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Ultra-compact nonvolatile phase shifter based on electrically reprogrammable transparent phase change materials



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Abstract

Optical phase shifters constitute the fundamental building blocks that enable programmable photonic integrated circuits (PICs)—the cornerstone of on-chip classical and quantum optical technologies [1, 2]. Thus far, carrier modulation and thermo-optical effect are the chosen phenomena for ultrafast and low-loss phase shifters, respectively; however, the state and information they carry are lost once the power is turned off-they are volatile. The volatility not only compromises energy efficiency due to their demand for constant power supply, but also precludes them from emerging applications such as in-memory computing. To circumvent this limitation, we introduce a phase shifting mechanism that exploits the nonvolatile refractive index modulation upon structural phase transition of Sb₂Se₃, a bi-state transparent phase change material (PCM). A zero-static power and electrically-driven phase shifter is realized on a CMOS-backend silicon-on-insulator platform, featuring record phase modulation up to 0.09 $\pi/\mu m$ and a low insertion loss of 0.3 dB/ π , which can be further improved upon streamlined design. Furthermore, we demonstrate phase and extinction ratio trimming of ring resonators and pioneer a one-step partial amorphization scheme to enhance speed and energy efficiency of PCM devices. A diverse cohort of programmable photonic devices is demonstrated based on the ultra-compact PCM phase shifter.

The ability to actively control the phase of light is the common denominator in applications ranging from optical switching in data communications [3], beam steering, [4] spectroscopy, [5] optical neural networks, [6] as well as quantum processing and computing with photons [7]. Phase modulation has been achieved with carrier injection or depletion in semiconductors, [8] via electro-optic (EO) effects in liquid crystals, [9] EO polymers, [10] and Pockels crystals, [11] and through the thermo-optic (TO) effect, [12] and mechanical displacement [13]. The small refractive index perturbation from these mech-



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anisms typically results in a large device footprint, often hundreds of microns or larger (Table 1). Other limitations include large optical loss penalty (carrier injection/depletion) and long switching time (liquid crystals), as well as challenges associated with standard foundry process integration (EO polymers, Pockels crystals, and mechanical displacement). Moreover, all the mechanisms above are volatile, mandating a constant power supply to operate. Such volatility significantly increases energy consumption, especially for applications where reconfiguration operations occur at millisecond intervals or even longer. Nonvolatile phase shifters for PICs have been achieved through charge-trapping effects, [14] mechanically latched micro-electromechanical systems, [15] and ferroelectric domain switching, [16] which thus far have suffered from high losses, severe long-term drift, large form factors, or incompatibility with standard Si processes. A compact, low-loss and nonvolatile optical phase shifter (OPS) compatible with Si photonic integration is therefore much coveted.

Chalcogenide phase change materials (PCMs) offer a promising solution to nonvolatile programmable PICs [17-19]. Traditional PCM devices, however, cannot provide phase-only modulation due to high optical losses associated with classical PCMs exemplified by Ge₂Sb₂Te₅ [20, 21]. Moreover, most PCM device prototypes to date still rely upon furnace annealing or an external laser stimulus to trigger the structural transition. This limitation has motivated the development of low-loss PCMs in recent years such as Ge₂Sb₂Se₄Te (GSST), [22] Ge₃Sb₂Te₆, [23] Sb₂S₃, [24–26] and Sb₂Se₃ [27]. In particular, Sb₂Se₃ offers vanishingly small losses at 1550 nm and an index contrast of $\Delta n \approx 0.77$, [27] making it an ideal PCM for programmable photonics in the telecommunication bands [28, 29]. Capitalizing on the unique attributes of Sb₂Se₃, this work demonstrates an OPS that simultaneously confers low insertion loss, small form factor, and zero-static power consumption. The OPS devices were fabricated in silicon-on-insulator (SOI) where silicon acts both as the light guide and resistive heater to actuate structural phase transition via single electrical pulses. Notably, the devices were processed on 200-mm wafers leveraging a 90-nm CMOS foundry, enabling them to be seamlessly integrated with industrystandard PIC platforms.

Results

The PCM-based phase shifter comprises a 30 nm thick Sb₂Se₃ cell, initially prepared in the crystalline state, resting on top of a 220 nm SOI waveguide with a 110 nm rib. This waveguide geometry is designed for transverse electric mode propagation and has been used in all the programmable photonic devices in this work. A section of the waveguide was phosphorus doped to act as a microheater (Fig. 1a). More details of the foundry fabrication process are described in the Methods and the Supplementary Section S1. An essential aspect of the microheater design is the choice of doping concentration given the trade-off between resistivity and optical losses (Fig. 1b). Lower resistivity reduces the pulse voltages required to reach the transition temperatures of Sb₂Se₃ at the expense of exacerbated free carrier absorption. After testing four different doping combinations, we found a good balance with a concentration of $n \sim 4 \times 10^{18}$ cm⁻³, which corresponds to an insertion loss of ~ 0.01 dB/µm. Details on the design and optimization of the microheater are described in Supplementary Section S2. Figure 1c shows a scanning electron microscope image of a phase shifter with a 10 µm-long bowtie-shaped doping region. With a refractive index contrast of $\Delta n \approx 0.77$ upon switching (Fig. 1d), mode simulation



Fig. 1 Sb₂Se₃-based phase shifter using a doped-silicon microheater. **a** Microheater design with doped-silicon profiles to create an electrical resistor in a half-etched 220 nm SOI waveguide. **b** FEM simulated propagation loss and sheet resistivity as a function of phosphorous doping concentration. The highlighted region corresponds to the range where our proposed optimum doping concentration lies. **c** SEM image of a fabricated 6 µm-long bowtie microheater with a 6 µm-long Sb₂Se₃. **d** Sb₂Se₃ refractive index in both amorphous and crystalline states with simulated optical modes for each state at 1565 nm wavelength. **e** FEM simulated transient temperature during and after the pulse excitation for both amorphization and crystallization. The three-dimensional (3D) plot shows the temperature profile at the end of the amorphization pulse, reaching the melting temperature of Sb₂Se₃.

indicates a change in effective refractive index of $\Delta n_{eff} \approx 0.071$ and a phase shift of 0.09 π/μ m at 1565 nm. Given the flat refractive index of Sb₂Se₃ between 1500 and 1600 nm range, the phase modulation in this range would only depend on the wavelength and lie within 0.089-0.094 π/μ m. This results in an ultra-compact OPS of only 11 μ m in length needed to attain full π phase delay, representing significant improvement over the state-of-the-art. The small device footprint also yields a low theoretical insertion loss of 0.01 dB/ μ m resulting from free carrier absorption. Experimentally, we measured a total insertion loss of 0.03 dB/ μ m. The loss is larger than the expected value due to other effects such as scattering. This loss change associated with Sb₂Se₃ crystallization is experimentally assessed to be 0.018 dB/ μ m, corresponding to a low 0.2 dB/ π excess loss, including 0.1 dB/ π scattering loss which can be circumvented with improved designs (Supplementary Section S5 and S6).

To reversibly switch the device in Fig. 1c, we use Joule heating induced by a voltage pulse that elevates the temperature of the central area of the microheater, which coincides with the waveguide, following similar phase transition dynamics demonstrated in Ref [30]. The pulse width and voltage depend on the microheater geometry. The high impedance of the microheaters (~1 k Ω) led to an approximated 1.94-fold higher voltage dropped on the devices than the voltage set on the pulse generator, which assumes a 50 Ω load. The latter value will be quoted in brackets for reference. For the bowtie geometry in Fig. 1c, which we will extensively use throughout this work for $\pi/2$ phase shift, we used 6.2 V [3.2 V] (3 mA) pulses of varying duration (up to 1 ms) to crystallize the device by heating to above the Sb₂Se₃ crystallization temperature, $T_x = 200$ °C. To amorphize, a single 21 V [10.8 V] (7.8 mA) × 400 ns pulse was employed to raise the temperature over the melting point, $T_m = 620$ °C (893 K). The crystallization pulse length correlates with the total phase shift; while we observed crystallization with 5 µs pulses, short pulses would induce smaller hotspots with $T > T_x$, crystallizing only small areas of Sb₂Se₃. To compensate this effect, we used longer pulses in the 0.1-1 ms range to heat up the entire cell. The large voltage required to amorphize results from the nonlinear I-V response of



Fig. 2 Sb₂Se₃-based phase shifters in balanced (2×2 switch) and unbalanced MZI devices. **a** Layout of an unbalanced MZI with an optical microscope image of a 6 μ m Sb₂Se₃ cell on a 10 μ m-long microheater. **b** Experimental demonstration of four interferograms corresponding to each phase state permutation in the device shown in **a**. Crystallization was achieved using 6.2 V [3.2 V] and 1 ms pulses. Amorphization was achieved with 21 V [10.8 V] and 400 ns pulses. Extinction ratio over 35 dB is achieved between opposite states with a nearly complete π phase shift modulation. **c** Reversible switching over 125 full cycles between two intermediate Sb₂Se₃ states of the top-arm phase shifter in a balanced MZI. The plotted changes in transmission correspond to the difference between each arm's intensity. **d** MZI 2×2 switch in the four different permutations: amorphous and crystalline states of each arm's phase shifter with 6 μ m Sb₂Se₃ cell for π /2 shift. Reversible switching is demonstrated for both cells. The Gaussian-like transmission curves result from the use of grating couplers

microheaters, given the strong dependence of the carrier mobility on temperature (see Supplementary Section S2). Figure 1e shows a finite-element method (FEM) simulation of the temperature evolution for both the amorphization and crystallization pulses as well as the 3D temperature profile at the end of the amorphization pulse. Given the large thermal conductivity of silicon, heat rapidly dissipates after the pulse excitation with a decay time of 400 ns, allowing the fast quench process required to amorphize Sb₂Se₃. We found that the most prolonged pulse duration able to amorphize was around 1 μ s, beyond which crystallization cannot be completely suppressed during cooling (Supplementary Sect. 2).

We initially demonstrate phase modulation using an unbalanced Mach-Zehnder interferometer (MZI) with a 6 μ m-long Sb₂Se₃ OPS in each arm. Figure 2b shows the four permutations for the extreme phase states corresponding to a total phase shift of nearly π . We attribute the difference in extinction ration to the small excess loss upon crystallization, as we discuss below, and the difference between am/am and cry/cry states to small variations in the area that is switched. Continuous (multi-state) switching by partially crystallizing amorphous Sb₂Se₃ and a MZI with four permutations and phase modulation exceeding 1.6 π with a longer OPS are also realized and discussed in the Supplementary Section S8. We further demonstrated a 2×2 switch – a fundamental building block for constructing programmable photonic architectures [1] – in which light can be switched between two outputs. Since using two 50/50 directional couplers (DC) means that switching between output ports is achieved with either both OPS in phase or with a π phase difference, we opted for "unbalanced" directional couplers that display



Fig. 3 Reversible and multi-level switching of Sb₂Se₃ phase shifters in micro-ring resonators. **a** Layout of a 120 µmradius, 200 nm-gap micro-ring resonator with a 3 µm-long Sb₂Se₃ cell. This device is used to experimentally demonstrate **b** one full reversible cycle featuring some of the partial crystallization states and a total of 0.28 π shift, in excellent agreement with the expected 0.09 π /µm, and **c** ten full cycles with excellent reproducibility. **d** Same ring resonator as in **a** featuring a drop port identical to the main coupling port. **e** Demonstration of phase shift of resonance dip modulation in the through port with up to 10 dB in ER modulation. **f** Drop port peaks for the same exact measurements shown in **e** with a total phase shift of 0.47 π . 3.2 V and 100 µs - 1ms pulses were used to partially crystallize and a single 21 V [10.8 V] and 400 ns pulse to fully amorphize

a total equal splitting at the switch output with both OPS in phase. This is achieved by using ~80/20 splitting in each DC, and as result, we could switch between output ports by applying $\pm \pi/2$ phase shift in both arms. This is clearly not an optimized MZI switch (design that is beyond the scope of this paper) but it allows us to visualize the resulting amplitude modulation by actuating the Sb₂Se₃ phase shifter. Using a 10 µm-long heater with 6 µm-long Sb₂Se₃, we show 250 switching events (125 complete cycles) of the MZI (Fig. 2c). Reversible and reliable switching of each phase shifter allows the four different 'full' phase permutations, including splitting equally or directing the light to each of the two output channels with up to 15 dB extinction ratio (Fig. 2d). An unbalanced directional coupler causes the difference in extinction ratio (ER) between the two opposite phase states at the output of the MZI, but the $\pi/2$ phase shift is verified in each arm (Supplementary Section S3). In Fig. S4, we further demonstrate 22 dB extinction ratio for a similar 2×2 MZI switch in time domain measurements.

We now introduce Sb₂Se₃-based phase shifters in micro-ring resonators (Fig. 3a). Like MZI devices, we demonstrate continuous, multi-state modulation of an all-pass resonator by partially crystallizing a 3 μ m-long Sb₂Se₃ cell (Fig. 3b). Figure 3c shows the reversible switching of the cell in 10 cycles between fully amorphous and fully crystalline states with remarkable reproducibility and excellent agreement with the theoretical value of 0.09 π/μ m (estimated considering only the Sb₂Se₃ cell length). The losses upon crystallization were 0.018 dB/ μ m, which is quantified from fitting the transmission spectra of the two states (Supplementary Section S5). We then measured micro-ring resonators with an add-drop configuration and a 6 μ m-long Sb₂Se₃ cell (Fig. 3d). Figure 3e



Fig. 4 Extinction ratio tuning via phase-shift triggered coupling coefficient modulation. **a** Schematic of the ring resonators under test with a drop port at the phase shifter position and the corresponding modulation of the coupling coefficients *k* and *r*. **b** Experimental results for the device's through port for two switching cycles of a Sb_2Se_3 cell placed on the ring waveguide only, as shown in the inset. **c** Same as **b** for a Sb_2Se_3 cell covering both the ring waveguide and the drop port with nearly 20 dB modulation in ER.

and f demonstrate the phase-only modulation changing the resonance wavelength of the through-port dip or drop-port peak and the $-\pi/2$ phase modulation of the ultra-compact device.

In addition to spectral detuning of the resonant peak, we also demonstrate that the phase shifter can be used to modulate the extinction ratio of an over-coupled ring resonator by tuning the waveguide effective index at the coupling region, which modifies the coupling coefficient between the ring and the drop port [31] (Fig. 4a and Supplementary Section S7). We explored the two different configurations shown in Fig. 4b and c with Sb_2Se_3 cells covering either the ring waveguide only or both waveguides, respectively. Large extinction ratio modulation up to 20 dB was demonstrated in the latter configuration, where amorphization of the Sb_2Se_3 cell results in light lost to the drop port, therefore increasing the losses inside the ring, which then switches from over-coupled to near-critically-coupled response. As we will show later in this paper, simultaneous amplitude and phase trimming are potential applications of Sb_2Se_3 -based phase shifters, allowing the correction of fabrication variations in the targeted resonance wavelength and in the intended coupling regime.

Up to this point, we have shown Sb_2Se_3 phase shifters capable of achieving intermediate states via partial recrystallization only - modulating the number and the width of the crystallization pulses. As a result, when reconfiguring the OPS to a state with lower crystallinity, at least two pulses are needed, where one fully amorphizes and 'resets' the material while the other partially crystallizes the PCM. While this has so far been the standard procedure in PCM photonics for continuous electrical switching, [30, 32–35] it not only complicates the switching scheme but also significantly compromises the switching speed and energy efficiency, given that crystallization pulses are inherently much longer and energy consuming than the amorphization pulse [36]. A switching approach that enables fast bidirectional tunability is therefore strongly preferred. Here we demonstrate a unique single-step partial amorphization scheme by engineering the temperature profile across a microheater. For this purpose, a microheater shown in Fig. 5a and b consisting of five *n*-doped 'bridges' each of 2 µm in width and connecting both n^{++} regions was employed. Sb₂Se₃ is only deposited directly on top of each bridge. Changing the pulse width, the temperature profile along the waveguide and across the 'bridges' leads to melting temperatures initially only in the central 'bridge,' but then



Fig. 5 Engineering microheater to achieve partial amorphization. **a** FEM heat transfer simulation of a microheater consisting of five 2 µm-wide 'bridges'. The temperature profile along the waveguide is plotted for different pulse widths. **b** Colored SEM image of a fabricated device. **c** Partial amorphization of the bottom arm Sb₂Se₃ in an unbalanced MZI. 10.8 V pulses with varying widths between 800 ns and 1050 ns were employed to amorphize and 3.2 V for 1 ms to fully crystallize. The transmission is measured in the top output waveguide where a maximum of $\pi/2$ phase shift is observed, corresponding to the full amorphization of the three central 'bridges'. Reversible switching is demonstrated over three cycles between the full crystalline state and the marked intermediate state (corresponding to a 900 ns pulse). **d** The same phase shifter with partial amorphization in a micro-ring resonator with continuous tuning of the resonance wavelength

expanding to the outer ones as the pulse width (i.e., energy dissipation) increases. This response, in turn, allows for partial amorphization of the Sb₂Se₃ cells, correspondingly leading to different optical responses and phase shifting. With a total resistance of 1025 $\pm 25 \Omega$, the pulse width to start inducing amorphization was 800 ns with 21 V [10.8 V]. By varying the pulse width, *W*, from 800 to 1050 ns; we demonstrate partial amorphization of Sb₂Se₃ in an unbalanced MZI (Fig. 5c) and a micro-ring resonator (Fig. 5d). Figure 5c shows reversible switching between an intermediate state achieved with a 900 µs pulse and the fully crystalline state. Bowtie-shaped microheaters in Figs. 1 and 2 require 176 nJ for full amorphization followed by a partial crystallization in the 4 µJ - 38 µJ range if the target intermediate state was less crystalline (more amorphous). Using microheaters with 'bridges', these two pulses are replaced by a single partial amorphization pulse of 388 nJ, for a total of one to two orders of magnitude lower energy consumption per switching event. The concept of leveraging temperature gradient to achieve precisely controlled partial amorphization can be readily extended to other heater designs with more tuning levels.

Leveraging the bidirectional tuning capability, we lastly demonstrate an application that exploits the nonvolatile nature of chalcogenide PCMs: optical device trimming. We use a device comprising two micro-ring resonators connected to a single bus waveguide (Fig. 6) to show the potential of PCMs in countering the optical response variations resulting from fabrication non-uniformity. By tuning the phase of one of the micro-ring resonators and keeping the second as a reference, we demonstrate that a nonvolatile phase shift can be used to either isolate each ring's response (Fig. 6a) or to make them



Fig. 6 Optical trimming of micro-ring resonators. **a** Detuning of two initially superimposed resonance dips corresponding to two micro-ring resonators with 100 μ m (blue line) and 120 μ m (red line) in radius connected to a common bus waveguide. The largest ring is left fixed while the smaller ring is phase-shifted via step amorphization to fully isolate the response of each ring. **b** Trimming of the 120 μ m micro-ring to match the resonance dip of the smaller ring (fixed), thus, correcting the detuning in **a** but with an overall wavelength shift of 0.6 nm. The inset shows an optical microscope image of the actual device. **c** Demonstration of combined amorphization and crystallization to properly trim one micro-ring resonator and compensate 'over-shifting'. Nonvolatile response is demonstrated in the last state of the device (bottom panel) which was measured 4×10^6 s (i.e. 47 days) later

coincide by precisely tuning the resonance wavelength until matching (Fig. 6b). Furthermore, Fig. 6c shows how both amorphization and crystallization processes can be combined to shift the peak position in both directions and correct any undesired over-shift if the goal is for the peaks to match. Once the trimming is completed, and thanks to the well-known nonvolatility of PCMs, [17, 37] the final optical response is held stable over time, as we demonstrate up to 4×10^6 s (i.e. 47 days) later, with a small shift of 0.01 nm, which is difficult to differentiate from that associated with temperature fluctuations (see Supplementary Information). More importantly, our low-loss phase shifter still offers the option of recalibrating or changing the configuration later without any power consumption in between.

Discussion

We have demonstrated electrothermal control of the low-loss phase change material Sb_2Se_3 as a nonvolatile phase shifter in various reconfigurable photonic devices. Our choice of phase change materials is based on the large bandgap and the fact that Sb_2Se_3 is a single-phase compound throughout the entire solid-liquid temperature range, which, in principle, means high endurance. Considering that electrical current does not flow through the material in our approach, electromigration is also prevented, which is a mechanism for atomic movement and failure. In this work, we tested the endurance of over 250 switching events, a number ultimately limited by our experimental setup; however, 4,000 and 2,000 switching cycles have been demonstrated using laser pulses [28] and graphene microheaters on PICs, [38] respectively.

From simulations, we expected a phase shift of 0.09 $\pi/\mu m$, which was in excellent agreement with the experimental demonstrations shown for smaller Sb₂Se₃ cells (Fig. 1). For longer cells, the total phase modulation was slightly smaller, down to 0.08 $\pi/\mu m$. We attribute this discrepancy to possible thickness variations of the Sb₂Se₃ film and non-uniformity of the heating profile. Nonetheless, we demonstrated that an ultra-compact device comprising only a 10 μ m-long microheater and a 6 μ m-long Sb₂Se₃ is enough to

introduce $\pi/2$ phase shift. In turn, two $\pi/2$ nonvolatile phase shifters in a balance MZI switch, one per arm, allowed us to demonstrate another fundamental building block of programmable photonic circuits, a 2×2 optical switch (Fig. 2). We have also demonstrated the phase tunability of all-pass and add-drop micro-ring resonators to control the signals in the through and drop ports (Fig. 3), and amplitude modulation by engineering the coupling efficiencies to a drop port (Fig. 4). The insertion loss of the phase shifter (0.03 dB/ μ m) can be lowered by further engineering the shape of the PCM cell to suppress scattering [39]. Additionally, the microheater doping profiles and geometry can also be optimized. A PIN configuration, which has demonstrated losses of 0.02 dB/ μ m³³ can be a good alternative; however, the width of the intrinsic region limits the lateral area of the microheater, which, in our approach, can be arbitrarily large. Single-doped heaters can therefore be used for waveguides, but also for larger PICs components such as MMIs and even to fit large metasurfaces. Other alternatives for transparent microheaters, already demonstrated to switch PCMs, might be considered to avoid doping losses overall and extend the transparency window to shorter wavelengths, such as graphene, [38, 40] fluorine-doped tin oxide, [41] and indium-tin-oxide; [39, 42] however, their full integration and reversible cyclability on integrated waveguides is yet to be demonstrated. We draw a direct comparison between our PCM approach and other phase-shifting approaches in Table 1 considering 10-90% rise time, the total change in the effective index, Δn_{eff} , the total length and voltage to achieve a π phase shift, L_{π} and V_{π} , and the insertion loss (IL).

The nonvolatility of our phase shifter guarantees zero-static power consumption – a new paradigm for PICs. Even if reconfigured every millisecond, our approach's energy efficiency is still superior to TO phase shifters. However, if we consider only applications in which programming is done sporadically, then the energy savings are orders of magnitude different. For example, let us consider a 32×32 optical switching fabric architecture, [22] consisting of 512 2×2 switches for a total of 1024 OPS (using the design in Fig. 2d), whose output is reconfigured every 1 s. Using our Sb_2Se_3 -based phase shifters, the average switching energy – calculated using the full crystallization pulse, which is the most energy-consuming process – would be 39.3 mJ. This is two orders of magnitude lower than a constant supply of 1.9 J to reconfigure the lowest reported 32×32 network with the same MZI switches but using TO phase shifters [43]. The energy-saving benefit becomes even more significant with longer durations between switching events. Such nonvolatility is also interesting for in-memory computing, [37, 44-46] and configure-once applications, such as optical trimming and transient coupler for wafer-scale photonic testing [47]. Moreover, in addition to the nonvolatile modulation by phase transitions of Sb₂Se₃, the doped-silicon microheater can provide conventional thermooptical modulation (see supplementary information), which further enhances the versatility of our approach by enabling dual nonvolatile and volatile response in a single device. We note that while the contact pads to electrically connect our devices are >1mm [2, they do not represent a limitation in terms of scalability. In 3D stacking architectures, vertical vias can be used to contact the doped-silicon while keeping on-silicon footprints of $20 \times 20 \mu m^2$.

In conclusion, we have experimentally demonstrated a new class of electrically-driven phase shifter that exploits the unique optical properties of phase-change materials to achieve zero-static power consumption in ultra-compact devices. We demonstrated

	${\it \Delta}n_{eff}$	IL(dB)	L _π (μm)	Ε _π	ν _π (V)	Non-volatility	Rise time ^d
Thermo-optical	~1.8×10 ⁻⁴	0.23	61.6	$24.8 \times t[s]$	4.36	No	2.69
(doped-Si) [12]	×ΔT[K]			mJ			μs
Thermo-optical (metal heater) [12, 48]	~1.8×10 ⁻⁴ ×∆T[K]	< 0.4	>200	20–25 × t[s] mJ	4	No	3.7 µs
Thermo-optical (ITO) [49]	$\sim 1.8 \times 10^{-4}$ $\times \Delta T[K]$	< 0.01	50	10 × <i>t</i> [s] mJ	9.1	Νο	5.2 µs
Thermo-optical with slow light [50]	~0.1	2	~10	$2 \times t[s] mJ$	4.5	No	100 ns
Electro-optical (depletion) [51]	1.5×10 ⁻⁴	3.6	1500	~ 2 × t[s] pJ ^a	10.7	No	15.5 ps
Electro-optical (injection) [52]	~ 10 ⁻³	2	400	1.7 × <i>t</i> [s] mJ	2	Νο	24 ns
Optoelectro- mechanical [13, 15]	~0.01	0.47	84	179 × <i>t</i> [s] nJ	2	Νο	123 ns
Liquid crystal [9]	0.016	0.25	49	1 × <i>t</i> [s] nJ	5	No	1 ms
Plasmonic nonlinear polymer [10] ^c	0.055	12	29	$4 \times t$ [s] mJ	45	No	15 ps
BTO Pockels effect [11]	7×10^{-4}	1	1000	8.2 × <i>t</i> [s] mJ	3.3	Νο	11.7 ps
BTO Ferroelectric domain switching [53]	~ 1 × 10 ⁻⁴ / pulse	0.07	150	4.6–26.7 pJ / pulse ^e	-10	Yes	300 ns
LiNbO ₃ [54]	1.5×10 ⁻⁴	2.5	5000	53 × t[s] mJ	5.1	No	3.5 ps
Transition metal dichalcogenides [55]	1.5×10 ⁻³	0.55	1000	0.64 × t[s] nJ ^a	8	Νο	1 ns
30 nm Sb ₂ Se ₃ cladding cry/am (this work)	0.07	0.36 ^b	11	38.4 μJ / 176 nJ	6.2/ 21.0	Yes	0.1-1 ms / 800

Table 1 Comparison of phase shifter platforms currently available in 220 nm SOI at 1550 nm. E_{π} is the energy required to switch and hold the state. ΔT is the difference in temperature, and t[s] is the time of operation

^a No current flows in this device, making the power dissipation extremely low. However, a constant supply of large voltage means a significant power consumption of the electronics. ^b Based on a 12 µm-long Sb₂Se₃ OPS – it can be further decreased if using lower doping concentrations. ^c These values are calculated from the supplementary material of the referenced paper.^d 10-90% rise time, where we used the single-pole approximation: $\tau = 0.35/f_{3dB}$ for results with no rise time reported. ^e Several pulses are required to achieve full π phase shift, and a constant DC signal is required to read out

several active photonic devices and applications as well as the potential of these devices for low-energy programmable photonic integrated circuits. Our results are promising for applications in optical trimming to correct fabrication variations and in photonic architectures with sporadic reprogramming, such as optical switching fabrics for telecommunications, phased arrays for particle trapping, photonic computing (inference with static matrices, for instance), and quantum networks, among others.

Methods

Device fabrication

Ten 220 nm SOI wafers were fabricated with varying *n*-doping concentrations using the 90-nm CMOS line in Lincoln Laboratory's 200 mm wafer foundry (Supplementary Section S1). The doping of silicon was carried out using ion implantation of phosphorous with varying dose and implantation energy. The n^{++} region was formed with a dose of 10^{16} cm [2] and an ion energy of 80 keV. The *n* region of $\sim 4 \times 10^{18}$ cm⁻³ was formed with a dose of 10^{14} cm⁻² and an ion energy of 80 keV. The contact was fabricated with a Ti/TiN barrier following a metallization with aluminum and passivation with SiO₂.

500 nm-wide SOI waveguides and Sb₂Se₃ cells were fabricated following two electron beam lithography fabrication steps on an Elionix ELS-F125 system (Supplementary Section S1). The waveguides were patterned using ZEP 520 A positive photoresist followed by chlorine etching of half the silicon thickness. A lift-off process with polymethyl methacrylate (PMMA) resist was used to open the windows for the subsequent thermal evaporation of 30 nm Sb₂Se₃. The film deposition was performed using thermal evaporation from Sb₂Se₃ bulk materials following previously established protocols [56, 57]. Bulk starting Sb₂Se₃ was synthesized using a standard melt-quench technique from high-purity (99.999%) raw elements [58]. The samples were annealed in an argon environment at 200 °C for 10–15 min before depositing 15 nm of Al_2O_3 by atomic layer deposition to obtain a conformal protective layer.

Experimental setup

A home-built wafer-scale automated photonic testing system was used to perform the optoelectronic measurements. A custom 18-channel SMF-28 Ultra Fiber Array with a 250 μ m pitch (PLC-Connections) was used to couple light in and out of the chip using silicon-etched grating couplers. A C+L band optical vector analyzer (LUNA OVA5000) was used to scan the spectrum and conduct time-domain measurements. The high voltage electrical pulses were generated with an analog 1 GHz, 40 V pulse generator (E-H Research 136 A) with a minimum of 0.5 ns raising and trailing edges, and the crystallization pulses with an Agilent 33,250 A 100 MHz pulse generator. An ACP40-AW-GSG-250 air-coplanar probe (FormFactor) was used to contact the electrical pads on the chip.

Microheater modeling

The Joule heating process and heat dissipation model were performed using a threedimensional finite-element method simulation in COMSOL Multiphysics. We used the Semiconductors module to simulate the carrier mobility in doped silicon with the same values and shapes as our fabricated devices. We coupled this module with the Heat Transfer in Solids, where surface-to-surface radiation and thermal boundary resistance were considered. The temperature in Heat Transfer was coupled back to the semiconductor module. The mode calculations shown in Fig. 1d were carried out in Lumerical Mode, considering the Sb₂Se₃ refractive index shown in Fig. 1b and the built-in refractive indices for Si, Al₂O₃, and SiO₂.

Supplementary information

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Supplementary Material 1

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Author contribution

C.Ríos performed material deposition, device design and electrooptical measurements of the photonic devices. Q.D. carried out the in-house fabrication of photonic devices. P.M., C.Roberts, and S.V. carried out the foundry-processed fabrication of the chips and material characterization. Y.Z., C-C.P., and M.Y.S. assisted in device fabrication. M.K. prepared the bulk materials. T.G., K.A.R., and J.H. supervised the research. All authors contributed to technical discussions and writing the paper.

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Data Availability

The data that support the findings of this study are available from the corresponding authors upon reasonable request.

Declarations

Competing interests

The authors declare that they have no competing interests.

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