol and per kg of hydrogen at $127^{\circ} \mathrm{C}$.
$[\mathrm{R}=8310 \mathrm{~J} / \mathrm{k} \mathrm{mol} \mathrm{K}$, molecular weight of hydrogen $=2$ ]
[3 Marks]
Ans: $4.986 \times 10^{6}$ J, $2.493 \times 10^{6} \mathrm{~J}$

### 3.8 Law of Equipartition of Energy

24. What is the law of equipartition of energy?
[3 Marks]
Ans: Refer Q. 36
25. What are degrees of freedom of a system? How many degrees of freedom a body moving in a plane has?
[2 Marks]
Ans: Refer Q. 37 and Q. 38
26. How many degrees of freedom a monatomic gas has?
[1 Mark]
Ans: Refer Q. 40
27. Describe the energy associated with each degree of freedom for a diatomic gas.
[3 Marks]
Ans: Refer Q. 41

### 3.9 Specific Heat Capacity

28. Calculate molar specific heat at constant volume and constant pressure of a diatomic gas.
[2 Marks]
Ans: Refer Q. 44 (ii)
29. Deduce the molar specific heat at constant volume and constant pressure of a monatomic gas.
[2 Marks]
Ans: Refer Q. 44 (i)
30. The difference between the two molar specific heats of a gas is $9000 \mathrm{~J} / \mathrm{kg} \mathrm{K}$. If the ratio of the two specific heats is 1.5 , calculate the two molar specific heats. [2 Marks] [Mar 22]
Ans: $\mathrm{C}_{\mathrm{V}}=1.8 \times 10^{4} \mathrm{~J} / \mathrm{kg} \mathrm{K}, \mathrm{C}_{\mathrm{P}}=2.7 \times 10^{4} \mathrm{~J} / \mathrm{kg} \mathrm{K}$.
31. The ratio of specific heats of a gas is 1.4. Its specific heat at constant volume is $4.96 \mathrm{k} \mathrm{cal} / \mathrm{k} \mathrm{mol} \mathrm{K}$. Calculate the universal gas constant. [ $\mathrm{J}=4200 \mathrm{~J} / \mathrm{k}$ cal]
[2 Marks]
Ans: $8332.8 \mathrm{~J} / \mathrm{k} \mathrm{mol} \mathrm{K}$
32. Specific heat of nitrogen at constant pressure and at constant volume is $0.24 \mathrm{kcal} / \mathrm{kg}-\mathrm{K}$ and $0.17 \mathrm{kcal} / \mathrm{kg}-\mathrm{K}$ respectively. Calculate the mechanical equivalent of heat.
[Molecular weight of nitrogen $=28$, $\mathrm{R}=8.31 \times 10^{3} \mathrm{~J} / \mathrm{k} \mathrm{mol} \mathrm{k} \mathrm{cal]}$ [2 Marks]
Ans: $4.24 \times 10^{3} \mathrm{~J} / \mathrm{K}$

### 3.10 Absorption, Reflection and Transmission of Heat Radiation

33. Derive the relation between $\mathrm{a}, \mathrm{r}$ and $\mathrm{t}_{\mathrm{r}}$. [2 Marks] Ans: Refer Q. 48
34. Define:
i. Coefficient of absorption
ii. Coefficient of transmission
[2 Marks]
Ans: Refer $Q .49$ (i) and (iii)
35. What are athermanous and diathermanous substances?
[2 Marks]
Ans: Refer Q. 50
36. What is coefficient of reflection? When can a body be said to be a perfect reflector? [2 Marks]
Ans: Refer Q. 49 (ii) and Q. 51 (ii)

### 3.11 Perfect Blackbody

37. What is perfectly blackbody?
[1 Mark] [Mar 19, July 22]
Ans: Refer Q. 52
38. Describe Ferry's blackbody with the help of a neat labelled diagram.
[3 Marks]
Ans: Refer Q. 53
39. Draw a neat labelled diagram for Ferry's perfectly blackbody. [2 Marks] [Mar 13; July 18]
Ans: Refer Q. 53 (Diagram only)
40. Explain what is cavity radiator.
[2 Marks]
Ans: Refer Q. 54

### 3.12 Emission of Heat Radiation

41. How does heat exchange takes place according to Prevost's theory?
[3 Marks]
Ans: Refer Q. 57
42. On which factors does amount of heat radiated by a body depend?
[2 Marks]
Ans: Refer Q. 58
43. What is emissive power?
[1 Mark]
Ans: Refer Q. 59
44. Define coefficient of emission.
[1 Mark] [July 22]
Ans: Refer Q. 60
45. The energy of 3000 J is radiated in 2 minutes by a body of surface area $100 \mathrm{~cm}^{2}$. Find emissive power of the body.
[2 Marks]
Ans: $2500 \mathrm{~J} / \mathrm{m}^{2} \mathrm{~s}$

### 3.13 Kirchhoff's Law of Heat Radiation and its Theoretical Proof

46. State Kirchhoff's law of heat radiations.
[1 Mark]
Ans: Refer Q.64(Statement only)
47. Prove Kirchhoff's law of radiation theoretically.
[2 Marks] [July 17]
Ans: Refer Q. 64 (Theoretical proof only)

### 3.14 Spectral Distribution of Blackbody Radiation

48. Explain energy distribution spectrum of a black body radiation in terms of wavelength.
[3 Marks] [Feb 20]
Ans: Refer Q. 65
49. For a perfectly blackbody at temperature of 4000 K , find the value of $\lambda_{\max }$.
(Take $\mathrm{b}=2.897 \times 10^{-3} \mathrm{~m} \mathrm{~K}$ )
[2 Marks]
Ans: $7242.5 \AA$
50. Calculate the value of $\lambda_{\text {max }}$ for radiations coming from a star with surface temperature of 6500 K . ( $b=2.897 \times 10^{-3} \mathrm{~m} \mathrm{~K}$ )
[2 Marks]
Ans: $4457 \AA$
51. Find the temperature of a blackbody if its spectrum has a peak at $\lambda_{\text {max }}=500 \mathrm{~nm}$ (visible)
[2 Marks]
Ans: 5794 K

### 3.15 Stefan-Boltzmann Law of Radiation

52. What does Stefan-Boltzmann law state?
[1 Mark]
Ans: Refer Q. 71
53. Obtain the expression for the rate of loss of heat by a blackbody in cooler surroundings. [2 Marks]
Ans: Refer Q. 73
54. Calculate the energy radiated in half a minute by a black body of surface area $200 \mathrm{~cm}^{2}$ at $127^{\circ} \mathrm{C}$.
[3 Marks] [July 22]
Ans: 875.5 J
55. A body of surface area $20 \mathrm{~cm}^{2}$ and temperature $527^{\circ} \mathrm{C}$ emits 400 J of energy per minute. Find its emissivity.
[Given: $\sigma=5.67 \times 10^{-8} \mathrm{watt} / \mathrm{m}^{2} \mathrm{~K}^{4}$ ] [2 Marks]
Ans: 0.1435
56. Compare the rates of emission of heat by a blackbody maintained at $327{ }^{\circ} \mathrm{C}$ and at $127^{\circ} \mathrm{C}$, if the blackbodies are surrounded by an enclosure (black) at $27^{\circ} \mathrm{C}$. What would be the ratio of their rates of loss of heat? [3 Marks]
Ans: $243: 35$ or $6.94: 1$

## Multiple Choice Questions

[1 Mark Each]
*1. In an ideal gas, the molecules possess
(A) only kinetic energy
(B) both kinetic energy and potential energy
(C) only potential energy
(D) neither kinetic energy nor potential energy
2. In the case of ideal gases,
(A) the molar specific heat at constant pressure is the same for all gases.
(B) the molar specific heat at constant volume is the same for all gases.
(C) the ratio of the molar specific heats at constant volume and at constant pressure is the same for all gases.
(D) the difference between the molar specific heats at constant pressure and at constant volume is the same for all gases.
*3. The mean free path $\lambda$ of molecules is given by
(A) $\sqrt{\frac{2}{\pi n d^{2}}}$
(B) $\frac{1}{\pi \mathrm{nd}^{2}}$
(C) $\frac{1}{\sqrt{2} \pi \mathrm{nd}^{2}}$
(D) $\frac{1}{\sqrt{2 \pi n d}}$
where n is the number of molecules per unit volume and $d$ is the diameter of the molecules.
*4. If pressure of an ideal gas is decreased by $10 \%$ isothermally, then its volume will
(A) decrease by $9 \%$
(B) increase by $9 \%$
(C) decrease by $10 \%$
(D) increase by $11.11 \%$
5. The average distance covered by a molecule between two successive collision is $\qquad$ .
(A) free path
(B) constant path
(C) mean free path
(D) free path per unit time.
6. ' P ' is the pressure and ' d ' is the density of gas at constant temperature, then
(A) $P \propto d$
(B) $\quad \mathrm{P} \propto 1 / \mathrm{d}$
(C) $P \propto \sqrt{d}$
(D) $\mathrm{P} \propto 1 / \sqrt{\mathrm{d}}$
7. The expression for pressure of gas and the gas equation shows that the absolute temperature of a gas is proportional to the average
(A) sum of vibrational, translational and rotational kinetic energies of molecules.
(B) translational kinetic energy of molecules.
(C) rotational kinetic energy of molecules.
(D) vibrational kinetic energy of molecules.
8. The root mean square speed of the molecules of a gas is proportional to $\qquad$ -.
( $\mathrm{T}=$ Absolute temperature of gas)
(A) $\sqrt{T}$
(B) $\frac{1}{\sqrt{\mathrm{~T}}}$
(C) T
(D) $\frac{1}{\mathrm{~T}}$
9. The absolute temperature of a gas increased three times. The root mean square velocity of the gas will be $\qquad$ -.
(A) 3 times
(B) 9 times
(C) $\frac{1}{3}$ times
(D) $\sqrt{3}$ times
10. The kinetic energy per molecule of a gas at temperature T is $\qquad$ [Mar 18]
(A) $\left(\frac{3}{2}\right) \mathrm{RT}$
(B) $\left(\frac{3}{2}\right) \mathrm{k}_{\mathrm{B}} \mathrm{T}$
(C) $\quad\left(\frac{2}{3}\right) \mathrm{RT}$
(D) $\left(\frac{3}{2}\right)\left(\frac{\mathrm{RT}}{\mathrm{M}}\right)$
11. The average K.E. of a gas is $\qquad$ . [July 22]
(A) directly proportional to absolute temperature of gas
(B) directly proportional to square of absolute temperature of gas
(C) directly proportional to square root of absolute temperature of gas
(D) inversely proportional to absolute temperature of gas
12. If the total kinetic energy per unit volume of gas enclosed in a container is $E$, the pressure exerted by the gas is $\qquad$ -.
[July 17]
(A) E
(B) $\frac{3}{2} \mathrm{E}$
(C) $\quad \sqrt{3} \mathrm{E}$
(D) $\frac{2}{3} \mathrm{E}$
13. The root mean square velocity of gas molecules is $10 \mathrm{~km} \mathrm{~s}^{-1}$. The gas is heated till its pressure becomes 4 times. The velocity of the gas molecules will now be
(A) $10 \mathrm{~km} \mathrm{~s}^{-1}$
(B) $20 \mathrm{~km} \mathrm{~s}^{-1}$
(C) $40 \mathrm{~km} \mathrm{~s}^{-1}$
(D) $80 \mathrm{~km} \mathrm{~s}^{-1}$
14. The law of equipartition of energy is valid for
(A) high temperatures
(B) extremely low temperatures
(C) only absolute zero temperature
(D) temperatures within the range -273 K to -100 K .
15. For hydrogen gas $\mathrm{C}_{\mathrm{P}}=4000 \mathrm{cal} / \mathrm{kg} \mathrm{K}$ and $\mathrm{C}_{\mathrm{V}}=3000 \mathrm{cal} / \mathrm{kg} \mathrm{K}$ and $\mathrm{R}=8300 \mathrm{~J} / \mathrm{k} \mathrm{mol} \mathrm{K}$.
The value of J will be [mol. wt. of $\mathrm{H}_{2}=2$ ]
(A) 4.18
(B) 4.17
(C) 4.16
(D) 4.15
16. According to the law of equipartition of energy, the average kinetic energy of one molecule of diatomic gas will be
(A) $3 \mathrm{k}_{\mathrm{B}} \mathrm{T} / 2$
(B) $5 \mathrm{k}_{\mathrm{B}} \mathrm{T} / 2$
(C) $3 \mathrm{RT} / 2$
(D) $5 \mathrm{RT} / 2$
17. If the degrees of freedom for polyatomic gas are f, then the average kinetic energy per molecule of the gas will be [ $\mathrm{N}_{\mathrm{A}}$ : Avogadro's number]
(A) $\mathrm{fk}_{\mathrm{B}} \mathrm{T} / \mathrm{N}$
(B) $\mathrm{fk}_{\mathrm{B}} \mathrm{T} / 2 \mathrm{~N}$
(C) $\mathrm{fk}_{\mathrm{B}} \mathrm{T} / 2$
(D) $\quad \mathrm{fk}_{\mathrm{B}} \mathrm{T}$
18. For polyatomic molecules having ' f ' vibrational modes, the ratio of two specific heats, $\frac{C_{P}}{C_{V}}$ is
$\qquad$ .
[Mar 16]
(A) $\frac{1+\mathrm{f}}{2+\mathrm{f}}$
(B) $\frac{2+\mathrm{f}}{3+\mathrm{f}}$
(C) $\frac{4+\mathrm{f}}{3+\mathrm{f}}$
(D) $\frac{5+\mathrm{f}}{4+\mathrm{f}}$
19. Coefficient of emission or emissivity (e) is defined as
(A) ratio of emissive power of a surface to that of a perfectly black surface at the same temperature.
(B) product of the emissive powers of the surface and perfectly blackbody at the same temperature.
(C) ratio of emissive power of the surface to that of perfectly blackbody.
(D) product of emissive powers of the surface and perfectly blackbody.
20. If the incident energy is 200 cal , absorptive power is 0.31 and reflection coefficient is 0.41 , then the amount of energy transmitted will be
(A) 48 cal
(B) 56 cal
(C) 58 cal
(D) 54 cal
21. A body which absorbs all the radiations incident over it is called a
(A) blackbody.
(B) perfectly blackbody.
(C) good absorber.
(D) good emitter.
22. The best ideal blackbody is
(A) lamp of charcoal heated to a high temperature.
(B) metal coated with a black dye.
(C) glass surface coated with coal tar.
(D) hollow enclosure blackened inside and having a small hole.
*23. If $a=0.72$ and $r=0.24$, then the value of $t_{r}$ is
(A) 0.02
(B) 0.04
(C) 0.4
(D) 0.2
24. The amount of thermal radiation emitted per unit time depends on
(A) nature of emitting surface
(B) temperature of emitting surface
(C) area of emitting surface
(D) all of these
25. Find the wavelength at which a blackbody radiates maximum energy, if its temperature is $427^{\circ} \mathrm{C}$. [Wien's constant $\mathrm{b}=2.898 \times 10^{-3} \mathrm{mK}$ ]
[Mar 17]
(A) $0.0414 \times 10^{-6} \mathrm{~m}$
(B) $4.14 \times 10^{-6} \mathrm{~m}$
(C) $41.4 \times 10^{-6} \mathrm{~m}$
(D) $414 \times 10^{-6} \mathrm{~m}$
26. "Good absorbers are good emitters" This statement is of
(A) Newton's law
(B) Kirchhoff's law
(C) Stefan's law
(D) Wien's law
27. Two stars emit maximum radiation at wavelength $3600 \AA$ and $4800 \AA$ respectively. The ratio of their temperature is
(A) $1: 2$
(B) $3: 4$
(C) $4: 3$
(D) $2: 1$
28. The mathematical relation of Stefan's law is
(A) $\mathrm{Q}=\sigma \mathrm{T}^{2}$
(B) $\mathrm{Q}=\sigma \mathrm{T}^{3}$
(C) $\mathrm{Q}=\sigma \mathrm{T}^{4}$
(D) $\mathrm{Q}=\sigma \mathrm{T}^{5}$
*29. The ratio of emissive power of perfectly blackbody at $1327^{\circ} \mathrm{C}$ and $527^{\circ} \mathrm{C}$ is
(A) $4: 1$
(B) $16: 1$
(C) $2: 1$
(D) $8: 1$
30. If wavelengths of maximum intensity of radiations emitted by the Sun and the moon are $0.5 \times 10^{-6} \mathrm{~m}$ and $10^{-4} \mathrm{~m}$ respectively, the ratio of their temperature is
(A) $1 / 100$
(B) $1 / 200$
(C) 100
(D) 200
31. The temperature of a blackbody is increased by $50 \%$. The amount of radiations emitted by the body increases by
(A) $50 \%$
(B) $100 \%$
(C) $225 \%$
(D) $500 \%$
32. The surface area of a blackbody maintained at $127^{\circ} \mathrm{C}$ radiating energy at the rate of $1459.2 \mathrm{~J} / \mathrm{s}$ is $\qquad$ . $\left[\sigma=5.7 \times 10^{-8}\right.$ S.I. unit $]$
(A) $4 \mathrm{~m}^{2}$
(B) $3 \mathrm{~m}^{2}$
(C) $2 \mathrm{~m}^{2}$
(D) $1 \mathrm{~m}^{2}$
33. The SI unit of Stefan's constant is
(A) $\mathrm{N} \mathrm{m} / \mathrm{s} \mathrm{K}^{4}$
(B) $\mathrm{W} / \mathrm{m} \mathrm{K}^{4}$
(C) $\mathrm{J} / \mathrm{s} \mathrm{m}^{2} \mathrm{~K}^{4}$
(D) $\quad \mathrm{erg} / \mathrm{s} \mathrm{m}^{3} \mathrm{~K}^{4}$

## Answers to Multiple Choice Questions

| 1. | (A) | 2. | (D) | 3. | (C) | 4. | (D) |
| :--- | :--- | :--- | :--- | :--- | :--- | :--- | :--- |
| 5. | (C) | 6. | (A) | 7. | (B) | 8. | (A) |
| 9. | (D) | 10. | (B) | 11. | (A) | 12. | (D) |
| 13. | (B) | 14. | (A) | 15. | (D) | 16. | (B) |
| 17. | (C) | 18. | (C) | 19. | (A) | 20. | (B) |
| 21. | (B) | 22. | (D) | 23. | (B) | 24. | (D) |
| 25. | (B) | 26. | (B) | 27. | (C) | 28. | (C) |
| 29. | (B) | 30. | (D) | 31. | (D) | 32. | (D) |
| 33. | (C) |  |  |  |  |  |  |

## Hints to Multiple Choice Questions

4. From ideal gas equation,
$\mathrm{PV}=\mathrm{nRT}$
For isothermal process, $\mathrm{T}=$ constant
$\therefore \quad \mathrm{PV}=$ constant
$\therefore \quad$ When pressure is decreased by $10 \%$,
$\mathrm{P}^{\prime}=\mathrm{P}-\frac{10}{100} \mathrm{P}$
$\therefore \quad \mathrm{P}^{\prime}=0.9 \mathrm{P}$
Now, $\mathrm{P}^{\prime} \mathrm{V}^{\prime}=\mathrm{PV}$
...[from (i)]
$\therefore \quad \frac{\mathrm{V}^{\prime}}{\mathrm{V}}=\frac{\mathrm{P}}{\mathrm{P}^{\prime}}=\frac{1}{0.9}$
...[from (ii)]
$\therefore \quad \frac{\mathrm{V}^{\prime}-\mathrm{V}}{\mathrm{V}}=\frac{0.1}{0.9}$.
$\therefore \quad \frac{\mathrm{V}^{\prime}-\mathrm{V}}{\mathrm{V}} \times 100=0.1111 \times 100=11.11 \%$
As, in isothermal process
$\mathrm{P} \propto \frac{1}{\mathrm{~V}}$
V will increase by $11.11 \%$
5. Law of equilibrium of energy cannot be applied where quantum effects become important.
6. $a+r+t_{r}=1$
$\therefore \quad 0.72+0.24+\mathrm{t}_{\mathrm{r}}=1$
$\therefore \quad \mathrm{t}_{\mathrm{r}}=1-0.96=0.04$
7. According to Wien's displacement law,
$\lambda_{\mathrm{m}}=\frac{\mathrm{b}}{\mathrm{T}}=\frac{2.898 \times 10^{-3}}{700}=4.14 \times 10^{-6} \mathrm{~m}$
8. Power $(P) \propto T^{4}$
$\therefore \quad \frac{\mathrm{P}_{1}}{\mathrm{P}_{2}}=\left(\frac{\mathrm{T}_{1}}{\mathrm{~T}_{2}}\right)^{4}=\frac{(1327+273)^{4}}{(527+273)^{4}}=\left(\frac{1600}{800}\right)^{4}=16: 1$

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