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Interface formation in ALD-based SnO₂/CeO_x heterostructures — •DOMINIC GUTTMANN¹, RUDI TSCHAMMER¹, CARLOS MORALES¹, MALGORZATA KOT², MICHAL MAZUR², DAMIAN WOJCIESZAK², PAULINA KAPUSCIK², WIKTORIA KOLODZINSKA², JAROSŁAW DOMARADZKI², and JAN INGO FLEGE¹ — ¹Applied Physics and Semiconductor Spectroscopy, BTU Cottbus-Senftenberg, Cottbus 03046, Germany — ²Faculty of Electronics, Photonics and Microsystems, WUST, 50-372 Wrocław, Poland

The electrical resistance of SnO₂ ultrathin films (< 20 nm) made by atomic layer deposition (ALD) strongly depends on thickness, due to intrinsic film defects at interfaces arising from changes in the ALD reaction mechanism during the first cycles and from the film/substrate interaction. Modifying interface properties in SnO₂/CeO_x heterostructures can enhance H₂ sensing performance. We studied the initial growth of SnO₂ by ALD on CeO_x substrates prepared by either electron beam evaporation (EBE) or ALD. Employing the commercial precursor tetrakis(dimethylamino)tin (TDMASn) and ozone (O₃) as well as in vacuo and near-ambient-pressure X-ray photoelectron spectroscopy, we examined how substrate preparation affects the SnO₂ nucleation behavior in the first cycles. SnO₂ growth on EBE-CeO_x indeed starts with the first precursor cycle, whereas ALD-CeO_x requires an additional conditioning step. Connecting these findings to surface chemistry, distinct C1s and N1s signatures attributed to TDMASn adsorption indicate a slow C/N buildup, consistent with previous reports on ALD-grown SnO₂ on Si, SiO₂, and Al₂O₃.

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