As traditional silicon technology is moving fast towards its fundamental limits, all-oxide electronics is emerging as a challenger offering principally different electronic behavior and switching mechanisms. This technology can be utilized to fabricate devices with enhanced and exotic functionality. One of the challenges for integration of complex oxides in electronics is the availability of appreciable low-temperature synthesis routes. Herein we provide a fundamental extension of the materials toolbox for oxide electronics by reporting a facile route for deposition of highly electrically conductive thin films of LaNiO$_3$ by atomic layer deposition at low temperatures. The films grow epitaxial on SrTiO$_3$ and LaAlO$_3$ as deposited at 225 °C, with no annealing required to obtain the attractive electronic properties. The films exhibit resistivity below 100 $\mu\Omega\text{cm}$ with carrier densities as high as $3.6 \cdot 10^{22}$ cm$^{-3}$. This marks an important step in the realization of all-oxide electronics for emerging technological devices.
Traditional silicon transistor technology is on the verge of reaching its fundamental limits, and nearly 60 years after the postulation of Moore’s law the tech industry is now struggling to maintain continuous evolution in transistor density and computing power. Oxide electronics is considered a viable future contender to Si-based circuitry. Architectures utilizing the plethora of functional properties offered by complex oxides are already important in, e.g., optoelectronics, magnetoelectrics, spintronics and thermoelectrics. Significant research effort is also currently put into the development of complex oxide field effect transistors (COFETs). Armed with fundamentally different switching mechanisms as compared to e.g. silicon-based MOSFETs, the COFETs will offer high-density electron systems that can be switched by lower potentials or different perturbing fields than what is applied in current technology. These novelities are proposed to lead to more energy conserving and faster devices with less heat dissipation due to lower internal resistance: Increasingly important in the search of greener technology.

One of the most notable advantages of complex oxide electronics is the possibility to integrate several functional properties via epitaxy between two or more structurally similar compounds. In addition, the interfaces created between the employed structures are also often found to exhibit novel functional properties themselves.

Complex oxides with the perovskite structure are ideal in terms of epitaxial integration of multiple layers. They are typically close to cubic with pseudocubic lattice parameters in the vicinity of 4 Å, offering the possibility of epitaxial interfaces. Compounds within or related to this structure type include ferromagnetic (e.g. (La,Sr)MnO$_3$), ferroelectric (e.g. Pb(Zr,Ti)O$_3$ and (K,Na)NbO$_3$), high-$\kappa$ dielectric (e.g. SrTiO$_3$ and LaAlO$_3$), superconductor (e.g. YBa$_2$Cu$_3$O$_{7-\delta}$) and multiferroic (e.g. BiFeO$_3$) materials.

Rare-earth nickelates ($\text{RE}_3\text{NiO}_5$, $\text{RE}$ = trivalent rare-earth) have seen extensive research interest over the last few decades. This is for a large part due to a sharp metal-to-insulator transition (MIT) that is correlated to the Ni–O–Ni bond angle, which varies with the size of the rare-earth cation. The Ni–O–Ni bond angle determines the O$_{2p}$–Ni$_{3d}$ orbital overlap, with bands widening when the angle is large (i.e. large orbital overlap). The underlying structural and physical mechanisms behind the MIT is previously extensively reviewed. Lanthanum nickelate, LaNiO$_3$ (LNO), is a special member of the $\text{RE}_3\text{NiO}_5$ group. The larger La$^{3+}$ cation facilitates a bond angle that favors the formation of a metallic rhombohedral structure ($R$-3c) even at absolute zero.

As a result, LNO remains a metal at all temperatures, and conducts electrons as such. Resistivity in single crystals is reported to be $\approx 10^{-4} \Omega \text{cm}$ at room temperature, with a temperature dependence following $\rho = \rho_0 + AT^n$ ($n = 2$). The resistivity and metallic behavior can, however, be tuned by strain engineering. In-plane compression (e.g. on LaAlO$_3$ (100)) leads to less orbital overlap and more narrow bands, while in-plane tensile strain (e.g. on SrTiO$_3$ (100)) leads to wider bands.

Electrically conducting perovskite oxides are rare, and hence LNO is thought to play a role in a wide variety of applications where electronic conductivity is required. Device integration requires LNO to take form as a thin solid layer, preferably epitaxially connected to a substrate. A technologically interesting example is LNO epitaxially integrated with SrTiO$_3$ (STO), effectively making a perovskite oxide stack with a conducting electrode and a high-$\kappa$ dielectric. This can enable the use of STO as a gate oxide in future transistors. Even further down the path this could become part of a fully integrated perovskite oxide electronics system, enabling faster and more energy-efficient electronic devices, e.g. COFETs.

The need for epitaxial integration is, however, also the main caveat for implementation of complex oxides in consumer electronic devices. Techniques used for the deposition of epitaxial complex oxide systems include molecular beam epitaxy and pulsed laser deposition, neither of which is straightforward to implement in a production line due to high-temperature, high-vacuum and limited area synthesis. Chemical techniques like chemical vapor deposition (CVD) are also employed, but the resulting films require post-annealing at $>600 \text{°C}$ to obtain epitaxy. This effectively hampers its applicability in monolithic device integration.

As a result of this, low-temperature-, low vacuum epitaxy of complex oxides on large areas is highly anticipated. The goal is simple: To develop a toolbox of functional materials that can be epitaxially integrated under conditions that are feasible for monolithic device integration. In addition, with increasing geometrical complexity of novel device architectures, synthesis technique that allows strong conformality control on high-aspect ratio substrates becomes more important.

Atomic layer deposition (ALD) is a derivative of CVD where the different precursors are separated as individual pulses in either time or space. This enables self-limiting saturating growth with sub-monolayer thickness control. ALD does not require line-of-sight, enabling deposition on high-aspect ratio and/or hollow substrates, which is important in novel device architectures. Although ALD traditionally have been used to deposit binary oxides, heavier chalcogenides and halides, there has been an increase in reports of functional complex oxides over the last 10 years.

Previous studies of ALD of LNO have resulted in amorphous films as deposited. These films have always required post-annealing to convert into crystalline/epitaxial films, with a lowest reported annealing temperature of 750 $\text{°C}$.

In this study, we report on the use of an alternative precursor chemistry enabling as-deposited epitaxial LNO at 225 $\text{°C}$ on a range of substrates and orientations. No post-annealing is required to obtain metallicity. The films exhibit long-range epitaxy as shown by scanning transmission electron microscopy (STEM) and reciprocal space mapping (RSM) by X-ray diffraction (XRD). The films conduct electrons with bulk-like resistivity and high carrier concentrations characterized by four-point probe and Hall measurements. The temperature dependence of the resistivity reveals that the films behave as metals at all temperatures, following a power law pointing towards Fermi liquid behavior. This novel low-temperature epitaxy of LNO creates a foundation for realization of a range of oxide-based devices where a highly electrically conductive perovskite template material is required, i.e. in COFETs. It extends the toolbox of available low-temperature deposited epitaxial complex oxides with appreciable functional properties.

### Results

**Atomic layer deposition process development.** Multi-cation ALD is traditionally carried out by sequentially employing processes for binary oxides. In the case of LNO, this can be done by combining the La$_2$O$_3$ (La(thd)$_3$ + O$_2$) and NiO (Ni(acac)$_2$ + O$_2$) processes. Note that dry Ni(acac)$_2$ is known to sublimate in the form of an octahedrally coordinated trimer, [Ni(acac)$_3$]$_3$. In this trimer, Ni–O–Ni bonds of similar length to those found in LaNiO$_3$ are already present, and this is believed to be of high importance with respect to epitaxial growth. An important feature of the process is that both binary recipes use ozone as the oxidant. This is especially important in the case of La-containing depositions, as La is prone to form La(OH)$_3$ upon reaction with water, effectively working as a water reservoir which can be detrimental to controlled growth. Also note that nickel is nominally in oxidation state 3+ in LNO, and that a strong oxidant is required to obtain this state.
The La(thd)₃ + O₃ process is known to exhibit controlled growth in the 200–350 °C temperature range, limited downwards by its applicable sublimation temperature (185 °C) and upwards by thermal decomposition of the precursor. The Ni(acac)₂ + O₃ process has a similar low-temperature limit, but Ni(acac)₂ starts to decompose already at around 300 °C. To minimize the thermal budget, but still facilitate controlled growth and a temperature gradient from the precursor source to the reaction chamber, we chose to employ a reactor temperature of 225 °C for the deposition of the La-Ni-O-system.

As both the size and reaction mechanism for the two cation precursors are different, resulting in a different GPC for the two processes, we expect that a 1:1 pulsed ratio of the two binary processes does not result in an equimetric mixture of La and Ni. To understand the relationship between the pulsed and deposited cation ratios, we mapped the cation ratio space by employing a variable La(thd)₃ + O₃ to Ni(acac)₂ + O₃ ratio and measuring the resulting cation stoichiometry by X-ray fluorescence (XRF) (Supplementary Fig. 1). A 5:2 pulsed ratio between La(thd)₃ + O₃ and Ni(acac)₂ + O₃ is found to result in films with a 1:1 cation ratio in the deposited films. The effective ALD-recipe used for the remainder of the study is thus:

\[ 5 \times (\text{La(thd)}_3 + \text{O}_3) + 2 \times (\text{Ni(acac)}_2 + \text{O}_3). \]  

This process facilitates a GPC of ~0.31 Å cycle⁻¹, where a cycle is considered to be one cation precursor pulse followed by one pulse of ozone. Self-limiting behavior was confirmed by employing variable pulse- and purge durations within the complex process (Supplementary Fig. 2).

**Structural analysis by X-ray diffraction.** Thin films of ~30 nm were deposited on a variety of single crystal substrates to determine the structure and functionality of LNO at variable strain. The films were amorphous as deposited on Si (100) (Supplementary Fig. 3), but is remarkably single-oriented crystalline as deposited on STO (100/110/111) and LAO (100/110/111) (Supplementary Fig. 4). This is an unexpected result, as multi-cation deposited films were amorphous as deposited on Si (100) (Supplementary Fig. 3), but is remarkably single-oriented crystalline as deposited on STO (100/110/111) and LAO (100/110/111) (Supplementary Fig. 4). This is an unexpected result, as multi-cation ALD films are typically amorphous and in need of post annealing to become crystalline due to the low deposition temperatures. Similar as-deposited epitaxy is recently shown for ferromagnetic NiFe₂O₄ and antiferromagnetic NiTiO₃ as well as for LaMn₂O₆.

LNO (100) is the most technologically interesting orientation for oxide electronics. Thus, in the remainder of the article, we focus on the STO(100)||LNO(100)pc and LAO(100)pc||LNO(100)pc systems (pc = pseudocubic). Considering the bulk pseudocubic lattice parameter of LNO to be 3.84 Å, LNO will observe a theoretical 1.6% tensile strain on STO (100) and 1.3% compressive strain on pseudocubic LAO (100)pc. This is previously shown to be important for the in-plane conductivity of the material, since the Ni—O—Ni bond angle (and thus the orbital overlap) is dependent on the lattice parameter (Fig. 1). The direction of strain can be confirmed by studying the out-of-plane lattice parameter of LNO in the two substrate systems. Assuming a (near) isovolumetric strain process, tensile in-plane strain leads to compression of the c-axis (out-of-plane), whereas compressive in-plane strain leads to expansion of the c-axis. This is what is observed for the STO(100)||LNO(100)pc and LAO(100)pc||LNO(100)pc systems (Fig. 2). On STO (100) the film exhibits an out-of-plane lattice parameter of 3.822 Å (0.5% compressive strain), while on LAO (100)pc the same parameter is 3.875 Å (0.9% tensile strain). In other words, the c-axis is more strained on LAO (100)pc than on STO (100).

We carried out a Williamson–Hall (WH) analysis of the broadening for three specular reflections to deconvolute broadening from out-of-plane strain and crystallite size. The out-of-plane crystallite size from the WH-analysis was estimated to be 38 nm on LAO (100)pc and 32 nm on STO (100), with a slightly larger c-axis strain on LAO (100)pc. This is consistent with the direct measurement of the c-axis by XRD. The crystallite size is WH-estimated to be close to the thin film thickness, pointing towards continuous crystals traversing all the way from the substrate to the film surface.

Specular XRD of select samples on SrTiO₃ (100) annealed at 650 °C in air was carried out to investigate any relaxation of the
imposed strain that was observed for the as-deposited samples (Supplementary Fig. 5). While the thin film c-axis was slightly compressed upon annealing ($\Delta 2\theta = +0.2^\circ$, $\Delta c = -0.013\,\AA$), very little additional crystallization was observed. Some additional fractional fringes are observed, indicative of a somewhat sharper interface and/or surface. Tuning the strain of the as-deposited films by post-deposition annealing is an intriguing possibility, but one that is left for later investigation.

We further carried out RSM of a specular (002) and an asymmetric reflection (103) to determine the epitaxial relationship (Fig. 3). The position of the reflections reveals the expected cube on cube orientation, with a stringent in-plane coordination (e.g. STO(100)||STO(100)||LNO (100)$_{pc}$||LNO(100)$_{pc}$). Also note that the film scatters at the same $ql$ as the substrate, showing that the film in-plane lattice parameters are strained to be equal to those of the substrate, as expected by the epitaxial relation. We finally carried out a $\varphi$-scan of the LNO (103)$_{pc}$ reflection to investigate the in-plane correlation between the substrate and the film (Supplementary Fig. 6). The (103)$_{pc}$ reflection is clearly fourfold and coupled to the substrate, once again confirming the stringent epitaxial relationship.

**Structural analysis by scanning transmission electron microscopy.** The determination of macroscopic epitaxy as observed by a variety of XRD-methods calls for a local investigation by transmission electron microscopy (TEM). High-angle annular dark field scanning TEM (HAADF-STEM) micrographs of the interface between STO and LNO show the local epitaxial quality (Fig. 4). These micrographs show that the interface is sharp, with well-defined epitaxy, and that there are very few structural defects. On low magnification, a few voids (approx. $\Omega 10\,\text{nm}$) can be observed in the film (Supplementary Fig. 7). The origin of these voids is not clear; however, this is quite commonly observed in thin films made by gaseous deposition and likely stem from encapsulation of either purging gas or byproducts during deposition\textsuperscript{37}. They do not seem to affect the functional properties of the films.

The existence of defects was evaluated by carrying out a direct Fourier transform and ensuing analysis in vertical, horizontal and diagonal directions, showing that the film is near defect free over large areas (Fig. 5). This means that the film has continuously grown epitaxially throughout the deposition, since we did not apply any post-deposition annealing that could have facilitated solid phase epitaxy. Furthermore, solid phase epitaxy on as-deposited amorphous films usually result in interface- and surface roughening as well as introduction of structural defects, which is often detrimental to the sharp interfaces that are needed in a device. The STEM analysis shows that our films are flat and exhibit sharp interfaces, which is essential in multilayer device engineering.

**Chemical analysis by X-ray photoelectron spectroscopy.** We carried out an X-ray photoelectron spectroscopy (XPS) analysis for films deposited on SrTiO$_3$ (100) of Ni 2p (intertwined with La 3d) in an attempt to study the Ni$^{3+}$ to Ni$^{2+}$ ratio (Fig. 6). In stoichiometric LNO the nominal oxidation state of Ni is 3+, but there will always be some reduction of the surface forming an oxygen-deficient LaNiO$_{3-\delta}$ phase. Unfortunately, the overlap between satellite peaks from La 3d make a quantitative analysis difficult. However, using prior literature on close to perfect bulk LNO, we were able to make a comparison pointing towards a high amount of Ni$^{3+}$, with a composition of at least LaNiO$_{2.8}$\textsuperscript{38}. Note that this is the composition in the top nanometers of the film, and that it is very probable that there is an even higher concentration of Ni$^{3+}$ towards the substrate interface.

We further used a peak fitting procedure on a high-quality survey spectra to confirm the composition of the films (Supplementary Fig. 8). This analysis points towards a composition of La$_{1.01}$Ni$_{0.99}$O$_{2.95}$, which again supports a high amount of Ni$^{3+}$ in the films. It should also be mentioned that the oxygen content here is likely to be slightly overestimated due to the existence of carbonate species on the surface, as there is a small fraction of carbon (<2 at.%) as analyzed by the survey spectrum.
Electrical properties. With confirmation that the structural and chemical quality of the LNO films are high, we turned towards an investigation of the electrical properties. Four-point probe measurements (point distance 1 mm) in room temperature reveal metallic electrical resistivity. STO(100)||LNO(100)_pc exhibits a resistivity of ~100 µΩ cm on average, with a lowest measured resistivity of 80 µΩ cm. This is below the reported values for bulk LNO, which is explained by the tensile strain that LNO is perturbed by an STO, increasing the Ni−O−Ni bond angle and widening the bands. Note that post-deposition annealing of these samples provides very little added functionality, with lowest measured resistivity of 70 µΩ cm. For LAO(100)_pc||LNO(100)_pc the average resistivity measured was ~300 µΩ cm, which is slightly higher than bulk LNO. This is explained by the compressive strain, decreasing the Ni−O−Ni bond angle and narrowing the bands. Note that films down to a thickness of 2.5 nm has been characterized and found to conduct with similar resistivity as the 30 nm films that are presented here.

To further investigate the electrical properties of the sample, we carried out a Hall analysis using a four-point setup where each corner of the 1·1 cm² sample is contacted to a probe (Fig. 7). The film is perturbed by a 1.02 T magnet, which allows for measurements of induced Hall currents, and deduction of carrier densities.

The resistivity was estimated to be 138 µΩ cm for STO (100)||LNO(100)_pc at ambient, which is slightly higher than measured by the four-point probe. This is likely because the Hall-set up uses a much larger probe distance (10× longer), which will be less tolerant to any grain boundaries that may exist in the film. It should still be noted that the resistivity is on par with that of bulk LNO (~120 µΩ cm)\(^{15}\).

The induced Hall voltage is measured to be ~2 mV A⁻¹. Using the film conductivity, sample thickness, current and magnetic field, a Hall coefficient of 0.0549 mm³ C⁻¹ and a charge carrier density of ~3.6·10²² cm⁻³ can be deduced. This is very close to the theoretical 3.4·10²² cm⁻³ carriers that would be present if each NiO₆-octahedron contributes with one carrier each. High carrier density is key for materials that are to be used as gates in oxide electronics.

We measured the temperature-dependent resistivity of the LNO films on STO using a PPMS cooled by liquid He. The temperature was swept from RT to 6 K and the resistivity was collected for every 2 K (Fig. 8).

The decrease in resistivity as a function of temperature underpins that the films are metallic across the whole temperature range. LNO with significant oxygen under stoichiometry will exhibit an MIT at a set temperature. Thus, the temperature-dependent resistivity complements our previous identification that our method can be used to deposit films with both high chemical and structural quality (i.e. low oxygen under stoichiometry). Note that resistivity measurements were carried out for pristine films and films that have been exposed to humidity over a long time, with no apparent degradation of the functionality.

Finally, we fitted the temperature dependence of the resistivity to a linear combination of fifth-, third- and second-order polynomials to identify the mechanism for resistivity, in compliance with Matthiessen’s rule. The dependence is strongly correlated to the second-order polynomial \(\rho = 27.6 + 0.012T^2 + 2.02·10^{-5}T^3 + 4.09·10^{-12}T^4\) (µΩ cm), indicating that the resistivity is mainly governed by electron—electron interactions. This is as expected in a correlated electron system like LaNiO₃. The constant term, \(\rho_0\), is found to be 27.6 µΩ cm, which is slightly higher than previously
reported for single crystals. This may be a result of the previously described oxygen vacancies, which has a higher concentration towards the film surface.

Discussion

We have developed an ALD process for LaNiO₃ that can be used to obtain highly epitaxial thin films on LaAlO₃ and SrTiO₃ substrates. The films are epitaxial as deposited at 225 °C and need no further post-deposition annealing to obtain the highly regarded functional properties. The films are nearly defect free as shown by XRD, X-ray photoelectron spectroscopy and TEM. They conduct electrons with low resistivity (~10⁻⁴ Ω cm) and with high charge carrier density (~3 × 10²² cm⁻³). The imposed strain from the substrate affects the electrical properties by changing the in-plane Ni−O−Ni bond angle, and thus the width of orbital bands.

The applicability of LaNiO₃ thin films in device construction has been significantly hampered by the need for high deposition temperatures or high-temperature post-deposition annealing. This new process maintains a low thermal budget that should be compatible with most device fabrication, and especially in monolithic device integration in transistor technology. The direct low-temperature deposition of LaNiO₃ on SrTiO₃ at 225 °C marks a fundamental step in realizing SrTiO₃ gate oxides, and furthermore as an electrode in future all-oxide electronics, e.g. in COFETs (Fig. 9).

Methods

Atomic layer deposition. Thin film depositions were carried out in an F-120 Sat ALD reactor (ASM Microchemistry). The deposition temperature was 225 °C with an operating pressure of 2.4 mbar, maintained by a 300 cm³ min⁻¹ primary flow rate of N₂. Nitrogen was supplied from gas cylinders (Praxair, 99.99%) and run through a Mykrolis purifier for removal of any oxygen or water contamination. La(thd)₃ (Volatex, 99%), and Ni(acac)₂ (Sigma Aldrich, 97%) were used as cation precursors (acac = acetylacetonate). Both precursors were supplied from open boats inside the reactor, and maintained at 185 °C throughout the deposition. Ni(acac)₂ was resublimated at 175 °C for purification prior to use in the reactor. The cation precursors were pulsed into the reaction chamber by means of inert gas valves. O₃ was used as the oxygen source, made from O₂ (Praxair, 99.5%) using an AC-2505 (In USA) ozone generator supplying 15 wt.% O₃ in O₂. Pulse durations were 2 and 4 s for La(thd)₃, Ni(acac)₂ and O₃, respectively. Purge durations were 2 s after cation precursor pulses, and 3 s after the ozone pulses. Confirmation of self-limiting behavior for the employed pulse- and purge scheme is shown in Supplementary Fig. 2.

The thin films were deposited on 1 · 1 cm² Si for routine characterization of thickness and 3 · 3 cm² Si for analysis of conformity and cation stoichiometry. Selected compositions were deposited on LaNiO₃ (LNO) (100) (Crystal GmbH), (110) and (111) (MTI Corp.) and SrTiO₃ (STO) (100) (Crystal GmbH), (110) and (111) (MTI Corp.) single crystals for facilitation of epitaxial growth.

Thickness analysis. Routine measurements of thickness were carried out on an alpha-SE spectroscopic ellipsometer (J.A. Woollam) in the 390–900 nm wavelength range. We successfully employed a Cauchy function to model the collected data. X-ray reflectivity (XRR) was used as a complementary method to determine thickness for very thin films (<10 nm). XRR was carried out on an Emmepi diffractometer (Panalytical) equipped with a Cu Kα₀ (λ = 1.5406 Å) source powered at 45 kV and 40 mA, and a parallel beam mirror.

Chemical and compositional analysis. Chemical composition was analyzed using an Axios Max Minerals X-ray fluorescence (XRF) system (Panalytical), equipped with a 4 kW Rh-tube. The system runs using Omnistep and Stratos options for standardless measurement of thin films. Complementary compositional analysis and evaluation of chemical environment was carried out by XPS, using a Theta Probe Angle-Resolved XPS (Thermo Scientific). This instrument is equipped with an Al Kα₀ (hv = 1486.6 eV) source and the analysis chamber was maintained at ~10⁻⁸ mbar during analysis. Pass energy values of 200 and 60 eV were employed for survey and detailed scans, respectively. The spectra were analyzed using the Thermo Advantage software suite.

X-ray diffraction. X-ray diffraction for structural analysis was carried out on an AXS D8 Discover (Bruker) diffractometer, equipped with a LynxEye strip detector and a Ge (111) focusing monochromator, providing CuKα₀ (λ = 1.5406 Å) radiation. Reciprocal space maps were collected with a Pixel 3D detector on an Empyrean diffractometer (Panalytical) equipped with a Cu Kα₀ (λ = 1.5406 Å) source powered at 45 kV and 40 mA, and a primary Barlths monochromator.

Scanning transmission electron microscopy. Scanning transmission electron microscopy was carried out using a JEOL 2100F microscope at 200 kV. Sample preparation was carried out by mechanical polishing and ion milling on a Gatan PIPS II. Collected high-angle annular dark field (HAADF) images were treated and Fourier-filtered using the Gatan Microscopy Suite software.

Electrical characterization. Room-temperature resistivity measurements were carried out using a four-point probe and aKeithley model 2400 SourceMeter. The sheet resistivity was recorded by measuring resistance in ten points from 1 to 10 µΩ. Variable temperature resistivity measurements were performed on a Model 4000 physical property measurement system (PPMS, Quantum Design). The samples were mounted on a puck and contacted with gold wires on gold pads deposited by evaporation. Resistivity was collected in a four-point setup, while the temperature was swept from 300 to 6 K.

Data availability

The data sets generated and analyzed during the current study are available from the corresponding author upon reasonable request.

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References

1. Colli, M. et al. Towards oxide electronics: a roadmap. Appl. Surf. Sci. 482, 1–93 (2019).
2. Zubiko, P., Gariglio, S., Gabay, M., Ghozse, P. & Triscone, J.-M. Interface physics in complex oxide heterostructures. Annu. Rev. Condens. Matter Phys. 2, 141–165 (2011).
3. Vaz, C. A. F., Hoffman, J., Ahn, C. H. & Ramesh, R. Magnetoelectric coupling effects in multiferroic complex oxide composite structures. Adv. Mater. 22, 2900–2918 (2010).
4. Rondinelli, J. M., Stengel, M. & Spaldin, N. A. Carrier-mediated magnetoelectricity in complex oxide heterostructures. Nat. Nanotechnol. 3, 46–50 (2008).
5. Bjaalie, L., Himmetoglu, B., Weston, L., Janotti, A. & Van De Walle, C. G. Oxide interfaces for novel electronic applications. N. J. Phys. 16, 025005 (2014).
6. Penicheva, R. & Pickett, W. E. Electronic phenomena at complex oxide interfaces: insights from first principles. J. Phys. Condens. Matter 22, 043001 (2010).
7. Yu, P., Chu, Y.-H. & Ramesh, R. Oxide interfaces: pathways to novel phenomena. Mater. Today 15, 320–327 (2012).
8. Hwang, H. Y. et al. Emergent phenomena at oxide interfaces. Nat. Mater. 11, 103–113 (2012).
9. Catalano, S. et al. Rare-earth nickelates RNiO$_3$: thin films and heterostructures. Rep. Prog. Phys. 81, 014501 (2018).
10. Varghese, J., Grigorieva, M. N., Itzuy, J., Barthelémy, A. & Bibes, M. Complete phase diagram of rare-earth nickelate from first-principles. npj Quantum Mater. 2, 21 (2017).
11. Mercy, A., Biedner, J., Itzuy, J. & Ghosez, P. Structurally triggered metal-insulator transition in rare-earth nickelates. Nat. Commun. 8, 1677 (2017).
12. Lee, S., Chen, R. & Balents, L. Metal-insulator transition in a two-band model for the perovskite nickelates. Phys. Rev. B 84, 165119 (2011).
13. Peil, O. E., Hampel, A., Ederer, C. & George, A. Mechanism and control parameters of the coupled structural and metal-insulator transition in nickelates. Phys. Rev. B 99, 245127 (2019).
14. Jaramillo, R., Ha, S. D., Silveit, D. M. & Ramanathan, S. Origins of bad-metal conductivity and the insulator-metal transition in the rare-earth nickelates. Nat. Phys. 10, 304–307 (2014).
15. Zhang, J., Zheng, H., Ren, Y. & Mitchell, J. F. High-pressure floating-zone growth of perovskite nickelate LaNiO$_3$ single crystals. Cryst. Growth Des. 17, 2730–2735 (2017).
16. Zhou, J. S., Marshall, L. G. & Goodenough, J. B. Mass enhancement versus Stoner enhancement in strongly correlated metallic perovskites: LaNiO$_3$ and LaCuO$_3$. Phys. Rev. B 89, 245138 (2014).
17. Cao, Y., Lin, B., Sun, Y., Yang, H. & Zhang, X. Sr-doped lanthanum nickelate nanofilbers for high energy density supercapacitors. Electrochem. Acta 174, 41–50 (2015).
18. Singh, R. N. et al. Preparation and characterization of thin films of LaNiO$_3$ for anode application in alkaline water electrolysis. J. Appl. Electrochem. 24, 149–156 (1994).
19. Hwang, D. K., Kim, S., Lee, J.-H., Hwang, I.-S. & Kim, I.-D. Phase evolution of perovskite LaNiO$_3$ nanofilbers for supercapacitor application and p-type gas sensing properties of LaOCo$_{1-x}$NiO$_x$ nano-composites. J. Mater. Chem. 21, 1959–1965 (2011).
20. Scherwitzl, R., Zubko, P., Lichtensteiger, C. & Tricsson, J. M. Electric-field tuning of the metal-insulator transition in ultrathin films of LaNiO$_3$. Appl. Phys. Lett. 95, 222114 (2009).
21. Aytug, T. et al. Growth and characterization of conductive SrRuO$_3$ and LaNiO$_3$ multilayers on textured Ni tapes for high-Jc YBa$_2$Cu$_3$O$_7$–Y. J. Mater. Res. 16, 2661–2669 (2001).
22. Golakhi, M. et al. Nature of the metal-insulator transition in few-unit-cell-thick LaNiO$_3$ films. Nat. Commun. 9, 2206 (2018).
23. Satyalakshmi, K. M. et al. Epitaxial metallic LaNiO$_3$ thin films grown by pulsed laser deposition. Appl. Phys. Lett. 62, 1233 (1993).
24. Kuprenaite, S. et al. Relationship processing–composition–structure–resistivity of LaNiO$_3$ thin films grown by chemical vapor deposition methods. Coatings 9, 35 (2019).
25. Miyazaki, H. et al. Preparation and evaluation of LaNiO$_3$ thin film electrode with chemical solution deposition. J. Eur. Ceram. Soc. 24, 1005–1008 (2004).
26. George, S. M. Atomic layer deposition: an overview. Chem. Rev. 110, 111–131 (2010).
27. Senstebey, H. B., Fjellvåg, H. & Nilsen, O. Functional perovskites by atomic layer deposition—an overview. Adv. Mater. Interfaces 4, 1600903 (2017).
28. Mackus, A. J. M., Schneider, J. R., Macisca, C., Baker, J. G. & Bent, S. F. Synthesis of doped, ternary, and quaternary materials by atomic layer deposition: a review. Chem. Mater. 31, 1142–1183 (2019).
29. Coll, M. & Napari, M. Atomic layer deposition of functional multicomponent oxides. APL. Mater. 7, 110901 (2019).
30. Seim, H., Mölså, H., Nieminen, M., Fjellvåg, H. & Niinistö, L. Deposition of LaNiO$_3$ thin films in an atomic layer epitaxy reactor. J. Mater. Chem. 7, 449–454 (1997).
31. King, P. J. et al. Toward epitaxial ternary oxide multilayer device stacks by atomic layer deposition. J. Vac. Sci. Technol. A 37, 020602 (2019).
32. Nieminen, M., Putkonen, M. & Niinistö, L. Formation and stability of lanthanum oxide thin films deposited from β-diketonate precursor. Appl. Surf. Sci. 174, 155–166 (2001).
33. Utirainen, M., Kröger-Laukkonen, M. & Niinistö, L. Studies of NiO thin film formation by atomic layer epitaxy. Mater. Sci. Eng. B 54, 98–103 (1998).
34. Nilsen, Ø., Rauwel, E., Fjellvåg, H. & Kjekshus, A. Growth of La$_2$-Ca$_x$MnO$_{3-y}$ thin films by atomic layer deposition. J. Mater. Chem. 17, 1466–1475 (2007).
35. Bratvold, J. E., Fjellvåg, H. & Nilsen, O. Atomic layer deposition of oriented nickel titanate (NITiO$_3$). Appl. Surf. Sci. 311, 478–483 (2014).
36. Bratvold, J. E., Senstebey, H. H., Nilsen, O. & Fjellvåg, H. Control of growth orientation on a-deposited epitaxial iron-rich nickel ferrite spinel. J. Vac. Sci. Technol. A 37, 021502 (2019).
37. Bazzi, C. et al. Role of nitrogen in defect evolution in zinc oxide: STEM–EELS nanoscale investigations. J. Phys. Chem. Lett. 10, 4725–4730 (2019).
38. Mickevičius, S. et al. Investigation of epitaxial LaNiO$_{3−x}$ thin films by high-energy XPS. J. Alloy. Compd. 423, 107–111 (2006).
39. Ha, S. D., Aydögü, G. H. & Ramanathan, S. Metal-insulator transition and electrically driven memristive characteristics of SmNiO$_3$ thin films. Appl. Phys. Lett. 98, 012105 (2011).
40. Shi, J., Ha, S. D., Zhou, Y., Schoofs, F. & Ramanathan, S. A correlated nickelate syanptic transistor. Nat. Commun. 4, 2676 (2013).
41. Dai, S. et al. Recent advances in transistor-based artificial synapses. Adv. Func. Mater. 29, 1903790 (2019).

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Author contributions
H.H.S. initiated and proposed the basis for the project, deposited several samples, performed PS characterization, and took part in XRD-characterization, resistivity measurements and Hall measurements, and wrote the bulk of the paper. E.S. deposited thin films and took part in structural characterization and physical property measurements, and took part in revising the manuscript. O.S.F. carried out TEM characterization, collection of reciprocal space maps and φ-scans, and took part in physical property measurements, and also took part in revising the manuscript. J.F.B. deposited thin films and took part in revising the manuscript. A.E.S. and J.O.N. supervised the project and took part in writing the manuscript. O.N. supervised the project and took part in writing the manuscript.

Competing interests
The deposition process (not the resulting films) is pending UK patent application 1910404.1, filed July 19, 2019. Inventors: Henrik H. Senstebey, Jon E. Bratvold, Ola Nilsen and Helmer Fjellvåg at the University of Oslo, Norway.

Additional information
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