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P. Su,1,a) Z. Han,1 D. Kita,1 P. Becla,2 H. Lin,1 S. Deckoff-Jones,1 K. Richardson,3 L. C. Kimerling,1 J. Hu,1,4 and A. Agarwal1,4

AFFILIATIONS
1Department of Materials Science and Engineering, Massachusetts Institute of Technology, Cambridge, Massachusetts 02139, USA
2CapeSym, Natick, Massachusetts 01760, USA
3CREOL, The College of Optics and Photonics, University of Central Florida, Orlando, Florida 32816, USA
4Materials Research Laboratory, Massachusetts Institute of Technology, Cambridge, Massachusetts 02139, USA

a)Author to whom correspondence should be addressed: peterxsu@mit.edu.

ABSTRACT

We demonstrate a monolithic waveguide sensor integrated with a detector on-chip for mid-infrared absorption spectroscopic sensing. The optical sensing element comprises a chalcogenide glass spiral waveguide, and the detector is a PbTe photodetector integrated directly with the chalcogenide waveguide. The limit of detection of the sensor for methane gas was experimentally assessed to be 1% by volume. Further optimization of the fabrication process and normalization of the laser power fluctuations should result in a maximum sensitivity of 330 ppmv.

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The ability to determine the chemical composition or the presence of a chemical in liquid and gaseous mixtures is important for many fields, such as industrial process monitoring, environmental monitoring, forensics, and medicine and biology. One method widely used to determine chemical compositions is optical absorption spectroscopy, which is typically performed in the infrared region of the electromagnetic spectrum where many chemicals have unique absorption spectra.1 The current equipment used to measure these spectra uses free-space optics that are bulky and expensive, but integrated photonics promises the ability to perform the same analysis with chips that are significantly smaller and that can be mass-produced via techniques similar to those used in the electronics industry.

These chips can generally be broken down into three main constituent parts: the light source, the sensing element, and the detector. Previous work has focused individually on the sensing element and the detector. In the near-infrared, spectroscopic detection of molecules relies on tracing their vibrational overtone absorption. The approach has been implemented using on-chip waveguides5–7 and resonators.8–10 Extending the working wavelength of these sensing elements to the mid-infrared (mid-IR), where strong and unique spectroscopic features reside, promises significantly enhanced selectivity and sensitivity,11 and this has been realized using various material systems including silicon,12–15 germanium,16 silicon-germanium,17 silicon nitride,18 III–V semiconductors,19 and chalcogenides.20–23 Nevertheless, these proof-of-concept sensors still use off-chip light sources and detectors, compromising the benefits of photon integration.

In this paper, we demonstrate a mid-IR waveguide sensor monolithically integrated with an on-chip detector on a silicon platform. In addition to being a critical step towards a fully integrated chip-scale sensing system, waveguide integrated detectors also offer reduced noise due to decreased size and increased speed.24,25 Our device borrows elements from our previous work on spiral chalcogenide glass waveguide sensors22 and waveguide-integrated PbTe detectors.25,26 PbTe provides a monolithic integration capability that represents a major advantage over previously demonstrated mid-IR waveguide-integrated photodetectors,27–30 which mostly rely on hybrid bonding or transfer of the active detector material. The use of chalcogenide glass, a well-known Kerr medium, as the sensor material also foresees seamless integration of the sensing element and the detector with on-chip nonlinear sources to enable broadband spectroscopic interrogation.31

The design of our sensor is schematically shown in Fig. 1. The sensing element is a Ge25Sb7S70 (GeSbS) chalcogenide glass...
spiral waveguide with a cross-section of 2 μm wide by 1 μm thick, providing single mode waveguiding at the absorption peak of methane at a 3.31 μm wavelength, as shown in Fig. 2. The spiral offers a long interaction length (5 mm) of light with the environment within a small footprint (550 μm by 530 μm), and the minimum bend radius is 100 μm, so the bending loss is negligible. Numerical simulations show 10^{-4} dB/cm, or 10^{-16} dB per 90° bend, of bending loss for the 100 μm minimum bend radius.) The gaseous analyte, composed of methane mixed with nitrogen, is transported to the spiral via a polydimethylsiloxane (PDMS) microfluidic channel. To prevent the PDMS microfluidic chamber wall from absorbing much of the guided light where it touches the waveguide, the input and output waveguide widths are adiabatically increased from 2 μm within the chamber to 15 μm as it approaches the PDMS chamber wall to reduce modal overlap with the elastomer and therefore reduce parasitic optical absorption. Light from the spiral sensing element is split using a multimode interferometer to an alignment waveguide and a waveguide with the integrated detector. The integrated detector comprises a 40 μm long PbTe photoconductive layer directly beneath the waveguide.

To fabricate our sensor, we started with a silicon substrate with a 3 μm thermally grown oxide. A 100 nm PbTe film and then a 300 nm thick Sn contact layer are deposited via thermal evaporation and patterned using photolithography and lift-off. The GeSbS layer was then deposited and patterned using an electron-beam-lithography-based lift-off procedure. The wafer was then cleaved to create the input and output waveguide facets, resulting in a chip of approximately 1 cm by 1 cm in size. The PDMS microfluidic chamber was prepared using replica molding and subsequently attached to the chip through plasma bonding to complete the sensor fabrication.

The gas sensing capabilities of the integrated spiral sensor and the PbTe detector were measured using the setup shown in Fig. 3. Light of 3.31 μm wavelength from a tunable mid-IR laser (wavelength tunable from 2.5 to 3.8 μm with a 5 nm linewidth, Firefly, M Squared, Ltd.) is modulated with a chopper at 1 kHz and edge-coupled to the input waveguide via an aspheric lens. Light from the alignment arm of the sensor is coupled via another lens to a cryogenically cooled InSb focal plane array to facilitate alignment. The gas flows of methane and nitrogen carrier gas are regulated using mass flow controllers to achieve different concentrations of methane in the PDMS gas chamber. The integrated PbTe detector is biased using a constant 100 μA current (from a Keithley 6220 current source), and the voltage across the detector is passed to a lock-in amplifier (SR810, Stanford Research Systems) synchronized with the modulation frequency of the chopper. In our experiments, the lock-in time constant is fixed at 1 s, corresponding to an effective noise bandwidth of 0.078 Hz.

The integrated detector signal was monitored under a flow of pure nitrogen for 30 s as a baseline reference, followed by 30 s measurements at different concentrations of methane with a balance of nitrogen. The signal with methane was normalized to the signal under pure nitrogen to obtain the transmittance. The sensor follows Beer’s law, which is given by the following equation:
\[ T = \frac{V_{CH_4}}{V_{noCH_4}} = \exp(-2\Gamma L C), \]  

where \( T \) is the transmittance, \( V_{CH_4} \) is the detector signal under the methane–nitrogen mixture, \( V_{noCH_4} \) is the detector signal under pure nitrogen, \( \alpha \) is the absorbance of pure methane at atmospheric pressure, \( \Gamma \) is the confinement factor in the gas (which is 12.5% based on our modal simulation), \( L \) is the length of the sensing element, and \( C \) is the methane volumetric concentration. Figure 4 plots the measured modal absorption coefficient \((\Gamma \alpha C)\) calculated using Beer’s law versus the methane concentration. Each concentration was measured three times to obtain the average and standard error. The response of the sensor agrees well with the fit calculated from Beer’s law.

The limit of detection (LOD) of the sensor can be determined by the following equation:

\[ \text{LOD} = \frac{\text{NEA} \cdot \text{SNR}}{2\alpha L}, \]

where NEA is the noise–equivalent absorbance and \( \text{SNR} \) is the signal-to-noise ratio. We use the commonly adopted 3-\( \sigma \) criterion, giving an SNR of 3. For our sensor, the detector noise was measured to be 8.13 \( \mu \text{V/Hz}^{1/2} \), while the detector signal under pure nitrogen was measured to be 1.09 mV. This results in a LOD of 3.5 vol. \%/Hz^{1/2}. Our measurement was performed using a noise bandwidth of 0.078 Hz; so, the minimum detectable concentration is 1 vol. %.

However, this is not the best sensitivity that can be expected from an integrated sensing element and detector. There are two primary considerations that govern the minimum detectable concentration: the waveguide transmission loss, which changes the ideal length of the spiral, and the detector noise.

The waveguide transmission loss was measured using the cutback method with several spiral waveguides of different lengths fabricated on the same chip. Using the shortest spiral as the baseline, the relative loss versus the relative length difference is plotted in Fig. 5. The slope of the best fit line is the transmission loss, which is 8 dB/cm. This transmission loss is higher than that in other works with chalcogenide glass waveguides, with Du et al. being able to achieve 0.5 dB/cm loss at 1550 nm.\(^{32}\)

The higher loss in this work is believed to be due to accumulated organic contamination from the fabrication protocol (being the third step in a multi-step process) and not roughness, since the average RMS sidewall roughness measured via AFM (2.3 nm) is lower than other waveguides fabricated using the same method which show lower loss.\(^{36,37}\) Improving the fabrication to decrease this loss to 0.8 dB/cm would allow an increase in the spiral length by 10 times, while maintaining the power at the integrated detector. This would enhance the detector signal, improving sensitivity by a factor of 10 to 0.35%/Hz^{1/2} or 0.1% (1000 ppmv) at our experimental noise bandwidth.

The noise measured at the detector can also be further decreased. Using a free space InAsSb detector (PDA10PT, Thorlabs), the laser power fluctuation noise was measured using the same effective noise bandwidth of 0.078 Hz, resulting in a noise of 0.20% of the power of the laser. This turns out to be the dominant contributor to our measured noise levels, and eliminating this by simultaneously measuring laser power along with the sensor signal decreases the measured noise to 2.74 \( \mu \text{V/Hz}^{1/2} \). Combined with the waveguide loss improvement, the minimum detectable concentration becomes 0.12%/Hz^{1/2} or 0.033% (330 ppmv) at our experimental noise bandwidth.

The theoretical minimum noise for a photoconductor like PbTe is the Johnson–Nyquist noise level, which is given by the following equation:

\[ V_{\text{noise}} = \sqrt{4k_B T R \Delta f}, \]

where \( k_B \) is the Boltzmann constant, \( T \) is the temperature, \( R \) is the resistance of the detector, and \( \Delta f \) is the noise bandwidth. The Johnson–Nyquist noise level of our 24 kΩ detector at room temperature is 20 nV/Hz^{1/2}, still several orders of magnitude smaller than the remaining noise measured. Further work is ongoing to characterize and understand the sources of the remaining noise to improve sensitivity.

In conclusion, we have demonstrated a fully integrated sensing element and detector on-chip for mid-infrared absorption
spectroscopic sensing at room temperature. The sensor is composed of a spiral waveguide sensing element and a waveguide–integrated PbTe detector. Using methane gas as a model gas, our integrated sensor is able to detect 1% methane at 3.31 GHz noise bandwidth of 0.078 Hz following the 3–σ criterion. The stability of the sensor is very good, with no apparent drift during measurements. Further improvements to the fabrication process and eliminating laser power fluctuation noise would result in a maximum sensitivity of 330 ppmv, while more work needs to be done to understand and eliminate the remaining noise sources from the detector.

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