

SOLID-PHASE-EPITAXY OF RUTILE-GeO₂ IN MOLECULAR BEAM EPITAXY (MBE)

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Ultrawide bandgap (UWBG) semiconductor rutile GeO₂ (r-GeO₂, E_g~4.7eV) shows tremendous prospects for the next-generation power electronics as it exhibits superior thermal conductivity, carrier mobility than β-Ga₂O₃. Additionally, it has been predicted to have bipolar dopability.

However, the synthesis of crystalline r-GeO₂ thin films is technically challenging due to the presence of competing metastable amorphous- and quartz- GeO₂ and the lack of r-GeO₂ substrate. Currently, thin films of r- GeO₂ are grown using various methods among which MBE suffers from limited growth rates (10nm/h) due to the high vapor pressure of GeO.

Developing a new strategy for growing r-GeO₂ films with suitable growth rates and high crystal quality is crucial in realizing its application potential. In this study, we create a r-GeO₂ template for quasi-homoepitaxial growth of r-GeO₂ films. It involves deposition of an amorphous GeO₂ layer (5nm) by MBE on an r-TiO₂ (110) substrate and in-situ crystallization at high temperature through solid phase epitaxy (SPE) which competes with the film desorption.

By monitoring surface evolutions (Fig. 1) during SPE using in-situ reflection high-energy electron diffraction, we observed a phase transition from amorphous GeO₂ to a crystalline layer. The presence of the desired stoichiometric ratio was confirmed by X-ray photoelectron spectroscopy (Fig. 2). While crystallinity details will be further studied by Transmission electron microscope.

The creation of GeO₂ crystal seeds enable templated epitaxial growth of GeO₂. Our advances in SPE potentially offer a new route for synthesizing thin film r-GeO₂ and provide important insights into addressing its challenging epitaxy problems.