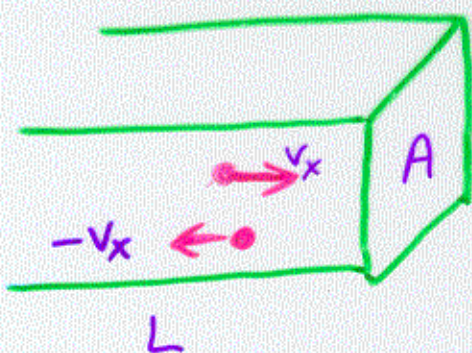


Pressure, Temp and Molecular Speeds

Gas container, side L , area of wall $= A$:



Molecule mass m
speed v
hits wall.

Momentum change $\Delta p = 2m v_x$ per collision

Time Δt between collisions with wall $= \frac{2L}{v_x}$

$$\Rightarrow \text{force on wall (Newton II)} \quad F = \frac{\Delta p}{\Delta t} = 2m v_x \cdot \frac{v_x}{2L} = \frac{m v_x^2}{L}$$

$$\text{Total pressure on wall } P = \frac{\sum F}{A} = \frac{\sum m v_x^2}{A \cdot L} = \frac{m \sum v_x^2}{V}$$

Now in 3-d, $v^2 = v_x^2 + v_y^2 + v_z^2$ so on average $v_x^2 = \frac{1}{3} v^2$

Define RMS speed as $v_{rms}^2 = \frac{\sum v^2}{N}$

$$\Rightarrow P = \frac{1}{V} \frac{Nm v_{rms}^2}{3} \Rightarrow \underline{P \cdot V = \frac{1}{3} Nm v_{rms}^2}$$

$$\text{c.f. } PV = N k_B T \Rightarrow v_{rms}^2 = \frac{3 k_B T}{m}$$

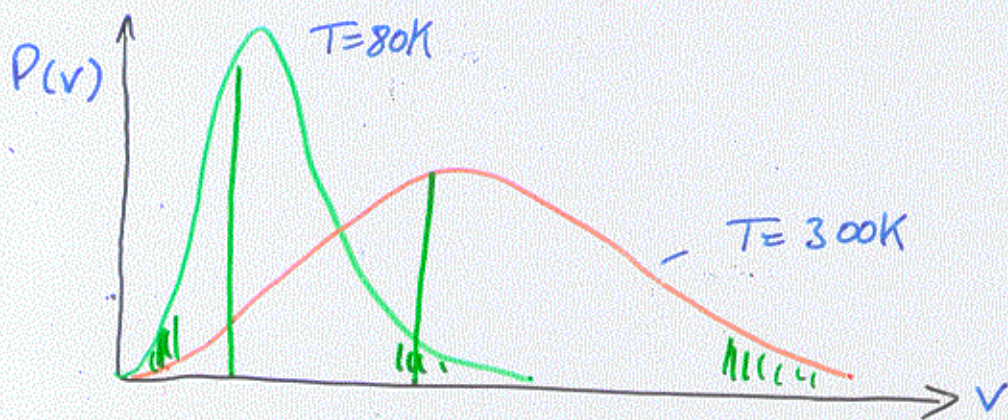
OR average k.e. of molecule $= \frac{3}{2} k_B T$
 $\frac{1}{2} m v_{rms}^2$

Details, Details1. Mean free path (sec 20.6)

- In fact molecules collide frequently, exchanging k.e. with each other.

"Mean free path" = av. distance traveled between collisions. $\lambda \propto \frac{1}{\text{density}}$: $\sim 0.1 \mu\text{m}$ at sea level
 $\sim 16\text{cm}$ ~~at~~ **100km**

2. Ideal gases have a distribution of molecular speeds :
 (sec 20.7, fig 20.7)



Maxwell speed distribution changes with T

$v_{\text{rms}}^2 = \frac{3k_B T}{m}$ but many molecules have $v \gg v_{\text{rms}}$

From Kinetic Theory: Average k.e. of molecule $\frac{1}{2} m v_{rms}^2 = \frac{3}{2} k_B T$

e.g. At room temp ($T = 300K$):

$$\text{k.e. of any molecule} = \frac{3}{2} k_B T = \frac{3}{2} \times 1.38 \times 10^{-23} \times 300 \\ = 6.2 \times 10^{-21} \text{ J.}$$

So for helium ^4He , $\frac{1}{2} m v_{rms}^2 = 6.2 \times 10^{-21} \text{ J}$

$$\Rightarrow v_{rms} = \sqrt{\frac{3 k_B T}{m}} = \underline{1366 \text{ m/s}}, \gg \text{escape speed from earth's gravity!}$$

$$\text{for } O_2 \text{ (molecular mass} = 32) \quad v_{rms} \propto \frac{1}{\sqrt{m}} = \underline{483 \text{ m/s}}$$

Internal Energy of a Monatomic Gas

E_{int} = total k.e. of molecules (no vibration, rotation)

$$E_{int} = \frac{3}{2} N k_B T = \frac{3}{2} n \underline{(N_A k_B)} T = \frac{3}{2} n \underline{R} T$$

So for 1 mole of any gas at room temp

$$\underline{E_{int} = 3738.4 \text{ J}} \quad (\text{monatomic})$$

E_{int}, Heat and Work: Example

e.g. 2 moles of gas heated at constant pressure $P = 200 \text{ kPa}$ from 300 to 400 K.

$$\underline{PV = nRT} \quad \underline{dE_{int} = Q - W} \quad \xrightarrow{=} \begin{matrix} = \frac{3}{2} PV \\ E_{int} = \frac{3}{2} nRT \end{matrix}$$

a) Initial state: $P_i = 200 \text{ kPa}, T_i = 300 \text{ K}, n = 2 \Rightarrow V_i = \frac{2 \times 8.31 \times 300}{200 \times 10^3} = 0.025 \text{ m}^3$

Final state: $V \propto T \Rightarrow V_f = 0.033 \text{ m}^3$

b) Work done $W = \int P \cdot dV = P(V_f - V_i) = 1666.7 \text{ J}$

c) $\Delta E_{int} = \frac{3}{2} nR (T_f - T_i) = \frac{3}{2} P (V_f - V_i) = 2500 \text{ J}$

d) $\therefore$ Heat input required $Q = \underline{dE_{int}} + W = 2500 + 1666.7 = 4167.7 \text{ J}$



So here, 2500 J of heat $\rightarrow$ internal energy (raises temp of gas)
 1667.7 J $\rightarrow$ does actual work. = mgh .