Mid-Infrared chemical sensing using a chalcogenide integrated transducer

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Mid-Infrared (2-20 μ m) spectroscopic techniques are widely used to identify chemicals substances, allowing quantitative real-time measurements in gases, liquids and solids. The current trend heads to the miniaturization of optical sensors, replacing bulky laboratory instruments (FTIR, monochromators, ATR etc..) by lab-on-chip devices providing portability, mechanical stability, immunity to electromagnetic noise and the potential for batch production [1]. In recent years, optical integrated devices have been fabricated using different technologies such as GaAs, Si, Si₃N₄ and Ge. Chalcogenide glasses have also emerged as good candidates to manufacture Mid-Infrared photonic integrated circuits thanks to their ability to be deposited as thin films, their broad transparency (up to 20 μ m), their potential to be doped with rare earth ions and their refractive index tunability obtained by varying glass composition [2].

In this paper, we present the optical detection of isopropanol solutions by evanescent field in the Mid Infrared using a chalcogenide single-mode ridge waveguide as integrated transducer. The optical absorption of isopropanol at 7.7 μ m (Fig. 1) is detected through the variation of the transmitted optical power as a function of analyte concentration.

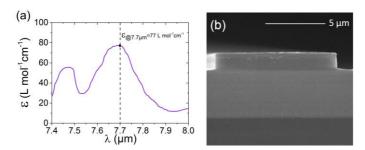


Fig. 1. (a) Molar absorption of isopropanol in the Mid-IR [3]. (b) Cross section of ridge waveguide (SEM image) made of two different compositions of chalcogenide glasses: Ge12.5Sb25Se62.5 as guiding layer, Ge28.1Sb6.3Se65.6 as confinement layer and the silicon substrate.

Chalcogenide multilayer structures were deposited on a silicon substrate by RF magnetron sputtering and processed as ridge waveguides using standard i-line photolithography and fluorine-based reactive ion etching (Fig. 1b) [2]. Refractive index values of 2.77 and 2.44 have been measured at λ =7.7 μ m, respectively, for the guiding (Ge_{12.5}Sb₂₅Se_{62.5}) and confinement (Ge_{28.1}Sb_{6.3}Se_{65.6}) layers of the integrated transducer.

Optimal dimensions of the single-mode ridge waveguide (width=10 μ m and height=1.7 μ m) led to an evanescent power factor η =5%. Intrinsic propagation losses α = 2.5 dB/cm were measured by a cutback method from S-shape waveguides [2].

The sensitivity of the device is represented by the variation of output optical power P as a function of the solute concentration C[1] according to the expression 1.

$$S = \left| \frac{dP}{dC} \right| = \varepsilon \eta L P_0 \exp \left(-\varepsilon \eta C L - \alpha_{prop} L \right)$$
 (1)

The evolution of the sensitivity S as a function of the waveguide length L was calculated and plotted in Fig. 2a taking into account the following parameters: ε =77 L mol⁻¹cm⁻¹, η =5 %, α =2.5 dB/cm, P₀= 0.5 mW. From this figure, it can be seen that the maximum transducer sensitivity S is obtained for a waveguide length $L_{\rm opt}$ equal to 0.63 cm.

Furthermore, the absorption of isopropanol at 7.7 μm is detected by measuring the attenuation of the transmitted power as a function of the analyte concentration. The performance of the evanescent optical integrated sensor was tested by depositing a droplet of 2 μL of the solution (isopropanol dissolved in cyclohexane) on the ridge waveguide surface. The intensity of guided mode, observed at the output of the waveguide by near-field at λ =7.7 μm , was measured by the Mid-IR camera for several concentrations between 2 and 10 v/v%, the results are displayed in Fig. 2b, 2c.

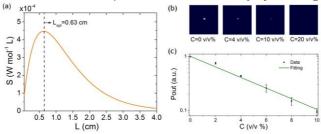


Fig. 2. (a) Evolution of the sensitivity S as a function of the waveguide length L for α =2.5 dB/cm. (b) Near field intensity of propagated light and (c) transmitted optical power at 7.7 μ m for different concentrations of isopropanol.

The optical transmitted power decreases by increasing isopropanol concentration following Beer-Lambert law. These experimental results demonstrating on-chip real-time detection of isopropanol at 7.7 μ m could be extended to other substances of interest and represent therefore an outstanding progress in lab-on-chip sensing schemes.

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- [3] See: vpl.astro.washington.edu/spectra/allmoleculeslist.htm for IR absorption cross-sections from PNNL (Last accessed December, 2016)