

Saha-Boltzmann calibration of the Harvard sequence

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Abstract. We discovered . . .

1. Introduction

Stellar spectra gives a lot of information about the conditions in different stars. Absorption lines in the spectra appears if the atoms in the outer layers of the stars absorbs photons emitted from the star. Therefore the absorption lines can tell something about the content of elements in the star. A classification method called the Harvard spectral sequence is based on this information from the stellar spectra. Figure 1 shows some stellar spectra illustrating the Harvard spectral sequence for wavelengths between around 3800-5000 Å. The spectra can be compared to energy level diagrams for different elements to recognize which elements that have made the different absorption lines.

The energy level diagram for hydrogen is shown in figure 2. According to this diagram, the Balmer β line should absorb wavelengths of 4861 Å, and is the only line with a wavelength within the same range as the spectra in figure 1. At this wavelength in the spectra there is clearly, strong black line as expected for the five uppermost stars. The stars further down has no line at all for the Balmer β line, or just a very weak one.

This article will study the Harvard spectral sequence and spectral lines. It will discuss the relationship between the population density of particles with different ionization level and the strength of the absorption lines in stellar spectra. The report will mainly focus on a fictional element called *Schadeenium*, but will also look at real elements.

1.1. The Boltzmann and Saha laws

The Boltzmann and Saha laws describe the population density of different ionization stages and over the discrete energy levels within each stage for a specific element. Saha's law describes how many particles of an element that occupies the different ionization stages (e.g. twice-ionized). Saha distribution is given by

$$\frac{N_{r+1}}{N_r} = \frac{1}{N_e} \frac{2U_{r+1}}{U_r} \left(\frac{2\pi m_e kT}{h^2} \right)^{3/2} e^{-\chi_r/kT}, \quad (1)$$

where T is the temperature, k is the Boltzmann constant, h is the Planck constant, N_e is the electron density, m_e is the electron mass, χ_r is the threshold energy needed to ionize a particle from stage r to $r+1$, and U_{r+1} and U_r are the partition functions of the the respective ionization stages. $N_r = \sum_s n_{r,s}$ is the total particle density in all levels of ionization stage r . At last, the partition function U_r is defined by

$$U_r \equiv \sum_s g_{r,s} e^{-\chi_{r,s}/kT} \quad (2)$$

The Boltzmann law describes the distribution of the different ionized particles over the discrete energy levels within each ionization (e.g. a neutral atom with one electron excited to the second level). The Boltzmann distribution is given by

$$\frac{n_{r,s}}{N_r} = \frac{g_{r,s}}{U_r} e^{-\chi_{r,s}/kT}. \quad (3)$$

Here r and s represent the level s of the ionization stage r , n is the number density in unit m^{-3} , g is the statistical weight and χ is the excitation energy for the given level measured from the ground state ($r, 1$).

1.2. Schadeenium

To analyze how the absorption lines are connected to the population density of particles with different ionization stages and excitation levels, a simple fictional element is studied. The element we call Schadeenium and it will be referred to as E. Its ionization energies are $\chi_1 = 7$ eV for neutral stage, $\chi_2 = 16$ eV for the first ionized stage, $\chi_3 = 31$ eV for the second ionized stage and $\chi_4 = 51$ eV for the this ionized stage. The excitation energies is given by $\chi_{r,s} = s - 1$ eV. This means that there is 7 excitation levels in the neutral stage, 16 in the first ionized stage etc. For Schadeenium it is also assumed that the statistical weights is $g_{r,s} = 1$ for all levels (r, s).

2. Observations

Before analyzing the fictional element Schadeenium, an assumption needs to be made. From observations, it seems like the strength of the absorption lines scales linearly with the population density of the lower level of the corresponding transition. If the stellar outer layers have a lot of atoms with the lower level of the corresponding transition, it is natural to conclude that there would be a lot of atoms which could excite and absorb photons. This is proven to not be correct, although absorption lines do get stronger at larger lower-level population. In this report we will anyways assume that the scaling is linear. With this assumption, we can find initial rough estimates of the strength ratios of different absorption lines.

Take a neutral hydrogen atom as an example. The ratio between the strengths for the different excitation lines for hydrogen can be found using equation 3. The statistical weight for a neutral hydrogen is given by $g = 2s^2$. If we then assume we have a temperature equal to the the Sun's surface temperature ($T = 5777$ K), we get a ration between the Lyman α and Balmer α line of

$$\frac{n_{1,1}}{n_{1,2}} = \frac{2e^0}{8e^{-10.20/kT}} \approx 2 \cdot 10^8.$$

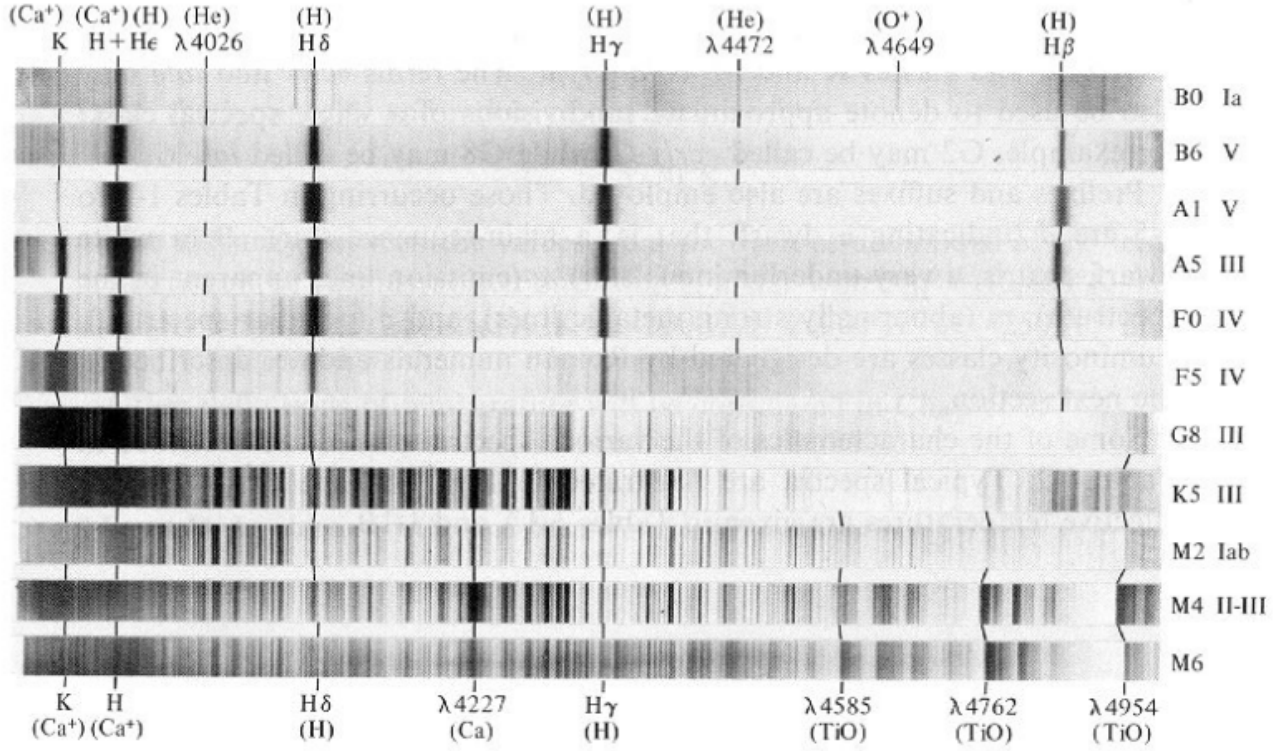


Fig. 1: Stellar spectra illustrating the Harvard spectral sequence. The dark lines represents the absorption lines for different wavelengths. The Harvard classification of the different stars, are written on the right. The stars are classified as either O, B, A, F, G, K or M, where the hottest stars are classified as O and the coolest as M. [Novotny \(1973\)](#)

The same calculation can be done for the Balmer α and Paschen α line strength ratio,

$$\frac{n_{1,2}}{n_{1,3}} = \frac{8e^{-10.20/kT}}{18e^{-12.09/kT}} \approx 20.$$

And for the Paschen α and Brackett α lines,

$$\frac{n_{1,2}}{n_{1,3}} = \frac{18e^{-12.09/kT}}{32e^{-12.75/kT}} \approx 2.$$

So on a sun-like star, the Lyman α line would be much stronger than the other lines.

3. Method

4. Results

4.1. Schadeenium

Table ?? shows the calculations of the partition function (equation 2) for Schadeenium for different ionization stages and temperature. It shows that the partition function is almost invariant to temperature and is almost constant for different ionization stages. It is also worth noting that the partition function re of order unity.

Figure 3 shows the Boltzmann distribution for the seven excitation levels in the ground state or neutral stage of Schadeenium. The figure shows that the lowest excitation level ($s=1$) is the most populated level at all temperatures. The energy for this

U_r	5000 K	10 000 K	20 000 K
U_1	1.11	1.46	2.23
U_2	1.11	1.46	2.27
U_3	1.11	1.46	2.27
U_4	1.11	1.46	2.27

level is $\chi_{1,1} = 0$ eV, so by looking at equation ??, we can see that for this level the Boltzmann distribution becomes temperature independent. This means that for this level the Boltzmann distribution is only dependent on the partition function, which also is almost invariant of temperature.

Figure 4 shows the Saha distribution for Schadeenium for the neutral or ground state and the three first ionization stages. From the distribution function we can see that there is at all temperatures maximum two stages of ionization. This is very different compared to the Boltzmann function, where there all the different energy states can be represented at the same temperature. The reason for their different dependence of temperature can be seen by looking at the equations for the Saha and Boltzmann distribution (equation 4 and equation 3. Here we can see that the Saha distribution is dependent on the temperature T to a factor of $3/2$ in addition to the temperature in the exponent, which they both have in common.

Another thing worth noting, is that the Saha distribution also depends on the electron pressure. In fact, with a lower pressure of electrons, the ionization happens at lower temperatures. If

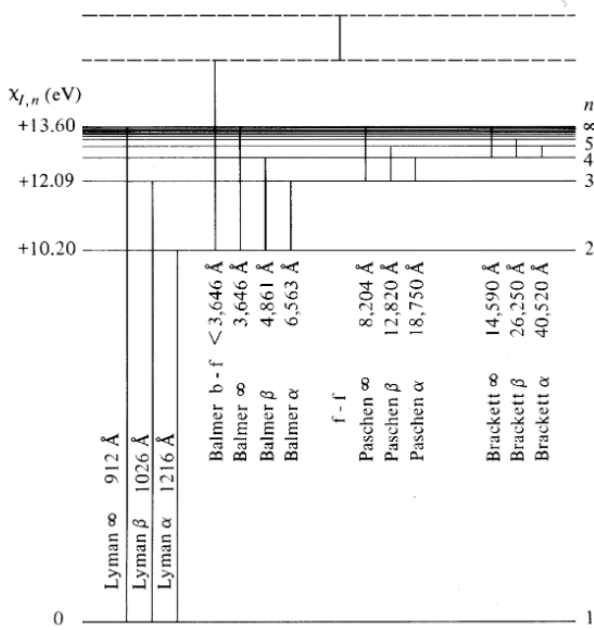


Fig. 2: Energy levels for hydrogen, each horizontal line represents one energy level. The excitation energies $\chi_{1,n}$ from the ground state to the respective energy levels are written in eV on the left. n denotes the energy level. The figure also shows the wavelengths the energy levels represent when light is emitted or absorbed from the atom. [Novotny \(1973\)](#)

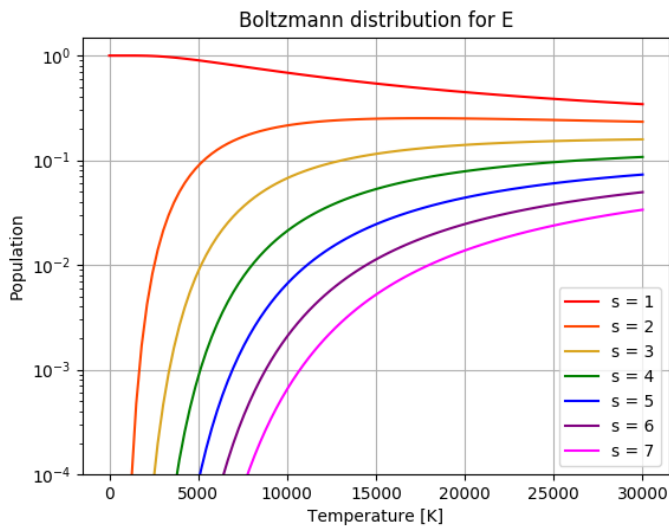


Fig. 3: Boltzmann's distribution for the seven excitation levels in the neutral/ground stage of Schadeenium.

we have more electrons in the gas, it would be harder for an atom to let go of another electron. But with a lower density of electrons, it is easier for the atom to "push away" a electron. A plot comparing the difference between Saha distributions for two different electron pressures, can be found in appendix A.

By making a plot of the Saha distribution multiplied with the Boltzmann distribution, we can obtain even more information about how the ionization and excitation change with tempera-

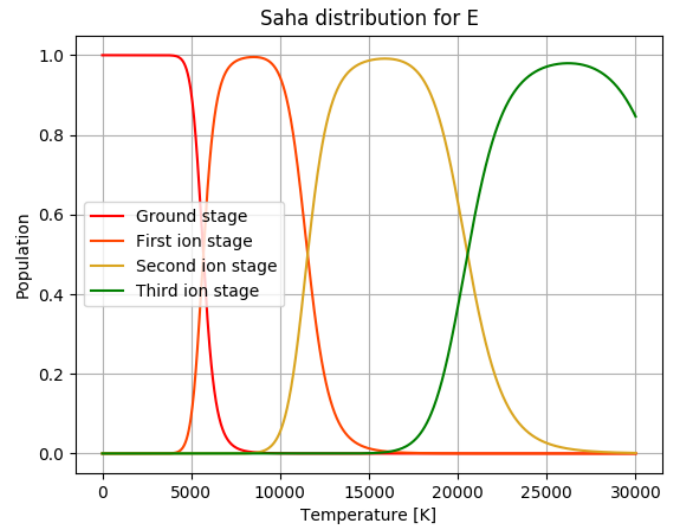


Fig. 4:

ture. This is done in figure 5. Here the Boltzmann distribution for the first level of excitation multiplied with different ionization stages is plotted. It is clearly that the ground stage dominates here, and even though the temperature increases, the population of the higher states is quite low, because we look at atoms with electrons in the first level of excitation. The steep flanks on each side of each peak is a result of the Saha distribution's shape.

In appendix B it is attached two more plots of the Saha-Boltzmann distributions of Schadeenium. These two are plotted for the the second level of excitation and the fourth level of excitation. For second level excitation, the population of higher ionized atoms decrease because the energy left to fully ionize the atom is now lower. For the forth level excitation, the electrons are excited in a high enough level to make more or less all the atoms ionized for temperatures over 5000 K.

For elements with higher ionization energies than E, there would be an even higher population of atoms in the ground stage, because it would require more energy to ionize the atom. For elements with lower ionization energies than E, the higher ionized stages would be more populated, and the ground stage would be less populated.

What causes the steep flanks on the left and the right side of each peak? What happens when $T \rightarrow \infty$ and $T \rightarrow 0$?

In Saha you can see that it is fully possible to ionize the atom, but in Boltzmann this probability is low (the ground state is always the most populated at all temperatures), but we know the atoms can be ionized. The partition function don't matter too much, can ignore it. exponent: more or less the same for high ionization energies, can ignore it. Given temperature $T = \text{constant}$. Electron density is the only parameter that can change. This is the difference between the two. Low N_e gives a higher likelihood to get to next ionization stage. If the electron density is low it is easy to give away electrons. S larger degree f ionization at lower electron pressure.

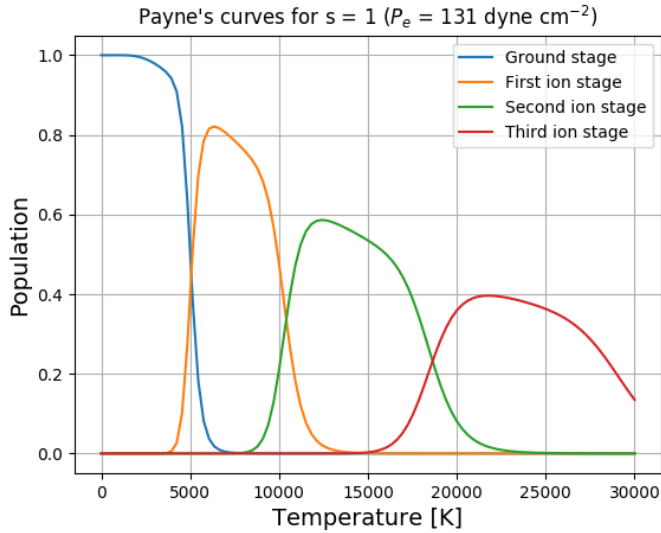


Fig. 5:

4.2. Hydrogen and calcium

Write something about 2.7? Further we will look at two more realistic elements, hydrogen and calcium. If the assumption we made in section 2 holds, we will find the strength ratio between the Ca^+K and $\text{H}\alpha$ Balmer absorption lines by plotting the ratio between the respective Saha-Boltzmann distributions. We would expect the Ca^+K line to be stronger at temperatures around 4000-6000 K because the Ca^+K line is in the ground level of excitation, but the $\text{H}\alpha$ Balmer line comes from the second to third transition of excitation. The Boltzmann distributions of Ca^+K and $\text{H}\alpha$ will behave similar to the Boltzmann distribution of E, even though this is a simplified atom, and therefore we would expect the ground level to be much more populated than the other levels as seen in figure 3.

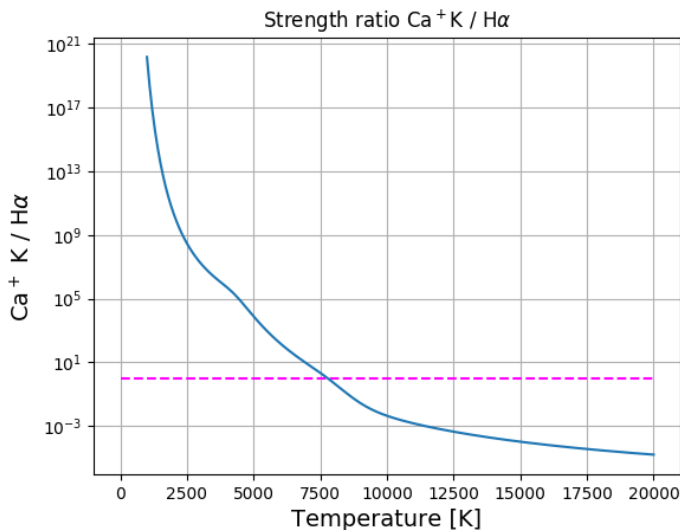


Fig. 6:

The ratio between the strength of the lines is plotted in figure 6. The figure shows that for temperatures lower than around

8000 K, the Ca^+K line should be stronger than the $\text{H}\alpha$ line as we assumed. Around 8000 K, the ratio is equal to 1, which means that the lines should be approximately at the same strength. At temperatures higher than 8000 K, the $\text{H}\alpha$ line is the strongest.

The Ca^+K line and the $\text{H}\alpha$ line, also differ much in their temperature sensitivity. The temperature sensitivity is shown in figure 7 together with the Saha-Boltzmann population of the two. At a bit over 6000 K, the Ca^+K curve drops to very small values. And a bit under 10 000 K the same happens for $\text{H}\alpha$ line. By comparing the temperature sensitivity to the population curves above in the figure, we can see that the populations have peaks at the same temperatures as where the temperature sensitivity drops. When the population reaches the maximum population, the temperature sensitivity is almost equal to zero for both of the elements. And when the population increases and decreases, the temperature sensitivity is bigger. So we can see that on the left side of the dips the population increases ($\Delta n > 0$), and on the right side of the dips the population decreases ($\Delta n < 0$).

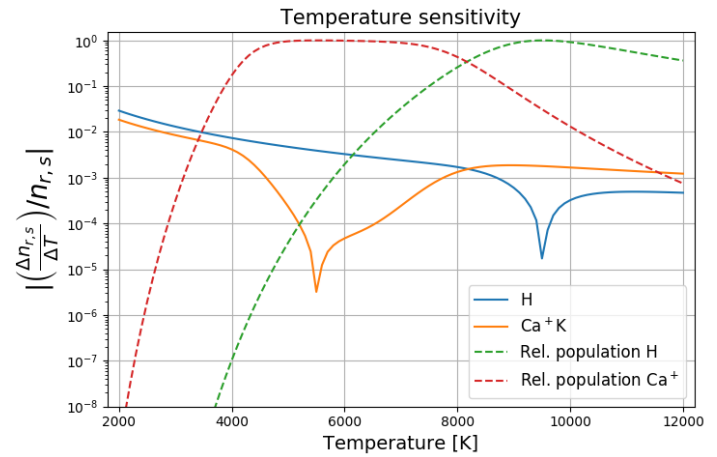


Fig. 7:

The Saha-Boltzmann distribution for neutral Hydrogen is shown in figure 8. The figure shows the fraction of hydrogen that is not ionized at different temperatures. The Hydrogen is half ionized at around 9200 K. This transition divides the "hotter" from the "cooler" stars. In hot stars, most of the hydrogen is ionized, resulting in a lot of free electrons. In cool stars almost none of the hydrogen atoms is ionized.

5. Conclusions

The answer is 42.

References

Novotny, E. 1973, Introduction to stellar atmospheres and interiors [ADS](#)

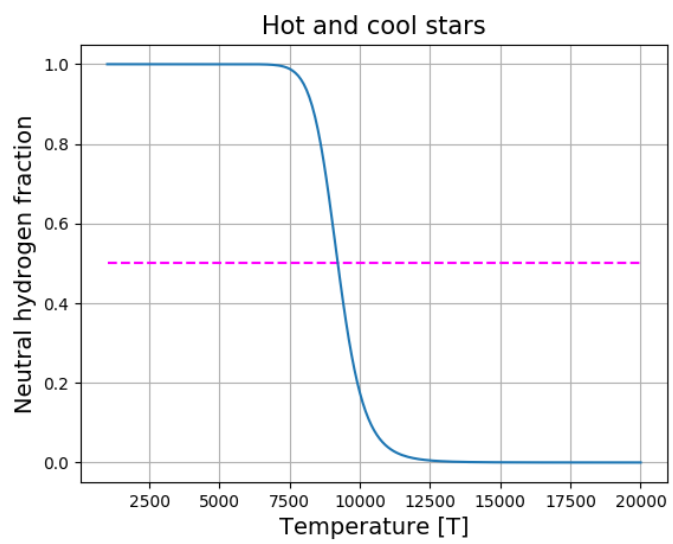


Fig. 8:

Fraunhofer line strengths and the curve of growth

1. Introduction

In the first part of this article, the Harvard classification was studied in terms of physics using the Saha and Boltzmann distributions for ionized and excited atoms. It was concluded that the variation in spectral line strength had a close connection to the change in temperature in stars. This part of the article will study how spectral lines form. This part will be less detailed and more figure oriented.

2. Theory

2.1. Planck's law

The radiation intensity emitted by a gas or body in thermodynamical equilibrium is given by

$$B_\lambda(T) = \frac{2hc^2}{\lambda^5} \frac{1}{e^{hc/\lambda kT} - 1}, \quad (1)$$

where h is the Planck constant, c is the speed of light, k is the Boltzmann constant, λ is the wavelength and T is the temperature.

In figure 1 Planck's function is plotted with respect to wavelength for different temperatures between 5000 and 8000 K. The figure shows that each temperature has a peak, and that these peaks shift to shorter wavelengths for higher temperatures as a result of *Wien's displacement law*. The highest temperature has the highest peak, so highest temperature emits the most intense radiation. The area under Planck's function for different temperatures increases a lot as the temperature increases. This means that the integrated Planck function is larger for higher temperatures.

We can also see that the function increases exponentially much faster for short wavelengths (at the left side of the figure) than at longer wavelengths. This is because when the $hc/\lambda kT$ -term is sufficiently large, that is when the wavelength is short, Planck's function can be approximated with the *Wien approximation* given by

$$B_\lambda \approx \frac{2hc^2}{\lambda^5} e^{-hc/\lambda kT}. \quad (2)$$

On the right side of the figure, the curves have a slope downwards. The same can be observed if we plot the Planck function with logarithmic y-axis or logarithmic x- and y-axis as shown in 2. This slope is caused by the longer wavelengths. When the wavelengths are longer $\exp(hc/\lambda kT) \approx hc/\lambda kT$, giving the *Rayleigh-Jeans approximation*,

$$B_\lambda \approx \frac{2ckT}{\lambda^4}. \quad (3)$$

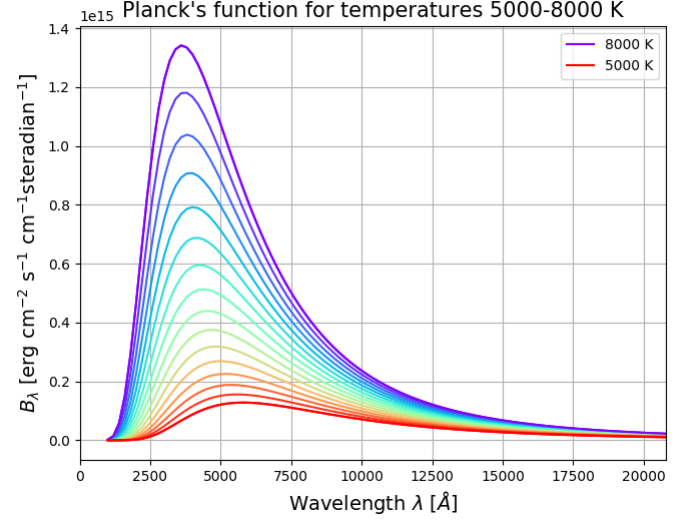


Fig. 1: Planck's function plotted with respect to wavelength for temperatures between 5000 and 8000 K. The red curve shows Planck's function for 5000 K, and the purple line shows Planck's function for 8000 K.

2.2. Radiation through an isothermal layer

When a beam of radiation with intensity $I_\lambda(0)$ passes through a layer of gas, it is attenuated. In addition the layer contributes to the beam with radiation originated within the layer. The total radiation emitting the layer is then given by

$$I_\lambda = I_\lambda(0)e^{-\tau} + \int_0^\tau B_\lambda[T(x)]e^{-(\tau-\tau(x))}d\tau(x), \quad (4)$$

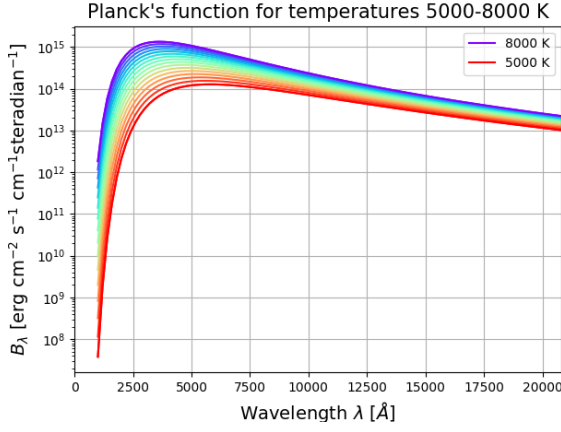
where τ is the optical thickness, B_λ is Planck's function and x is the location with the layer. For an isothermal layer the temperature T is constant, and therefore B_λ is also constant, giving

$$I_\lambda = I_\lambda(0)e^{-\tau} + B_\lambda \int_0^\tau e^{-(\tau-\tau(x))}d\tau(x)$$

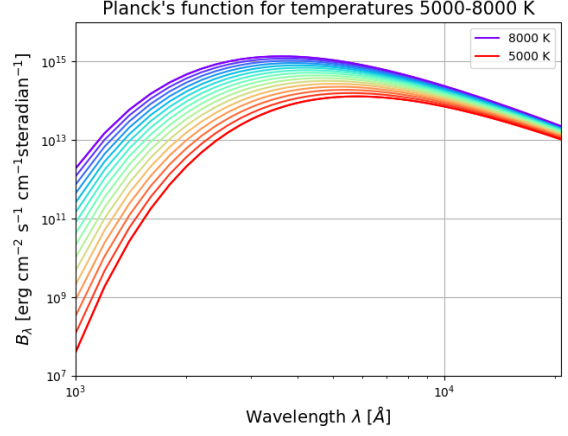
Substituting $u = -(\tau - \tau(x))$ and $du = d\tau$ gives

$$\begin{aligned} I_\lambda &= I_\lambda(0)e^{-\tau} + B_\lambda \int_0^\tau e^u du \\ &= I_\lambda(0)e^{-\tau} + B_\lambda \left[e^{-\tau+\tau(x)} \right]_0^\tau \\ &= I_\lambda(0)e^{-\tau} + B_\lambda(e^0 - e^{-\tau}) \\ &= I_\lambda(0)e^{-\tau} + B_\lambda(1 - e^{-\tau}). \end{aligned} \quad (5)$$

Figure 3 shows the emergent intensity through an isothermal layer where equation 5 holds and $B_\lambda = 2$. Figure 4 shows the same case with logarithmic x- and y-axis. Here we can see that when the optical thickness is low, the emergent intensity is almost the same as the radiation intensity before the beam passes the medium. Therefore such a layer is called optically



(a) Logarithmic y-axis.



(b) Logarithmic x- and y-axis.

Fig. 2: Planck's function plotted with respect to wavelength for temperatures between 5000 and 8000 K. The red curve shows Planck's function for 5000 K, and the purple line shows Planck's function for 8000 K. The left figure has logarithmic y-axis. The right plot has logarithmic x- and y-axis.

thin. For higher optical thickness, the emergent intensity approaches the radiation intensity emitted by the medium itself (I_λ). So the initial intensity decreases if $I_\lambda(0) > B_\lambda$. Therefore such a layer is called optically thick. Even when no intensity is sent through the medium, the radiation intensity out of the medium will increase against the Planck function.

For very large optical thickness, the emergent intensity becomes independent of τ . This is because if the medium is very optically thick, no radiation will emit through the medium. So the medium will only emit with radiation originated within itself.

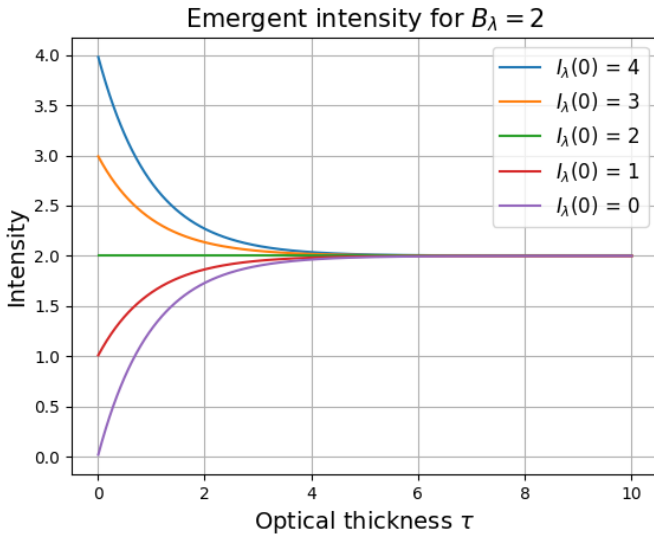


Fig. 3: The emergent intensity through an isothermal layer with $B_\lambda = 2$ plotted against the optical thickness τ .

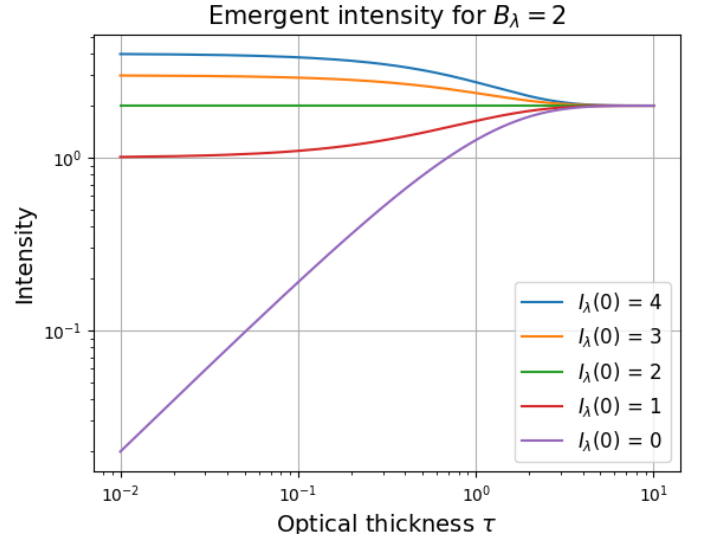


Fig. 4: The emergent intensity through an isothermal layer with $B_\lambda = 2$ plotted against the optical thickness τ with logarithmic x- and y-axis.

2.3. Spectral lines from a solar reversing layer

A Voigt function represents the convolution of a Gauss profile with a Lorentz profile. The function has a typical Gaussian shape close to the center of a spectral line. It is often approximated by taking the sum instead of the convolution of the two profiles

$$V(a, u) \approx \frac{1}{\Delta\lambda_D \sqrt{\pi}} \left[e^{-u^2} + \frac{a}{\sqrt{\pi}u^2} \right] \quad (6)$$

Figure 6 shows the Voigt profile with different damping parameters plotted against u . u measures the wavelength separation $\Delta\lambda = \lambda - \lambda(0)$ from the center of the spectral line at $\lambda = \lambda(0)$ in dimensionless units

$$u \equiv \Delta\lambda / \Delta\lambda_D \quad (7)$$

where $\Delta\lambda_D$ is the *Doppler width*. Stellar atmospheres typically have a damping parameter between $a \approx 0.01 - 0.5$. The figure shows that larger damping parameters gives a lower and more broad peak. Smaller damping factors on the other hand, gives higher, narrower peaks. Use eq 6 to explain why and something about fig 6.

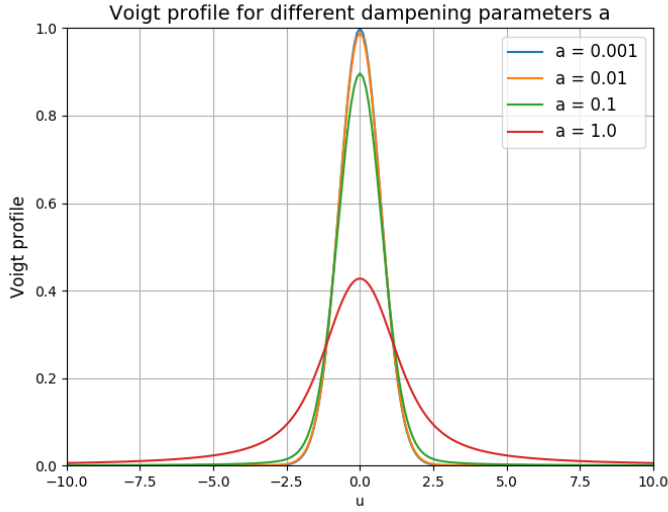


Fig. 5: The Voigt profile with different damping parameters plotted against u , where u is a measure on the wavelength separation from the center of the spectral line.

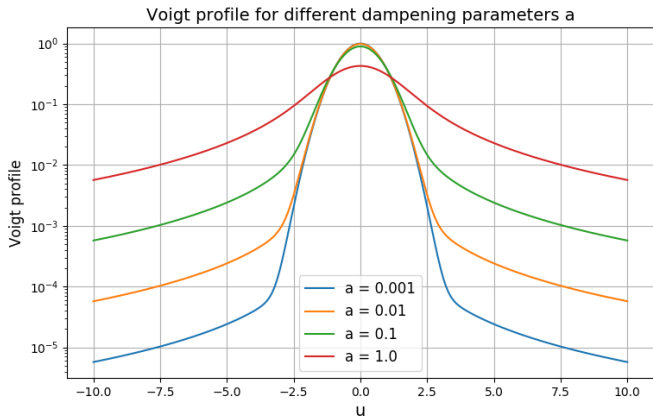


Fig. 6: The Voigt profile with different damping parameters plotted against u , where u is a measure on the wavelength separation from the center of the spectral line. Here it is used a logarithmic y-axis.

Write about what Schuster-Schwarzschild line profiles is. Figure 7 shows the Schuster-Schwarzschild line profile for a solar like photosphere.

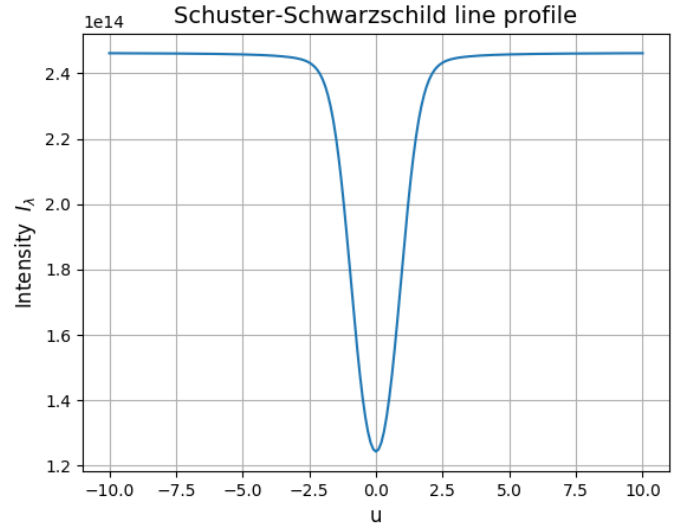


Fig. 7: The Schuster-Schwarzschild line profile for a solar like photosphere plotted with respect to u .

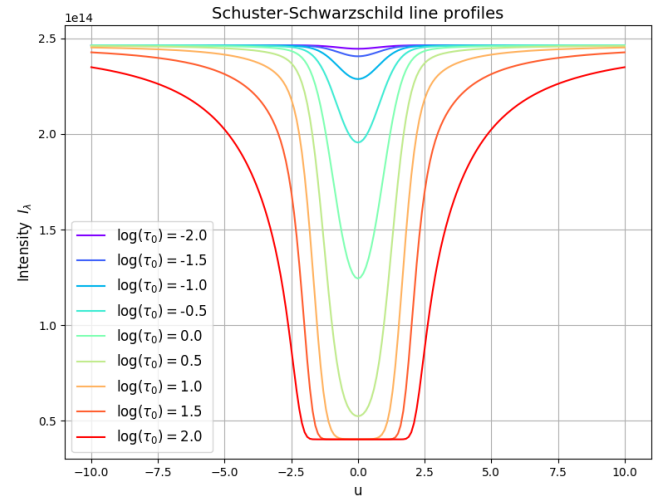


Fig. 8: The Schuster-Schwarzschild line profile for a solar like photosphere for different optical thickness τ plotted with respect to u .

Appendix A: Saha distribution for different electron pressures

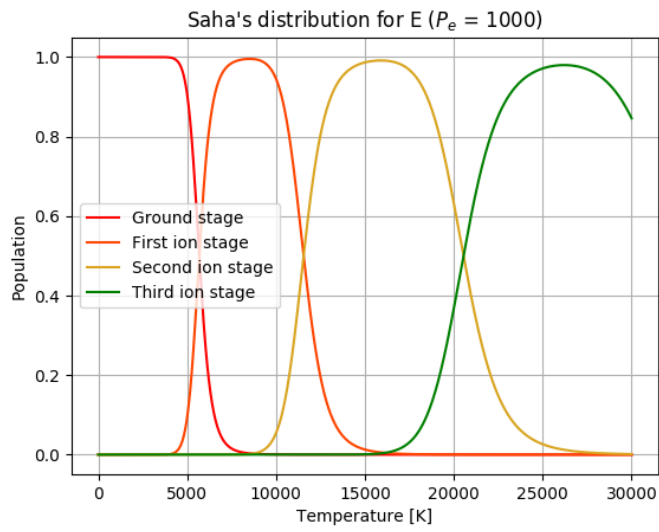


Fig. A.1:

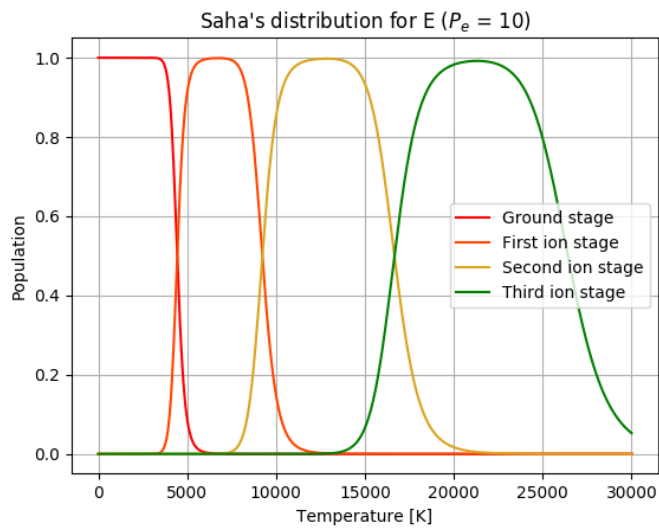


Fig. A.2:

Appendix B: Saha-Boltzmann distribution with excitation level $s=2$ and $s=4$

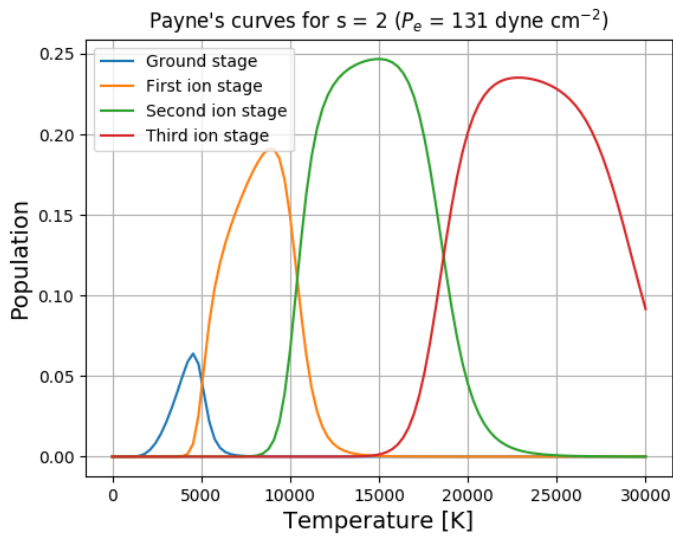


Fig. B.1:

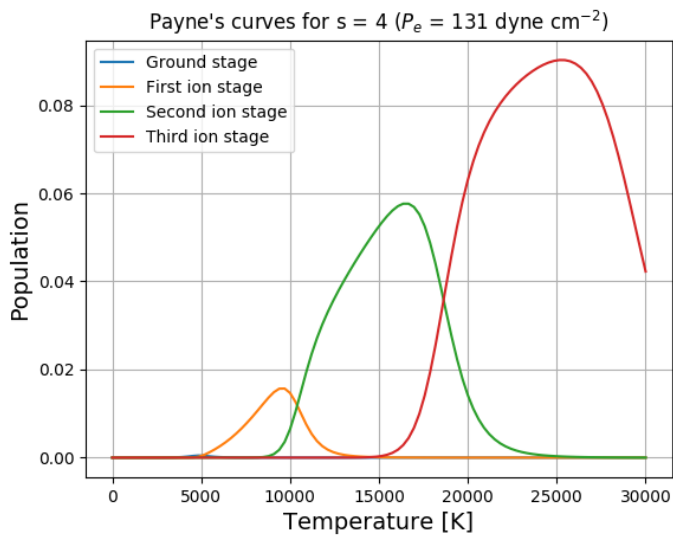


Fig. B.2: