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HfO₂-based platform for high-index-contrast visible and UV integrated photonics

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Ultraviolet and visible integrated photonics enable applications in quantum information, sensing, and spectroscopy, among others. Few materials support low-loss photonics into the UV, and the relatively low refractive index of known depositable materials limits the achievable functionality. Here, we present a high-index integrated photonics platform based on HfO₂ and Al₂O₃ composites deposited via atomic layer deposition (ALD) with low loss in the visible and near UV. We show that Al₂O₃ incorporation dramatically decreases bulk loss compared to pure HfO₂, consistent with inhibited crystallization due to the admixture of Al₂O₃. Composites exhibit refractive index n following the average of that of HfO₂ and Al₂O₃, weighted by the HfO₂ fractional composition x . At $\lambda = 375$ nm, composites with $x = 0.67$ exhibit $n = 2.01$, preserving most of HfO₂'s significantly higher index, and 3.8(7) dB/cm material loss. We further present fully etched and cladded waveguides, grating couplers, and ring resonators, realizing a single-mode waveguide loss of 0.25(2) dB/cm inferred from resonators of 2.6 million intrinsic quality factor at $\lambda = 729$ nm, 2.6(2) dB/cm at $\lambda = 405$ nm, and 7.7(6) dB/cm at $\lambda = 375$ nm. We measure the composite's thermo-optic coefficient (TOC) to be $2.44(3) \times 10^{-5}$ RIU/°C near $\lambda = 397$ nm. This work establishes (HfO₂) _{x} (Al₂O₃)_{1- x} composites as a platform amenable to integration for low-loss, high-index photonics spanning the UV to NIR. © 2025 Optica Publishing Group. All rights, including for text and data mining (TDM), Artificial Intelligence (AI) training, and similar technologies, are reserved.

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Integrated photonics in the visible and UV is essential for a variety of applications [1] spanning bio-chemical spectroscopy [2], neural stimulation and probing [3], and quantum control of trapped-ion [4–9], neutral atom [10], and solid-state quantum systems [11]. Visible and UV functionality often presents a challenge however [12,13] since commonly used integrated photonics materials absorb at short wavelengths, and scattering loss increases with decreasing λ [14,15], e.g., as $\sim 1/\lambda^4$ for Rayleigh scattering.

Si₃N₄ is a mature and commonly used CMOS-compatible platform with a refractive index $n \sim 2.1$ at $\lambda = 405$ nm enabling

passive and thermo-optic functionality across much of the visible. However, material loss limits performance below $\lambda \sim 450$ nm [6,16], with SM waveguides exhibiting >10 dB/cm loss near $\lambda \sim 405$ nm that further rapidly increases with decreasing wavelength [17]. Crystalline AlN waveguides on sapphire [18] offer loss as low as ~ 8 dB/cm at $\lambda \sim 390$ nm [19], but more flexibly integrable AlN deposited on amorphous substrates typically exhibits high loss in the visible and UV due to its polycrystalline form, e.g., 650 dB/cm at $\lambda \sim 400$ nm [20]. Deposited amorphous aluminum oxide (Al₂O₃) [21] has enabled single-mode (SM) propagation as low as ~ 1.35 dB/cm at $\lambda = 369$ nm [7,22]. Despite its utility for low-loss UV photonics, its relatively low index ($n = 1.68$ at $\lambda = 405$ nm) and hence low index contrast with respect to SiO₂ are key limitations. High index contrast is critical for strong optical confinement, small bend radii and device footprints, compact and efficient gratings whose strengths scale approximately as $(n_{\text{core}}^2 - n_{\text{clad}}^2)^2$ [23,24], photonic bandgap functionality [24], and efficient acousto-optic interaction (with acousto-optic figure of merit scaling as n^6) [25]. A CMOS-compatible, deposited, high-index material transmissive for blue/UV wavelengths would hence alleviate key limitations of Al₂O₃ for a variety of integrated photonics functionalities.

A promising candidate is hafnia (HfO₂), having a bandgap of approximately 5.65 eV [26], high refractive index ($n = 2.09$ at $\lambda = 405$ nm), and well studied CMOS compatible deposition form via both sputtering and ALD [27,28]. However, its propensity to crystallize in films of thickness beyond a few nanometers [29] results in significant optical loss in waveguides. Its use in photonics has so far been limited to optical coatings or metasurfaces with optical interaction lengths of hundreds of nm [30,31]; significantly lower losses are usually required in waveguide photonics where centimeter-scale propagation lengths are typical. Here, we show that incorporation of Al₂O₃ in a primarily HfO₂ layer results in dramatic decrease in bulk material loss, which we attribute to inhibition of crystallization, consistent with similar observations in the context of electronic applications [32]. This results in an effective composite with index well described by the fractional average composition of the film, preserving most of HfO₂'s advantage in index despite drastically lower loss. We also present methods to lithographically pattern these films, demonstrating single-mode waveguide transmission of 2.6(2) dB/cm at $\lambda = 405$ nm and ring resonators with 2.55×10^6 intrinsic Q at $\lambda = 729$ nm. Our work establishes (HfO₂) _{x} (Al₂O₃)_{1- x}

composites for high-index visible and UV photonics, as well as fabrication methods for low-loss photonic structures. The resulting platform offers low-loss propagation in a single core material over much of the range offered separately by Si_3N_4 and Al_2O_3 to date.

Composite films were deposited at IBM Zurich using plasma-enhanced ALD (PEALD) at 300°C with trimethylaluminum (TMA) and tetrakis(ethylmethylamino)hafnium (TEMAH) precursors. Nanolaminate films were grown with layer periodicity P , each period consisting of $(1-x)P$ layers of Al_2O_3 followed by xP layers of HfO_2 (Fig. 1(a) shows an example with $x = 0.67$ and $P = 3$). Films of ~ 80 nm thickness of varying x were grown on Si wafers, and their n was inferred via an isotropic Cauchy fit to spectroscopic ellipsometry data. At $\lambda = 405$ nm, the measured n agrees with $x n_{\text{HfO}_2} + (1-x) n_{\text{Al}_2\text{O}_3}$, the weighted average of the two constituent material indices. For propagation loss measurements via a prism coupling method, ~ 75 nm thick films were grown on Si wafers with $2.7\ \mu\text{m}$ of thermal SiO_2 , with a fixed composition near $x = 0.67$ and varying period from 3 to 100 layers. We calculate the power confinement factor ($C_{\text{slab}} = P_{\text{core}}/P_{\text{total}}$) and estimate the bulk absorption as $\alpha_{\text{bulk}} = \alpha_{\text{slab}}/C_{\text{slab}}$, where α_{slab} is the measured slab loss, neglecting the loss arising from surface scattering. As shown in Fig. 1(c), inferred material loss decreases substantially with decreasing P . Atomic force microscopy shows the root mean square roughness decreases from ~ 2.63 nm for pure HfO_2 to ~ 0.35 nm for a configuration of $x = 0.67$ and $P = 3$. Both observations suggest the films are more prone to crystallization as the number of continuous HfO_2 layers increases, consistent with other observations [33]. Films with $x = 0.67$ and $P = 3$ achieve $n = 1.96$ and $\alpha_{\text{bulk}} \approx 4.2$ dB/cm at $\lambda = 405$ nm with no annealing. This represents a reduction of 9% in the index of refraction from pure HfO_2 and a substantial decrease in bulk loss from well above the measurement limit of the prism coupling method used (>30 dB/cm) observed for pure HfO_2 at these thicknesses. We refer to this configuration as the composite material for the rest of the article.

The deposition was reproduced at the Cornell NanoScale Science and Technology Facility (CNF), where after conditioning the PEALD chamber with ~ 50 nm of composite material and annealing in N_2 ambient at 800°C for 1 h, we infer $\alpha_{\text{bulk}} = 2.57(8)$ dB/cm at $\lambda = 405$ nm. After annealing, the material is expected to be isotropic, with ALD layering serving only as a means to achieve the mixture. Figure 2(a) shows the measured refractive indices for Si_3N_4 , Al_2O_3 , HfO_2 , and the composite. We also fit data to a Cody–Lorentz model in order to extract the extinction coefficient. These data demonstrate an improvement in UV transparency (Fig. 2(b)) compared to Si_3N_4 and HfO_2 , with a comparable refractive index. To measure the extinction coefficients well below the sensitivity limit of the ellipsometry employed ($k \sim 0.002$, $\alpha \sim 300$ dB/cm) [16,34] as relevant to integrated waveguide devices, as well as to explore utilization of this material platform in integrated photonic devices and systems, we fabricate and measure the propagation loss in etched waveguides.

Device fabrication begins with annealed 80 or 100 nm thick composite films. Patterns are defined via electron-beam lithography (JEOL9500) in 300 nm of ZEP520-A resist. Patterns are etched into the film via inductively coupled plasma reactive ion etching with a BCl_3/Ar chemistry, resulting in a sidewall angle of $\sim 11^\circ$ (Fig. 3(a)). Following a resist strip with remover 1165 and a standard RCA clean, we deposit an ~ 4 nm layer of Al_2O_3

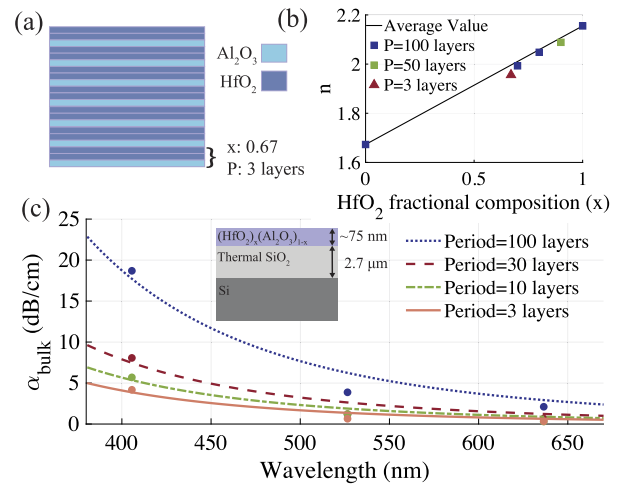


Fig. 1. (a) Al_2O_3 (blue) and HfO_2 (dark blue) layers are deposited with a period $P = 3$ and a fractional composition $x = 0.67$. (b) Measured refractive index n from an isotropic Cauchy model. (c) Measured material loss (α_{bulk}) inferred from a slab mode with a fixed fractional composition near $x = 0.67$ (data points) with $1/\lambda^4$ fits (lines), corresponding to the assumption of scattering originating from within the bulk. Inset: film stack-up supporting the slab mode.

and anneal again at 800°C for 1 h. Finally, we deposit ~ 600 nm of plasma-enhanced chemical vapor deposition (PECVD) SiO_2 at 300°C using a TEOS precursor as a waveguide cladding.

We found that direct deposition of SiO_2 with LPCVD at 800°C or PECVD SiO_2 at 300°C without the ~ 4 nm of Al_2O_3 significantly increased propagation loss as compared to etched air-clad structures. This degradation may be due either to diffusion of oxygen vacancies across the HfO_2 – SiO_2 interface [35] or the creation of surface states as suspected for Si_3N_4 [36] or Si [37]. We interpret the thin Al_2O_3 as acting as a diffusion barrier or passivation layer, with little impact on the optical modes.

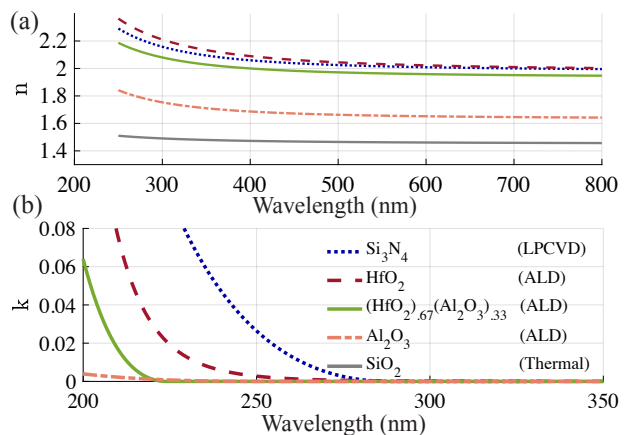


Fig. 2. Measured refractive index n (a) and extinction coefficient k (b) of common materials used in photonic devices. Note that k values inferred from ellipsometry indicate the onset of strong absorption associated with the optical bandgap but do not resolve the range of $k \ll 0.002$ ($\alpha \ll 1300$ dB/cm at $\lambda = 405$ nm) relevant to integrated waveguide devices.

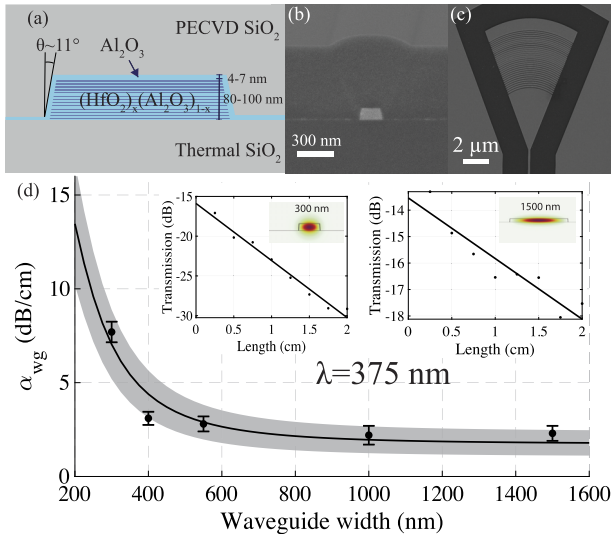


Fig. 3. (a) Schematic cross section of composite waveguides realized. (b) SEM cross section of a 100 nm thick waveguide used to measure waveguide loss at $\lambda = 405$ nm. (c) SEM image of a representative grating used for fiber coupling at $\lambda = 405$ nm. (d) Measured waveguide loss at $\lambda = 375$ nm as a function of width for an 80 nm thick waveguide. The gray line represents the model fit with one standard deviation. Inset: measured transmission of 300 nm and 1500 nm wide waveguides versus test length.

We employ a cutback method to measure waveguide propagation loss α_{wg} and power coupling efficiency η , via fits to $P_{out}/P_{in} = \eta^2 e^{-\alpha_{wg}z}$ with z the straight waveguide length varied up to 2 cm. We use surface grating couplers [38,39] for coupling fabricated waveguides to optical fibers angled 25° from the surface normal. Simulations show the grating coupling efficiency η for the quasi-TE mode is $\sim 10\times$ that for the quasi-TM mode; thus, we selectively excite the former by adjusting input polarization to maximize transmission. Measurements at $\lambda = 405$ nm employ light from a fiber-coupled laser diode (Thorlabs LP405-SF10). For a SM waveguide of 80×300 nm² cross section, we obtain $\alpha_{wg} = 2.6(2)$ dB/cm. The coupling loss is found to be $\eta = 10.8(1)$ dB (simulated to be ~ 5.5 dB) at the nominal design angle without experimental angle optimization. At $\lambda = 375$ nm (Toptica iBeam smart), we measure 7.7(6) dB/cm for the same waveguide geometry, and $\eta = 7.9(3)$ dB (simulated to be 4.1 dB).

Both bulk material loss and sidewall surface scattering contribute to α_{wg} . For large widths, the sidewall mode overlap decreases, and α_{wg} is dominated by the bulk. We model α_{wg} for waveguides of different widths, taking into account the power confinement factor $C_{wg} = P_{core}/P_{total}$, and surface scattering loss arising from the mode's sidewall overlap (Supplement 1). We observe that α_{wg} 's dependence on the waveguide width aligns with the predicted trend (Fig. 3(d)). We upper-bound the material loss α_{bulk} by assuming all loss arises from the bulk in a 1500 nm wide waveguide so that $\alpha_{bulk} \approx \alpha_{wg}/C_{wg}$. At $\lambda = 375$ nm, we measure $\alpha_{wg} = 2.3(4)$ dB/cm, inferring $\alpha_{bulk} = 3.8(7)$ dB/cm. A similar analysis for $\lambda = 405$ nm (Supplement 1) gives an upper bound for material loss of $\alpha_{bulk} = 3.7(3)$ dB/cm. The similar material losses obtained at $\lambda = 375$ nm and 405 nm suggest these wavelengths are not near the absorption band edge.

In the visible/NIR at $\lambda = 729$ nm, waveguide loss is low enough to pose a challenge for practical cutback measurement.

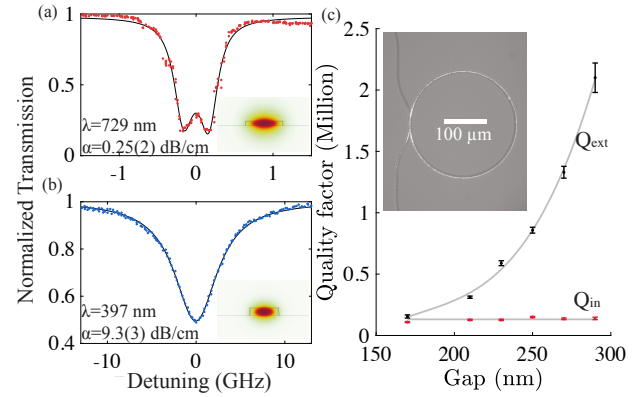


Fig. 4. (a) Measured and fit transmission for a nearly critically coupled ring near $\lambda = 729$ nm and (b) for $\lambda = 397$ nm for a 250 nm coupling gap. (c) Measured Q_{in} and Q_{ext} at $\lambda = 397$ nm. With increasing coupling gap, Q_{in} is relatively constant, and Q_{ext} increases, indicating reliable extraction of Q_{in} . Gray lines are polynomial fits to the data as guides to the eye. Inset: micrograph of a ring resonator operating near critical coupling.

We fabricate ring resonators in a symmetric coupling configuration with a radius of 250 μm and waveguides (ring and bus) with a width of 750 nm ($C_{wg} = 21\%$), near the threshold of SM operation. Transmission optimization via polarization control through grating coupled waveguide structures ensures quasi-TE mode excitation similar to the above. We sweep the frequency of a CW Ti-sapphire laser (M-squared SolisTiS) across the resonance and monitor the frequency with a wavemeter (HighFinesse WS-7). We fit the transmission to obtain the intrinsic (Q_{in}) and external quality factor (Q_{ext}). α_{wg} is related to Q_{in} by

$$\alpha_{wg} = 10 \log_{10}(e) \frac{2\pi n_g}{\lambda Q_{in}} \quad [\text{dB/cm}], \quad (1)$$

where n_g is the group index of the bend mode and is simulated to be $n_g = 1.683$. A gap of 800 nm results in near critical coupling, exhibiting $Q_{loaded} = 1.34 \times 10^6$ and $Q_{in} = 2.55 \times 10^6$ (Fig. 4(a)), corresponding to $\alpha_{wg} = 0.25(2)$ dB/cm. We again upper-bound the material loss to be $\alpha_{bulk} < 1.25$ dB/cm near $\lambda = 729$ nm.

We also measure ring resonators with a 300 nm width and a 180 μm radius at $\lambda = 397$ nm, using a tunable external cavity laser (Toptica DL Pro). Devices with varying coupling gaps were measured, showing that the intrinsic quality factor remains constant, while the external quality factor increases, as expected in an under-coupled regime (Fig. 4(c)). A group index of $n_g = 2.03$ together with the fit $Q_{in} = 1.50(5) \times 10^5$ gives $\alpha_{wg} = 9.3(3)$ dB/cm, $\sim 2\text{--}3\times$ higher than expected based on the SM propagation measurements, despite comparable sidewall overlap of the bend mode for this resonator geometry. We believe this discrepancy is due to suboptimal exposure and proximity correction in the bent waveguides and coupling region, resulting in visible excess loss in this region with the present dosage.

We measure the composite's TOC ($\frac{dn_{core}}{dT}$) by tuning the temperature and monitoring transmission near $\lambda = 397$ nm. The measured temperature dependence of the resonance wavelength λ_{res} is related to the material refractive indices comprising the waveguide (n_{core} and n_{clad}) via

$$\frac{d\lambda_{res}}{dT} = \frac{\lambda_{res}}{n_g} \left[\frac{\partial n_{eff}}{\partial n_{core}} \frac{dn_{core}}{dT} + \frac{\partial n_{eff}}{\partial n_{clad}} \frac{dn_{clad}}{dT} \right]. \quad (2)$$

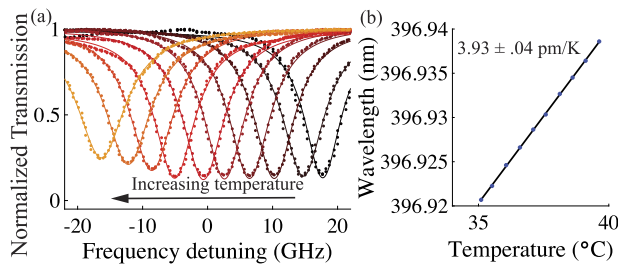


Fig. 5. (a) Microring transmission near $\lambda = 397$ nm for temperatures between 35.1°C and 39.5°C. (b) Resonance wavelength versus temperature. A linear fit shows $\frac{d\lambda_{\text{res}}}{dT} = 3.93(4)$ pm/K, corresponding to $\frac{dn_{\text{core}}}{dT} = 2.44(3) \times 10^{-5}$.

Figure 5 shows the measured resonance temperature shifts, with fit resulting in $\frac{d\lambda_{\text{res}}}{dT} = 3.94(4)$ pm/K. Taking SiO_2 's TOC to be 1×10^{-5} RIU/K [40] and considering the mode's confinement factor, we find $\frac{dn_{\text{core}}}{dT} = 2.44(3) \times 10^{-5}$ RIU/K, close to the measured values for Al_2O_3 and Si_3N_4 [21,41].

These results show that $(\text{HfO}_2)_x(\text{Al}_2\text{O}_3)_{1-x}$ enables low-loss visible and UV photonics, preserving most of HfO_2 's advantage in refractive index compared to platforms in pure Al_2O_3 . We have demonstrated fabrication of SM waveguides, microresonators, and grating couplers (see also [42]). The composite's broad transmission range can enable significant process simplifications, potentially allowing for devices operating over required bandwidths with just one or multiple layers of this composite, rather than both Al_2O_3 and Si_3N_4 [17]. Further work will explore alternatives to PEALD to avoid potential plasma damage [43], such as sputtering, which may enable lower material loss [22], ultimate short-wavelength transmission and power-handling limits for these composites, as well as routes to alleviating surface roughness currently limiting UV SM waveguide loss. Other deposited wide-bandgap materials including ZrO_2 and AlN could be incorporated in similar amorphous composites, for broad transparency and tunable index.

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Data availability. Data are available from the authors upon request.

Supplemental document. See Supplement 1 for supporting content.

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