

Molecular Clouds

- radiative trapping
- optically thick mol. emission lines
- measuring molecular gas mass

Mass is dominated by -unobservable- H_2 , but we have CO as proxy. \rightarrow Can we use line strength to probe mass? Yes, ~~at~~ even if the molecular cloud is optically thick.

Easy

Molecular clouds are self-gravitating objects. CO has spectral lines with upper level $E_{up} \propto J^2$ and critical density $n_{crit} \propto J^3$. Thus higher J lines probe denser / hotter gas. Works for other simple linear molecules as well, but have higher energy levels.

CO(1-0) : the probe for coolish / low density gas.

$$\nu = 115 \text{ GHz}$$

$$\lambda = 2.6 \text{ mm}$$

$$A_{10} = 6.78 \cdot 10^{-8} \text{ s}^{-1}$$

Typical molecular cloud :

$$n_H \sim 10^3 \text{ cm}^{-3}$$

$$R \approx 10^9 \text{ cm} \quad (3 \text{ pc})$$

$$\frac{n_{CO}}{n_H} \approx 7 \cdot 10^{-5} \quad (\text{for } 25\% \text{ freely available CO, rest is locked up in dust grains})$$

$$\tau_\nu = \int \kappa_\nu dl = \kappa_\nu R$$

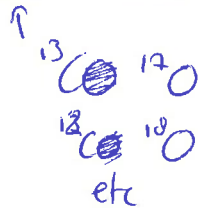
$$\hookrightarrow = n_0 \left(1 - \frac{n_1}{n_0} \frac{g_0}{g_1} \right) \frac{\lambda^2}{8\pi} \frac{g_1}{g_0} A_{10} \underbrace{\frac{1}{\sqrt{n}} \frac{\lambda}{\sqrt{2}\sigma}}_{\text{line profile}} e^{-\frac{v^2}{2\sigma^2}}$$

$$\text{Peak optical depth } \tau_0 = \kappa_0 R = n_0 R \underbrace{\left(1 - \frac{n_1}{n_0} \frac{g_0}{g_1} \right)}_{\text{assuming } T_{ex}} \frac{\lambda^3}{8\pi^{3/2} \sqrt{2}\sigma} \underbrace{\frac{g_1}{g_0}}_{\text{assuming } \sigma} A_{10}^{\text{line profile}}$$

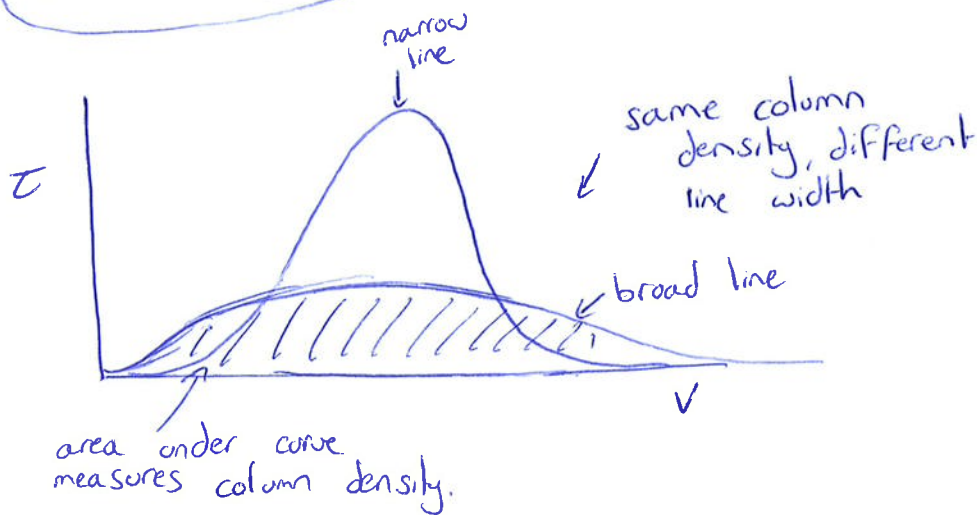
For cold molecular cloud, Tex not k.

$$\tau_0 = 15 \left(\frac{n_H}{1000 \text{ cm}^{-3}} \right) \left(\frac{R}{10^{19} \text{ cm}} \right) \left(\frac{n_{CO}/n_H}{7 \cdot 10^{-5}} \right) \left(\frac{1.4 \text{ km s}^{-1}}{\sigma} \right)$$

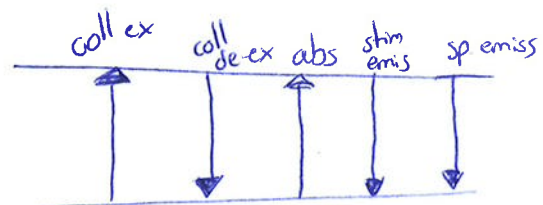
So in general we have optically thick lines. For rare molecules, we have the same problem. To get optically thin lines, we can only use optically ~~thin~~ rare isotopes. eg. ^{13}CO instead of ^{12}CO



$$\tau_0 \propto \frac{n(\text{CO})}{\sigma}$$



$$\frac{dn_U}{dt} = (n_C k_{LU} + \bar{n}_X \frac{g_U}{g_L} A_U) n_L - [n_C k_{UL} + (1 + \bar{n}_X) A_{UL}] n_U$$



$$= n_C k_{LU} n_L - n_C k_{UL} n_U - A_{UL} n_U + n_L \frac{g_U}{g_L} A_{UL} \bar{n}_X \left(1 - \frac{n_U g_L}{n_L g_U} \right)$$

In molecular cloud, n_X will change from place to place

↳ conditions at every point are coupled to conditions at every other point. That is not the case for optically thin emission.

We observe photons escaping from the cloud. So what we want to use is a photon escape probability.



Probability for photon to escape is

$$p(\vec{r}, \hat{n}) = e^{-\tau_\nu(\vec{r}, \hat{n})}$$

$$= \beta(\vec{r}, \hat{n})$$

Averaging over directions;

$$\bar{\beta}(\vec{r}) = \frac{1}{4\pi} \int \beta(\vec{r}, \hat{n}) d\Omega$$

$$\langle \bar{\beta}(\vec{r}) \rangle = \frac{\int \bar{\beta}_\nu(\vec{r}) \phi_\nu d\nu}{\int \phi_\nu d\nu} = \int \bar{\beta}_\nu(\vec{r}) \phi_\nu d\nu$$

So escape probability averaged over direction ~~at~~ and frequency.
Now

- 1) Assume excitation temperature is not function of temperature. (This will not be the case if mol. cloud is very inhomogeneous)
- 2) Use on-the-spot approximation \rightarrow if photon does not escape, it is absorbed immediately again.

Now we have a local problem, which makes things much easier

$$\begin{cases} I_\nu = I_\nu^{(0)} e^{-\tau_\nu} + B_\nu(T_{\text{ex}})(1 - e^{-\tau_\nu}) \\ n_\gamma = \frac{c^2}{2h\nu^3} I_\nu \end{cases} \quad \frac{n_u}{n_l} = \frac{g_u}{g_l} e^{-\frac{h\nu_{\text{ex}}}{kT_{\text{ex}}}}$$

$$\rightarrow n_\gamma(\nu) = n_\gamma^{(0)} e^{-\tau_\nu} + \frac{1 - e^{-\tau_\nu}}{\frac{n_l g_u}{n_u g_l} - 1}$$

We assume fixed β , which is good approximation except for edge of cloud, but there that has very small ~~occupation~~ volume filling / occupation.

$$n_\gamma(v) = n_\gamma^{(0)} \bar{\beta}_v + \frac{1 - \bar{\beta}_v}{\frac{n_L g_u}{n_u g_L} - 1}$$

Average
Average over line profile

$$\langle n_\gamma(v) \rangle = \langle \bar{\beta} \rangle n_\gamma^{(0)} + \frac{1 - \langle \bar{\beta} \rangle}{\frac{n_L g_u}{n_u g_L} - 1}$$

$$\frac{dn_u}{dt} = n_c k_{uv} n_L - n_c k_{ul} n_u - \langle \bar{\beta} \rangle A_{ul} n_u$$

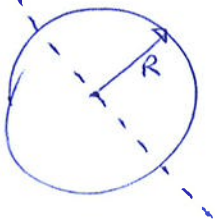
$$+ n_L \frac{g_u}{g_L} \langle \bar{\beta} \rangle A_{ul} n_\gamma^{(0)} \left(1 - \frac{n_u g_L}{n_L g_u} \right)$$

Because of on-the-spot approximation, we have no internally generated radiation field.

We have to calculate the escape probability $\langle \bar{\beta} \rangle$

The energy that would normally be in the radiation field ^{stays} is now inside the molecule.

Consider spherical cloud with radius R



$$\langle \bar{\beta} \rangle = \frac{1}{1 + 0.5 \tau_0}$$

← numerical fit !

Escape probability is quite small, at order of 0.04. An ~~an~~ numeric fit is also possible for uniformly expanding cloud

↳ $v \propto R$, e.g. Hubble flow

Every layer / radius in the cloud has its own velocity. Thus line is wide and peak optical depth is low.

$$\hookrightarrow \frac{dv}{dR} \sim \frac{\Delta v}{R}$$

Real molecular clouds are turbulent, therefore an ~~LGV~~ LGV (large velocity gradient) model is a much better approximation. (as above)

$$n_{\text{crit}} = \frac{A_{ul}}{k_{ul}} \rightarrow \langle \bar{\beta} \rangle \frac{A_{ul}}{k_{ul}} \text{ and thus lower.}$$

$$\text{CO}(1-0) : \tau \ll 1 \quad n_{\text{crit}} \sim 1100 \left(\frac{T}{10\text{K}} \right)^{-0.2} \text{ cm}^{-3}$$

Temperature dependent because collision-coefficient is T-dependent.

$$\tau \gg 1 \quad n_{\text{crit}} = \langle \bar{\beta} \rangle 1100 \left(\frac{T}{10\text{K}} \right)^{-0.2} \text{ cm}^{-3} \approx 50 \left(\frac{T}{10\text{K}} \right)^{-0.2} \text{ cm}^{-3}$$

Photon trapping

So we can do radiative transfer in optically thick lines