Q02-Lecture 9

So for, we have only paid attention to the atoms.

What about the light? How do we relate population levels to things like the index of refraction?

Recall that:

Fis the polarization induced in an atomic sample resulting from the imposition of an electric field Eocos(wt+0).

Assumptions:

1. Medium is isotropic (P is in the same direction as E)

Hopefully, you remember from E+M that this medium has a complex index of refraction given by

$$n^2 = 1 + \chi(\omega)$$

And if X(w) <<1, then

$$n^2 | + \frac{\chi'}{2} + i \frac{\chi''}{2}$$

and the wavevector is:

where
$$k' = (1 + \frac{\chi_{E}}{2}) \frac{\omega}{c}$$

$$k' = \frac{\chi'' \omega}{2}$$

Finally, the field evolves like:

$$\vec{E}(z,t) = \vec{E}_0 e^{i(kz-\omega t)} + c.c.$$

$$\left(\vec{E}(z,t) = \vec{E}_{o} e^{-\vec{k}z}\right)$$

So, k' gives us dispersion and k' gives us absorption.

So how do ne figure out X (and thus k' + k")?!

We calculate the dipole moment (B) of each individual atom, the multiply by the density of the medium to get

the maroscopic polarization!

Remember density matrix theory-

and remember that Di= <ilDli)=0

so we have!

$$\langle \hat{D} \rangle = \sum_{j \in \mathcal{D}} \rho_{jk} \mathcal{D}_{kj}$$

For a two level system, in have:

But me have up until non assumed <elôlg>=<glôle>=d,

Now, recall from homework that for:
- atoms initially in ground state
- in the rotating frame
- in steady state.

$$\langle G_2 \rangle^2 = \frac{\delta^2 + (\frac{1}{2})^2}{\delta^2 + \Omega n_2^2 + (\frac{1}{2})^2}$$

And also recall (from lecture 6):

Then we should be able to substitute the steady state values into the equation for (D), compare this to \vec{P} and find X!

But WE FORGOT! these stendy state values are in the rotating frame.

Recall that to go into the rotating frame, we made the transformation: |14) = Rz(wt) 14')

then solved for 14').

To get the lab frame picture bach, we must now use the transformation!

or rather

Given that R2(wt) = e

$$= \begin{pmatrix} e^{-i\omega t/2} & 0 \\ 0 & e^{i\omega t/2} \end{pmatrix}$$

and the previous expression for p', it is easy to see that:

$$P = R_2(ut) p' R_2(ut) = \frac{1}{2} \left(\frac{1 + 262}{e} - iwt \frac{1}{2} \left(\frac{1}{2} + 262 \right) - iwt \frac{1}{2} \left(\frac{1}{2} + 262 \right) \right)$$

$$= \frac{1}{2} \left(\frac{1}{2} + 262 \right) \frac{1}{2} \left(\frac{1}$$

Substituting this into the equation for IDD, we have:

$$\angle \hat{D} > = \frac{-i\omega t}{e} \frac{d}{d} (\langle r_{x} \rangle - i\langle r_{y} \rangle) + c.c.$$

 $= \frac{e}{e} \frac{d}{d} \frac{Q_{R}}{d} (\delta - i \frac{r_{z}}{2}) + c.c.$
 $\frac{2(\delta^{2} + D_{R}^{2}/2 + (\tilde{N})^{2})}{2(\delta^{2} + D_{R}^{2}/2 + (\tilde{N})^{2})}$

But renumber, to SR= -dEo 1 50

$$(3)^{2} \frac{d^{2} e E_{0}}{2 + (5^{2} + \frac{2n^{2}}{2} + (5^{2})^{2})} (-5 + i \frac{\pi}{2}) + c.c.$$

This is the steady state dipole moment for each individual atom. To get the polarization of the nedium, we must multiply by the density $\binom{N}{V}$.

$$P = (N)(\frac{d^{2}}{h})(\frac{-S+iN_{2}}{5^{2}+\frac{\Omega k^{2}}{2}+(N_{2})^{2}}) = e^{-i\omega t} + c.c.$$

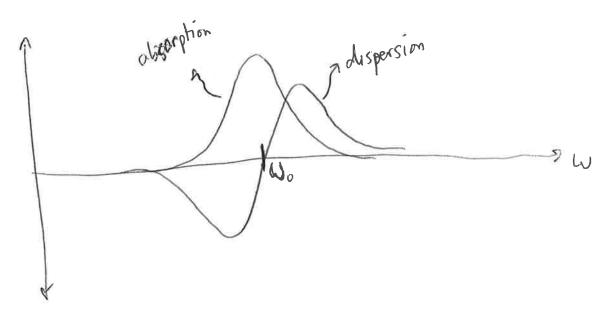
$$= (8) \chi = e^{-i\omega t} + c.c.$$

$$\delta \cdot \chi = \frac{N}{V} \frac{d^2}{8 t} \left(\frac{-5 + i \frac{7}{2}}{5^2 + \frac{10}{2} + (\frac{7}{2})^2} \right)$$

The real part (dispersion) is:

and the imaginary part (absorption) is:

$$\chi'' = \frac{N}{V} \frac{d^2}{5^2 + \frac{\Omega^2}{2} + (\frac{7}{2})^2}$$



Note 1: Linewidths The linewidths of these functions are \ \frac{12n_2}{2} + (\frac{n}{2})^2 The first turn (52) is called POWER BROADENING turning the loser up broadens the transition - The second term (1/2)2) is called the NATURAL LINEWIDTH. Lo syontanoous emission broaders the transition (the faster the level decays, the broader the transition). Note 2: Solution. If we define $S = \frac{97/2}{1}$ $5^2 + (\frac{7}{2})^2$ and $\chi_1 = \frac{N}{V} \frac{d^2}{\xi_0 \hbar} \left(\frac{-5 + i \frac{\pi}{2}}{5^2 + (\frac{\pi}{2})^2} \right) = \frac{1}{1 \text{ riear}}$ susceptibility" then $\left(\chi = \frac{\chi_1}{1+5}\right)$ - the susceptibility tends to O as Sir - so. - the transition is "saturated" Note 3: Other broadening mechanisms Homogeneons: Collisional Broadening

- Atoms collide with each other and can cause different effects. Non-radiative transitions can occur (mostly in solids) causing a modification of M (it: 1 -> M+ [coll). This bronders the curve (though in a somewhat trivial manner). - During collisions, we will fluctuate causing dephasing effects. This is a non-trivial effect: the Bloch equations are modified! d (0x) = 5(0x) -8(0x) dery = - 5(6x) - 2r6027 - 7(6y) d(6) = er(6) - 1 (1527+1) Note the P + 7's! with no dephosing, Y= 1/2. Otherwise 8= 1/2 + Tduph. The absorption is modified: $\chi'' = \frac{N}{V} \frac{d^2}{8t} \left(\frac{\delta^2}{\delta^2 + \delta^2 + \Omega_n^2(\frac{\gamma}{D})} \right)$ no dephasing dephasing

Inhomogeneous: Doppler Brondening

In a gas, the particles are moving at different speeds, with relocity distribution (in the x-direction):

$$f(v_x) = \frac{1}{\sqrt{12\pi}} e^{\frac{-\sqrt{2}}{2\sqrt{2}}}, \quad \overline{V} = \sqrt{\frac{2k_BT}{M}}$$

At rest, the atoms see a field:

$$E(r,t) = F_0 cos(wt-k-r)$$

But the atoms have trajectories like $r = r_0 + vt$ and therefore experience a field

$$E = E_0 \cos \left[(\omega - k \cdot v)t - k \cdot r_0 \right]$$

ie) the resonance condition becomes

or for $k = k_x \hat{x}$ and $|k| = \frac{\omega}{c}$,

$$\sqrt{\frac{1}{c}} = \frac{\omega - \omega_0}{\omega}$$

The absorption profile is modified. Each "relocity class" has a different resonance frequency and the total absorption is just the weighted owerage:

The profile is a mixture between the old LORENTZIAN lireshape, and a GAUSSIAN LINESHAPE introduced by the doppler effect. It is broader than each lineshape alone.

Colled the "VOIGT PROFILE"