A coherent stimulated phonon spectrometer

Joel N. Johnson[1,1*], Peter T. Rakich[1], Co Authors[1], Ryan O. Behunin[1,1*]

[1*]Department of Applied Physics and Materials Science, Northern Arizona University, Flagstaff, 86011, AZ, USA.
[2]Center for Materials Interfaces in Research and Applications, Flagstaff, 86011, AZ, USA.
[3]Department of Applied Physics, Yale University, New Haven, 06520, CT, USA.
[4]Department, Organization, Street, City, 610101, State, Country.

*Corresponding author(s). E-mail(s): joel.johnson@nau.edu; ryan.behunin@nau.edu;

Abstract

We present a coherent stimulated phonon spectrometer that utilizes a pump-probe design to measure Brillouin scattering with unprecedented sensitivity. By employing three mechanisms to segregate the pump and probe light, we achieve sub-10 femptowatt sensitivity, enabling phonon spectroscopy on scales not previously possible. We demonstrate the capabilities of the instrument by observing Brillouin scattering in 1 centimeter and 1 millimeter of UHNA3 fiber, and 4 millimeters of bulk carbon disulfide, each at room temperature with sub-Watt optical power. This instrument paves new avenues for materials characterization and the development of novel nano-acousto-optic devices.

Keywords: Brillouin, femptowatt, nano-acousto-optic, phonon spectrometer

1 Introduction

[better first sentence] Brillouin scattering is a [less definition-y] fundamental process in which light interacts with acoustic phonons in a material, leading to a shift in the scattered light's frequency. [annihilation of ...] This process has been extensively studied for its potential applications in fields such as sensing, communications, and the development of optomechanical devices. However, [spontaneous, stimulated, would be nice if dedicated Stokes pump, but such an instrument would need to overcome the conflation of Stokes pump and retro-reflected Stokes-shifted probe signal..] conventional methods for measuring Brillouin scattering have been limited by their sensitivity, hindering the exploration of new phenomena and applications at the nanoscale. In this paper, we introduce a coherent stimulated phonon spectrometer that overcomes these limitations, enabling measurements with sub-10 femptowatt sensitivity.

2 Results

To demonstrate the capabilities of the instrument we choose two example targets, one fiber-coupled and one bulk material. In each case, our instrument exhibited high sensitivity and enabled the observation of Brillouin scattering at scales not previously achievable.

To demonstrate the capabilities of the instrument we perform coherent Brillouin spectroscopy on 1 mm of UHNA3 fiber and 4 mm of carbon disulfide liquid, serving as representative examples of the fiber-coupled and bulk materials regimes, respectively. In each case, our instrument exhibited high sensitivity and enabled the observation

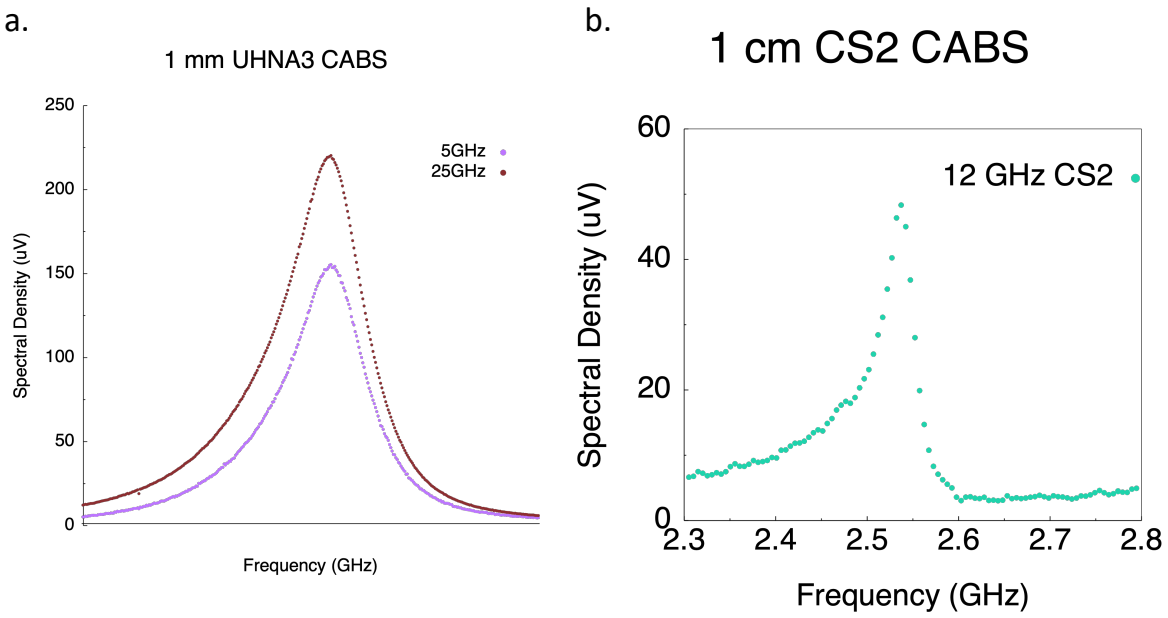

Fig. 1 This is a fig. This is an example of long caption this is an example of long caption this is an example of long caption this is an example of long caption

of Brillouin scattering at scales not previously achievable.

Fig. 1 shows the spectral measurements achieved by the instrument, overlaid with finite-difference simulation data. In Fig. 1a we see the expected lorentzian spectral shape in good alignment with simulation data for guided longitudinal modes ine the core of the UHNA3 fiber. In Fig. 1b, however, we see a distortion of this lorentzian shape. This is expected for partially unguided longitudinal modes, such as is the case for a bulk liquid filling the volume of a ...

First, we measured Brillouin scattering in a 1-centimeter-long UHNA3 fiber at room temperature and with sub-Watt optical power (Fig. 1a). Figure 1a, clearly displays the Brillouin scattering features with remarkable signal-to-noise ratio, highlighting the effectiveness of our apparatus in isolating the backscattered probe light. This observation serves as one of the main showcases of the instrument's capability.

Next, we performed Brillouin scattering measurements on a 4-millimeter-thick bulk carbon disulfide sample in a free-space optics arrangement. The observed spectrum, presented in Figure 2, exhibits well-resolved Brillouin scattering peaks. This successful measurement in a bulk sample demonstrates the versatility and adaptability of our instrument to various experimental configurations, further emphasizing the instrument's capability.

Lastly, we conducted a measurement in a 1-millimeter-long UHNA3 fiber under low-power conditions, with only 10 microwatts of power at the sample. Despite the reduced sample length and low power, the instrument's sensitivity allowed us to observe distinct Brillouin scattering features in the spectrum, as illustrated in Figure 3. This result underscores the potential of our spectrometer for nanoscale measurements and serves as a demonstration of the instrument's sensitivity, defining the sensitivity floor of the apparatus.

These three observations collectively showcase the high sensitivity, broad applicability, and impressive capabilities of our coherent stimulated phonon spectrometer in measuring Brillouin scattering across different sample types, scales, and power levels.

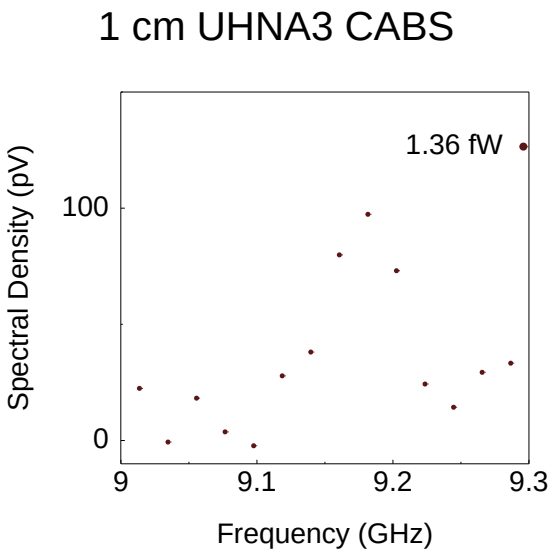

Fig. 2 This is a fig. This is an example of long caption this is an example of long caption this is an example of long caption this is an example of long caption

3 Theory

Here we derive the coupled wave equations that describe coherent stimulated Brillouin scattering involving a pump, Stokes, probe, and backscattered optical field given respectively by

$$\tilde{E}_P(z,t) = A_P e^{i(k_P z - \omega_P t)} + c.c. \quad (1)$$
$$\tilde{E}_S(z,t) = A_S e^{i(k_S z - \omega_S t)} + c.c. \quad (2)$$
$$\tilde{E}_{Pr}(z,t) = A_{Pr} e^{i(k_{Pr} z - \omega_{Pr} t)} + c.c. \quad (3)$$
$$\tilde{E}_{Sig}(z,t) = A_{Sig} e^{i(k_{Sig} z - \omega_{Sig} t)} + c.c. \quad (4)$$

and a common acoustic field given by

$$\tilde{\rho}(z,t) = \rho_0 + \rho(z,t) e^{i(qz - \Omega t)} + c.c., \quad (5)$$

where $\Omega = \omega_P - \omega_S$ and $q = k_P - k_S = 2k_P$.

3.1 Acoustic Field

We start by assuming that the material obeys the acoustic wave equation,

$$\frac{\partial^2 \tilde{\rho}}{\partial t^2} - \Gamma' \nabla^2 \frac{\partial \tilde{\rho}}{\partial t} - v^2 \nabla^2 \tilde{\rho} = \nabla \cdot \vec{f}, \quad (6)$$

where v is the sound speed in the material and Γ' is a damping parameter given by

$$\Gamma' = \frac{1}{\rho}\left[\frac{4}{3}\eta_s + \eta_b + \frac{\kappa}{C_p}(\gamma - 1)\right], \quad (7)$$

where η_s and η_b are the shear and bulk viscosity coefficients of the material, respectively. The source term on the right side of Eq. (6) is the divergence of the electrostrictive force:

$$\vec{f} = \nabla p_{st} = \nabla \cdot \left[-\frac{1}{2}\epsilon_0 \gamma_e \left(\langle \tilde{E}_P \cdot \tilde{E}_S \rangle + \langle \tilde{E}_{Pr} \cdot \tilde{E}_{Sig} \rangle\right)\right], \quad (8)$$

which yields, after applying the slowly varying amplitude approximation,

$$\nabla \cdot \vec{f} = \epsilon_0 \gamma_e q^2 (A_P A_S^* + A_{Pr} A_{Sig}^* e^{i\Delta k z}), \quad (9)$$

Where $\Delta k = (k_{Pr} - k_{Sig}) - (k_P - k_S)$ is the phase mismatch between the four optical fields. Only two electrostrictive terms survive terms after accounting for the orthogonal polarization of the pump and Stokes fields with respect to that of the probe and backscattered signal. Inserting this electrostrictive force term and the acoustic field (Eq. (5)) into Eq. (6) and assuming a slowly varying acoustic amplitude we find

$$-2i\Omega \frac{\partial \rho}{\partial t} - \Gamma' 2iq^2 \Omega \rho - 2iqv^2 \frac{\partial \rho}{\partial z} = \epsilon_0 \gamma_e q^2 (A_P A_S^* + A_{Pr} A_{Sig}^* e^{i\Delta k z}), \quad (10)$$

which can be restated in terms of the Brillouin linewidth, $\Gamma_B = q^2 \Gamma'$, as

$$-2i\Omega \frac{\partial \rho}{\partial t} - 2i\Omega \Gamma_B \rho - 2iqv^2 \frac{\partial \rho}{\partial z} = \epsilon_0 \gamma_e q^2 (A_P A_S^* + A_{Pr} A_{Sig}^* e^{i\Delta k z}). \quad (11)$$

Given the phonon dispersion relations $\Omega_B = |q_B| v$ and $\Omega^2 = q^2\left(v^2 - i\Omega\Gamma'\right)$, Eq. (11) can be rewritten as

$$-2i\Omega \frac{\partial \rho}{\partial t} + \left(\Omega^2 - \Omega^2 - i\Omega\Gamma_B\right)\rho - 2iqv^2 \frac{\partial \rho}{\partial z} = \epsilon_0 \gamma_e q^2 (A_P A_S^* + A_{Pr} A_{Sig}^* \quad (12)$$

We take the common assumption that the phonon propagation distance is small compared to the distance over which the source term varies significantly, which allows the spatial derivative term in Eq. (12). We further assume steady-state conditions such that the time derivative term also vanishes, leaving

$$(\Omega_B^2 - \Omega^2 - i\Omega\Gamma_B)_\rho = \epsilon_0 \gamma_e q^2 (A_P A_S^* + A_{Pr} A_{Sig}^* e^{i\Delta k z}). \quad (13)$$

We thus find the acoustic field amplitude to be

$$\rho(z,t) = \epsilon_0 \gamma_e q^2 \frac{A_P A_S^* + A_{Pr} A_{Sig}^* e^{i\Delta kz}}{\Omega_B^2 - \Omega^2 - i\Omega\Gamma_B}. \quad (14)$$

3.2 Optical Fields

We now turn to the spatial evolution of the optical fields, described by the wave equation,

$$\frac{\partial^2 \tilde{E}_i}{\partial z^2} - \frac{n^2}{c^2}\frac{\partial^2 \tilde{E}_i}{\partial t^2} = \frac{1}{\epsilon_0 c^2}\frac{\partial^2 \tilde{P}_i}{\partial t^2}, \quad (15)$$

where i denotes the four optical fields, namely: pump, Stokes, probe, and backscattered signal. The total nonlinear polarization that gives rise to the source term in the wave equation is given by

$$\tilde{P} = \epsilon_0 \Delta\chi \tilde{E} = \epsilon_0 \Delta\epsilon \tilde{E} = \epsilon_0 \rho^{-1} \gamma_e \tilde{\rho} \tilde{E}. \quad (16)$$

The parts of $\tilde{P}$ that can act as phase-matched source terms for the optical fields are

$$\tilde{P}_P = p_P e^{i(k_P z - \omega_P t)} + c.c. = \frac{1}{2}\epsilon_0 \rho_0^{-1} \gamma_e \rho A_S e^{i(k_P z - \omega_P t)} \quad (17)$$

$$\tilde{P}_S = p_S e^{i(-k_S z - \omega_S t)} + c.c. = \frac{1}{2}\epsilon_0 \rho_0^{-1} \gamma_e \rho^* A_P e^{i(-k_S z - \omega_S t)} \quad (18)$$

$$\tilde{P}_{Pr} = p_{Pr} e^{i(k_{Pr} z - \omega_{Pr} t)} + c.c. = \frac{1}{2}\epsilon_0 \rho_0^{-1} \gamma_e \rho A_{Sig} e^{i(k_{Pr} z - \omega_{Pr} t)} e^{i\Delta kz} \quad (19)$$

$$\tilde{P}_{Sig} = p_{Sig} e^{i(-k_{Sig} z - \omega_{Sig} t)} + c.c. = \frac{1}{2}\epsilon_0 \rho_0^{-1} \gamma_e \rho^* A_{Pr} e^{i(-k_{Sig} z - \omega_{Sig} t)} e^{-i\Delta kz}. \quad (20)$$

Inserting the optical fields (Eqs. 1-4) and phase-matched source terms (Eqs. 17-20) into Eq. (15), we obtain

$$\frac{\partial A_P}{\partial z} + \frac{n}{c}\frac{\partial A_P}{\partial t} = \frac{i\omega_P \gamma_e}{2nc\rho_0}\rho A_2 \quad (21)$$

$$-\frac{\partial A_S}{\partial z} + \frac{n}{c}\frac{\partial A_S}{\partial t} = \frac{i\omega_S \gamma_e}{2nc\rho_0}\rho^* A_P \quad (22)$$

$$\frac{\partial A_{Pr}}{\partial z} + \frac{n}{c}\frac{\partial A_{Pr}}{\partial t} = \frac{i\omega_{Pr} \gamma_e}{2nc\rho_0}\rho A_{Sig} \quad (23)$$

$$-\frac{\partial A_{Sig}}{\partial z} + \frac{n}{c}\frac{\partial A_{Sig}}{\partial t} = \frac{i\omega_{Sig} \gamma_e}{2nc\rho_0}\rho^* A_{Pr} \quad (24)$$

We again assume steady-state conditions, allowing the time derivative term to be dropped. Plugging in the acoustic field amplitude (Eq. 14), we arrive at the coupled-amplitude wave equations for the optical fields,

$$\frac{\partial A_P}{\partial z} = \frac{i\epsilon_0 \omega_P q^2 \gamma_e^2}{2nc\rho_0}\frac{A_P|A_S|^2 + A_{Pr}A_{Sig}^* A_S e^{i\Delta kz}}{\Omega_B^2 - \Omega^2 - i\Omega\Gamma_B} \quad (25)$$

$$\frac{\partial A_S}{\partial z} = -\frac{i\epsilon_0 \omega_S q^2 \gamma_e^2}{2nc\rho_0}\frac{|A_P|^2 A_S^* + A_{Pr}A_{Sig}^* A_P e^{i\Delta kz}}{\Omega_B^2 - \Omega^2 - i\Omega\Gamma_B} \quad (26)$$

$$\frac{\partial A_{Pr}}{\partial z} = \frac{i\epsilon_0 \omega_{Pr} q^2 \gamma_e^2}{2nc\rho_0}\frac{A_P A_S^* A_{Sig} + A_{Pr}|A_{Sig}|^2 e^{i\Delta kz}}{\Omega_B^2 - \Omega^2 - i\Omega\Gamma_B} \quad (27)$$

$$\frac{\partial A_{Sig}}{\partial z} = -\frac{i\epsilon_0 \omega_S q^2 \gamma_e^2}{2nc\rho_0}\frac{A_P A_S^* A_{Pr} + |A_{Pr}|^2 A_{Sig}^* e^{i\Delta kz}}{\Omega_B^2 - \Omega^2 - i\Omega\Gamma_B} \quad (28)$$

References