

Probability Distributions and Gases

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1 Probability Distributions

1.1 Statistics on distributions

- (a) We simply integrate the distribution over all possible values of v and enforce that it must equal one (to be a valid probability distribution):

$$\int_{-\sqrt{a/b}}^{\sqrt{a/b}} f(v) dv = 1 \implies \left[Cav - \frac{Cb}{3} v^3 \right]_{-\sqrt{a/b}}^{\sqrt{a/b}} = 1 \quad (1)$$

Plugging in the numbers, we have

$$Ca(2\sqrt{a/b}) - \frac{Cb}{3} \frac{a}{b} (2\sqrt{a/b}) = 1$$
$$C = \frac{3}{4a} \sqrt{\frac{b}{a}}$$

- (b) To compute average velocity, we integrate the distribution weighted by the velocity over the full range, using the value for C that we found in part (a):

$$\begin{aligned} \bar{v} &= \int_{-\sqrt{a/b}}^{\sqrt{a/b}} v f(v) dv \\ &= \int_{-\sqrt{a/b}}^{\sqrt{a/b}} (Cav - Cbv^3) dv \\ &= \left[\frac{Ca}{2} v^2 - \frac{Cb}{4} v^4 \right]_{-\sqrt{a/b}}^{\sqrt{a/b}} \\ &= \frac{Ca}{2} \left(\frac{a}{b} - \frac{a}{b} \right) - \frac{Cb}{4} \left(\left(\frac{a}{b} \right)^2 - \left(\frac{a}{b} \right)^2 \right) \\ &= 0 \end{aligned}$$

Equivalently, we could have immediately concluded $\bar{v} = 0$ m/s by simply saying integrating an odd function over a domain symmetric about the origin always yields zero.

- (c) We proceed exactly as in part (b), except with v^2 as our weight on the probability distribution:

$$\begin{aligned}
 \bar{v}^2 &= \int_{-\sqrt{a/b}}^{\sqrt{a/b}} v^2 f(v) dv \\
 &= \int_{-\sqrt{a/b}}^{\sqrt{a/b}} (Cav^2 - Cbv^4) dv \\
 &= \left[\frac{Ca}{3} v^3 - \frac{Cb}{5} v^5 \right]_{-\sqrt{a/b}}^{\sqrt{a/b}} \\
 &= C \left[\frac{a}{3} \sqrt{\frac{a}{b}} \frac{2a}{b} - \frac{b}{5} \sqrt{\frac{a}{b}} \frac{2a^2}{b^2} \right] \\
 &= \left(\frac{3}{4a} \sqrt{\frac{b}{a}} \right) \left(\frac{2a^2}{b} \sqrt{\frac{a}{b}} \right) \left(\frac{1}{3} - \frac{1}{5} \right) \\
 &= \frac{3a}{2b} \frac{2}{15} \\
 &= \frac{a}{5b}
 \end{aligned}$$

- (d) Quite simply, the number distribution is the probability distribution of the velocities multiplied by the number of particles:

$$N(v) = Nf(v) \quad (2)$$

1.2 Concrete numbers

- (a) Proceeding formulaicly:

$$\begin{aligned}
 \bar{v} &= \frac{2 * 10 + 4 * 12 + 2 * 14 + 1 * 15 + 1 * 17}{10} \\
 &= 12.8 m/s \\
 \sqrt{\bar{v}^2} &= \sqrt{\frac{2 * 10^2 + 4 * 12^2 + 2 * 14^2 + 1 * 15^2 + 1 * 17^2}{10}} \\
 &= 12.96 m/s \\
 &\approx 13.0 m/s
 \end{aligned}$$

- (b) The average kinetic energy is related to both temperature and the average squared speed velocity via

$$\begin{aligned}
 \bar{K} &= \frac{1}{2} m \bar{v}^2 = \frac{3}{2} k_B T \\
 T &= \frac{1}{3} \frac{m}{k_B} \bar{v}^2 \\
 &\approx \frac{13^2}{3} \frac{m}{k_B} \times \frac{m^2}{s^2}
 \end{aligned}$$

Note that I explicitly keep the units of $m^2 s^{-2}$ since otherwise they would be lost with the 13^2 being put into v^2 .

1.3 Setting up expressions

- (a) Let $f(v)$ be the usual Maxwell distribution. Then,

$$0.1 = \left(\int_{\alpha}^{\infty} f(v) dv \right) / N \quad (3)$$

- (b) This is given by

$$p_E = \frac{\int_{\alpha}^{\infty} v^2 f(v) dv}{\int_0^{\infty} v^2 f(v) dv} \quad (4)$$

- (c) $N \rightarrow 0.9N$ and $E \rightarrow 0.72E$, so with $T_0 = \frac{2}{3k_B} \frac{E}{N}$ and $T_1 = \frac{2}{3k_B} \frac{0.72E}{0.9N}$, we have

$$\begin{aligned} T_1 &= \frac{0.72}{0.9} T_0 \\ &= \frac{8}{10} T_0 \end{aligned}$$

- (d) We want $\left(\frac{8}{10}\right)^n T_0 < \frac{1}{2} T_0$, where n is the number of evaporative cooling cycles:

$$\begin{aligned} \left(\frac{8}{10}\right)^n &< \frac{1}{2} \\ n \log\left(\frac{8}{10}\right) &< \log\left(\frac{1}{2}\right) \\ -n \log\left(\frac{10}{8}\right) &< -\log(2) \\ n &> \log(2) / \log\left(\frac{10}{8}\right) \\ n &> 3.1 \end{aligned}$$

Hence, we must go through at least four cycles.

1.4 Puddles!

Puddles evaporate because the molecules move with a distribution of velocities and the ones with the top velocities could have enough to escape the surface tension of the puddle and go into the atmosphere. In a sealed jar, the water cannot evaporate because those fast molecules that escape will “bounce” off the walls and go right back in.

1.5 Building intuition

As N increases (keeping T constant), the distribution just gets vertically stretched since it is a number distribution, so this really isn't terribly insightful. As T increases (keeping N constant), the distribution gets a smaller peak but the tail increases, so you get on average faster molecules as the tail with the high speed ones becomes more probable/populated.