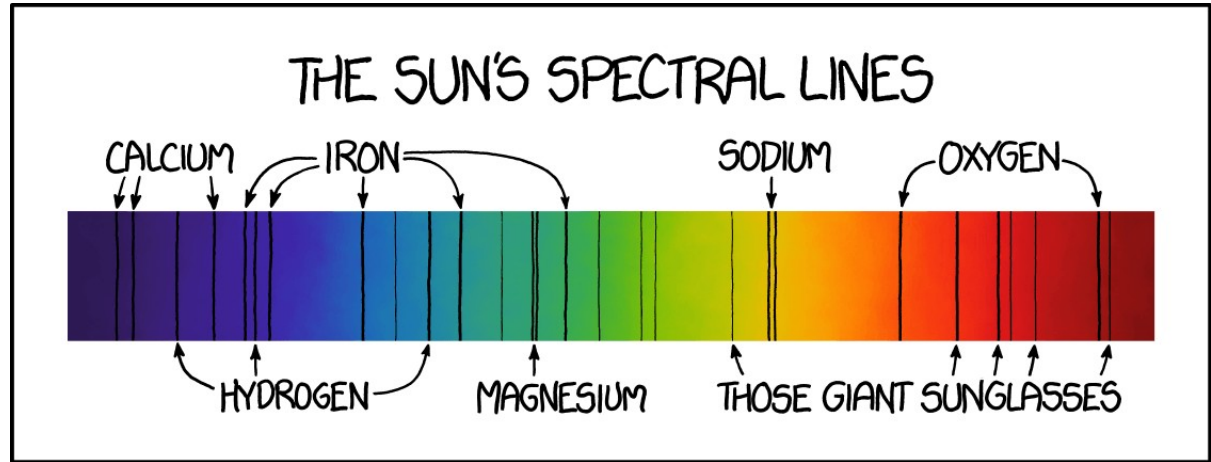
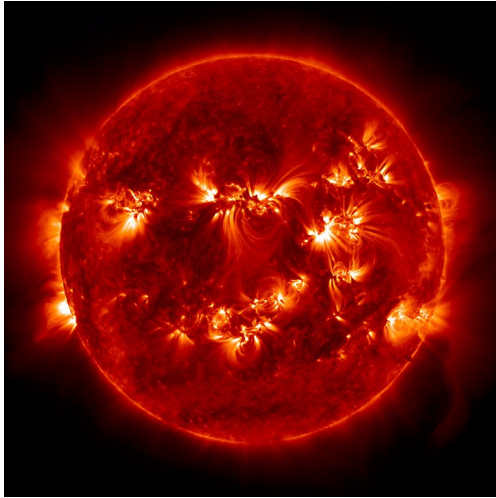


# PHYS 7810 / COLLAGE2021: Solar Spectral Line Diagnostics



## Lecture 05: Opacity and Emissivity in Spectral Lines (cont)

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## Plan for today:

- On Tuesday we entered the discussion talking about emissivity and deriving an expression
- Today we are first going to do the same for opacity (but much faster)
- Then we are going to investigate some relationships between the two
- And to analyze, through equations how each physical parameter ( $T$ ,  $p$ ,  $\mathbf{v}$ ,  $\mathbf{B}$ ), influences opacity/emissivity.

Once again, this is equation that includes it all

This whole equation tells us how the things behave on big scales. ( $dl$  is geometrical length, and light propagates over very large distances inside of astrophysical objects.

$$\frac{dI(\vec{r}, \mathbf{n}, \lambda)}{dl} = j(\vec{r}, \mathbf{n}, \lambda) - \chi(\vec{r}, \mathbf{n}, \lambda)I(\vec{r}, \mathbf{n}, \lambda)$$

These two coefficients, on the other side, depend on specific physics of emission and absorption. **We talk about them now**

# Emissivity due to spectral line transitions

$$j_{\lambda} = \frac{hc}{4\pi\lambda} n_u A_{ul} \phi_{ul,\lambda}$$

- This is only due to **bound-bound** processes, other sources would look differently
- Finding the number density of the atoms that are in the upper energy state (population of the upper level) will be the most cumbersome task.
- Also, line emission/absorption profile (often the same) contains many dependencies

# The absorption coefficient (opacity)

- We can have a very similar story here:  $\frac{dI_{\lambda}^{\text{abs}}}{dl} = -\chi_{\lambda}I_{\lambda}$
- The intensity should have units of inverse length. What if we relate it somehow to number density of absorption events (absorbers?)

$$\chi_{\lambda}[\text{m}^{-1}] = n^{\text{absorbers}}[\text{m}^{-3}] \times \sigma[\text{m}^2]$$

- This is more “classical” than the previous argument, but it very intuitively tells us what opacity depends on!

Let's write an equation for the opacity by analogy

$$\chi_{\lambda} = \frac{hc}{4\pi\lambda} n_l B_{lu} \phi_{ul, \lambda}$$

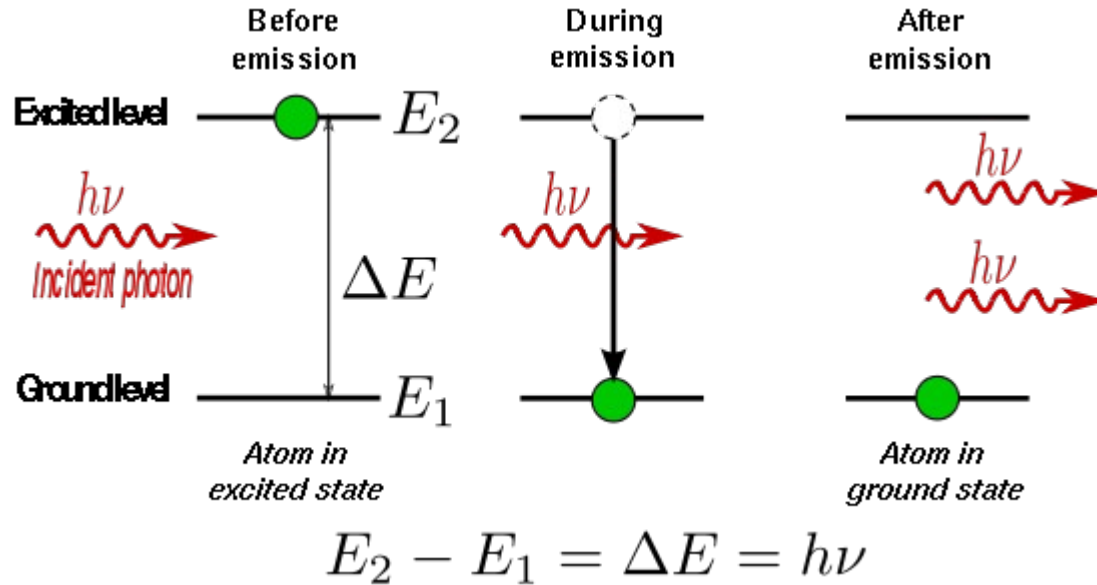
- Does this make sense?
- What changed with respect to emissivity?
- Can you figure out what units for  $B_{lu}$  have to be?

Let's write an equation for the opacity by analogy

$$\chi_{\lambda} = \frac{hc}{4\pi\lambda} n_l B_{lu} \phi_{ul, \lambda}$$

- Does this make sense?
- What changed with respect to emissivity?
- Units for  $B_{lu}$  have to be  $1/(\text{s} \times \text{Units of Intensity})$       $B_{lu} \times I = R_{\text{abs}}$

# What about stimulated emission though?



- Should we not include this as an emission source too?
- Still, it's a bit awkward, it depends on the direction, it scales with incoming intensity... it's almost as if it...



Stimulated emission is negative absorption!

$$\chi_{\lambda} = \frac{hc}{4\pi\lambda} (n_l B_{lu} - n_u B_{ul}) \phi_{\lambda}$$

- Wait Ivan, does that mean that the absorption can be negative?
- Can it?
- Can it?

# Yes! This is how lasers work!

- To obtain lasing we need to obtain the so called population inversion (not necessary inversion, depends on the values for B)
- We also have naturally occurring lasers, they are called MASERs (Microwave Amplification by Stimulated Emission of Radiation).
- This does not happen (at least not to my knowledge) in the atmosphere of the Sun that much

Now we know our opacity and emissivity

$$j_{\lambda} = \frac{hc}{4\pi\lambda} n_u A_{ul} \phi_{\lambda}$$

$$\chi_{\lambda} = \frac{hc}{4\pi\lambda} (n_l B_{lu} - n_u B_{ul}) \phi_{\lambda}$$

- We can now calculate optical depth and the source function, right?

Let's start with the source function

$$S = \frac{n_u A_{ul}}{n_l B_{lu} - n_u B_{ul}}$$

- The Source function in the line does not depend on the wavelength
- (This is generally not true as we have other op/em sources, but can be really good approximation)
- So, why don't we talk about these level populations a little bit? How to calculate them?

# Boltzmann excitation distribution

$$n_i = n \frac{g_i e^{-E_i/kT}}{Z}$$

Total number density of the atom (ion)

Energy with respect to the ground level

Population of the level i

Partition function

- If we know the total number of the atom (ion), and the temperature, we can calculate the population of each of the atomic levels. Easy enough.

# Boltzmann excitation distribution

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- If we know the total number of the atom (ion), and the temperature, we can calculate the population of each of the atomic levels. Easy enough.
- There is a slight problem. What we usually work with is total number of particles, say, hydrogen atoms. However... what figures in the above equation is the number of **neutral** hydrogen atoms.

So I am going to slightly re-write it

Total number density of the atoms in ionization state j

Energy of the state i in the ionization state j

$$n_{j,i} = n_j \frac{g_{j,i} e^{-E_{j,i}/kT}}{Z_j}$$

Population of the level i, in the ionization state j

- But this leads to another the question: How to find the number density of given ion.

*Example: How many of the total hydrogen atoms are neutral?*

# Saha ionization equation

Number density of the ions in the next ionization state

Energy needed to ionize ion j

$$n_j = C_I \times n_{j+1} n_e \frac{Z_j}{Z_{j+1}} \frac{e^{E_j/kT}}{T^{3/2}}$$

Number density of the ions in the ionization state j

Electron number density

- Let's spend a couple of minutes to see if this equation makes sense
- It of course complicates things, now there is this annoying electron density here.



Let's try and figure out how to apply this to an example:

- Number density of atomic hydrogen atoms is  $10^{22} \text{ m}^{-3}$ . Temperature of the plasma is 6000 K. Using Saha ionization equation try to find the number density of electrons and protons. You can symbolically represent Saha equation as:

$$n_j = n_{j+1}n_e \times f(T)$$

Let's try and figure out how to apply this to an example:

- Number density of atomic hydrogen atoms is  $10^{22} \text{ m}^{-3}$ . Temperature of the plasma is 6000 K. Using Saha ionization equation try to find the number density of neutral hydrogen and protons. You can symbolically represent Saha equation as:

$$n_j = n_{j+1} n_e \times f(T)$$

$$n_0 + n_+ = 10^{22}$$

$$n_+ = n_e$$

Does this help a bit?

# Solving Saha ionization equilibrium

- However, in general, we don't have only one element, but many
- At first it might seem that Hydrogen is the most important one but....

# Solving Saha ionization equilibrium

- However, in general, we don't have only one element, but many
- At first it might seem that Hydrogen is the most important one but:  
**Metals, despite being much less abundant, have much smaller ionization potential, and are thus more likely ionized. So, metals are the ones who are determining the electron density of the gas.**
- Technically all the species are "coupled", and for given point in the atmosphere we have to solve Saha equation for all the elements (species) simultaneously

# Equation of state

- Generally, by equation of state, I mean a relationship that connects, **pressure, mass density and temperature**
- From any two of these we should be able to calculate the third. Say we have **pressure and temperature:**
- Use  $p = nkT$ , to find the total number density of all the particles
- Use Saha ionization equation to calculate number densities of each constituent
- Add them with their masses to find **mass density**.

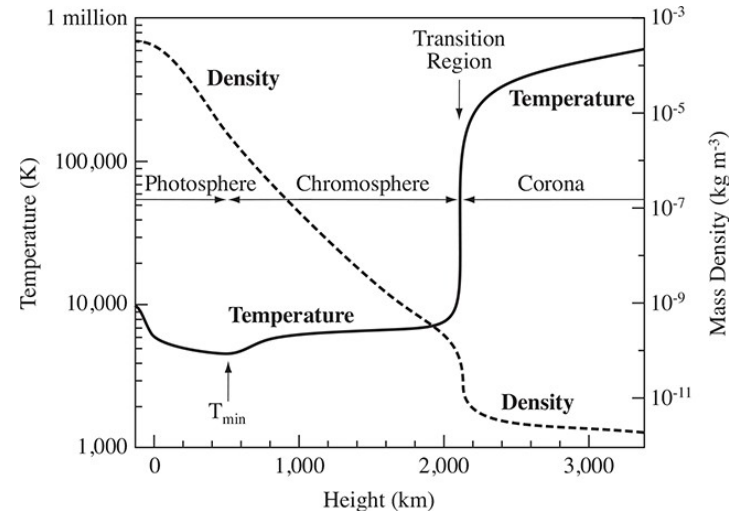
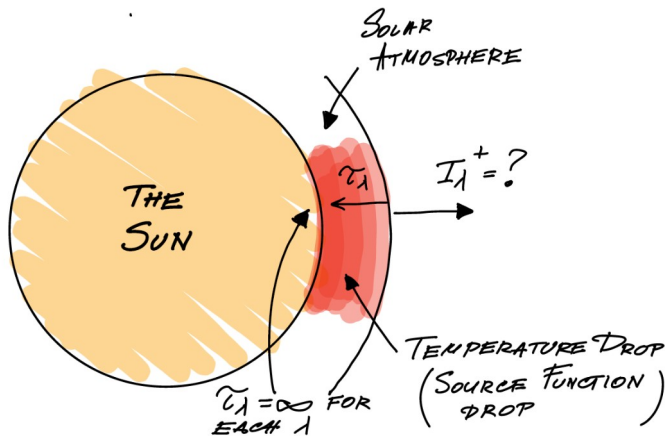
So, to recoup:

If we know the temperature and the pressure in the gas, we can solve Saha and then Boltzmann equation to find population of each relevant level.

It's cumbersome, but doable.

# Disclaimer

- Saha-Boltzmann equation only works in equilibrium.
- We need sufficient number of collisions between the particles to establish that equilibrium
- This means we need “dense” plasma (well, dense, for astrophysical contexts ;)
- In tenuous plasma, situation is even more complicated.



But, I told this long story because of this:

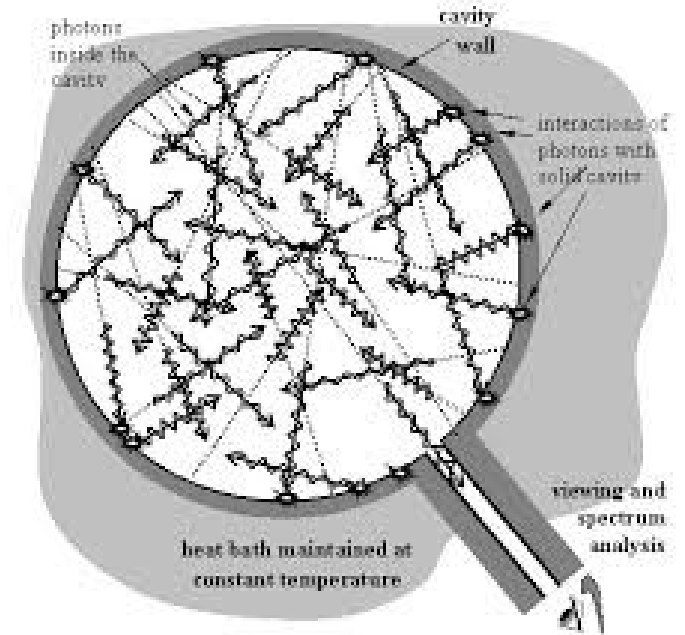
$$S = \frac{n_u A_{ul}}{n_l B_{lu} - n_u B_{ul}}$$

- We will now forget about the levels and everything else and go back to what Gustav Kirchhoff concluded about this (it seems like already he knew about RTE!)



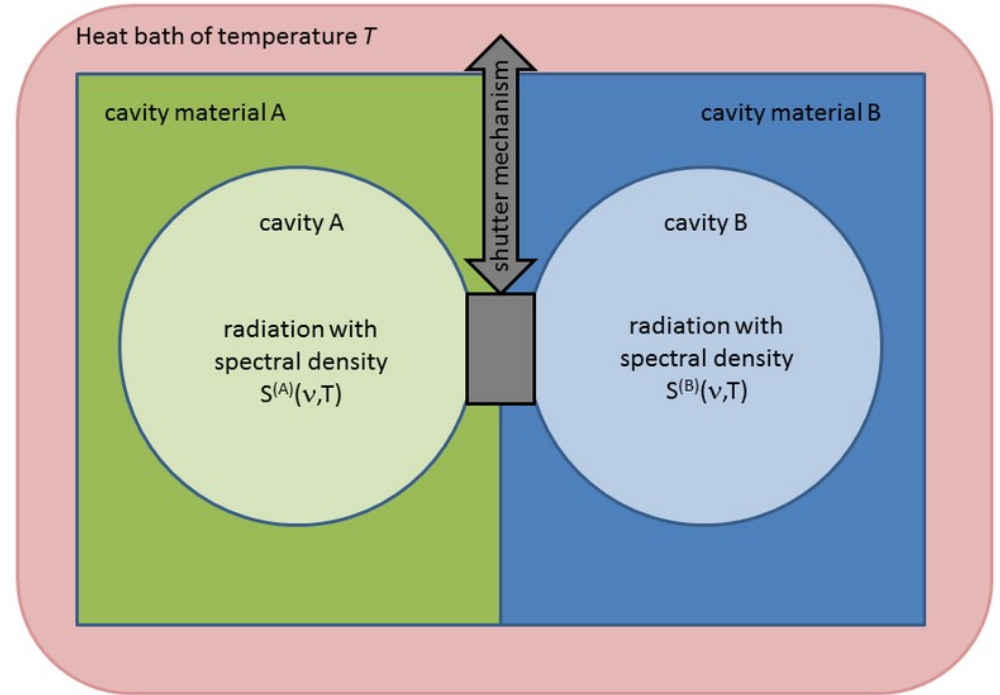
# Blackbody radiation

- Usually imagined as a “cavity”, an empty hole where photons are trapped and brought to equilibrium with surrounding walls
- The distribution of the photons according to energy (wavelength), and thus the emergent specific monochromatic intensity must depend on the temperature only



# Blackbody radiation

- Kirchoff proved this by a thought experiment (how else :-))
- Imagine an obstacle that allows only a specific wavelength to pass by,  $T$  is the same in both halves
- If this “spectral density” (intensity), was different, there would be net flow of the photons, and thus energy.
- But then one part would heat up!



So Kirchhoff said:

- It seems that the intensity is universal function of wavelength and T:

$$I_\lambda = B(\lambda, T)$$

- And then he substituted this in the RTE:

$$\frac{dI_\lambda}{dl} = j_\lambda - \chi_\lambda I_\lambda$$

$$0 = \frac{dB_\lambda}{dl} = j_\lambda - \chi_\lambda B_\lambda$$

$$\frac{j_\lambda}{\chi_\lambda} = B_\lambda$$

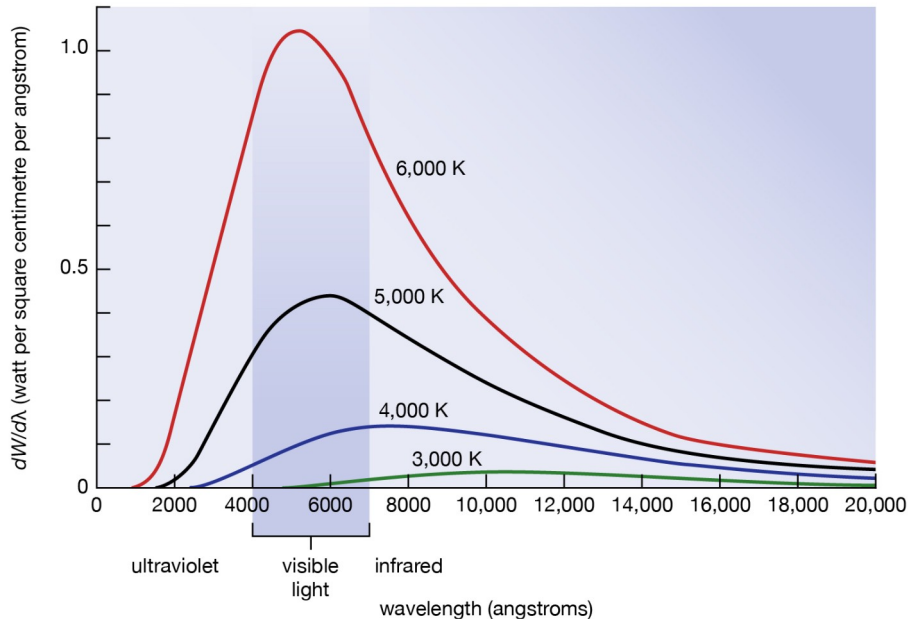
Kirchhoff's law. Does not care about where em/op come from!

So what Kirchhoff actually said:

“It is a highly important task to find this function”

# Which Max Planck (and Bose, and Einstein, and ...) did

- Blackbody radiation describes distribution of photons over energies, but also the source function of the matter in the (local) equilibrium
- This might require some unpacking ;)



$$B_{\lambda} = \frac{2hc^2}{\lambda^5} \frac{1}{e^{hc/\lambda kT} - 1}$$

## Putting together Boltzmann distribution and Planck Law:

$$S_\lambda = \frac{n_u A_{ul}}{n_l B_{lu} - n_u B_{ul}} \quad n_i = n \frac{g_i e^{-E_i/kT}}{Z} \quad S_\lambda = B_\lambda = \frac{2hc^2}{\lambda^5} \frac{1}{e^{hc/\lambda kT} - 1}$$

- We can find relationships between the Einstein parameters
- (This will actually be your homework)
- But there is a deeper reasoning here...

# Local thermodynamic equilibrium

- If the matter is locally in the equilibrium (Saha-Boltzmann distribution is valid), the source function is equal to the Planck function.
- This **does not** mean that the intensity is equal to Planck function too!
- This is quite some assumption, which is valid only when collisions dominate and we can neglect influence of the radiation on the matter (so only matter influences the radiation)
- We will talk about relaxing this. That is called **non-local thermodynamic equilibrium** (weird name since non- negates the whole statement)

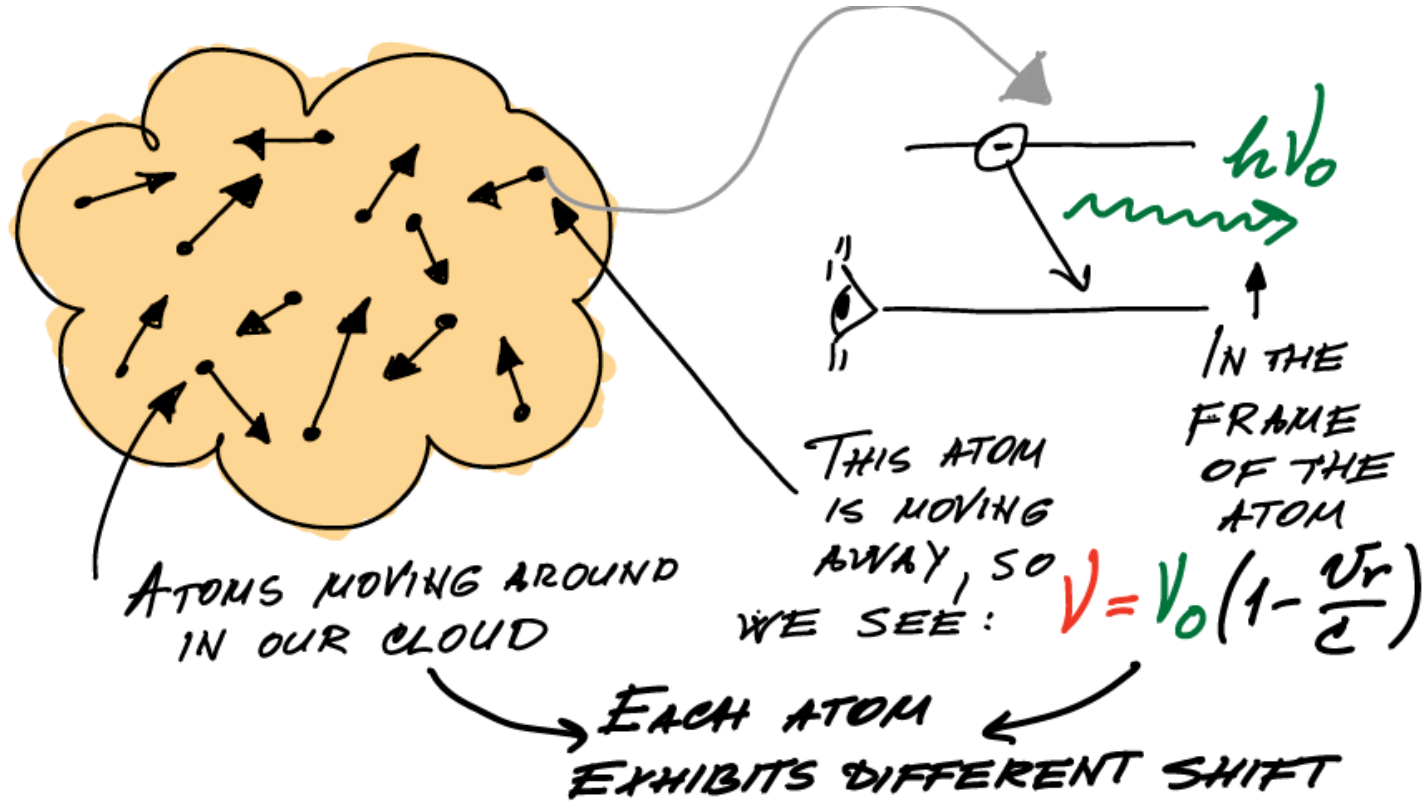
# Line absorption (and emission) profile

- We are covered on the source function part, and it seems simple enough now.
- But we need to calculate the optical depth, for that we need opacity, and for that we need the line absorption profile:

$$\chi_\lambda = \frac{hc}{4\pi\lambda} (n_l B_{lu} - n_u B_{ul}) \phi_\lambda$$



# Simple spectral line broadening



The line-of-sight velocity distribution follows Gaussian distribution. Then, when we sum up emissions (absorptions) by all the particles, Doppler-shifted frequency (and wavelength) will follow Gaussian Distribution too!

# The other way to write this would be:

- Each atom absorbs as a delta function (infinitely thin levels)
- But we look at large ensembles of atoms:

Delta function shifted  
due to Doppler effect

$$\phi_\lambda = \int_{-\infty}^{\infty} \delta\left(\lambda_0\left(1 - \frac{v}{c}\right)\right) p(v) dv$$

“Average” line profile

Probability of the atom  
having that velocity (this  
is radial, or los velocity)

- This is how you would get Gaussian!

## But then, the levels are **not** completely sharp

- **Natural (radiative) broadening** : Levels themselves are not perfectly sharp. The shorter is the lifetime of the level, the more smeared it is.
- **Pressure broadening (damping)**: Other particles exert force on our atom and “deform” its levels, this leads to so called pressure or collisional damping. This can even make the level be “asymmetrical” in a way
- Combination of the two yields the infamous **Voigt profile**

Lorentzian profile, due to damping,  
shifted due to Doppler effect

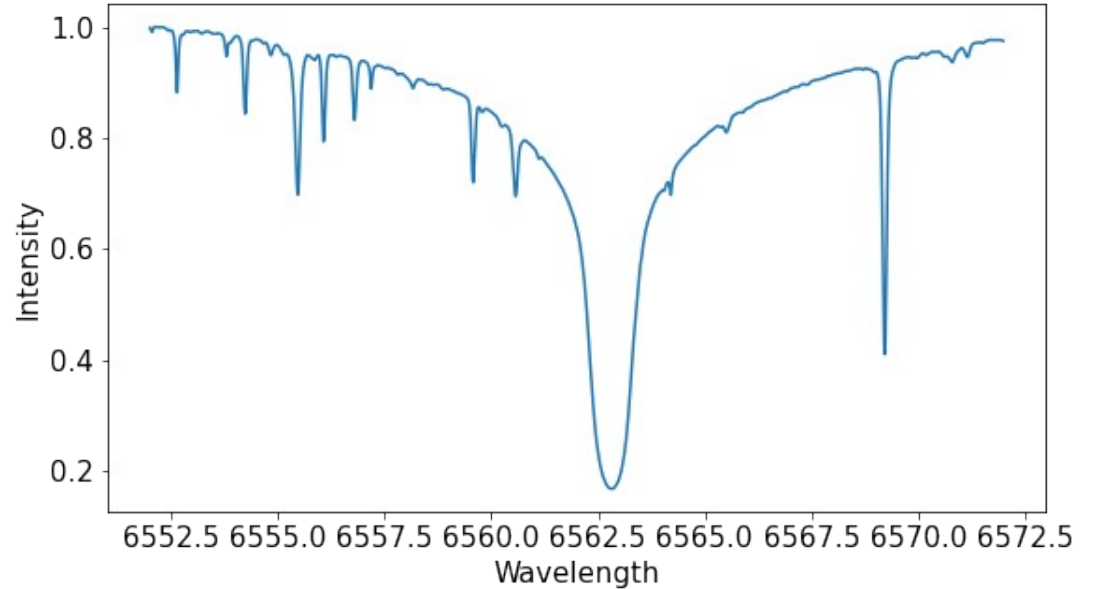
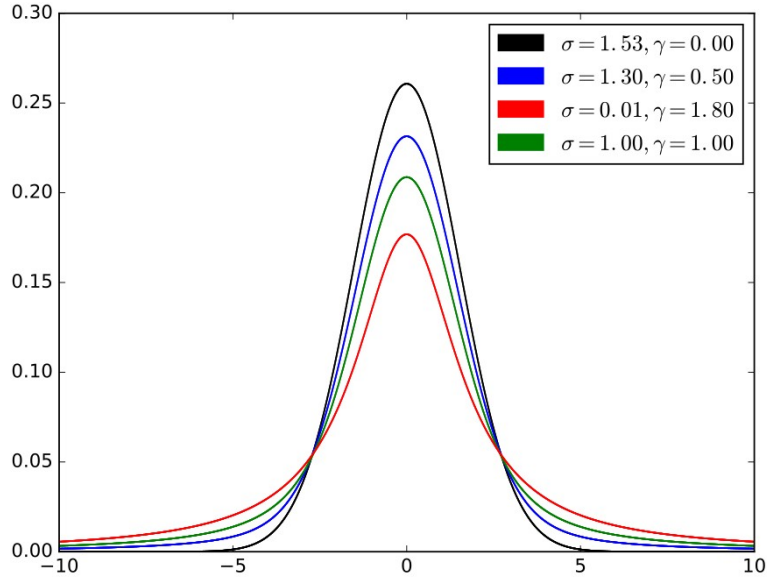
$$\phi_\lambda = \int_{-\infty}^{\infty} L\left(\lambda_0\left(1 - \frac{v}{c} - \frac{v_z}{c}\right)\right) p(v) dv$$

Final line profile, Voigt

Velocity distribution - Gaussian

# So finally

- Voigt profile enhances the line “wings”, and it is very visible in some observed spectral lines:



# Where do physical quantities come in:

- **Temperature:** ionization, excitation, thermal broadening, amount of collisions
- **Pressure:** constrains total particle density, ionization, damping
- **Large Scale velocity:** shifts the profile as a whole
- **Turbulent velocity (TBC) :** additionally broadens the line in a quasi-thermal fashion
- **Magnetic field:** a lot of things, wait for it!

# The end (for now)

- We know now (in principle), how to solve for the ionization and excitation state of the gas
- We know how to find the wavelength dependence of the line absorption (and emission)
- We know how to put them together to solve the radiative transfer equation.
- We will do these things next week!