Structurally and morphologically engineered chalcogenide materials for optical and photonic devices

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Abstract. We discuss several recent advances in the development of methodologies and techniques used to structurally and morphologically engineer chalcogenide (ChG) materials. We introduce two ChG patterning techniques both offering spatial resolution beyond the classical single-photon diffraction limit: multiphoton lithography and thermal scanning probe lithography (TSPL). The former was applied to produce nanoscale modifications in thermally deposited As2S3, and we realized gradient refractive index (GRIN) effective medium lens fabrication in multilayer As2S3-As2Se3 films with features as small as 120 nm using this approach. The GRIN lens was shown to be optically functional. ChG Ge-Sb-Se-Te (GSST) material was also explored for its potential as a phase-change material (PCM). We demonstrated nanoscale feature patterning using TSPL in PCMs with critical dimensions below 100 nm. In addition, new patterning methods, we also report solution processing of GSST PCMs as an alternative route for ChG film deposition. These new material processing and structuring techniques will offer new pathways for creating functional planar optical and photonic devices.© The Authors. Published by SPIE under a Creative Commons Attribution 4.0 Unported License. Distribution or reproduction of this work in whole or in part requires full attribution of the original publication, including its DOI. [DOI: 10.1117/1.JOM.1.1.013502]

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1 Introduction

Chalcogenide glasses (ChGs) are formed when one or more chalcogens (S, Se, or Te) are melt-processed with one or more elements from groups XIII to XV.¹ They are well known for their useful optical properties, which has led to many infrared (IR) applications in the fields of medicine, photonics, environmental sensing, and security.² ChGs can be compositionally tuned and processed to have useful properties such as a tailorable coefficient of thermal expansion, a suitably large refraction index or dispersion modification, combined with the ability to maintain a low thermo-optical coefficient (dn/dT), increased mechanical strength, and improved radiation hardness.³ Compositions can be crystalline, as in the mineral orpiment arsenic trisulfide (As2S3), but many form IR-transparent, homogeneous optical-quality glasses (in select cases transformable to optical glass ceramics), that can be engineered to be thermodynamically and/or kinetically stable over a wide range of compositional variations.⁴,⁵ ChGs can be used as bulk, fiber, or as thin films via thermal deposition or solution processing.⁶

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This paper discusses recent advances in the development of methodologies and techniques used to structurally and morphologically engineer ChG thin films for optical and photonic device applications. Multiphoton lithography (MPL) is used for nanoscale modifications in thermally deposited ChG multicomponent compositions such as As$_2$S$_3$, Gradient refractive index (GRIN) effective medium lenses were created in As$_2$S$_3$ and characterized for optical function. Laser writing in Ge-Sb-Se-Te (GSST) phase-change materials (PCMs) allows for simple area-selective phase transitions; however, its resolution is bound by the optical diffraction limit. An alternative atomic force microscopy (AFM)-related technique, thermal scanning probe lithography (TSPL) is introduced for high-resolution lithography. TSPL and large area thermal conversion are used for large and nanoscale modifications in GSST PCMs. Processing and characterization of high-performance solutions derived GSST PCMs are demonstrated. Switching performance and prospects of this system as a functional metalens are explored.

2 Multiphoton Lithography in Arsenic Sulfide As$_2$S$_3$ Films

As$_2$S$_3$ is a ChG material with excellent IR transparency (620 nm to 11 μm), an optical band gap at 2.35 eV (λ ~ 517 nm), and large nonlinear refractive indices. Thermal deposition of bulk As$_2$S$_3$ fragments the network solid, creating molecular clusters that can be deposited as photosensitive thin films. Photoexcitation through processes such as the MPL (laser writing) cross-links the homopolar bonds in the film back to the glassy network solid. Laser written areas remain intact after being submersed in a polar solvent [typically a mixture of diisopropanolamine (DIPA) and dimethyl sulfoxide (DMSO)] while the rest of the unexposed As$_2$S$_3$ film is etched away, resulting in a remaining structure that has the shape of the laser written pattern. Although other lithographic techniques for patterning As$_2$S$_3$ are available, MPL is the standout method for achieving highly controllable As$_2$S$_3$ material processing at the microscale without the use of clean rooms and masks. Due to its nonlinear optical nature in the laser–matter interaction when ultra-short laser pulses are used, MPL can achieve material modifications that are confined to the focal volume and can reach the micro- and nanoscales without adverse thermal effects. MPL in thermally deposited As$_2$S$_3$ films has been shown to produce high-precision three-dimensional (3D) micro- and nanostructures, such as waveguides, woodpile photonic crystals, and nanowires. In Sec. 2.1, we show how the optimization of processing conditions is used to photopattern 3D microstructures in 4-μm-thick As$_2$S$_3$ films thermally deposited on a ZnSe substrate. In Sec. 2.2, we use MPL to fabricate As$_2$S$_3$ microstructure-based metalenses that offer reduced size, weight, and cost as compared to traditional bulk IR counterparts.

2.1 Optimization of MPL and Chemical Etch Processing Conditions for the Creation of Microstructures in As$_2$S$_3$ Films

A femtosecond Mai Tai Ti:sapphire laser centered at 800 nm with a repetition frequency of 80 MHz is used for MPL in 4-μm-thick As$_2$S$_3$ films thermally deposited on a ZnSe substrate. The beam is passed through a variable attenuator to control the average power and a shutter, both of which are controlled externally though a computer. The beam is focused by either a 50×/0.55 NA or 100×/0.75 NA objective onto the As$_2$S$_3$ film. The substrate is secured onto a Physik Instrumente H-840 6-Axis Hexapod (programmable stage), and a camera is used to image the sample plane during irradiation. The entire system is controlled by a LabVIEW program that automates the laser writing of predetermined patterns.

A series of experiments were initially conducted in order to dial in on the optimal power, write velocity, and focus depth of laser into films. Dose arrays of raster scans, grids, and concentric circles are laser written into the film at varying average powers, write velocities, and focus depths of laser into film to test for the best etch selectivity as a function of these parameters. Examples of these laser written patterns are shown in Fig. 1. A range of powers is defined as a percentage of maximum possible average power at the sample plane from the variable attenuator. For the 50×/0.55 NA objective, 100% power corresponds to 40 mW and for the 100×/0.75 NA objective, 100% power corresponds to 35 mW. The etch results from these experiments indicate that a write velocity of 0.25 mm/s at an average power in the 40% to 60% range induces the desired
molecular change to the film without ablation. Higher powers and lower write velocities were usually found to ablate the film, whereas lower powers and higher write velocities did not result in good contrast between exposed and unexposed regions after etching. Further experimentation is in progress to determine the depth of focus into film for optimal etch selectivity, as indicated by the $+3 \mu m$ legend in Fig. 1. In this scenario, the laser is focused $3 \mu m$ deep into the film.

The postirradiation molecular structure of the film was investigated using micro-Raman spectroscopy. Figure 1(d) shows the Raman spectra collected at $5 \mu m$ intervals over a raster scan (60% power, 0.25 mm/s write velocity). A Horiba Jobin Yvon confocal micro-Raman system with an excitation wavelength of 785 nm was used. The beam was focused by a 100x/0.75 NA objective and the Raman signal was collected for 60 s at a low optical power (reduced by optical density filter), to prevent any undesired photoinduced change to the film. The laser-induced film cross-linking causes the exposed regions of the film to exhibit a different Raman signature from the surrounding pristine film.11,13 Most notably, the As$_4$S$_4$ molecular clusters that are present in the as-deposited/unirradiated film, characterized by the Raman peaks at 222 and 363 cm$^{-1}$ [as illustrated by the red spectra of Fig. 1(d)] emerge during the laser exposure process (blue spectra), leading to good etch selectivity between these two regions.

After the patterns are written using the laser, the sample is submerged into a solvent mixture of DIPA and DMSO for the etching process. After the sample is initially immersed for an hour, the resulting structure is analyzed using a ZYGO 6300 white-light interferometer (Fig. 2). Subsequent etching is done in 30-min increments until the profile of the structure is fully developed. Etch times vary depending on laser power and target resolution of laser-written structures, but it typically takes between 1 to 4 h of immersion. Good etch selectivity has been shown repeatedly down to $7.5 \mu m$ feature spacing, as shown in Fig. 2(a). The pictured $7.5 \mu m$ grid structure in Fig. 2(a) (irradiated at 60% power or 21 mW, 0.25 mm/s, and 100x objective) was achieved with 1 h of etching. At $5 \mu m$ feature spacing, etch selectivity takes longer to develop. This can be attributed to undesired partial exposure of the film in the regions between the laser write paths.

The height map shown in Fig. 2(b) indicates that the structure did not fully resolve after 1 h of etching. Subsequent etching on this sample resulted in the desired structure, as evident by the height map in Fig. 2(c). This $5 \mu m$-spaced structure (irradiated at 60% power or 21 mW, 0.25 mm/s, and 100x objective) took 240 min of etching to fully resolve.

In summary, MPL is shown to be effective technique for photopatterning 4-$\mu m$-thick thermally deposited As$_4$S$_4$ films. Experimentation through repeated dose arrays indicate that an average irradiation power between 40% and 60% at a write speed of 0.25 mm/s induces the desired molecule change in the films. Good etch selectivity for these conditions were shown repeatedly for $7.5 \mu m$ feature spacing after 1 h of etching and for $5 \mu m$ feature spacing after 240 min of etching.
2.2 Multiphoton Lithography of Functional GRIN Lenses

Single-layer As$_2$S$_3$ films are inadequate for nanostructure MPL, where the incident wavelength is on the order of, or smaller than, the thickness of the film, because of the standing wave caused by reflection at the substrate interface. This effect leads to the production of unintended nanobead structures with large sizes and poor substrate adhesion. In this section, we investigated the properties and processing of MPL in single- and multilayered films of As$_2$S$_3$ thermally deposited onto an underlying thin film of arsenic selenide (As$_2$Se$_3$) and the creation and optical characterization of a GRIN effective medium lens.

Thermal deposition was used to create a multilayered film of As$_2$S$_3$ on an anti-reflective (AR) layer of As$_2$Se$_3$. A layer of As$_2$Se$_3$ was deposited first on polished silicon wafers followed by thermal evaporation of a layer of As$_2$S$_3$. The thermal deposition process has been described previously. The As$_2$S$_3$ layer was deposited at a rate of 30 Å s$^{-1}$, and the thickness of the resulting film was 896 ± 13 nm. The As$_2$Se$_3$ layer was deposited at a rate of 4 Å s$^{-1}$ and the thickness of the resulting film was 203 nm ± 3 nm. The As$_2$Se$_3$ serves as an AR layer, which reduces reflection of the incident laser beam back into the focal volume during laser patterning.

MPL was performed using a system and process described previously. MPL was conducted using the output from a mode-locked Ti:sapphire laser ($\lambda = 800$ nm, 76 MHz repetition rate, and 120 fs pulse duration). An acousto-optic modulator was used for the electronic control of the intensity of the laser beam. The laser pulses were focused into the film with a 100×/1.4 NA oil-immersion objective (Nikon). GRIN lenses were photopatterned by translating the film along programmed paths in the $x$, $y$, and $z$ directions at a speed of 50 μm s$^{-1}$ while keeping the laser beam fixed, and then opening and closing a shutter in synchrony with the movements to control local exposure. The average focused laser power was varied from 0.05 to 0.50 mW. The detailed chemical etch processing has been described elsewhere.

Incorporating the etch-resistant As$_2$Se$_3$ AR-layer between the ChG fabrication layer and the substrate reduces the amplitude of a standing wave caused by reflection at the substrate.
This reduction resulted in the generation of pillars with uniformly cylindrical shapes with increased material–substrate adhesion that were more likely to obtain a smaller size before delamination than the bead-shaped features created in the single layered film [Fig. 3(a)]. The uniform shapes also allowed for more predictable fill fractions when compared to stacked bead shaped nanostructures.8,15 Predictable fill fractions are important for GRIN lens design when determining effective index and optical function. Large pillar arrays of uniformly cylindrical shapes were demonstrated in the previous works.8,15

GRIN effective-medium lenses with diameters of 20 and 100 μm were successfully fabricated in the multilayered film. Figure 2(c) shows scanning electron microscope images of the top-down view of the GRIN lens having a diameter of 20 μm. The inset to Fig. 2(c) shows the top-down view of the GRIN lens having a diameter of 100 μm.

The GRIN lenses consisted of cylindrical pillar features separated by a pitch of 500 nm. The lenses were designed to have a spherical refractive index profile that ranged from $n = 1.06$ to 2.45, the latter representing a region completely filled in with As$_2$S$_3$. The corresponding total optical phase delay from center to edge of the lens would be $1.26\pi$. The index profile was created by fabricating pillars having diameters that decreased with radial distance from the center of the lens, so that the corresponding decrease in local fill-fraction of the high-index ChG creates a local refractive index that follows that targeted spherical refractive index profile. Across the lens, the pillars had consistent height, and within a small designed region, the diameters of the pillars

![Fig. 3](image_url)

**Fig. 3** (a) Comparison of pillars fabricated in single-layered film (top) and multi-layered film with an AR coating (bottom). (b) Diagram of the source beam interaction with sample. (c) GRIN effective-medium lens (diameter = 20 μm) designed to have a spherical refractive index profile and fabricated by MPL in thermally deposited As$_2$S$_3$. This device consists of cylindrical pillars, which increased in fill fraction (effective refractive index) from the outside diameter to the inner. (d) Transverse intensity scans within focal plane of the GRIN effective medium lens (diameter = 100 μm) designed to have a spherical refractive index profile.

Schwarz et al.: Structurally and morphologically engineered chalcogenide materials...
were similar. Widths of the pillars ranged from 120 nm ± 14 nm fabricated at a power of 0.16 mW to pillars large enough to overfill the entire area. The pillar heights were measured to be a 907 nm ± 10 nm across the structure and were fixed based by the original film height.

The GRIN lenses were characterized using the output from an Er:YAG laser ($\lambda = 2.94 \, \mu\text{m}$, 170-µs pulse duration) with the system shown schematically in Fig. 3(b). The laser beam over-filled the sample area so that the intensity distribution across the microlens was uniform. An optical fiber coupled to an IR photodetector (ThorLabs PDA 20H) was mounted on a three-axis translation stage and positioned behind the sample. Transmitted light collected by the fiber provided a relative measure of the local intensity as a function of position behind the lens. Measurements of transmitted light far from the lens provided a baseline measurement that can be used to correct for loss due to reflection from the supporting silicon substrate.

Figure 3(c) shows data from the optical characterization of the 100-µm diameter GRIN lens shown in the inset. The black trace is a line scan of light intensity transmitted through the silicon substrate far from the lens. The transmission is roughly 50%, as expected due to reflection losses at both the optical interfaces of the front- and backsides of the silicon substrate. The transmission versus position is uniform within the shot-to-shot variation of the pulsed laser, which confirms that the lens will be uniformly illuminated when positioned in the probe beam. Figure 3(d) shows the transverse intensity scans within the focal plane of the GRIN lens. Irradiance profiles generated by scanning in the $x$ or $z$ direction of the lens resulted in peaks of ~250% of $I_0$.

A homogenous phase shifting aperture “block” with a 100-µm diameter was created and characterized in the same manner as the lens. This was done in order to benchmark the transmission of the lens and account for increased throughput on the front-interface of the substrate that may result from refractive-index matching caused by the nanostructures. At the focal point of the lens, the intensity was found to be two times more intense than that transmitted through the uniform block, and four times as intense as light transmitted through the unpatterned substrate away from the lens. The transmitted light behind the lens had a clear focal structure with an FWHM of 32.1 µm ± 1.9 µm for the $x$ scan and 28.6 µm ± 1.5 µm for the $z$ scan. The effective width of the incident beam may be taken to be equivalent to the effective aperture set by the width of the GRIN lens, which was ~140 × 250 µm$^2$ at 90% intensity.

The addition of an AR layer enabled the fabrication of robust reproducible pillars with better material substrate adhesion and smaller dimensions via MPL in As$_2$S$_3$ film. GRIN lenses with spherical index profiles were created and IR characterization indicated optical functionality. This work opens new routes to ChG-based electronics and optical devices such as detectors, sensors, photonics waveguides, and acousto-optics.

### 3 Processing and Nanopatterning of Chalcogenide Phase Change Materials

We have shown in preceding sections that MPL is a powerful technique to generate large-area and high-aspect-ratio patterns in ChG films via modulating the glass local network structures and hence their chemical solubility. Here, we turn our attention to another class of ChG compounds known as phase change materials (PCMs). PCMs possess an ability to reversibly transform between the amorphous and crystalline states via electrical, optical, thermal, or a combination of these stimuli. The switching property of PCMs has enabled optical storage media devices including rewritable DVDs and Blu-ray discs while also opening the way toward next generation nonvolatile memory applications such as phase-change random access memory. Recently, PCMs are entering the arena of integrated photonic applications as platforms to arbitrarily tailor light–matter interactions.

A ChG alloy of Ge$_2$Sb$_2$Te$_5$ is notably the most industrially adopted PCM due to its rapidly transformed response, exhibiting a large contrast in optical and electrical properties upon phase transitions as well as thermal stability over a wide range of temperatures. This unique attribute qualifies PCMs as promising material candidates for innovative reconfigurable optoelectronic devices. Therefore, different from As$_2$S$_3$ and As$_2$Se$_3$ whose photosensitive structural changes are more subtle, photonic structures can be directly written into PCM films by inducing crystallization in a spatial-selective manner without resorting to subsequent chemical etching.
In the following studies, we focus on Ge$_2$Sb$_2$Se$_{5-x}$Te$_x$ ($x = 0.5$ to 1.0) (GSST), an emerging composition exhibiting exceptional bistate broadband transparency. The addition of Se leads to a marked decrease in absorption-induced optical loss while the remaining Te allows the PCM to retain a large contrast in refractive index upon phase transitions. The optical constants of GSST are shown in Fig. 5(a). The ability to produce subwavelength features in GSST is, therefore, of significant interest for applications such as photonic switching and active metasurfaces. Although laser patterning provides a facile route for triggering area-selective phase transition in PCMs, its resolution is ultimately bound by the optical diffraction limit. In Sec. 3.1, we introduce a stylus-based writing technique based on AFM enabling nanoscale patterning in thermally deposited GSST well below the wavelength. In Sec. 3.2, we explore solution processing of GSST materials as an alternative to thermal deposition and as a route to films that can be deposited and patterned on large-area nonplanar platforms.

3.1 Nanopatterning of Chalcogenide Phase Change Material via Thermal Scanning Probe Lithography

Stylus-based writing based on AFM techniques offers significantly improved resolution not constrained by optical diffraction. So far, AFM-based patterning of PCMs has been limited to external laser-assisted heating or using conductive atomic force microscopy (c-AFM). However, the c-AFM method is limited to substrates that can form a conductive loop with the AFM tip to

Fig. 4 Thermal scanning probe patterning of the phase-change material GSST. (a) Optical micrograph of a thermal probe (NanoFrazor, Swisslitho AG) that offers simultaneous in situ imaging and patterning capabilities. (b) Raman spectra of as-deposited GSST film and the same film in regions patterned with TSPL. (c) Depth map displaying a 100-nm-wide crystalized line resulting from volume contraction upon GSST crystallization. (d) A snapshot cross-sectional temperature profile in a 30-nm-thick GSST film on Si after 300 μs thermal AFM tip dwell time, simulated using the finite-element method. This profile corresponds to the maximum crystallization depth for GSST, before going over the melting temperature (~890 K) on the surface.
facilitate Joule heating. An alternative AFM-related technique, TSPL, has recently demonstrated its promise for high-resolution lithography processes in nanoelectronics, thermochemistry, and biomolecular nanodevices. The approach enables simultaneous in situ patterning and AFM imaging [Fig. 4(a)], and unlike c-AFM, it does not require a closed circuit, making it compatible with a diverse range of substrates. Using this technique, we demonstrated nanoscale patterning of an amorphous GSST film with 30-nm thickness. Micro-Raman measurements confirmed that the thermal AFM writing process leads to local crystallization of GSST, evidenced by a shift in the Raman signal peak from near 160 to 120 cm$^{-1}$, associated with the crystalline phase of GSST [Fig. 4(b)]. In Fig. 4(c), a TSPL-written 100-nm-wide serpentine line pattern is imaged by tracing the changes of GSST volume upon switching. The result indicates the potential of the thermal AFM technique for nanoscale patterning of PCMs.

To further elucidate the PCM patterning mechanism, we performed finite-element simulation of the transient temperature profile in GSST during thermal nanopatterning. The model was implemented in the commercial COMSOL heat module. The thermal probe is approximated as a point heat source with $4 \times 10^{-6}$ W power following our experimental parameters. Figure 4(d) shows the temperature distribution inside a 30-nm-thick GSST film after a tip-dwelling time of 300 μs. The highly localized temperature profile suggests that nanoscale patterning with resolution better than 20 nm is viable with our TSPL technique. The simulation also indicates that only the top 15 nm of the film is crystallized, largely because the thermally conductive Si substrate serves as an efficient heat sink. By modulating the power on the tip, we envision a modulation on the crystallization depth by reaching different amorphous/crystalline areal distributions, which enables applications such as color depth modulation for grayscale patterning and continuously phase tuning for holographic and metasurface applications. The result validates the potential of TSPL for nanopatterning of PCM ultrathin films, providing a pathway toward realizing topologically optimized, reconfigurable photonic devices.

### 3.2 Solution-Deposited Chalcogenide Ge-Sb-Se-Te Films as a Route Toward Low-Loss Phase-Change Optical Materials on Large-Area Nonplanar Platforms

Although GSST PCMs have been fabricated using either a standard melt-quench (bulk) or thermal evaporation (film/waveguides), an ability to conformally deposit PCM layers on large-area nonplanar platforms such as lenses or substrates with complex surface topography has been increasingly demanded. Employing previously demonstrated methods of dispersing chalcogen-containing alloys into solution, one can form films of these alloys following composition-specific protocols. Solution deposition (SD) is a cost-effective and scalable method where substrates with virtually any shape and size are amenable to forming conformally coated layers, as illustrated in Fig. 5(a). Here, we discuss the ongoing progress made on processing high-performance Ge$_2$Sb$_2$Se$_{5-x}$Te$_x$ PCM alloys, using SD.

Figure 5(b) illustrates the step-wise SD process flow. The process begins with the preparation of bulk Ge$_2$Sb$_2$Se$_{5-x}$Te$_x$ ($x = 0.5$ to 1.0) formed using a standard melt-quench method. The bulk alloys are ground and filtered through a 120-μm sieve to realize fine powder and then dissolved into a homogeneous solvent mixture of ethanethiol (thiol) and ethylenediamine (EDA). Polished and plasma cleaned ZnSe substrates are dipped into the solution using a speed-controlled robotic arm to create films of uniform thickness and surface quality. Finally, the as-coated films undergo soft- and hard-heat treatment to remove residual solvents. To realize optimal performance, SD films must maintain parent alloy stoichiometry, exhibit spatially uniform surface quality, be optically thick ($t \geq 1 \mu m$), be chemically compatible with the underlying IR transparent substrates, exhibit low optical loss in the mid-wave IR, and convert to desired crystalline phases upon thermal or laser treatment. These key attributes assure minimum degradation to the film’s phase change response.

Here, we discuss key process conditions that have deterministic impacts on the resulting quality of coated films. Specifically, the atomic percentages of constituent elements in resulting films are significantly associated with the thiol-EDA mixing ratio. Although Ge and Sb elements have been shown to be easily soluble using amine solvents such as propylamine and EDA, Se and Te are inherently more challenging to dissolve due to their preferential dissolution kinetics.
necessitating the introduction of an additional solvent such as thiol, as shown in prior efforts.\textsuperscript{56,59} Specifically, Te has very low solubility in amine solvents, and thus is partially transferred to a substrate from the solution, disrupting the stoichiometry of resulting films from that of the target parent alloy. This variation, as a function of solvent chemistry, is illustrated in Fig. 5(c). Meanwhile, the composition data extracted using energy dispersive spectroscopy (EDS) on resulting films in Fig. 1(c) indicate that an increase (decrease) in the volume percentage of thiol (EDA) leads to an increase (decrease) in the atomic percentages of Se and Te (Ge and Sb) in a resulting film. Their atomic percentages match those of a parent alloy when the thiol-EDA mixing ratio becomes 50\% to 50\%. This indicates that the dissolutions of Se/Te and Ge/Sb are likely facilitated using thiol and EDA, respectively. To ensure that resulting films maintain the parent alloy stoichiometry in a spatially uniform manner, thiol and EDA are required to be homogeneously mixed. The top $x$ axis of Fig. 5(d) shows intersolvent solubility as a function of thiol-EDA mixing ratio where H and S correspond to homogeneous and separated, respectively, indicating the presence of an immiscibility zone across 20\% to 40\% volume percentages of thiol to avoid. The $y$ axis of Fig. 5(d) shows the maximum solute loading level within solutions with varying thiol-EDA mixing ratios where $Y$ and $N$ correspond to yes (complete dissolution) and no (partial dissolution), indicating the maximum loading level achieved at a thiol-EDA mixing ratio of 50\% to 50\%, which facilitates material transfer throughput from a parent alloy to a coated film. Therefore, the 50\% to 50\% ratio, represented by the green window in Fig. 5(d), is an optimum condition, which would induce the stoichiometry of resulting films to be spatially homogeneous and consistent with that of a parent alloy while maximizing a film deposition rate.

To assess the phase change behavior of SD films, Ge$_2$Sb$_2$Se$_4$Te$_1$ films underwent \textit{in situ} grazing-incidence x-ray diffraction (GIXRD) at elevated temperatures ranging from room temperature to 325°C in an Ar environment, as shown in Fig. 6(a). The GIXRD data collected at room temperature show weak yet noticeable peaks corresponding to Te crystals. Those collected at elevated temperatures up to 325°C show the evolution of peaks corresponding to hexagonal Sb$_2$Te$_3$ crystals, which give rise to an increase in refractive index of the film,\textsuperscript{19} whereas the Te peaks are carried over to high-temperature data. Given the fact that \textit{in situ} GIXRD data collected
from thermally deposited Ge$_2$Sb$_2$Se$_4$Te$_1$ films show Sb$_2$Te$_3$ peaks with a relatively higher intensity, Te crystals existing in SD films are estimated to partially suppress the formation of desirable Sb$_2$Te$_3$ crystals upon heat treatment. This necessitates a method to reduce inherent Te crystals in starting films, thereby promoting the formation of desirable Sb$_2$Te$_3$ crystals. To achieve this, Te content was reduced from Ge$_2$Sb$_2$Se$_4$Te$_1$ to Ge$_2$Sb$_2$Se$_4.5$Te$_{0.5}$ in the parent alloys. Figure 6(b) shows the EDS-extracted atomic percentages of constituent elements on various locations of parent alloy, a coated film with the adjusted composition of Ge$_2$Sb$_2$Se$_4.5$Te$_{0.5}$, and a picture of the sample in the inset. Here, the uniform film coverage, spatial composition homogeneity, and compositional match from a parent alloy to a coated film are apparent, which are further cross confirmed by homogeneous elemental maps at a micron scale, as shown in Fig. 6(c). Figure 6(d) shows that the GIXRD data collected from the SD Ge$_2$Sb$_2$Se$_4.5$Te$_{0.5}$ films clearly indicate no presence of Te crystals in their as-coated form, while the intensity of Sb$_2$Te$_3$ peaks is observed to be stronger and sharper, indicating a higher volume fraction and uniform size distribution of the crystals as compared to those from Ge$_2$Sb$_2$Se$_4$Te$_1$ films. The IR transmission spectra (not corrected for Fresnel loss) of the SD films are shown in Fig. 6(e) where both as-coated and heat-treated films exhibit broadband transparency from mid-wave to long-wave IR. However, the short-wave IR transparency noticeably decreases upon heat treatment likely due to Sb$_2$Te$_3$ crystals-induced optical scattering. Preliminary laser writing to access a tendency toward laser-induced crystallization was carried out on the as-coated film, and visible arrays of patterns on film surfaces illustrate the films’ photo sensitivity and phase change, as shown in Fig. 6(f).

In summary, the approach which demonstrates SD Ge$_2$Sb$_2$Se$_4.5$Te$_{0.5}$ films as a candidate PCM is presented as a route toward conformal PCM media. Such strategies are expected to greatly increase design flexibility providing new opportunities for optical applications such as reconfigurable metalenses and other advanced devices. As shown, the process’ scalability and applicability to diverse composition and substrate material systems is also advantageous. Finally, further insight into the switching behavior of the SD films to understand the limitations in refractive index and electrical conductivity modifications upon either heat treatment or laser exposure will extend integration potential in advanced devices.
4 Conclusions

In this paper, recent advances in material response, methods, and design strategies aimed at improving nanoscale modifications and writing in ChG IR-transparent materials are presented. Multiphoton lithography and etch processing conditions were optimized for creating microstructures in ChG As$_2$S$_3$ films. By incorporating an AR layer between the ChG As$_2$S$_3$ fabrication layer and the Si substrate, the standing wave interference was significantly reduced, which resulted in high-quality pillar structures with more controllable dimensions. Gradient effective index nanostructured lenses with diameters of 100 μm were fabricated and found to be optically functional. Processing and characterization of high-performance Ge-Sb-Se-Te (GSST) PCM alloys is also demonstrated. TSPL was shown to be a good alternative to standard AFM techniques as well as a route to the creation of high-resolution nanostructures by overcoming the optical diffraction limit associated with laser writing in GSST materials. This technique enabled simultaneous in situ patterning and AFM imaging and does not require a closed circuit, making it compatible with a diverse range of substrates. Using TSPL, nanoscale patterning of the ChG PCM GSST with features down to 100 nm was realized. Solution-deposited ChG GSST films were shown to be a good route toward low-loss phase-change optical materials on large-area nonplanar platforms. Switching performance and prospects of this system as a functional metalens are also explored with laser exposures. These developments are of importance to the IR materials community as a route to next-generation optical and photonic systems.

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