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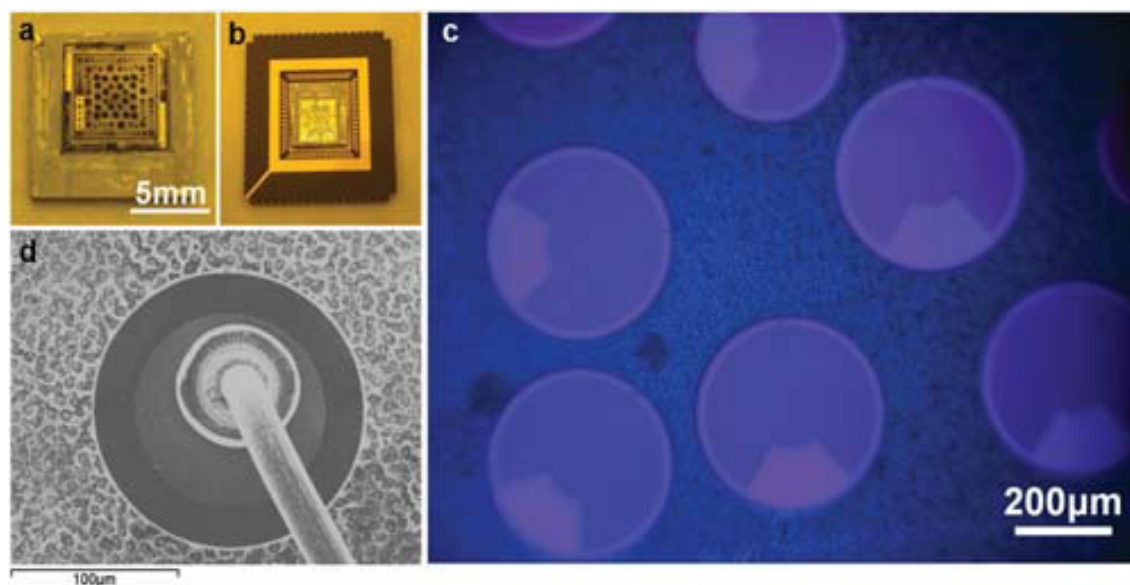
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(a) An array of single-element LWIR $\text{InAs}/\text{AlAs}_{1-x}\text{Sb}_x/\text{InAs}_{1-x}\text{Sb}_x$ T2SL photodetectors on Si after the standard processing. (b) Wire-bonded pixels in a 68-pin leadless ceramic chip carrier. (c) An optical photo of transferred LWIR T2SL photodetectors on Si. (d) A SEM image of a wirebonded transferred device ($150\ \mu\text{m}$) ready for measurement. (Zhang *et al.*, Article #4000207).

Thin-Film Antimonide-Based Photodetectors Integrated on Si

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Abstract—Monolithic integration of antimonide (Sb)-based compound semiconductors on Si is in high demand to enrich silicon photonics by extending the detection range to longer infrared wavelengths. In this paper, we have demonstrated the damage-free transfer of large-area ($1 \times 1 \text{ cm}^2$) narrow-bandgap Sb-based type-II superlattice (T2SL)-based thin-film materials onto a Si substrate using a combination of wafer-bonding and chemical epilayer release techniques. An array of Sb-based T2SL-based long-wavelength infrared (LWIR) photodetectors with diameters from 100 to 400 μm has been successfully fabricated using standard "top-down" processing technique. The transferred LWIR photodetectors exhibit a cut-off wavelength of $\sim 8.6 \mu\text{m}$ at 77 K. The dark current density of the transferred photodetectors under 200 mV applied bias at 77 K is as low as $5.7 \times 10^{-4} \text{ A/cm}^2$ and the $R \times A$ reaches $66.3 \Omega \cdot \text{cm}^2$, exhibiting no electrical degradation compared with reference samples on GaSb native substrate. The quantum efficiency and peak responsivity at 6.75 μm (@77 K, 200 mV) are 46.2% and 2.44 A/W, respectively. The specific detectivity (D^*) at 6.75 μm reaches as high as $1.6 \times 10^{11} \text{ cm} \cdot \text{Hz}^{1/2}/\text{W}$ under 200 mV bias at 77 K. Our method opens a reliable pathway to realize high performance and practical Sb-based optoelectronic devices on a Si platform.

Index Terms—Type-II superlattice infrared photodetectors, epitaxial lift-off, photonics integration.

I. INTRODUCTION

LONG-WAVELENGTH infrared (LWIR) photodetectors are particularly suitable for terrestrial-based infrared imaging since emission for room temperature objects tends to peak in the 8 to 12 μm atmospheric window according to Planck's blackbody law. Over the course of the last decade, LWIR imaging has seen tremendous improvement thanks to advances in detector technologies. InAs/GaSb/AlSb Type-II superlattices (T2SLs) have emerged as a highly suitable candidate for LWIR detection due to the ease with which the bandgap can be tuned while retaining closely lattice-matched conditions [1]. They are now directly competing with the state-of-the-art technologies such as mercury cadmium telluride (HgCdTe) compounds [2], [3]. By spatially separating electrons and holes into different quantum wells, the type-II

band alignment enables full control of the electronic band structure and allows for great flexibility in the design of heterostructure devices [4]–[14]. In addition, because T2SLs are a III-V semiconductor material system, the mass production of T2SL-based imagers has the potential to leverage existing commercial III-V foundries to further drive down costs. Another T2SL advantage is its spatial uniformity, already proven in the LWIR band, which is a critical element for the yield of larger high-resolution imagers. Most recently, InAs/AlAs_{1-x}Sb_x/InAs_{1-x}Sb_x T2SLs have been introduced, and they have been shown to have significantly longer minority carrier lifetimes than more conventional InAs/GaSb/AlSb T2SLs; they are actively being pursued as an alternative to InAs/GaSb/AlSb T2SL-based infrared photodetectors and imagers [15]–[22]. Even though they share the same type-II band alignment, these two types of T2SLs exhibit differences in superlattice design. GaSb, AlSb, and InAs all belong to the 6.1Å family, which allows large scale of freedom in the variation of layer thicknesses as well as allowing a variety of novel hetero-structures such as the W-structure [23] or M-structure [24] for pMp [25] or nBn [26] device architectures. On the contrary, AlAs_{1-x}Sb_x, InAs_{1-x}Sb_x, and InAs can have very different lattice constants; thus, the constituent layer thicknesses and the antimony molar composition (x) in the AlAs_{1-x}Sb_x and InAs_{1-x}Sb_x layers need to be taken into account in order to control the strain. This is, especially, true when going to the LWIR and VLWIR regimes where a large antimony molar fraction in the InAs_{1-x}Sb_x layers is required [27]. Facing these constraints, the device designers face more challenges in the InAs/AlAs_{1-x}Sb_x/InAs_{1-x}Sb_x T2SLs material system.

Recently, significant efforts have been made to integrate III-V photonic or electronic devices on Si substrates [28]–[32]. There is no doubt that hybrid integration of Sb-based T2SL-based photodetectors on a mature Si photonic platform can extend the applications and utility of the Sb-based material system [33], [34]. Specifically, it will allow for the heterogeneous integrated LWIR detectors on Si. However, it is very difficult to obtain high-performance devices by direct epitaxial growth of Sb-based materials on Si substrates because of the large mismatches in both the lattice constants and thermal expansion coefficients [35], [36]. Therefore, it is desirable to have a simple but reliable method that can lift-off and transfer the Sb-based epi-layers while maintaining the superior performance of the devices.

In the present work, we report the hybrid integration of high-quality large-area LWIR Ga-free

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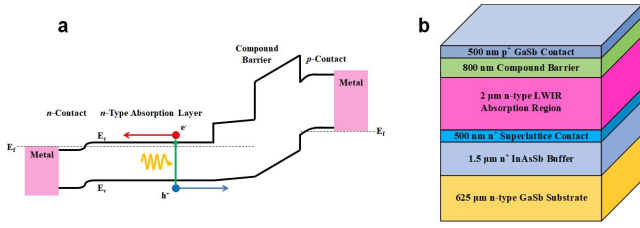


Fig. 1. Schematic diagrams of (a) the device design and (b) the epi-structure for the LWIR InAs/AlAs_{1-x}Sb_x/InAs_{1-x}Sb_x T2SL photodetectors.

InAs/AlAs_{1-x}Sb_x/InAs_{1-x}Sb_x T2SL thin films and photodetectors onto Si substrates using an epitaxial lift-off method assisted by wafer-bonding techniques. We first introduce the experimental procedures which include the device design and epitaxial growth structure used for the transfer as well as the thin-film transfer process and photodetector fabrication process. Then we will present the detailed characterization of both optical and electrical performance of transferred thin-film photodetectors on Si. Finally, we will discuss the optical response spectrum and the observed oscillations by using simulations and theoretical calculations.

II. EXPERIMENTAL PROCEDURE

A. Device Design and Materials Growth

Typical device designs for infrared photodetectors utilize a single electron barrier layer of constant bandgap, whose band structure is specifically chosen to produce a near-zero valence-band offset in order to obtain high-efficiency collection of photo-generated minority carriers. In such a device, the depletion region extends to the narrow-bandgap absorption region which is an important source of generation-recombination dark current. The photodetectors for this study have a C_pDBn architecture (shown in Fig. 1(a)). It consists of an n-doped LWIR T2SL-based absorption region (n) and a large-bandgap p-doped GaSb top contact (C_p). Adjacent to the top contact is a thin double electron barrier (DB) which has zero valence band discontinuity with respect to the n-type absorption region. This compound electron barrier is designed to only allow transport of minority carriers (holes) from the absorption region to the top contact; it blocks hole transport in the other direction with a step barrier. To ensure flat-band conditions in the absorber, a compound barrier was used in this experiment such that the electric field in the absorber can be tailored by adjusting the doping in the barrier layer located adjacent to the absorber. This results in a device with reduced dark current and increased sensitivity [37]. The compound barrier consists of a large-bandgap barrier (>1.2 eV) adjacent to the GaSb p-contact and a second smaller electron barrier layer (~300meV) between the first barrier layer and the absorber layer.

The LWIR superlattice absorption region design consists of 26/7 mono-layers (MLs) of InAs/InAs_{0.52}Sb_{0.48}, respectively, per period with a ~8 μm nominal cut-off wavelength at 77K. The small-bandgap electron barrier design consists of 4/3/4/3/4/9 MLs of InAs/AlAs/InAs/AlAs/InAs/InAs_{0.52}Sb_{0.48}, respectively, per period with a nominal cut-off wavelength of ~4 μm.

We call this structure a saw-tooth superlattice. An AlAs_{0.52}Sb_{0.48}/GaSb superlattice is used as the large-bandgap electron barrier. The addition of this second barrier pushes part of the depletion region into the larger bandgap region, decreasing the generation-recombination (G-R) current. The larger bandgap barrier also helps suppress surface band-bending and associated surface leakage [37]. Ternary AlAs_{0.52}Sb_{0.48} was used in the superlattice design because it is lattice-matched to the GaSb substrate and has antimony (Sb) atoms in common with GaSb; this provides a great deal of flexibility in the superlattice design and eliminates the need for any special interface design or strain balancing. Because of the common-anion rule of band lineups, the valence-band offset between AlAs_{0.52}Sb_{0.48} and GaSb is small, which is the desired situation to allow smooth hole transport. The electron quantum well in an AlAs_{0.52}Sb_{0.48}/GaSb superlattice is deep (~1.19 eV), however, which allows for large bandgaps in the range of ~0.8 to ~1.6 eV. Furthermore, using an AlAs_{0.52}Sb_{0.48}/GaSb superlattice as the electron barrier layer results in a smoother surface morphology compared to the case in which the barrier layer consists of a simple ternary AlAsSb layer. Finally, the growth conditions for AlAs_{0.52}Sb_{0.48}/GaSb superlattices are exactly the same as for the InAs/ InAs_{0.52}Sb_{0.48} absorption region, which allows for straightforward integration of these components. We call this structure an H-structure superlattice. The electron barrier H-structure superlattice design consists of 5/2 MLs of AlAs_{0.10}Sb_{0.90}/GaSb, respectively, with a bandgap energy of ~1.2 eV at 77 K.

The device was grown on a Te-doped n-type (10¹⁷cm⁻³) GaSb wafer using a solid source molecular beam epitaxy (SSMBE) reactor equipped with group III SUMO® cells and group-V valved cracker cells. The growth started with a 100 nm GaSb buffer layer to smooth out the surface, then, a 1.5 μm n-doped InAs_{0.91}Sb_{0.09} buffer layer (10¹⁸cm⁻³) was grown, which was followed by the 0.5 μm n-contact (10¹⁸cm⁻³), the 2 μm-thick n-type absorption region (10¹⁵cm⁻³), the 0.8 μm compound electron barrier, and the 0.5 μm p-contact (see Fig. 1(b)). Silicon (Si) and Beryllium (Be) were used as the n- and p-type dopants, respectively.

B. Thin-Film Transfer Procedure

Fig. 2 shows schematic diagrams of the process flow for transferring the LWIR InAs/AlAs_{1-x}Sb_x/InAs_{1-x}Sb_x T2SL photodetectors onto a Si substrate. First, the samples were degreased with ultrasonic cleaning in organic solvents. A layer of Ti/Au (40nm/120nm) was then deposited on the both the Si and InAs/AlAs_{1-x}Sb_x/InAs_{1-x}Sb_x T2SL wafers by electron-beam evaporation. The Ti/Au layer on the T2SL serves both as a bottom contact and as a metallic reflector for the incident light. Both Ti/Au layers were then covered with a 1.5 μm-thick indium film which was then thermally evaporated on this Ti/Au layer to be used as a bonding layer. After that, the InAs/AlAs_{1-x}Sb_x/InAs_{1-x}Sb_x T2SL and Si wafers were diced into 1×1 cm² and 2×2 cm² pieces,

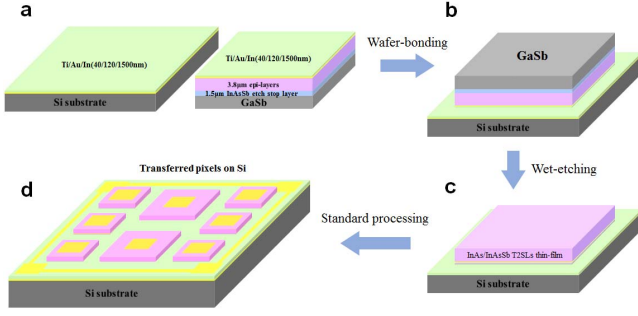


Fig. 2. Schematic diagram of the process flow for transferring the LWIR InAs/AlAs_{1-x}Sb_x/InAs_{1-x}Sb_x T2SL photodetectors to a Si substrate (a–c) and then fabricating detectors via standard processing (d).

respectively. A wafer-bonding process was then carried out for 20 min at room temperature with a force of 160 kg applied; this allows the wafers to be bonded firmly. After wafer bonding, wax was applied around the periphery of the InAs/AlAs_{1-x}Sb_x/InAs_{1-x}Sb_x T2SL wafer to protect the indium and edges of the wafer while leaving the back of the GaSb substrate exposed. The protected hybrid-wafer was then emerged into a CrO₃/HF etching solution to remove the GaSb substrate. This etchant offers excellent etch selectivity between GaSb substrate and InAs_{1-x}Sb_x buffer layer, which effectively terminates the wet-etching process at the substrate/buffer layer interface. After the GaSb substrate was totally removed, the wafer was taken out and rinsed in deionized (DI) water and the wax was removed with hot trichloroethylene (TCE) (80 °C). The wafer was then cleaned in an ultrasonic bath with acetone and then blown dry with nitrogen. After this GaSb substrate removal process, the transferred thin-film was assessed using high resolution X-ray diffraction (HR-XRD) and atomic force microscopy (AFM).

C. Device Fabrication and Testing

The transferred T2SL material was then processed into single-element InAs/AlAs_{1-x}Sb_x/InAs_{1-x}Sb_x T2SL unpassivated circular and square-shaped photodetectors with sizes ranging from 100 × 100 μm² to 400 × 400 μm² using standard processing techniques, including a two-step (dry followed by wet etching) mesa definition process and deposition of metallic top/bottom electrodes (Ti/Au) [19], [20]. The two-step mesa definition process consists of dry-etching via inductively-coupled plasma (ICP) in an Oxford Plasma Lab System 100, followed by a 90 second citric acid-based wet-etch to remove the ion-induced damage and surface traps on the pixel sidewalls. Silicon pieces containing processed detector pixels were placed in 68-pin leadless ceramic chip carriers (LCCC), and individual photodetectors were wire bonded for testing. Chip carriers were then loaded into a cryostat for both optical and electrical characterizations at 77 K. The optical characterization of these LWIR photodetectors was performed under front-side illumination with no anti-reflection coating. A Bruker IFS 66v/S Fourier transform infrared spectrometer (FTIR) was used to measure the spectral response and a calibrated blackbody source at 1000 °C was used to calibrate the responsivity and quantum efficiency.

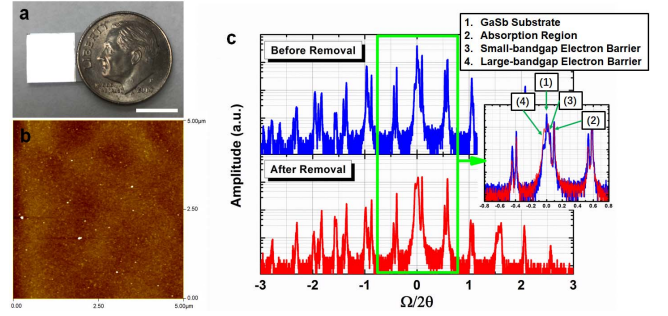


Fig. 3. (a) A 1 × 1 cm² InAs/AlAs_{1-x}Sb_x/InAs_{1-x}Sb_x T2SL-based thin-film transferred onto a Si substrate (The scale bar is 1 cm and a US dime is also shown for size comparison). (b) Atomic force microscopy of a 5 × 5 μm² surface area of the transferred material with an RMS roughness value of 4.47 Å. (c) High-resolution X-ray diffraction of the materials before/after the transfer (inset: detail view showing just the zero-order peaks of the T2SLs, compound barriers and GaSb substrate/contact layer).

III. RESULTS AND DISCUSSIONS

A large-area (1 × 1 cm²) InAs/AlAs_{1-x}Sb_x/InAs_{1-x}Sb_x T2SL-based thin-film transferred to a Si substrate is shown Fig. 3(a), with a US dime shown for size comparison. The Si surrounding the transferred Sb-based thin-film has been removed via dicing. The mirror-like surface of the transferred thin-film is continuous and crack-free, which shows the excellent quality of the wafer-bonding and chemical substrate removal. Atomic force microscopy shows a good surface morphology with a root mean squared (RMS) roughness of 4.47 Å over a 5 × 5 μm² area (see Fig. 3(b)). High-resolution X-ray diffraction (HR-XRD) scans show that the films are almost identical before and after transfer (see Fig. 3(c)). The position of all the peaks for the transferred thin-film are completely consistent with the peaks from the as-grown thin-film, suggesting that no relaxation has occurred after the transfer process. The inset of Fig. 3(c) overlays both scans in a narrow range and shows the expected decrease in the zero-order GaSb peak intensity due to the GaSb substrate having been completely removed during the substrate removal process. The remaining GaSb peak after transfer is due to the GaSb p-contact layer. The lattice mismatch to the GaSb substrate of absorption region, small-bandgap electron barrier, and larger bandgap barrier is -1500, -480, and 490 ppm, respectively, as is expected.

Fig. 4(a) shows an optical image of an array of InAs/AlAs_{1-x}Sb_x/InAs_{1-x}Sb_x T2SL-based pixels on Si after processing. Even with 1 hour of ultrasonic cleaning, there is no loss in the number of hybridized pixels with sizes ranging from 100 × 100 μm² to 400 × 400 μm², demonstrating that the transferred pixels are bonded very well to the Si substrate. This is significant for the integration of IR imaging devices onto a Si platform, which will require an enormous number of micro-pixels for high resolution. Moreover, the sidewalls of the transferred pixels are totally fresh and clean after the mesa definition and cleaning process, suggesting that good electrical properties can be obtained. Fig. 4(c) shows a closeup view of the mesa-isolated LWIR InAs/AlAs_{1-x}Sb_x/InAs_{1-x}Sb_x T2SL photodetectors on Si substrate. A SEM image of a 150 μm wire-bonded pixel on the chip carrier is presented in Fig. 4(d).

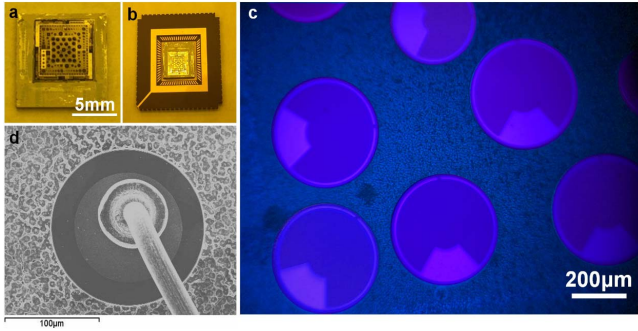


Fig. 4. (a) An array of single-element LWIR InAs/AlAs_{1-x}Sb_x/InAs_{1-x}Sb_x T2SL photodetectors on Si after the standard processing. (b) Wire-bonded pixels in a 68-pin leadless ceramic chip carrier. (c) An optical photo of transferred LWIR T2SL photodetectors on Si. (d) A SEM image of a wire-bonded transferred device (150 μm) ready for measurement.

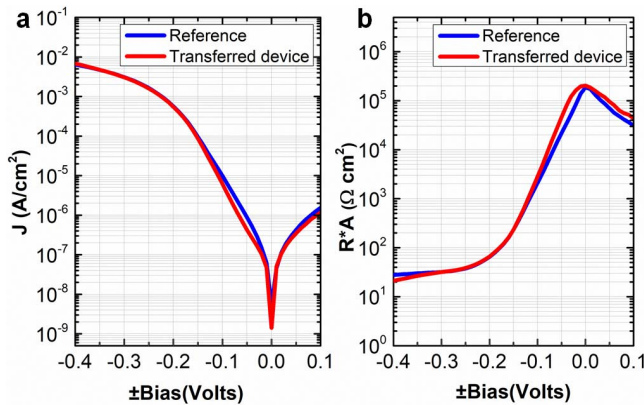


Fig. 5. (a) Dark current density and (b) differential resistance \times area ($R \times A$) product vs. applied bias voltage of a typical transferred photodetector on Si and a reference sample on native GaSb substrate at 77 K.

Fig. 5(a)-5(b) shows the current-voltage characteristic and differential resistance \times area ($R \times A$) product of the transferred LWIR T2SL photodetectors on Si and reference sample on native GaSb substrate at 77K. Unlike other transfer methods resulting in conspicuous degradation in the electrical performances, [38], [39] by using the proposed epitaxial liftoff (ELO) method in this work, the dark-current characteristics of the transferred LWIR T2SL photodetectors on Si are almost identical to that of reference samples on native substrate, demonstrating that the electrical properties do not degrade after the InAs/AlAs_{1-x}Sb_x/InAs_{1-x}Sb_x T2SL photodetectors are transferred. Specifically, the dark current density of the transferred LWIR T2SL photodetectors at a forward bias of 200mV is as low as 5.7×10^{-4} A/cm² and the corresponding $R \times A$ product reaches 66.3 $\Omega \cdot \text{cm}^2$. The low dark current density is attributed to the good cleaning of the sidewalls during the processing. Because of the flipped structure, the reference devices are driven in the opposite polarity to the transferred devices, but the bias has been reversed in the graphs to make a better comparison.

Fig. 6 shows the quantum efficiency for the transferred photodetectors and reference sample on the native GaSb substrate. The cut-off wavelength is around 8.6 μm , which is consistent with the material design. The peak quantum

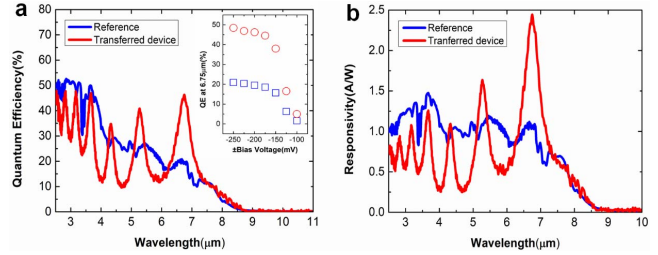


Fig. 6. (a) Saturated quantum efficiency spectra of the transferred device and reference sample at 200mV applied bias voltage in front-side illumination configuration without any anti-reflection coating. Inset shows quantum efficiency of the devices at 6.75 μm in front-side illumination configuration as a function of applied bias voltage.

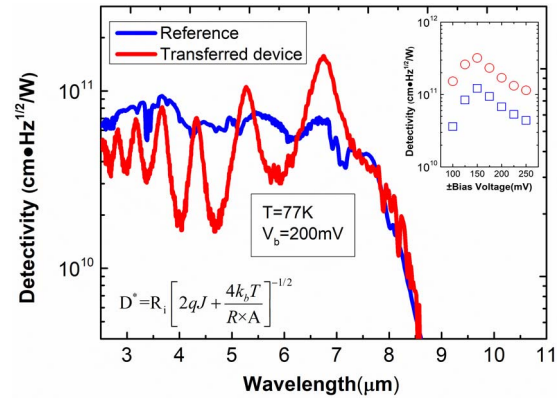


Fig. 7. Specific detectivity (D^*) spectrum of the transferred devices on Si and of a reference sample on native GaSb substrate at 200 mV applied bias voltage, at 77K. Top-right inset shows the detectivity versus bias at 6.75 μm . The specific detectivity is calculated based on the equation in the bottom-left inset, where R_i is the device responsivity, J is the dark current density, $R \times A$ is the differential resistance \times area product, k_b is the Boltzmann constant, and T is the operating temperature.

efficiency and responsivity (@200mV) emerging at a wavelength of 6.75 μm are about 46.2% and 2.44 A/W, respectively, which are 2.3 times higher than that of the reference sample. As the bias increases from 200 to 250mV, a maximum quantum efficiency of $\sim 49\%$ is obtained.

After performing both optical and electrical characterization, the specific detectivity (D^*) was calculated for the transferred LWIR photodetectors at 77K. Fig. 7 shows the calculated detectivity for the transferred LWIR T2SL photodetectors on Si and for a reference sample on native GaSb substrate. The peak detectivity at 6.75 μm is 1.6×10^{11} $\text{cm} \cdot \text{Hz}^{1/2} / \text{W}$ at a forward bias of 200 mV, which is 2.4 times higher than that of the reference sample. The maximum detectivity reaches as high as 3.2×10^{11} $\text{cm} \cdot \text{Hz}^{1/2} / \text{W}$ at a forward bias of 150mV, as shown in the top-right inset of Fig. 7.

For these substrate-transferred thin-film Sb-based photodetectors, there is a clear oscillation throughout the optical response spectrum due to Fabry-Perot resonances within the thin-film device. Fig. 8(a) shows the photoresponse and simulated transmission spectra from a 3.81 μm -thick pixel transferred on Si. The height of 3.81 μm is obtained from the measurement with an optical profiler, as shown in inset of Fig. 8(a). The oscillations observed in the photoresponse spectra are consistent with the Fabry-Perot modes confined

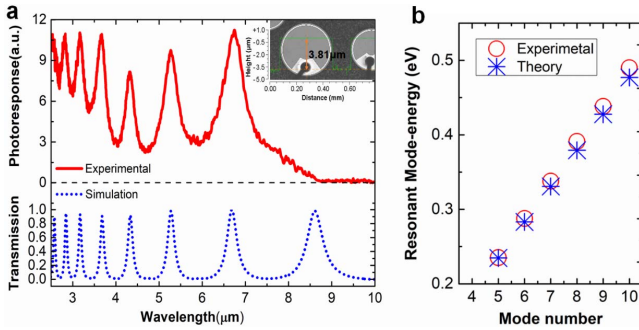


Fig. 8. (a) Photoresponse and simulated transmission spectra from a $3.81\mu\text{m}$ -thick pixel transferred on Si. (b) Experimental photoresponse and calculated transmission oscillations for incoming light. Inset shows the height of the transferred pixels from optical profilometry.

within $\text{InAs}/\text{AlAs}_{1-x}\text{Sb}_x/\text{InAs}_{1-x}\text{Sb}_x$ T2SL thin-film photodetectors on Si from the two-dimensional finite-difference time-domain (2D-FDTD) simulation. Here, the 2D-FDTD optical simulations were carried out for the Sb-based T2SL thin-film structure, which has a measured thickness of $3.81\mu\text{m}$ and a width of $100\mu\text{m}$. A plane-wave with a wavelength range of $2\text{--}10\mu\text{m}$ was used as the incident light source, which is placed $15\mu\text{m}$ from the Sb-based materials. A frequency domain field and power monitor was placed inside the $\text{InAs}/\text{AlAs}_{1-x}\text{Sb}_x/\text{InAs}_{1-x}\text{Sb}_x$ T2SL thin-film to check the transmission spectra. A Ti/Au ($40\text{nm}/150\text{nm}$) layer sequence was placed at the bottom of the Sb-based T2SL thin-film. The simulation domain was set with a width of $100\mu\text{m}$ and with $50\mu\text{m}$ in thickness, with periodic boundary conditions and perfectly matched layers. The effective refractive index of the Sb-based T2SL was chosen from fitting experimental data. No optical absorption was considered in the simulations. The confinement originates from the reflections that occur between the Ti bottom contact and the InAsSb/air interface. In general, the resonant wavelengths for different mode numbers ($m=4, \dots, 10$) can be calculated with the relationship:

$$m\lambda/2 = n(\lambda) \times L,$$

where $n(\lambda)$ is the dispersive refractive index and L is the thickness of the T2SL photodetectors transferred onto Si. Fig. 8(b) shows that the calculated transmission peaks match well with the experimental results, which further confirms that the oscillations in the photoresponse spectra are attributed to Fabry-Perot resonances. It is worth noting that these Fabry-Perot oscillations will be reduced and a nearly flat response spectrum from these substrate-transferred on-chip thin-film photodetectors can be expected if a broadband anti-reflection coating is applied to the InAsSb/air interface. In addition, one probable reason that it cannot be observed such enhancement at shorter wavelengths is due to the Ti/Au reflector at the bottom has a lower reflectivity for the short-wavelength infrared range. Therefore, the performance of these substrate-transferred detectors will be further improved with better optical designs to reduce the loss at the metallic contact interface with the Sb-based materials and application of an anti-reflection coating to the top surface.

IV. CONCLUSION

In summary, an array of high-performance LWIR Ga-free $\text{InAs}/\text{AlAs}_{1-x}\text{Sb}_x/\text{InAs}_{1-x}\text{Sb}_x$ T2SL photodetectors were successfully integrated onto Si substrate through wafer bonding of $1\times 1\text{ cm}^2$ thin films, substrate removal, and standard device processing. These transferred devices displayed a cut-off wavelength of $8.6\mu\text{m}$ at 77 K . At the peak response of $6.75\mu\text{m}$, the transferred photodetector revealed a quantum efficiency and responsivity of 46.2% and 2.44 A/W at 200 mV bias, respectively, when it was under front-side illumination and with no anti-reflection coating. The dark current density of the transferred photodetectors under 200 mV bias at 77 K reached $5.7\times 10^{-4}\text{ A/cm}^2$, exhibiting no electrical degradation compared to reference samples on native GaSb substrates. The devices also demonstrated a saturated dark current shot noise limited specific detectivity of $1.6\times 10^{11}\text{ cm}\cdot\text{Hz}^{1/2}/\text{W}$ at $6.75\mu\text{m}$, under 200 mV bias, at 77 K . Being compatible with traditional semiconductor processing techniques, this reliable large-area transfer method and high-yield fabrication of high-performance Sb-based T2SL micro-pixel photodetectors on Si substrates will allow new applications for this material system. This technology offers a straightforward, yet simple, technique for hybrid integration of Sb-based T2SL photodetectors with silicon readout integrated circuits (ROIC) for focal plane array (FPA) imaging applications. In addition, this work also presents a feasible solution for heterogeneous integration of high-performance Sb-based electronic devices like high electron mobility transistors (HEMTs) and metal-oxide-semiconductor field-effect transistors (MOSFETs) on a Si microelectronics platform.

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