Boise State University

ScholarWorks

[Materials Science and Engineering Faculty](https://scholarworks.boisestate.edu/mse_facpubs) [Publications and Presentations](https://scholarworks.boisestate.edu/mse_facpubs)

[Micron School for Materials Science and](https://scholarworks.boisestate.edu/mse) **Engineering**

10-1-2006

Photonic Band Tuning in 2D Photonic Crystals by Atomic Layer Deposition

Elton Graugnard Boise State University

Davy P. Gaillot Georgia Institute of Technology

Simon N. Dunham Georgia Institute of Technology

Curtis W. Neff Georgia Institute of Technology

Tsuyoshi Yamashita Georgia Institute of Technology

See next page for additional authors

©2006 IEEE. Personal use of this material is permitted. However, permission to reprint/republish this material for advertising or promotional purposes or for creating new collective works for resale or redistribution to servers or lists, or to reuse any copyrighted component of this work in other works must be obtained from the IEEE. DOI: [10.1109/LEOS.2006.278973](https://doi.org/10.1109/LEOS.2006.278973)

Authors

Elton Graugnard, Davy P. Gaillot, Simon N. Dunham, Curtis W. Neff, Tsuyoshi Yamashita, and Christopher J. Summers

This conference proceeding is available at ScholarWorks: https://scholarworks.boisestate.edu/mse_facpubs/13

Photonic band tuning in 2D photonic crystals by atomic layer deposition

Elton Graugnard, Davy P. Gaillot, Simon N. Dunham, Curtis W. Neff, Tsuyoshi Yamashita and Christopher J. Summers *School of Materials Science & Engineering Georgia Institute of Technology, Atlanta, GA 30332-0245 USA*

Atomic layer deposition (ALD) has become a powerful tool for the fabrication of high quality 3-dimentional photonic crystals (PCs) from both inorganic (opal) and organic (holographically patterned polymer) templates [1,2]. With ALD, highly conformal films can be grown with a precision of 0.05 nm, which, when combined with the availability of a wide range of low temperature film growth protocols, enables a high degree of control over material and structural properties to precisely tune optical properties [3]. Two-dimensional photonic crystals have been developed extensively for applications in optical interconnects, beam steering, and sensor devices; and are predominantly fabricated by electron-beam lithography. The optical properties of 2D photonic crystal slab waveguides are determined by the precision of the lithography process, with limited post fabrication tunability.

Here were report the application of ALD to 2D photonic crystal silicon slab waveguides as a means to precisely tune the dispersion properties of the as-fabricated device. A triangular lattice of air holes in a 300 nm thick single crystal silicon slab, shown in Fig. 1 (lower section), was fabricated on a silicon-on-insulator substrate by ebeam lithography and inductively-coupled-plasma etching. The periodicity of the lattice was 360 nm with a hole radius of 123 nm. The optical properties of the as-fabricated device were characterized with a resonant-coupling reflectance technique [4] along the main PC lattice symmetry directions, Γ-M and Γ-K, over a wavelength range from 700- 1700 nm. Amorphous $TiO₂$ (refractive index of 2.31 at 800 nm) layers were then conformally deposited by ALD at 100 ºC in steps of 20 nm (392 cycles at 0.051 nm per cycle) using TiCl₄ and H_2O precursors. The upper section of Fig. 1 shows the same PC after coating with 40 nm of $TiO₂$ and confirms the conformality and precision of ALD in uniformly reducing the structure hole radii. The device was optically probed between each deposition to monitor the progressive shifts in band structure. 3D finite-difference

Figure 1. Scanning electron microscope images of a bare triangular lattice PC (lower section) and after being coated with 40 nm of $TiO₂$ (upper section).

time-domain calculations were performed to model the coated device using a conformally-defined dielectric function to accurately represent the $TiO₂$ coatings. Figure 2 shows both the experimental data and FDTD calculations along the Γ-K direction for the uncoated (left) and 40 nm $TiO₂$ coated (right) sample. The arrows in Fig. 2 illustrate the shifts in the measured and also calculated bands from the uncoated to coated sample. The FDTD data are observed to be in excellent agreement with the optical measurements for the lowest band and in very close agreement for all higher bands. As expected, the $TiO₂$ coating shifts the bands to lower frequency by increasing the average dielectric constant of the device. These results demonstrate a technique capable of simultaneously enabling very large static adjustments $(\sim 20\%)$ in the photonic band structure with very high precision $(\sim 0.09\%$ per nm or $\sim 0.004\%$ per ALD cycle). It follows that this simple and unique process, which can be applied to any 2D structure, enables precise control over the optical properties of photonic crystal devices.

Figure 2. Measured (dots) and FDTD calculations (lines) of photonic band structure in the uncoated (left) and 40 nm $TiO₂$ coated (right) Si slab waveguide.

In summary, we have demonstrated a technique in which the photonic bands of two-dimensional triangular lattice photonic crystal slab waveguides can be statically tuned by nanoscale modifications, thereby enabling unprecedented adjustment to the dispersion properties of any 2D photonic crystal. Additionally, the technique facilitates the formation of layered and composite 2D PC waveguides thus opening new fabrication routes to control dispersion, propagation and dielectric contrast.

We acknowledge support from the Army Research Office under contract DAAD19-01-1-0603.

 \overline{a}

¹ J. S. King, E. Graugnard, and C. J. Summers, Adv. Mater. **17** 1010 (2005).

² J. S. King, E. Graugnard, O. M. Roche, D. N. Sharp, J. Scrimgeour, R. G. Denning, A. J. Turberfield, and C. J. Summers, Adv. Mater. **18** 1561 (2006).

³ E. Graugnard, J. S. King, D. P. Gaillot and C. J. Summers, Adv. Func. Mater. **16** 1187 (2006).

⁴ V.N. Astratov, D.M. Whittaker, I.S. Culshaw, R.M. Stevenson, M.S. Skolnick, T.F. Krauss, and R.M. De La Rue, Phys. Rev. B **60** R16255 (1999).