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Optical bistability with a repulsive optical force in coupled silicon photonic crystal membranes

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We demonstrate actuation of a silicon photonic crystal membrane with a repulsive optical gradient force. The extent of the static actuation is extracted by examining the optical bistability as a combination of the optomechanical, thermo-optic, and photo-thermo-mechanical effects using coupled-mode theory. Device behavior is dominated by a repulsive optical force which results in displacements of \( \approx 1 \text{ nm/mW} \). By employing an extended guided resonance which effectively eliminates multi-photon thermal and electronic nonlinearities, our silicon-based device provides a simple, non-intrusive solution to extending the actuation range of micro-electromechanical devices. 

Rapid developments in the field of optomechanics have opened up avenues for fundamental research on quantum state manipulation with macroscopic structures and show promise for optomechanical sensors and technologies for both radio-frequency and telecom applications. While most attention has been devoted to compact structures featuring low (picogram) mass and ultrahigh-frequency (gigahertz) mechanical modes, the technological implications of static deformation due to optical forces have been less explored.

In coupled photonic waveguide geometries, bonding and anti-bonding optical modes are supported and the corresponding attractive and repulsive optical forces exerted on a pliant structure (low mechanical frequency) could serve to broaden the range of motion of integrated microelectromechanical devices. This translates to improvement in the detection range of pressure and displacement sensors and the actuation range of electrostatic actuators. In particular, the pull-in limit of electrostatic actuators could be extended by increasing the plate separation with a repulsive optical force. Additionally, schemes for preventing stiction, which occurs when attractive forces like the Casimir force and electrostatic force become overwhelming large compared to the mechanical restoring force, have been proposed using a real-time monitoring of the structure’s displacement and a counteracting feedback repulsive force (of the order of nano-Newton and linear with excitation power). In this paper, we demonstrate nanometer-pulling of a thin silicon photonic crystal (PhC) membrane under low vacuum with a repulsive optical gradient force and an attractive photo-thermo-mechanical force. Furthermore, optical bistability induced by optical forces and thermo-optic effect is observed at large excitation powers.

Our devices, illustrated in Fig. 1(a), consist of a square silicon PhC slab suspended by four support arms \( \approx 250 \text{ nm} \) above a Silicon-on-Insulator (SOI) substrate. They are fabricated (Fig. 1(b)) from a double-SOI platform, formed by oxide-oxide bonding of two thermally oxidized SOI wafers. A sacrificial silicon dioxide layer between the two silicon layers is \( s_0 = 265 \text{ nm} \) thick. Electron-beam lithography is performed on a layer of resist (ZEP-520A) to define the pattern. To combat the strong buckling of the silicon device layer by the compressive stress and upward turning moments of the oxide layer underneath, stress management techniques were incorporated to obtain structures with lithographically determined membrane-substrate gaps. After developing, a fluorine-based reactive-ion etch is employed to transfer the patterns to the top silicon layer. The device is then released by undercutting the patterned silicon layer with the vapor-phase hydrofluoric acid etch. Finally, an annealing step was performed at 500 °C for 1 hour in a nitrogen environment to limit surface losses and maximize optical and mechanical quality factors. The separations of the released membranes from the substrate are characterized by a confocal microscope (Olympus LEXT OLS-4000).

The structure was designed to support an optical anti-bonding mode in the wavelength range of 1480–1680 nm, which results from the hybridization of waveguide modes in the membrane and substrate. The precise spectral location of the resonance is determined by the optomechanical coupling between the two modes, the strength of which is defined as \( g_{OM} \equiv d\omega_0/ds \), where \( \omega_0 \) is the optical resonant frequency. The distribution of the x-component of the electric field in the top membrane is out-of-phase from that in the bottom membrane, as depicted in the simulation results of the whole structure in Fig. 1(c), which corresponds to the generation of a repulsive gradient force. Additionally, the field symmetries along the \( x-z \) and \( y-z \) planes indicate that we are operating with a “dark” mode, which theoretically does not couple to normally incident light because of mismatch in field symmetry. However, by breaking the periodicity of the
full structure, we can couple to the dark mode and achieve high $Q_{\text{opt}}$. Such devices have been the subject of numerous theoretical and experimental investigations on subjects ranging from the lowering of the laser thresholds to increasing the sensitivity of photonic-crystal-based sensors. Here, the dark mode is made accessible due to the finite size of the membrane and slight fabrication imperfection. The high $Q_{\text{opt}}$ of the dark mode, together with the mode’s large optomechanical coupling coefficient $g_{\text{OM}}/2\pi = -23$ GHz/nm (at $s_0 = 220.6$ nm), boosts the strength of the optical force and hence the range of actuation. A low power (25 $\mu$W) wavelength sweep is shown in the inset of Fig. 1(d), revealing a cavity resonance centered at $\lambda_0 = 1581.55$ nm. To account for interference fringes from parasitic reflections, we carefully fit the optical resonance and the oscillating background (black line) to an expression which has the form:

$$R = |r|^2 = \left| r_d(\lambda)e^{-i\phi} + \frac{K_c}{i\Delta_0 + \kappa/2} \right|^2,$$  \hspace{1cm} (1)

where $r_d(\lambda)$ is the background reflectivity, $\phi$ is the relative phase between the underlying background reflection and the optical cavity, $\kappa$ is the full-width half-max linewidth of the optical resonance, $K_c$ is the external coupling rate, $\Delta_0$ is the detuning (expressed in frequency) of the sweep wavelength from $\lambda_0$, Fitting parameters correspond to an optical cavity with $K_c = 0.3 \kappa$ and $Q_{\text{opt}} = 3400$.

As previously described, the potential of a mechanical harmonic oscillator with equilibrium position $s_0$ when perturbed by the potential of an optical “spring” centered at $s_1$ for a laser wavelength $\lambda_i$ can create a multi-well potential with two stable mechanical equilibria. The transition between these mechanical equilibria is reflected by the occurrence of optical bistability, due to the dependence of the resonance frequency on $s$. Yet the direct observation of the optomechanically induced optical bistability can easily be obscured in actual systems by other competing mechanisms including the thermo-optic effect due to two-photon absorption, free-carrier dispersion, and the Kerr nonlinearity. We designed our geometry to minimize these effects by exciting a guided resonance which is delocalized throughout the PhC membrane. We estimate the total mode volume to be $\approx 260(\lambda/n_g)^3$ from simulation. Due to its large modal volume, the thermal and electronic nonlinearities (which scale inversely with the modal volume) are dramatically reduced. This is in contrast with many of the optomechanical structures being studied, which have modal volumes $\approx (\lambda/n_g)^3$ and where thermal nonlinearities could be readily observed at even modest input
powers. Here, with the coupled PhC membrane of the current separations, optomechanical detuning is larger than thermo-optic detuning that originates from linear absorption due to defects introduced during the fabrication processes, which is two orders of magnitude larger than the intrinsic material absorption of bulk silicon.

We solve for the optical and mechanical equilibria in the presence of the thermo-optic effect within the coupled-mode theory framework. In particular, the stored optical energy in the system $|a|^2$ is given by

$$|a|^2 = \frac{\kappa_c}{(\kappa/2)^2 + \Delta^2} P_{in},$$

(2)

where $P_{in}$ is the power incident on the structure, and $\kappa_c/\kappa$ represents the fraction of incident power coupled into the cavity. The detuning $\Delta$ of the laser excitation frequency $\omega_l$ from the perturbed optical resonant frequency can be written as

$$\Delta = \omega_l - \left[\omega_0 + (d\omega_l/dT)\Delta T + g_{OM}\Delta x\right].$$

(3)

The third term in Eq. (3) is the thermo-optic detuning, with $d\omega_l/dT = (dn/dn)(dn/dT)$, $n$ is the refractive index of silicon, $dn/dn$ is obtained from simulations and approximately $-2\pi \times 10^{14}$ Hz, and $dn/dT$ is the thermo-optic coefficient of silicon equal to $2 \times 10^{-4}$ K$^{-1}$. The absorbed optical power and hence the temperature change of the system is given by

$$\Delta T = \frac{\Gamma_{abs}|a|^2}{C_h\kappa_i},$$

(4)

where $\Gamma_{abs}$ is the absorption coefficient of the system, $C_h$ is the heat capacity, and $\kappa_i$ is the thermal diffusion rate. The fourth term in Eq. (3) is the optomechanical detuning. In particular, the displacement of the membrane due to the respective photo-thermo-mechanical force and the repulsive gradient force is given by

$$\Delta x = \frac{DA}{K} + \frac{|a|^2 g_{OM}}{\omega_0 K},$$

(5)

where $K$ is the spring constant of the mechanical resonator and $D$ is the thermal-mechanical force coefficient in units of Newtons per Kelvin. We neglect the Duffing nonlinearity in our mechanical model as the extent of the optical actuation is still well within the linear regime for our structures: the amplitude is much less than the membrane thickness (185 nm) and the compressive stress in the silicon device layer is alleviated by thin accordion structures as shown in Fig. 1(b). The above equations can be solved self-consistently to yield $\Delta$ and hence the perturbed optical resonant frequency $\omega_0 = \omega_0 + (d\omega_l/dT)\Delta T + g_{OM}\Delta x$ at a given $\omega_l$ and $P_{in}$.

The values of $\omega_0 = 2\pi/\lambda_0$ at which solutions of Eq. (3) exist are plotted in Fig. 2, as a function of laser wavelength $\lambda_l = 2\pi/\omega_l$ for incident powers of 0.275 mW (green line), 0.775 mW (blue line), 1.275 mW (red line), 1.525 mW (purple line), 1.775 mW (orange line), and 2.275 mW (black line). The unperturbed optical resonance occurs at $\lambda_0 = 1581.55$ nm. The dashed portions of the curves correspond to unstable equilibria. At high powers, a clear bistable region exists in which there are two stable configurations of the membrane for fixed power and laser wavelength, due to both optomechanical and thermo-optic detunings whose magnitudes are comparable. The boundaries of the bistable region are denoted by $\lambda_f$ and $\lambda_b$, representing the hysteretic transition wavelengths for a laser swept forward (left to right) and backward (right to left) across the resonance.

We can model the reflectance of the system as a function of laser wavelength at multiple powers ($P = 0.275$ mW to 2.275 mW—same powers as in Fig. 2) with Eq. (1) (with $\Delta_0$ replaced by $\Delta_0$) and the respective equilibria calculated in Fig. 2. The results are shown in Fig. 3(a) (offset for clarity), and are compared to our experimental observations shown in Fig. 3(b). The experimental data were collected by sweeping the tunable laser output from short to long wavelength (red curve) and then back (blue curve) at a fixed tuning speed of 1 nm/s. We find excellent agreement between experiment and theory, which displays an overall redshift of the mode and increasing hysteresis at higher powers. In particular, we directly compare the locations of the forward and backward bistable jumps, $\lambda_f$ and $\lambda_b$, respectively, in Fig. 3(c). The locations of these transitions were extracted from the data shown in Fig. 3(b) by finding the minima of the reflectivities of forward and backward wavelength sweeps at each power. We see strong agreement between experiment (red/blue circles) and theoretical predictions from Eq. (3) (red/blue line) on the locations of $\lambda_f$ and $\lambda_b$.
rupt hop back to the pulled initial equilibrium position. The transition wavelength during the forward sweep \( \lambda_b \) (red circles) is linear and matches well to calculations from Eq. (3) (red line). The backward transition wavelength \( \lambda_a \) (blue circles) also show good agreement in the transition wavelength locations and the onset power for hysteresis.

Alternately, we can investigate the range of actuation of the optical force by sweeping the laser power up and down at fixed wavelengths slightly red-detuned from the unperturbed cavity resonance. Theoretical predictions and experimental results are plotted in Figs. 4(a) and 4(b), respectively, showing the reflected power plotted against the incident laser power at nine red-detuned wavelengths: 1581.9 nm (magenta line), 1581.95 nm (violet line), 1582.0 nm (green line), 1581.9 nm (magenta line), 1581.95 nm (violet line), 1582.05 nm (red line). Solid lines represent the power output as a function of increasing laser power, while dashed lines represent power output as a function of decreasing input power. Modeling predicts hysteresis will occur at all wavelengths longer than 1581.95 nm.

When we decompose the perturbation to the optical resonance into its constituent parts, we find optomechanically induced bistability to be the dominant effect. For example, at \( P_{in} = 2.275 \text{ mW} \), optomechanical effects correspond to a peak resonance shift \( \Delta \lambda_{OM} = 0.44 \text{ nm} \), while thermo-optic contributions lead to \( \Delta \lambda_{PT} = 0.23 \text{ nm} \) and photo-thermal-mechanical contributions lead to \( \Delta \lambda_{PTM} = -0.01 \text{ nm} \). This corresponds to a membrane which is mechanically pushed upward 2.3 nm by the optical gradient force and 0.1 nm downward by the photo-thermal-mechanical force, resulting in a net upward displacement of 2.2 nm. These results hold promise for large actuation range with repulsive optical forces by designing membranes which are less mechanically stiff and generate larger repulsive forces per mW of incident optical power by increasing \( Q_{opt} \). For instance, \( Q_{opt} \) is currently limited by fabrication imperfections and the finite size effect of the PhC and could be boosted by simply increasing the number of unit cells in the membrane.16 To maintain the same compactness of the structure which is related to its dynamic range, the optical design could be modified with a smaller lattice constant and/or graded hole modulation.23

In conclusion, we demonstrated actuation of a micron-scale membrane with a repulsive optical force using an extended guided resonance in a coupled silicon PhC membrane. The net red-shift displayed in the optical resonance of our doubly bonded SOI platform is a result of an optomechanically induced red-shift, a thermo-optic red-shift, and a photo-thermo-mechanically induced blue-shift. Furthermore, simulations indicate that absorption in our system is dominated by surface defects and adsorbents, resulting in a linear absorption coefficient two orders of magnitude larger than
that expected from bulk silicon. By minimizing these effects through fabrication process and design modifications, we can further isolate and exploit the unique optomechanical properties of this platform. Since multi-photon nonlinearities do not occur until the excitation power exceeds $\approx 1 \text{ W}$ with the use of a delocalized optical mode, the extent of pulling of the PhC membrane can be many tens of nanometers. Our silicon-based device provides a simple, non-intrusive solution to extending the actuation range of MEMS devices.

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