

High-mobility two-dimensional electron gas with quantized states in polar-discontinuity doped LaInO₃/BaSnO₃ heterostructure grown by molecular beam epitaxy

G. Hoffmann,¹ A. Hartl,² M. Zupancic,³ A.A. Riaz,⁴ V.N. Strocov,² M. Albrecht,³
A. Regoutz,⁴ O. Bierwagen¹

¹ Paul-Drude-Institut für Festkörperelektronik, Leibniz-Institut im Forschungsverbund Berlin e.V.,
Hausvogteiplatz 5–7, 10117 Berlin, Germany

² Swiss Light Source, Paul Scherrer Institute, CH-5232 Villigen-PSI, Switzerland

³ Leibniz-Institut für Kristallzüchtung, Max-Born-Straße 2, 12489 Berlin, Germany

⁴ University College London, 20 Gordon Street, London WC1H 0AJ, UK

Transistor applications of semiconducting oxides require, both high room-temperature electron mobilities (μ_{RT}) and high charge carrier densities (CCDs), ideally realized with a two-dimensional electron gas (2DEG). So far, prototype oxide 2DEG systems have either high μ_{RT} but limited CCD such as modulation-doped (Al_xGa_{1-x})₂O₃/Ga₂O₃, or a high CCD but low μ_{RT} such as the polar-discontinuity doped LaAlO₃/SrTiO₃ interface. Interfacing the more suitable, wide-bandgap, nonpolar semiconductor BaSnO₃ (BSO), having high bulk μ_{RT} (up to 320 cm²/Vs), with polar LaInO₃ (LIO) is predicted to create and confine a 2DEG with CCD up to 2×10^{14} cm⁻² for the SnO₂/LaO interface termination.

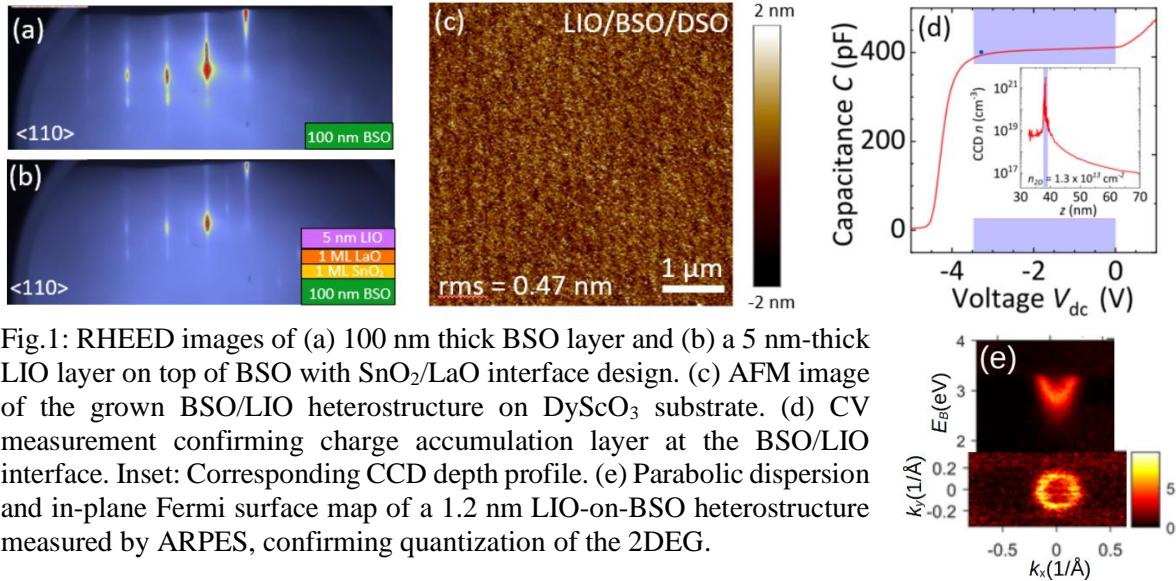


Fig.1: RHEED images of (a) 100 nm thick BSO layer and (b) a 5 nm-thick LIO layer on top of BSO with SnO₂/LaO interface design. (c) AFM image of the grown BSO/LIO heterostructure on DyScO₃ substrate. (d) CV measurement confirming charge accumulation layer at the BSO/LIO interface. Inset: Corresponding CCD depth profile. (e) Parabolic dispersion and in-plane Fermi surface map of a 1.2 nm LIO-on-BSO heterostructure measured by ARPES, confirming quantization of the 2DEG.

We demonstrate the adsorption-controlled growth of the LIO[1] on BSO heterostructure by molecular beam epitaxy using a shutter sequence to control the SnO₂/LaO interface termination. The films were analyzed by reflection high-energy electron diffraction (RHEED) [Figs. 1(a) and 1(b)], x-ray diffraction, atomic force microscopy (AFM) [Fig. 1(c)]. The interface structure is investigated by cross-sectional transmission-electron microscopy. The formation of the quantized 2DEG at their interface is confirmed by capacitance-voltage (CV) [Fig. 1(d)] and angular-resolved photo-electron spectroscopy (ARPES) [Fig. 1(e)]. Van der Pauw-Hall measurements confirm CCD>10¹³cm⁻² and $\mu_{RT}>100$ cm²/Vs.

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⁺ Author for correspondence: bierwagen@pdi-berlin.de