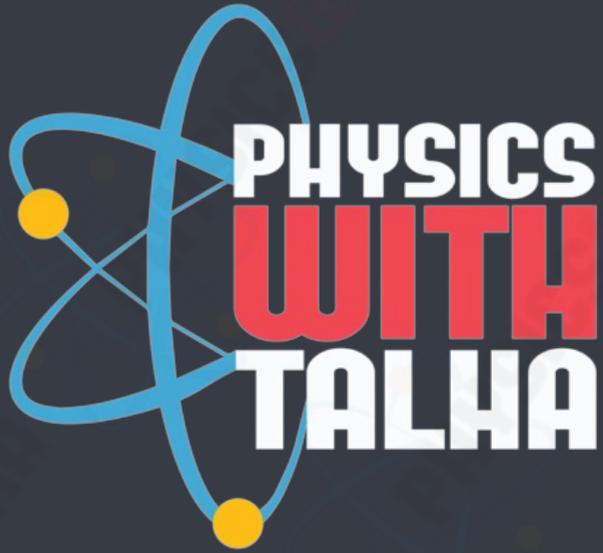


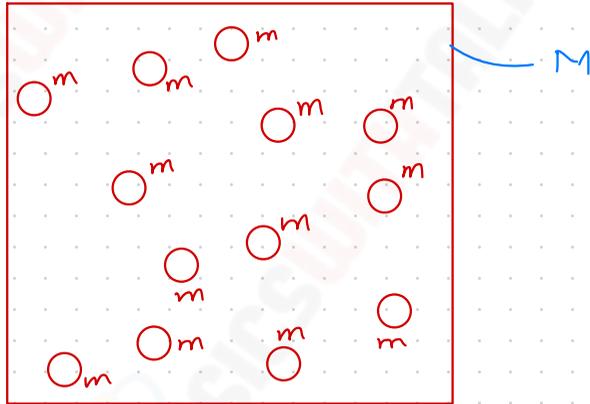
Ideal gases





The Mole

Suppose a gas is stored in a container



$m \rightarrow$ mass of each particle

$M \rightarrow$ mass of entire gas in the container

$N \rightarrow$ number of particles of gas

$n \rightarrow$ number of moles [unit = mol]

$N_A \rightarrow$ number of particles in one mole of a gas (Avogadro's constant)

$$N_A = 6.02 \times 10^{23} \text{ mol}^{-1}$$

We will look at gases made of a single element.
(e.g.: He, Ar, Ne, Xe, Kr, Rn)

Ar \rightarrow relative atomic mass : mass of one mole of a gas
(molar mass) [in grams]

$$n = \frac{M}{A_r}$$

e.g. Argon gas has a molar mass of 40 g.

A container has 150 g of Argon gas. How many moles of the gas are there in the container?

$$n = \frac{150}{40} = 3.75 \text{ mol}$$

$$M = m \times N$$

$$A_r = m \times N_A$$

$$N_A = 6.02 \times 10^{23}$$

$$\text{if } n = \frac{M}{A_r}$$

$$\text{then } n = \frac{m \times N}{m \times N_A}$$

$$\rightarrow n = \frac{N}{N_A}$$

Avagadro constant ($N_A = 6.02 \times 10^{23}$) : " the number of particles in 12 g of Carbon-12. "

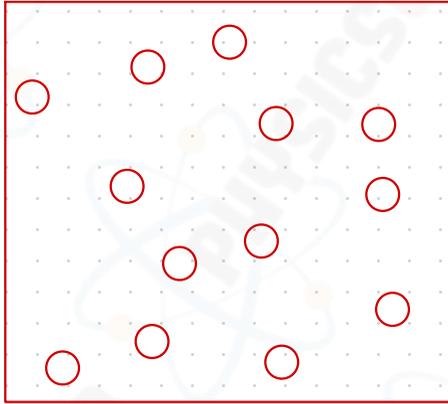
Mole : " amount of substance containing N_A (6.02×10^{23}) particles "

or " amount of substance containing the same number of particles as 12 g of carbon-12. "

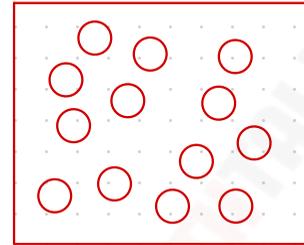
Gas laws

Three macro properties of gas are its pressure, volume, and temperature (in Kelvin)

1) Pressure vs Volume



P_1, V_1
 $1000 \text{ Pa}, 100 \text{ cm}^3$



P_2, V_2
 $2000 \text{ Pa}, 50 \text{ cm}^3$

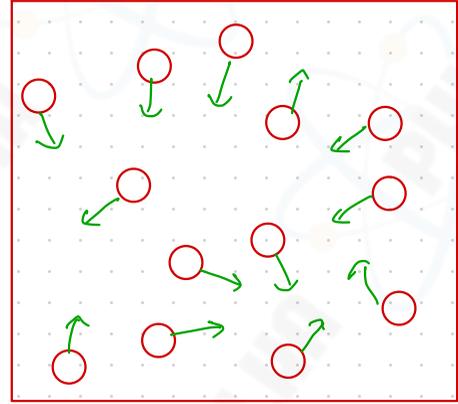
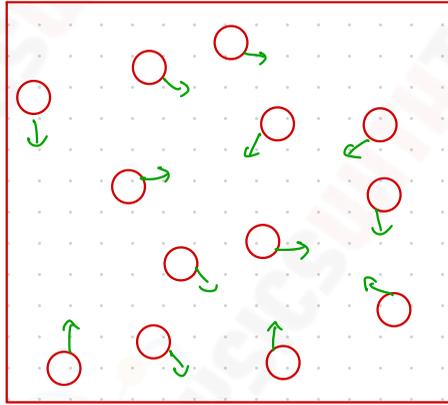
Assumption : Temp constant , mass constant .

$$P \propto \frac{1}{V} \longrightarrow PV = \text{constant}$$

$$P_1 V_1 = P_2 V_2$$



2) Pressure vs Temperature (in K)



$$1000 \text{ Pa} = P_1, \quad T_1 = 200 \text{ K}$$

Assumptions: Volume constant, mass constant

$$P \propto T$$



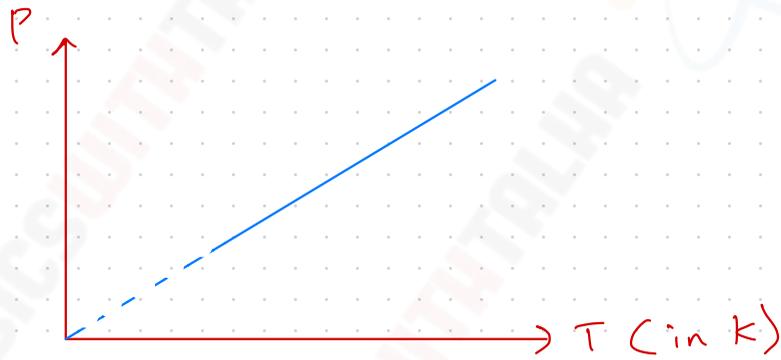
$$\frac{P}{T} = \text{constant}$$

$$\boxed{\frac{P_1}{T_1} = \frac{P_2}{T_2}}$$

↑↑↑
heat

$$P_2, T_2$$

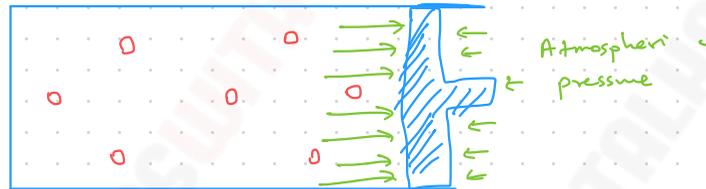
↓ ↓
2000 Pa, 400 K



3) Temperature (in K) vs volume



V_1, T_1
 $100\text{cm}^3, 300\text{K}$



↑↑↑
heat

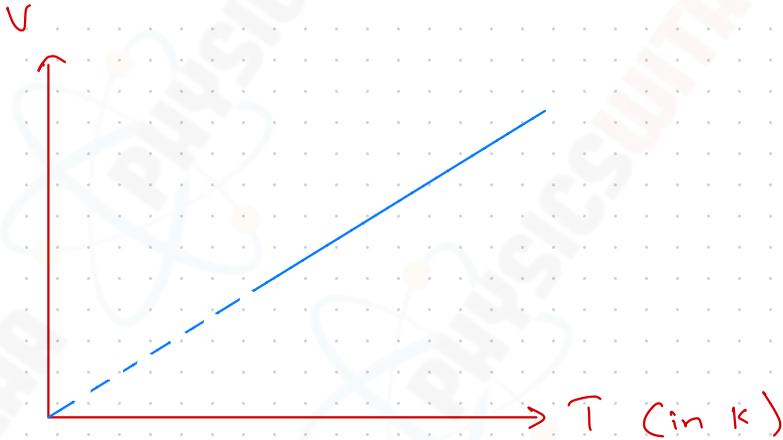


V_2, T_2 600K
 200cm^3

Assumption: pressure constant, mass constant.

$$V \propto T \quad \longrightarrow \quad \frac{V}{T} = \text{constant}$$

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



→ if mass (or number of moles) of gas is constant.

$$P_1 V_1 = P_2 V_2$$

$$, \frac{P_1}{T_1} = \frac{P_2}{T_2} ,$$

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

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

Ideal gas equation

→ 3 gas law can be combined to form an equation

$$PV = nRT$$

$R = 8.31 \text{ J K}^{-1} \text{ mol}^{-1}$ → universal gas constant
Temperature must be in Kelvin.

$$T (\text{K}) = T (^{\circ}\text{C}) + 273.15$$

→ we know that $n = \frac{N}{N_A}$

$$PV = \left(\frac{N}{N_A} \right) R T$$

R and N_A are both constants.
So $\frac{R}{N_A}$ is also a constant.

$$\frac{R}{N_A} = k \rightarrow \text{Boltzmann constant}$$

so

$$PV = NkT$$

$$k = 1.38 \times 10^{-23} \text{ J K}^{-1}$$

"Ideal gas is defined as a gas that obeys the ideal gas equation at all pressures, volumes, and temperatures."

→ Real gases are not ideal.

→ Ideal gas is just a model.

→ Real gases behave like ideal gas when :

1) Low density

2) Pressure is low

3) Temperatures are well above the liquification temperatures.

Kinetic Theory of gases

→ Gas is made up of tiny particles

→ The particles are always in motion.

→ There are 4 assumptions of kinetic theory of ideal gases.

- 1 ● All molecules behave as identical, hard, perfectly elastic spheres. (elastic collisions)
- 2 ● The volume of the molecules is negligible compared with the volume of the containing vessel. (dist. b/w particles \gg diameter of particles)
- 3 ● There are no forces of attraction or repulsion between molecules. (particles have no P.E)
- 4 ● There are many molecules, all moving randomly. (so averaging is possible)

→ Consider a particle of mass, m , moving with horizontal component of velocity c_x .

→ When it hits the wall, the velocity would be equal and opposite after rebounding. (1)

$$-c_x$$

→ To make a round trip and collide with the same wall again, the time needed:

$$t = \frac{\text{dist}}{\text{speed}} = \frac{2L}{c_x}$$

→ The change in momentum (x-component) of the particle:

$$\Delta p = p_f - p_i = (-mc_x) - (mc_x) = -2mc_x$$

for magnitude only, neg. sign is ignored

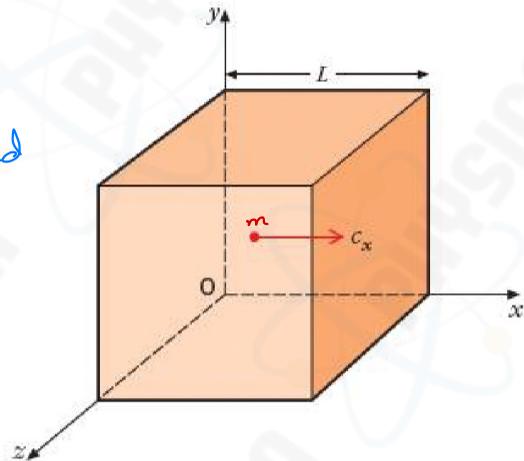


Figure 10.5 A gas molecule in a cubic container

The rate of change of momentum of the particle:

$$\frac{\Delta p}{\Delta t} = \frac{2mcx}{t} = \frac{2mcx}{\frac{2L}{cx}} = \frac{m}{L} cx^2$$

Force (x-comp.) exerted on (and by) the particle:

$$F = \frac{\Delta p}{\Delta t} = \frac{m}{L} cx^2$$

→ If no. of particles is N , the total average force exerted in x-component is:

$$\text{Total } F = \frac{N m c x^2}{L}$$

$$T_1 = 200^\circ\text{C}, 473\text{K}$$

$$T_2 = 210^\circ\text{C}, 483\text{K}$$

$$\Delta T = 10^\circ\text{C or K}$$

→ Area of the wall

$$A = L^2$$

Pressure on the wall

$$P = \frac{\text{Total } F}{A} = \frac{\left(\frac{Nm c_x^2}{L} \right)}{L^2} = \frac{Nm c_x^2}{L^3}$$

Volume of container

$$V = L^3$$

So pressure becomes:

$$P = \frac{Nm c_x^2}{V}$$

* Above expression is for x-component only. The particle will have x, y, and z component of velocity

$$c^2 = c_x^2 + c_y^2 + c_z^2 \quad (\text{Pythagoras Theorem})$$

Because we are dealing with a large number of particles and their motion is random (4), so we can assume their averages of squares are equal

$$\langle C_x^2 \rangle = \langle C_y^2 \rangle = \langle C_z^2 \rangle$$

So average square velocity will be :

$$\begin{aligned}\langle C^2 \rangle &= \langle C_x^2 \rangle + \langle C_y^2 \rangle + \langle C_z^2 \rangle \\ &= \langle C_x^2 \rangle + \langle C_x^2 \rangle + \langle C_x^2 \rangle\end{aligned}$$

$$\langle C^2 \rangle = 3 \langle C_x^2 \rangle$$

↙
mean square velocity

Rearranging :

$$\frac{\langle C^2 \rangle}{3} = \langle C_x^2 \rangle$$

So pressure becomes :

$$P = \frac{Nm \langle c^2 \rangle}{3V}$$

→

$$PV = \frac{Nm \langle c^2 \rangle}{3}$$

Total mass = Nm

$$\text{Density } \rho = \frac{\text{Total mass of gas}}{\text{Volume}} = \frac{Nm}{V}$$

$$P = \frac{Nm \langle c^2 \rangle}{3V}$$

So pressure becomes

$$P = \rho \frac{\langle c^2 \rangle}{3}$$

The avg K.E is :

$$\langle K.E \rangle = \frac{1}{2} m \langle c^2 \rangle$$

$$\text{So total K.E is} = \frac{1}{2} N m \langle c^2 \rangle$$

$$30^\circ\text{C}$$

1

$$30^\circ\text{C}$$

2

Ideal gas equation was : $PV = NkT$

So r.h.s of pressure equation and ideal gas equation can be equated

$$NkT = \frac{N m \langle c^2 \rangle}{3}$$

divide both sides by 2

$$\frac{1}{2} kT = \frac{1}{2} \times \frac{m \langle c^2 \rangle}{3}$$

$$\frac{1}{2} kT = \frac{\langle K.E \rangle}{3}$$

* Avg K.E of a molecule

$$\langle K.E \rangle = \frac{3}{2} kT$$

$$\text{Total K.E} = \frac{3}{2} NkT$$

Also ;

$$\frac{1}{2} m \langle c^2 \rangle = \frac{3}{2} kT$$

$$\sqrt{\langle c^2 \rangle} = \sqrt{\frac{3kT}{m}}$$

↳ root mean square velocity

$$\text{r.m.s speed} = \sqrt{\frac{3kT}{m}}$$

$$\text{r.m.s speed} \propto \sqrt{T}$$

$$\frac{\text{r.m.s speed}_A}{\sqrt{T_A}} = \frac{\text{r.m.s speed}_B}{\sqrt{T_B}}$$