Before we looked at radiative energy transport In the context of stars. The energy per volume (energy density) is $l(\hat{n}, \vec{x}, \nu, t) d^2 \hat{n} d\nu$. There were 4 processes that can change the energy density:

- · Transport (but nothing happens to photons)
 · Absorption (uncorrelated photon motion)

 - · Scattering (correlated pheton motion)
 - · Emissim (thermal and nuclear)

Scattered prietons do not travel in a fixed direction and transport does not affect photons. We will focus then on Scatternabsorption and emission, the phenomena that will affect what an observer looking in a straight line would observe. Remember that the fraction of absorvers is $H_{abs}(V, \vec{x}) p(\vec{x})$. we will call this $K(v,\vec{x})$. The rate of emission $j(v,\vec{x})p(\vec{x})$ will be J(U, x'). The time-independent radiation transport equation is then

 $\hat{n} \cdot \nabla L(v, \hat{n}, \vec{x}) = -K(v, \vec{x}) L(v, \hat{n}, \vec{x}) + J(v, \vec{x}) / 4\pi c$ If we take only a straight bline, with fixed direction n towards the observer, we can drop no Let $\vec{x} = s\hat{n}$, then $\frac{d}{ds}l(u,s) = -K(u,s)l(u,s) + J(u,s)/4\pi c$

$$\frac{d}{ds}l(\nu,s) = J(\nu,s)/4\pi c$$

$$\int_{s_1}^{s} dl(\nu,s) = \frac{1}{4\pi c} \int_{s_1}^{s} J(\nu,s) ds$$

$$l(\nu,s) = l(\nu,s_1) + \frac{1}{4\pi c} \int_{s_1}^{s} J(\nu,s) ds$$

SI is a convenient reference point. It could be the location of a light source, or the back of an interstellar cloud.

If the absorption is not negligible, define the optical deplant

$$\tau(\nu,s) = \int_{S_1}^{S} \kappa(\nu,s')ds' \qquad \underline{W3.1.4} \qquad \tau \approx 0 \text{ if } s \approx S_1$$

$$d\tau(v,s) = k(v,s^{\circ})ds^{\circ} \Rightarrow k(v,s) = \frac{d\tau(v,s)}{ds}$$

$$\frac{dl(v,s)}{ds} = -K(v,s)l(v,s) + J(v,s)/4\pi c$$

$$\frac{d}{ds} L(v,s) = -\frac{d\tau(v,s)}{ds} L(v,s) + J(v,s)/4\pi c$$

$$\frac{d}{ds}g(v,s) = d\tau(v,s) \left[-\frac{L(v,s)}{ds} + \frac{J(v,s)/4\pi c}{d\tau(v,s)} \right]$$

$$\frac{d}{dr} l(v,s) = ds \left[-\frac{l(v,s)}{ds} + \frac{J(v,s)/4\pi c}{d\tau(v,s)} \right]$$

$$\frac{d}{d\tau} l(\nu,s) = \lambda s \left[-\frac{l(\nu,s)}{\lambda s} + \frac{J(\nu,s)/4\pi c}{K(\nu,s) \lambda s} \right]$$

$$e^{-\tau(v,s)} \frac{dl(v,s)}{d\tau(v,s)} = -e^{-\tau(v,s)}l(v,s) + \frac{J(v,s)/4\pi c}{K(v,s)} e^{-\tau(v,s)}$$

But
$$\frac{d}{d\tau(v,s)} \left[e^{-\tau(v,s)} l(v,s) \right] = l(v,s) \frac{d}{d\tau(v,s)} e^{-\tau(v,s)} \frac{d}{d\tau(v,s)}$$

$$\frac{d}{d\tau(v,s)} \left[e^{-\tau(v,s)} l(v,s) \right] = l(v,s) \frac{d}{d\tau(v,s)} e^{-\tau(v,s)} \frac{d}{d\tau(v,s)}$$

$$= -e^{-\tau(v,s)} l(v,s) + e^{-\tau(v,s)} \frac{d}{d\tau(v,s)}$$

$$\Rightarrow e^{-\tau(v,s)} \frac{d}{d\tau(v,s)} = \frac{d}{d\tau(v,s)} \left[e^{-\tau(v,s)} \frac{1}{e^{-\tau(v,s)}} + e^{-\tau(v,s)} \frac{1}{e^{-\tau(v,s)}} \right]$$

$$\frac{d}{d\tau(v,s)} \left[e^{-\tau(v,s)} l(v,s) \right] = \frac{J(v,s)/4\pi c}{k(v,s)} e^{-\tau(v,s)}$$

$$\begin{cases} \int d \left[e^{-\tau(u,s)} \right] = \int_{s_1}^{s} \frac{J(u,s)/4\pi c}{k(u,s)} e^{-\tau(u,s)} d\tau(u,s) \end{cases}$$

$$= \int_{s_1}^{s} \frac{J(u,s)/4\pi c}{k(u,s)} e^{-\tau(u,s)} d\tau(u,s) \approx \int_{s_1}^{s_2} \frac{J(u,s)/4\pi c}{k(u,s)} e^{-\tau(u,s)} d\tau(u,s) \approx \int_{s_2}^{s_2} \frac{J(u,s)/4\pi c}{k(u,s)} e^{-\tau(u,s)} d\tau(u,s)$$

Assume that, even though J(U,S) and K(U,S) (127) depend explicitely on S, their ratio does not, So $\frac{J(\nu,s)/4\pi c}{K(\nu,s)} = \frac{J(\nu)}{4\pi c K(\nu)}$ can be taken out of the integral

$$L(v,s) = L(v,s_1) e^{-\tau(v,s)} + \frac{J(v)}{4\pi c K(v)} \int_{s_1}^{s} e^{-\tau(v,s)} d\tau(v,s')$$

Eventually, .

$$l(v,s) = e^{-\tau(v,s)}l(v,s) + \frac{J(v)}{4\pi c K(v)} \left[1 - e^{-\tau(v,s)}\right] w_{3.1.6}$$

In practice, $J(\nu)$ is an average of $J(\nu,s)$ K(ν)

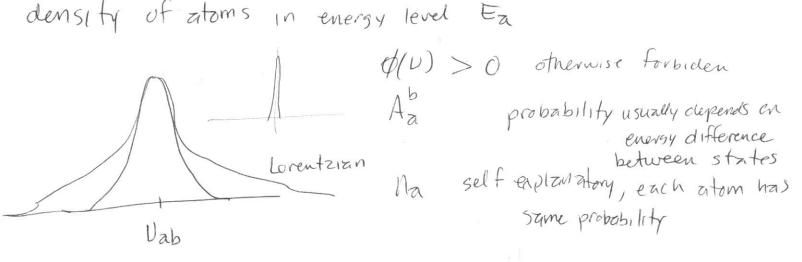
In 1917, Einstein defined a quantity Aa as the rate at which an atom spentaneously makes a transition from oversy level Ea to lower energy level Eb emitting a frequency $Vab = \frac{Ea-Eb}{h}$. The quantities are called Eißstein coefficients.

$$\Rightarrow \Delta E \ge \frac{t}{2\Delta t}$$

In order for the frequency to be perfectly sharp, $\Delta t \rightarrow \infty$, but photons are not produced continuosly, the transition occurs over some finite Δt and hence $\Delta E \neq 0$.

The atoms or molecules producing the radiation are moving in random directions, so they will have a small Dopler shift that also contributes to the linewidth of the emission.

If the probability that the phothen is emitted with frequency in the range [U, U+dV] is $\phi(U)dU$, sharply peaked at U=Vab with $\int \phi(U)dU=1$, the rate of emission per volume, per solid angle, and per frequency interval is $\phi(U)A_a^b n_a/4\pi$ where n_a is the number density of atoms in energy level E_a



It takes a photon a time $t = \frac{ds}{c}$ to travel [129] a distance ds, so the amount of additional

radiant energy per distance is $J(\nu)$ per solid angle pur frequency is $J(\nu) = \frac{1}{2} \frac{1}{4\pi c}$ per solid angle pur frequency is $J(\nu) = \frac{1}{4\pi c} \frac{1}{4\pi c}$, so

J(V)= hup(V) Aana

Applying Popler shift: $U = V_{ab} \left[1 - V/c \right]$ where $v = V_{ab} \left[1 - V/c \right]$ where $v = V_{ab} \left[1 - V/c \right]$

AWHAT IS THE DISTRIBUTION OF THE VELOCITIES WILL ISE? MAXWELL-BOLTZMAN

In I+D $p(v) = \left(\frac{m}{2\pi k_B T}\right)^{1/2} \exp\left(-m\left(v-\overline{v}\right)^2/2K_B T\right)$

So $\phi(\nu) = \frac{c}{\nu_{ab}} P(c(1-\nu/\nu_{ab}))$

it is the same for absorbemption, althoug that one includes stimulated emission, so

K(v) = hup(v) (Banb-Bana)/c

To work out the coefficients ne = gae - Ea/KBT

Nb gbe-Fb/KBT

Boltzman distribution