

## Chapter # 4

### STATE OF MATTER

#### GASES

**Q1. How many states of matter exist in nature?**

**Ans:** Matter consists of three fundamental states, the gases, liquid and solid. Besides these three states, the fourth one is plasma state (a partially ionized gas at very high temperature)

**Q2. Define and explain the term Brownian movement.**

**Ans:** The word gas is derived from chaos. This is because the gaseous particles have chaotic (random) motion. The molecules travel in a straight line until they collide with each other or with the walls of the container. Such motion of the molecules is called Brownian Movement (Robert Brown 1827).

**Q3. How can you explain the effect of pressure on gas molecules?**

**Ans:** In a gas the average distance between the molecules is quite large as compared to their diameters. This shows that the intermolecular forces among the molecules are negligible at the lowest pressure. However, these forces increase with the increase of pressure. For example, the forces are negligible at one atmosphere (lowest) significant at about 10 atmospheres and very important at higher pressures (about 30 atmospheres). It is found by experiment that when the pressure is sufficiently low, all gases satisfy the perfect gas law  $PV = nRT$

**Q4. Earth's atmosphere is a mixture of gases how can you explain it by giving daily life examples?**

**Ans:** The earth's atmosphere is a mixture of gases. Oxygen (21%) and Nitrogen (78%) are its two major components, both support life. In fact, the earth's atmosphere is an immense system of gases where many chemical reactions occur. For example, photo-synthesis, combustion etc. Natural gas, the major component of which is methane, is used to heat homes in winter, to cook food and to drive vehicles etc.

**Kinetic Molecular Theory of Gases:**

The behavior and properties of gases can be theoretically explained using the Kinetic Molecular Theory. This theory is a model that explains the behavior of gases using generalization about random moving molecules within a gas.

The Kinetic Molecular Theory was first postulated by Daniel Bernoulli, a Swiss mathematician. The Kinetic Theory of gases was developed by a number of well-known physicists such as James Maxwell (1859) and Boltzmann in (1870). The research was contained on the Kinetic Theory making it a cornerstone of our present view of molecular substances. In 1857, Clausius derived the Kinetic equation

Physical theories are often given in terms of postulates, the basic statements from which all conclusions or prediction of theory is deduced.

#### **Q5. Write and explain kinetic equation?**

**Ans: Kinetic equation:**

R J Clausius deduced an equation for the pressure of an ideal gas from the postulates of kinetic theory. This equation is called kinetic equation.

$$\text{Mathematically, } pv = \frac{1}{3}mnc^2$$

Where  $p$  = Pressure

$v$  = volume

$m$  = mass of one molecule of the gas

$n$  = number of moles of the gas

$c^2$  = mean square velocity of gas molecules

#### **Mean square velocity:**

Since the gas molecules under the given condition do not have the same velocity, so mean square velocity is considered. If  $n_1$  molecules have velocity  $c_1$ ,  $n_2$  molecules have velocity  $c_2$  and so on. Then,

$$c^{\bar{}} = \frac{n_1 c_1^2 + n_2 c_2^2 + \dots}{n_1 + n_2 + \dots}$$

Where  $c^{\bar{}}$  is the average of all the possible velocities

Root mean square velocity:

The square root of is called root mean square velocity.

**Q6. How many types of motion occur between gas molecule also elaborate these motions?**

**Ans: Motion of Particles of a Gas:**

In gases, the molecular motion is of three types

- I. Translational Motion
- II. Rotational M.
- III. Vibrational Motion

A mono atomic molecule (0.9. He) will show translational motion while a diatomic ( $H_2$ ) and polyatomic molecules ( $CO_2$ ,  $NH_3$ , etc.) will undergo, in addition to translational motion, the rotational and vibrational motions too.

**I. Translational motion**

The motion imparted to the gaseous molecules due to their motion in all possible directions is called translational motion and the energy as kinetic translation energy. In this case the entire molecules move from place to place.

**II. Rotational Motion:**

The motion imparted to the gaseous molecules as a result of net angular momentum about their center of gravity is called rotational motion and the energy as kinetic rotational energy. In this case the molecule spins like a propeller.

**III. Vibrational Motion:**

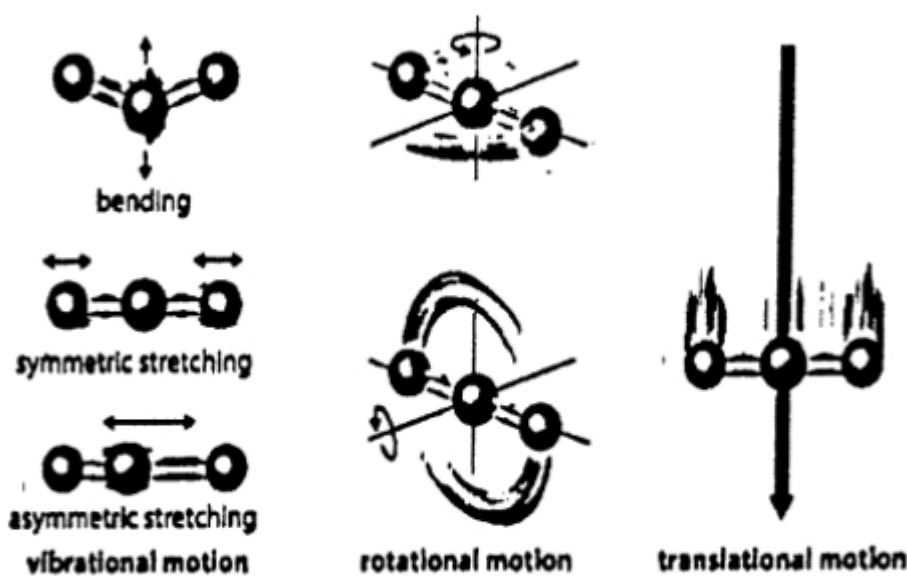


The motion imparted to the gaseous molecules due to the oscillations is called vibrational motion and the energy as kinetic vibrational energy. In this case the molecules vibrate back and forth about the same fixed location.

In such an oscillating system, there is a continuous Interchange between vibrational kinetic energies and potential energies. As a result, both KE and PE is possessed by the gaseous molecules Thus,

$$\text{Total KE} = \text{KE (Vib)} + \text{PE (Vib)}$$

The vibrational motions in polyatomic molecules (like  $\text{CO}_2$ ,  $\text{NH}_3$  etc) are quite complicated



**Figure 4.4 Different motions of molecules**

**Example 1:**

Convert the following Celsius temperatures to Kelvin temperatures:

- (a)  $-132^{\circ}\text{C}$    (b)  $96^{\circ}\text{C}$    (c)  $0^{\circ}\text{C}$    (d)  $-12^{\circ}\text{C}$    (e)  $148^{\circ}\text{C}$

**Solution:**

Using the formula:  $K = ^{\circ}\text{C} + 273$

(a)  $\text{TK} = -132 + 273 = 141 \text{ TK}$

(b)  $\text{TK} = 98^{\circ}\text{C} + 273 = 369 \text{ TK} \times$

(c)  $\text{TK} = 0^{\circ}\text{C} + 273 = 273 \text{ TK}$

(d)  $\text{TK} = -12^{\circ}\text{C} + 273 = 261 \text{ TK}$

(e)  $\text{TK} = 148^{\circ}\text{C} + 273 = 421 \text{ TK}$

**Example 2:**

Convert the following Kelvin temperatures to Celsius degree centigrade temperatures.

(a) 340 TK    (b) 200 TK    (c) 10TK    (d) 405 TK

**Solution:**

Using the formula:  $^{\circ}\text{C} = \text{TK} - 273$

$$(a) \text{ } t^{\circ}\text{C} = 340 \text{ TK} - 273 = 67^{\circ}\text{C}$$

$$(b) \text{ } t^{\circ}\text{C} = 200 \text{ TK} - 273 = -73^{\circ}\text{C}$$

$$(c) \text{ } t^{\circ}\text{C} = 10 \text{ TK} - 273 = -263^{\circ}\text{C}$$

$$(d) \text{ } t^{\circ}\text{C} = 405 \text{ TK} - 273 = 132^{\circ}\text{C}$$

**Q7. What is the total Kinetic Energy of translation in ergs of two moles of perfect gas at 27°C in calories?**

**Ans: Solution:**

$$\text{No of moles of perfect gas (n)} = 2$$

$$T = 27^{\circ}\text{C} + 273 = 300\text{K}$$

$$R = 8.314 \times 10^7 \text{ ergs mole}^{-1}\text{K}^{-1}$$

$$\text{Now, K.E of translation (K.E)} = \frac{3}{2}n RT$$

$$\text{or } \text{K.E} = \frac{3}{2} \times 2 \text{ moles dm}^{-3} \times 8.314 \times 10^7 \text{ ergs mole}^{-1}\text{K}^{-1} \times 300\text{K}$$

$$\text{K.E} = \frac{3}{2} \times 2 \times 8.314 \times 10^7 \times 300 \text{ moles dm}^{-3} \times \text{ergs mole}^{-1} \times \text{K}^{-1} \times \text{K}$$

$$= 7483.5 \times 10^7 \text{ ergs dm}^{-3}$$

**Q8. Briefly explain gas pressure in the Light of Kinetic Molecular Theory.**

**Ans: Gas Pressure in the Light of Kinetic Molecular Theory:**

According to the kinetic molecular theory, gas molecules are in constant random motion. They move in straight line until they collide with another molecule or the walls of the container. The pressure of a gas in a container is due to the force exerted by its molecules on the walls of the container. The average kinetic energy of gas molecules is directly proportional to the Kelvin

temperature. So, the average kinetic energy of a collision when a gas molecule collides with the walls of a container will not change at constant temperature.

**Q9. Define gaseous pressure and its various units.**

**Ans: Pressure:**

A force being exerted over a unit area is called Pressure

So,  $P = \frac{F}{A}$  where P is measured in Newton Force in joules and the area in  $m^2$

**Pascal (Pa):**

A Pascal (Pa) is defined as the force of one Newton (N) spread over an area of  $1m^2$ .

**Units of Pressure:**

**SI Units:**

SI unit for pressure is Pascal (Pa), in S.I. system, unit of force is Newton (N) and the unit of area is metre square ( $m$ ). Thus Pascal is one  $Nm$ .

$$1 \text{ atm} = 101325\text{Pa} = 101325 \text{ Nm}^2$$

$$1 \text{ atm} = 14.7 \text{ Psi (Pounds per square inch)}$$

$$1 \text{ atm} = 101325\text{Pa}$$

$$1 \text{ atm} = 101.325 \text{ KPa}$$

$$1 \text{ atm} = 760 \text{ torr} = 760\text{mm of Hg}$$

$$1\text{J} = 1\text{Nm} = 10^7\text{ergs} = \text{kgm}^2\text{s}^{-2}$$

$$1 \text{ Cal} = 4.18\text{J}$$

Interconversion of Pressure in Pascals, Kilo Pascals, atmosphere and bar.

**Example 4:**

**Convert 10 Pascals into (1) Kilo Pascals (ii) atmosphere (iii) bars**

**Solution:**

I. One Kilo Pascal = 1000 Pascals

or 1000 Pascals = one Kilo Pascal

$$10 \text{ Pa} = \frac{1}{1000} \times 10 = 0.01 \text{ Kilo Pascal}$$

II. One atmosphere = 101325 Pa

or 101325 Pa = one atm

$$10 \text{ Pa} = \frac{1}{101325} \times 10 = 0.000098 \text{ atm}$$

**Activity for Students**

**Convert 35 bars into (i) Pascal (ii) Kilo Pascal (iii) Atmosphere**

**Solution:**

i. bar = 101325 pa

$$1 \text{ bar} = \frac{101325}{1.01325} \text{ Pa}$$

$$35 \text{ bar} = \frac{101325}{1.01325} \times 35 = 35 \times 10^6 \text{ pa}$$

ii. 1.01325 bar = 101.325 kpa

$$1 \text{ bar} = \frac{101.325}{1.01325} \text{ kPa}$$

$$35 \text{ bar} = \frac{101.325}{1.01325} \times 35 = 3.5 \times 10^3 \text{ kpa}$$

iii. 1.01325 bar = 1 atm

$$35 \text{ bar} = \frac{1}{1.01325} \times 35 = 34.54 \text{ atm}$$

**Q10. Briefly explain Boyle's Law. And why the Isotherm goes away from both the axis in the graphical explanation of Boyle's Law.**

**Ans: Boyle's Law:**

It states that the pressure of a fixed amount of a gas varies inversely with its volume, when the temperature is maintained constant.

**Mathematical form of Boyle's Law:**

Mathematically,  $P \propto \frac{1}{V}$  or  $P = K \times \frac{1}{V}$  where K is proportionality constant

The value of K is different for different amounts of the same gas

Again  $P = K \times \frac{1}{V}$  or  $PV = K$ , thus the product of volume and Pressure remains constant provided the temperature is kept constant.

$$P_1 V_1 = K \text{ (For gas at pressure } P_1)$$

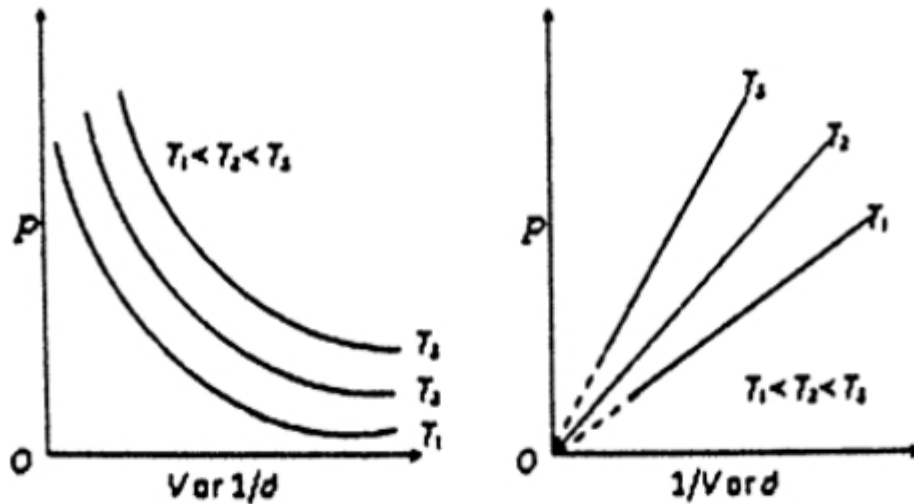
$$P_2 V_2 = K \text{ (For gas at pressure } P_2)$$

Therefore,  $P_1 V_1 = P_2 V_2$  where  $V_1$  and  $P_1$  are initial volume and Pressure while  $P_2 V_2$  are their final Pressure and volume.

**Graphical Representation of Boyle's Law:**

**Isotherm:**

When the Pressure of a gas is plotted against volume at different temperatures, we get a family of curves as shown in the figure. Each curve is a hyperbola with different values of K. Each curve is known as Isotherm (constant temperature plot). As the temperature is increased, the Isotherm goes away from both the axis. This is because at higher temperature, the volume of the gas is increased.



**Q11. Example 5:** An ideal gas occupies a volume of  $0.300 \text{ dm}^3$  at a pressure of  $1.80 \times 10^5 \text{ Pa}$ . What is the volume of the gas maintained at the same temperature if the pressure is reduced to  $1.15 \times 10^5 \text{ Pa}$ .

**Solution:**

$$V_1 = 0.300 \text{ dm}^3 \quad V_2 = ?$$

$$P_1 = 1.80 \times 10^5 \text{ Pa} \quad P_2 = 1.15 \times 10^5 \text{ Pa}$$

According to the Boyle's Law

$$V_1 P_1 = V_2 P_2$$

$$V_2 = \frac{V_1 P_1}{P_2}$$

$$\frac{0.300 \text{ dm}^3 \times 1.80 \times 10^5 \text{ Pa}}{1.15 \times 10^5 \text{ Pa}}$$

$$\frac{0.300 \times 1.80}{1.15} \text{ dm}^3 = 0.469 \text{ dm}^3$$

**Q12. Example 6:** At sea level, where the pressure was 775 torr, the gas in a balloon occupied  $2 \text{ m}^3$ . What volume will the balloon expand when it has risen to an altitude where the pressure is  
 (a) 100 torr (b) 10 torr (temperature remains constant)

**Solution:**

(a)  $P_1 = 755 \text{ torr}$   $P_2 = 100 \text{ torr}$

$$V_1 = 2\text{m}^3 \quad V_2 = ?$$

According to the Boyle's Law

$$V_1 P_1 = V_2 P_2$$

$$2\text{m}^3 \times 755 \text{ torr} = V_2 \times 100 \text{ torr}$$

Or

$$V_2 = \frac{2\text{m}^3 \times 755\text{torr}}{100\text{torr}}$$
$$= \frac{2 \times 755}{100} \text{m}^3 = 15.10\text{m}^3$$

(a)  $P_1 = 755 \text{ torr}$   $P_2 = 10 \text{ torr}$

$$V_1 = 2\text{m}^3 \quad V_2 = ?$$

According to the Boyle's Law

$$V_1 P_1 = V_2 P_2$$

$$2\text{m}^3 \times 755 \text{ torr} = V_2 \times 10 \text{ torr}$$

Or

$$V_2 = \frac{2\text{m}^3 \times 755\text{torr}}{10\text{torr}}$$
$$= \frac{2 \times 755}{10} \text{m}^3 = 151\text{m}^3$$

### Activity for Student

**375cm<sup>3</sup> of a given gas has a pressure of 770mm of Hg. What will be the volume if the pressure is reduced to 750mm of Hg? (Ans: 385 cm)**

**Solution:**

$$P_1 = 770 \text{ mm of Hg} \quad P_2 = 750 \text{ mm of Hg}$$

$$V_1 = 375 \text{ cm}^3 \quad V_2 = ?$$



According to the Boyle's law

$$P_1V_1 = P_2V_2$$

$$375 \times 770 = V_2 \times 750$$

$$V_2 = \frac{375 \times 770}{750} = 385 \text{ cm}^3$$

**Q13. Briefly explain Charles's Law.**

**Ans: Charles's Law:**

It states that at constant pressure, the volume of a given mass of a gas varies linearly with the absolute temperature of the gas

**Critical definition:**

Charles's law can also be stated as "at constant pressure the volume of given mass of gas increases or decreases by  $\frac{1}{273}$  times of its original volume at  $0^\circ\text{C}$  for every  $1^\circ\text{C}$  rise or fall in temperature respectively

From his preliminary investigations, Charles observed that the gases such as  $\text{H}_2$ ,  $\text{CO}_2$ ,  $\text{O}_2$ , etc. expanded equally on heating from  $0$  to  $80^\circ\text{C}$  at constant pressure

**Derivation of critical form of Law:**

Suppose the volume of a gas at  $0^\circ\text{C} = V_0$

$$\text{Volume at } 1^\circ\text{C} = V_0 + V_0 \frac{1}{273}$$

$$\text{Volume at } 2^\circ\text{C} = V_0 + V_0 \frac{2}{273}$$

$$\text{Volume at } t^\circ\text{C} = V_0 + V_0 \frac{t}{273}$$

$$V_t = V_0 \left[ 1 + \frac{t}{273} \right] = V_0 \left[ \frac{273 + t}{273} \right]$$

$$= V_0 \left[ \frac{T}{273} \right]$$

Where  $t^\circ\text{C} + 273 = T$  (Absolute temperature)

$$\text{So } V_t = \frac{V_o}{273} T$$

**Another definition:**

Thus the Law states that the volume of a definite quantity of a gas at constant pressure is directly proportional to the absolute temperature

Again  $V \propto T$  or  $V = KT$ , where T is the temperature on absolute scale.

Thus the Law may also be defined as "the ratio between volume and temperature is always a constant quantity

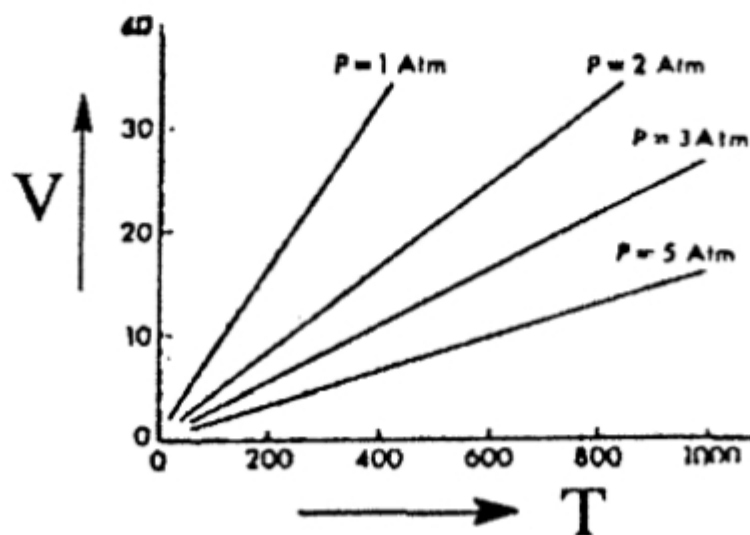
$$\text{So, } \frac{V_1}{T_1} = K \text{ and } \frac{V_2}{T_2} = K$$

$$\frac{V_1}{T_1} = \frac{V_2}{T_2}$$

Where  $V_1$  and  $T_1$  are initial volume and absolute temperature.  $V_2$ ,  $T_2$  are final volume and absolute temperature

**Graphical Representation of Charles's Law:**

The equation for Charles's Law gives a straight line proved by plotting a graph between volume and temperature. Different straight lines are obtained with different pressures, which are a constant quantity. Each constant pressure line is called an "Isobar".



**Graphical verification of Charles law**

**Q14. How centigrade or Celsius scale can be converted to the absolute scale or Kelvin scale?**

**Ans:** The temperature on degree centigrade or Celsius scale can be converted to the absolute scale or Kelvin scale (after the name lord Kelvin 1824-1907) by adding 273.15.

Thus  $K = t^{\circ}\text{C} + 273.15$

Now  $V = KT$       or  $\frac{V}{T} = K$

**Q15. What is absolute zero? Explain it graphically also write its significance.**

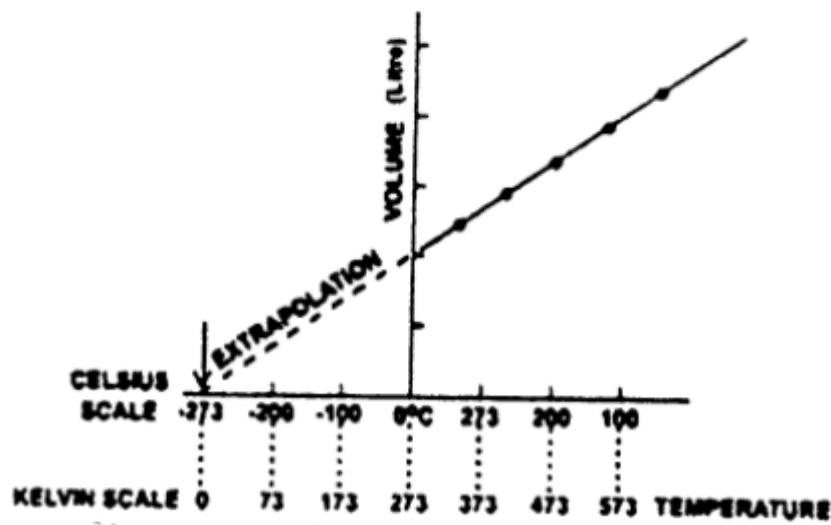
**Ans: Absolute Zero:**

According to critical definition of Charles's Law, at constant pressure, the volume of given mass of a gas increases or decreases by  $\frac{1}{273}$  of its original volume at  $0^{\circ}\text{C}$  by  $1^{\circ}\text{C}$  rise or fall of temperature.

Let the volume of an ideal gas at  $0^{\circ}\text{C}$  by  $V_0 \text{ cm}^3$

Volume at  $-1^{\circ}\text{C} = V_0 - \frac{1}{273}V_0$

Volume at  $-273^{\circ}\text{C} = V_0 - \frac{273}{273}V_0 = 0$



Effect of decrease of temp on a definite volume of a gas at constant pressure

Thus at (exact – 273°C), the volume of a given mass of a gas reduces to zero. A graph between volume and temperature gives a straight line and cuts the temperature axis at 273.15°C.

Thus the temperature at which the given volume of a gas reduces to zero is called Absolute zero for this reason a new temperature scale has been developed.

This new scale is called Absolute or Kelvin scale. It starts from -273.15°C which is marked as zero Kelvin. - 273.15°C = Zero Kelvin

This is equal for all other gases as well and is the lowest possible in the gaseous state but actually it does not happen because all the gases liquefy or solidify before they reach this temperature (-273.15°C) This temperature is considered as the lowest possible temperature.

### Significance of Absolute Zero:

- I. It is not possible to have a gas at Absolute zero because it changes into liquid or solid states
- II. At -273.15°C, the gas particles do not move and volume of gas becomes minimum.
- III. Absolute zero may be initialized to get the Kelvin scale of temperature
- IV. Zero Kelvin (-273.16°C) is called as Absolute zero which is the lowest possible temperature.

**Q16. Example 7: If 50 cm<sup>3</sup> of a gas in a syringe at 15°C is heated to 50°C and the piston of syringe is allowed to move outwards against constant atmospheric pressure, calculate the new volume of the hot gas.**

**Solution:**

$$V_1 = 50\text{cm}^3 \quad V_2 = ?$$

$$T_1 = 15^\circ\text{C} + 273 = 288\text{K} \quad T_2 = 50^\circ\text{C} + 273 = 323\text{K}$$

According to Charles's Law

$$\begin{aligned} \frac{V_1}{T_1} &= \frac{V_2}{T_2} \text{ or } \frac{50\text{cm}^3}{288\text{K}} = \frac{V_2}{322} \text{ or } \frac{50 \times 323}{288} \times \frac{\text{cm}^3 \times \text{K}}{\text{K}} \\ &= 56\text{cm}^3 \end{aligned}$$

Therefore, the new volume of the hot gas = 56cm<sup>3</sup>

**Q17. Example 8: At 17°C, a sample of H<sub>2</sub>, gas occupies 125 cm<sup>3</sup>. What would be the volume at 100°C? (Pressure remains constant)**

**Solution:**

$$V_1 = 125\text{cm}^3 \quad V_2 = ?$$

$$T_1 = 17^\circ\text{C} + 273 \quad T_2 = 100^\circ\text{C} + 273$$

$$= 290\text{K} \quad = 373\text{K}.$$

According to Charles's Law

$$\frac{V_1}{T_1} = \frac{V_2}{T_2} \text{ or } \frac{125\text{cm}^3}{290\text{K}} = \frac{V_2}{373\text{K}} \text{ or } \frac{125 \times 373}{290} \times \frac{\text{cm}^3 \times \text{K}}{\text{K}}$$

Thus the new volume will be 161 cm<sup>3</sup>

### Activity for Student

**A Volume-Temperature experiment which was performed on a sample of M gas produced the Charles's Law Relationship  $V = 0.167 T$  (V is in dm<sup>3</sup>). At what temperature in degree Celsius would this sample of gas occupy 50 dm<sup>3</sup>, if the pressure remains constant? (Ans: 26.40°C)**

**Solution: As given relation**

$$V = 0.167 T$$

Putting the  $V = 50 \text{ dm}^3$

$$\frac{50}{0.167} = T$$

$$299.4 \text{ K} = T$$

$$t = 299.4 - 273$$

$$t = 26.40 \text{ }^{\circ}\text{C}$$

**Q18. Explain Avogadro's Law with the help of examples and what is its significance?**

**Ans: Avogadro's Law:**

It states that equal volumes of all gases at the same temperature and pressure must contain equal number of molecules.

It means volume is directly proportional to the number of molecules at constant pressure and temperature.

$$V \propto n$$

**Examples:**

1 mole of  $\text{N}_2$  (28g) at S.T.P. =  $22.4 \text{ dm}^3 = 6.022 \times 10^{23}$  molecules

1 mole of  $\text{Cl}_2$  (71 g) at S.T.P. =  $22.4 \text{ dm}^3 = 6.022 \times 10^{23}$  molecules

1 mole of  $\text{O}_2$  (32g) at S.T.P. =  $22.4 \text{ dm}^3 = 6.022 \times 10^{23}$  molecules

1 mole of  $\text{CO}_2$  (44g) at S.T.P. =  $22.4 \text{ dm}^3 = 6.022 \times 10^{23}$  molecules

**Significance of Avogadro's Law:**

Avogadro's law can be applied to determine:

- i. Mass                      ii. Volume                      iii. Number of molecules of a gas

**Q19. By using Avogadro's Law explain molar volume.**

**Ans: Molar volume:**

1 gram molecule or one mole of a gas will have  $22.4 \text{ dm}^3$  at S.T.P. This volume of  $22.4 \text{ dm}^3$  at STP is called Molar volume.

1 mole of  $\text{N}_2$  (28g) at S.T.P. =  $22.4 \text{ dm}^3 = 6.022 \times 10^{23}$  molecules

1 mole of  $\text{Cl}_2$  (71 g) at S.T.P. =  $22.4 \text{ dm}^3 = 6.022 \times 10^{23}$  molecules

1 mole of  $O_2$  (32g) at S.T.P. =  $22.4 \text{ dm}^3 = 6.022 \times 10^{23}$  molecules

1 mole of  $CO_2$  (44g) at S.T.P. =  $22.4 \text{ dm}^3 = 6.022 \times 10^{23}$  molecules

The standard temperature is  $0^\circ\text{C}$  and pressure as one atmosphere.

**Q20. Example 9: Calculate the (I) number of molecules and atoms**

**(ii) Number of mole**

**(iii) Volume in  $\text{dm}^3$  of 10g of ammonia at S.L.P.**

**Solution:**

**i. Number of molecules and atoms:**

According to Avogadro's Law, one mole of any gas at STP contains  $6.022 \times 10^{23}$  molecules in case of di-atomic and poly-atomic molecules For example, in the case of ammonia one mole of it contains  $6.022 \times 10^{23}$  molecules

Now mass of  $NH_3 = 10\text{g}$

$$\text{Number of moles of } NH_3 = \frac{\text{mass}}{\text{molecular mass}} = \frac{10}{17}$$

$$= 0.588 \text{ moles}$$

One mole of  $NH_3$  at S.T.P. =  $6.022 \times 10^{23}$  molecules

0.588 moles of  $NH_3$  at STP =  $0.588 \times 6.022 \times 10^{23}$  molecules

$$= 3.54 \times 10^{23} \text{ molecules}$$

Now number of atoms = no of molecules x atomicity (Number of atoms in one molecule of  $NH_3$ )

$$= 3.54 \times 10^{23} \times 4$$

$$= 14.16 \times 10^{23} \text{ atoms}$$

**ii. Volume in  $\text{dm}^3$**

We know that



22.4dm<sup>3</sup> of any gas at S.T.P = 1 mole

22.4dm<sup>3</sup> of NH<sub>3</sub> at S.T.P = 17g

or 17g NH<sub>3</sub> = 22.4 dm<sup>3</sup>

10g NH<sub>3</sub> =  $\frac{22.4}{17} \times 10 = 13.176$  dm<sup>3</sup>

### Self-Check Exercise 4.4

Calculate the number of molecules in (i) 1 dm<sup>3</sup> of oxygen (ii) 2 dm of Hydrogen (iii) 2.5 dm<sup>3</sup> of Nitrogen All gases are at S.T.P. (Ans: (i)  $2.7 \times 10^{22}$  (ii)  $5.4 \times 10^{22}$  (iii)  $6.75 \times 10^{22}$ )

**Solution:**

i. 22.414 dm<sup>3</sup> of oxygen at STP = 1 mole of oxygen

$$= 6.023 \times 10^{23} \text{ molecules}$$

$$1 \text{ dm}^3 \text{ of oxygen at STP} = \frac{6.023 \times 10^{23}}{22.414}$$

$$= 2.7 \times 10^{22} \text{ molecules}$$

ii. 22.414 dm<sup>3</sup> of hydrogen at STP = 1 mole of hydrogen

$$= 6.023 \times 10^{23} \text{ molecules}$$

$$= 6.023 \times 10^{23}$$

$$2 \text{ dm of hydrogen at STP} = 2 \times \frac{6.023 \times 10^{23}}{22.414}$$

$$= 5.4 \times 10^{22} \text{ molecules}$$

iii. 22.414 dm<sup>3</sup> of nitrogen at STP = 1 mole of nitrogen

$$= 6.023 \times 10^{23} \text{ molecules}$$

$$2.5 \text{ dm}^3 \text{ of nitrogen at STP} = 2.5 \times \frac{6.023 \times 10^{23}}{22.414}$$

$$= 6.67 \times 10^{22}$$

**Q21. Derive General Gas Equation or Ideal Gas Law Equation with the help of Boyle's Law, Charles's Law and Avogadro's Law.**

**Ans: Derivation of Ideal Gas Equation:**

The ideal Gas Equation is a combination of three Laws

1. Boyle's Law
2. Charles's Law
3. Avogadro's Law

Now according to Boyle's Law

$$V \propto \frac{1}{P} \text{ (at constant... and } n) \dots\dots\dots (1)$$

According to Charles's Law

$$V \propto T \text{ (at constant... } P \text{ and } n) \dots\dots\dots (2)$$

According to Avogadro's Law

$$V \propto n \text{ (at constant. } P \text{ and } T) \dots\dots\dots (3)$$

Where n is the number of moles of the gas.

Combining equations (1), (2) and (3). we get

$$V \propto \frac{1}{P} \times T \times n$$

$$V \propto \frac{Tn}{P}$$

$$V = R \frac{Tn}{P}$$

$$PV = nRT$$

Where R is a constant called General Gas Constant. This equation is called General Gas Equation or Ideal Gas Law Equation.

If  $n = 1$  Then  $PV = RT$  or  $\frac{PV}{T} = R$

So  $\frac{P_1V_1}{T_1} = R$  (For first condition of gas)

And  $\frac{P_2V_2}{T_2} = R$  (For second condition of gas)

$$\frac{P_1V_1}{T_1} = \frac{P_2V_2}{T_2}$$

**Q22. Example 10: A certain mass of gas occupies 1000 cm<sup>3</sup> at 57°C and 723 mm of Hg. What will be the volume at S.T.P.?**

**Solution:**

$$V_1 = 1000\text{cm}^3$$

$$V_2 = ?$$

$$P_1 = 723\text{mm of Hg}$$

$$P_2 = 760\text{mm of Hg}$$

$$T_1 = 57^\circ\text{C} + 273 = 330\text{K} \quad T_2 = 273\text{K}$$

Now apply the general gas equation

$$\frac{P_1V_1}{T_1} = \frac{P_2V_2}{T_2}$$

$$V_2 = \frac{723\text{mm of Hg} \times 1000\text{cm}^3}{330\text{K}} \times \frac{273\text{K}}{760\text{mm of Hg}}$$

$$V_2 = \frac{723 \times 1000 \times 273}{330 \times 760} \times \text{cm}^3$$

$$V_2 = 790.3 \text{ cm}^3$$

So the volume of the gas at STP = 790.3 cm<sup>3</sup>

### Activity for Student

**Q.1 A given mass of gas occupies 850 cm<sup>3</sup> at 320K and 0.92x10<sup>5</sup> Nm<sup>-2</sup> pressure. Calculate the new volume of the gas at STP**  
**(Ans: 660cm)**

**Solution:**

$$V_1 = 850 \text{ cm}^3$$

$$V_2 = ?$$

$$P_1 = 0.92 \times 10^5 \text{ Nm}^{-2}$$

$$P_2 = 1 \text{ atm} = 101325 \text{ Nm}^{-2}$$

$$T_1 = 320 \text{ K}$$

$$T_2 = 273 \text{ K}$$

Now apply the general gas equation.

$$\frac{P_1 V_1}{T_1} = \frac{P_2 V_2}{T_2}$$

$$\frac{0.92 \times 10^5 \text{ Nm}^{-2} \times 850 \text{ cm}^3}{320 \text{ K}} = \frac{101325 \text{ Nm}^{-2} \times V_2}{273 \text{ K}}$$

$$V_2 = \frac{0.92 \times 10^5 \text{ Nm}^{-2} \times 850 \text{ cm}^3 \times 273 \text{ K}}{101325 \text{ Nm}^{-2} \times 320 \text{ K}}$$

$$V_2 = 658.410 \text{ cm}^3$$

**Q.2 A container contains 2.5m<sup>3</sup> of Helium gas at 15°C and 98 Nm<sup>-2</sup> pressures. What volume would this gas occupy at S.T.P?**  
**(Ans: 2.3cm')**

**Solution:**

$$V_1 = 2.5 \text{ m}^3$$

$$V_2 = ?$$

$$P_1 = 98 \text{ Nm}^{-2}$$

$$P_2 = 1 \text{ atm} = 101325 \text{ Nm}^{-2}$$

$$T = 15^\circ \text{C} + 273 \text{ K} = 288 \text{ K} \quad T = 273 \text{ K}$$

Now apply the general gas equation

$$\frac{P_1 V_1}{T_1} = \frac{P_2 V_2}{T_2}$$

$$\frac{98Nm^{-2} \times 2.5m^3}{288K} = \frac{101325Nm^{-2} \times V_2}{273K}$$

$$V_2 = \frac{98Nm^{-2} \times 2.5m^3 \times 273K}{101325Nm^{-2} \times 288K}$$

$$V_2 = 0.0023m^3$$

**Q23. List the significance of General Gas Equation.**

**Ans: Significance of General Gas Equation:**

With the help of the Ideal Gas equation, we can determine

(a) Molecular mass of the gas. (b) Density of the gas

**Q24. Derive a relation for molecular mass of the gas with the help of general gas equation.**

**Ans: Molecular Mass of the Gas:**

According to the general gas equation  $PV = nRT$  ..... (1)

But  $n$  (number of moles)  $= \frac{W}{M}$ , Where  $W$  = mass of the gas

$M$  = molecular mass of the gas

Putting the value of  $n$  in

$$PV = \frac{W}{M}RT$$

$$MPV = WRT \quad \text{..... (2)}$$

or  $M = \frac{WRT}{PV} \quad \text{..... (3)}$

So molecular mass ( $M$ ) can be calculated if  $P$ ,  $V$ ,  $W$  and  $T$  are known.

**Q25. Derive a relation for density of the gas with the help of general gas equation.**

**Ans: Density of the gas ( $dm^3$ ):**

According to the General Gas Equation  $PV = nRT$  ..... (1)

But  $n = \frac{W}{M}$ , therefore,  $PV = \frac{W}{M}RT$  or  $PM = \frac{W}{V}RT$

As,  $\frac{W}{M} = d$  (density).

So  $MP = d RT$

Or  $d = \frac{MP}{RT}$

So knowing the values of P, T and M, the density can be calculated.

**Q26. Calculate the numerical value of R in different ways with the help of general gas equation.**

**Ans: Numerical Value of R:**

The value of R can be calculated in different ways as shown below.

**(1)** If one mole of a gas is taken at S.T.P (273 K ,1 atm), then the volume occupied by it is 22.4dm. Now according to the general gas equation.

$PV = nrt$  ..... (1)

Where  $V = 22.4\text{dm}^3, P = 1 \text{ atm}, n = 1 \text{ mole}, T = 273 \text{ K}$  Then  $R = ?$

From the equation (1)

$PV = Nrt$

$$R = \frac{PV}{nT} = \frac{22.4\text{dm}^3 \times 1\text{atm}}{1\text{mole} \times 273\text{K}}$$

$$R = \frac{22.4 \times 1}{1 \times 273} \text{dm}^3 \text{atm} \times \text{mole}^{-1} \text{K}^{-1}$$

$$R = 0.0821 \text{dm}^3 \text{atm} \times \text{mole}^{-1} \text{K}^{-1}$$

**(2)** if pressure is measured in mm of Hg or torr and V in  $\text{cm}^3$  then

$$R = 0.0821 \text{dm}^3 \times \text{atm} \text{mole}^{-1} \text{K}^{-1}$$

$$= 62.4 \text{ dm}^3 \text{ mm mole}^{-1} \text{ K}^{-1}$$

$$= 62.4 \text{ dm}^3 \text{ torr mole}^{-1} \text{ K}^{-1} \text{ (one mm of Hg = one torr)}$$

$$= 62.4 \times 1000 \text{ cm}^3 \text{ torr mole}^{-1} \text{ K}^{-1}$$

$$= 62.4 \times 100 \text{ cm torr mole}^{-1} \text{ K}^{-1}$$

$$= 62400 \text{ cm}^3 \text{ torr mole}^{-1} \text{ K}^{-1}$$

**Q27. Calculate the numerical value and SI unit of R with the help of general gas equation.**

**Ans:** In SI Units, pressure is expressed in  $\text{Nm}^{-2}$  and volume in  $\text{m}^3$ , then

$$V = 0.0224 \text{ m}^3 \text{ (1 dm}^3 = 100 \text{ m}^3)$$

$$P = 101325 \text{ Nm}^{-2}$$

$$n = 1 \text{ mole}$$

$$T = 273 \text{ K}$$

$$R = ?$$

$$R = \frac{PV}{nT} = \frac{0.0224 \text{ m}^3 \times 101325 \text{ Nm}^{-2}}{1 \text{ mole} \times 273 \text{ K}}$$

$$R = \frac{0.0224 \times 1}{1 \times 273} \text{ m}^3 \text{ Nm}^{-2} \times \text{mole}^{-1} \text{ K}^{-1}$$

$$R = 0.0821 \text{ dmm} \times \text{mole}^{-1} \text{ K}^{-1}$$

Now we know that  $1 \text{ Nm} = 1 \text{ J}$

$$R = 8.3143 \text{ J mole}^{-1} \text{ K}^{-1}$$

**Q28. Example 11: Calculate the average molar mass of air at sea level and  $0^\circ\text{C}$  if the density of air is  $1.29 \text{ Kg m}^{-3}$ .**

**Solution:**

At sea level the pressure may be taken equal to 1 atm or 101325 Pa. We know that



$$D = \frac{MP}{RT} \text{ or } M(\text{average molar mass}) = \frac{dRT}{P} \dots\dots\dots (1)$$

where  $d = 1.29 \text{ Kgm}^{-3}$

$R = 8.31453 \text{ K}^{-1} \text{ mole}^{-1}$

$T = 273.15 \text{ K}$

$P = 101325 \text{ Nm}^{-2} = \text{kgm}^{-1}\text{s}^{-1}$

Now putting the values of these in equation .....(1)

$$M = \frac{1.29 \text{ Kgm}^{-3} \times 8.31453 \text{ K}^{-1} \text{ mole}^{-1} \times 273.15 \text{ K}}{101325 \text{ kgm}^{-1}\text{s}^{-1}}$$

$$M = \frac{1.29 \times 8.31453 \times 273.15}{101325} \cdot \frac{\text{Kgm}^{-3} \times \text{K}^{-1} \text{ mole}^{-1} \text{ K}}{\text{kgm}^{-1}\text{s}^{-1}}$$

$$M = 0.0289 \text{ Kgmole}^{-1}$$

**Example 12: A certain gas occupies a volume of 6 dm<sup>3</sup> under a pressure of 720 mm of Hg at 25°C. What volume will this gas occupy under standard conditions of Temperature and Pressure (S.T.P)?**

**Solution:**

$V_1 = 6 \text{ dm}^3$	$V_2 = ?$	$P_1 = 720 \text{ mm Hg}$	$P_2 = 760 \text{ mm Hg}$
$T_1 = 25^\circ\text{C} - 273 = 298 \text{ K}$		$T_2 = 273 \text{ K}$	

According to the General Gas Equation

or

$$\begin{aligned}
 P_1 &= \frac{n_1 RT}{V} \\
 &= \frac{6 \times 720 \times 298}{760 \times 273} \frac{\text{dm}^3 \times \text{mm of Hg} \times \text{K}}{\text{mm of Hg} \times \text{K}} \\
 &= 6.204 \text{ dm}^3
 \end{aligned}$$

**Q30. Example 13: Four grams of CH<sub>4</sub> at 27°C and a pressure of 2.5 atm occupy a volume of 2.46 dm. Calculate the value of R.**

**Solution:**

$$\text{No of moles of CH}_4 = \frac{\text{mass}}{\text{molecular mass}}$$

$$N = \frac{4}{16} = \frac{1}{4} \text{ moles}$$

$$T = 27^\circ\text{C} + 273 = 300\text{K}$$

$$P = 2.5 \text{ atm}$$

$$V = 2.46 \text{ dm}^3$$

$$R = ?$$

According to the General Gas Equation,

$$VP = nRT$$

$$R = \frac{VP}{nT}$$

$$= \frac{2.46 \text{ dm}^3 \times 2.5 \text{ atm}}{\frac{1}{4} \text{ mole} \times 300 \text{ K}}$$

$$= \frac{2.46 \times 2.5 \text{ dm}^3 \text{ atm}}{\frac{1}{4} \times 300 \text{ mole K}}$$

$$R = 0.0821 \text{ dm}^3 \text{ atm mole}^{-1} \text{ K}^{-1}$$

**Q31. Example 14:** Find the density of ammonia gas at  $100^\circ\text{C}$  when confined by a pressure of 1600 mm of Hg.

**Solution:**  $d = ?$

$$T = 100^\circ\text{C} + 273 = 373\text{K}$$

$$P = 1600 \text{ mm of Hg} = \frac{1600}{760} \text{ atm} = 2.105 \text{ atm}$$

$$M_{\text{NH}_3} = 17 \text{ g mole}^{-1}$$

$$R = 0.0821 \text{ dm}^3 \text{ atm mole}^{-1} \text{ K}^{-1}$$

Now according to the General Gas Equation

MP = dRT (Derived from General Gas Equation)

$$d = \frac{MP}{RT} = \frac{17 \text{ gmole}^{-1} \times 2.105 \text{ atm}}{0.0821 \text{ dm}^3 \text{ atm mole}^{-1} \text{ K}^{-1} \times 273 \text{ K}}$$

$$d = \frac{17 \times 2.105}{0.0821 \times 273}$$

$$d = 1.598 \text{ g dm}^{-3}$$

**Q32. Example 15:** At 27°C a volume of 500 cm<sup>3</sup> of H<sub>2</sub> measured under a pressure of 400 mm of Hg and 1000 cm<sup>3</sup> of N<sub>2</sub>, measured under a pressure of 600 mm of Hg are introduced into an evacuated 2 dm<sup>3</sup> flask. Calculate the resulting Pressure.

**Solution:**

We will apply Boyle's Law as the temperature is constant.

**For H<sub>2</sub>:**

$$V_1 = 2 \text{ dm}^3 \quad V_2 = 500 \text{ cm}^3 = \frac{500}{1000} \text{ dm}^3 = \frac{1}{2} \text{ dm}^3$$

$$P_1 = ? \quad P_2 = 400 \text{ mm} = \frac{400}{760} \text{ atm} = 0.526 \text{ atm}$$

$$V_1 P_1 = V_2 P_2 \text{ or } 2 \times P_2 = \frac{1}{2} \times 0.526 \quad P_2 = 0.1315 \text{ atm}$$

**For N<sub>2</sub>:**

$$V_3 = 1000 \text{ cm}^3 = 1 \text{ dm}^3$$

$$P_3 = ? \quad P_4 = \frac{600}{760} \text{ mm} = 0.789 \text{ atm}$$

$$V_3 P_3 = V_4 P_4 \text{ or } 2 \times P_3 = 1 \times 0.789 = 0.3945 \text{ atm}$$

$$\text{Total pressure} = 0.1315 \text{ atm} + 0.3945 \text{ atm} = 0.5260 \text{ atm}$$

**Q33. Example 16:** Find the total pressure exerted by 2g of ethane and 3g of CO<sub>2</sub>, contained in a 5dm<sup>3</sup> vessel at 50°C.

**Solution:**

44 g of CO<sub>2</sub> = 1 mole

$$3\text{g CO}_2 = \frac{1}{44} \times 3 = 0.068 \text{ moles}$$

Number of moles of CO<sub>2</sub> = 0.068 moles

$$V = 5\text{dm}^3 \quad T = 50^\circ\text{C} - 273 = 323\text{K}$$

$$R = 0.082\text{dm}^3\text{atm K}^{-1} \text{ mole}^{-1}$$

Now according to the General Gas Equation

$$VP = nRT \quad \text{or} \quad P = \frac{nRT}{V}$$

$$P_{\text{CO}_2} = \frac{0.068\text{mole} \times 0.0821\text{dm}^3\text{atmmole}^{-1}\text{K}^{-1} \times 323\text{K}}{5\text{dm}^3}$$

$$P_{\text{CO}_2} = \frac{0.082 \times 0.068 \times 323}{5} = 0.678\text{atm}$$

30g Ethane (C<sub>2</sub> H<sub>6</sub>) = 1 mole

$$2\text{g Ethane (C}_2\text{H}_6) = \frac{1}{30} \times 2 = 0.06 \text{ moles}$$

$$V = 5\text{dm}^3$$

$$n = 0.6 \text{ moles}$$

$$T = 50^\circ\text{C} + 273 = 323 \text{ K}$$

$$R = 0.082\text{dm}^3\text{atm K}^{-1} \text{ mole}^{-1}$$

$$VP = nRT \quad \text{or} \quad P = \frac{nRT}{V}$$

$$P = 0.06 \text{ moles} \times R = 0.082\text{dm}^3\text{atm K}^{-1} \text{ mole}^{-1} \times 323\text{K}$$

$$= 0.06 \times 0.082 \times 323 = 6.317 \text{ atm}$$

**Q34. Briefly explain why Real Gases Deviate from the Gas Laws explain it with the help of example.**

**Ans: Deviation of Real gases from the Gas Laws:**

At low pressures, the molecules are generally very far apart and the attractions between the molecules are insignificant at temperatures above the condensation point, the Kinetic Energy of the gaseous molecules is so great that it overwhelms the attractive forces between them.

The example of Boyle's Law ( $PV = a$  (constant)) be quoted as follows e.g. for  $H_2$ .

Pressure (atm)	Volume ( $dm^3$ )	PV
0.1	224.1	22.41
50	0.4634	23.71
1000	0.384	38.34

The result (PV) is quite different, temperature remains constant.

**Q35. Briefly explain the causes of Deviation of Real Gases.**

**(OR)**

**Gases show non-ideal behavior at low temperature and high pressure. Explain.**

**Ans: Causes of Deviations (Limitations of Boyle's Law and Charles's Law):**

The ideal condition is that the ideal gas obeys Gas Laws at all temperature but at low temperature and high-pressure deviations caused by gases is due to the two faulty assumptions of kinetic molecular theory.

**(1) Inter-molecular Forces of attraction:**

There are no attractive or repulsive forces between the gas molecules

**(2) Volume:**

The volume of the gas molecules is negligible as compared to the volume of the container.

**Explanation of deviations:**

**(1) At lower temperature:**

At high temperature the kinetic energy of the gaseous molecules is very high. As a result, the attractive forces between them are negligible. But when temperature is decreased, the Kinetic

Energy of the molecules decreases. The intermolecular forces become significant. It means that the molecules come close to each other.

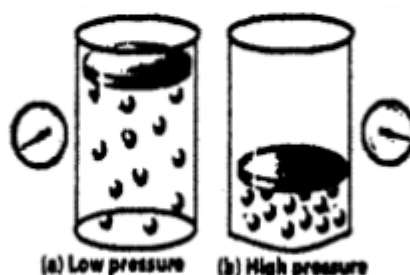
At a certain very low temperature the gases change into the liquid state. Therefore, attractive forces between the gas molecules become significant near liquefying temperature. That is why the ideal gases deviate from their original behavior at low temperature

**Example:**

$\text{SO}_2$ , liquefies at  $-10^\circ\text{C}$  while  $\text{H}_2$  at  $-252.7^\circ\text{C}$ . Therefore, attractive forces between  $\text{SO}_2$  molecules cannot be considered negligible at room temperature. For this reason,  $\text{SO}_2$  gas shows non-ideal behavior at room temperature as compared to  $\text{H}_2$  gas.

**(2) At high pressure:**

At low pressure actual volume of gas molecules is very small as compared to the volume of the container. However, this volume does not remain negligible at high pressure. This can be understood from the following figure.



**Suppose:**

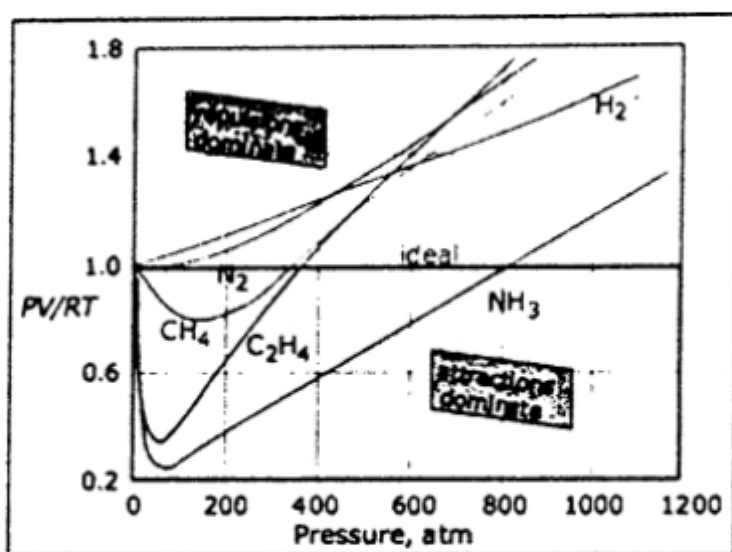
When pressure is one atm, individual volume of one mole of gas molecules say  $32\text{ cm}^3$  is negligible as compared to the total volume of gas  $22400\text{ cm}^3$  but if it is subjected to a pressure of 100 atm, the volume of the gas is reduced to  $224\text{ cm}^3$ . Under this pressure, individual volume of gas molecules ( $32\text{ cm}^3$ ) is not negligible as compared a volume of the gas ( $224\text{ cm}^3$ ).

**Q36. Describe Graphical Representation of Deviation of Gases from ideal behavior.**

**Ans: Graphical Representation:**

For one mole of a gas, a graph is plotted between  $\frac{PV}{RT}$  and pressure then for an ideal gas, it must give a straight line the temperature remains at 0°C. But actually, it's seen that the gases do not give a straight line. They deviate from their original behavior as shown in the figure

Actually, when the pressure is initially increased, it pushes the molecules closer and increases the intermolecular forces. Due to these forces, the volume of the gas shrinks more than that predicted from the gas laws (i.e: Boyle's Law). This will decrease the value of  $\frac{PV}{RT}$ . That is why the gases do not show a straight line.  $\frac{PV}{RT}$  is called compressibility factor. For ideal gases it should be one, whereas for real gases it is not equal to one.



The sudden decrease in the curve of  $C_2H_4$ , and  $NH_3$ , shows that these gases liquefy at the given pressure

**Q37. Explain and derive Van Der Waal's Equation for Real Gases.**

**Ans: Van Der Waal's Equation: (Correction Factors to Ideal Gas Equation);**

The general gas equation is not applicable to real gases at all temperature and pressure. In order to make it applicable to real gases, a Dutch Scientist Van der Waals in 1873 made correction in volume and pressure of the gases and derived an equation known as Vander Waals equation.

#### **Volume Correction:**

Vander Waal thought that some of the volume of the vessel is occupied by the molecules of the gas. Hence, the volume is not available for the free movement of the molecules. Actually,



we need the free volume of the gas and that is obtained when we subtract the individual volume of gas molecules from the volume of the vessel

$$V = V_{\text{vessel}} - V_{\text{molecule}}$$

$V$  is free volume

$$V_{\text{molecule}} = nb$$

where  $b$  is individual volume of one mole of the gas molecules and  $n$  is total number of moles of the gas molecules

$$\text{Then, } V = V_{\text{vessel}} - nb$$

Here ' $b$ ' is effective volume of gas molecules. Keep it in mind that ' $b$ ' is not the actual volume of gas molecule, but is roughly equal to 4 times their molar mass volumes. If we have one mole of gas then  $b = 4V_m$ . Where,  $V_m$  is the actual volume of gas molecules for one mole of the gas.

### **Pressure correction:**

The pressure which is exerted on the walls of the vessels is due to collisions of molecules of gas. Since there are some weak forces of attractions among the gas molecules, so the molecules cannot hit the walls of the vessels with that much force with which they should have been in the absence of these attractive forces.

It means that the pressure being observed on the walls of the vessels is a little bit lesser than the ideal pressure

$$\text{Therefore, } P_{\text{observed}} = P_{\text{ideal}} - P_{\text{lessened}}$$

If the ideal pressure is denoted by  $P_i$  and the pressure lessened due to molecular attractions is denoted by  $P'$  then

$$P_{\text{observed}} = P_i - P'$$

$$\text{and } P_i = P_{\text{observed}} + P'$$

But the value of  $P'$  is inversely proportional to the density of the gas so,

$$P \propto (\text{density})^2$$

$$P' = \frac{an^2}{V^2}$$

where  $n$  = number of moles,  $V$  = volume,  $a$  = proportionality constant

$$P_i = P + P' = \frac{an^2}{V^2}$$

After the correction of volume and pressure.  $PV = nRT$  becomes

$$\left(P + \frac{an^2}{V^2}\right)(V - nb) = nRT$$

For 1 mole of gas,  $n = 1$ , then

$$\left(P + \frac{a}{V^2}\right)(V - b) = RT$$

**Q38. List the Van der Waals constants for some gases.**

**Ans:**

Van der Waals constants for some gases.		
Substance	$a(\text{dm}^3\text{atm mole}^{-1})$	$b(\text{dm}^3 \text{mole}^{-1})$
H <sub>2</sub>	0.0247	0.0266
He	0.0034	0.0237
N <sub>2</sub>	0.1408	0.0391
O <sub>2</sub>	0.1378	0.0318
Cl <sub>2</sub>	0.6579	0.0562

Ar	0.1355	0.0322
Kr	0.2349	0.0398
CO	0.1505	0.0399
NO	0.1358	0.0279
CO <sub>2</sub>	0.3640	0.0427
HCl	0.3716	0.0408
SO <sub>2</sub>	0.6803	0.0564
H <sub>2</sub> O	0.5536	0.0305
NH <sub>3</sub>	0.4225	0.0371
CH <sub>4</sub>	0.2283	0.0428
CCl <sub>2</sub> F <sub>2</sub>	0.1066	0.0973

**Q39. Example 17: One mole of methane gas is maintained at 300 K. Its volume is 250 cm<sup>3</sup>. Calculate the pressure exerted by the gas under the following conditions.**

- (i) When the gas is ideal
- (ii) When the gas is non ideal, and at

$$a = 2.253 \text{ dm}^3 \text{ atm mole}^{-1}$$

$$b = 0.0428 \text{ dm}^3 \text{ mole}^{-1}$$

**Solution:**

(i) **When the gas is ideal**

General gas Equation is applied here.

(a) **Gas=Methane:**

$$n = 1 \text{ mole}$$

$$T = 300\text{K}$$

$$V = 250\text{cm}^3 = 0.25\text{dm}^3$$

$$R = 0.0821 \text{ dm}^3 \text{ atm Mole}^{-1}$$

$$p = ?$$

Now applying the General Gas Equation,  $PV = nRT$ .

$$P = nRT/V$$

$$= 98.52\text{atm}$$

(ii) **When the gas is non-ideal (Here Vander Waals' equation is applied)**

$$(P + n^2a/V^2) (V - nb) = nRT$$

Now dividing by  $(V - nb)$  on both sides Putting the following values in this equation

$$n = 1 \text{ mole,}$$

$$R = 0.0821 \text{ atm mole}^{-1}\text{K.},$$

$$V = 0.25\text{dm}^3,$$

$$T = 300\text{K,}$$

$$= 2.253\text{dm}^3\text{atmmole}^{-1},$$

$$b = 0.0428\text{dm}^3\text{mole}^{-1}$$

$$1\text{M} \times 0.0821\text{dm}^3\text{mole}^{-1} \text{K}^{-1} \times 300\text{K}$$

$$1\text{M} \times 2.25\text{dm}^3\text{atmK}^{-1}$$

$$0.25\text{dm}^3 - (1\text{M} \times 0.0428\text{dm}^3\text{mole}^{-1})$$

$$(0.25)^3\text{dm}^3$$

$$= 118.87 - 3.6048 = 115.2652 \text{ atm}$$

**Q40. Example 18: Estimate the molar volume of  $\text{N}_2$  at 300K and 100atm by treating it as a Van der Waals' equation. ( $R = 0.0821 \text{ dm}^3 \text{ atm K}^{-1} \text{ mole}^{-1}$ )**

$$a = 1.39\text{dm}^2 \text{ atm Mole}^{-2}$$

$$b = 3.913 \times 10^{-2} \text{ dm}^3\text{mole}^{-1}$$

$$\text{(Ans: Molar Volume} = 0.5 \text{ dm}^3\text{)}$$

For 1 mole of a gas ( $n = 1$ ) Van der Waals equation is given by

$$(P + n^2a/V)(V - b) = RT$$

$$(p + \frac{a}{V^2})(v - b) = RT$$

$$(v - b) = \frac{RT}{P + \frac{n^2r^2}{P^2}}$$

Since  $PV=RT$ ,  $V = \frac{RT}{P}$ ,  $V^2 = \frac{n^2r^2}{P^2}$  putting in equation (i)

$$V = \frac{RT}{P + \frac{n^2r^2}{P^2}} + b$$

$$V = \frac{RT}{P + \frac{aP^2}{n^2r^2}} + b$$

$$V = \frac{0.0821 \times 500}{100 + \frac{1.39 \times 100^2}{(0.0821)^2 \times (500)^2}} + 3.913 \times 10^{-2}$$

$$V = \frac{0.0821 \times 500}{100 + \frac{1.39 \times 10000}{(0.0067 \times 250000)}} + 3.913 \times 10^{-2}$$

$$V = \frac{41.05}{100 + \frac{13900}{1685.1025}} + 3.913 \times 10^{-2}$$

$$V = \frac{41.05}{100 + 8.23} + 3.913 \times 10^{-2}$$

$$V = 0.4 + 3.913 \times 10^{-2}$$

$$V = 0.4 + 0.03913$$

$$V = 0.439$$

$$V = 0.5 \text{ dm}^3$$

**Q41. Define Dalton's Law of Partial Pressure and its applications.**

**Ans: Dalton's Law of Partial Pressure and its Applications:**

An English Chemist John Dalton (1766-1844) showed in 1801 that "The total pressure exerted by a gaseous mixture is equal to the sum of partial pressures of each gas present in the mixture."

**Conditions for Dalton's Law of Partial Pressure:**

The law is only obeyed, if the component gases do not chemically react with each other. Moreover, the gases must behave ideally. The partial pressure is the pressure exerted by one component of the gaseous mixture.

**Mathematically:**

$$\text{Total Pressure } (P_1) = P_A + P_B + P_C$$

Whereas  $P_A$ ,  $P_B$  and  $P_C$  are the partial pressures of individual gases.

**Example of Air in terms of Dalton's Law of Partial Pressure:**

Air is a mixture of non-reacting gases. Percentage of each gas in the gaseous mixture is  $N_2$  (78.08%),  $O_2$  (20%), Ar (0.93%),  $CO_2$  (0.03%). Traces of Ne, He, Kr,  $H_2$ , along with various amounts of water vapors and pollutant (which can pollute the atmosphere) gases such as oxides of Sulphur and Nitrogen are also found in air. The total pressure exerted by the air will be the sum of partial pressure of each gas.

**Q42. Derive Relationship between pressure and number of moles of a gas with the help of Dalton's Law of Partial Pressure.**

**Ans: Relationship between pressure and number of moles of gases:**

If  $n_A$  and  $n_C$  are the number of moles of the gases and  $P_A$ ,  $P_B$  and  $P_C$  be their partial pressure respectively.

Then according to the general gas equation

$$PV = nRT$$

**For gas A:**

$$P_A = n_A RT / V$$

**For gas B:**

$$P_B = n_B RT / V$$

**For gas C:**

$$P_C = n_C RT / V$$

$$P_t = n_t RT / V$$

Where  $n_t$  (total number of moles of all the gases) =  $n_a + n_b + n_c$  Dividing equation (1) by (4)

$$P_A / P_t = n_A RT / V \dots n_t RT / V$$

$$P_A / P_t = n_A RT / V \times V / n_t RT$$

$$P_A / P_t = n_A / n_t = P_A = n_A / n_t = P_A \times n_A$$

Partial Pressure of any gas / total no of moles of all gases =

No of moles of that gas / total no of moles of all gases

**Q43. Derive a Relationship between pressure and mole fraction of a gas with the help Dalton's Law of Partial Pressure.**

**Ans: Relationship between pressure and mole fraction of a gas:**

We know that

Partial Pressure of any gas = No of moles of that gas

Pressure of all gas's                      Total no of moles of all gasses

$$\text{Or } \frac{p_t}{P_t} = \frac{n_t}{n_t} \quad p_t = \frac{n_t}{n_t} \times P_t$$

$\frac{n_t}{n_t} = X$  where  $X$ , is called mole fraction of the component

$$\text{So } p_t = P_t X$$

Thus, the partial pressure of any component is equal to the product of total pressure of all the components and mole fraction of that component.

**Q44. Give at least three Applications of the Law of Partial Pressure.**

**Ans: Applications of the Law of Partial Pressure:**

**i. Collection of gases over water:**

The Dalton's Law is particularly Useful when a gas is generated and subsequently collected over water provided the gas is insoluble in water. The total pressure consists of the pressure of the water vapors in addition to the pressure of the gas that is generated. The pressure due to water vapor is called aqueous tension.

$$\text{Thus } P_{\text{total}} = P_{\text{gas}} + P_{\text{water}} \quad \text{or } P_{\text{gas}} = P_{\text{total}} - P_{\text{water}}$$



The gas contaminated with water may be called a moist gas

### **ii. Respiration at higher altitudes:**

The respiration process in living things depends upon the differences in partial Pressure eg. Partial pressure of  $O_2$  in the outside air is higher ( $159 \text{ g/cm}^2$ ) than in the lungs where the partial pressure of  $O_2$  is lower ( $116 \text{ g/cm}^2$ ). However at high altitudes, the pressure of oxygen decreases and the pilots may have uncomfortable breathing in a non-pressurized cabin where the partial pressure of  $O_2$  is about  $150 \text{ g/cm}^2$

### **Respiration by deep sea divers:**

Deep sea divers breathe air under increased pressure. At a depth of 40 meters, the pressure increases 5 times than the normal pressure. Therefore, regular air cannot be used in diver's tanks because the partial pressure of  $O_2$  would be  $795 \text{ mm of Hg}$  ( $159 \times 5 = 795 \text{ mm of Hg}$ ) in that case. Therefore, deep sea divers use to breathe a mixture of 96%  $N_2$  and 4%  $O_2$  in the respiration tank, the scuba. A scuba (self-contained breathing apparatus having oxygen used by divers underwater) contains compressed air to breathe. As diver returns to the surface, it becomes hazardous for him if not handled properly. As the diver comes up, the pressure of the surrounding water drops. Consequently, the compressed air in the lungs expands. Surfacing must be done very slowly so that the compressed air may escape out from the lungs without causing damage to them.

### **Q45. How deep-sea divers respire on return from sea?**

**Ans:** The percentage of  $N_2$  in air is about 80%. If  $N_2$  is present in diver's tank, the solubility of  $N_2$  in blood can increase with increase in pressure in very deep sea. And blood of a diver may be saturated with  $N_2$  during a dive.



If the diver returns quickly and relatively at low pressure at the surface, he will face life threatening condition called the "bends" Deep sea divers must either use different solutions of gases such as He in O<sub>2</sub> or spend many hours in a decompression chamber after a dive. The pressure in a compression chamber is slowly lowered over many hours. Nitrogen gas comes out of the blood and disposed of safely for normal breathing

**Q46. Example 19: 1.00 mole of N<sub>2</sub> and 3.00 moles of H<sub>2</sub> are enclosed in a container of volume 10.0 dm<sup>3</sup> at 298K. What are the partial pressures and the total pressure?**

**Solution:**

No of moles of N<sub>2</sub> (n<sub>1</sub>) = 1.00 mole

No of moles of H<sub>2</sub> (n<sub>2</sub>) = 3.00 moles

Total number of moles (n<sub>total</sub>) = 1 + 3 = 4 moles

V = 10.0 dm<sup>3</sup>

T = 298K,

R = 0.082 dm<sup>3</sup> atm K Mole<sup>-1</sup>

According to the General Gas Equation,

PV = nRT

$$= \text{or } P_{total} = \frac{n_{total}RT}{V}$$

$$= \frac{4 \times 0.082 \times 298}{10} \cdot \frac{dm^3 \text{atmos} K^{-1} \text{Mole}^{-1} \times K}{dm^3 \text{Mole}}$$

$$= \frac{4 \times 0.082 \times 298}{10} \text{atm}$$

$$= 9.77 \text{atm}$$

$$P_{N_2} = \frac{n_{N_2}}{n_{total}} \times P_{total} = \frac{1}{4} \times 9.77 = 2.44 \text{atm}$$

$$P_{H_2} = \frac{n_{H_2}}{n_{total}} \times P_{total} = \frac{3}{4} \times 9.77 = 7.33 \text{atm}$$

**Q47. Example 20:** The composition of dry air at sea level is approx.  $N_2=75.5$ ,  $O_2=23.2$ .  
 $Ar=1.3$  in percentage by weight. What is the partial pressure of each component when the total pressure is 1.000 atm.

**Solution:**

Let the total number of moles of air = 100 then

No of moles of  $N_2$  ( $n_{N_2}$ ) = 75.5

No of moles of  $O_2$  ( $n_{O_2}$ ) = 23.2

No of moles of  $Ar$  ( $n_{Ar}$ ) = 1.3

$P_{total} = 1.00$  atm.

$$P_{N_2} = \frac{n_{N_2}}{n_{total}} \times P_{total}$$

$$= \frac{75.52}{100.00} \times 1.00 = 0.752 \text{ atm}$$

$$P_{N_2} = \frac{n_{O_2}}{n_{total}} \times P_{total} = \frac{23.2}{100.00} \times 1.00 = 0.232 \text{ atm}$$

$$P_{N_2} = \frac{n_{Ar}}{n_{total}} \times P_{total}$$

$$= \frac{1.3}{100.00} \times 1.00 = 0.013 \text{ atm}$$

**Q48. Example 21:** A certain mass of  $H_2$  gas collected over water at  $6^\circ C$  and 765mm of Hg pressure occupied a volume of  $35 \text{ cm}^3$

Calculate its dry volume at S.T.P. (V.P. of water at  $6^\circ C = 7 \text{ mm. of Hg}$ ) Solution:

$V_1 = 35 \text{ cm}^3$ ,  $V_2 = ?$   $P_1 = 765 - 7 = 758 \text{ mm}$

$T_1 = 6 + 273 = 279 \text{ K}$ ,  $T_2 = 273 \text{ K}$   $P_2 = 760 \text{ mm.}$

$$\frac{V_1 P_1}{T_1} = \frac{V_2 P_2}{T_2} \text{ Or } \frac{V_1 P_1 T_2}{T_1 P_2} = \frac{35 \text{ cm}^3 \times 758 \text{ mm} \times 273 \text{ K}}{279 \text{ K} \times 760 \text{ mm of Hg}} = 34.2 \text{ cm}^3$$

### Activity for students

A gas was collected over water at 10°C and 803mm of Hg. If the gas occupies 73cm<sup>3</sup> calculate the volume of dry gas at S.T.P. (aqueous tension at 10°C - 9.2mm).

(Ans: = 73.55cm<sup>3</sup>)

**Solution:**

$P_{\text{moist}} = P_{\text{dry}} + \text{Aqueous tension}$

Volume of gas at 10 °C =  $V_1 = 73 \text{ cm}^3$

$T_1 = 10 \text{ °C} + 273 = 283\text{K}$

Values at STP:  $P_2 = 760$

$T_2 = 273 \text{ K}$

$V_2 = ?$

According to the general gas equation

$$\frac{V_1 P_1}{T_1} = \frac{V_2 P_2}{T_2}$$

$$V_2 = \frac{V_1 P_1 T_2}{T_1 P_2}$$

$$V_2 = \frac{706 \times 73 \times 273}{760 \times 283}$$

$$V_2 = 0.002\text{m}^3$$

$$V_2 = 65.41 \text{ cm}^3$$

**Q49. Define diffusion with the help of example.**

**Ans: Diffusion:**

The spontaneous intermixing of molecules of one gas with another at a given temperature and pressure is called Diffusion. Diffusion is the random movement of a gas from an area of higher concentration to an area of lower concentration.

**Example:**

i. Fragrance of rose.

ii. Intermixing of  $\text{NH}_3$  and  $\text{HCl}$  gives dense white fumes of  $\text{NH}_4\text{Cl}$ .  $\text{NH}_3 + \text{HCl} \rightarrow \text{NH}_4\text{Cl}$

iii. Reddish brown bromine ( $\text{Br}_2$ ) gas taken in a test tube will combine with air taken in another test tube. Red coloration of  $\text{Br}_2$  gas will spread in the test tube containing air.

iv. If liquid ink is left in the bottom of test tube containing water, it will spread evenly throughout the water.

#### Q50. Define Effusion.

**Ans: Effusion:**

The movement of gaseous molecules through extremely small pores in a region of low pressure is called Effusion. This escaping of molecules is not due to collisions but due to their tendency to escape one by one.

**Q51. Example 22: Equal volumes of  $\text{HCl}$  and  $\text{SO}_2$  are confined in a porous container. What would be the comparative rates of diffusion of these gas  $\text{HCl}$  through the porous walls? The molecular masses of  $\text{HCl}$  and  $\text{SO}_2$  are 36.5 and 64 respectively.**

**Solution:**

Molecular mass of  $\text{HCl}$  ( $M_1$ ) = 36.5

Molecular mass of  $\text{SO}_2$  ( $M_2$ ) = 64

Rate of diffusion of  $\text{HCl}$  ( $r_1$ ) = ?

Rate of diffusion of  $\text{SO}_2$  ( $r_2$ ) = ?

According to the Graham's Law of Diffusion

$$\frac{r_1}{r_2} = \sqrt{\frac{M_2}{M_1}} = \sqrt{\frac{64}{36.5}}$$

$r_1(\text{HCl}) > r_2(\text{SO}_2)$

1.33:1

Hence rate of diffusion of  $\text{HCl}$  will be 1.33 times the rate of diffusion of  $\text{SO}_2$ .

**Q52. Example23:** If 280 cm<sup>3</sup> of H<sub>2</sub> diffuse in 40 seconds, how long will it take 490 cm<sup>3</sup> of a gas (X), where vapor density is 25 to diffuse under the same conditions (molecular mass of H<sub>2</sub> is 2).

**Solution:**

For H<sub>2</sub>:

$$V = 280 \text{ cm}^3 \quad M_1 = 2 \text{ gm mole}^{-1}$$

T<sub>1</sub> = 40 sec (time for the diffusion)

velocity 280 cm<sup>3</sup>

$$r_1 = \text{rate of diffusion of H}_2 = \frac{\text{velocity}}{\text{time}} = \frac{280 \text{ cm}^3}{40 \text{ sec}} = 7 \text{ cm}^3 \text{ sec}^{-1}$$

**For Gas (X):**

V = 490 cm<sup>3</sup> Vapor density (V.D.) = 25

T<sub>2</sub> = ? M<sub>2</sub> (Molar mass) = 2 x V.D. = 2 x 25 = 50 gm mole<sup>-1</sup>

$$\text{(Rate of diffusion of gas X)} = \frac{\text{velocity}}{\text{time}} = \frac{490 \text{ cm}^3}{t_2 \text{ sec}} = \frac{490}{t_2} \text{ cm}^3 \text{ sec}^{-1}$$

Now according to Graham's Law of Diffusion

$$\frac{r_1}{r_2} = \sqrt{\frac{M_2}{M_1}}$$

$$= \sqrt{\frac{50 \text{ gm Mole}^{-1}}{2 \text{ gm Mole}^{-1}}}$$

$$\frac{7}{490} \times t_2 = \sqrt{\frac{25}{1}} = \frac{5}{1} \quad \text{or} \quad 7 \times t_2 \times 1 = 490 \times 5 = 350 \text{ sec}$$

Hence time for the diffusion of un-known gas = 350 sec

### ACTIVITY

If 465 cm<sup>3</sup> of 502 can diffuse through porous partition in 30 seconds. How long will 620 cm<sup>3</sup> of H<sub>2</sub>S take to diffuse through the same partition (H = 1, S = 32, O = 16) (Ans: 29.15 sec)

**Solution:**

$$V_{SO_2} = 465 \text{ cm}^3$$

$$r_{SO_2} = \frac{\text{velocity}}{\text{time}} = \frac{465}{30} = 15.5 \text{ cm}^3 \text{ sec}^{-1}$$

$$M_{SO_2} = 64 \text{ gmol}$$

$$V_{H_2S} = 620 \text{ cm}^3$$

$$M_{H_2S} = 34 \text{ gmol}^{-1}$$

$$r_{H_2S} =$$

$$\frac{15.5 \text{ cm}^3 \text{ sec}^{-1}}{0.72} = \sqrt{\frac{34}{64}}$$

$$r_{H_2S} = \frac{15.5 \text{ cm}^3 \text{ sec}^{-1}}{0.72}$$

$$r_{H_2S} = 21.29 \text{ cm}^3 \text{ sec}^{-1}$$

$$r_{H_2S} = \frac{\text{velocity}}{\text{time}} = \frac{465}{21.29} = 29.15 \text{ sec}$$

$$T_{H_2S} = 29.15 \text{ sec}$$

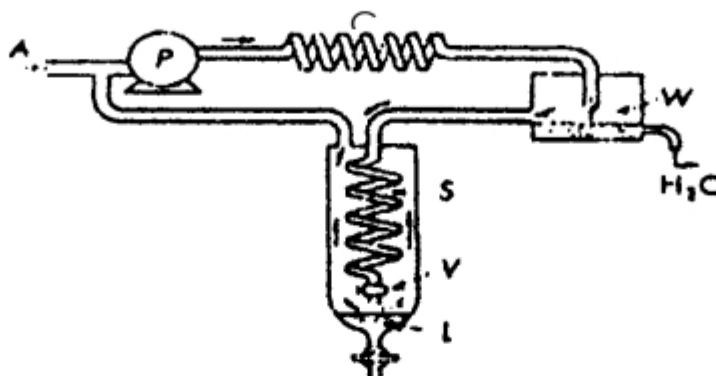
**Q53. Explain Joule-Thomson's effect and Liquefaction of Gases.****Ans: Liquefaction of Gases:****Principle:**

The gases can be liquefied by Joule-Thomson's effect. The liquefaction of a gas requires high pressure and low temperature. When a highly compressed gas is allowed to escape out through a throttle (small hole), the temperature falls to such an extent that it changes into the liquid form. At high pressure, the gaseous molecules come close to each other with the result that the molecular attractions increase. When it is allowed to escape through a nozzle (small hole) into a region of low pressure, the molecules move apart. In doing so, the intermolecular attractions must be overcome. This energy is taken from the molecules themselves. Therefore, the gas is cooled. This process is repeated for many times until the gas completely changes into the liquid form.

**Q54. Describe principle and working of Linde's Method for liquefaction of gases.**

### Ans: Lynde's Method (1895):

According to Joule-Thomson's effect "When a highly compressed gas is allowed to escape out through a throttle" the temperature of the gas falls to such an extent, that it changes into the liquid form". This is the basic principle of the adiabatic expansion which is thermally isolated from its environment, Joule Thomson expansion and subsequent cooling of the air.



Lynde's method for liquefaction of air

### Working Steps:

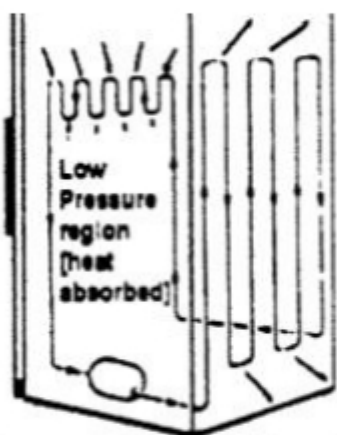
- Air is compressed to 200 at map prox.
- Most of the water in the air condenses and is removed.
- The heat generated as a result of compression is removed by passing the gas through coils C.
- The dry gas is then passed through a copper spiral coil S.
- It is then expanded to almost atmospheric pressure through a controlled valve V.
- The issuing gas, cooled now due to Joule-Thomson's effect is passed over the copper spiral and cool further the incoming compressed gas.
- The cycle is repeated several times.
- The temperature of the expanding gas finally drops and the remaining air liquefied.
- The liquid air collects in the chamber L and can be drawn off.
- Any uncondensed air is re-circulated.



**Q55. Gives the different application of Linde's Method.**

**Ans: I. Construction of appliances:**

The construction of appliances such as refrigerators, heat pumps and air conditioners all work on the principle of Joule-Thomson's effect. e.g. In a refrigerator a compressor, a condenser, a restrictor valve and an evaporator are used. A gas such as Freon is compressed. Thus, an increase in temperature takes place. The hot Freon gas is then moved to a condenser on the outside of the refrigerator. Air at room temperature cools the gas down and as a result condenses into liquid.



**Cooling gas path in a refrigerator**

The liquid Freon, now at room temperature, is then passed through a small hole in a restriction valve into the low pressure tube which are inside the refrigerator. There the Freon liquid vaporizes rapidly. The heat of vaporization needed to do this is drawn from the kinetic energy of the Freon molecules becoming very cold. This cold gas absorbs heat from the refrigerator and its contents, thereby cooling. Then the Freon gas is once again fed into the compressor and the cycle starts over again.

**Self-cooling pop can(container):**

A container holding liquid  $\text{CO}_2$  is built right into the can.

When the can is opened, the liquid  $\text{CO}_2$  escapes

out of the top of the can. The heat absorbed by the vaporizing  $\text{CO}_2$  can lower the temperature of the POP by about  $16^\circ\text{C}$  in a few seconds. Thus, the temperature is lowered considerably.



**Q56. Describe different Applications of Kinetic Molecular Theory.**

**Ans: Velocity of molecules:**



The equation for the root mean square velocity deduced from kinetic equation

$$C_{rms} = \sqrt{\frac{3RT}{M}}$$

**C<sub>rms</sub>** = root mean square velocity

**M** = molecular mass of the gas

T = absolute temperature.

This equation gives a quantitative relationship between the absolute temperature and the velocities of gas molecules. Higher the temperature of a gas, greater would be the velocities of the molecules.

## ii. **Graham's Law of diffusion of gases:**

Thus Kinetic Molecular theory predicts that the average velocity depends upon the molecular mass. Further the average velocity of the particles is inversely proportional to the square root of their molecular mass.

$$E_k = \frac{1}{2}mv^2 \Rightarrow \frac{2E_k}{m} = \sqrt{\frac{2E_k}{m}} \Rightarrow v = \sqrt{\frac{1}{m}}\sqrt{2E_k}$$

As at constant temperature K.E will remain constant so  $\sqrt{2E_k} = \text{constant}$

Therefore  $v \propto \sqrt{\frac{1}{m}}$

Graham's Law states that the rate of diffusion of a gas is inversely proportional to the square root of the mass of its molecules. This law is direct experimental evidence for the Kinetic Molecular Theory.

## iii. **Use of combined gas laws in calculations**

**(Applying):**

**Combined Gas Laws:**

The combination of Boyle's Law and Charles' Law is called "Combined Gas Law."

**Applications:**

i. It is applied to solve problems when volume of a fixed amount of gas is involved, when both the temperature and pressure change. Thus, two conversion factors are involved.

a. Conversion factor for pressure

Boyle's Law

b. Conversion factor for temperature

Charles' Law

**Calculations:**

$V_2 = V_1$  (Conversion factor due to temperature change)  $\times$  (Conversion factor due to Pressure change)

$$\text{Or } V_2 = V_1 \times \frac{T_2}{T_1} \times \frac{P_1}{P_2}$$

The ratio between  $T_1$  and  $T_2$  is due to Charles law ( $V \propto T$ ) and that of  $P_1$  and  $P_2$  is due to Boyle's law

$$(V \propto \frac{1}{P})$$

$$\text{Therefore } V_2 = V_1 \times \frac{T_2}{T_1} \times \frac{P_1}{P_2}$$

In combined gas law problems, which involve a change in temperature and pressure conditions, the mass of the gas must remain constant.

The combined gas law is used when a fixed number of moles of a gas are maintained at different conditions of temperature and pressure.

**iv. Ideal Gas Laws to calculate the Pressure or Volume of a gas:**

Ideal Gas Laws i.e. Boyles' Law and Charles' Law when combined together give Ideal gas law equation at S.T.P.

The equation is  $PV = nRT$

If one of the variables is known, the other three can be calculated.

**Q57. Example 124: A certain mass of  $H_2$  gas collected over water at  $6^\circ C$  and 755mm of Hg pressure occupied a volume of  $35\text{cm}^3$ . calculate its dry volume at S.T.P.**

**Solution:**

$$V_1 = 35\text{cm}^3 \quad V_2 = ?$$

$$P_1 = 765 - 7 = 758\text{mm}$$

$$P_2 = 760\text{mm}$$

$$T_1 = 6 + 273 = 279\text{K} \quad T_2 = 273\text{K}$$

$$\frac{V_1 P_1}{T_1} = \frac{V_2 P_2}{T_2} \quad \text{or } V_2 = 34.2 \text{ cm}^3$$

**Q58. Example 25:** A fire fighters air tank contains 12.0 dm<sup>3</sup> of air compressed to 1.40 x 10<sup>4</sup> KPa at 22°C. What volume of air will this tank provide when it is used in a hot, smoke-filled building where the temperature is 42°C and pressure 102 KPa?

**Solution:**

$$V = 12 \text{ dm}^3$$

$$V_2 = ?$$

$$P = 1.40 \times 10^4 \text{ KPa}$$

$$T_1 = 22^\circ\text{C} + 273 = 295\text{K}$$

Apply the equation for conversion factors

$$P_2 = 102 \text{ KPa}$$

$$T_2 = 42^\circ\text{C} + 273 = 315\text{K}$$

For comparison purposes, the Ideal Law equation can also be written as

$$\begin{aligned} V_2 &= \frac{V_1 P_1 T_2}{T_1 P_2} \text{ or } V_2 = V_1 \times \frac{T_2}{T_1} \times \frac{P_1}{P_2} \\ &= 12 \text{ dm}^3 \times \frac{315 \text{ K}}{295 \text{ K}} \times \frac{1.40 \times 10^4 \text{ KPa}}{10^2 \text{ KPa}} \\ &= 12 \times \frac{315}{295} \times 1.40 \times 10^4 \times 10^{-2} \text{ dm}^3 \\ &= 12 \times \frac{315}{295} \times 140 \text{ dm}^3 = 1793.89 \text{ dm}^3 \\ &= 1.79 \times 10^3 \text{ dm}^3 \end{aligned}$$

Result: The fire-fighter's air tank will provide 1.79 x 10<sup>3</sup> dm<sup>3</sup> of air in the hot building.

**Q59. Example 26:** Calculate the volume that 6.30 moles of CO<sub>2</sub> gas occupy at 23°C and 550 KPa pressure.

**Solution:**

$$V = ? \quad R = 8.31 \text{ KPa dm}^3 \text{ mole}^{-1} \text{ K}^{-1}$$

$$n = 6.30 \text{ moles}$$

$$T = 23^\circ\text{C} + 273 = 296\text{K}$$

$$P = 550\text{KPa}$$

Now according to the Ideal gas law equation

$$VP = nRT$$

$$= \frac{6.30 \times 8.31 \times 296 \text{ mole}^{-1} \cdot \text{KPa} \cdot \text{dm}^3 \cdot \text{K}^{-1} \cdot \text{Mole}^{-1} \times \text{K}^{-1}}{550} \cdot \frac{\text{KPa} \cdot \text{dm}^3 \cdot \text{K}^{-1} \cdot \text{Mole}^{-1} \times \text{K}^{-1}}{\text{KPa}}$$

$$V = 28.2 \text{ dm}^3$$

**Result:**

6.30 moles of  $\text{CO}_2$  occupy a volume of  $28.2 \text{ dm}^3$  at  $23^\circ\text{C}$  and  $550\text{KPa}$  pressure.

### Activity for students

**A person can inhale a maximum of 0.115 moles of air per breath. Calculate the maximum volume of air a person can inhale in one breath if the atmospheric pressure is  $100\text{KPa}$ , and the person's body temperature is  $31^\circ\text{C}$ .**

**Solution:**

$$n = 0.115 \text{ moles}$$

$$V = ?$$

$$P = 100\text{kPa} = 0.986\text{atm}$$

$$T = 37^\circ\text{C} + 273 = 310\text{K}$$

According to the general gas equation

$$VP = nRT$$

$$V = \frac{RT}{P} = \frac{0.115 \times 0.0821 \times 310}{0.986} = 2.96 \text{ dm}^3$$

**Q60. Example 27: If a  $0.75 \text{ dm}^3$  cylinder holds 6.5 moles of  $\text{N}_2$  gas at  $23^\circ\text{C}$ , calculate the Pressure in the cylinder.**

**Solution:**

$$V = 0.75 \text{ dm}^3$$

$$N = 6.5 \text{ moles}$$

$$T = 23^\circ\text{C} + 273 = 296\text{K}$$

$$R = 8.31 \text{ KPa dm}^3 \text{ K}^{-1} \text{ mole}^{-1}$$

$$p = ?$$

According to the Ideal Gas Law equation

$$VP = nRT$$

$$P = \frac{nRT}{V}$$

$$= \frac{6.5 \times 8.31 \times 296}{0.75} \cdot \frac{\text{KPa dm}^3 \text{ K}^{-1} \text{ Mole}^{-1} \times \text{K}^{\cancel{-1}} \times \text{K}^{\cancel{+1}}}{\text{dm}^3}$$

$$= 21318 \text{ KPa} \text{ Now } 101.325 \text{ KPa} = 1 \text{ atm}$$

$$21318 \text{ KPa} = \frac{1}{101.325} \times 21318.0 \text{ atm}$$

$$= 210.392 \text{ atm}$$

**Activity for students**

1.  $130 \text{ cm}^3$  of a gas exerts a pressure of 750 mm of Hg at  $20^\circ\text{C}$ . Calculate its pressure if its volume is increased to  $150 \text{ cm}^3$  at  $35^\circ\text{C}$ . (Ans: 683.28mm)

**Solution:**

$$V_1 = 130 \text{ m}^3$$

$$V_2 = 150 \text{ m}^3$$

$$P_1 = 750 \text{ mm Hg}$$

$$P_2 = ?$$

$$T_1 = 20^\circ\text{C} + 273\text{K} = 293\text{K}$$

$$T_2 = 35^\circ\text{C} + 273\text{K} = 308\text{K}$$

$$\frac{V_1 P_1}{T_1} = \frac{V_2 P_2}{T_2} \text{ or } P_2 = P_1 \times \frac{T_2}{T_1} \times \frac{P_1 V_1}{V_2}$$

$$P_2 = \frac{750 \times 130 \times 308}{293 \times 150} = 68.328$$

**2. find the total pressure exerted by 2 grams of ethane and 3grams of CO<sub>2</sub> contained in a dm<sup>3</sup> vessel at 50°C. (R = 0.082dm<sup>3</sup> atm K<sup>-1</sup> Mole<sup>-1</sup>)**

**Solution:**

Mass of Ethane given= 2g

Number of moles of ethane = 0.066 mol

Mass of CO<sub>2</sub> given= 3g

Number of moles of C= 0.068 mol

$$P_{\text{total}} = P_{\text{ethane}} + P_{\text{CO}_2}$$

$$P_{\text{ethane}} V = nRT$$

$$P_{\text{ethane}} = \frac{nRT}{V}$$

$$P_{\text{ethane}} = \frac{0.66 \text{ mol} \times 0.0821 \times 323 \text{ K}}{5} = 0.350 \text{ atm}$$

$$P_{\text{CO}_2} V = nRT$$

$$P_{\text{CO}_2} = \frac{nRT}{V}$$

$$P_{\text{CO}_2} = \frac{0.68 \times 0.0821 \times 323}{5} = 0.360 \text{ atm}$$

$$P_{\text{total}} = 0.350 + 0.360$$

$$P_{\text{total}} = 0.71 \text{ atm}$$

**Q61. Example 28: (a) Determine the volume that 2.5 moles of Cl<sub>2</sub> gas occupies at S.T.P**

**(b) How many individual chlorine molecules are there in the 2.5 moles of Cl<sub>2</sub>.**

**Solution:**

(a). One mole of  $\text{Cl}_2$  molecules at S.T.P =  $22.4 \text{ dm}^3$

$$2.5 \text{ mole of } \text{Cl}_2 \text{ molecule at S.T. P} = 22.4 \times 2.5 \\ = 56 \text{ dm}^3$$

(b). 1 mole of  $\text{Cl}_2$  molecules at S.T. P =  $6.022 \times 10^{23}$  molecules

$$2.5 \text{ moles of } \text{Cl}_2 \text{ molecule at S.T. P} = 2.5 \times 6.022 \times 10^{23} \text{ molecules} \\ = 1.51 \times 10^{23} \text{ molecules}$$

**Q62. Example 29: A  $2.60 \text{ dm}^3$  container was evacuated and its mass was found to be  $655.53 \text{ g}$ . The container was then filled with  $\text{CO}_2$  gas at a pressure of  $100 \text{ KP}$ , and a temperature of  $24^\circ\text{C}$ , the container and gas were found to have a mass of  $660.13 \text{ g}$ . Calculate the standard molar volume of  $\text{CO}_2$ .**

**Solution:**

$$V_{\text{CO}_2} = 2.60 \text{ dm}^3$$

$$m_1 \text{ (mass of container)} = 655.539 \text{ g}$$

$$m_2 \text{ (mass of container + gas)} = 660.13 \text{ g}$$

$$P_{\text{CO}_2} = 100 \text{ KPa}$$

$$T_{\text{CO}_2} = 24^\circ\text{C} + 273 = 297 \text{ K}$$

$$m_{\text{CO}_2} \text{ (mass of } \text{CO}_2) = 660.13 - 655.53 = 4.60 \text{ g}$$

$$\text{mass of } \text{CO}_2 = 660.13 - 655.53 - 4.60 \text{ g}$$

Convert the gas ( $\text{CO}_2$ ) to S.T.P by applying General Gas Equation

$$(V_1) V_{\text{CO}_2} = 2.60 \text{ dm}^3 \quad v_2 = ?$$

$$(P_1) P_{\text{CO}_2} = 100 \text{ KP}$$

$$(T_1) T_{\text{CO}_2} = 293 \text{ K}$$

$$P_2 = 101.3 \text{ KP} \quad T_2 = 273 \text{ K S.T.P}$$

Conversion from volume ( $V_2$ ) to the molar volume ( $V$ )

Standard Molar Volume ( $V_m$ ) = Molar Volume  $\times$  volume of  $\text{CO}_2$  at S.T.P.

$$= 9.5652 \text{ moles} \times 2.36 \text{ dm}^3$$

$$= 9.5652 \times 2.36 \text{ Moles} \cdot \text{dm}^3$$

$$= 22.6 \text{ dm}^3 \text{ Moles}$$

The accepted value for Molar value =  $22.4 \text{ dm}^3$

$$V_m = \frac{\text{Molar mass of CO}_2}{\text{Mass of CO}_2}$$

Mass of  $\text{CO}_2$

Thus, the conversion of volume of a gas to molar volume can be obtained by using the rates

**An empty container is filled with butane gas at 102KP. and  $20^\circ\text{C}$ . It was then found to have a mass of 66.184gm. Determine the Molar mass of butane. (Ans:  $272 \text{ dm}^3$ )**

**Solution**

$$p = 102 \text{ KP}_a = \frac{102000}{101325} = 1.006$$

$$T = 20^\circ\text{C} + 273 = 293 \text{ k}$$

$$m = 66.184 \text{ g}$$

$$n = 1.141 \text{ mol}$$

$$PV = nRT$$

$$V = \frac{nRT}{P}$$

$$V = 21.2 \text{ dm}^3$$

**Calculate the number of molecules atmospheric pressure and 10 grams of ammonia at 1.5 (Ans:  $3.60 \times 10^{23}$  molecules)**

**Solution:**

Mass of ammonia given = 10 g

Molar mass of ammonia = 17 g/mol

Number of moles of ammonia = 0.588 moles



1 mole of ammonia contains =  $6.02 \times 10^{23}$  molecules

0.588 moles of ammonia contains =  $0.588 \times 6.02 \times 10^{23}$  molecules

0.588 moles of ammonia contain =  $3.60 \times 10^{23}$  molecules

**Q63. Describe the effect of heating gases to extremely high temperature.**

**Ans: Effect of Heating Gases to Extremely High Temperature:**

A fourth type of matter i.e. Plasma State is formed when an extremely high temperature is applied to a molecular gas. The molecular gas is first converted to Atomic gas and ultimately to ions.

Molecular gas  $\rightarrow$  atoms gas  $\rightarrow$  ions.

Resulting Plasma consists of ions, positive or negative. The negatively charged ions i.e. electrons are mostly called as Plasma.



