

Lab 2 - DFT - Problems

February 7, 2018

1 1 Fourier transform of Gaussian Functions

Recall the Gaussian function we defined in lab 1:

$$g(t) = \frac{1}{\sqrt{\pi}t_H} e^{-(t/t_H)^2}$$

where t_H is the half duration.

1. Plot $g(t)$ for $t_H = 10$ and $t_H = 20$ sec on the same graph.
2. (Bonus, 0.5 pt) derive the analytical formula for the Fourier transform of $g(t)$

$$G(\omega) = e^{-\frac{\omega^2 t_H^2}{4}}$$

3. Compute the discrete Fourier transform (DFT) for both sampled $g(t)$ time series, and compare them to the analytical $G(\omega)$ for both t_H 's on the same graph.

Hint: as numpy fft assumes signal starts from time 0, you can use the shift property of Fourier transform to first shift the $g(t)$ to start from zero, and after `fftshift(fft())` operations, multiply the spectrum by complex exponential sinusoid function.

4. Comment on the effect of filtering a general input time function $f(t)$ by $g(t)$ (i.e. convolution of $f(t)$ with $g(t)$), and explain the difference in filtered output after applying Gaussian functions with $t_H = 10$ or 20 secs.
5. Comment on how this is related to the time-frequency uncertainty principle (a signal cannot be infinitesimally sharp both in time and frequency).

2 2 Fourier transform of Window Functions

A continuous time signal $f(t)$ can be truncated into a signal of finite length T by window functions $b(t)$:

$$g(t) = f(t)b(t)$$

Typical window functions include:

- Boxcar function

$$b(t) = \begin{cases} 1 & 0 \leq t \leq T \\ 0 & \text{else} \end{cases}$$

- Hann window

$$b(t) = \begin{cases} \frac{1}{2} (1 - \cos \frac{2\pi t}{T}) & 0 \leq t \leq T \\ 0 & \text{else} \end{cases}$$

Now let $T = 10\text{sec}$, and sample both window functions by $\Delta t = 0.01\text{sec}$:

1. Plot both window functions on the same graph.
2. Calculate the Fourier transform of both functions by `numpy.fft()`. Pay extra attention to how you interpret the corresponding frequencies of output results from python. (*Hint: `fftshift()` may be useful. Also pay attention to the length of the input signal ($> 10\text{ sec}$), as it dictates the frequency resolution for the spectrum.*)
3. Plot the Fourier transform of both functions in the appropriate frequency range on the same graph.
4. Based on the FTs, comment on the effect of truncating a continuous time series by either window on its frequency spectrum $G(\omega)$ compared to the original spectrum $F(\omega)$.
5. Speculate on the advantages and disadvantages of boxcar and Hann window functions for truncation.

3 Radial Distribution Function

3.1 Background

Liquids have no fixed internal structure. Yet they do have some short range order in the sense that they have preferred intermolecular spacings, which are determined by the locations of minima in the intermolecular potentials. The microscopic structure of liquids is often characterized by a quantity known as the Radial Distribution Function $g(r)$, which is essentially the probability (Relative to the average probability, which means that $g(r)$ tends to 1 at large r , where the neighbour is too far away to feel any interaction.) that a molecule has a neighbouring molecule at distance r . Typically $g(r)$ shows a value that approaches zero at small r since molecules cannot occupy the same space; it also shows a peak at the preferred distance of nearest neighbours, and secondary peaks at preferred distances of more distant neighbours. If a suitable collimated beam of particles (e.g. X-rays or neutrons) is sent through a sample of the liquid, some of the particles are scattered. The number of particles scattered through a given angle is related to the Fourier Transform of $g(r)$ evaluated at the wavenumber k corresponding to the transverse momentum transfer associated with that scattering angle. Kittel derives this relationship in Chapter 17 of Introduction to Solid State Physics.

If this all sounds complicated, all you need to know here is that something called the Structure Factor $S(k)$ is effectively measured by looking at the scattered intensity as a function of scattering transverse wavenumber k (proportional to scattering angle), and that the Radial Distribution Function is related to it by

$$g(r) = 1 + \frac{1}{2\pi^2\rho r} \int_0^\infty k (S(k) - 1) \sin(kr) dk$$

where ρ is liquid number density (number of atoms per unit volume, computable from the three constants mentioned in the introduction), k is wavenumber, and r is radius.

1. You may have noticed some resemblance between expression (6) and the Fourier transform. First show that the integration part $\int_0^\infty k(S(k) - 1) \sin(kr) dk$ can be rewritten as

$$p(r) = \int_{-\infty}^{\infty} \frac{1}{2i} k (S(k) - 1) e^{ikr} dk.$$

- Now we can make some connections between the Radial Transfer Function and the Fourier Transform, if we substitute $r \rightarrow t$ and $k \rightarrow \omega$. What is the Fourier transform $P(k)$ of $p(r)$? Is $P(k)$ a real, imaginary or general complex function? Is it even or odd? **How will these affect $p(r)$? Is that what you expect?** Plot $P(k)$ as a function of k ranging from -15 \AA^{-1} to 15 \AA^{-1} based on `argon.py` (i.e. import and use the variables defined there).

- Write a Python function `[gn, rn] = RDFcalc(S, dk, rho)` to calculate Radial Distribution Function $g(r)$ from Structure Factor $S(k)$ data, sampled at dk , and density ρ , and output the RDF vector g_n and its corresponding radial distance vector r_n .

Hint: for Python `fft()` and `ifft()` functions, realize that the values of the Fourier Transform corresponding to negative frequencies are stored in the second half of the arrays given to (`ifft`) or obtained from it (`fft`). You also have to study the difference between the DFT and FT to multiply the right factors.

- With the data provided in `argon.py`, compute the corresponding Radial Distribution Function $g(r)$. Plot your results for r from 0 to 25 \AA . Over what range of radius can you trust your result?

Hint: To check if your results make sense, consider that $g(r)$ is the probability that a molecule has a neighbouring molecule at distance r , therefore, should be close to 0 when $r \rightarrow 0$, i.e. two molecules can not occupy the same space, and you can set $g(r = 0) = 0$. Also note the unit ρ used in $g(r)$ formula (6).

- From the $g(r)$ you computed, estimate the average molecular radius R_a of liquid argon. Give your reasoning and state what accuracy you can justify for your estimation.
- Now we explore the effect of sampling range. Yan sampled in wavenumber k out to $k_{max} = 15.24 \text{ \AA}^{-1}$, and he could have saved himself work by not collecting as much data, i.e., reducing k_{max} . But how much could he have reduced the sampling length k_{max} , while still see distinct peaks in the Radial Distribution Function? Also explain theoretically what you observe.

Hint: Plot on top of the $g(r)$ obtained in part 4, the $g(r)$'s you compute for a series of k_{max} values. You can try half k_{max} each time to look for changes. For the theoretical explanations for part 6 and 7, realize the interchangeability of $t \leftrightarrow \omega$ ($r \leftrightarrow k$).

- To explore the effect of data sampling, let's assume Yan decided to save his work by sampling less often (i.e. increasing dk). How large dk can he use to be able to still recover the first two peaks clearly? State your answers and a theoretical justification for what you expect to see if you increase dk too much.

Hint: Plot on top of the $g(r)$ obtained from `argon.py` data, the $g(r)$'s you obtain when you subsample the same dataset. Try double dk each time to observe the effect of coarser sampling.