

Thin film engineering in Er-doped CeO₂ for quantum memory

Kathryn E. Sautter,^{1,2} Gregory D. Grant,^{1,2} Sean E. Sullivan,¹ P. Vamsi K. Nittala,^{1,2} Cheng Ji,^{1,2} Manish K. Singh,^{1,2} F. Joseph Heremans,^{1,2} Supratik Guha^{1,2}

¹ Materials Science Division, Argonne National Laboratory, Lemont, IL 60439, USA.

² Pritzker School of Molecular Engineering, University of Chicago, Chicago, IL 60637, USA.

Developing a commercially available quantum memory device is essential for building wide-area ground-based quantum networks, which will yield applications like guaranteed cryptographic security that uses existing telecommunication infrastructure. However, a functional quantum memory requires the discovery of the optimal materials for these devices, and the optimization of their processing parameters. Researchers have investigated a multitude of materials that could be useful, especially diamond nitrogen-vacancy centers, silicon carbide, and rare-earth doped oxides. A particularly promising candidate for quantum memory is erbium-doped ceria (Er:CeO₂), which theoretically could yield long quantum coherence times of up to 47 ms [1]. Er:CeO₂ is also favorable due to erbium's ⁴I_{15/2} to ⁴I_{13/2} optical transition in the telecom-C band (~1.5 μm), which would allow for direct incorporation of Er:CeO₂ quantum memory into already-existing telecom optical fiber infrastructure with minimal signal attenuation. Recent research on MBE-grown Er:CeO₂ on Si(111) suggests that the emission lifetime of Er:CeO₂ can be further improved through thin film engineering and the reduction of Er³⁺ concentration below 1% [2].

We build on this research by doping CeO₂(111) with low Er³⁺ concentrations (<1%). We use Raman spectroscopy, atomic force microscopy (AFM), and photoluminescence excitation (PLE) spectroscopy to investigate the effects of MBE growth parameters, grown host material quality, and low Er³⁺ concentrations on Er:CeO₂ optical linewidths and emission lifetime. We expand this investigation from Si(111) to two additional substrates, GaAs(111)A and yttria-stabilized zirconia(111) (YSZ), to identify the effects of different substrates and compressive/tensile strain on PLE spectra.

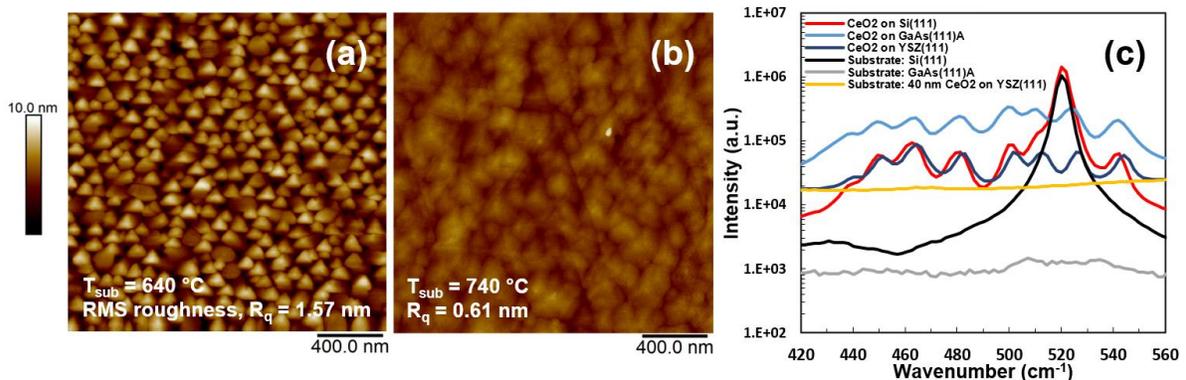


Figure: (a),(b) 2 × 2 μm² AFM images of Er-doped CeO₂ grown at different substrate temperatures (T_{sub}): (a) T_{sub} = 640 °C, (b) T_{sub} = 740 °C. (c) Raman spectra for Er-doped CeO₂ grown on Si(111), GaAs(111)A, and YSZ(111), with each substrate included for reference.

[1] Kanai *et al.*, arXiv:2102.02986 [quant-ph], (2021).

[2] Inaba *et al.*, *Opt. Mater. Express*, 8(9), (2018).

+ Authors for correspondence: ksautter@anl.gov, grantg@anl.gov, sguha@anl.gov