Heterogeneously-Grown Tunable Tensile Strained Germanium on Silicon for Photonic Devices

Michael Clavel,‡ Dzianis Saladukha,‡§ Patrick S. Goley,‡ Tomasz J. Ochalski,‡§ Felipe Murphy-Armando,‡ Robert J. Bodnar,¶ and Mantu K. Hudait*†

‡Advanced Devices & Sustainable Energy Laboratory (ADSEL), Bradley Department of Electrical and Computer Engineering, Virginia Tech, Blacksburg, Virginia 24061, United States
§Tyndall National Institute, Lee Maltings, Dyke Parade, Cork, Ireland
¶Centre for Advanced Photonics and Process Analysis, Cork Institute of Technology, Cork, Ireland
*Electrical Power Engineering Institute, Department of Electrical Engineering, Virginia Tech, Blacksburg, Virginia 24061, United States

ABSTRACT: The growth, structural and optical properties, and energy band alignments of tensile-strained germanium (ε-Ge) epilayers heterogeneously integrated on silicon (Si) were demonstrated for the first time. The tunable ε-Ge thin films were achieved using a composite linearly graded In,Ga1−As/GaAs buffer architecture grown via solid source molecular beam epitaxy. High-resolution X-ray diffraction and micro-Raman spectroscopic analysis confirmed a pseudomorphic ε-Ge epitaxy whereby the degree of strain varied as a function of the In,Ga1−As buffer indium alloy composition. Sharp heterointerfaces between each ε-Ge epilayer and the respective In,Ga1−As strain template were confirmed by detailed strain analysis using cross-sectional transmission electron microscopy. Low-temperature microphotoluminescence measurements confirmed both direct and indirect bandgap radiative recombination between the Γ and L valleys of Ge to the light-hole valence band, with L-hh bandgaps of 0.68 and 0.65 eV demonstrated for the 0.82 ± 0.06% and 1.11 ± 0.03% strained Ge on Si, respectively. Type-I band alignments and valence band offsets of 0.27 and 0.29 eV for the ε-Ge/In0.17Ga0.83As (0.82%) and ε-Ge/In0.17Ga0.83As (1.11%) heterointerfaces, respectively, show promise for ε-Ge carrier confinement in future nanoscale optoelectronic devices. Therefore, the successful heterogeneous integration of tunable tensile-strained Ge on Si paves the way for the design and implementation of novel Ge-based photonic devices on the Si technology platform.

KEYWORDS: germanium, optical domain, efficient cost-effective light sources, laser integration, molecular beam epitaxy, heterostructure integration

INTRODUCTION

The shrinking feature size of silicon (Si) transistors has enabled an exponential increase in transistor density, resulting in increased compute power. However, current Si-based complementary metal-oxide-semiconductor (CMOS) technology is nearing the physical limits of its scaling potential, and with the end in sight of the traditional technology roadmap, only a radical departure from Si-based technologies can ensure continued technological progress. New material innovations, novel device architectures, and heterogeneous technology cointegration, new functionalities, and their monolithic integration onto Si are projected to continue transistor miniaturization beyond the Si CMOS era. Moreover, interconnect bottlenecks for both interchip and intrachip communication are projected to be major impediments to energy-efficient performance scaling. Therefore, there is an urgent need for large bandwidth, low-resistance interconnects in high-end computing applications as copper-based electrical interconnects are rapidly becoming inefficient in meeting essential bandwidth requirements. It will become increasingly challenging to transmit signals electrically while maintaining low power consumption, low delay, and a high signal-to-noise ratio. An enticing approach would be the integration of photonic devices with Si technology, hence the monolithic integration of Si-based optoelectronics would be an obvious choice. However, the indirect bandgap of Si limits the realization of Si-based photonic devices. Thus, the hybrid integration of germanium (Ge) and III–V materials-based optoelectronic devices with traditional Si CMOS technology would revolutionize technology needs in the near future. The superior transport properties and large modulation bandwidth of Ge and III–V compound semiconductor material systems make them ideal candidates for integration on Si. Besides, Ge-based light sources on Si are important for the realization of future, nanoscale optical on-chip communication. Furthermore, optical interconnects compatible with Si process technology are needed to provide for ever-increasing future bandwidth needs. Moreover, the electrical-to-optical inter-
connect transition for chip-to-chip communication is expected to be a gradual process depending upon specific application requirements and cost-performance trade-offs with current copper-based interconnect technologies.26

Currently, the III-V laser bonded to Si waveguide approach27,28 imposes strict design and process constraints, with limited scalability of size (because of the evanescently coupled lasing mode) and cost (because of III-V wafer area utilization in large channel count transceivers). A heterogeneously grown active material with the scope for a tunable wavelength laser would alleviate these constraints and provide a more scalable solution as device size decreases and integration density increases. Very recently, lasing from tensile strained direct-bandgap Ge1−xSnx alloys on Si3 and its potential applications in optoelectronics as well as transistors30,31 has gained widespread research interest due to the modulation of the Ge1−xSnx bandgap both by tin (Sn) incorporation and strain modulation. The direct bandgap Ge1−xSnx (x > 0.1) material can be used as a gain medium in tunable wavelength lasers due to the modulation of the direct Γ valley bandgap for increasing Sn compositions; however, the thermal stability of Ge1−xSnx because of Sn segregation11 as well as the lower Ge1−xSnx growth temperature imposes great difficulties for process integration with state-of-the-art Si device fabrication processes. Therefore, there is a critical need to develop a material system that will be compatible with industry-standard Si process flows, including higher process temperature stability, and provides a feasible template for strain-engineered bandgap modulation for on-Si light sources. Furthermore, light sources with directly grown active materials are needed for the large-scale integration of highly integrated photonic devices for optical interconnects in future high-capacity datacenters, warehouse-scale computing, and high-performance computing applications. In this paper, we have demonstrated a heterogeneous integration scheme for tunable tensile-strained Ge (ε-Ge) on Si using variable surface-terminated lattice constant buffering of III-V compound semiconductor metamorphic layers that shows great promise for realizing modular group-IV based lasers on Si. In our hybrid Ge and III-V on Si integration approach, we have simultaneously combined a substrate architecture that offers a greater design flexibility with diverse material choices to produce scalable, heterogeneously integrated tunable light sources on Si that target the possibility of realizing hybrid electronic and photonic systems on Si for ultralow power computing, interchip communication, and optoelectronic applications. We are able to achieve these attributes by carefully controlling the modular tensile strain in the Ge epitaxial layers during growth, and in tandem with the strain analysis, defect examination, morphological, and band alignment properties of these strained Ge material systems, provide a path to achieve group-IV-based lasers on Si.

RESULTS AND DISCUSSION

Strain Analysis via XRD. Figures 1a and 1b show the schematics of the ε-Ge/In0.17Ga0.83As heterostructures that were heterogeneously grown on off-cut (100) Si substrates and studied in this work. For the 1.11% ε-Ge sample, a 15 nm constant composition In0.17Ga0.83As capping layer was grown in order to study the materials and optical properties of In0.17Ga0.83As cladded ε-Ge optical cavities. The strain relaxation properties of these structures were investigated using high-resolution X-ray diffractometry (HR-XRD). Figure 2 shows the high-resolution triple axis symmetric (004) X-ray rocking curves (RCs) from the 30 to 40 nm ε-Ge epilayers grown on (i) 750 nm In0.11Ga0.89As/1.4 μm graded In0.35Ga0.65As/2 μm GaAs (0.82% tensile strain, green) and (ii) 600 nm In0.17Ga0.83As/1.9 μm graded In0.35Ga0.65As/2 μm GaAs (1.11% tensile strain, blue) metamorphic buffers on Si, respectively. Additionally, each diffraction peak in Figure 2 is labeled with its corresponding epilayer. Likewise, Figure 3 shows the symmetric (004) and asymmetric (115) RSMs of the ε-Ge/In0.11Ga0.89As on Si structure, while Figure 4 similarly shows the (004) and (115) RSMs taken from the ε-Ge/In0.11Ga0.89As on Si structure.

Figure 1. Cross-sectional schematic of (a) 0.82% ε-Ge/In0.11Ga0.89As and (b) 1.11% ε-Ge/In0.17Ga0.83As structures heterogeneously integrated onto Si.

Figure 2. X-ray rocking curves from the 0.82% (green) and 1.11% (blue) ε-Ge on Si structures showing movement of the Ge Bragg angle with increasing tensile strain. Inset demonstrates the effect of heteroepitaxial mismatch on the in-plane lattice constant of the top-lying epilayer.
Also shown in Figures 3 and 4 are the (004) and (115) RSMs recorded for each respective metamorphic buffer structure after removal of the ε-Ge (In_{0.17}Ga_{0.83}As/ε-Ge) topmost epilayer(s) using dilute NH_4OH:H_2O_2:H_2O (2:1:200 volume ratio). For all RSMs, each layer has been labeled to its corresponding reciprocal lattice point (RLP) for clarity. One can find from the symmetric (004) RCs in Figure 2 and RSMs in Figures 3a and 4a that the Ge RLP exhibits a larger Bragg angle as compared to the GaAs buffer RLP, indicating a smaller out-of-plane lattice constant and thus the presence of tensile strain in the Ge epilayer. As shown in the inset of Figure 2, because of the larger lattice constant of the In_{0.11}Ga_{0.89}As (x = 0.11, 0.17) strain template, the in-plane lattice constant of the heteroepitaxial Ge is stretched to accommodate the in-plane lattice constant of the In_{0.11}Ga_{0.89}As layer, resulting in a biaxial tensile-strained, pseudomorphic Ge epilayer with an expanded in-plane lattice constant (labeled as a) and reduced out-of-plane lattice constant (labeled as c). One can find from Figures 2–4 that...
the diffraction peak and RLP of ϵ-Ge is partially suppressed by the intensity of the GaAs substrate as a result of minor Bragg angle modulation due to the moderate strain levels studied in this work. To mitigate error in the experimental strain relaxation analysis resulting from the partial superposition of the ϵ-Ge and GaAs diffraction peaks, only the centroid of the ϵ-Ge (004) RLP taken from the symmetric (004) RSM was used for quantitative analysis of the out-of-plane lattice spacing. Moreover, so as to more clearly distinguish the superimposed ϵ-Ge and GaAs peaks, Figure 5 shows the symmetric (004) RSM of the ϵ-Ge/In0.11Ga0.89As structure plotted along ω-2θ (a) prior to and (b) after removal of the ϵ-Ge layer. As seen in the inset of Figure 5a, magnification of the Bragg angles surrounding the GaAs layer revealed the distinct outline of the ϵ-Ge peak, thus enabling a more accurate determination of the ϵ-Ge ω-2θ prior to conversion to reciprocal lattice units. Furthermore, Figures 3b and 3d (4b and 4d) show the (004) and (115) RSMs for the In0.11Ga0.89As/GaAs (In0.17Ga0.83As/GaAs) metamorphic buffers following wet etching of the terminating ϵ-Ge (In0.11Ga0.89As/ϵ-Ge) epilayer(s). One can find by comparing Figures 3b and 3d (4b and 4d) to Figures 3a and 3c (4a and 4c) that the ϵ-Ge RLP indeed contributes significantly to the broadening and intensity of the region in reciprocal space in proximity to the GaAs virtual substrate peak. The ability to clearly differentiate between the ϵ-Ge and GaAs diffraction peaks thus provides ancillary support for the strain-dependent shift in the ϵ-Ge RLP observed in Figures 3 and 4. Further movement of the ϵ-Ge RLP can be achieved by providing increased tensile strain to the Ge layer through growth on higher indium (In) content InGa1−xAs virtual substrates, thereby increasing the accuracy of the measured strain data.

As shown in Figures 3c and 4c by the vertical alignment of the InGa1−xAs and ϵ-Ge RLPs (orange dotted lines), the in-plane lattice constant of ϵ-Ge was found to be closely matched with the in-plane lattice constant of the In0.11Ga0.89As and In0.17Ga0.83As constant composition layers, signifying that the tensile strain transferred to the Ge was successfully modulated by varying the underlying InGa1−xAs In alloy composition. The detailed analyses of the relaxation and strain states of the epitaxial Ge and InGa1−xAs layers were evaluated by measuring the in-plane and out-of-plane lattice constants, a and c, for both Ge and InGa1−xAs using the recorded symmetric (004) and asymmetric (115) RSM data. The relaxed lattice constant (a_r) of each layer was calculated using a, c and Poisson’s ratio for each material, noting that a relaxed Ge lattice parameter of 5.658 Å was assumed.32 Table 1 summarizes the in-plane, out-of-plane, and relaxed lattice constants of the InGa1−xAs (x = 0.11, 0.17) and ϵ-Ge layers as well as the amount of strain, relaxation, and epitaxial tilt determined via these measurements. Using the relaxed lattice constant of InGa1−xAs and Vegard’s law, the In alloy composition of each InGa1−xAs virtual substrate was determined to be 11.4% and 16.6%, which was consistent with the design criteria. The GaAs and InGa1−xAs metamorphic buffers on Si were found to be ~90% and 82–87% relaxed, respectively. The detailed methodology used in the strain relaxation analysis is reported elsewhere.35 Additionally, the uncertainty in the experimental strain values for each ϵ-Ge/InGa1−xAs heterostructure was derived using the measured effect of epitaxial tilt on the calculated in-plane and out-of-plane lattice parameters and thereby the strain held by the Ge epilayer. For the measured results presented in Table 1, the data used in the strain relaxation analysis were collected from RSMs in which the tilt of the InGa1−xAs and ϵ-Ge layers was minimized with respect to the Si substrate. The resulting uncertainties were found to be ±0.06% and ±0.03% for the 0.82% and 1.11% ϵ-Ge, respectively. Consequently, the experimentally demonstrated tensile strain modulation due to increasing In alloy composition in InGa1−xAs strain templates is expected to modify the Ge bandgap, an essential step toward achieving tunable wavelength Ge-based photonic devices, as will be discussed in the microphotoluminescence (µ-PL) analysis below.
Strain Analysis via Raman Spectroscopy. For several decades, the semiconductor industry has implemented strain to boost device performance, modify material bandgaps, and enhance carrier confinement. Thus, alternative strain measurement techniques to independently confirm material stress have been important technological considerations. Most recently, Raman spectroscopy has been utilized in determining the Ge composition in Si$_{1-x}$Ge$_x$ layers for source/drain stressors in nanoscale Si CMOS. Figure 6a shows the shift in tensile stress must be accounted for when analyzing the Raman shift of ε-Ge heterogeneously integrated onto Si utilizing III–V buffer architectures. While it is well documented that the Ge–Ge phonon vibration mode is approximately 300 cm$^{-1}$ and that a shift in phonon vibration mode with respect to this number is a result of crystallographic strain, an exact relationship between strain and wavenumber shift is only approximate. Recent work has utilized the relation $\Delta \omega = -b \epsilon_t$ to analyze Raman shift as a function of strain in tensile Ge thin films, where $\Delta \omega$ is the wavenumber shift (in cm$^{-1}$) and $b$ is a material parameter dependent on the material’s phononic and elastic constants. Using the reported literature value of 415 cm$^{-1}$ for Ge, the wavenumber shift versus In alloy composition as well as strain versus In alloy composition relationships are shown in Figure 6b. One can find from Figure 6b that tensile strain (green) determined from the Raman shift (blue) corresponds accurately to the theoretical misfit (red) for the In alloy compositions studied, thereby independently confirming the pseudomorphic nature of the strained Ge epitaxy. Moreover, the strain analysis demonstrated utilizing Raman spectroscopy was found to be in good agreement with the strain relaxation properties of the ε-Ge/In$_{0.11}$Ga$_{0.89}$As heterostructures on Si as determined via X-ray diffraction analysis, thereby highlighting the suitability of Raman spectroscopy for studying ε-Ge thin films with moderate strain levels.

Defect Analysis by TEM. Additional investigation into the material quality and nature of the ε-Ge/In$_{0.11}$Ga$_{0.89}$As (x = 0.11, 0.17) heterointerface was performed by high-resolution cross-sectional transmission electron microscopy (TEM) analysis. Figures 7a–f and 8a–e show the bright-field cross-sectional TEM micrographs of the ε-Ge grown on In$_{0.11}$Ga$_{0.89}$As/In$_{0.11}$Ga$_{0.89}$As/GaAs/Si and In$_{0.11}$Ga$_{0.89}$As/In$_{0.17}$Ga$_{0.83}$As/In$_{0.11}$Ga$_{0.89}$As/GaAs/Si, respectively. Cross-sectional TEM measurements were performed on various locations of the TEM specimen and the representative results are shown in Figures 7 and 8. Moreover, all layers are labeled in each figure. The GaAs and linearly graded In$_{0.11}$Ga$_{0.89}$As buffers on Si were found to effectively mitigate the lattice mismatch induced defects and dislocations between the ε-Ge/In$_{0.11}$Ga$_{0.89}$As (x = 0.11, 0.17) layers of interest and the Si substrate. One can find from Figure 7a (8a) that the majority of dislocations were confined within the linearly graded In$_{0.11}$Ga$_{0.89}$As buffer. Moreover, the residual strain within the top In$_{0.11}$Ga$_{0.89}$As (In$_{0.17}$Ga$_{0.83}$As) layer was minimized due to the nominal accommodation of mismatch induced epitaxial strain via misfit dislocation formation and subsequent dislocation glide, corroborating the previously discussed strain relaxation properties for each In$_{0.11}$Ga$_{0.89}$As virtual substrate. Similarly, the apparent absence of threading dislocation propagation into the ε-Ge active region suggests a device-quality active region, further reinforcing the structural data found via reciprocal space mapping. Hence, the composite GaAs and linearly graded In$_{0.11}$Ga$_{0.89}$As buffer provided a high-quality virtual substrate for the tunable tensile-strained Ge heterogeneously integrated onto Si. Furthermore, high resolution lattice indexing shown in Figure 7c (8b) revealed the lattice line extending uninterrupted between the ε-Ge and In$_{0.11}$Ga$_{0.89}$As (In$_{0.17}$Ga$_{0.83}$As) epilayers, indicating that the Ge in-plane lattice constant was internally registered with the in-plane lattice constant of the In$_{0.11}$Ga$_{0.89}$As (In$_{0.17}$Ga$_{0.83}$As) strain template and demonstrating pseudomorphic tensile-stained Ge epitaxy in good agreement with the X-ray and Raman analysis presented earlier. Figures 7d–f and 8c–e show Fast Fourier Transform (FFT) patterns representative of the regions.
denoted by arrows. The indexing of these FFT patterns indicates that the electron beam was parallel to the [011] orientation. Moreover, the FFT patterns obtained from the ε-Ge/In\textsubscript{0.11}Ga\textsubscript{0.89}As heterointerface and detailed lattice line indexing (including FFT patterns) demonstrating the tensile nature of Ge with respect to the underlying In\textsubscript{0.11}Ga\textsubscript{0.89}As.

Accordingly, the cross-sectional TEM analysis suggests optical-quality film structures with atomically smooth interfaces, which is crucial to minimizing cavity losses in Ge-based photonic devices.

To further evaluate the atomic species profile across each heterointerface, energy dispersive X-ray spectroscopy (EDS) was utilized to perform an elemental mapping of the ε-Ge/In\textsubscript{0.11}Ga\textsubscript{0.89}As heterostructure. EDS is a powerful technique that has been extensively used for the analysis of the structural and chemical modulation of nanoscale heterostructures and multilayer heterostructures.\textsuperscript{41} Figure 7b shows the elemental mapping of the ε-Ge/In\textsubscript{0.11}Ga\textsubscript{0.89}As structure where the Ge layer (green) was found to be uniformly distributed and the In (blue) exhibits a variable composition, as expected in the linearly graded In\textsubscript{0.11}Ga\textsubscript{0.89}As buffer layer, in addition to the composition profile of Si (red). The EDS analysis exhibits a uniform and sharp heterointerface between the ε-Ge and the In\textsubscript{0.11}Ga\textsubscript{0.89}As layer, with no apparent interdiffusion across any of the structure’s heterointerfaces despite the increased thermal budget as a result of experiencing extended growth temperature conditions during the strained layer epitaxy of Ge. Therefore, the atomic scale imaging and microstructural analysis feedback loop is necessary for the large-scale heterointegration of tunable tensile-strained Ge on Si.

**Strain-Induced Ge Bandgap Modulation Analysis via Photoluminescence.** Photoluminescence spectroscopy (PL) has been extensively used to investigate strained-induced bandgap shifts,\textsuperscript{42} the competition between direct and indirect emission characteristics,\textsuperscript{43,44} radiative and nonradiative recombination properties,\textsuperscript{45} and the ratio between direct and indirect bandgap recombination with increasing strain in Ge.\textsuperscript{36} In this work, low-temperature micro-PL (μ-PL) measurements were performed at 7.5K utilizing an incident power density of 860 kW/cm\textsuperscript{2} and excitation wavelength of 800 nm. This high power density is required to obtain a reasonable PL emission intensity.\textsuperscript{46,47} Figure 9a shows the PL spectra obtained from 0.82% (blue) and 1.11% ε-Ge with (red) and without (green) In\textsubscript{0.17}Ga\textsubscript{0.83}As capping layer as well as quasi-lattice matched Ge on Al\textsubscript{0.17}Ga\textsubscript{0.83}As (gray). The PL spectra have been shifted vertically for clarity. One can see from Figure 9a that the main peak position is red-shifted toward lower photon energies (higher wavelengths) due to the increase in tensile strain and corresponding reduction in ε-Ge bandgaps. Furthermore, from Figure 9b and c, one can see that each sample exhibits a convolved emission spectra containing a single, intense peak.
and a shoulder whose relative intensity is strongly strain-dependent.

The deconvolution of these spectral features into accurate transition energies requires considering several coinciding factors, most importantly measurement temperature and strain incorporation. Although it has been reported that experimental collection geometries normal to the sample surface (i.e., $\hat{z}$) favor conduction band coupling with the heavy-hole (hh) valence band,\textsuperscript{36} cryogenic sample temperatures result in a rapid depopulation of the hh valence band due to its higher energy states. Consequently, radiative recombination between the conduction band minima and light-hole (lh) valence band increases as heavy-holes undergo rapid nonradiative relaxation into lower energy states within the lh valence band. The dominance of the L-lh and $\Gamma$-lh coupling at low measurement temperatures is seen experimentally through direct observation of the lower energy transitions, i.e. measurement of the lower optical energy gaps within a material, which were indeed of the lower energy transitions, i.e. measurement of the lower temperatures is seen experimentally through direct observation.

Figure 9. (a) Red-shift (plotted in photon energy) due to strain in the $\epsilon$-Ge epilayers as measured by $\mu$-PL at 7.5 K. (b and c) Gaussian-Lorentzian fitting of the $\epsilon$-Ge/In$_{0.11}$Ga$_{0.89}$As (b) and $\epsilon$-Ge/In$_{0.17}$Ga$_{0.83}$As (c) emission.

The wavelength-dependence of the absorption coefficient in Ge films as a function of strain proposed by Virgilio and co-workers is found to be in good agreement with prior experimental results,\textsuperscript{54,55} investigating the temperature- and strain-dependence of Ge optical absorption. From these previous results, one would expect that the lower energy spectral features would exhibit higher relative PL intensities when compared with the higher energy features, which is observed in Figures 9b and 9c. Nonetheless, one might anticipate that due to the low-temperature nature of the sample during optical characterization, $\Gamma$ valley recombination would dominate due to reduced phonon-assisted momentum conservation between the L- and $\Gamma$-points in momentum space. This interpretation, however, fails to address the need for additional energy in order to surmount the L-$\Gamma$ conduction band minima separation, which is in addition to the required momentum preservation. For unstrained, bulk Ge, the separation between the L and $\Gamma$ minima is approximately 150 meV. Thus, although momentum conservation may be met via exciton-generated LA phonons at cryogenic measurement temperatures,\textsuperscript{36–38} the aforementioned L-$\Gamma$ energy difference restricts radiative recombination primarily to L valley pathways for low-to-moderate strains and doping concentrations in which the L-$\Gamma$ separation remains prohibitively large. It is worth noting that previous work investigating similar $\epsilon$-Ge/In$_{x}$Ga$_{1-x}$As heterointerfaces revealed an integrated PL intensity power-dependence suggestive of exciton recombination for type-I $\epsilon$-Ge/In$_{x}$Ga$_{1-x}$As heterointerfaces. The detailed analysis of the interfacial energy band alignment for the structures studied in this work can be found in a subsequent section, suffice to say the structures indeed exhibited a type-I band alignment, thus reinforcing the notion that exciton–phonon interaction underlies the observed indirect radiative recombination.\textsuperscript{60} Moreover, as higher tensile strains are incorporated, the more rapid lowering of the $\Gamma$ valley minimum results in a similarly rapid decrease in the L-$\Gamma$ separation energy, thereby enhancing direct gap recombination as validated by the increase in relative intensity and integrated peak area of the $\Gamma$-lh transition between Figure 9b and c.

Lastly, it is generally understood that the reduction in effective mass in the lh valence band due to strain correlates with a decreasing density of states and an increasingly restricted range of transition energies. Hence, it is expected that the full-width at half-maximum (fwhm) of the more prominent L-lh transition as well as the ratio of the integrated peak areas between L-lh and $\Gamma$-lh transitions (i.e., $I_{L\text{-lh}}/I_{\Gamma\text{-lh}}$) will both decrease as the biaxial tensile strain held by the Ge epilayer increases. Figures 8b and c demonstrate that for increasing tensile strain, the fwhm of the L-lh transition indeed decreases from 54.5 meV at $\varepsilon = 0.82\%$ to 31.7 meV at $\varepsilon = 1.11\%$. Likewise, the ratio of the L-lh to $\Gamma$-lh integrated peak areas also decreases from 1.95 at $\varepsilon = 0.82\%$ to 0.30 at $\varepsilon = 1.11\%$, indicating that the increase in biaxial tensile strain strongly enhances $\Gamma$-lh recombination via significant $\Gamma$-lh bandgap reduction as well as restricts the energy range for L-lh radiative recombination. Figure 10 compares the experimentally observed bandgaps (symbols) and strain determined from $\mu$-PL and XRD analysis, respectively, with the predicted dependence of the low-temperature Ge bandgap on increasing
strain (solid lines) calculated using a 30 × 30 \( \mathbf{k} \cdot \mathbf{p} \) formalism.\(^{61}\) Additionally, the effect of decreasing \( \varepsilon \)-Ge epilayer thickness \((t_{\text{Ge}})\) on the \( \varepsilon \)-Ge bandgap because of energy level quantization is shown via the dashed \((t_{\text{Ge}} = 40 \text{ nm})\) and dotted \((t_{\text{Ge}} = 30 \text{ nm})\) lines. The measured L−lh \((\Gamma−lh)\) bandgaps were found to be 0.684 (0.735 eV) and 0.645 eV (0.704 eV) for the \( \varepsilon \)-Ge/In\(_{0.11}\)Ga\(_{0.89}\)As and \( \varepsilon \)-Ge/In\(_{0.17}\)Ga\(_{0.83}\)As structures, respectively. This is in excellent agreement with the theoretical bandgap-strain relation shown in Figure 10, noting that the quantization effect induced from decreasing \( \varepsilon \)-Ge layer thickness has a negligible impact on the measured bandgaps for the \( \varepsilon \)-Ge film thicknesses studied in this work. Furthermore, the demonstration of tunable wavelength \( \varepsilon \)-Ge epitaxial layers heterogeneously integrated onto Si is a key first step toward the monolithic integration of Ge-based photonic devices and optically active layers with state-of-the-art CMOS technology as well as the development of energy-efficient light sources for future on-chip optical interconnects.

**Energy Band Offset Determination via XPS.** Understanding of the energy band alignment between \( \varepsilon \)-Ge and the underlying constant composition In\(_{x}\)Ga\(_{1−x}\)As layer is essential for evaluating carrier confinement in future \( \varepsilon \)-Ge-based optical devices on Si. The valence band and conduction band offsets at the \( \varepsilon \)-Ge/In\(_{x}\)Ga\(_{1−x}\)As heterointerface were determined using X-ray photoelectron spectroscopy (XPS) by measuring the atomic core level (CL) binding energies in \( \varepsilon \)-Ge \((\text{Ge 3d})\) and In\(_{x}\)Ga\(_{1−x}\)As \((\text{As 3d}_{5/2})\), the valence band maxima (VBM) of each material, and the interfacial CL binding energy shifts in each material. In order to accurately determine the band offsets, three samples were selected for XPS analysis: (i) 40 (30 nm) \( \varepsilon \)-Ge on In\(_{0.11}\)Ga\(_{0.89}\)As (In\(_{0.17}\)Ga\(_{0.83}\)As) was used to measure the CL and VBM binding energy spectra for \( \varepsilon \)-Ge; (ii) 750 nm (600 nm) In\(_{0.11}\)Ga\(_{0.89}\)As (In\(_{0.17}\)Ga\(_{0.83}\)As) without the top \( \varepsilon \)-Ge layer was used to measure the CL binding energy of arsenic \((\text{As})\) and the VBM of In\(_{0.11}\)Ga\(_{0.89}\)As (In\(_{0.17}\)Ga\(_{0.83}\)As) without the top \( \varepsilon \)-Ge layer was used to measure the CL binding energy of arsenic \((\text{As})\) and the VBM of In\(_{0.11}\)Ga\(_{0.89}\)As (In\(_{0.17}\)Ga\(_{0.83}\)As); and (iii) ∼2 nm \( \varepsilon \)-Ge/In\(_{0.11}\)Ga\(_{0.89}\)As (In\(_{0.17}\)Ga\(_{0.83}\)As) was used to measure the CL binding energy shifts of Ge and As at the \( \varepsilon \)-Ge/In\(_{x}\)Ga\(_{1−x}\)As heterointerface. The binding energy was corrected by adjusting the carbon (C) 1s CL peak position to 285.0 eV for each sample surface. Residual native oxide was removed from the surface of Ge and In\(_{x}\)Ga\(_{1−x}\)As using dilute NH\(_4\)OH:H\(_2\)O\(_2\):H\(_2\)O (2:1:200 volume ratio) prior to loading into the XPS chamber. The valence band offset \((\Delta E_v)\) between the \( \varepsilon \)-Ge and the In\(_{x}\)Ga\(_{1−x}\)As layer was determined using Kraut’s method,\(^{62}\) which is widely used in the analysis of heterointerface band discontinuities at semiconductor/semi-

![Figure 10. \( \mu \)-PL determined bandgaps of 0.82% and 1.11% \( \varepsilon \)-Ge in comparison to the theoretical bandgap-strain dependence for Ge calculated using a 30 × 30 \( \mathbf{k} \cdot \mathbf{p} \) model taking into quantization-induced bandgap enhancement at decreased \( \varepsilon \)-Ge layer thicknesses. Good agreement is found between the measured (symbols) and predicted (lines) bandgaps as a function of increasing tensile strain.](image)

![Figure 11. Representative photoelectron spectra used in the valence band analysis for the \( \varepsilon \)-Ge/In\(_{0.11}\)Ga\(_{0.89}\)As (a) and \( \varepsilon \)-Ge/In\(_{0.17}\)Ga\(_{0.83}\)As (b) structures, corresponding to (i) thick (>10 nm) \( \varepsilon \)-Ge emission, (ii) bulk In\(_{x}\)Ga\(_{1−x}\)As emission, and (iii) interfacial core level binding energy shifts.](image)
conductor\textsuperscript{53–66} as well as dielectric/semiconductor\textsuperscript{57,68} heterojunctions. The positive charges generated during the XPS measurements were neutralized by flowing electrons through the sample stage to prevent uncompensated electron loss that could potentially affect the interfacial band bending results due to sample charging. The valence band offset can be written as\textsuperscript{62}

\[
\Delta E_V = (E_{V_{\text{Ge}}} - E_{V_{\text{BVM}}}) - (E_{\Gamma_{\text{As3d}}} - E_{\Gamma_{\text{Ga3s}}})
\]

where \(E_{V_{\text{Ge}}}\) and \(E_{\Gamma_{\text{As3d}}}\) are the CL binding energies of \(\varepsilon\)-Ge Ge 3d and In\(_{0.11}\)Ga\(_{0.89}\)As 3d\(_{3/2}\) CLs, respectively, and \(E_{\Gamma_{\text{Ga3s}}}\) is the VBM of the corresponding material. \(E_{\Gamma_{\text{As3d}}}\) is determined by linearly fitting the leading edge of the valence band (VB) spectra to the photoemission background line. \(E_{\Gamma_{\text{As3d}}}\) and \(E_{\Gamma_{\text{Ga3s}}}\) are the CL binding energies of As 3d\(_{5/2}\) and Ge 3d measured at the heterointerface. The conduction band offset \((\Delta E_c)\) can be estimated by\textsuperscript{62–66}

\[
\Delta E_c = E_{\Gamma_{\text{Ga3s}}} - E_{\Gamma_{\text{Ge}}} - \Delta E_V
\]

where \(E_{\Gamma_{\text{Ga3s}}}\) and \(E_{\Gamma_{\text{Ge}}}\) are the bandgap energies of In\(_{0.11}\)Ga\(_{0.89}\)As (In\(_{0.17}\)Ga\(_{0.83}\)As) and \(\varepsilon\)-Ge, respectively. The \(\varepsilon\)-Ge bandgap was extracted from \(\mu\)-PL measurements as discussed above and assumes the experimentally determined \(\Gamma\)-lh bandgap in calculating \(\Delta E_c\), whereas the bandgap energy of In\(_{0.11}\)Ga\(_{0.89}\)As \((x = 0.11, 0.17)\) was estimated from ref 69. Figures 11a–b show the CL and VB spectra from (i) \(\varepsilon\)-Ge, (ii) In\(_{0.11}\)Ga\(_{0.89}\)As and In\(_{0.17}\)Ga\(_{0.83}\)As, and (iii) the \(\varepsilon\)-Ge/In\(_{0.11}\)Ga\(_{0.89}\)As and \(\varepsilon\)-Ge/In\(_{0.17}\)Ga\(_{0.83}\)As (b) interfaces, respectively. The inset shows the schematic layer diagram of the sample used for each measurement. From these spectra, the values of \(E_{\Gamma_{\text{As3d}}} - E_{\Gamma_{\text{Ga3s}}}\) and \(E_{\Gamma_{\text{As3d}}} - E_{\Gamma_{\text{Ga3s}}}\) were determined to be 29.45, 40.73, and 11.55 eV, respectively, for the \(\varepsilon\)-Ge/In\(_{0.11}\)Ga\(_{0.89}\)As and \(\varepsilon\)-Ge/In\(_{0.17}\)Ga\(_{0.83}\)As structure and 29.45, 40.72, and 11.56 eV for the \(\varepsilon\)-Ge/In\(_{0.11}\)Ga\(_{0.89}\)As structure. \(\Delta E_c\) was determined to be 0.27 ± 0.05 eV \((0.29 ± 0.05 \text{ eV})\) using eq 1 and the measured binding energy differences. The uncertainty value is taken from the scattering of the VB data with respect to the linear fitting of the VBM position. \(\Delta E_c\) was determined using eq 2 and the bandgap energy of intrinsic In\(_{0.11}\)Ga\(_{0.89}\)As \((x = 0.11, 0.17)\) at 300 K, that is, 1.26 eV \((1.19 \text{ eV})\).\textsuperscript{59} The tensile-strain-modified Ge \(\Gamma\)-lh bandgap energies of \(\sim 0.70\) and \(\sim 0.73 \text{ eV}\) were determined from \(\mu\)-PL measurements and resulted from the simultaneous lifting of the light-hole/heavy-hole valence band degeneracy and reduction of the \(\Gamma\)-point conduction band minima. Using these data, \(\Delta E_c\) was calculated to be 0.26 ± 0.1 eV \((0.20 ± 0.1 \text{ eV})\). It is worth noting that the fitting of the \(\Gamma\)-lh bandgap energy from the experimental \(\mu\)-PL spectra becomes less precise at lower strain levels due to the decreased intensity of the \(\Gamma\)-lh transition (see \(\mu\)-PL analysis section), resulting in an increased convolution between the \(L\)-lh and \(\Gamma\)-lh emission spectra and therefore increased error in the calculated \(\Delta E_c\) value. Additionally, the temperature dependence of the optical bandgaps in Ge will correlate to an increase in \(\Delta E_c\) as a function of increasing temperature because of the reduction in the Ge bandgap. Hence, the calculated \(\Delta E_c\) values reported here provide guidance on the effect of increasing tensile strain at the \(\varepsilon\)-Ge/In\(_{0.11}\)Ga\(_{0.89}\)As heterointerface, however are associated with increased uncertainty as compared to the measured \(\Delta E_c\) values.

Figure 12 shows the schematic band alignment diagrams for the (a) \(\varepsilon\)-Ge/In\(_{0.11}\)Ga\(_{0.89}\)As and (b) \(\varepsilon\)-Ge/In\(_{0.17}\)Ga\(_{0.83}\)As heterointerfaces utilizing the measured \(\Gamma\)-lh bandgaps.

**CONCLUSIONS**

In summary, tunable tensile-strained Ge heterogeneously integrated on Si using composite GaAs and linearly graded In\(_{0.11}\)Ga\(_{0.89}\)As buffers grown by solid source molecular beam epitaxy opens the opportunity for strain-engineered group-IV-based photonic devices. Moreover, the shift in \(\Delta E_c\) as a function of increasing tensile strain in Ge and In alloy composition in the underlying strain template corroborates the trend previously reported for low-to-high strain \(\varepsilon\)-Ge/In\(_{0.11}\)Ga\(_{0.89}\)As heterojunctions grown on GaAs substrates,\textsuperscript{57} indicative of a successful integration scheme for tensile-strained Ge active layers on III–V metamorphic buffer architectures implemented on Si.
all ε-Ge strain states studied in this work. Furthermore, the μ-PL measurements demonstrated an effective, strain-induced bandgap modulation, with L-h (Γ-h) bandgaps of 0.684 eV (0.735 eV) and 0.645 eV (0.704 eV) for the 0.82 ± 0.06% and 1.11 ± 0.03% strained Ge layers, respectively. A type-I energy band alignment with valence band offsets of 0.27 and 0.29 eV for the ε-Ge/In0.11Ga0.89As and ε-Ge/In0.17Ga0.83As heterojunctions were demonstrated via X-ray photoelectron spectroscopy analysis, suggesting that the ε-Ge/InGaAs heterointerface is feasible for carrier confinement in optical devices utilizing increased strain. This is the first experimental and comprehensive study of strain-engineered tensile-strained epitaxial Ge layers heterogeneously integrated on Si. Consequently, the tunable tensile-strained Ge materials growth, structural and optical properties, and band offset parameters pave the way for the realization of novel Ge-based photonic devices integrated on Si.

**MATERIALS AND METHODS**

**Material Synthesis.** Unintentionally doped tensile-strained epitaxial 30–40 nm thick Ge layers were grown on off-cut (100) Si substrates using an in situ growth process utilizing separate solid-source molecular beam epitaxy (MBE) growth chambers for the Ge and III–V materials, connected via an ultrahigh vacuum transfer chamber. The effect of substrate off-cut on the suppression of antiphase domain boundary and stacking fault formation at the GaAs/Si interface is well-supported in the literature and was utilized to achieve device quality active layers in this work. GaAs and linearly graded InGaAs buffers were grown on Si to mitigate the defects and dislocations due to the significant lattice and thermal mismatch between Ge and Si. An InxGa1-xAs constant composition virtual substrate was grown prior to Ge layer growth, immediately followed by strained-layer Ge epitaxy at a growth temperature of 400 °C and growth rate of ~0.1 Å/s, respectively. During the substrate oxide desorption and periodically throughout each layer growth, reflection high energy electron diffraction (RHEED) was used to monitor the surface reconstruction for growth-induced defect patterns. After the GaAs/InGa1-xAs layer growth, the sample was transferred under ultrahigh vacuum to the Ge MBE chamber for subsequent ε-Ge growth. The ε-Ge on Si samples were characterized by high-resolution X-ray diffraction, atomic force microscopy, and low-temperature magneto-transport measurements.

**Materials Characterization.** To determine the crystalline quality and relaxation state of the tensile-strained epitaxial Ge layers grown on Si, high-resolution triple axis X-ray rocking curves and reciprocal space maps were recorded from each sample using a Panalytical Xpert Pro system equipped with both PIXel and proportional detectors and a monochromated Cu Krα (λ = 1.540598 Å) X-ray source. Select samples underwent further X-ray diffraction characterization following the baseline of the valence band spectra recorded for the ε-Ge and InGaAs heterointerface. The Ge 3d core level was characterized via high-resolution X-ray photoelectron spectra (HERED) and InGaAs heterointerface. HR-TEM imaging was performed utilizing a JEOl 2100 transmission electron microscope. For this purpose, electron transparent foils of thin film cross sections of each sample structure were prepared by standard polishing techniques, i.e. mechanical grinding, dimpling and low-temperature Ar⁺ ion beam milling. To analyze the elemental composition of the structure, energy dispersive X-ray spectroscopy (EDS) was performed using a JEOl 2100 system operating in scanning TEM mode. The bandgap as a function of strain within each ε-Ge epilayer was characterized via microphotoluminescence spectroscopy using an 800 nm laser source and a thermoelectric cooled InGaAs detector for optical pumping and detection, respectively. The band alignment at each ε-Ge/InGaAs heterointerface was investigated using a PHI Quantera SXM XPS system with a monochromatic Al-Kα (E = 1486.7 eV) X-ray source. Samples with the InGaAs heterointerface were decapped using an NH₄OH:H₂O₂:H₂O (2:1:200 volume ratio) wet etch prior to loading in the XPS chamber. The queue time was minimized between the cleaning of each ε-Ge layer or removal of the InGaAs capping layer and loading into the XPS chamber. Each ε-Ge layer was thinned down to 1.5–2 nm for XPS measurements requiring photoelectron collection from the ε-Ge/InGaAs heterointerface. The Ge 3d and As 3d/2 CL binding energy spectra, as well as ε-Ge and InGaAs valence band binding energy spectra were collected with a pass energy of 26 eV and an exit angle of 45°. Curve fitting was performed by CasaXPS v2.3.14 using a Lorentzian convolution with a Shirley-type background. The CL energy position was defined to be the center of the peak width at half of the peak height. The VBM values were determined by linear extrapolation of the leading edge to the baseline of the valence band spectra recorded for the ε-Ge and InGaAs heterointerfaces. The VBM value is sensitive to the choice of points on the leading edge used to obtain the regression line, resulting in an uncertainty in the range of 0.05–0.1 eV for ΔE, and ΔEF, respectively, as determined by regressions analysis of selected data over the linear region.

**AUTHOR INFORMATION**

**Corresponding Author**

Tel: (540) 231-6663. Fax: (540) 231-3362. E-mail: mantu.hudait@vt.edu.

**Notes**

The authors declare no competing financial interest.

**ACKNOWLEDGMENTS**

M.C. acknowledges financial support from the NSF under grant number ECCS-1348653. P.G. acknowledges support from an NSF Graduate Research Fellowship Grant No. DGE. 0822220. The authors also acknowledge Dr. Luke F. Lester for several insightful technical discussions related to this work. Lastly, the authors acknowledge the NCFL-Institute for Critical Technology and Applied Science, Virginia Tech Nanofabrication facilities, and Charles Farley for assistance in materials characterization.

**REFERENCES**

(1) International Technology Roadmap for Semiconductors, 2013. h t t p : / / w w w . i t r s . n e t / I T R S % 201 9 9 9 - 2 0 1 4 % 2 0 M t g s , % 2 0 P r e s e n t a t i o n s % 2 0 & % 2 0 L i n k s / 2 0 1 3 I T R S / S u m m a r y 2 0 1 3 . h t m l


