Thermalization properties at mK temperatures of a nanoscale optomechanical resonator with acoustic-bandgap shield

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Optical measurements of a nanoscale silicon optomechanical crystal cavity with a mechanical resonance frequency of 3.6 GHz are performed at sub-kelvin temperatures. We infer optical-absorption-induced heating and damping of the mechanical resonator from measurements of phonon occupancy and motional sideband asymmetry. At the lowest probe power and lowest fridge temperature \( T_f = 10 \) mK, the localized mechanical resonance is found to couple at a rate of \( \gamma_s/2\pi = 400 \) Hz \( (Q_m = 9 \times 10^5) \) to a thermal bath of temperature \( T_s \approx 270 \) mK. These measurements indicate that silicon optomechanical crystals cooled to millikelvin temperatures should be suitable for a variety of experiments involving coherent coupling between photons and phonons at the single quanta level.

Introduction. - The coupling of a mechanical object’s motion to the electromagnetic field of a high finesse cavity forms the basis of various precision measurements [1], from large-scale gravitational wave detection [2] to microscale accelerometers [3]. Recent work utilizing both optical and microwave cavities coupled to mesoscopic mechanical resonators has shown the capability to prepare and detect such resonators close to their quantum ground-state of motion using radiation pressure back-action [5–7]. Optomechanical crystals (OMCs), in which bandgaps for both optical and mechanical waves can be introduced through patterning of a material, provide a means for strongly interacting nanomechanical resonators with near-infrared light [8]. Beyond the usual paradigm of cavity-optomechanics in which mechanical motion is confined to single objects, such as movable end-mirrors [9] and intra-cavity membranes [10], OMCs can be fashioned into planar circuits for photons and phonons, and arrays of optomechanical elements can be interconnected via optical and acoustic waveguides [11]. Such coupled OMC arrays have been proposed as a way to realize quantum optomechanical memories [12], nanomechanical circuits for continuous variable quantum information processing [13] and phononic quantum networks [14], and as a platform for engineering and studying quantum many-body physics of optomechanical meta-materials [15,17].

The realization of optomechanical systems in the quantum regime is predicated upon the ability to limit thermal noise in the mechanics while simultaneously introducing large coherent coupling between optical and mechanical degrees of freedom. In this regard, laser back-action cooling has recently been employed in simple OMC cavity systems [6,15] consisting of a one-dimensional (1D) nanobeam resonator surrounded by a two-dimensional (2D) phononic bandgap. In this work, we optically measure the properties of such an OMC cavity system in a helium dilution refrigerator down to base temperatures of \( T_f = 10 \) mK.

The device studied here is formed from the top silicon (Si) device layer of a silicon-on-insulator (SOI) wafer using a combination of electron beam lithography, plasma etching, and wet etching (see also App. B). Figure 1a
shows a scanning electron microscope (SEM) image of a suspended device after processing. This device consists of two nanobeam OMC cavities, optically coupled to a common central waveguide (Fig. 1a). As described in Ref. [9], the nanobeam cavities are patterned in such a way as to support an optical resonance in the 1550 nm wavelength band and a 'breathing' mode mechanical resonance at 3.6 GHz. The highly localized optical and acoustic resonances couple strongly via radiation pressure, with a theoretical vacuum coupling rate of $g_0/2\pi = 870$ kHz, corresponding physically to the optical resonance shift due to zero-point fluctuations of the mechanical resonator. Surrounding the waveguide and nanobeam structure on three sides is a 2D 'cross' pattern [11] which has a full phononic bandgap for all acoustic waves in the frequency range from 3 – 4 GHz. The fourth side is left open, and a final set of etches is used to clear the buried oxide and silicon handle wafer so as to allow close approach of an optical fiber. The SOI sample is mounted to the mixing chamber plate of a dilution refrigerator, and a set of position encoded 'slip-stick' stages are used to align an anti-reflection-coated tapered lensed fiber (beam waist=2.5 $\mu$m; focal distance=14 $\mu$m) to the coupling waveguide of a given device under test (see Fig. [11], and App. [C]). In order to aid efficient optical coupling, the Si waveguide is tapered down to a tip of width 225 nm, providing mode-matching between fiber and waveguide [19].

Optical spectroscopy of a fiber-coupled device at a fridge temperature of 4 K is shown in Fig. [11]. In this measurement, the frequency of a narrow linewidth external-cavity diode laser is swept across the fundamental optical resonance of one of the OMC cavities centered at $\lambda_c = 1545$ nm. The input laser light is reflected from a photonic crystal mirror at the end of the Si waveguide, and when resonant, evanescently couples in to one or the other of the nanobeam cavities (the other nanobeam cavity coupled to this waveguide has a resonance several THz to the red). This geometry allows for the extrinsic coupling rate, $\kappa_c$, to the OMC cavities to be adjusted via the waveguide-cavity gap size, and eliminates optical loss in transmission past the cavity. Some of the light entering the cavity decays through intrinsic loss channels (i.e., ones we do not detect externally) at rate $\kappa_i$, while the remainder couples back into the central waveguide and is collected in reflection by the lensed optical fiber. For this particular cavity, we observe a total optical energy decay rate of $\kappa/2\pi = 529$ MHz, an external coupling rate $\kappa_c/2\pi = 153$ MHz, and an intrinsic decay rate $\kappa_i/2\pi = 376$ MHz. From the normalized reflection signal level ($R$), the fraction of optical power reflected by the OMC cavity, collected by the lensed fiber, and detected on a photodetector is estimated to be $\eta_{pol} = 34.7\%$, which after accounting for loss in external fiber-optic components, corresponds to a single pass fiber-to-waveguide coupling of $\eta_c = 50\%$ (see App. [D] for more details).

The optical heterodyne detection scheme illustrated in Fig. [11] is used to measure the motion of the localized 'breathing' mode at frequency $\omega_m/2\pi = 3.6$ GHz, which is coupled to the fundamental optical resonance of the nanobeam. A high-power (~0.7 mW) local oscillator (LO) sets the gain of the heterodyne receiver, and a low-power (2 nW - 20 $\mu$W) optical signal beam is used to probe the OMC cavity. In order to selectively detect either the upper or lower motional sidebands generated on the optical signal beam, the LO frequency ($\omega_{LO}$) is shifted relative to that of the signal beam ($\omega_s$) using a combination of a phase modulator and tunable optical filter. For the measurements presented here, $\omega_{LO}$ is adjusted such that the mechanical modulation beat frequency $\Omega = \omega_{LO} - (\omega_s + \omega_m) \approx 2\pi \times 50$ MHz, placing a single motional sideband within the bandwidth (100 MHz) of the balanced photodetectors of the heterodyne receiver. In the case of a signal beam red-detuned from cavity resonance (frequency $\omega_s$) by $\Delta \equiv \omega_c - \omega_m = \omega_m$, the resultant noise power spectral density (NPSD) as transduced on a spectrum analyzer yields a Lorentzian component proportional to

![FIG. 2. $T = 4$ K data. a, Measured mechanical linewidth $\gamma$ for $\Delta = \omega_m$ (red) and $-\omega_m$ (blue) at a fridge temperature of $T_f = 4$ K. The vertical blue dashed line indicates the threshold $n_c$ beyond which the mechanical resonance self-oscillates. Black circles indicate the values of $\gamma$ obtained by taking the average of the detuned data. The inset shows the optomechanical damping rate $\gamma_{OM}$ determined by two methods: in circles $\gamma_{OM}$ is extracted by subtracting $\gamma$ from the red-detuned $\gamma$, and in squares $\gamma_{OM}$ is found by determining cooperativity $C$ using the calibrated $\langle n \rangle$. A linear fit (red line) yields $g_0/2\pi = 735$ kHz. b, Calibrated mechanical mode occupancy $\langle n \rangle$ versus intracavity photon number $n_c$. Blue and red circles are measured with probe laser detunings $\Delta = \mp \omega_m$, respectively. The mode occupancy $n_f$ corresponding to $T_f = 4$ K is indicated by a black solid line. c, Series of red-detuned NPSD for range of $n_c$. Here the NPSD is plotted as $S_{xx} = x_{2yy}^2 S_{hh}$, where $x_{2yy} = 4.1$ fm is the zero-point amplitude of the mechanical resonator.](image-url)
$S_{bb}(\omega) = \langle n \rangle \gamma / (\omega - \Omega)^2 + (\gamma / 2)^2)$ [20]. Here $\langle n \rangle$ is the phonon occupancy of the mechanical mode, and the total mechanical damping rate is given by $\gamma = \gamma_1 + \gamma_{OM}$, where $\gamma_1$ is the intrinsic damping rate of the mechanical resonator and $\gamma_{OM} = 4g_0^2n_c/\kappa$ is the optomechanically induced damping rate produced by an intracavity photon number $n_c$. For a blue-detuned probe ($\Delta = -\omega_m$), the NPSD Lorentzian is proportional to $n_c$ and $\gamma = \gamma_1 - \gamma_{OM}$.

Mechanical spectroscopy is first performed at a fridge temperature of $T_f = 4$ K in order to calibrate the optomechanical transduction. The coupling rate $g_0$ is determined by observing the dependence of the mechanical linewidth on $n_c$ for both red ($\Delta = \omega_m$) and blue ($\Delta = -\omega_m$) laser-cavity detunings, as shown in Fig. 2. A threshold value, $n_{thr} > n_{thr} \approx 1.5$, optical amplification and self-oscillation of the mechanical resonator occurs for blue detuning. Below this value, the optomechanical damping $\gamma_{OM}$ can be found from the difference between the red and blue detuned linewidths. A linear fit of the derived $\gamma_{OM}$ versus $n_c$ yields a coupling rate of $g_0/2\pi = 735$ kHz. The mechanical mode occupancy versus $n_c$, plotted in Figure 2a, is calibrated from the area under the Lorentzian part of the measured NPSD (see Fig. [2]) using this value of $g_0$, along with calibration of the total system efficiency, photodetector gain, and LO power. At high $n_c$ significant cooling of the mechanical mode is measured for increasing $n_c$, whereas at low $n_c$ the calibrated occupancy saturates to a constant value, indicating thermalization to the fridge temperature. For these low probe powers, the calibrated mode occupancy is larger than the 4 K fridge occupancy ($n_f$) by a factor of $\beta = 1.3$, which represents an unknown systematic error in our calibration procedure. This correction factor is applied to all subsequent mode occupancies presented below.

As the fridge temperature is lowered into the sub-kelvin range, a very different dependence of measured linewidth and mode occupancy on optical probe power is observed. In particular, the measured mechanical linewidth versus $n_c$, shown in Fig. 2b, for $T_f = 185$ mK, increases with decreasing probe power below an apparent minimum at $n_c \approx 1$. The measured linewidth at low power is also too large to explain the observed threshold of self-oscillation, $n_{thr} \approx 0.1$. These inconsistencies indicate that the linewidth associated with the true energy decay rate of the mechanics is likely obscured in the time-averaged spectrum of Fig. 2b due to frequency jitter [21]. Due to the long averaging times (minutes to hours) required at low optical probe power, direct observation of the frequency jitter in the ‘breathing’ mode is not possible. Indirect confirmation and quantification of the frequency jitter, however, is possible by studying the detuning ($\Delta$) dependence of the mode occupancy and linewidth for a fixed $n_c \approx 2$. Such a measurement keeps constant any effects such as optical heating or frequency jitter that might depend on the intra-cavity photon number. From a fit to the integrated NPSD area shown in Fig. 2b, we extract a cooperativity at detuning $\omega_m$ of $C = \gamma_{OM}(\Delta = \omega_m)/\gamma_1 = 3.9$. This value of $C$ is then used as a constraint in fitting $\gamma(\Delta)$ assuming a Voigt lineshape (Fig. 2b), where the energy damping Lorentzian linewidth $\gamma_L = \gamma_1 + \gamma_{OM}(\Delta)$ is fit with an additional random Gaussian frequency jitter term $\gamma_G$. The resulting fit (red curve) yields $\gamma_1/2\pi = 2.3$ kHz, $\gamma_G/2\pi = 6.1$ kHz, and $g_0/2\pi = 715$ kHz consistent with the $T_f = 4$ K value. In what follows we use additional on-resonance heating measurements to determine $\gamma_1$ over the entire range of $n_c$, including at the lowest optical probe powers where back action is weak.

Heating of the mechanical mode by optical absorption becomes significant at sub-kelvin temperatures due to the sharp drop in thermal conductance with temperature [22]. The source of optical absorption in our structures is most likely due to electronic defect states at the surface of Si [23, 24], with phonon-assisted relaxation from these mid-gap states heating the mechanics. This heating mechanism is investigated at $T_f = 10$ mK, for which $n_f$ is negligible, by measuring $\langle n \rangle$ and $\gamma$ using an optically resonant probe ($\Delta = 0; \gamma_{OM} = 0$). We observe in Fig. 4a that $\langle n \rangle \propto n_c^{1/4}$. This weak power law dependence is consistent with indirect coupling of
the ‘breathing’ mode to an optically-generated bath with thermal conductance scaling as $G_{th} \sim T^3$. High frequency phonons with wavelengths small relative to the dimensions of the cavity structure should have this conductance scaling \cite{26}, although their expected fast escape time ($\gamma_{THz}^{-1} \sim 1–10$ ns) suggests that they quickly come into thermal equilibrium. Given the slow rates of most bulk relaxation processes, such as anharmonic three-phonon mixing at low temperatures \cite{26}, this fast thermalization is likely due to a relatively large degree of diffusive and inelastic scattering at the surfaces of the patterned nanobeam \cite{27}.

Based on these on-resonance observations, a proposed microscopic model for the optical absorption heating and damping is illustrated in Fig. 4b. Here the long-lived ‘breathing’ mode is weakly coupled ($\gamma_0$) through the phononic shield to the exterior fridge environment, and is locally coupled via phonon-phonon scattering ($\gamma_p$) \cite{28} to the optically-generated high frequency phonons within the acoustic cavity. A phenomenological model based upon this microscopic picture is shown schematically in Fig. 4c. We parameterize the coupling of the mechanical resonator to the separate thermal baths by decomposing the mechanical damping rate into $\gamma_c = \gamma_0 + \gamma_p + \gamma_{OM}$, where the fridge bath (occupancy $n_c$) couples at rate $\gamma_0$, the optical-absorption-induced bath (temperature $T_p$, occupancy $n_p$ at $\omega_m$) couples at rate $\gamma_p$, and the intracavity laser field (effective zero-temperature bath) couples at rate $\gamma_{OM}$. The resulting average mechanical mode occupation is then given by $\langle n \rangle_{n_c} = (\gamma_0 n_c + \gamma_p(T_p)n_p(T_p))/(\gamma_0 + \gamma_p(T_p) + \gamma_{OM}(n_c))$, where $T_p(n_c)$.

The calibrated mechanical mode occupation for a red detuned ($\Delta = \omega_m$) probe is plotted against $n_c$ in Fig. 4a for $T_f = 10$ mK (purple) and 635 mK (green). Both curves exhibit a series of heating and cooling trends, and in fact coincide for $n_c \gtrsim 1$. At the lowest optical probe powers ($n_c = 0.021$) and lowest fridge temperature ($T_f = 10$ mK), the calibrated phonon occupancy reaches a minimum $\langle n \rangle = 0.98 \pm 0.11$, corresponding to $T \approx 270$ mK. The complex behavior of these two cooling curves can be understood by comparing to the proposed phenomenological model. For this model $n_f$ is taken to correspond to the measured $T_f$, $n_p(T_p)$ is ascertainment by extrapolating the on-resonance measurement of $\langle n \rangle$ (Fig. 4a)., and $\gamma_{OM}(n_c)$ is found from the fit value $g_0/2\pi = 735$ kHz to the high power region of the red detuned mechanical linewidth (red circles in Fig. 4a). Assuming a common $\gamma_p(n_c)$ and $\gamma_0$, the resulting $\gamma_c(n_c)$ curve that best fits the measured $\langle n \rangle$ data for both $T_f = 10$ mK and 635 mK fridge temperatures is plotted in Fig. 4b (blue circles). Also shown in Fig. 4b are the best fit value of the coupling to the fridge bath $\gamma_0/2\pi = 306 \pm 28$ Hz (dashed black horizontal curve) and a smooth spline curve fit to the inferred values of $\gamma_0(n_c)$ (red solid curve). A plot of the best fit model is shown alongside the measured $\langle n \rangle$ cooling curves in Fig. 4c. In addition to the good agreement of the model for both fridge temperatures, the inferred intrinsic energy damping rate at $n_c \sim 2$ is consistent with that from the above analysis of the detuning dependence of the transduced
the absolute mode occupancy and to the sum of \( \pm \) detuning \( \Delta = I_{\text{Lorentzian}} \). For an optical probe with \( \xi \) as inset to Fig. 4e. The sideband asymmetry is defined also measured the sideband asymmetry, \( \xi \) plays a significant role in heating and damping of the long range quality of the Si devices, optical absorption still in the devices of this work due to native oxide formation, TLS, common in amorphous materials, can be expected along with increasing temperature as \( T \). The asymmetry is sensitive to both the absolute mode occupancy and to the sum of \( \gamma_0 \) and \( \gamma_p \) through the cooperativity \( C = \gamma_{\text{OM}} / (\gamma_0 + \gamma_p) \). Good correspondence can also be seen between the best fit model (solid curves) and the measured \( \xi \) (circles).

Although significant work remains to determine the exact microscopic details of the optical absorption heating, thermalization properties, and frequency jitter observed in the measurements of the quasi-1D OMC cavity studied here, there are nonetheless several interesting points that can already be noted. First, from the measured time-averaged mechanical linewidth and optically-induced bath temperature \( T_p \) in Fig. 4 we find that the frequency noise of the mechanical resonator drops with increasing temperature as \( T_p^{-0.9} \) (see App. C for details). Such a frequency noise behavior is similar to that found for two level systems (TLS) coupled to superconducting microwave resonators \( \xi \). The presence of TLS, common in amorphous materials, can be expected in the devices of this work due to native oxide formation at the Si surfaces. Secondly, despite the very high optical quality of the Si devices, optical absorption still plays a significant role in heating and damping of the long range ‘breathing’ mode at sub-kelvin temperatures. Here, the phononic shield provides excellent mechanical isolation of the ‘breathing’ mode, yet still provides good mechanical coupling to the fridge bath for heat carrying phonons above the acoustic bandgap. Thirdly, although lower phonon occupancies could have been measured using thinner phononic shields, effectively increasing the coupling rate \( \gamma_0 \) to the fridge bath at \( T \), this would come with a commensurate reduction in cooperativity \( C = \gamma_{\text{OM}} / \gamma_0 \). Ultimately, coherent quantum interactions between the optical cavity field and the mechanical resonator require \( C > 1 \) and \( \langle n \rangle < 1 \). The devices of this work closely approach this limit, however, quasi-2D Si OMC devices such as those recently realized in Ref. \( \xi \) should have orders of magnitude larger thermal conductance, enabling coherent coupling between light and mechanics in the quantum regime as envisioned in recent proposals \( \xi \).

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The geometry of the optomechanical crystal (OMC) studied in this work is numerically optimized for optical and mechanical quality, as well as optomechanical coupling, via finite-element method (FEM) simulation in COMSOL Multiphysics [32]. In Fig. 7a, a top view of the OMC shows the center defect in the 600 nm wide by 220 nm thick Si nanobeam. The larger holes on the ends of the nanobeam support simultaneous bandgaps for 1550 nm band light and 3 µm thick Si nanobeam. The smaller holes in the center of the nanobeam perturb the bandgaps such that optical and mechanical modes are co-localized [33]. The fundamental optical mode has a nominal wavelength of 1535 nm (Fig. 5b), and the acoustic breathing mode has a nominal resonance frequency of 3.6 GHz, and $g_0 = 735$ kHz, as discussed in the main text.

Appendix B: Fabrication

The devices are fabricated from a silicon-on-insulator (SOI) wafer (SOITEC, 220 nm device layer, 3 µm buried oxide) using electron beam lithography followed by reactive ion etching (RIE/ICP). The Si device layer is then masked using ProTEK PSB photoresist to define a mesa region of the chip to which a tapered lensed fiber can access. Outside of the protected mesa region, the protected region is removed with a plasma etch and a trench is formed in the underlying silicon substrate using tetramethylammonium hydroxide (TMAH). The devices are then released in hydrofluoric acid (49% aqueous HF solution) and cleaned in a piranha solution (3-to-1 H$_2$SO$_4$:H$_2$O$_2$) before a final hydrogen termination in diluted HF.

Appendix C: Experimental setup

The full experimental setup for heterodyne spectroscopy and mechanical thermometry is shown in Fig. 8. A fiber-coupled, wavelength-tunable external cavity diode laser is used as the light source, and a small percentage of the laser output is sent to a wavemeter ($\lambda$-meter) for frequency stabilization. The remaining laser power is split into a high-power ($\approx 0.7$ mW) local oscillator (LO) path and a low-power ($\approx 20$ µW) signal path. The signal beam is sent through an electro-optic modulator ($a$-$m$) to stabilize the signal intensity and a variable optical attenuator to allow control of the probe power sent to the cavity. The signal is sent into an optical circulator which directs the signal beam to the dilution refrigerator in which the fiber terminates with a lensed tip for end-fire coupling to the device. The cavity reflection then circulates to a variable coupler (VC) and mixes with the LO before being detected on a pair of balanced photodiodes (BPD). The difference photocurrent is then amplified and its noise power spectral density (NPSD) is measured on a real-time spectrum analyzer (RSA). The LO is first sent through an electro-optic phase modulator ($\phi$-$m$) which produces optical sidebands at $\pm (\omega_m/2\pi - 50$ MHz) for the purpose of mixing the mechanically induced signal modulation down to within the 100 MHz bandwidth of the BPD circuit. An erbium-

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Fig. 5. **Optical and Mechanical FEM Simulations.**

a, Silicon OMC geometry. 

b, Optical mode FEM simulation, showing the electric field $E_y$ component (polarization in the plane of the page and transverse to the long axis of the nanobeam). 

c, FEM simulation of the localized ‘breathing’ mode at frequency $\omega_m/2\pi = 3.6$ GHz. Here the displacement mode profile is shown as a distortion of the structure and as a color-coded local displacement amplitude.

Fig. 6. **Experimental setup.**

a, Fiber-based optical heterodyne setup. $\lambda$-meter: wavemeter, FPC: fiber polarization controller, a-m: electro-optic amplitude modulator, $\varphi$-m: electro-optic phase modulator, VOA: variable optical attenuator, SW: optical switch, EDFA: erbium-doped fiber amplifier, PM: optical power meter, VC: variable coupler, BPD: balanced photodiode pair, RSA: real-time spectrum analyzer. 

b, SEM image of the OMC device, with acoustic radiation shield and end-fire fiber coupling indicated. 

c, Zoom in SEM image showing details of the waveguide–cavity coupling region, indicating the extrinsic cavity-waveguide coupling rate $\kappa_e$ and the intrinsic loss rate $\kappa_i$. 

d, Edge view SEM image of the central waveguide tip. 

e, Normalized optical cavity reflection spectrum ($R$). Detunings from resonance of $\Delta = \pm \omega_m/2\pi = \pm 3.6$ GHz are denoted by the red and blue arrows, respectively.

doped fiber amplifier (EDFA) and a variable optical attenuator (VOA) are used to set the power of the LO sidebands, and the appropriate sideband is selected by a high-finesse tunable Fabry-Pérot filter before recombining it with the signal.

Fig. 6b shows a scanning electron micrograph (SEM) of a typical device, consisting of a central Si waveguide side-coupled to two nanobeam OMCs, all of which is surrounded by an acoustic radiation shield phononic crystal. Light coupled into the central waveguide either reflects from a photonic crystal mirror or, when resonant, evanescently couples to one of two side-coupled OMCs [6, 34], where the coupling rate $\kappa_e$ is controlled by the design of the gap size between the waveguide and the cavity (Fig. 6c). Some cavity light decays through parasitic scattering at a rate $\kappa_i$, while the remainder couples back into the central waveguide and is collected from the central waveguide tip (Fig. 6d) by the lensed fiber to...
be guided to the photodetector in Fig. 6a. By tuning
the laser over the cavity mode, we observe in the re-
lected signal on the output of the dilution refrigerator
(Fig. 6a) a fiber collection efficiency of \( \eta_{\text{cpl}} = 34.7\% \) and
an optical linewidth of \( \kappa/2\pi = 529 \text{ MHz} \), composed of
\( \kappa_e/2\pi = 153 \text{ MHz} \) and \( \kappa_i/2\pi = 376 \text{ MHz} \).

Appendix D: Fiber coupling in the dilution
refrigerator

The microchip sample is mounted to the mixing cham-
ber of the dilution refrigerator, and we utilize an end-
fiber coupling scheme to probe individual devices with an
anti-reflection-coated tapered lensed fiber. The lensed
fiber tip is clamped down on a position encoded piezo
xyz-stage inside the dilution refrigerator (Fig. 7a), which
allows nanopositioning with respect to the sample. When
mounting the fiber and sample, the fiber is only roughly
aligned to within a few millimeters (Fig. 7b). After cool-
ing the experiment from room temperature to 4 K, we
monitor the reflected optical power on a slow photode-
tector as we carefully lower the fiber tip to match the
height of the device layer. The distinct reflection of the
device layer allows us to iteratively adjust the fiber posi-
tion and optimize the coupling to each device (Fig. 7c).

The fiber-tip launches the light to free-space and fo-
cuses it to a beam waist of 2.5 \( \mu \text{m} \) at a focal distance
of 14 \( \mu \text{m} \). We position the fiber such that the beam
waist aligns to a silicon waveguide tip (Fig. 7d) that
matches the convergence of the optical field, which be-
comes a guided mode in the waveguide [19, 35, 36].

The design of the tapered waveguide coupler is simi-
lar to that presented in Ref. 36, where the tip of the
waveguide is mode matched to an input Gaussian field
of the appropriate width and adiabatically tapered up to
the full width of the photonic crystal mirror section. The
major distinction is the use of end-fire coupling utilizing
a lensed fiber rather than butt-coupling with a cleaved
single-mode fiber. In contrast to the high-stress Si\(_3\)N\(_4\)
utilized in Ref. 36 a Si waveguide will begin to sag if
the waveguide is made too long, even in the presence
of a supporting tether, which will lead to misalignment
of the waveguide tip and bending losses that inhibit the
coupling efficiency. As such, it is necessary to substan-
tially reduce the length of the waveguide taper, which
necessitates a smaller overall change in waveguide width
to maintain the adiabaticity of the taper. Consequently
we are forced to use a larger waveguide width at the tip
and a correspondingly smaller mode field diameter such
as can be obtained with a lensed fiber.

The broadband reflection spectrum of the coupler,
calculated using finite-difference-time-domain simula-
tion [37], is shown for an ideal coupler in Fig. 8a. The
presence of fringes in the reflection spectrum are con-
sistent with the estimated weak reflectivity of the
waveguide–air interface (\( R \approx 0.5\% \)), which forms a low-
finesse Fabry-Pérot cavity with the high reflectivity pho-
tonic crystal mirror at the other end of the waveguide.
From the fringe visibility we can back out the expected
single-pass coupling efficiency \( \eta_{\text{cpl}} \) shown in Fig. 8b [36].

The calculated reflection spectrum and single-pass effi-
ciency for the device used in this work (using dimensions
determined from SEM images) is also shown in Fig. 8b
and is expected to be \( \eta_{\text{cpl}} \approx 60\% \) at the cavity resonance.
The actual single-pass efficiency of the measured device
is found to be \( \eta_{\text{cpl}} \approx 50\% \). The difference from simula-
tions is attributed to the difficulty of measuring the exact
dimensions of the sample using the SEM, as a small dif-
fERENCE of even 10 nm in the width of the waveguide tip
can have a significant effect on the mode matching and
thus on the overall coupling efficiency.
Appendix E: Calibration of the optical transduction of mechanical motion

To perform accurate thermometry of the mechanical mode, it is necessary to calibrate the detection efficiency of the setup. We first measure the efficiency of transmission in the circulator from port 1 to port 2 ($\eta_{12} = 88\%$), and from port 2 to port 3 ($\eta_{23} = 84\%$). These values are measured once when the optical components are connected and do not change. These calibrations are used to determine the reflection efficiency of the device and the overall detection efficiency of the heterodyne setup.

To measure device efficiency, the laser is tuned off-resonance from the optical mode (where the device should act as a near-perfect mirror) and a continuous-wave signal of input power $P_{in}$ is sent into port 1 of the circulator, leading to a power $\eta_{12}P_{in}$ exiting port 2 of the circulator. The optical losses incurred in the path from port 2 to the device-under-test are accumulated into an efficiency factor $\eta_{cpl}$, which includes signal loss in the fiber path through the fridge, mode-mismatch between the lensed-fiber to tapered-waveguide tip, and mode-scattering from the waveguide tip to the photonic-crystal mirror (Fig. 7). These losses are incurred twice in reflection back to the circulator, so a power of $\eta_{cpl}^2\eta_{12}P_{in}$ propagates into port 2 and $\eta_{23}\eta_{cpl}^2\eta_{12}P_{in}$ propagates out of port 3 of the circulator. An optical switch (SW3) is used to send this signal to a power meter (PM), and thus the coupling efficiency is determined as

$$\eta_{cpl} = \sqrt{\frac{P_{PM}}{\eta_{23}\eta_{12}P_{in}}} = 34\%. \quad (E1)$$

To calibrate the overall detection efficiency, we must also determine the efficiency of the heterodyne detector itself, which includes the intrinsic quantum efficiency of the BPD, the alignment of the polarization between the LO and the signal, and the degree to which the LO power overcomes the electronic noise of the detector. This is accomplished by using the amplitude modulator to create optical sidebands detuned from the signal by the mechanical frequency while the laser is tuned off-resonance from the optical mode. The optical switches SW1 and SW2 are used to route the signal through a tunable filter to select a single sideband which is sent through the device and onto the BPD. The power $P_{cal}$ in this sideband is directly measured on the PM at SW3, and the photocurrent NPSD ($S_{II}(\omega)$) as transduced on the RSA is given by

$$S_{II}(\omega) = S_{dark} + \frac{G_e^2 \bar{s}_{SN}^2}{P_L} \left( 1 + \frac{\eta_{VC}\eta_{det}S_{cal}(\omega)}{\hbar\omega_o} \right), \quad (E2)$$

where $S_{dark}(\omega)$ is the electronic NPSD of the detector, $S_{SN} = \sqrt{2\hbar\omega_o P_{LO}}$ is the optical shot-noise NPSD arising from $P_{LO}$ of LO optical power at optical frequency $\omega_o$, which lies an order of magnitude above the electronic noise, and $S_{cal}$ is the NPSD of the signal, where $\int_{-\infty}^{\infty} S_{cal}(\omega) d\omega = P_{cal}$. The gain factor $G_e$ represents the conversion from optical power to voltage while $R_L$ is the input impedance of the RSA. The total noise floor $S_{noise} = G_e^2 \bar{s}_{SN}^2 + S_{dark}$ is measured with the signal beam blocked, while $S_{dark}$ is measured independently with both signal and LO beams blocked. When referenced back to the PM, the calibration tone (with NPSD $S_{cal}(\omega)$) picks up losses in the VC and BPD, as parametrized in $\eta_{VC}$ and $\eta_{det}$, respectively. The efficiency of the heterodyne receiver is extracted as

$$\eta_{VC}\eta_{det} = \frac{\hbar\omega_o}{P_{cal}} \int_{-\infty}^{\infty} \frac{S_{II}(\omega) - S_{noise} d\omega}{S_{noise} - S_{dark}} \frac{d\omega}{2\pi} = 56\%. \quad (E3)$$
This, combined with the measured device coupling efficiency, yields the overall measurement efficiency \( \eta \) used for calibrated mechanical thermometry as

\[
\eta = \eta_{\text{cpl}} \eta_{\text{2a}} \eta_{\text{CV}} \eta_{\text{det}} = 16\%.
\]  

(E4)

Appendix F: Heating and damping via three-phonon scattering processes

Though a detailed microscopic calculation of the additional heating and damping due to optical absorption is beyond the scope of this work, some qualitative insight into the nature of the locally heated mechanical bath and its coupling to the mechanical mode of interest can be obtained from consideration of a simplified model of the phonon-phonon interactions. At the temperatures considered in this work (\( T < 10 \) K) the mean free path of the thermal phonons is expected to be much larger than the wavelength of the mechanical mode of interest. Consequently, the damping should be described by the Landau-Rumer theory, where losses occur primarily due to three-phonon mixing with the local thermal environment due to anharmonicity in the Si lattice \([22, 38]\).

Consider first a toy model where two high-frequency modes, with frequencies \( \omega_1 \) and \( \omega_2 \) respectively, are coupled with the mode of interest at frequency \( \omega_m (\omega_1 - \omega_2 = \omega_m) \). To first order in perturbation theory, the scattering rates into and out of the mechanical cavity mode due to the lowest order anharmonic interaction can be given by \([38]\) \( \Gamma_\pm = A(n_m + 1)(n_2 + 1)n_1 \) and \( \Gamma_e = A(n_1 + 1)n_m n_2 \), where \( A \) is a constant which depends on the matrix element of the anharmonic potential, and \( n_1, n_2 \) and \( n_m \) are the number of quanta in each of the three mechanical modes. Thus, in the absence of other dissipative processes a simple rate equation for the population of the mechanical cavity mode is given by

\[
\dot{n}_m = \Gamma_+ - \Gamma_e = -A(n_2 - n_1)n_m + A(n_2 + 1)n_1, \tag{F1}
\]

which has the same form as the equation for a harmonic oscillator interacting with a bath of occupation \( n_p \) with a coupling rate \( \gamma_p \), where

\[
n_p = \frac{n_2(n_1 + 1)}{n_2 - n_1}, \quad \gamma_p = A(n_2 - n_1). \tag{F2}
\]

If the two high-frequency modes are both in equilibrium with each other at some elevated temperature \( T_p \), such that \( n_1 = (\exp(\frac{\hbar \omega_1}{k_B T_p}) - 1)^{-1} \), it is easy to show that \( n_p \) is simply given by the Bose-Einstein occupation factor for the mechanical mode at temperature \( T_p \)

\[
(n_p = (\exp(\frac{\hbar \omega_m}{k_B T_p}) - 1)^{-1}). \tag{F3}
\]

We can also see that the scattering rate \( \gamma_p \) will depend on \( T_p \) through the temperature dependence of the population difference \( n_2 - n_1 \). This dependence can be approximately linear or exponential in \( T_p \), depending on the value of \( (\hbar \omega_2)/(k_B T_p) \).

More realistically, the optical absorption process will populate a range of high-frequency phonon modes above some cutoff frequency \( (\omega_c) \), which can contribute to the heating. On the assumption that these modes come into equilibrium with each other at some elevated temperature \( T_p \), we can easily show that the expression for \( n_p \) is unchanged, and the effective bath occupancy \( n_p \) will be given by the above expression. The scattering rate will now be given by

\[
\gamma_p = \int_{\omega_c}^{\infty} d\omega \rho(\omega) A(\omega, \omega_m)(n(\omega, T_p) - n(\omega + \omega_m, T_p), \tag{F4}
\]

where \( \rho(\omega) \) is the density of modes at frequency \( \omega \), the matrix element \( A \) now can depend explicitly on frequency, and \( n(\omega, T) \) is just the Bose-Einstein occupation at frequency \( \omega \) and temperature \( T \). In general, we can obtain limited analytical results if we assume that the product of the density of states and the matrix element obeys some power law as a function of frequency, \( \rho(\omega)A(\omega, \omega_m) \propto \omega^a \). For example, under a simple continuum elastic model the product would be given by \( \rho(\omega)A(\omega, \omega_m) \propto \omega_m(\omega_1 + \omega_m)^b \) \([35]\). Making a simple change of variables \( x = \frac{\hbar \omega_c}{k_B T_p} \), and assuming \( \omega_m \ll \omega_c \), we then arrive at the approximate relation

\[
\gamma_p \propto \omega_m T_p^{a+1} \int_{x_c}^{\infty} dx \frac{x^a e^x}{(e^x - 1)^2}; \quad x_c = \frac{\hbar \omega_c}{k_B T_p} \tag{F5}
\]

Considering first the limiting case \( x_c \gg 1 \), we note that the integral can be approximated by

\[
\int_{x_c}^{\infty} dx \frac{x^a e^x}{(e^x - 1)^2} \approx \int_{x_c}^{\infty} dx \frac{x^a}{e^x} = \Gamma(a + 1, x_c) \tag{F6}
\]

where \( \Gamma(a, z) \) is the upper incomplete Gamma function. For real values of \( z \), this function has the asymptotic behavior \( \Gamma(a, z) \rightarrow z^{a-1} e^{-z} \) as \( |z| \rightarrow \infty \). This leads to the approximate scaling law

\[
\gamma_p \propto T_p^{a+1} x_c^{a-1} e^{-x_c} = T_p \rho(\omega_c) e^{-\frac{\hbar \omega_c}{k_B T_p}}, \tag{F7}
\]

for \( T_p \ll \frac{\hbar \omega_c}{k_B} \). In the limiting case \( x_c \ll 1 \), the lower limit of the integral can be extended to 0, and a simple integration by parts shows that, for \( a > 1 \),

\[
\int_{0}^{\infty} dx \frac{x^a e^x}{(e^x - 1)^2} = a \int_{0}^{\infty} dx \frac{x^{a-1}}{e^x - 1} = a \Gamma(a) \zeta(a), \tag{F8}
\]

where \( \Gamma(a) \) is the gamma function and \( \zeta(a) \) is the Riemann zeta function. As this is a simple constant, we get the scaling law \( \gamma_p \propto T_p^{a+1} \), when \( T_p \gg \frac{\hbar \omega_c}{k_B} \) and \( a > 1 \).
In Figure 9b the low temperature end of the inferred $\gamma_p(T_p)$ data is seen to fit well to a curve $\propto T_p \exp (-T_c/T_p)$, where the cut-off frequency is $T_c \approx 2$ K ($\omega_c/2\pi \approx 35$ GHz). Although this simple model does not capture all of the features of the measured $\gamma_p(T_p)$ curve (in particular, the kink at $T_p \approx 3$ K), it does show that the damping rate due to 3-phonon mixing can vary substantially as a function of $T_p$, particularly at low temperatures in the vicinity of $T_p \lesssim \frac{\hbar \omega_c}{k_B}$.

**Appendix G: Temperature dependence of frequency noise**

At sub-kelvin fridge temperatures, frequency noise of the 'breathing' mode resonance is seen to dominate the time-averaged measured linewidth for low optical probe powers (Fig. 9a). In the case of the red-detuned probe, frequency noise is observed for $n_c < 1$ before optomechanical damping kicks in, while the on-resonance measurement shows frequency noise for $n_c < 10$ before the intrinsic energy damping ($\gamma_p$) begins to dominate. Combining the sections of both data sets which are dominated by frequency noise, a power law fit shows linewidth scaling as $n_c^{-0.23}$. By comparing to the on-resonance measured mode occupancy (Fig. 4a in main text), the frequency noise can also be plotted with respect to the optical absorption driven temperature $T_p$, as shown in Fig. 9a. A power law fit of the time-averaged measured linewidth in the frequency jitter dominated regime, shows a frequency jitter scaling $\propto T_p^{-0.9}$.

Similar inverse temperature scaling of frequency noise has been observed in superconducting microwave resonators [30], with substantial evidence indicating the source to be fluctuations in two-level tunneling states (TLS) of near-field amorphous materials [39, 40]. These TLS also couple to phonons, contributing to not only the dielectric properties of the material, but also the elastic properties. As such, the microwave mechanical modes of the devices studied here can be expected to couple to TLS in a similar fashion as for the microwave electromagnetic resonators. The decrease in frequency noise with temperature can then be attributed to the thermal excitation and saturation of the TLS. The presence of TLS in the Si devices of this work likely stems from the formation of a native oxide on the Si surfaces of the patterned nanobeam. Careful measures, involving a final wet etch in HF acid [24], were made to passivate the Si surfaces of the nanobeam in order to reduce the optical absorption from sub-bandgap surface electronic states. This surface cleaning should have been effective in removing the surface oxide as well, however, a one hour procedure was required to load the sample into the dilution refrigerator, during which time a native oxide would at least partially reform on the Si surfaces of the device. This native oxide is likely to be the source of the TLS. Future work will look to substantially reduce the device loading time after HF surface cleaning, hopefully reducing both the optical absorption and TLS density.