Silane-Mediated Expansion of Domains in Si-Doped \(\kappa\)-Ga\(_2\)O\(_3\) Epitaxy and its Impact on the In-Plane Electronic Conduction

Piero Mazzolini,*, Zsolt Foga\'rassy, Antonella Parisini, Francesco Mezzadri, David Diercks, Matteo Bosi, Luca Seravalli, Anna Sacchi, Giulia Spaggiari, Danilo Bersani, Oliver Bierwagen, Benjamin Moritz Janzen, Marcella Naomi Marggraf, Markus R. Wagner, Ildiko Cora, B\'ela P\'ecz, Abbas Tahraoui, Alessio Bosio, Carmine Borelli, Stefano Leone, and Roberto Fornari

1. Introduction

For almost a decade gallium oxide (Ga\(_2\)O\(_3\)) has summoned significant interest in the scientific community because of its potential in the field of power electronics.\([1,2]\) This material has five different known polymorphs,\([1]\) all of them characterized by an ultra-wide bandgap of \(\approx 5\) eV. The monoclinic \(\beta\)-Ga\(_2\)O\(_3\) is the thermodynamically stable crystal structure; for this reason, it has been so far the most investigated one with different proposed device architectures, most of them based on homoepitaxy\([1]\) due to the possibility to synthesize bulk \(\beta\)-Ga\(_2\)O\(_3\) from the melt.\([2,3]\) Moreover, the ability for i) tuning its bandgap through Al- and In-alloying,\([4,5]\) ii) controlling its electrical properties through n-type extrinsic doping (e.g., Si,\([6]\) Sn\([6]\)), and iii) depositing epitaxial layers with low densities of structural defects and smooth surfaces (because...
of the presence of perfectly matched substrates, i.e., $\beta$-Ga$_2$O$_3$ interfaces.\cite{9, 10, 11, 12} Concurrently, the experimental demonstration of the confinement of 2D electron gases (2DEGs) at $\beta$-Ga$_2$O$_3$/Al$_2$O$_3$ interfaces.\cite{13} Nonetheless, the low symmetry of the monoclinic cell poses serious obstacles for future device development, i.e., the challenging heteroepitaxial\cite{14, 15} as well as homoepitaxial\cite{9, 10, 16, 17} growth of high quality $\beta$-Ga$_2$O$_3$ thin films and the intrinsic anisotropy of some of its functional properties (e.g., thermal conductivity).\cite{18} For these reasons, other polymorphs of Ga$_2$O$_3$ are recently gaining attention. Particularly, $\kappa$-Ga$_2$O$_3$ for power electronic applications. Both of these polymorphs in fact have a higher unit cell symmetry compatible with relatively cheap crystalline substrates like sapphire. Moreover, for both of these metastable polymorphs a relatively wide window of synthesis parameters have been identified for various physical- as well as chemical- vapor phase epitaxial deposition techniques (PVD and CVD, respectively) that allow for their single phase stabilization.\cite{19}

In particular, orthorhombic $\kappa$-Ga$_2$O$_3$ possesses a peculiar spontaneous polarization along the (001) direction, predicted to be larger than the one of GaN (23–26 $\mu$Ccm$^{-2}$), which should allow for the confinement of high mobility 2DEGs at $\kappa$-Ga$_2$O$_3$/Al$_2$O$_3$ heterointerfaces characterized by higher carrier densities ($n \approx 10^{14}$ cm$^{-2}$) with respect to the ones obtainable in $\beta$-Ga$_2$O$_3$ heterostructures and in principle without the need of modulation doping.\cite{20, 21} Nonetheless, despite the possibility of depositing high quality (001)-oriented $\kappa$-Ga$_2$O$_3$/Al$_2$O$_3$ heterostructures on cheap c-plane sapphire substrates, no 2DEG has been so far experimentally reported.\cite{22} This is probably linked to the real structure of this orthorhombic polymorph\cite{23} and in particular to the presence of rotational domains commonly found in heteroepitaxial films despite different employed substrates\cite{24} with the exceptions of recently reported single domain layers obtained on an epitaxially matched GaFeO$_3$ substrate\cite{25} and on an offcut c-plane sapphire.\cite{26} In fact, epitaxial (001)-oriented $\kappa$-Ga$_2$O$_3$ thin films are characterized by a peculiar columnar structure with $\approx 10$ nm width.\cite{27} This morphology is related to the presence of 120°-rotated orthorhombic $\kappa$-Ga$_2$O$_3$ domains, that should be nominally free of dangling bonds,\cite{28} and initially caused an erroneous attribution to a pure hexagonal cell of the $\varepsilon$-Ga$_2$O$_3$ polymorph. For this reason, both the $\varepsilon$ and $\kappa$ nomenclatures can be found in literature even if referring to the very same polymorph. In this manuscript we refer to the orthorhombic polymorph with the correct $\kappa$ nomenclature. However, it is not trivial to obtain a fine control of the electrical properties in $\kappa$-Ga$_2$O$_3$ thin films: to the best of our knowledge, no PVD extrinsically doped $\kappa$-Ga$_2$O$_3$ electrically conducting layer has ever been reported in literature. On the other hand, CVD deposited layers, and in particular metal-organic vapour phase epitaxy (MOVPE), already demonstrated the possibility to extrinsically dope the material during the deposition process with the employment of a silane (SiH$_4$) flow, or ex-situ via thermally-induced Sn diffusion.\cite{29} Particularly, in the case of Si, a partial solubility into $\kappa$-Ga$_2$O$_3$ is expected to take place mostly as $\text{Si}_{\text{Ga}}$ substitutional in the tetrahedral Ga$_3$ lattice sites resulting in a shallow donor state.\cite{30} Nonetheless, the extrinsically doped layers suffered from a limited electron mobility (at room temperature $\mu \approx 1$ cm$^2$V$^{-1}$s$^{-1}$) due to a hopping transport mechanism.\cite{29, 31} As recently suggested by Kneiß et al.\cite{10} and some authors of this work,\cite{11} this could be linked to the nanometric-sized rotational domains representing an effective barrier for the in-plane electronic conduction, i.e., a perturbation in the lateral crystal periodicity comparable to the electrons’ mean free path.\cite{29} Therefore, finding synthesis conditions that could at the same time allow the extrinsic doping of the material and a domain size control would represent a major step forward for the understanding of the real $\kappa$-Ga$_2$O$_3$ intrinsic material properties, which is in turn fundamental for its breakthrough in the field of new generation electronic devices.

In this article we demonstrate that the typical rotational domain size in electrically conducting Si-doped $\kappa$-Ga$_2$O$_3$ layers on c-plane sapphire substrates can increase from $\approx 10$ to $\approx 300$ nm. Particularly, we show that this is closely related to the addition of silane during the MOVPE deposition, which could either increase the mobility of the adsorbed species during growth or decrease the initial nuclei density, resulting in a larger domain formation. Moreover, we provide i) non-destructive characterization techniques and procedures (X-ray diffraction XRD and Raman spectroscopy) that allow relative comparisons of the mean rotational domain size in $\kappa$-Ga$_2$O$_3$ epitaxial films and ii) a direct experimental proof (i.e., in-plane versus out-of-plane resistance measurements performed on insulated mesa structures with well-defined geometries) that the mean domain dimension is itself playing a fundamental role—in combination with planar defects—in the determination of the in-plane electronic conduction. We believe that these findings are of paramount importance for the full physical understanding of the $\kappa$-Ga$_2$O$_3$ material system, paving the way for overcoming its defects-driven limitations which have so far inhibited its application in the field of power electronics.

Moreover, the general occurrence of rotational domains in heteroepitaxy is common for the growth of lower symmetry (epilayer) over higher symmetry (substrate) materials. Their presence and size distribution can generally result in significant consequences on the functional properties of the epilayer. In particular, as discussed by Grundmann et al.\cite{32} the rotational domains could affect the impurity distribution and the concentration of deep levels in the layer, cause reduced carrier diffusion length, mobility and lifetime, or lower the efficiency of radiative recombination; this can in turn detrimentally affect the properties of various devices, e.g., breakdown voltage, leakage current, or quantum efficiency. Apart from the $\kappa$-Ga$_2$O$_3$ here investigated, also the heteroepitaxy of the $\beta$ polymorph on c-plane sapphire is affected by the presence of rotational domains that negatively reflects on its functional properties.\cite{14, 15} Similarly, in other heteroepitaxial material systems like the orthorhombic SrRuO$_3$ deposited on SrTiO$_3$,\cite{13} LiNbO$_3$ and LiTaO$_3$ on Si,\cite{14} Heusler Ni–Mg–Ga alloys on MgO,\cite{15} or the monoclinic BiFeO$_3$ on LaSrAlO$_3$ and SrTiO$_3$,\cite{33} the domain engineering plays a fundamental role with application fields ranging from spintronics and memory devices, radio-frequency acoustic filters and photonic devices, memory shape alloys and magnetocaloric effect, or piezoelectricity. The experimental approach here proposed combines several nondestructive as well as destructive techniques that could be generally advantageous for the thorough characterization of various
domain-structured epilayers, a key requirement when these structural defects can be considered as functional elements.

2. Results

2.1. X-Ray Diffraction

Figure 1a reports the typical XRD symmetric out-of-plane diffraction pattern obtained for the investigated (001)-oriented \(\kappa\)-Ga\(_2\)O\(_3\) layers on c-plane sapphire, which has been found to be qualitatively similar for all the different silane fluxes (i.e., no additional diffraction peaks were identified). The crystal structure is purely orthorhombic and all the deposited thin films are characterized by the presence of 120° in-plane rotated domains as exemplarily shown in Figure 1b by the \(\phi\)-scan performed on the (122) reflection.[37] For sake of clarity, the four expected peaks associated with each of the three rotational domains were calculated and highlighted (labelled with Roman numbers I, II, and III) in Figure 1b and perfectly match the experimental data. The full width at half maximum in 2\(\theta\)-\(\theta\) scans is not significantly affected by the silane flow \(\Phi_{\text{SiH}_4}\) [the mean FWHM for the most intense (004) peak is \(\approx 0.057^\circ\) with scattered variations among the samples below 10%, see Figure S1, Supporting Information]. The incorporation of Si in the \(\kappa\)-Ga\(_2\)O\(_3\) matrix should preferably occur as a Ga substitution (Si\(\rightarrow\)Ga) in the tetrahedral sites as previously shown via electron paramagnetic resonance.[29] The c unit cell parameter (evaluated from 2\(\theta\)-\(\theta\) XRD scans) does not seem to be majorly affected by the \(\Phi_{\text{SiH}_4}\), in our samples: the recorded c variations (between \(\approx 9.260\) and 9.275 Å) are well in line with previously reported data for the orthorhombic Ga\(_2\)O\(_3\) polymorph.[38] This is likely related to the limited amount of provided Si, as well as the non-trivial role of the Si-Ga aliovalent substitution and related defects (see following subsections).

Noteworthy, the increment of the \(\Phi_{\text{SiH}_4}\) provided during the deposition process is also accompanied by a significant reduction of the rocking curve FWHM values [\(\omega\)-scans of the most intense (004) peak, see Figure 1c], indicating a decrease of the crystal mosaicity with respect to the substrate normal. In particular, a decrease from a maximum of 0.371° (nominally

![Figure 1. XRD a) 2\(\theta\)-\(\theta\) scan and b) \(\phi\)-scan of a (001) \(\kappa\)-Ga\(_2\)O\(_3\) layer deposited on c-plane sapphire with 10 sccm silane flux; in b), the Roman numbers I, II, III, and respective orange, green, and cyan lines refer to the calculated and expected positions for the three 120° rotated domains. c) Full width at half maximum (FWHM) in \(\omega\)-scans (rocking curve) of the (004) peak as a function of the \(\Phi_{\text{SiH}_4}\). The green arrows in c) highlight the samples that have been further investigated by TEM.](image-url)
undoped, i.e., $\Phi_{\text{SiH}_4} = 0$) down to a minimum of $0.133^\circ$ ($\Phi_{\text{SiH}_4} = 15$ sccm) is recorded. The displayed set of rocking curve values obtained in these $\kappa$-Ga$_2$O$_3$ layers deposited with silane (mostly $< 0.2^\circ$ for $\Phi_{\text{SiH}_4} \geq 7.5$ sccm, see Figure 1c) are among the best reported values in literature for this material system, regardless of the presence of an extrinsic dopant/ligand, the employment of different substrates, or various CVD or PVD growth techniques.[23,25,39–44] The only sensibly lower rocking curve value published so far ($0.08^\circ$) refers to an undoped $\kappa$-Ga$_2$O$_3$ layer deposited on an offcut c-plane sapphire substrate and referred to as a single-domain.[27]

Three selected samples (0, 5, and 15 sccm $\Phi_{\text{SiH}_4}$, among the highlighted by green arrows in Figure 1c) were further investigated through XRD reciprocal space mapping (RSM) in the vicinity of the (122) orthorhombic $\kappa$-Ga$_2$O$_3$ reflection (see Figure S2 and Table S1, Supporting Information). In line with the symmetric out-of-plane measurements of the (004) reflection (Figure 1c), the (122) $\omega$ FWHM showed a clear difference between the two Si-doped samples and the nominally undoped one (Table S1, Supporting Information). It should be noted that the [122] direction is $\approx 54^\circ$ away from the surface normal, so a relevant component of the scattering vector is in-plane. Similarly, the 2$\theta$ FWHM values extracted from the RSM measures show a significant reduction as $\Phi_{\text{SiH}_4}$ is introduced during growth (from 0.667$^\circ$ for the layer without $\Phi_{\text{SiH}_4}$ to 0.412$^\circ$ and 0.428$^\circ$ for the 5 and 15 sccm samples, see Table S1, Supporting Information).

### 2.2. In-Plane Transport

Similar to the highlighted trend in the rocking curve values (Figure 1c), also the room-$T$ resistivity $\rho$ shows a pronounced decrease between 0 ($\approx 10^{-1}$–$10^{-4}$ $\Omega$cm) and 7.5 sccm $\Phi_{\text{SiH}_4}$ ($\approx 10^{-3}$ $\Omega$cm), while a further increase of the $\Phi_{\text{SiH}_4}$ did not result in any other sensitive $\rho$ variations, i.e., $\rho$ plateaus at $\approx 0.5$ $\Omega$cm apart from a single value ($\rho \approx 2$ $\Omega$cm) recorded for the highest tested silane flow ($\Phi_{\text{SiH}_4} = 17.5$ sccm) (see Figure 2a). Particularly, a minimum value of $\rho \approx 0.2$ $\Omega$cm is reached for $\Phi_{\text{SiH}_4} = 15$ sccm. The $T$-dependent ($T = 300–50$ K) resistivity $\rho(T)$ of the investigated layers points toward the electronic conduction

![Figure 2](https://example.com/figure2.png)

Figure 2. Room temperature, a) electrical resistivity $\rho$, b) mobility $\mu$, and c) charge carrier density $n$ extracted from van der Pauw–Hall electrical measurements of Si-doped $\kappa$-Ga$_2$O$_3$ layers as a function of the $\Phi_{\text{SiH}_4}$ provided during the deposition. The green arrows highlight the samples that have been further investigated by TEM.
Figure 3. Plan view TEM images of samples deposited at $\Phi_{\text{SiH}_4} = 0 - 5 - 15 - 17.5$ sccm. A schematical representation of the typical plane defects identified in top view images is highlighted by blue (inside the domains) and red (domain walls) arrows for the 15 sccm sample. An EDS line profile crossing various extended defects with corresponding Si and Ga concentrations extracted for the same sample is reported; circles and arrows help to identify the corresponding image-to-atomic fraction positions for the presented EDS line profiles.

being dominated by a variable-range hopping transport mechanism (VRH) as previously shown for this material system,$^{[28,29]}$ despite the employment of different $\Phi_{\text{SiH}_4}$ (Figure S3, Supporting Information). When $\ln(\rho)$ is plotted versus $T^{-0.5}$, i.e., VRH for independent electrons (Mott-VRH),$^{[45]}$ the natural logarithm of $\rho$ shows linear dependencies (Figure S3b, Supporting Information), whose slopes and absolute values significantly decrease from nominally undoped to doped samples. It is worth noting that $\ln(\rho)$ seems to show linear behavior even when plotted versus $T^{-0.25}$, i.e., ES-VRH for interacting electrons (Efros-Shklovskii ES-VRH).$^{[46]}$ Unfortunately, the variation of the resistivity magnitude along a linear trait is not wide enough to permit to unambiguously distinguish between the two exponents, i.e., ES-VRH or Mott-VRH. Nonetheless, the proximity to the semiconductor to metal transition$^{[29]}$ and going toward higher temperatures$^{[47]}$ should favor the Mott- over the ES-VRH transport mechanism (see Supporting Information for additional details).$^{[48]}

Interestingly, the RT mobility $\mu$ increases about one order of magnitude while increasing the provided $\Phi_{\text{SiH}_4}$ (from $\approx 1$ up to $\approx 10$ Vcm$^{-2}$s$^{-1}$, Figure 2b). Such mobility values are typical of a transport between localized states, about one-to-two orders of magnitude lower than room-temperature mobilities of $\beta$-Ga$_2$O$_3$ (i.e., in the range of 50 to 100 Vcm$^{-2}$s$^{-1}$ for $\beta$ with similar $n$ of about $10^{18}$ cm$^{-3}$)$^{[49]}$ The comparison between the monoclinic and the orthorhombic polymorph is relevant due to their similar electron effective masses.$^{[29,50,51]}$ Conversely, the RT $\tau$ is found to follow an opposite trend with the increment of $\Phi_{\text{SiH}_4}$, decreasing from $n = 3 - 5 \times 10^{18}$ down to $n = 1 \times 10^{18}$ cm$^{-3}$ while increasing the $\Phi_{\text{SiH}_4}$ (Figure 2c).

2.3. Transmission Electron Microscopy–TEM

Four selected samples deposited at different silane flows (i.e., $\Phi_{\text{SiH}_4} = 0 - 5 - 15 - 17.5$ sccm, highlighted with green arrows in Figure 1c and Figure 2) were further investigated via TEM and EDS. Plan view images (Figure 3) confirm that the orthorhombic structure is affected by the presence of 120° rotated domains in all samples (XRD in Figure 1b, TEM Fourier transform patterns in Figure S6, Supporting Information). Of note, the increment of the provided $\Phi_{\text{SiH}_4}$ results in an evident qualitative enlargement of the domains (Figure 3). Nonetheless, a quantitative assessment of their mean size is not easily achievable due to their irregular shape. Therefore, the area of the domains for each analyzed sample has been evaluated for the reported plan view images in Figure 3. The evaluated areas were translated into regular circles and their diameters were considered as the grain size (full analysis reported in Figure S4, Supporting Information). In line with previous reports,$^{[24]}$ the typical domain size of a nominally undoped sample is $\approx 5$ to 20 nm wide. Nonetheless, a clear trend between the maximum domain size and the $\Phi_{\text{SiH}_4}$ is identified (Figure 3). In particular, the maximum domain size is observed passing from $\approx 20$ nm up to $\approx 320$ nm for $\Phi_{\text{SiH}_4}$ of 0 and 175 sccm, respectively.

The HRTEM plan view images at higher magnification allow the identification of domain boundaries and plane defects inside the domains (exemplarily highlighted by arrows in a plan image of a 15 sccm layer in Figure 3). EDS line profiles from the TEM plan views allowed for roughly (due to the Si signal being close to the detection limit) estimating the amount of Si with respect to Ga ($\approx 1\%$ for $\Phi_{\text{SiH}_4}$ of 15 sccm, see exemplary line profile scan in Figure 3; Figure S5, Supporting Information). For the 17.5 sccm sample the value was fluctuating over a large range ($\approx 2.5-3.5\%$, Figure S5, Supporting Information). Moreover, the investigated line profiles suggest a slight Si accumulation located in the vicinity of defects, i.e., domain boundaries and plane defects (e.g., red arrow in the EDS line profile of the 15 sccm sample in Figure 3 highlighting a domain boundary). Nonetheless, the possible Si-accumulation on the defect sites is not clearly present in all the investigated line.
profiles (e.g., see additional 15 and 175 sccm TEM plane scans and EDS line profiles in Figure S5, Supporting Information).

Additional cross-sectional TEM images (Figure 4) allowed the identification of a γ-Ga2O3 interlayer at the sapphire/κ-Ga2O3 interface with an irregular thickness ranging between 2 and 25 nm for all the investigated samples (see Fourier transform patterns of selected interface areas in Figure S6, Supporting Information). The presence of a γ interlayer has been already highlighted for this material/substrate system deposited via MOVPE.[24] Amorphous SiOx precipitates with a triangular shape were present in all the investigated samples deposited in the presence of a ΦSiH4 (Figure 4; Figure S7, Supporting Information). Interestingly, i) the presence of these precipitates seems to be limited to just the first 200–300 nm of the layer (i.e., closer to the substrate interface) and ii) their mean size diminishes with increasing ΦSiH4 (from ≈ 50 nm down to ≈ 2–5 nm base length for 5 and 17.5 sccm, respectively—see Figure 4), while their density increases.

Moreover, the combined plan-view and cross-sectional TEM investigation shows that generally the domain walls run straight from the bottom (γ-Ga2O3/sapphire interface) up to the layer surface. Nonetheless, the plane defects identified inside the domains are mostly not perfectly columnar and in the case of the employment of ΦSiH4, the presence of triangular SiOx precipitates is found to affect the formation/distribution of them (see blue arrows in Figure 4 for a 17.5 sccm layer). As a result, despite the generally larger width of the rotational domains while increasing the ΦSiH4, the concentration of plane defects is also suggested to increase (Figure 4).

2.4. Atom Probe Tomography

Since the combined TEM-EDS analysis intrinsically suffers from a large uncertainty in the determination of the Si concentration and its spatial distribution, APT has been performed on two focused ion beam FIB-prepared specimens of samples deposited at ΦSiH4 of 5 and 15 sccm (same layers investigated via TEM-EDS). These specimens were obtained close to the film surfaces, i.e., far from the substrate interface were the γ phase and the presence of SiOx precipitates was identified by TEM investigation (Figure 4). The TEM dark field images of the two analyzed APT specimens displayed in Figure 5 show a larger density of straight defects for the 5 sccm with respect to the 15 sccm ΦSiH4, reasonably associated to rotational domain boundary walls. Therefore, these data qualitatively confirm the previous independent TEM investigation (Figure 3). The mass spectra of the whole analyzed volumes via APT allow for quantification of the atomic percentage of Si with respect to Ga, that is found to be 0.54% and 1.02% for ΦSiH4 of 5 and 15 sccm respectively (Figure S8, Supporting Information). These values are well above a minimum detection level of 52 ppm determined using a 95% confidence interval and the method of reference,[52] and generally in good agreement with TEM-EDS data (Figure 3).

To investigate the presence of Si-doping clusters inside the orthorhombic crystal matrix and/or SiOx precipitates small enough to be possibly not highlighted by TEM imaging, a silicon nearest neighbor distribution statistical analysis has been performed on both of the APT analyzed samples (Figure S8, Supporting Information). If there were Si-clustering, the nearest neighbor distance of Si atoms from each other would be different than the nearest neighbor distance between a randomized assignment of the atoms. In neither of the two analyzed sample volumes does that appear to be the case, i.e., the Si dopant is evenly spread in the κ-Ga2O3 matrix independently of its overall percentage. As already pointed out, this does not appear to be the case for the first 200–400 nm of the doped films (thickness region probed exclusively by the TEM investigation, see Figure 4) where the presence of SiOx precipitates has

Figure 4. Cross sectional TEM and HRTEM for samples deposited at different ΦSiH4. The white arrows highlight some of the SiOx precipitates and blue arrows plane defects located at the apexes of the triangular precipitates.
already been highlighted. Therefore, different from the qualitative indications coming from EDS line profiling in plan TEM view (Figure 3; Figure S5, Supporting Information), no Si accumulation at defects sites has been highlighted by APT. Nonetheless, just in the 15 sccm sample it has been possible to spot two distinct regions with homogeneous but slightly different Si content, i.e., 0.95% and 1.25% for the low and high concentration sides of the analyzed volume (Figure 5). It is reasonable to assume that such variation could be related to a slightly different content of Si in adjacent orthorhombic domains.

2.5. Raman Spectroscopy

Undoped and doped κ-Ga2O3 layers were further analyzed by confocal Raman spectroscopy in back-scattering geometry from the sample surface and edge. Surface back-scattering measurements were performed on all the samples investigated by TEM and APT (ΦSiH4 = 0.5, 15, 17.5 sccm), with two different laser wavelengths (633 nm), with the first being the one reported in Figure S5, Supporting Information), producing similar results. Cross-sectional (edge) measurements were performed on the same ΦSiH4 = 0.5, 15 sccm samples and on an additional 10 sccm one, with the 532 nm laser wavelength in parallel polarization to the c-plane. All spectra were collected in the 50–800 cm−1 range. Their analysis is performed after air spectrum subtraction, to remove artifacts from its undesired contributions at low Raman shift (60–80 cm−1), and after normalization with respect to the most intense peak from the sapphire substrate (417 cm−1).

Only modes typical of the orthorhombic polymorph of Ga2O3 were identified (aside from the ones of the c-plane sapphire), consistent with the XRD and TEM investigations. Despite the different analysis configurations, both surface and edge spectra equivalently highlight similar differences among the analyzed samples, especially when undoped and doped layers are compared (Figure 6a; Figure S9a, Supporting Information).

In particular, the peak-fitting analysis of selected peaks in both configurations, suggests the presence of the Si dopant (i.e., from the introduction of a ΦSiH4 during deposition) consistently results in a variation of the intensity ratio among some of the Raman modes (e.g., 82 and 113 cm−1, insets in Figure 6a; Figure S9a, Supporting Information). Noteworthy, the increasing domain size of samples deposited at larger ΦSiH4 (Figure 3) is accompanied by a strong decrease of the Raman peaks’ FWHM (Figure 6b,c; Figure S9b,c, Supporting Information). In cross-sectional spectra, where substrate contribution is less prominent and κ-Ga2O3 peaks are better resolved, this evidence is stronger (Figure 6). Moreover, the largest reduction of the line width of all phonon modes is observed between the undoped and 5 sccm sample.
2.6. Out-of-Plane versus In-Plane Transport Measurements

Finally, to enable a direct measurement on the possible defect-induced (e.g., domains) conductivity anisotropy in epitaxial κ-Ga$_2$O$_3$ thin films, ≈110 and 1100 nm thick, nominally undoped films (i.e., $\Phi_{\text{SiH}_4} = 0$) were deposited on top of C-doped and Si-doped GaN epitaxial templates on c-plane sapphire substrates (insulating and conducting templates, respectively; XRD reported in Figure S10, Supporting Information). The MOVPE chamber has been conditioned before the deposition of these layers so to avoid the possible role of Si unintentional incorporation after growth runs in presence of $\Phi_{\text{SiH}_4}$ (potentially leading to lower resistivities, see Supporting Information).

The samples deposited on insulating GaN templates as well as a 590 nm thick layer directly deposited on c-plane sapphire under the very same deposition conditions have been used for the determination of the in-plane transport properties of the κ-Ga$_2$O$_3$ layers, resulting in a mean $\rho_{\text{in-plane}}$ of $≈8 \times 10^5$ Ωcm (measurements performed on similar TLM contact structures, see Figure S11, Supporting Information).

To measure the out-of-plane transport, squared mesa structures of κ-Ga$_2$O$_3$ with well-defined geometries ($200 \times 200 \times 150 \times 150 \times 100 \times 70 \times 70 \mu$m$^2$, height equal to the κ-Ga$_2$O$_3$ layer thicknesses, i.e., 1100 and 110 nm) were then isolated via dry etching on semiconducting GaN templates (66).

Several mesas for both κ-Ga$_2$O$_3$ deposited layer thicknesses (i.e., 1100 and 110 nm) were electrically characterized via 2-point probe configuration. Considering the case of anisotropic resistance, the total area through which the current is supposed to flow during the out of plane measurements is limited by the top contact size (see schematic representation of a mesa structure in Figure 7a), i.e., $A_{\text{top contact}} = 180 \times 180 \times 130 \times 130 \times 80 \times 80 - 20 \times 20 - 55 \times 55 \mu$m$^2$.

As pointed out in Figure 7c, several series contributions add up to the experimentally measured resistance $R_{\text{measured}}$, out of which we are interested in the intrinsic out-of-plane resistance ($R_{\kappa \text{-Ga}_2\text{O}_3}$) of the κ-Ga$_2$O$_3$ mesa. Given the

![Figure 6](https://example.com/figure6.png)

**Figure 6.** a) Raman spectra in cross sectional (edge) configuration (parallel polarization) of κ-Ga$_2$O$_3$ samples deposited at $\Phi_{\text{SiH}_4} = 0 \text{–} 5 \text{–} 10 \text{–} 15$ sccm. In the inset the two peaks at $≈82$ and $113$ cm$^{-1}$ are highlighted and compared for the undoped and 15 sccm samples. In b) and c), the fitted peak widths (color/shape correspondence to the symbols reported on the Raman spectra) are reported as a function of the $\Phi_{\text{SiH}_4}$; this analysis is performed on selected peaks (i.e., choosing those that are clearly visible in the spectra collected in both configurations, see Figure S9, Supporting Information) and displaced in two separate graphs for sake of clarity.
well-defined geometries of the isolated mesas, $R_{\kappa\text{-Ga}_2\text{O}_3}$ could be translated into the out-of-plane resistivity, $\rho_{\text{out-plane}} = (R_{\kappa\text{-Ga}_2\text{O}_3} A_{\text{top contact}})/(\kappa\text{-Ga}_2\text{O}_3$ layer thickness), which can be readily compared to the in-plane resistivity $\rho_{\text{in-plane}}$ (evaluated for the material deposited under the very same growth conditions on the insulating GaN templates and sapphire, see Figure S11, Supporting Information). Despite the order of magnitude difference in the two investigated $\kappa\text{-Ga}_2\text{O}_3$ layer thicknesses, the $R_{\text{measured}}$ for the different mesa dimensions did not scale, i.e., $R_{\text{measured}, \ 1100 \text{ nm}} \approx R_{\text{measured}, \ 110 \text{ nm}}$ (blue squares and orange circles respectively in Figure 7b). This evidence points toward the series resistances being dominant in the 2-point-probe measurements, i.e., not allowing for a direct determination of $R_{\kappa\text{-Ga}_2\text{O}_3}$. Notwithstanding, it is unquestionable that $R_{\text{measured}}$ provides an upper-bound estimate for $R_{\kappa\text{-Ga}_2\text{O}_3}$, i.e., $R_{\kappa\text{-Ga}_2\text{O}_3} < R_{\text{measured}}$.

Therefore, $R_{\text{measured}}$ can be used to provide the upper-bound estimate of the out-of-plane resistivity, i.e., $\rho_{\text{out-of-plane upper bound}} = (R_{\text{measured}} A_{\text{top contact}})/(\kappa\text{-Ga}_2\text{O}_3$ layer thickness) $> \rho_{\text{out-of-plane}}$. Figure 7d compares the resulting upper-bound estimate $\rho_{\text{out-of-plane upper bound}}$ for the 1100 nm thick $\kappa\text{-Ga}_2\text{O}_3$ film (i.e., being limited by the series resistances in this estimate) to the $\rho_{\text{in-plane}}$, giving a lower bound estimate for the transport anisotropy. It is therefore experimentally found that the out-of-plane resistivity

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**Figure 7.** a) Schematic representation (not to scale) of a mesa structure obtained for the $\kappa\text{-Ga}_2\text{O}_3$ thin films deposited on the conductive epitaxial GaN layer. In (b), the experimental out-of-plane $R_{\text{measured}}$ (mediated data among several measured mesa structures; full set of data in Figure S12, Supporting Information) as well as the estimated $R_{\text{isotropy}}$ (from $\rho_{\text{in-plane}}$ measurements, see Figure S11, Supporting Information) are reported for both the investigated $\kappa\text{-Ga}_2\text{O}_3$ thicknesses as a function of the squared lateral top contact size. In (c), the different series resistance contributions embedded in $R_{\text{measured}}$ are reported in the current two-terminal probe configuration. In (d), the $\rho_{\text{out-of-plane upper bound}}$ evaluated for the measured mesa structures with different dimensions is reported for the 1100 nm thick $\kappa\text{-Ga}_2\text{O}_3$; in the same graph the straight black line is highlighting the $\rho_{\text{in-plane}}$ with the top and bottom lines (gray shaded area) representing the standard deviation (data coming from Figure S11, Supporting Information).
is at least one order of magnitude lower with respect to the in-plane resistivity in such \( \kappa \)-Ga\(_2\)O\(_3\) layers.

3. Discussion

The size of the 120° rotated orthorhombic domains in (001) \( \kappa \)-Ga\(_2\)O\(_3\) epitaxial layers on c-plane sapphire substrates (Figure 1) is usually limited to 5–20 nm width. Our results showed that the addition of a silane flow \( \Phi_{\text{sil}} \), during the MOVPE deposition results in n-type Si extrinsic doping while at the same time helping to increase the domain size by more than one order of magnitude (Figure 3), both of them contributing to a clear decrease of in-plane resistivity \( \rho_{\text{in-plane}} \) and increase of in-plane electron mobility \( \mu \) (Figure 2a,b). The reason behind the domain size control in \( \kappa \)-Ga\(_2\)O\(_3\) samples as a function of the \( \Phi_{\text{sil}} \), could be related to a silane-induced increase of the surface mobility of the adsorbed species during the growth process (especially during nucleation of the orthorhombic phase). In the MOVPE growth of Ga\(_2\)O\(_3\) with TMG metallorganic precursor and H\(_2\) as gas carrier, hydrogen is supposed to replace the leaving methane resulting in GaH to be adsorbed on the surface.[31] The oxidizing agent (i.e., H\(_2\)O in this study) was provided in excess with respect to the TMG flow. Several Ga-O clusters could then possibly form and diffuse on the growth surface (i.e., Ga\(_2\)O\(_2\), Ga\(_2\)O\(_4\), Ga\(_3\)O\(_5\), and GaO\(_2\)), some of which might also desorb (e.g., Ga\(_2\)O\(_4\)), although in H\(_2\)O excess the latter desorption should not play an important role.[31] In the MOVPE deposition of \( \kappa \)-Ga\(_2\)O\(_3\) the growth temperature can not usually exceed 650 °C in order to prevent the stabilization of the monoclinic \( \beta \) polymorph (i.e., \( T_g \) can be just varied on a narrow window to increase the surface mobility).[32] Our samples were deposited at the same \( T_g = 610 \) °C and TMG/H\(_2\)O flow ratio, and the employment of various \( \Phi_{\text{sil}} \), was the only tuned parameter. A precise discussion of the growth process (e.g., unambiguous identification of the mobile adsorbed species on the surface during growth, role of the surface they are diffusing and nucleating on, i.e., c-plane sapphire or \( \gamma \)-Ga\(_2\)O\(_3\), see Figure 4) would require a dedicated study that is out of the scope of this article. Nevertheless, based on the presented results, it is reasonable to assume that silane could be capable of increasing the mobility of the adsorbed species on the growth surface, resulting in an overall lower density of nuclei. This could be triggered by Si or SiH\(_4\) accumulating on the surface during growth and preventing in this way the formation of critical size nuclei, i.e., resulting in an increasing diffusion length of the adsorbed species. Such a mechanism has already been proposed for the growth of GaN nanorods by MOCVD in the presence of silane[34] and could qualitatively explain the increasing domain size in \( \kappa \)-Ga\(_2\)O\(_3\) while increasing \( \Phi_{\text{sil}} \), (Figure 3). The partial incorporation of Si is also suggested by the APT-measured Si concentration on the two samples deposited with a threecfold different \( \Phi_{\text{sil}} \), (5 and 15 sccm) resulting in a just twofold different Si concentration in the probed volume (0.54% and 1.02%, respectively see Figure S8, Supporting Information). On the other hand, the growth rate of the layers remained unaltered with respect to the \( \Phi_{\text{sil}} \).

In this framework, we could also interpret the presence of the SiO\(_x\) precipitates highlighted in the first 200–400 nm of the samples deposited in the presence of a \( \Phi_{\text{sil}} \), (Figure 4). Since the \( \Phi_{\text{sil}} \), is continuously provided during growth, once it overcame a critical density on the surface, this could result in the precipitation of SiO\(_x\) inclusions (see HRSTEM, HAADF images and EDS maps in Figure S7, Supporting Information) in the \( \kappa \)-Ga\(_2\)O\(_3\) matrix. The precipitates are amorphous and characterized by a not perfectly triangular shape (HRTEM micrographs reported in Figure 4), with their flat side being mostly parallel to the substrate/surface, and their size and density being related to the \( \Phi_{\text{sil}} \). In particular, larger silane flows result in higher density of smaller triangular precipitates (Figure 4). Nonetheless, the SiO\(_x\) precipitates being just limited to the region close to the substrate surface remains an open point that requires further dedicated investigations (TEM, Figure 4; APT, Figure 5).

The electrical properties of the \( \kappa \)-Ga\(_2\)O\(_3\) layers are affected by the addition of the \( \Phi_{\text{sil}} \), during growth for a threefold reason: i) Si being a shallow donor in this material system,[29] ii) the domain walls and iii) structural (e.g., plane defects) as well as compensation/self-compensation defects both playing a role in limiting the in-plane electronic conduction. Regarding i), the presented results suggest that not only Si, but also H is likely to play the role of a shallow donor in \( \kappa \)-Ga\(_2\)O\(_3\), since the nominally undoped layers (\( \Phi_{\text{sil}} = 0 \)) deposited with H\(_2\) as a carrier gas show an in-plane room temperature \( \rho \) in the order of \( 10^{3−10^4} \) Ω cm (Figure 2a; Figure S11, Supporting Information). It should be noted that different MOVPE synthesis parameters (TMG/H\(_2\)O flux ratios, H\(_2\) carrier gas flows, and especially the use of different gas carriers, e.g., He)[35] can result in much higher resistivities for layers deposited without \( \Phi_{\text{sil}} \), (\( \rho \geq \rho_{\text{VRH}} \)). In all the deposited layers the trend of \( \rho \) in \( T \) (Figure S3, Supporting Information) is suggested to be ruled by a variable range hopping transport (VRH, see Figure S3 and related discussion in the Supporting Information). Nonetheless, the RT \( \rho \) is found to decrease while increasing \( \Phi_{\text{sil}} \), during the deposition (Figure 2a). In particular, a significant drop of \( \rho \) is recorded up to \( \Phi_{\text{sil}} \), while afterward its value seems to go toward saturation being mostly scattered around mid-to-low \( 10^{-3} \) Ω cm. While ii) the domain size is found to significantly increase together with the silane flow at least up to \( \Phi_{\text{sil}} = 15 \) sccm (Figure 3), it is reasonable to assume that the iii) disorder—i.e., compensation/self-compensation and/or plane defects—would also increase. The occurring \( \rho \) saturation could be related to these two competing effects. Particularly, together with the rotational domain size increase with \( \Phi_{\text{sil}} \), the SiO\(_x\) precipitates are also increasing in density, with the latter also interacting with plane defects (Figure 4). The precipitates/plane defects and possible (self-) compensation effects can both contribute to increase the disorder, in turn possibly affecting the VRH transport mechanism.

Also non-uniform Si distribution could play a role in the electronic transport. TEM-EDS line profiles seem to suggest possible Si accumulation at defects interfaces (i.e., both domain walls and plane defects, see Figure 3; Figure S5, Supporting Information). On the other hand, the more sensitive and spatially resolved APT technique excluded the presence of Si inhomogeneities/aggregates in presence of defects in the analyzed volumes (i.e., close to the surface of the deposited films, see Figure 5; Figure S8, Supporting Information). Nonetheless, APT performed on a layer deposited at relatively high \( \Phi_{\text{sil}} \),
(15 sccm) reveals local Si concentration differences (e.g., 0.95% and 1.25%, Figure 5) that have been attributed to different domains, although the Si concentrations are homogeneously distributed inside the domains themselves (Figure S8, Supporting Information).

In the Hall effect characterization (Figure 2b,c), the Hall voltage must be handled with caution in the case of hopping conduction.\[45\] The experimental Hall density \(n\) is related to the density of donor sites \(N_d(T)\) taking part to the hopping process, which is usually considered evenly distributed over the film thickness. Surprisingly, it is here reported that \(n\) tends to decrease while increasing \(\Phi_{\text{sat}}\) (Figure 2c). A \(n\) reduction implies an increased compensation and/or a reduction of the donor sites. An effect of reduction of effective thickness of the conductive channel due to the SiO\(_x\) precipitates (Figure 4; Figure S7, Supporting Information) and possible depleted regions around them, even if possible, cannot explain a reduction of the Hall density of a factor of about five (Figure 2c). Differently, the mobility \(\mu\) increase (Figure 2b) would not be affected by the eventual overestimation of the conduction channel; therefore, this should be related to a rising hopping probability which can be qualitatively associated with the widening of the domains.

The Hall-effect absolute values of \(n\) should be interpreted with caution in this particular case (i.e., VRH transport mechanism and not well-defined thickness of the conductive channel). Nonetheless, a comparison among the extracted \(n\) and the APT measurements (Figure 2c; Figure S8, Supporting Information) allows to highlight a high level of compensation. For example, the APT measured layer deposited at 15 sccm \(\Phi_{\text{sat}}\), is characterized by a Si concentration of \(\approx 1\%\). For a full dopant activation mechanism once overcoming a certain threshold Si concentration) resulting in the favorable formation of deep acceptor defects (e.g., Ga vacancies \(V_{\text{Ga}}\), and b) by the incorporation of Si in sites different from the Ga substitutional ones (e.g., interstitial, or as SiO\(_x\) precipitates as highlighted in the first 200–400 nm of the doped layers). In (a), also a self-passage mechanism once overcoming a certain \(E_f\) level (e.g., a threshold Si concentration) resulting in the favorable formation of deep acceptor defects (e.g., \(V_{\text{Ga}}\), to maintain the crystal charge neutrality could be considered, like in the case of the \(\beta\)-Ga\(_2\)O\(_3\) polymorph.\[56\] In light of the VRH transport, a significant increase of compensating defects while increasing the \(\Phi_{\text{sat}}\) can have detrimental effects on the density of hopping carriers, while the hopping mobility could be mostly beneficially affected by the concuring increase of the domain size. This simplified picture could explain the puzzling behavior of the Hall density and mobility with the silane flow. In this direction at least on very different domain size distributions (i.e., from \(< 20 \text{ nm}\) to \(> 100 \text{ nm}\)), it is worth mentioning that the \(\kappa\)-Ga\(_2\)O\(_3\) material system has significantly decreased when the \(\Phi_{\text{sat}}\) is in-plane. In this light both the different \(\omega\) and \(\theta\) FWHM extracted values (Table S5, Supporting Information) significantly decreased when the \(\Phi_{\text{sat}}\) was introduced during growth. Particularly, the smaller FWHM values in the \(\omega\) direction indeed can be directly linked to the increased mean crystallite size, in agreement with TEM characterization (Figure 3; Figure S4, Supporting Information). Nonetheless, it should be also considered that the \(\theta\) FWHM could be as well related to strain and/or microstructural effects. In this framework it is difficult to evaluate the role of the inhomogeneously distributed SiO\(_x\) precipitates (just limited to the region close to the substrate surface as highlighted by TEM—Figure 4—and electronic conduction results in the experimentally demonstrated defect-driven conduction anisotropy in \(001\) epitaxial \(\kappa\)-Ga\(_2\)O\(_3\) layers. In particular, the data reported in this work quantitatively show an at least (lower estimate due to series resistances, see Figure 7) one order of magnitude higher in-plane resistivity (i.e., charge carriers crossing domain interfaces) than out-of-plane resistivity (i.e., charge carriers flowing along the vertically oriented domains), in line with previously reported qualitative experimental data.\[30\] In this framework it is important to stress out that theoretical\[50,57\] and experimental\[26\] works on \(\kappa\)-Ga\(_2\)O\(_3\) as well as the comparison among similarly structurally anisotropic oxides (e.g., \(\beta\)-Ga\(_2\)O\(_3\)\[58\] and SnO\(_2\)[59]) exclude the possibility that the extent of this highlighted (lower bound) anisotropy could be possibly related to the pure orthorhombic structure of Ga\(_2\)O\(_3\). The measurements here reported have been performed on nominally undoped samples (\(\Phi_{\text{sat}} = 0\), maximum domain dimension around 20 nm, see Figure 3) deposited on GaN templates; this has been done to avoid the presence of SiO\(_x\) precipitates (confined in the first 200–300 nm close to the substrate interface in layers deposited with \(\Phi_{\text{sat}}\), see Figure 4) that could potentially interfere in the out-of-plane R measurements. The demonstrated defects-driven anisotropy in the \(\kappa\)-Ga\(_2\)O\(_3\) material system is most likely the reason that so far prevented the isolation of 2DEGs in \(\kappa\)-Ga\(_2\)O\(_3\)/Al\(_x\)Ga\(_{2-x}\)O\(_3\) heterostructures,\[23\] and more generally so far inhibited the application of this material system in the field of power electronics. The deep understanding of the growth process and the employment of proper substrates\[26,27\] will soon together open the attainment of single domain \(\kappa\)-Ga\(_2\)O\(_3\) semiconducting layers with tunable electrical properties.

As pointed out, the characterization of the domain structure—i.e., both their presence and typical associated width—is fundamental for the \(\kappa\)-Ga\(_2\)O\(_3\) epitaxial layers. Therefore, it is important to highlight the thorough non-destructive structural characterization here proposed via XRD and Raman spectroscopy. At first, the significantly reduced FWHM in the XRD \(\omega\)-scans of the out-of-plane (004) reflection as a function of the \(\Phi_{\text{sat}}\) (Figure 1c) highlights its clear relation with the different orthorhombic domain size in these samples. Moreover, the RSM measurements in the vicinity of the (122) reflection confirmed the possibility of getting qualitative information at least on very different domain size distributions (i.e., from \(< 20 \text{ nm}\) to \(> 100 \text{ nm}\)), it should be noted that the (122) direction is \(\approx 54^\circ\) away from the surface normal, so a relevant component of the scattering vector is in-plane. In this light both the different \(\omega\) and \(\theta\) FWHM extracted values (Table S5, Supporting Information) significantly decreased when the \(\Phi_{\text{sat}}\) was introduced during growth. Particularly, the smaller FWHM values in the \(\omega\) direction indeed can be directly linked to the increased mean crystallite size, in agreement with TEM characterization (Figure 3; Figure S4, Supporting Information). Nonetheless, it should be also considered that the \(\theta\) FWHM could be as well related to strain and/or microstructural effects. In this framework it is difficult to evaluate the role of the inhomogeneously distributed SiO\(_x\) precipitates (just limited to the region close to the substrate surface as highlighted by TEM—Figure 4—and
APT—Figure S8, Supporting Information) with size and concentration being a function of the $\Phi_{\text{SiH}_4}$.

Finally, a clear indication of increasing mean domain size is independently provided also by Raman spectroscopy. In particular, the consistent trend of the Raman peaks narrowing highlighted in both confocal and cross-sectional configurations (Figure 6; Figure S9, Supporting Information) may be generally related to phonon lifetimes being affected by the different domain sizes of the investigated layers. More in detail, phonons in the samples with the smallest domains of $\approx 5-20 \text{ nm}$ (i.e., no $\Phi_{\text{SiH}_4}$) are significantly affected by boundary scattering at the domain wall, resulting in a reduction of the phonon mean free path and phonon lifetimes. Consequently, the Raman modes experience a significant line width broadening. Upon silane introduction, the increased domain size reduced the contributions of domain boundary scattering, thus a strong reduction in the Raman line width is observed (see Figure 6b,c; Figure S9b,c, Supporting Information). The effect becomes less significant for even larger domains $>100 \text{ nm}$ (i.e., $\Phi_{\text{SiH}_4} > 5 \text{ sccm}$) as other scattering mechanisms contribute to the limitation of the phonon lifetimes. Moreover, Raman has also shown high potential for the investigation of the dopant role in this material system, i.e., intensity change of specific peaks (e.g., 82 and 113 cm$^{-1}$, see insets in Figure 6a; Figure S9a, Supporting Information). This aspect should be further investigated in a dedicated work.

4. Conclusions

The rotational domain size is among the major issues affecting the in-plane electronic transport in $\kappa$-Ga$_2$O$_3$, a polymorph with significant, but so far unrealized potential in the field of new generation power electronics. In this framework, the ability to control their size upon silane-mediated growth demonstrated in this work (i.e., from about tens to about hundreds of nanometers) represents a fundamental step forward for the obtainment of an ultra-wide bandgap material with tailored semiconducting properties. Being of particular significance for this material system, we have here demonstrated and provided experimental guidelines to get fine and qualitative information for the characterization of the domain size distribution in $\kappa$-Ga$_2$O$_3$ (001)-oriented epitaxial layers with two widely employed and non-destructive characterization techniques, i.e., XRD and Raman spectroscopy. Moreover, in-versus out-of-plane electrical measurements experimentally allowed to measure the defect-mediated lower bound limit of resistivity anisotropy in epitaxial $\kappa$-Ga$_2$O$_3$ layers. In particular, the out-of-plane resistivity has been found to be at least one order of magnitude lower than the in-plane resistivity. This i) once more highlights the detrimental role of domains in $\kappa$-Ga$_2$O$_3$ while ii) suggests that their enlargement/suppression together with the limitation of plane and compensation defects—is of paramount importance to improve and tailor the in-plane conduction, therefore highlighting the full potential of this material. Moreover, the thorough characterization of a domain structured epilayer here proposed is of wider relevance considering different material systems where such structural defects rule the resulting functional properties.

5. Experimental Section

The $\kappa$-Ga$_2$O$_3$ layers were grown by MOVPE at a partial pressure of 60 mbar and substrate temperature $T_s = 610 \degree \text{C}$ on nominally oriented c-plane sapphire substrates. Trimethylgallium (TMG) and ultrapure $\text{H}_2\text{O}$ stored in dedicated stainless-steel bubblers at 1 and 30 $\degree \text{C}$ respectively, were used as metal precursor and oxidizing gas. $\text{H}_2$ was used as gas carrier with a flow of 2000 standard cubic centimeters per minute (sccm). The ratio of the precursors pressures ($P_{\text{TMG}}/P_{\text{TMH}} \approx 350$), as well as the deposition time ($t_{\text{dep}} = 100 \text{ min}$, resulting deposition thickness $\approx 800-900 \text{ nm}$) (as determined using a JVC LV-1700, via V-530 spectrophotometer$^{[60]}$) had been kept constant for all the growth runs. Different flows of an $\text{H}_2$-diluted mixture of 0.05% $\text{SiH}_4$ were provided while maintaining the other synthesis parameters in different growth runs ($\Phi_{\text{SiH}_4}$ from 0 to 17.5 sccm, where $\Phi_{\text{SiH}_4}$ refers to the $\text{H}_2$-diluted $\text{SiH}_4$ mixture). The investigation ratio between the partial pressure of effective $\text{SiH}_4$ and partial pressure of TMG ($P_{\text{SiH}_4}/P_{\text{TMG}}$) ranged from 0 to $8.7 \times 10^{-2}$.

The deposited layers were characterized by XRD using a Rigaku Smartlab XE diffractometer with CuK$_\alpha$ wavelength, equipped with crossed beam optics unit to achieve parallel beam geometry, Eulerian cradle and HyPix3000 detector. Symmetric out-of-plane 2$\theta$ and $\omega$ rocking curves as well as $\varphi$-scans on asymmetric reflections were performed to get information on the crystal quality and domain structure of the orthorhombic $\kappa$-Ga$_2$O$_3$ layers.$^{[58]}$ Reciprocal space mapping (RSM) had been performed in order to qualitatively investigate the mean size of the columnar domains among samples deposited with different $\Phi_{\text{SiH}_4}$. Confocal Raman spectroscopy measurements were obtained using a Horiba LabRAM HR Evolution confocal spectrometer (focal 800 mm), using a configuration with a 600 lines mm$^{-1}$ grating, a set of Bragg ultra-low frequency filters for the rejection of the Rayleigh scattering and a liquid nitrogen-cooled CCD detector. The 633 and 532 nm laser lines were used for excitation ($P \leq 1 \text{ mW}$). An optical 100x objective was used to obtain a spatial resolution of $\approx 1 \mu \text{m}$, with $0.5 \text{ cm}^{-1}$ resolution. Typical collection time for the confocal configuration was 60 s for each acquisition, with 10 repetitions. Additional measurements in cross-sectional configuration$^{[39]}$ with a $\lambda = 532 \text{ nm}$ were performed with another LabRAM HR 800 spectrometer (Horiba Jobin-Yvon).

The (scanning) transmission electron microscopy (TEM) was carried out in an aberration corrected Thermo Fischer Scientific THEMIS microscope at 200 kV on selected samples. The cross-sectional TEM samples were prepared by focused ion beam (FIB) technique. The plan view TEM samples were prepared by conventional Ar-ion beam milling. For the energy dispersive X-ray spectroscopy (EDS) mapping a Super-X detector was used in STEM mode. In order to get quantitatively reliable information on the amount of Si ($N_s$) in this layers and its distribution with a nanometric scale resolution, atom probe tomography (APT) had been employed on two selected samples deposited at different $\Phi_{\text{SiH}_4}$. The APT specimens were prepared using a FIB lift-out method in an FEI Helios Nanolab 600i and a Tescan SB252G instrument. The specimens were mounted on tungsten half-grids in holders compatible with the FIB, TEM, and APT instruments.$^{[81]}$ The FIB-prepared APT specimens were analyzed by dark and bright field TEM imaging at 200 kV in an FEI Talos F200X TEM before and after the APT analysis in order to identify and spatially locate the presence of structural defects in the probed volume of the material and assist in generating the reconstructions. Laser pulsed APT analysis was performed in a Cameca LEAP 4000X Si using a 90 mm flight path length at a temperature of 50 K. A laser energy of 40 pJ at repetition rates of 333–500 kHz was used. The detection rate was fixed at 6 ions per 1000 pulses that resulted in a bias range of 3200–6300 V during the analyses. A systematic energy deficit correction was applied to the resulting data to improve the mass spectral resolution by correcting for electrostatic effects.$^{[82]}$ Reconstructions were generated using Cameca’s IVAS 3.6.18 software with the TEM images of the specimens before and after analysis providing guidance for constraining the reconstruction parameters.$^{[81]}$

All the $\kappa$-Ga$_2$O$_3$ layers were electrically characterized at room temperature (RT) via 4-point probe van der Pauw configuration,
performing resistivity as well as Hall measurements (Keithley Hall-measurement setup, magnetic field of 0.8 T). For a selected number of samples, an additional investigation in the 10–300 K temperature range was performed. For this purpose Ti/Au[28] or SnO2/ITO/Au[4] Ohmic contacts were fabricated with a sputtering technique on the corners of squared shaped samples.

Additional κ-Ga2O3 layers were deposited on electrically conducting (Si-doped) as well as insulating (C-doped) GaN epitaxial templates on c-plane sapphire substrates, which were deposited according to reference.[8] Ohmic Ti/Au contacts (20 nm/100 nm) with different geometries and spacings had been deposited by electron-beam evaporation and a lift-off process on top of the κ-Ga2O3/insulating GaN epitaxial layers to allow in-plane transport measurements through the transfer length TLM method (linear TLM 200 µm wide, gaps: 70, 60, 50, 40, 30, and 20 µm—circular TLM inner contact diameter 80 µm, gap spacing: 5, 10, 15, 25, 50, 75, and 100 µm). A mesa structure with well-defined and isolated κ-Ga2O3 squared columns (200 × 200, 150 × 150, 100 × 100, and 70 × 70 µm²) had been obtained via dry etching processing the samples (inductively couple plasma – reactive ion etching ICP-RIE process details reported in reference,[10] 1.1 nm s⁻¹ etch rate for κ-Ga2O3 deposited on top of the conducting GaN epitaxial templates. Ohmic Ti/Au contacts had been deposited around the mesa (on the Si:GaN layer) as well as on top of the κ-Ga2O3 mesas (180 × 180, 130 × 130, 80 × 80, and 55 × 55 µm²) so to allow 2-point-probe out-of-plane resistance measurements to provide information on the possible structurally-driven conduction anisotropy (i.e., in- versus out-of-plane) in the material. For this purpose, two different κ-Ga2O3 thicknesses (1100 and 110 nm) were tested.

Supporting Information

Supporting Information is available from the Wiley Online Library or from the author.

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Conflict of Interest

The authors declare no conflict of interest.

Data Availability Statement

The data that support the findings of this study are available in the supplementary material of this article.

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gallium oxides, rotational domains, semiconducting oxides, structural and point defects, thin films epitaxy, transport anisotropy

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