Dispersion & Attenuation for EM Waves in Dissipative Media [Secs 7.547.7].

1) As we have noted for the Fresnel Formulas, the index of refraction, namely $n = \sqrt{\mu E} = \sqrt{E(\omega)} \int_{medium, \mu=1}^{nm-permeable} 1)$ is of central importance in describing how an EM wave propagates in a material. In actual materials, the atoms/molecules can resonate at many different frequencies under stimulation by EM waves; at these resonant frequencies, Where the atoms absorb the EM wave energy, the character of the wave propagation must change radically. We expect n=n(w) becomes a fon of frequency, and Should Show absorption features at specific frequencies.

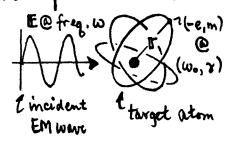
Also, the propagation velocity $V=c/n \rightarrow v(\omega)$, Jand So a "wavepacket" containing some spread of to to to frequencies W± AW will spread and, or <u>disperse</u>, as it propagates, as sketched. Finally, we have only briefly noted the effects of attenuation due to the medum, through its conductivity or. We now want to account in more detail

tory E + E(W) [ABSORPTION], U+V(W)[DISPERSION], O+O(W)[ATTENUATION]. (2)

The place to begin is to construct a <u>model</u> for the delectric const E=E(w). This will tell us how the medium is polarized by the EM were field E(x,t) during its passage; this polarization is the basic interaction between the wave E and the medium's electrons (-e,m). Simplest is a damped SHO model:

 $\rightarrow \mathbf{m} \left[\mathbf{r} + \gamma \mathbf{r} + \omega_0^2 \mathbf{r} \right] = -e \mathbb{E}(\mathbf{x}, \mathbf{t}).$

Here the electron's coupling to E is characterized by the two costs Wo & Y ... Wo is it's bound orbital frequency in its parent atom, and I is a damping onst represent-



ing an inhibition of the electron motion due to interaction with neighboring e's.

Now if $E = E_0 e^{-i\omega t}$ is minochromatic at freq. ω , then $E = F_0 e^{-i\omega t}$ (after transients) and Eq. (3) yields - upon division by $e^{-i\omega t}$...

$$\rightarrow mr_{o}\left[-\omega_{o}^{2}+i\gamma\omega+\omega_{o}^{2}\right]=-eE_{o}\Rightarrow \frac{Dirole}{moment}\} p=-er_{o}=\frac{(e^{2}/m)E_{o}}{(\omega_{o}^{2}-\omega^{2})-i\gamma\omega},$$

i.e.,
$$p = \alpha E_0$$
, $W_0 = \frac{e^2}{m}/[(\omega_0^2 - \omega^2) - i\gamma \omega]$ [polarizability for one e 4] in orbit with (ω_0, γ) .

<u>Generalize</u> this alone e) result to a for Z electrons per atom and N atoms per unit volume by simple addition (assumes all e's act independently)...

$$\alpha \longrightarrow N \frac{e^2}{m} \frac{2}{3!} f_i / [(\omega_i^2 - \omega^2) - i \gamma_i \omega] \int (\omega_i, \gamma_i) \text{ characterize}$$
 (5)

The fig are called "oscillator strengths": they are numbers, whom I fig = Z, that specify the relative polarization contribution of the jt electron (e's that are strongly bound have a small fig, while weakly bound e's have a lunge fig.).

The dielectric const for a medium with the polarizability in Eq. (5) is then:

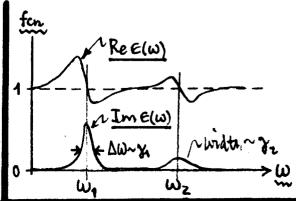
$$E(\omega) = 1 + 4\pi\alpha = 1 + \omega_p^2 \frac{2}{5i} \frac{g_i}{[(\omega_i^2 - \omega^2) - i\gamma_i\omega]} \int \frac{\text{Jackson}}{\text{Eq. (7.51)}}$$

$$\omega_p^2 = 4\pi N Z e^2 / m \begin{cases} \text{collidathe} \\ \text{"plasma freq."} \end{cases}, \quad g_i^2 = f_i / Z \begin{cases} g_i \leq 1 \text{ is a "fractional} \\ \text{oscillator strength"} \end{cases}$$

$$\underbrace{\text{Re } \varepsilon(\omega)} = 1 + \omega_{P}^{2} \underbrace{\sum_{j} g_{j} \left[\frac{\omega_{j}^{2} - \omega^{2}}{Q_{j}(\omega)} \right]}_{j}, \quad \underbrace{\text{Im } \varepsilon(\omega)}_{j} = \omega_{P}^{2} \underbrace{\sum_{j} g_{j} \left[\frac{\gamma_{j} \omega}{Q_{j}(\omega)} \right]}_{j};$$

$$\left(w_{i}^{2} \Delta_{i}(\omega) = (\omega_{i}^{2} - \omega^{2})^{2} + (\gamma_{i} \omega)^{2}. \right)$$

The frequency dependence of this model is sketched at right. Both ReE(W) & ImE(W) Show rapid variation with w near each of the atomic resonant frequencies Wj.



3) The functional form of E(W) connects to actual wave propagation as follows. Recall that the wave number k for propagation in the medium is...

$$\rightarrow k = \frac{\omega}{4r} = \frac{\omega}{c} \sqrt{\mu \epsilon} \dots \epsilon$$
, and so k, is now complex; set $\mu = 1$ (nonpermentle). (8)

• ... put:
$$k = \beta + \frac{1}{2}i\alpha$$
, and define: $k_0 = \frac{\omega}{c} \int free space$.

$$\frac{k^2 = (\beta^2 - \frac{\alpha^2}{4}) + i \alpha \beta}{k^2 = k_0^2 \in (\omega)} \int width : \beta = Rek, \qquad (9)$$

Eq. (9) can be solved for at & B separately, with the following results...

$$\begin{cases} \text{effective} \\ \text{wave #} \end{cases} \underbrace{\beta = k_0 / \epsilon_R U(\epsilon)}_{\text{to wave #}}, \text{ coefficient} \end{cases} \underbrace{\alpha = \frac{\beta}{[U(\epsilon)]^2} (\epsilon_I / \epsilon_R)}_{\text{to wave #}};$$

$$\begin{cases} \text{to wave #} \\ \text{to wave #} \end{cases} \epsilon_R = \text{Re } \epsilon(\omega), \epsilon_I = \text{Im } \epsilon(\omega), \text{ and } U(\epsilon) = \left[\frac{1}{2} \left(1 + \sqrt{1 + (\epsilon_I / \epsilon_R)^2}\right)\right]. \end{cases}$$

The parameter &= 2 Imk is clearly connected with attenuation of the wave, as

The physics of B= Rek is less apparent, but we can define a phase velocity:

$$\rightarrow V_{ph} = \omega/\beta = c/U(\epsilon)\sqrt{\epsilon_R} = v_{ph}(\omega).$$

The frequency dependence of the attenuation coefficient $\alpha(\omega)$ & wave phase velocity $\nu_{\mu}(\omega)$ is sketched at right. α & ν_{μ} roughly repeat the behavior of Ime & 1/Re & per the sketch last page -- they show rapid varia-

tron near the atomic resonance fregs. Wy [NOTE: U(E)=1 except near the resonances]. Near the resonances, the wave is both absorbed & dispersed.

CONCLUSION: When E-> Elwi, the propagation problem is seriously attend.

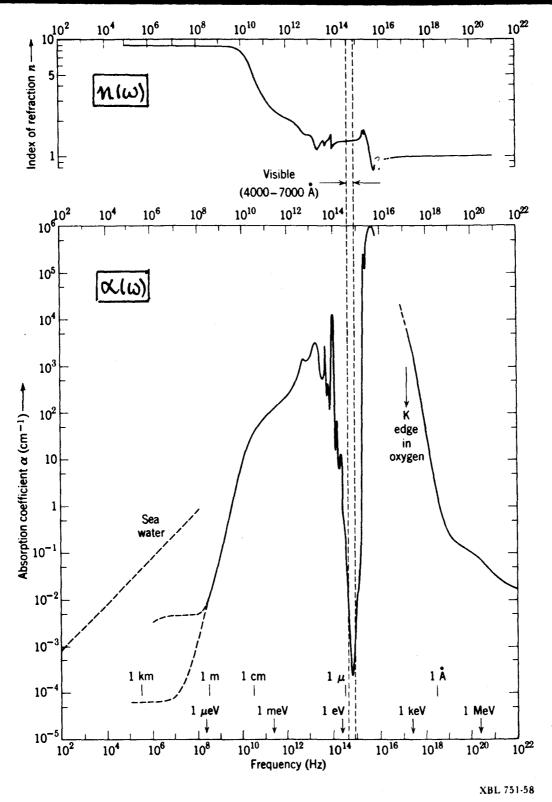


Fig. 7.9 The index of refraction (top) and absorption coefficient (bottom) for liquid water as a function of linear frequency. Also shown as abscissas are an energy scale (arrows) and a wavelength scale (vertical lines). The visible region of the frequency spectrum is indicated by the vertical dashed lines. The absorption coefficient for sea water is indicated by the dashed diagonal line at the left. Note that the scales are logarithmic in both directions.