

Chapter 1 – Math Toolbox

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Objectives

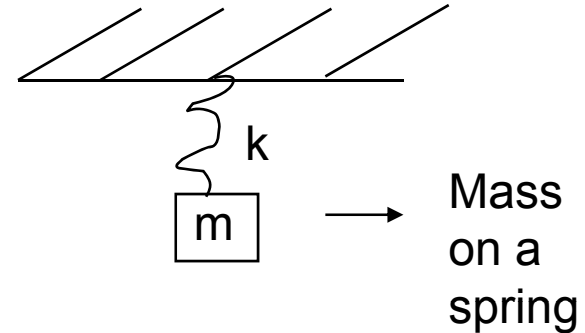
- Develop a set of tools useful throughout the course



1.1 Linear Systems

- Consider a simple system:
- Equation of motion:

$$m \frac{d^2 x}{dt^2} + \gamma m \frac{dx}{dt} + m \omega_o^2 x = f(t) \quad (1.1)$$



$$\omega_o = \sqrt{\frac{k}{m}}$$

- Define Operator: (linear differential eqs)

$$L = m \frac{d}{dt^2} + \gamma m \frac{d}{dt} + m \omega_o^2 \quad (1.2)$$

$$\rightarrow \boxed{L(x) = F(t)} \quad (1.3)$$



1.1 Linear Systems

- Operator L has important properties:

$$\begin{aligned} \text{a) } L(ax) &= m \frac{d^2(ax)}{dt^2} + \gamma m \frac{d(ax)}{dt} + m\omega_o^2(ax) = \\ &= a \left[m \frac{d^2(x)}{dt^2} + \gamma m \frac{d(x)}{dt} + m\omega_o^2(x) \right] = \\ &= aL(x) \end{aligned} \tag{1.4}$$

$$\begin{aligned} \text{b) } L(x + y) &= m \frac{d(x + y)}{dt} + \gamma m \frac{d(x + y)}{dt} + m\omega_o^2(x + y) = \\ &= L(x) + L(y) \end{aligned} \tag{1.5}$$



1.1 Linear Systems

- Definition: An operator obeying properties $L(ax) = aL(x)$ and $L(x+y)=L(x)+L(y)$ is called linear
- Most of the system in nature are linear; well, at least to the first approximation
- They are mathematically tractable \rightarrow analytic solutions
- Consider equations:

$$\begin{cases} L(x_1) = 0 \\ L(x_2) = 0 \end{cases} \quad (1.6)$$

$\rightarrow x_1, x_2$ are solutions



1.1 Linear Systems

- Continuing:

$$\begin{aligned}
 \rightarrow L(ax_1 + bx_2) &= L(ax_1) + L(bx_2) \\
 &= aL(x_1) + bL(x_2) \\
 &= 0 + 0 \qquad (1.7)
 \end{aligned}$$

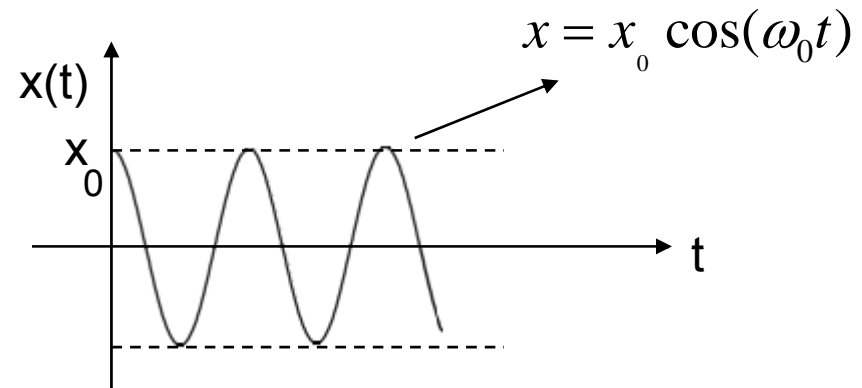
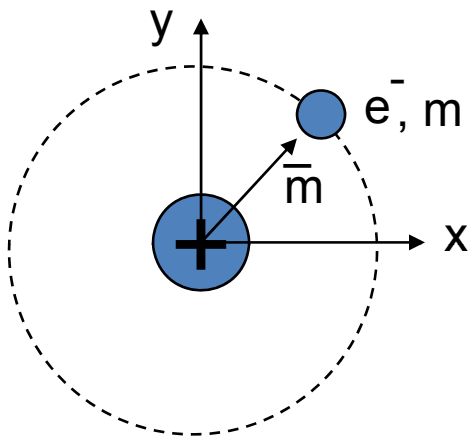
- Any linear combination of solutions: x_1, x_2 is also a solution
- The number of independent solutions = degrees of freedom
 X_1, X_2, \dots, X_N = independent solutions if

$$X_i \neq \sum_{j \neq i} \alpha_j x_j, \text{ for any } \alpha_j \qquad (1.8)$$
- Linear Differential eqs of order N allow for N independent solutions



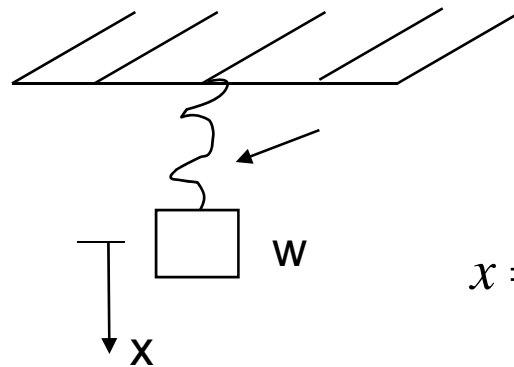
1.2 Light-matter interaction

- Classic model of atom: e^- rotating around $N \approx$ planets



→ Lorentz Model

- Analogy :



$$x = x_0 \cos(\omega_0 t) \quad \rightarrow \quad \omega_0 = \sqrt{\frac{k}{m}}$$



1.2 Light-matter interaction

- ■ So, motion of charge follows the same eq (1.1)

$$m \frac{d^2 x}{dt^2} + \gamma m \frac{dx}{dt} + m \omega_o^2 x = F(t)$$

- Incident field drives the charge: $\bar{F}(t) = q\bar{E}(t)$ (1.9)

- For e^- , $q = -e$!

- Monochromatic field: $E(t) = E_o e^{-i\omega t}$

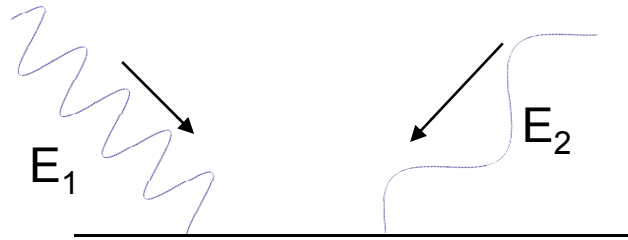
$$\rightarrow m\ddot{x} + \gamma m\dot{x} + m\omega_o^2 x = qE_o e^{-i\omega t} \quad (1.10)$$

- This is the eq of motion for electric charge under incident EM field. Can explain most of Optics!



1.3 Superposition principle

- Suppose we have 2 fields simultaneously interacting with the material (Eg. ω_1, ω_2):



$$\begin{aligned} E_1 &= |E_1| e^{-i\omega_1 t} ; qE_1 = F_1 \\ E_2 &= |E_2| e^{-i\omega_2 t} ; qE_2 = F_2 \end{aligned} \quad (1.11)$$

- Let x_1, x_2 be solutions of displacements for the two forces F_1 and F_2

$$\begin{cases} L(x_1) = F_1(t) \\ L(x_2) = F_2(t) \end{cases} \quad (1.12)$$

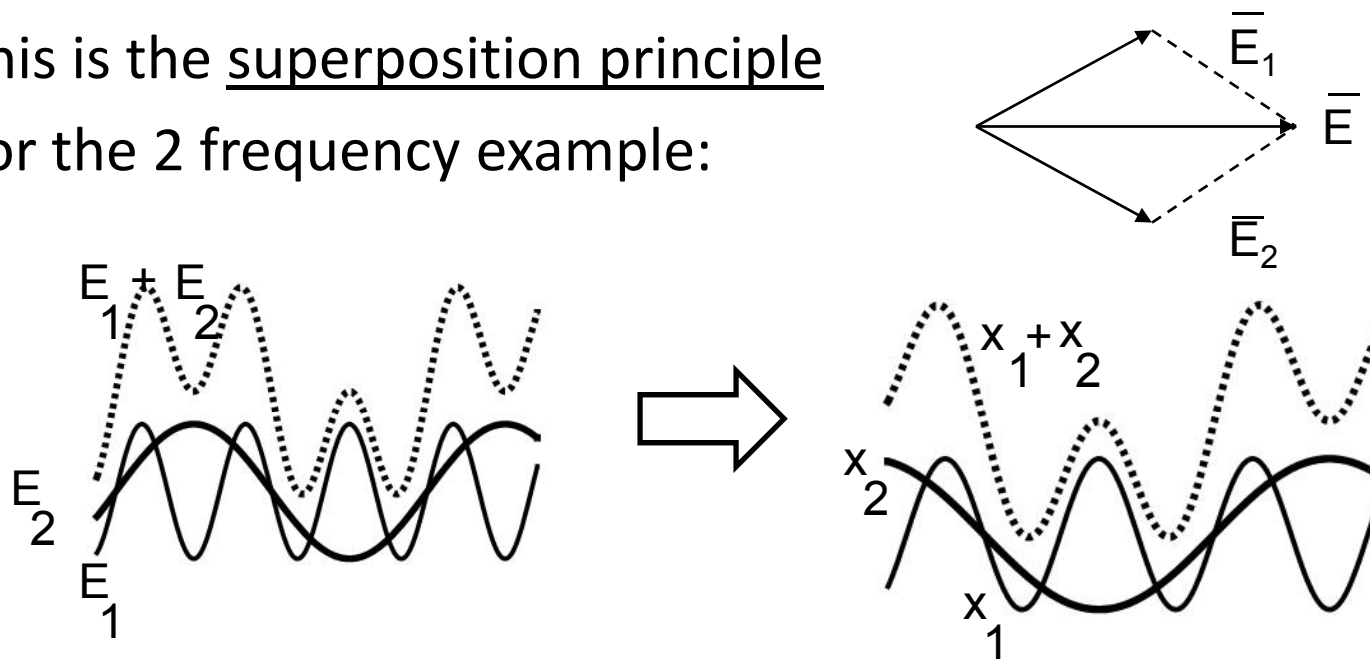


1.3 Superposition principle

- Consider the same solution:

$$\begin{aligned} L(x_1 + x_2) &= L(x_1) + L(x_2) \\ &= F_1(t) + F_2(t) \end{aligned} \quad (1.13)$$

- So, final solution is just the sum of individual solutions. Nice!
- This is the superposition principle
- For the 2 frequency example:

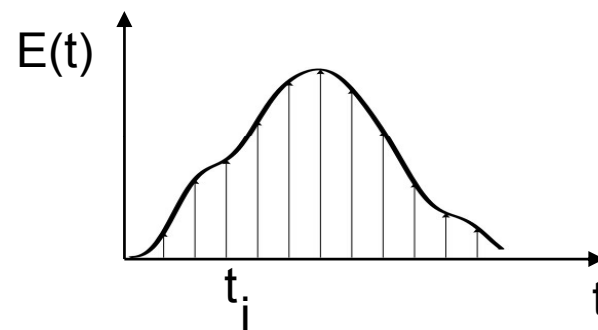


- It's as if one applies the fields one by one and sums their results



1.4 Green's function/impulse response

- Let the incident field, i.e driving field, have a complicated shape



→ arbitrary

- $E(t)$ can be broken down into a succession of short pulses, i.e Dirac delta functions:

$$\rightarrow \delta(t) = \begin{cases} 1, & t = 0 \\ 0, & \text{otherwise} \end{cases} \quad (1.14)$$

$$\rightarrow E(t) = \int_{-\infty}^{\infty} E(t') \delta(t - t') dt' \quad (1.15)$$



1.4 Green's function/impulse response

- If we know the response of the system to a short pulse, $\delta(t)$, the problem is solved
- Let $h(t)$ be the solution to $\delta(t)$
- The final solution for an arbitrary force $\bar{F}(t) = q\bar{E}(t)$ is:

$$x(t) = \int_{-\infty}^{\infty} E(t')h(t - t')dt' \quad (1.16)$$

- This is the Green's method of solving linear problems
- **$h(t)$ = Green's function or impulse response of the system**
- Complicated problems become easily tractable!



1.5 Fourier Transforms

- Very efficient tool for analyzing linear (and non-linear) processes

- Definition:
$$\mathfrak{F}[f(x)] = \int_{-\infty}^{\infty} f(x)e^{-i2\pi x f_x} dx \quad (1.17)$$

$$= F(f_x) = \tilde{f}(\xi)$$

- F is the Fourier transform of f
- $f : \Delta \rightarrow \Delta; \Delta \in \mathbb{C}$, f must satisfy:

$$(x, y, z) \xrightarrow{\mathfrak{F}} (\xi, \eta, \zeta)$$

- $\int |f| < \infty$ - modulus integrable
 - f has finite number of discontinuities in the finite domain Δ
 - f has no infinite discontinuities
- In practice, some of these conditions are sometimes relaxed



1.5 Fourier Transforms

- Inverse Fourier Transforms:

$$\begin{aligned}\mathfrak{F}^{-1}[\mathfrak{F}(f(x))] &= \int_{-\infty}^{\infty} \tilde{f}(\xi) e^{+i2\pi x f_x} df_x \\ &= f(x)\end{aligned}\tag{1.18}$$

$$\rightarrow \mathfrak{F}^{-1}[\mathfrak{F}(f)] = f\tag{1.19}$$

- Meaning of F.T: reconstruct a complicated signal by summing sinusoidals with proper weighting



1.5 Fourier Transforms

- Fourier transform is a linear operator:

$$\begin{aligned}\mathfrak{F}[af(x) + bg(x)] &= \\ &= \int_{-\infty}^{\infty} [af(x) + bg(x)]e^{-i2\pi x\xi} dx = \\ &= a \int_{-\infty}^{\infty} f(x)e^{-i2\pi x\xi} dx + b \int_{-\infty}^{\infty} g(x)e^{-i2\pi x\xi} dx \\ &= a\mathfrak{F}[f(x)] + b\mathfrak{F}[g(x)]\end{aligned}\tag{1.20}$$



1.6 Basic Theorems with Fourier Transforms

a) Shift Theorem: if $\tilde{f}(\xi) = \mathfrak{F}[f(x)]$

$$\mathfrak{F}\{f(x - a)\} = \tilde{f}(\xi)e^{-i2\pi\xi a} \quad (1.21)$$

- Easy to prove using definition
- Eq 1.21 suggest that a shift in one domain corresponds to a linear phase ramp in the other (Fourier) domain



1.6 Basic Theorems with Fourier Transforms

b) Parseval's theorem: if $\mathfrak{F}[f(x)] = \tilde{f}(\xi)$

$$\int_{-\infty}^{\infty} |f(x)|^2 dx = \int_{-\infty}^{\infty} |\tilde{f}(\xi)|^2 d\xi \quad (1.22)$$

- Conservation of total energy



1.6 Basic Theorems with Fourier Transforms

c) Similarity theorem: if

$\mathfrak{F}[f(x)] = \tilde{f}(f_x)$, i.e. \tilde{f} is the F.T of f

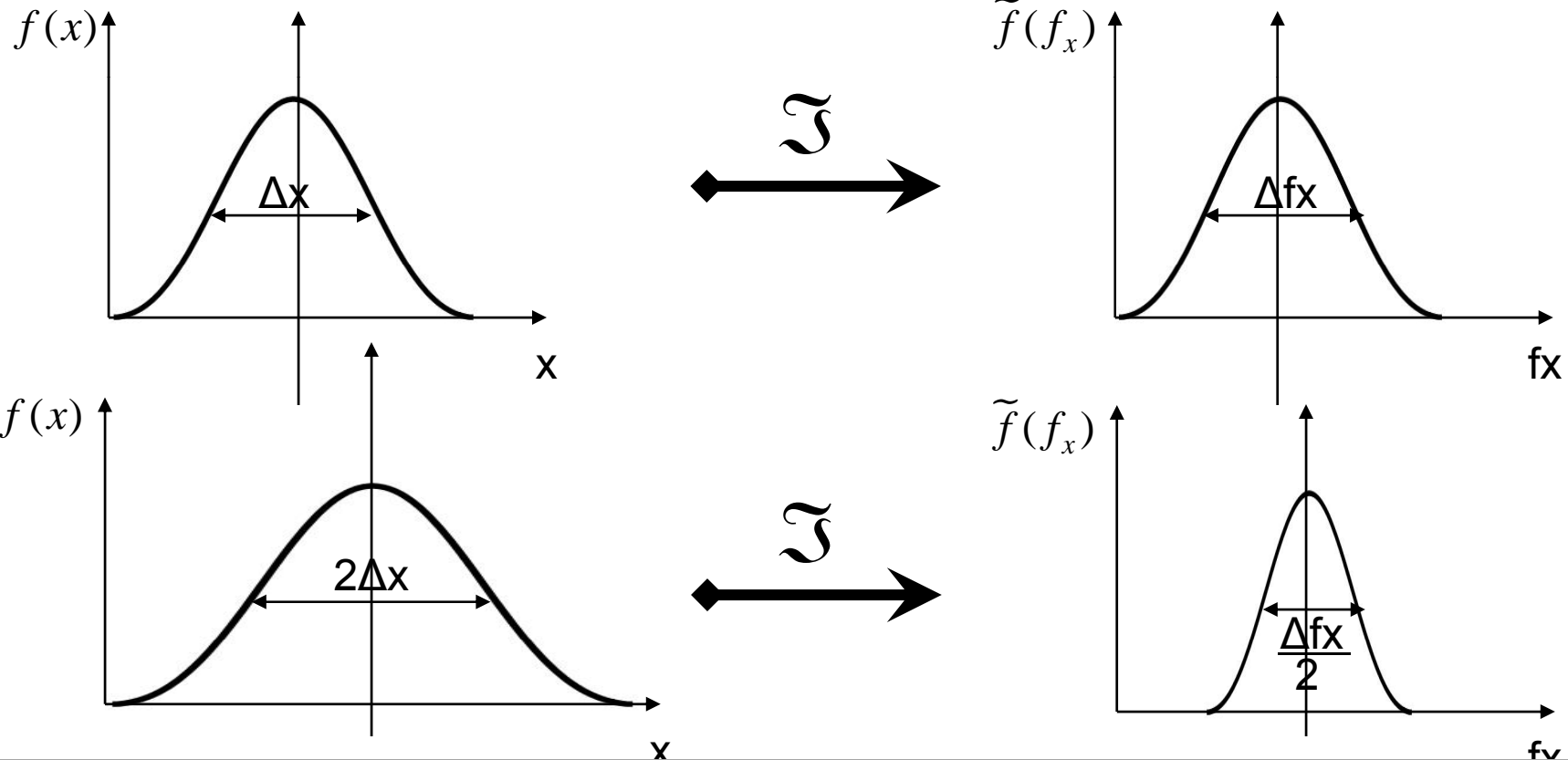
$$\mathfrak{F}[f(ax)] = \frac{1}{|a|} \tilde{f}\left(\frac{\xi}{a}\right) \quad (1.23)$$

- Theorem 1.23 provides intuitive feeling for F.T



1.6 Basic Theorems with Fourier Transforms

- Let's consider:



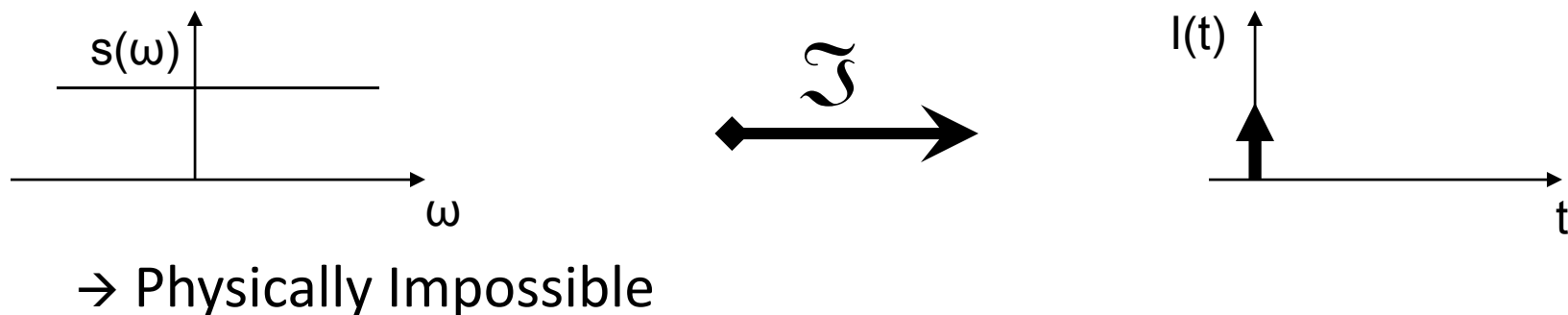


1.6 Basic Theorems with Fourier Transforms

! Broader functions in one domain imply narrower functions in the other and vice-versa

▪ Eg. To obtain short temporal pulses of light, one needs a broad spectrum (Ti: Saph laser)

! Only an infinite spectrum allows for δ -function pulses





1.6 Basic Theorems with Fourier Transforms

- Before we present the last theorems, we introduce the definitions of convolution and correlation

- Let

$$g(x) \xrightarrow{\mathcal{F}} G(\xi)$$

$$h(x) \xrightarrow{\mathcal{F}} H(\xi)$$

- Convolution of g and h:

$$g \otimes h = \int_{-\infty}^{\infty} g(x')h(x - x')dx' \quad (1.24)$$

- Correlation of g and h

$$g \otimes h = \int_{-\infty}^{\infty} g(x')h(x' - x)dx' \quad (1.25)$$



1.6 Basic Theorems with Fourier Transforms

- Difference between \otimes and \circledast is $h(x-x')$ vs $h(x'-x)$, i.e. flip vs non-flip of h
- Particular case:
 - Autocorrelation: $g=h$

$$g \otimes g = \int_{-\infty}^{\infty} g(x') g(x' - x) dx' \quad (1.26)$$

- Exercise: Use PC to show:

$$\text{Rect} \otimes \text{Rect} = \text{Triang}$$

$$\text{Sin} \otimes \text{Sin} = \text{Sin}$$

$$\text{Gauss} \otimes \text{Gauss} = \text{Gauss}$$



1.6 Basic Theorems with Fourier Transforms

d) Convolution theorem:

$$\mathcal{F}[g \otimes h] = GH \quad (1.27)$$

$$\text{i.e. } \mathcal{F}\left[\int_{-\infty}^{\infty} g(x')h(x-x')dx'\right] = G(\xi)H(\xi)$$

- Convolution in one domain corresponds to a product in the other. Nice!
- Multiplication is always easy to do
- Recall Green's function: $h(t)$ = the response to a δ -function light pulse



1.6 Basic Theorems with Fourier Transforms

- We found (Eq 1.16):

$$x(t) = \int_{-\infty}^{\infty} E(t')h(t-t')dt'$$

i.e the response to an arbitrary field $E(t)$ is the convolution $E \otimes h$!

- Let's take the F.T:

$$x(\omega) = E(\omega)h(\omega) \quad (1.28)$$

→ It doesn't get any simpler than this

i.e if we know the impulse response $h(t)$, (or the Green's function) take F.T → $h(\omega) \equiv$ transfer function → response to any field E is:

$$x(t) = \mathfrak{I}[E(\omega)h(\omega)] \quad (1.29)$$



1.6 Basic Theorems with Fourier Transforms

e) Correlation theorem:

- \otimes differs from \odot only by minus sign \rightarrow similar theorem:

$$\mathfrak{F}[g \otimes h] = GH^* \quad (1.30)$$

$$\text{i.e. } \mathfrak{F}\left[\int_{-\infty}^{\infty} g(x')h(x' - x)dx'\right] = G(\xi)H(\xi)^*$$

\rightarrow Particular case: $g = h$ (auto correlation):

$$\mathfrak{F}[g \otimes g] = GG^* = |G|^2 \quad (1.31)$$



1.6 Basic Theorems with Fourier Transforms

- Eg: F.T of an auto correlation is the power spectrum
- Very important for both time and space fluctuating fields:

$$\left[\begin{array}{l} \Gamma(t) = \int_{-\infty}^{\infty} E(t')E(t'-t)dt = \text{auto correlation} \\ \mathfrak{F}[\Gamma(t)] = E(\omega)E^*(\omega) = S(\omega) = \underline{\text{power spectrum}} \end{array} \right. \quad (1.32)$$

(Wiener–Khinchin theorem)

- We'll meet them again later!



1.7 Differential equations and Fourier Transforms

- Let f be a function of time:

$$f(t) = \int_{-\infty}^{\infty} F(\omega) e^{+i\omega t} d\omega = \mathfrak{F}^{-1}(F) \quad (1.33)$$

- What is $\frac{\partial f}{\partial t}$?

$$\begin{aligned} \frac{\partial f}{\partial t} &= \frac{\partial}{\partial t} \left[\int_{-\infty}^{\infty} F(\omega) e^{i\omega t} d\omega \right] = \\ &= \int_{-\infty}^{\infty} F(\omega) \frac{\partial}{\partial t} [e^{i\omega t}] d\omega = \\ &= \int_{-\infty}^{\infty} [i\omega F(\omega)] e^{i\omega t} d\omega \\ &= \mathfrak{F}^{-1} [i\omega F] \end{aligned} \quad (1.34)$$

- So, $f \rightarrow F$ & $\frac{\partial f}{\partial t} \rightarrow i\omega F \rightarrow$ Very Useful!



1.7 Differential equations and Fourier Transforms

- Great:

$$\left\{ \begin{array}{l} \mathfrak{F}[f(t)] = F(\omega) \rightarrow \text{Then} \\ \mathfrak{F}\left[\frac{\partial f(t)}{\partial t}\right] = i\omega F(\omega) \end{array} \right. \quad (1.34)$$

- Now $\mathfrak{F}\left[\frac{\partial^2 f}{\partial t^2}\right] = \mathfrak{F}\left[\frac{\partial}{\partial t}\left(\frac{\partial f}{\partial t}\right)\right] = i\omega i\omega F(\omega)$
 $= -\omega^2 F(\omega)$

In others words: $\mathfrak{F}\left[\frac{\partial^n f}{\partial t^n}\right] = i^n \omega^n F(\omega)$ (1.35)

- Differentiation theorem



1.7 Differential equations and Fourier Transforms

- Why 1.35 result is important? Because linear differential equations are resolved in the frequency domain more easily
- Eg: Recall our e^- revolving around nucleus under field illumination $E(t)$

$$m \frac{d^2 x(t)}{dt^2} + \gamma m \frac{dx(t)}{dt} + m \omega_o^2 x(t) = qE(t) \quad (1.36)$$

- The solution is $x(t)$. But we can solve for $x(\omega) = \mathfrak{F}[x(t)]$ and take \mathfrak{F}^{-1} in the end



1.7 Differential equations and Fourier Transforms

- So, let's take F.T of 1.36, using the differentiation theorem:

$$m[-\omega^2 x(\omega)] + i\omega\gamma m x(\omega) + m\omega_o^2 x(\omega) = qE(\omega)$$

$$x(\omega)[-m\omega^2 + i\omega\gamma m + m\omega_o^2] = qE(\omega)$$

Since $q=-e$:

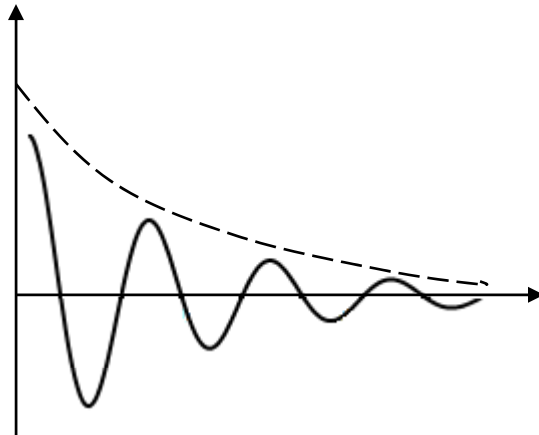
$$x(\omega) = \frac{\frac{e}{m} E(\omega)}{\omega^2 - i\gamma\omega - \omega_o^2}$$

(1.37)



1.7 Differential equations and Fourier Transforms

- Exercise: use PC to take \mathfrak{F}^{-1} of Eq. 1.37



- “damped” oscillation, γ = damping factor
! Problem solved



1.8 Refraction and absorption. Dispersion

- Given the electron displacement as a function of frequency, $x(\omega)$, we can define the dipole moment:

$$\bar{p} = q\bar{x}$$

$$\boxed{p = -ex} \quad (1.38)$$

- The dipole moment is a microscopic quantity; we need a macroscopic counterpart:

$$\bar{P} = N \langle \bar{p} \rangle = \frac{-\frac{Ne^2}{m} E}{\omega^2 - \omega_o^2 - i\gamma\omega} \quad (1.39)$$



1.8 Refraction and absorption. Dispersion

- $\bar{P} \equiv$ induced polarization
- $\bar{N} \equiv$ concentration [m^{-3}]
- But \bar{P} relates to the macroscopic response of the material χ , i.e. electric susceptibility:

$$\bar{P} = \epsilon_0 \chi \bar{E} \quad (1.40)$$

- ϵ_0 = permeability of vacuum
- Finally, $\chi = \epsilon_r - 1 = n^2 - 1$ (1.41)

- ϵ_r = relative permeability

- $n =$ refractive index
- If $\chi \in \mathbb{R}$, as opposed to $\chi = \begin{pmatrix} \chi_{11} & \chi_{12} & \chi_{13} \\ \chi_{21} & \chi_{22} & \chi_{23} \\ \chi_{31} & \chi_{32} & \chi_{33} \end{pmatrix}$, material is isotropic



1.8 Refraction and absorption. Dispersion

- So, combining 1.39 and 1.40:

$$\chi = \frac{Ne^2 / m\epsilon_0}{(\omega^2 - \omega_o^2) - i\gamma\omega} = n^2 - 1 \in \mathbb{C} \quad (1.42)$$

- For low-n materials, such as rarefied gases,

$$n^2 - 1 = (n - 1)(n + 1) \approx 2(n - 1)$$

$$\begin{aligned} \rightarrow n &= 1 + \frac{Ne^2}{2m\epsilon_0} \frac{1}{(\omega^2 - \omega_o^2) - i\gamma\omega} \\ &= n' + in'' \end{aligned} \quad (1.43)$$



1.8 Refraction and absorption. Dispersion

$$\rightarrow \begin{cases} n' = 1 + \frac{Ne^2}{2m\epsilon_0} \frac{\omega^2 - \omega_o^2}{(\omega^2 - \omega_o^2) - \gamma^2 \omega^2} & (1.44a) \\ n'' = \frac{Ne^2}{2m\epsilon_0} \frac{\gamma\omega}{(\omega^2 - \omega_o^2) - \gamma^2 \omega^2} & (1.44b) \end{cases}$$

- $n' = \text{Re}(n) =$ refractive index
- $n'' = \text{Im}(n) =$ absorption index



1.8 Refraction and absorption. Dispersion

- Eg: Plane wave:

$$E = E_o e^{i\vec{k}\cdot\vec{r}}; k = nk_o$$

$$\begin{aligned} E &= E_o e^{ink_o r} = \\ &= E_o e^{ik_o r (n' + in'')} \end{aligned}$$

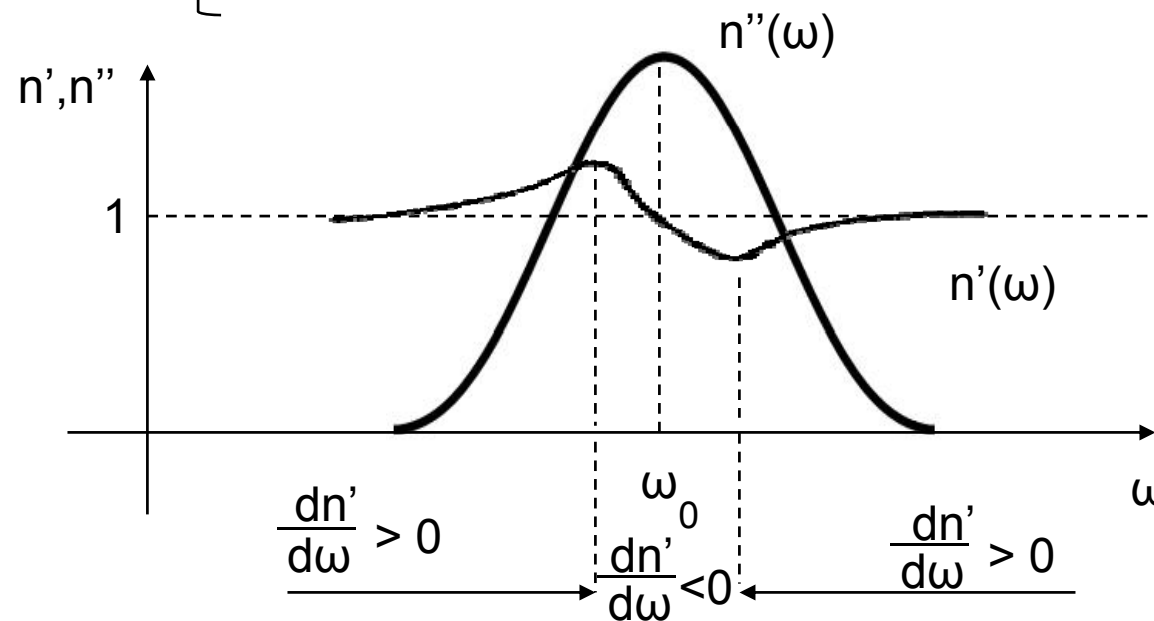
$$E = E_o \underbrace{e^{-n''k_o r}}_{\text{absorption}} \underbrace{e^{in'k_o r}}_{\text{refraction}}$$

$$\alpha = n''k_o = \text{absorption coefficient}$$



1.8 Refraction and absorption. Dispersion

- Definition:
 - $n'(\omega)$ = variation of refractive index with frequency
 - = dispersion
 - $n''(\omega)$ = absorption line shape





1.8 Refraction and absorption. Dispersion

- Note the line shape:

$$\frac{\gamma\omega}{(\omega^2 - \omega_o^2) + \gamma^2\omega^2} \approx \frac{1}{\gamma\omega} \frac{1}{1 + \left(\frac{(\omega - \omega_o)2\omega_o}{\gamma\omega}\right)^2} = \frac{1}{\gamma\omega} \frac{1}{1 + \left(\frac{\omega - \omega_o}{\gamma\omega / 2\omega_o}\right)^2}$$

- Lorentz function:

$$\mathfrak{L}(\omega) = \frac{1}{a} \left(\frac{1}{1 + (\omega / a)^2} \right) ; a = \text{width}$$

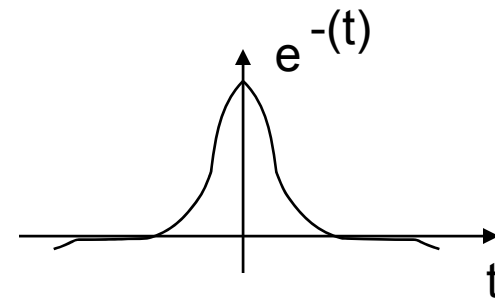


1.8 Refraction and absorption. Dispersion

- Thus the absorption line is a Lorentzian:

$$\alpha(\omega; \omega_o) = \frac{1}{\Delta\omega} \frac{1}{1 + \left(\frac{\omega - \omega_o}{\Delta\omega}\right)^2}; \Delta\omega \sim \gamma$$

$$\mathfrak{F}[\alpha(\omega)] = e^{-\Delta\omega|t|}$$



- The Fourier transform of a Lorentzian is an exponential!



1.8 Refraction and absorption. Dispersion

Connect to quantum mechanics:

- 2 level system:

$$1 \rightarrow E_1$$

$$0 \rightarrow E_0$$

$$\Delta E = E_1 - E_0 = \hbar(\omega_1 - \omega_0)$$

- Probability of spontaneous emission/absorption:

- $p(t) \sim e^{-t/\text{lifetime}} \rightarrow$ exponential decay

- Linewidth is Lorentz = natural linewidth

! The model of e^- on springs was introduced by... Lorentz



1.9 Maxwell's Equations

- Fully describe the propagation of EM fields
- Quantify how \hat{E} and \hat{H} generate each other

$$\left\{ \begin{array}{ll} \nabla \times \bar{E} = -\frac{\partial \bar{B}}{\partial t} & \text{I} \\ \nabla \times \bar{H} = \frac{\partial \bar{D}}{\partial t} + \bar{j} & \text{II} \\ \nabla \bar{D} = \rho & \text{III} \\ \nabla \bar{B} = 0 & \text{IV} \end{array} \right. \quad (1.45a)$$



1.9 Maxwell's Equations

- Plus material equations

$$\left\{ \begin{array}{l} \bar{D} = \epsilon_0 \bar{E} + \bar{P} \quad \text{V} \\ \quad = \epsilon \bar{E} \\ \bar{B} = \mu_0 \bar{H} + \bar{M} \quad \text{VI} \end{array} \right.$$

- Definitions:

\bar{E} = Electric field vectors ρ = charge density

\bar{H} = Magnetic field vectors $\bar{j} = \sigma \bar{E}$ = current density

\bar{D} = Electric displacement \bar{P} = polarization

\bar{B} = Magnetic inductance \bar{M} = magnetization



1.9 Maxwell's Equations

- Let's combine I and II (assume no free charge: $\rho = 0$, $\vec{j} = 0$)

- Use property: $\nabla \times (\nabla \times \vec{E}) = \nabla(\nabla \cdot \vec{E}) - \nabla^2 \vec{E}$

- Since $\rho = 0 \Rightarrow \nabla \cdot \vec{E} = 0 \Rightarrow \boxed{\nabla \times (\nabla \times \vec{E}) = -\nabla^2 \vec{E}}$

- Take $\nabla \times (EqI)$:

$$\nabla \times (\nabla \times \vec{E}) = -\nabla \times \left(\frac{\partial \vec{B}}{\partial t} \right) \quad (1.46)$$

$$\begin{aligned} \rightarrow -\nabla^2 \vec{E} &= \frac{\partial}{\partial t} (\nabla \times \vec{B}) = \\ &= -\frac{\partial}{\partial t} (\mu_0 \nabla \times \vec{H}) = \\ &= -\frac{\partial}{\partial t} \left(\mu_0 \frac{\partial \vec{\Delta}}{\partial t} \right) = \end{aligned}$$

(see next slide)



1.9 Maxwell's Equations

$$\begin{aligned}\rightarrow -\nabla^2 \bar{E} &= \frac{\partial}{\partial t} (\nabla \times \bar{B}) = \\ &= -\frac{\partial}{\partial t} (\mu_o \nabla \times \bar{H}) = \\ &= -\frac{\partial}{\partial t} \left(\mu_o \frac{\partial \bar{\Delta}}{\partial t} \right) = \\ &= -\epsilon \mu \frac{\partial^2 E}{\partial t^2}\end{aligned}$$

Thus: $\boxed{\nabla^2 \bar{E} - \epsilon \mu \frac{\partial^2 E}{\partial t^2} = 0}$ (1.47)

- Wave Equation



1.9 Maxwell's Equations

Note:

$$\left\{ \begin{array}{l} \epsilon\mu = \frac{1}{v^2}; \quad v = \frac{c}{n}; \quad c = \text{speed of light in vacuum} \\ n = \sqrt{\frac{\mu\epsilon}{\mu_0\epsilon_0}} \end{array} \right.$$

- The wave equation describes the propagation of a time-dependent field (eg. pulse)
- Solution: plane wave: $E = E_0 e^{-i(\omega t - \vec{k} \cdot \vec{r})}$
 - $k = \frac{2\pi}{\lambda} = \frac{\omega}{v} = n \frac{\omega}{c}$; $k = \text{wave equation}$



1.9 Maxwell's Equations

- Phase of the field:

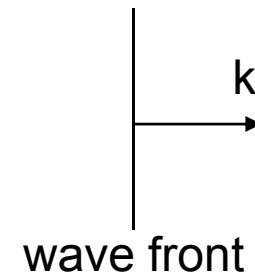
$$\varphi = \omega t - \bar{k} \cdot \bar{r} \quad (1.48)$$

- Note: $\varphi = \text{constant}$ describes a surface that moves with a certain velocity

$$\boxed{\omega t - \bar{k} \cdot \bar{r} = \text{constant}} \quad \text{eq of planes } \perp \bar{k}$$

$$\rightarrow \omega dt - k dr = 0$$

$$\rightarrow \boxed{\frac{dr}{dt} = \frac{\omega}{k} = v_p} \quad (1.49)$$



The surface of constant phase is traveling with velocity:

$$v_p = \frac{\omega}{k} = \underline{\text{phase velocity}}$$



1.9 Maxwell's Equations

- What is the counterpart of the wave equation in the frequency domain?

- Well, remember $\frac{\partial}{\partial t} \xrightarrow{\mathfrak{F}} i\omega$

- Upon Fourier transforming, Eq. 1.47 becomes:

$$\nabla^2 \bar{E} - \frac{1}{v^2} (i\omega \cdot i\omega) E(\omega) = 0$$

$$\rightarrow \nabla^2 \bar{E} + \frac{1}{v^2} (\omega^2) E(\omega) = 0$$

- Note: $k = \frac{\omega}{v}$

$$\rightarrow \nabla^2 E(\omega) + k^2 E(\omega) = 0 \quad (1.50)$$



1.9 Maxwell's Equations

$$\rightarrow \nabla^2 E(\omega) + k^2 E(\omega) = 0$$

- The equation above is the “Helmholtz equation”
- Describes how each frequency ω propagates