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Thermal Brillouin noise observed in silicon optomechanical waveguide

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Abstract

Stimulated Brillouin scattering was recently observed in nanoscale silicon waveguides. Surprisingly, thermally-driven photon–phonon conversion in these structures had not yet been reported. Here, we inject an optical probe in a suspended silicon waveguide and measure its phase fluctuations at the output. We observe mechanical resonances around 8 GHz with a scattering efficiency of 10^{-5} m⁻¹ and a signal-to-noise ratio of 2. The observations are in agreement with a theory of noise in these waveguides as well as with stimulated measurements. Our scheme may simplify measurements of mechanical signatures in nanoscale waveguides and is a step towards a better grasp of thermal noise in these new continuum optomechanical systems.

Keywords: optomechanics, nonlinear optics, silicon photonics, Brillouin scattering

(Some figures may appear in colour only in the online journal)

1. Introduction

The concept of photon-phonon interaction was conceived in the 1920s by Raman, Brillouin and Mandelstam [1-3] in the context of thermal scattering of photons by optical and acoustic phonons. Many previously predicted phenomena became experimentally accessible with the invention of lowloss fibers. In fact, backward Brillouin scattering-a stimulated process initiated by thermal phonons-is a main power limitation in fiber optical networks [4, 5] but has also been used to delay and store light at room temperature [6, 7]. Far afield, Braginsky studied analogous physical processes in gravitational wave detectors-realizing that photon-phonon coupling creates a mechanical instability similar to that in fibers at high optical powers [8]. The field has progressed to observations of photon-phonon conversion in fiber loops [9], photonic crystal fibers [10, 11] and whispering gallery resonators [12-14]. In the last decade, micro- and nano-scale confinement allowed for ever-increasing photon-phonon coupling rates in sub-wavelength fibers [15], chalcogenide rib waveguides [16, 17] and silicon optomechanical crystals [18, 19].

A recent branch of optomechanics seeks to demonstrate efficient coupling between photons and acoustic phonons in nanoscale silicon *waveguides* [20, 21]. These devices require milli- to centimeter interaction lengths and milliwatts of optical input power to observe mechanical action. They possess a continuum of optical and mechanical modes, enabling broadband functionality [22-24] that is otherwise not easily accessible in cavity optomechanics. As the mechanical propagation loss of these waveguides far exceeds their optical propagation loss [25], these waveguides typically provide optical gain-often called Brillouin amplification. Within the field of silicon photonics this effect was first seen in silicon/silicon-nitride hybrid suspended waveguides [26] and in silicon pedestal wires [27]. These observations soon led to reconfigurable microwave filters [22, 23]. Cascades of suspended wires enabled gain exceeding the propagation losses last year [28], which was improved recently in suspended silicon rib waveguides [29].

The goal of using these types of coupling to realize quantum and classical information processing, communications and sensing applications has motivated theoretical studies of classical and quantum noise in these structures [24, 30, 31]. Despite the exciting progress, a hallmark of



Figure 1. Thermal Brillouin scattering in suspended silicon waveguide. (a) Impression of a silicon-on-insulator waveguide that consists of a series of suspensions and socket anchors (left) and microscope image of a fabricated waveguide (right). The silicon socket layer masks the hydrofluoric etch of the silicon dioxide substrate. The silicon film is 220 nm thick and the wire width varies between 450 and 750 nm. Each suspension is 25 μ m long and the number of suspensions ranges from 30 to 120. Standard grating couplers provide waveguide access at in- and ouput. (b) Electric field norm of the quasi-TE polarized optical mode (top) and horizontal displacement of the acoustic mode (bottom). (c) Thermal motion of the waveguide down- and up-converts probe photons of frequency ω_p into sidebands of frequency $\omega_p \pm \omega_m$ with $\frac{\omega_m}{2\pi} \approx 8$ GHz.

photon-phonon coupling—the thermal Brownian motion has not yet been observed in these waveguides. Previous work focused either on optical instead of acoustic phonons [32–34], on megahertz vibrations in silicon slots [35, 36] and silica fibers [37, 38] or on gigahertz thermal-initiation of Brillouin amplification in chalcogenide waveguides [39]. In stark contrast, measurements of thermal occupation are standard in cavity optomechanics [40] where the goal is to read out and control the state of a mechanical oscillator. As a step toward understanding noise in these new continuum waveguide systems [24, 25, 30, 31, 41–43], we present the first observation of gigahertz Brownian motion in a silicon waveguide.

2. Results

We fabricated a series of silicon wire waveguides through a europractice-ePIXfab multi-project wafer run at imec (figure 1(a)). The waveguides' thickness was 220 nm and their widths were swept from 450 to 750 nm. The silicon was fully etched in trenches next to most of the waveguide, leaving a silicon core surrounded only by air and silicon dioxide. However, we periodically tapered a thin silicon 'socket' layer of thickness 70 nm (figure 1(a)) next to the silicon core. Finally, we removed the oxide substrate with a buffered hydrofluoric etch (etching for 11 min at 70 nm min⁻¹)—letting the socket layer serve as a mask. This produced a cascade of 25 μ m long suspended silicon beams. The resulting devices are identical to those of [28] except for the anchors, which now consist of the thin silicon socket layer

(figure 1(a)). In our collection of waveguides, we swept the number of suspensions from 30 to 120 at each width. We accessed the wires optically via standard grating couplers [44] for the quasi-TE polarized optical mode (figure 1(b)).

2.1. Experiment

We injected an optical probe at 1550 nm into the waveguide, aiming to observe Brownian mechanical motion of a Fabry– Pérot-like acoustic mode (figure 1(b)) that was previously observed by us in stimulated Brillouin measurements [28, 45]. The Brownian motion phase-modulates the optical probe, generating weak red- and blue-shifted sidebands (figure 1(c)) called the Stokes and anti-Stokes signals. We derived a new theoretical model for the Stokes- and anti-Stokes intensities and spectra in the appendix. A fiber Bragg filter rejected the anti-Stokes signal, leading only the probe and Stokes signals into a high-speed photodetector. Thus the optical phase fluctuations were transformed into intensity and photocurrent fluctuations. The probe was pre- and post-amplified by erbium-doped fiber amplifiers to have sufficient power at the photodetector (figure 2(b)).

We observed thermal Brillouin resonances between 5 and 10 GHz with a signal-to-noise ratio (SNR) of around 2 (figure 2(a)). Our model (see appendix) predicts the scattering efficiency η [1/m] to be

$$\eta = rac{\omega_{\mathrm{p}}}{4} rac{ ilde{\mathcal{G}}}{Q_{\mathrm{m}}} k_{\mathrm{B}} T pprox 10^{-5} \,\mathrm{m}^{-1}$$

with $\frac{\omega_p}{2\pi} = 193$ THz the probe frequency, $\frac{\tilde{g}}{Q_m} \approx 12 \text{ W}^{-1} \text{ m}^{-1}$ the non-resonant Brillouin gain coefficient measured in stimulated measurements [28, 45] and $k_B T = 25$ meV. In other words, in a millimeter of propagation length about 10 probe photons per billion are scattered by thermal phonons. Our model agrees with another recently developed treatment of noise [47] in forward Brillouin interactions.

We ensured that the noise background of the electrical spectrum analyzer scaled with optical power. Assuming this background was set by the erbium-doped fiber amplifier, we obtain (see appendix) an estimated SNR of

$$\mathrm{SNR} = \frac{4\Phi_{\mathrm{s}}}{F_{\mathrm{n}}\kappa_{\mathrm{m}}} \approx 2$$

with $\Phi_s = \Phi_{as} = \eta L \Phi_p$ the (anti-)Stokes photon flux $[s^{-1}]$, *L* the suspended waveguide length, Φ_p the probe photon flux at the ESA, κ_m the acoustic decay rate and F_n the noise factor of the fiber amplifier. Above we inserted typical parameters for our set-up:

$$\begin{split} \hbar \omega_{\rm p} \Phi_{\rm p} &\approx 1 \text{ mW} \\ \Phi_{\rm p} &\approx 10^{16} \text{ s}^{-1} \\ \eta &\approx 10^{-5} \text{ m}^{-1} \\ L &\approx 1 \text{ mm} \\ \frac{\kappa_{\rm m}}{2\pi} &\approx 10 \text{ MHz} \\ F_{\rm p} &\approx 3. \end{split}$$

The estimated SNR agrees roughly with the measured SNRs (figure 2(a)).



Figure 2. Thermal Brillouin spectrum in suspended silicon waveguide. (a) An example of a thermal Brillouin resonance in a suspended silicon waveguide of width w = 750 nm consisting of N = 120 suspensions each 25 μ m long. The resonance has a signal-to-noise ratio of SNR = 2.7 and a quality factor of $Q_m = 918$. The RF spectrum was normalized with respect to both the shot noise background and the response of the electrical spectrum analyzer (ESA). (b) We inject an optical probe into the waveguide. It is preamplified by an erbium-doped fiber amplifier (EDFA). The mechanical noise phase-modulates the optical probe as it travels along the waveguide. This optical phase modulation is transducted to intensity modulation by a fiber Bragg grating (FBG) that usually rejects the anti-Stokes signal. We observe identical spectra when the FBG rejects the Stokes instead of the anti-Stokes signal. Thus the Brownian mechanical motion of the waveguide is read out optically on an ESA. A switch determines whether we send the waveguide transmission to the ESA or a power meter (PM). To measure the thermal spectra, the switch is in the 'cross' state—leading waveguide transmission to the ESA. The set-up is simpler than gain and cross-phase modulation experiments [26, 28, 29, 45, 46] previously used on such devices. The power dropped in the waveguide was typically around 20 mW, of which 1 mW remained after the output grating coupler. The FBG and BPF had 6 and 4 dB insertion loss, which was compensated for by the post-EDFA. Thus the ESA photodetector received a couple of milliwatts of probe power.



Figure 3. Thermal Brillouin spectrum as a function of probe wavelength. The RF spectra depend strongly on the probe wavelength relative to the FBG rejection band. The red curve sketches the estimated FBG position as a function of transmission (unitless) versus probe wavelength. It is parallel to the wavelength axis at an arbitrary frequency (5.56 GHz) and z-range (1-1.5). Within the FBG rejection band the acoustic resonance is not visible at all. As the probe wavelength increases, the probe and the redshifted Stokes sideband make it through the FBG and next the post-EDFA. Then the interplay between EDFA gain and post-FBG probe power determine the SNR, as long as the blue-shifted anti-Stokes sideband remains rejected by the FBG. We find RF spectra with maximum SNR close to the FBG's band edge. Figures 2 and 4 are based on the maximum SNR spectra. The FBG's band edge is 2.75 GHz (0.22 nm) wide and it has an extinction ratio of 30 dB. This figure concerns a waveguide of width w = 750 nm with N = 120 suspension, the same one as in figure 2(a).

Further, the RF spectra depend heavily on the probe wavelength with respect to the fiber Bragg grating's (FBG's) rejection band (figure 3). Within the FBG rejection band, the thermal Brillouin resonances are not visible at all, as too little optical power reaches the photodetector. As the probe wavelength reaches the FBG's band edge, the probe and its red-shifted Stokes sideband make it through to the photodetector (figure 2(b)). Then the interplay between the EDFA gain—which depends on the probe power—and the post-FBG probe power determine the SNR, as long as the blue-shifted anti-Stokes sideband remains rejected in order to convert phase- to intensity-fluctuations. We find a maximimum SNR close to the FBG's band edge, and use these spectra to extract the acoustic frequencies $\frac{\omega_m}{2\pi}$, linewidths $\frac{\kappa_m}{2\pi}$ and quality factors Q_m with a Lorentzian fit (figure 2(a)).

The acoustic resonance frequencies scale inversely with the wires' width (figure 4(a)), in agreement with finite-element simulations and previous stimulated Brillouin measurements [28, 45]. The scaling can be understood if we consider the acoustic mode to be the fundamental excitation of a Fabry-Pérot resonator-with a transverse wavelength equal to twice the waveguide width w. This directly implies $\frac{\omega_{\rm m}}{2\pi} \propto \frac{1}{w}$ [28, 45]. Further, we extract the acoustic quality factors $Q_{\rm m}$ for a range of wire widths w and number of suspensions N. The quality factors are highly dependent on these two parameters, but the pattern is not monotonic. Overall we see higher quality factors in wider waveguides with less suspensions (figure 4(b)). This trend is likely caused by geometric disorder: the wire width fluctuates along the waveguide, inhomogeneously broadening the acoustic resonance. We saw such effects also in stimulated measurements [28]. A theoretical treatment of inhomogeneous broadening is given in [48]. With a frequency sensitivity of 20 MHz nm^{-1} (figure 4(a) and [28, 45]), the acoustic resonance is vulnerable to even atomic-scale disorder. This is the cost of miniaturization and is a major challenge that must be resolved. Similar effects have been seen in other nanoscale devices, such as



Figure 4. Mechanical frequencies and quality factors deduced from thermal spectra. (a) The resonance frequencies are in agreement with the 1/w Fabry–Pérot model of the acoustic resonance [28, 45]. (b) The acoustic quality factors are highly dependent on the waveguide width and number of suspensions. The pattern is not monotonic, but generally we find higher quality factors in wider and shorter waveguides. This behavior was also seen in stimulated measurements and is likely caused by geometric disorder along the waveguide [28].

silicon rib waveguides [29] and snowflake optomechanical crystals [18].

3. Conclusion

We observed thermal Brillouin scattering in a silicon photonic waveguide and presented a new model for the scattering efficiencies and noise spectra. Our theory is in exact agreement with another model [47] and in rough agreement with our observations. We analyzed the acoustic frequencies and quality factors for a range of waveguide widths and suspension lengths, finding close correspondence to earlier stimulated Brillouin measurements [28, 45]. The acoustic quality factors are generally higher in shorter and wider waveguides, likely because of atomic-scale geometric disorder. The measurement scheme is simpler than previous stimulated Brillouin set-ups. It is a step towards a better understanding of noise in continuum waveguide systems and may generate alternative approaches to on-chip thermometry [49, 50].

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Appendix

Phase-shift model for the thermal spectrum

We consider a translationally invariant and lossless waveguide of length *L*. We inject a weak optical probe at frequency $\frac{\omega_p}{2\pi}$. The probe is weakly phase-modulated by the thermal motion of the waveguide. Assuming the waveguide has a mechanical mode at frequency $\frac{\omega_m}{2\pi}$, this generates sidebands at angular frequency $\omega_{\rm p} \pm \omega_{\rm m}$. Here we seek the spectrum and intensity of the scattered light—neglecting optical driving of the mechanical mode. We denote the optical field at the output of the waveguide by $a_{\rm out}(\tau)$. The mechanical motion phase-shifts the input field $a_{\rm p}(\tau)$:

$$a_{\text{out}}(\tau) = a_{\text{p}}(\tau)e^{i\phi(\tau)} + \xi(\tau)$$
$$\approx a_{\text{p}}(\tau)(1 + i\phi(\tau)) + \xi(\tau)$$

with $\phi(\tau)$ a random variable with vanishing ensemble average ($\langle \phi(\tau) \rangle = 0$) and $\xi(\tau)$ the vacuum noise. Here we treat the phase shift ϕ as induced by a lumped element. This is a good approximation since in forward intra-modal Brillouin scattering momentum conservation dictates that the acoustic phase velocity equals the optical group velocity [26, 45]. Thus $\Lambda = \frac{2\pi v_g}{\omega_m} \approx 1 \text{ cm} \gg L \approx 1 \text{ mm}$ with Λ the acoustic wavelength and $v_g = 7 \times 10^7 \text{ m s}^{-1}$ the optical group velocity. Therefore the finite acoustic wavevector can be considered zero.

Field spectral density

First we look at the spectrum of the output *field*, which would result from a spectrally selective photon-counting experiment. The autocorrelation of output field is

$$\begin{aligned} \langle a_{\text{out}}^{\dagger}(\tau) a_{\text{out}}(0) \rangle &= \langle a_{\text{p}}^{\dagger}(\tau) a_{\text{p}}(0) \rangle \\ &+ \sqrt{\Phi_{\text{p}}} \langle \phi(\tau) \phi(0) \rangle + \langle \xi^{\dagger}(\tau) \xi(0) \rangle \end{aligned}$$

with Φ_p the probe photon flux (s⁻¹) and where we treated the phase shift ϕ and the probe field a_p as uncorrelated and ϕ as a real observable. We took $a_p(\tau) = \sqrt{\Phi_p}$ to be a constant in the second term, neglecting noise in the probe. The output field spectral density $S_a(\omega)$ equals

$$S_a(\omega) = \int_{-\infty}^{\infty} d\tau \ e^{i\omega\tau} \langle a_{out}^{\dagger}(\tau) a_{out}(0) \rangle$$

= $S_p(\omega) + \Phi_p S_{\phi}(\omega)$

with $S_p(\omega)$ the probe spectrum, $S_{\phi}(\omega)$ the spectral density of the phase and we used $\langle \xi^{\dagger}(\tau)\xi(0)\rangle = 0$ for optical frequencies with negligible thermal occupation. We also have

$$\phi(\tau) = k_{\rm p} \,\partial_q n_{\rm eff} \,q(\tau)I$$

with $k_p = \frac{\omega_p}{c}$ the vacuum probe wavevector, *c* the speed of light, $\partial_q n_{\text{eff}}$ the sensitivity of the optical effective index n_{eff} to motion *q* and $q(\tau)$ a coordinate describing the average motion along the entire waveguide of length *L*. Its ensemble average vanishes $\langle\langle q(\tau) \rangle = 0\rangle$ and it can be written as $q(\tau) = \frac{1}{L} \int_0^L q(z, \tau) dz$ with q(z, t) the actual motion at point *z* along the waveguide. Note that $q(\tau)$ represents a single degree of freedom if the curvature of the phonon band is neglected—an excellent approximation here³. It is therefore an effective harmonic oscillator that corresponds to the average motion along the entire waveguide. Thus the spectral density of the phase is

$$S_{\phi}(\omega) = \int_{-\infty}^{\infty} \mathrm{d}\tau \; \mathrm{e}^{\mathrm{i}\omega\tau} \langle \phi(\tau)\phi(0) \rangle$$
$$= (k_{\mathrm{p}}\partial_{q}n_{\mathrm{eff}}L)^{2}S_{q}(\omega).$$

Further, the spectral density of the motion $S_a(\omega)$ is [51]

$$\begin{split} S_q(\omega) &= q_{\rm zpf}^2 \\ &\times \left(\frac{\kappa_{\rm m} \overline{n}}{(\omega_{\rm m} + \omega)^2 + \kappa_{\rm m}^2/4} + \frac{\kappa_{\rm m} (\overline{n} + 1)}{(\omega_{\rm m} - \omega)^2 + \kappa_{\rm m}^2/4} \right) \end{split}$$

in thermal equilibrium for a weakly damped oscillator and with $q_{\rm zpf} = \sqrt{\frac{\hbar}{2m_{\rm eff}L\omega_{\rm m}}}$ the zero-point fluctuation, $m_{\rm eff}$ the effective modal mass per unit length and \bar{n} the thermal phonon occupation in the degree of freedom described by $q(\tau)$. At 300 K we have $\bar{n} \approx \frac{k_B T}{\hbar\omega_{\rm m}} = 624$ for a 10 GHz acoustic mode. This implies the two sidebands are of nearly equal intensity. Besides, \bar{n} is the number of phonons in the degree of freedom captured by the coordinate $q(\tau)$ as defined above. This phonon occupation \bar{n} does not scale with waveguide length L, which means that $S_q(\omega) \propto \frac{1}{L}$, and therefore $S_{\phi}(\omega) \propto L$ as expected from a random walk: the root-mean-square of the phase $\sqrt{\langle \phi^2 \rangle}$ goes as \sqrt{L} . Thus the blue- and red-shifted sideband intensities increase linearly with length.

Previously we showed that [45]

$$\tilde{\mathcal{G}} = 2\omega_{\rm p} \frac{Q_{\rm m}}{k_{\rm eff}} \left(\frac{1}{c} \partial_q n_{\rm eff}\right)^2$$

³ Consider the waveguide of length *L* as a series of independent oscillators. The phase shift then becomes $\phi \propto \sum_z q_z$ with q_z the motion of the oscillator at position *z*. Applying periodic boundary conditions and with Fourier decomposition $q_k = \frac{1}{\sqrt{L}} \sum_z e^{-ikz} q_z$, we see that $\phi \propto q_{k=0}$, representing a single degree of freedom that appears quadratically in the energy. If the phonon band is flat, the precise boundary conditions do not impact this conclusion. The band of the mechanical mode studied here remains flat if the acoustic wavelength is longer than just a couple of microns.

⁴ The scaling $S_q(\omega) \propto \frac{1}{r}$ can be understood from $\langle q(z', \tau)q(z, \tau) \rangle \propto \delta(z'-z)$: the thermal Brownian motion is delta-correlated along the waveguide, unlike in stimulated Brillouin schemes with quadratic build-up [45, 47].

with $k_{\rm eff} = m_{\rm eff} \omega_{\rm m}^2$ the effective mechanical stiffness per unit length, $\tilde{\mathcal{G}}$ the Brillouin gain coefficient (W⁻¹m⁻¹) and $Q_{\rm m} = \frac{\omega_{\rm m}}{\kappa_{\rm m}}$. So the total output spectral density of the field is

$$S_a(\omega) = S_p(\omega) + \Phi_p L \eta \left(\frac{\kappa_m}{(\omega_m + \omega)^2 + \kappa_m^2/4} + \frac{\kappa_m}{(\omega_m - \omega)^2 + \kappa_m^2/4} \right)$$

with η the scattering efficiency (m⁻¹) and

$$\eta = \frac{\omega_{\rm p}}{4} \frac{\tilde{\mathcal{G}}}{Q_{\rm m}} k_B T.$$

This result is in agreement with another recently developed noise model [47] for forward Brillouin interactions. Since

$$\int_{-\infty}^{\infty} \frac{\mathrm{d}\omega}{2\pi} \frac{\kappa_{\mathrm{m}}}{(\omega_{\mathrm{m}} \pm \omega)^2 + \kappa_{\mathrm{m}}^2/4} = 1$$

we have

$$\Phi_{\rm s} = \Phi_{\rm as} = \Phi_{\rm p} L \eta$$

with Φ_s and Φ_{as} the Stokes and anti-Stokes photon flux.

Current spectral density

Instead of doing a photon-counting experiment, we use a high-speed photodetector. Photodetectors cannot directly detect the phase $\phi(\tau)$. Therefore, we first send the output field through a fiber Bragg filter, rejecting either the Stokes or anti-Stokes sideband. Thus the symmetry between the sidebands is broken and the Stokes (or anti-Stokes) signal produces a beat note at ω_m in the photocurrent $I(\tau)$. Here we briefly discuss the properties of the current spectral density. The filtered output field is

$$a_{\text{out}}(\tau) = a_{\text{pf}}(\tau) + a_{\text{s}}(\tau) + \xi(\tau)$$

with the Stokes (or anti-Stokes) field $a_s(\tau) = i\sqrt{\Phi_p}\phi_f(\tau)$ and where 'f' stands for the filtered fields after the fiber Bragg grating. Note that the field spectrum is asymmetric because of the filter. The current $I(\tau)$ is

$$I(\tau) = a_{\text{out}}^{\dagger}(\tau) a_{\text{out}}(\tau) = I_{\text{pf}}(\tau) + \sqrt{\Phi_{\text{p}}} (a_{\text{s}}(\tau) + a_{\text{s}}^{\dagger}(\tau) + \xi(\tau) + \xi^{\dagger}(\tau))$$

with $I_{\rm pf}(\tau)$ the probe current and where we neglected weak currents associated with the Stokes field and the shot noise. So the current spectral density $S_{II}(\omega)$ is

$$\begin{split} S_{II}(\omega) &= \int_{-\infty}^{\infty} \mathrm{d}\tau \; \mathrm{e}^{\mathrm{i}\omega\tau} \langle I(\tau)I(0) \rangle \\ &= S_{I_{\mathrm{pf}}}(\omega) + \Phi_{\mathrm{p}}^{2}(S_{\phi_{\mathrm{f}}}(\omega) + S_{\phi_{\mathrm{f}}}(-\omega)) + F_{\mathrm{n}}\Phi_{\mathrm{p}}, \end{split}$$

where we used $\langle \xi(\tau)\xi^{\dagger}(0) \rangle = \delta(\tau)$. We multiplied the shot noise (third term) by the noise factor $F_{\rm n} \approx 3$ of the EDFA. During the experiment, we ensured that the background on the electrical spectrum analyzer scaled with the optical power. Further, the spectral density $S_{\phi_{\rm r}}(\omega)$ of the filtered phase $\phi_{\rm f}$ is

$$S_{\phi_{\rm f}}(\omega) = L\eta \frac{\kappa_{\rm m}}{(\omega_{\rm m} \pm \omega)^2 + \kappa_{\rm m}^2/4},$$

where the \pm indicates whether the fiber Bragg filter rejected the Stokes (+) or anti-Stokes (-) – which is determined by the probe wavelength. Assuming the EDFA sets the background, the SNR equals

$$\mathrm{SNR} = \frac{\Phi_{\mathrm{p}} S_{\phi_{\mathrm{f}}}(\omega_{\mathrm{m}})}{F_{\mathrm{n}}} = \frac{4\Phi_{\mathrm{s}}}{F_{\mathrm{n}}\kappa_{\mathrm{m}}}.$$

References

- [1] Raman C V and Krishnan K S 1928 Nature 121 501-2
- [2] Brillouin L 1922 Ann. Phys. 17 21
- [3] Mandelstam L 1926 Zh. Russ. Fiz-Khim. Ova 58 381
- [4] Agrawal G 2013 *Nonlinear Fiber Optics* (New York: Academic)
- [5] Boyd R, Rzałewski K and Narum P 1990 Phys. Rev. A 42 5514–21
- [6] Zhu Z, Gauthier D and Boyd R 2007 Science 318 1748-50
- [7] Song K, Herráez M and Thévenaz L 2005 *Opt. Express* 13 82–8
- [8] Braginsky V, Strigin S and Vyatchanin S 2001 Phys. Lett. A 287 331–8
- [9] Hill K O, Kawasaki B S and Johnson D C 1976 Appl. Phys. Lett. 28 608–9
- [10] Beugnot J C, Sylvestre T and Maillotte H 2007 Opt. Lett. 32 2006–8
- [11] Kang M, Nazarkin A, Brenn A and Russell P 2009 Nat. Phys. 5 276–80
- [12] Kippenberg T, Rokhsari H, Carmon T, Scherer A and Vahala K 2005 Phys. Rev. Lett. 95 033901
- [13] Grudinin I, Matsko A and Maleki L 2009 Phys. Rev. Lett. 102 043902
- [14] Lee H, Chen T, Li J, Yang K Y, Jeon S, Painter O and Vahala K J 2012 Nat. Photon. 6 369–73
- [15] Beugnot J C, Lebrun S, Pauliat G, Maillotte H, Laude V and Sylvestre T 2014 Nat. Commun. 5 5242
- [16] Kabakova I 2013 Opt. Lett. 38 3208–11
- [17] Casas-Bedoya A et al 2016 Frontiers in Optics 2016
- [18] Safavi-Naeini A H, Hill J T, Meenehan S, Chan J, Gröblacher S and Painter O 2014 Phys. Rev. Lett. 112 153603
- [19] Chan J, Alegre T, Safavi-Naeini A H, Hill J, Krause A, Gröblacher S, Aspelmeyer M and Painter O 2011 Nature 478 89–92
- [20] Rakich P, Reinke C, Camacho R, Davids P and Wang Z 2012 Phys. Rev. X 2 011008
- [21] Qiu W, Rakich P T, Shin H, Dong H, Soljačić M and Wang Z 2013 Opt. Express 21 31402
- [22] Casas-bedoya A, Morrison B, Pagani M, Marpaung D and Eggleton B J 2015 Opt. Lett. 40 4154–7
- [23] Shin H, Cox J A, Jarecki R, Starbuck A, Wang Z and Rakich P T 2015 Nat. Commun. 6 6427

- [24] Zoubi H and Hammerer K 2016 arXiv:1610.03355
- [25] Van Laer R, Baets R and Van Thourhout D 2016 Phys. Rev. A 93 053828
- [26] Shin H, Qiu W, Jarecki R, Cox J, Olsson R, Starbuck A, Wang Z and Rakich P 2013 Nat. Commun. 4 1944
- [27] Van Laer R, Kuyken B, Van Thourhout D and Baets R 2014 Opt. Lett. 39 1242–5
- [28] Van Laer R, Bazin A, Kuyken B, Baets R and Van Thourhout D 2015 New J. Phys. 17 115005
- [29] Kittlaus E A, Shin H and Rakich P T 2016 Nat. Photon. 10 463–7
- [30] Sipe J E and Steel M J 2016 New J. Phys. 18 045004
- [31] Rakich P and Marquardt F 2016 arXiv:1610.03012
- [32] Dhakal A, Subramanian A Z, Wuytens P, Peyskens F,
- Thomas N L and Baets R 2014 *Opt. Lett.* **39** 4025–8 [33] Claps R, Dimitropoulos D, Han Y and Jalali B 2002 *Opt. Express* **10** 1305–13
- [34] Peyskens F, Dhakal A, van Dorpe P, le Thomas N and Baets R 2016 ACS Photonics 3 102–8
- [35] Roels J, de Vlaminck I, Lagae L, Maes B, Van Thourhout D and Baets R 2009 *Nat. Nanotechnol.* 4 510–3
- [36] Li M, Pernice W and Tang H 2009 Nat. Photon. 3 464-8
- [37] Shelby R M, Levenson M D and Bayer P W 1985 *Phys. Rev.* B 31 5244
- [38] Shelby R M, Levenson M and Bayer P 1985 Phys. Rev. Lett. 54 939–42
- [39] Pant R, Poulton C G, Choi D Y, Mcfarlane H, Hile S, Li E, Thevenaz L, Luther-Davies B, Madden S J and Eggleton B J 2011 Opt. Express 19 8285–90
- [40] Aspelmeyer M, Kippenberg T and Marquardt F 2014 Rev. Mod. Phys. 86 1391–452
- [41] Zoubi H and Hammerer K 2016 Phys. Rev. A 94 1-16
- [42] Wolff C, Steel M J, Eggleton B J and Poulton C G 2015 Phys. Rev. A 92 013836
- [43] Sarabalis C J, Hill J T and Safavi-Naeini A H 2016 APL Photonics 1 071301
- [44] Van Laere F, Claes T, Schrauwen J, Scheerlinck S, Bogaerts W, Taillaert D, O'Faolain L, Van Thourhout D and Baets R 2007 IEEE Photonics Technol. Lett. 19 1919–21
- [45] Van Laer R, Kuyken B, Van Thourhout D and Baets R 2015 Nat. Photon. 9 199–203
- [46] Kittlaus E A, Otterstrom N T and Rakich P T 2016 arXiv: 1611.03556
- [47] Kharel P, Behunin R O, Renninger W H and Rakich P T 2016 Phys. Rev. A 93 063806
- [48] Wolff C, Van Laer R, Steel M J, Eggleton B J and Poulton C G 2016 New J. Phys. 18 025006
- [49] Safavi-Naeini A H, Chan J, Hill J T, Gröblacher S, Miao H, Chen Y, Aspelmeyer M and Painter O 2013 New J. Phys. 15 035007
- [50] Purdy T P, Grutter K E, Srinivasan K and Taylor J M 2016 arXiv:1605.05664
- [51] Clerk A A, Devoret M H, Girvin S M, Marquardt F and Schoelkopf R J 2010 Rev. Mod. Phys. 82 1155–208