

This writing focuses on 2 aspects of the 2011 paper written by Dawkins and others on "dispersive optical interface based on nanofiber-trapped atoms":

1. reproducing some of their equations, like Eqn.(4), using our quantitative theory.
2. following their line of experimental setup, derive more quantitative equations for atom number estimation with completely mixed state of atoms. → Also extend to clock-state case.

Part I: Reproduce the phase shift formula (Eqn.4) in Dawkins' paper) using our method.

In their experimental setup, atoms are initially prepared in a completely mixed state of  $F=4$  ground state manifolds.

The phase shift for a forward-propagating guided mode,  $u$ , can be given by

$$\begin{aligned}\phi_u &= 2\pi R \left( \frac{c}{v_g} \right) \sum_{\vec{r}} \text{Re} \left[ \frac{|\langle \text{eig} | \vec{r} \rangle \cdot \vec{u}_u(\vec{r})|^2}{\hbar(\Delta_{FF'} + i\Gamma/2)} \right] \leftarrow \text{ground state: } F=4, 6S_{1/2} \\ &= 2\pi R n_g \sum_{F'} \text{Re} \left[ \frac{|\langle J'F' | \vec{u}_u(\vec{r}) \rangle|^2}{\hbar(\Delta_{FF'} + i\Gamma/2)} \right] C_{J'F'F}^{(0)} |\vec{u}_u(\vec{r})|^2 \\ &= 2\pi R n_g |\vec{u}_u(\vec{r})|^2 \sum_{F'} C_{J'F'F}^{(0)} \frac{|\langle J'F' | \vec{u}_u(\vec{r}) \rangle|^2}{\hbar} \frac{\Delta_{FF'}}{\Delta_{FF'}^2 + \Gamma^2/4} \\ &= \frac{2\pi R}{A_{\text{eff}}} \frac{|\langle J'F' | \vec{u}_u(\vec{r}) \rangle|^2}{\hbar} C_{J'F'F}^{(0)} \sum_{F'} \frac{2\tilde{\Delta}_{FF'}}{\tilde{\Delta}_{FF'}^2 + 1}\end{aligned} \quad (1)$$

where  $\tilde{\Delta}_{FF'} = \Delta_{FF'}/\Gamma$ .

In deriving, we have used the fact that

$$\text{polarizability } \alpha(\Delta) = \alpha_0(\Delta) \sum_{F'} C_{J'F'F}^{(0)} = \frac{|\langle J'F' | \vec{u}_u(\vec{r}) \rangle|^2}{\hbar} \sum_{F'} \frac{C_{J'F'F}^{(0)}}{\Delta_{FF'} + i\Gamma/2}$$

↳ only scalar polarizability is considered here.

The effective mode area is defined as  $A_{\text{eff}} = \frac{1}{n_g |\vec{u}_u(\vec{r})|^2}$ .

On the other hand, the scattering cross section can be understood in the following way:

The energy attenuation for a collective spin ensemble can be given by

$$a = 4\pi k \text{Im}(\alpha) \rho = \rho \sigma(\Delta)$$

↳ coefficient → scattering cross section  
↳ atomic density.

For a single atom case, we treat  $\rho=1$ , and hence the scattering cross section

$$\sigma(\Delta) = 4\pi k \text{Im}(\alpha) = -4\pi k \frac{|\langle J'F' | \vec{u}_u(\vec{r}) \rangle|^2}{\hbar} \sum_{F'} \frac{\Gamma/2 \cdot C_{J'F'F}^{(0)}}{\Delta_{FF'}^2 + \Gamma^2/4} = \frac{8\pi k}{\Gamma} \frac{|\langle J'F' | \vec{u}_u(\vec{r}) \rangle|^2}{\hbar} \sum_{F'} \frac{C_{J'F'F}^{(0)}}{\tilde{\Delta}_{FF'}^2 + 1} = \sum_{F'} \frac{\sigma_{F'}}{\tilde{\Delta}_{FF'}^2 + 1}$$

As a special case, the resonant cross-section is defined at  $\Delta=0$  (usually only for 2-level systems):

$$\sigma_0 = \sigma(\Delta=0) = -8\pi k \frac{d_{ij}}{\hbar\Gamma} = -\frac{8\pi k}{\hbar\Gamma} \sum_{F'} |\langle J'F' | \vec{u}_u(\vec{r}) \rangle|^2 C_{J'F'F}^{(0)} \quad \text{with } \Gamma = \frac{4}{3}\hbar \deg k^3 \Rightarrow \sigma_0 = \frac{6\pi}{k^2} = \frac{3\lambda^2}{2\pi}$$

neg. not suitable for multi-level atoms.

We substitute the expression of  $\sigma_0$  into Eq.4 of  $\phi_u$ , and obtain

$$\text{resonant cross-section for } F \leftrightarrow F' \text{ transition: } \sigma_{F'} = \frac{8\pi k}{\hbar\Gamma} |\langle J'F' | \vec{u}_u(\vec{r}) \rangle|^2 C_{J'F'F}^{(0)}$$

$$\phi_u = \frac{-\sigma_0}{4A_{\text{eff}}} \sum_{F'} \frac{\sigma_{F'}}{\sigma_0} \frac{2\tilde{\Delta}_{FF'}}{\tilde{\Delta}_{FF'}^2 + 1} = -\phi_{\text{max}} \sum_{F'} \frac{\sigma_{F'}}{\sigma_0} \frac{2\tilde{\Delta}_{FF'}}{\tilde{\Delta}_{FF'}^2 + 1}$$

$$\text{with } \phi_{\text{max}} = \frac{\sigma_0}{4A_{\text{eff}}}, \quad \frac{\sigma_{F'}}{\sigma_0} = \frac{C_{J'F'F}^{(0)}}{C_{J'F'F}^{(0)}} \rightarrow \frac{\sigma_4}{\sigma_0} (F=4) = \frac{21}{44}, \quad \frac{\sigma_3}{\sigma_0} (F=4) = \frac{7}{44}$$

Therefore, with  $H$  &  $V$  modes, the rotation angle on the Poincaré sphere is

$$\theta_H, \theta_V, \theta$$

Therefore, with H & V modes, the rotation angle on the Poincaré sphere is

$$\boxed{\begin{aligned}\phi &\doteq \phi_H - \phi_V \\ &= -\frac{\sigma_S}{4A_{\text{eff}}} \cdot \sum_{F'} \frac{\sigma_{F'}}{\sigma_S} \frac{2\tilde{\Delta}_{FF'}}{\tilde{\Delta}_{FF'}^2 + 1} = -\phi_{\text{max}} \sum_{F'} 2 \frac{\sigma_{F'}}{\sigma_S} \frac{\tilde{\Delta}_{F'}}{\tilde{\Delta}_{F'}^2 + 1}\end{aligned}} \quad \leftarrow \text{Eq. (4) of Dawkin's paper! for } D_2 \text{ line transitions.}$$

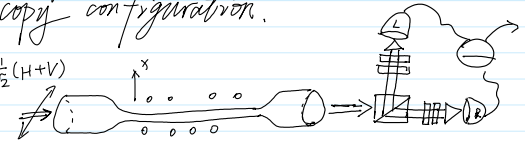
with  $A_{\text{eff}} = \frac{1}{n_g (|\vec{u}_H(\vec{r}_S)|^2 + |\vec{u}_V(\vec{r}_S)|^2)}$

$\phi_{\text{max}} = \frac{\sigma_S}{4A_{\text{eff}}} = \frac{OD}{4}$  with optical depth  $OD = \frac{\sigma_S}{A_{\text{eff}}}$ .

$\phi$  reaches maximum when  $\tilde{\Delta}_{FF'} = -1$  or  $\Delta_{FF'} = -\frac{\Gamma}{2}$ .

Part II: Minimum detectable atom number using a mixed state of atoms on the ground states. We continue to use our usual H-V birefringence spectroscopy configuration.

With  $N$  atoms, the signal power measured is

$\Delta P_S = P_0 \sin \phi_n \approx N P_0 \phi$  ( $P_0$  is the input power,  $\phi$  is the rotation/atom.) 

The shot noise is

$\Delta P_{SN} = \sqrt{\frac{P_0 \hbar \omega_0}{2\eta \tau_{pd}}}$  ( $\eta$  is the quantum efficiency,  $\tau_{pd}$  is the response time of the photon detector.)

In the shot-noise limit,  $\Delta P_S = \Delta P_{SN}$ .

We set  $\eta = 1$ , and  $\tau_{pd} = \frac{1}{\gamma_S}$  equals the photon scattering time.

$$\left\{ \begin{aligned} \gamma_S &= \sigma(\Delta) \frac{I(\vec{r}_S)}{\hbar \omega_0} \approx \frac{I(\vec{r}_S)}{\hbar \omega_0} \sum_{F'} \frac{\sigma_{F'}}{1 + \tilde{\Delta}_{FF'}^2} \Rightarrow \Delta P_{SN} = P_0 \sqrt{\frac{1}{2A_{\text{in}}}} \sqrt{\sum_{F'} \frac{\sigma_{F'}}{\tilde{\Delta}_{FF'}^2 + 1}} \\ I(\vec{r}_S) &= \frac{c \epsilon_0 n}{2} |E|^2|_{\vec{r}_S} = \frac{P_0}{A_{\text{in}}(\vec{r}_S)} \end{aligned} \right.$$

Aside:  $\left\{ \begin{aligned} &\text{For a Gaussian beam, the peak intensity } I_P = \frac{P_0}{\pi w^2/2}, A_{\text{in}} = \frac{\pi W^2}{2}. \\ &\text{For our nanofiber } \begin{cases} I(\vec{r}_S) = \frac{c \epsilon_0}{2} |E_0|^2 \frac{(\vec{u}_H(\vec{r}_S) + \vec{u}_V(\vec{r}_S))^2}{\sqrt{2}} = \frac{c \epsilon_0}{2} \frac{|E_0|^2}{2} (|\vec{u}_H(\vec{r}_S)|^2 + |\vec{u}_V(\vec{r}_S)|^2) = \frac{1}{2} (I_H(\vec{r}_S) + I_V(\vec{r}_S)) \text{ for } \phi' = 0 \text{ \& } \pi. \\ P_0 = \frac{c \epsilon_0}{2} \int n(\vec{r}_S) |E_0|^2 |\vec{u}_H(\vec{r}_S) + \vec{u}_V(\vec{r}_S)|^2 d\vec{r}_S \\ = \frac{c \epsilon_0}{2} \frac{|E_0|^2}{2} \left[ \int n(\vec{r}_S) (|\vec{u}_H(\vec{r}_S)|^2 + |\vec{u}_V(\vec{r}_S)|^2) d\vec{r}_S + \underbrace{\int_0^{2\pi} \int_0^\infty n(\vec{r}_S) [|\vec{u}_H|^2 + |\vec{u}_V|^2 - |\vec{u}_\phi|^2] \sin 2\phi \, d\phi \, r_S dr_S}_{=0 \text{ for odd function } \sin 2\phi} \right] \\ = \frac{1}{2} (P_H + P_V). \end{cases} \end{aligned} \right.$

Alternatively, using the circular polarization basis,

$$\left\{ \begin{aligned} \vec{u}_H &= \frac{1}{\sqrt{2}} (\vec{u}_+ + \vec{u}_-) \\ \vec{u}_V &= \frac{1}{i\sqrt{2}} (\vec{u}_+ - \vec{u}_-) \Leftrightarrow \begin{cases} \vec{u}_+ = \frac{1}{\sqrt{2}} (\vec{u}_H + i\vec{u}_V) \\ \vec{u}_- = \frac{1}{\sqrt{2}} (\vec{u}_H - i\vec{u}_V) \end{cases} \Rightarrow \begin{cases} I(\vec{r}_S) = \frac{c \epsilon_0}{2} \frac{|E_0|^2}{2} (|\vec{u}_H(\vec{r}_S)|^2 + |\vec{u}_V(\vec{r}_S)|^2) = \frac{1}{2} [I_+(\vec{r}_S) + I_-(\vec{r}_S)] \text{ for } \phi = 0 \text{ \& } \pi \\ P_0 = \frac{1}{2} (P_+ + P_-), \text{ with } P_\pm = \frac{c \epsilon_0}{2} \int n(\vec{r}_S) |E_0|^2 |\vec{u}_\pm(\vec{r}_S)|^2 d\vec{r}_S \end{cases} \\ \Rightarrow A_{\text{in}} &= \frac{P_H + P_V}{I_H(\vec{r}_S) + I_V(\vec{r}_S)} = \frac{P_+ + P_-}{I_+(\vec{r}_S) + I_-(\vec{r}_S)} \end{aligned} \right.$$

Therefore, in the shot-noise limit, the minimum detectable atom number

$$\boxed{\begin{aligned} N_{\text{min}} &= \sqrt{\frac{1}{2A_{\text{in}}}} \sqrt{\sum_{F'} \frac{\sigma_{F'}}{\tilde{\Delta}_{FF'}^2 + 1}} / |\phi| \\ &= 2A_{\text{eff}} \sqrt{\frac{1}{2A_{\text{in}}}} \sqrt{\sum_{F'} \frac{\sigma_{F'}}{\tilde{\Delta}_{FF'}^2 + 1}} / \left( \sum_{F'} \sigma_{F'} \frac{\tilde{\Delta}_{FF'}}{\tilde{\Delta}_{FF'}^2 + 1} \right) \\ &= \sqrt{\frac{2A_{\text{eff}}^2}{A_{\text{in}}}} \sqrt{\sum_{F'} \frac{\sigma_{F'}}{\tilde{\Delta}_{FF'}^2 + 1}} / \left( \sum_{F'} \frac{\tilde{\Delta}_{FF'} \sigma_{F'}}{\tilde{\Delta}_{FF'}^2 + 1} \right) \approx \sqrt{\frac{2A_{\text{eff}}}{A_{\text{in}} \sigma_0}} \end{aligned}}$$

Part III. To extend this result to the QND measurement case with atoms in clock states, we can use the solution for using magic frequencies that

$$\phi = \chi_{\text{eff}} = -\frac{\sigma_0}{A_{\text{eff}}} \frac{\Gamma}{4\Delta_{\text{eff}}}$$

$$\text{where } \begin{cases} \frac{1}{A_{\text{eff}}} = \frac{2}{\pi g} \frac{|\vec{u}_V(\vec{r}_0)|^2 \cdot |\hat{e}_\pi \cdot \vec{u}_H(\vec{r}_0)|^2 - |\vec{u}_H(\vec{r}_0)|^2 \cdot |\hat{e}_\pi \cdot \vec{u}_V(\vec{r}_0)|^2}{(|\vec{u}_H(\vec{r}_0)|^2 + |\vec{u}_V(\vec{r}_0)|^2)} \\ \frac{\Gamma}{4\Delta_{\text{eff}}} = \sum_{F'} \left( \frac{10 C_{J'F'F}^{(2)}}{\Delta_{4F'}} - \frac{6 C_{J'F'F}^{(2)}}{\Delta_{2F'}} \right) \frac{\Gamma}{4} \equiv \frac{1}{2} \frac{1}{\Delta_{\text{eff}}} \end{cases}$$

Using the formulas developed earlier, the minimum detectable atom number becomes

$$\begin{aligned} N_{\min} &= \sqrt{\frac{1}{2A_{\text{in}}}} \sqrt{\sum_{F,F'} \frac{\sigma_{FF'}}{\Delta_{FF'}^2 + 1}} / |\phi| \\ &= \sqrt{\frac{2A_{\text{eff}}^2}{A_{\text{in}}}} \frac{\tilde{\Delta}_{\text{eff}}}{\sigma_0} \sqrt{\sum_{F,F'} \frac{\sigma_{FF'}}{\Delta_{FF'}^2 + 1}} \\ &\simeq \sqrt{\frac{2A_{\text{eff}}^2}{A_{\text{in}} \sigma_0}} \tilde{\Delta}_{\text{eff}} \sqrt{\sum_{FF'} \frac{1}{\Delta_{FF'}^2 + 1}} \end{aligned}$$

For the clock-state atoms,

$$\sigma_{FF'} = -\sum_q \frac{8\pi R}{\hbar F} |\langle J' || d || J \rangle|^2 |O_{JF}^{J'F'}|^2 |C_{F',q}^{F,0;1,q}|^2$$

$U_0$

$\sim$   
 $\Delta_{\text{eff}}$