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Etching of the SiGaN_x passivation layer for full emissive lateral facet coverage in InGaN/GaN core-shell nanowires by MOVPE

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Abstract

To strongly enhance the vertical growth rate in MOVPE-grown GaN core-shell wires, large quantities of silane (SiH_4) need to be introduced during the growth of the wire core. This results in the formation of a SiGaN_x layer that acts effectively as a dielectric mask on the sidewalls of the GaN core, thereby promoting the vertical growth. While useful during core growth, its presence precludes the formation of homogeneous core-shell heterostructures, whose coverage and optical quality tend to be maximized at the top of the wires. In this paper, we propose three different strategies to remove this passivating layer once its initial role accomplished. They are based on either chemical, physical or thermal etching. Their effect in the optical quality of subsequent core-shell InGaN/GaN heterostructures, including single and multiple-quantum well heterostructures, has been analyzed. Overall, an *ex-situ* chemical etching of SiGaN_x by H_3PO_4 results in an enhanced emissive coverage and a stronger overall luminescence intensity from the active regions, while simultaneously removing deep-defect emission arising from the high growth temperature of the core.

Introduction

InGaN/GaN heterostructures are a material of choice for their applications in fields such as, but not limited to, lasers^{1,2}, lighting^{3,4}, and displays⁵. In this context, the growth of planar nitride heterostructures has been documented for more than 25 years^{6,7}. However, because of the polarity of GaN, the growth of heterostructures along the *c*-axis leads to the so-called Quantum-Confined Stark Effect (QCSE), in which the quantum well (QW) band structure is distorted by the internal

polarization fields, leading to a decrease in the wavefunction overlap between electrons and holes. The QCSE is responsible for the power dependence of the wavelength of the emitted light, for the reduction of the overall efficiency and for longer carrier lifetimes (which could be beneficial in the absence of omnipresent non-radiative defects)⁸. To overcome these problems, one way is to grow QWs on non-polar GaN planes. One possibility that does not require expensive non-polar GaN substrates is to grow nanowires (NWs) along the *c*-axis and directly use the non-polar $\{1\bar{1}00\}$ *m*-plane sidewalls to grow the QWs. This “core-shell” structure also presents the advantage of having fewer or no dislocations (thanks to dislocation bending^{9,10,11} and mask filtering^{11,12} in the case of selective area growth). The high structural quality of the core and the absence of QCSE allows the growth of an InGaN shell with a larger thickness, reducing thereby the efficiency droop for a given light emission intensity thanks to the need of a reduced current density^{13,14}. The last advantage of the nanowire geometry is that with proper aspect ratios and spacing, the emissive surface is larger than that of standard planar LEDs.

Two main growth techniques, i.e. pulsed growth mode and continuous-flow mode have been reported for the growth of GaN nanowires by Metal-Organic Vapor Phase Epitaxy (MOVPE) and the impact of the growth parameters have been analyzed^{15,16,17,18}. In the case of continuous flow mode, temperature and V/III ratio appear as key parameters to promote GaN growth along the *c* axis^{17,18}. However, when playing with these parameters, GaN nanowires height is usually limited to few microns (1-3 μm). In order to significantly increase the vertical growth rate and reach >10 μm height nanowires, large quantities of SiH_4 must be introduced into the reactor^{10,19,20,21,22,23}. The introduction of SiH_4 during MOVPE growth leads to the formation of a thin passivation layer of SiGaN_x on the lateral facets of the wires²⁴ as reported by Tessarek et al.²⁵. This layer acts as a dielectric mask and promotes the diffusion of the reacting species along the sidewalls up to the top

of the wire thus highly enhancing the vertical growth^{23,25}; the second effect is the enhanced doping of the GaN core, which is beneficial for subsequent core-shell LEDs fabrication²⁶. The presence of this SiGaN_x layer has since then been evidenced in several other reports^{22,27,28,29,30,31,32,33,34,35}. Unfortunately, this passivation layer impacts the structural and optical quality of the InGaN shell grown on it by degrading the sticking of species and the resulting morphology^{27,31}. To avoid this issue, often the core growth is finished by a “non-intentionally doped” (nid) GaN portion during which SiH₄ is no longer introduced in the reactor^{28,30,36}. The surface of the nid GaN is thus not covered by the SiGaN_x passivation layer. When the active InGaN shell region is subsequently grown, the shell covers only the top nid part of the wire, which limits the overall light emitting area^{25,30}. In addition, the active region grown in the top part can be strongly degraded by a defective ultra-thin SiGaN_x underlayer, due to the residual SiH₄ in the reactor chamber³⁰.

Another method to circumvent the negative effect of the SiGaN_x layer consists in burying it. One possibility is to use AlGaIn as an underlayer before the growth of the LEDs active region, given its ability to nucleate on SiGaN_x^{28,36}. However, while AlGaIn underlayers have been used previously for reducing leakage current^{37,38} and for burying defects and optimizing crystalline quality^{39,40}, only few reports show a successful increase of the InGaIn QWs intensity at the NW base⁴¹. Besides, AlGaIn can also lead to cracks in the shell,^{28,36} as reported in the case of N-polar GaIn NWs. Similarly, a GaIn layer grown at low temperature reduces the Ga atom mobility and therefore enhances similarly the nucleation on top of the SiGaN_x layer, thus burying it. Still this method does not allow full emissive coverage³⁰, and very few articles, one to our knowledge, show a luminescence along the entire length of the nanowires grown by continuous MOVPE⁴¹.

In this report, we propose another approach that consists in removing completely this SiGaN_x passivation layer by a chemical or physical treatment^{23,27,29}, exposing the whole non-polar facet of

the NW and hence enabling full emissive coverage of the wire by the subsequent InGaN shell. This paper investigates first the impact of the treatment on the InGaN shell: we show that the luminescence of a single QW can be enhanced depending on the chemical or physical treatment. Second, the effects of these treatments on the surface of the underlying GaN are studied through the growth of a sample with an InGaN underlayer and a sample with three QWs. This provides a first understanding of the effects of the tested treatments on the GaN material quality and how it translates to the quality of the shell heterostructure. Last, a study of the emission on the entire length of the nanowire was realized, showing a clear impact on the emissive coverage of the treatments.

Experimental Section

The NWs were grown using selective area growth (SAG) on a GaN template grown on *c*-plane sapphire substrate covered with a 35 nm thick SiO₂ mask deposited by PECVD. The pattern on the mask was fabricated using Displacement Talbot lithography and inductively coupled plasma etching⁴², resulting in openings with a diameter of 300 nm and a pitch of 1.5 μm. The SAG of NWs was performed in a MOVPE Thomas Swan showerhead reactor with 1790 μmol/min NH₃ and 110 μmol/min Trimethylgallium (TMGa), giving a low V/III ratio of 15 typical of NW growth. Silane was added during the whole core growth at a flux of 0.4 μmol/min. Other growth parameters were as follows: a setpoint temperature of 1175°C, a pressure of 100 Torr, a growth time of 1800 s and a carrier gas ratio of H₂/N₂ = 3. In total, six identical samples were obtained. One of them is kept as reference while the others are used for the following experiments.

Three kinds of treatments have been investigated to remove the SiGaN_x layer: wet etching, dry etching and thermal etching. For wet etching, 3 different chemical solutions have been tested. Diluted Buffered Oxide Etch (BOE), potassium hydroxide (KOH)⁴³ and phosphoric acid (H₃PO₄)⁴⁴ have been chosen as they were all reported to etch SiGaN_x. Moreover, KOH is able to etch GaN, especially N-polar *c*-planes and semi-polar planes^{45,46,47,48}. H₃PO₄ was also reported to etch semipolar planes in GaN NWS^{29,48,49,50}. More precisely Deeb et al.²⁹ showed by a surface photovoltage measurements that H₃PO₄ treatment removes the surface states and reveals bulk-like properties of the underlying GaN. The exact protocols were as follows: The BOE was diluted 80 times, and the sample was treated for 10 seconds at room temperature. The KOH etching was performed with 45% KOH at 75°C for 5 minutes. Last, the H₃PO₄ etching was realized with 85% H₃PO₄ at 125°C for 10 minutes. Regarding the dry etching, one sample was treated using reactive ion etching in an Oxford system 100 ECR-RIE apparatus with SF₆ at 200W for 20s. Last, a method avoiding taking the samples out of the growth reactor was tested, and consisted in using an *in-situ* annealing under H₂ only at 1175°C for 15 seconds as a method of thermal etching.

After the treatments, the samples were reintroduced in the MOVPE reactor for the shell growth. Three different shell structures were grown and are represented on figure 1. For each sample, the regrowth started by growing a spacer of GaN at 1175°C during 100s with NH₃ and TMGa fluxes being respectively 0.13mol.min⁻¹ and 55μmol.min⁻¹. A first structure containing one single InGaN quantum well (SQW) was grown on all the samples at 850°C and 100 Torr during 360s. The NH₃, TMGa and Trimethylindium (TMIn) fluxes were 0.13mol.min⁻¹, 58μmol.min⁻¹ and 32μmol.min⁻¹, respectively, with N₂ as the carrier gas. QWs were subsequently capped using a layer of GaN grown at 1060°C, under the same fluxes for NH₃ and TMGa, and without any TMIn introduction. The second series contained a shell structure made of an InGaN underlayer (UL) followed by a

single QW. The underlayer was grown at 880°C and 100 Torr for 600s. The precursor fluxes were the same as for the QWs except for TMIIn, which had a reduced flux of 18 $\mu\text{mol}\cdot\text{min}^{-1}$. A third series of shell structures containing three QWs (MQWs) was also fabricated. The QW growth conditions were the same for all structures. A summary of all samples highlighting the different treatments and the regrown shells is given on table 1. For simplicity, the samples will be referred by the nature of the treatment to which they were exposed (reference, SF₆, H₃PO₄, KOH, BOE, H₂ annealing), as well as the shell structure used (SQW, UL+SQW, MQW). Last, the emission properties of the samples were measured at room temperature using a scanning electron microscope (SEM) equipped with a cathodoluminescence (CL) setup, operated at 5kV and with a constant probe current of 5.4nA.

Results and discussion

Single Quantum Well (SQW). SEM images of the NWs have been taken systematically before and after the shell deposition in order to monitor the effects of the treatment and assess the shell quality, as depicted in figure 2. The morphology of the NWs in the SF₆ and BOE treated sample (figure 2b and 2e) is similar to the reference sample (figure 2a), which indicates that any effect of the treatments should be localized close to the top surface. In the case of both H₃PO₄ (figure 2c) and KOH (figure 2d) treatments, a smoothing of the edges between neighboring non-polar facets as well as between the lateral non-polar and the top semi-polar facets is observed. Further, in the case of H₃PO₄ the top part of the sidewalls at the edges between *m*-plane facets seems to have been etched (which is more visible for a longer time of etching, as seen in **Supporting information S1**). Last, in the case of the H₂ annealing (figure 2f), the top surface has been thermally etched

away while the lateral non-polar facets seem unaffected, suggesting that in these conditions GaN is preferentially etched compared to the SiGaN_x passivation layer. Images at much larger magnification of a layer around the GaN core (**Supporting information S2**) suggest a SiGaN_x thickness of around 25nm, which is much thicker than the SiGaN_x layer reported by Tessarek *et al.*²⁵. This could mean that this layer might contain some GaN that stayed unetched by the treatment.

Figures 2(g) to (l) show the morphology of the NWs after the subsequent growth of the SQW shell. The reference sample (figure 2g), i.e. without SiGaN_x removal, shows a high surface roughness on both the lateral *m*-planes, due to the direct nucleation on the SiGaN_x layer, and the topmost semipolar planes, which display an inhomogeneous and uncomplete coverage. On the other hand, the SF₆ SQW sample (figure 2h) displays relatively smooth *m*-plane facets, with a minor growth disruption on the top *c*-plane of the truncated pyramid. This suggests that the surface of the core has been modified by the etching treatment. A similar result can be seen with the H₃PO₄ SQW sample (figure 2i), with a smooth and continuous *m*-plane, suggesting an extended emissive coverage. The NWs on the KOH SQW (figure 2j) sample have a “match-head” shape and some visible roughness at the edges between the *m*-plane facets, just below the “match-head”. This is consistent with the smoothing of the upper part of the wires after KOH treatment, suggesting that the QW containing shell has only been deposited on the top part of the wire. The BOE sample (figure 2k) shows similar rough features at the edges between the *m*-plane facets but with a covering of the shell being more homogeneous in thickness along the wire. Finally, the H₂ SQW samples (figure 2l) shows a high roughness on the *m*-plane facets, similar to the reference sample, and a clear overgrowth of a pyramid on the top part of the wire. This confirms the assumption that

only the upper part of the NWs consisting of the semi-polar plane facets was etched away by the H₂ treatment, while the SiGaN_x layer survives the H₂ treatment.

Figure 3a shows cathodoluminescence (CL) measurements made at room temperature on a single representative NW for each sample. These spectra have been obtained by integrating the emission spectra of hyperspectral images along the entire NW length. Four emission bands can be identified. The first one, around 364nm, is attributed to the GaN band edge. The red shifted emission compared to bulk GaN has already been reported in the case of VLS-grown Si-doped GaN NWs⁵¹. This phenomenon has already been studied in Si-doped planar polar *c*-plane⁵² and non-polar *a*-plane⁵³ GaN and could be explained by both bandgap narrowing and tensile strain introduced by the Si atoms⁵⁴. The second is a large band centered at around 550nm, identified as the yellow band^{55,56,57}. Finally, two other emission bands centered at around 400nm and 450nm can be identified and attributed to the InGaN QW luminescence. The 400nm band is generally identified as the *m*-plane luminescence from the QW, while the 450nm band is caused either by the red-shifted luminescence of *r*-planes QWs, or by In-rich clusters at the junctions between *m*-planes, which will be further discussed later in this document.

The GaN band edge signal is visible on all the samples and in particular for the reference sample and H₂ SQW sample. However, the luminescence peak at 400nm is observed only on the H₃PO₄, SF₆ and KOH SQW samples. The signal at 450nm is visible on all samples except for the BOE sample, whose general luminescence intensity is very weak. On the contrary, a strong luminescence is observed on the H₂ SQW sample but it is largely due to the emission of the yellow band. Finally, the H₃PO₄ sample shows the strongest luminescence at 400nm and a much weaker YB emission as compared to all other samples.

CL maps at 400nm and 450nm (figure 3b) allow to visualize the spatial distribution of these emission bands along the whole NWs length and, thus, evaluate the impact of the treatment on the shell's light emission. Some small disparities in height can be observed in SEM images and CL maps, which may be due to size inhomogeneity but more likely to the manual method used to remove the nanowires from their host substrate which does not guarantee to cut them precisely at their base. It should be noted that the intensity of the CL maps is arbitrary, and one should refer to figure 3a for a quantitative intensity comparison between the samples. In particular, monochromatic hyperspectral maps at 400nm allow to illustrate the efficiency of the employed treatment to grow a radiative SQW on *m*-plane sidewall facets. Note that a similar reasoning could be applied to the monochromatic images at 450nm, though the conclusions could be misleading due to the strong contribution of the yellow band, even at this wavelength, for some of the samples.

One should note first that for the reference, BOE and H₂ SQW samples, the intensity at 400nm is very low compared to all other three samples (figure 3a,b) and, thus, they can be excluded from the following analysis. Interestingly, while the signal at 400nm for the other samples is comparatively strong, its spatial distribution is very different: SF₆ and H₃PO₄ samples clearly display a SQW emission extending over the topmost half of the NWs, while in the KOH treated NWs the SQW emission comes preferentially from the top 0.75μm of the nonpolar planes, which is consistent with the SEM observations (fig 1j).

Moreover, the CL maps in figure 3b show different origins for the emission at 450nm. Indeed, reference, SF₆ and H₂ SQW samples shows a 450nm emission coming from the top pyramidal part of the wire with no signature detected on the non-polar sidewalls. Semi-polar planes are known to incorporate more indium⁵⁸, which has already been observed in NWs. The higher intensity observed for the H₂ SQW sample could be explained by the larger surface of the pyramid

overgrown at the top of the wire (see figure 2(l)). For the three other samples, the emission at 450nm is more evenly distributed along the non-polar facets. A remarkable point concerns the H₃PO₄ sample: indeed, the 450nm signal is coming from the edges between *m*-plane facets and its spatial distribution is complementary to the emission at 400nm. One hypothesis is that the 450nm emission is due to a greater incorporation of indium on the edges, which has been previously detected and ascribed to a combination of strain relaxation occurring at the edges⁵⁹ along with the formation of nanometric *a*-plane facets^{60,61}. Another explanation could be that H₃PO₄ etches preferentially the *a*-planes over *m*-planes in GaN, which is supported by a recent study on etching barrier index calculations⁵⁰, as introduced in Ref 47 to approximately predict relative resistance to etching of each crystallographic plane. A similar conclusion could be made for the top of the KOH SQW sample, but not for most of its sidewall. Indeed, a consistent 450nm signal is visible on the whole sample, despite no 400nm emission. Several explanations could be at play here (inhomogeneous etching leading to inhomogeneous emission, impurities introduced by the KOH, etc...), but it is difficult to conclude on the precise origin of this behavior. Last, the BOE sample also seems to exhibit 450nm emission from the sidewalls, but this emission can be ascribed to the tail of the yellow luminescence band of the sample as suggested by the corresponding spectrum on figure 3a.

These measurements evidence the impact of the treatments on the shell luminescence efficiency and on its coverage: measurements on the reference sample clearly show that the regrowth of the InGaN shell on the SiGaN_x passivation layer results in an excessive amount of defects, which prevents any QW emission from *m*-plane sidewalls. Similarly, BOE etching or H₂ annealing do not give satisfactory results. On the other hand, a more homogeneous distribution of the shell is obtained by chemical treatment with SF₆, KOH and in particular H₃PO₄. This last chemical

treatment allows a strong enhancement of the CL emission while reducing the YB emission, which suggests a better structural quality and, potentially, less incorporation of deep defects.

Effect on heterostructures containing InGaN underlayer (UL) and multi quantum well (MQW) samples. It has been shown in Ref 62 that in the case of planar GaN grown at a temperature higher than 900°C some point defects appear in the GaN and segregates at the surface. According to the authors, this defect could be a nitrogen vacancy (v_N) that would form non-radiative centers in quantum wells by reacting with In. One way to solve the problem is to grow a so-called “underlayer” (UL) with a small In composition below the QWs in order to bury these defects⁶³. This result is further supported by the fact that it is possible to observe the defects accumulating in the first quantum well in samples with MQWs⁶⁴. However, most reports on this subject focus on *c*-plane planar GaN and similar studies on core-shell structures are scarce. Among them, Kapoor et al.³⁰ observed that an UL with indium seemed to have no effect on the efficiency of the QW. In this section different core-shell structures have been studied to evidence the effect of etching treatment on the potential point defects associated to a defective GaN surface layer.

Both SQW+UL and MQW structures were grown on reference, SF₆ and H₃PO₄ samples. The choice of these two last samples among others described in the previous section, along with the reference sample, is motivated by the fact that they are the most luminescent samples. In addition, these two chemicals are known to attack the materials differently, because the SF₆ is expected to etch only the SiGaN_x passivation layer, while H₃PO₄ is expected to also etch the GaN core. The UL was chosen with a low-In content InGaN in order to compare the results with the literature³⁰. The spatially-integrated spectra are given in Figure 4 and a CL-mapping of a representative NW

of each sample is also provided in Figure 5. The CL spectra and maps presented for the SQW samples in both figures are the same as the ones presented in figure 3 and were included for comparison.

In the case of the reference sample, an increase of one order of magnitude of the QW peak is observed with the addition of an UL. On the other hand, the intensity of the MQWs sample is comparable to the one of the UL sample, but redshifted by 50 nm. The CL-mapping shows that the UL sample is the only one that exhibits *m*-plane luminescence.

For the SF₆ treatment, the 400nm peak associated with the *m*-plane QW emission is noticeable on every sample, and along a significant part of the wire, which is a signature of the SiGaN_x passivation layer. Here again, the addition of an UL results in an increase of one order of magnitude of the luminescence intensity. A threefold increase can be noticed with the use of several QWs.

Last, the samples treated with H₃PO₄ behaves differently. First, on the SQW spectrum, the dominant emission is the 450 nm band and, as shown on figure 4, it comes from the entire length of the NW. Second, the UL addition does not increase the 400 nm *m*-plane emission, however, a suppression of the emission at 450nm can be noticed. In the MQW sample, the intensity of the 400 nm peak is increased by one order of magnitude.

The results will now be discussed more in depth, and an explanation to the observed phenomena will be provided. Regarding the reference sample, a comparable luminescence increase is observed with respect to the SQW case both on the UL and on the MQWs samples. One possible explanation is that this improvement is due to the burial of non-radiative centers; in the first case (MQWs) it would be achieved by the addition of QWs, the first QWs being used to bury the non-radiative centers without providing further intensity, while in the second case it would be achieved by the

addition of an UL which traps defects, but also increases the emissive coverage on the m -plane of the NW, resulting in an intensity similar to the one in the MQW sample. Lastly, no QW emission can be seen at wavelengths shorter than 450 nm, whereas most other samples have an m -plane QW signal at 400 nm. The SQW+UL sample also exhibits some interesting m -plane luminescence, which is further discussed in **Supporting Information S3**.

For the SF₆ sample, the same mechanism is at play, and confirms that there is a defect responsible for non-radiative centers in the QW that can be suppressed by the addition of an underlayer. However, as said before, the luminescence of the MQWs sample here is greater than the one with the underlayer, which is expected since one can roughly consider that the first QW buries or traps the defects, and the second and third one contributes effectively to the overall emission, thus resulting in a total intensity larger than that of just one QW. This contrasts with the reference sample, where the first two QWs seem to be needed to bury the non-radiative centers⁶⁴, which is consistent with a higher defect density (see Figure 2g). It is important to note that the obtained results seem contradictory with the ones obtained in Ref. 30, in which the combination of an UL with a GaN spacer appears to have no effect on the luminescence intensity. A possible explanation is that the residual SiGaN_x ultra-thin layer on the wire sidewalls reported in the Ref. 30 can trap the point defects and prevent any diffusion into the shell. The subsequent spacer growth is realized at a temperature of 900°C, which is the low temperature threshold for point defect formation⁶². There are therefore few or no defects trapped in the UL. In the present case, since the SiGaN_x layer has been etched, the point defects are free to diffuse in the shell, even at low shell growth temperature. An underlayer may therefore be required to achieve a satisfactory luminescence in NWs where the core passivation layer has been removed.

The behavior is different for the H_3PO_4 samples. First, the addition of an UL results in a disappearance of the 450nm emission. This could be explained by the fact that the growth rate of the a -planes responsible for the emission at this wavelength in the H_3PO_4 SQW is faster than that of m -planes so that the a -planes disappear during the UL growth⁶⁵, resulting in the 400nm GaN m -plane emission being the only emission measured in the SQW+UL sample. A weak 450nm shoulder can be noticed in the spectrum of the MQW sample and is visible on the fig 4c which could be due to an emission from the first QW. Second, while the UL is expected to trap non-radiative centers and thus to increase the luminescence intensity, as seen in the two other samples, this is not the case here. One possible explanation for this is the etching of the GaN by the H_3PO_4 . According to Ref. 62, the defects responsible for non-radiative centers are segregated at the surface, and their incorporation in bulk GaN is negligible. On the other hand, it has been observed in Ref. 29, that a short H_3PO_4 treatment on GaN nanocolumns made them exhibit bulk-like properties. An interpretation of the result presented in this study is that the H_3PO_4 has been able to etch the surface of the GaN, and particularly the surface defects responsible for non-radiative centers. The UL thus no longer plays its role as an NRC trap, which explains why no intensity increase can be noticed. On the other hand, the MQW sample has a more intense 400nm signal because all the QWs are free of NRC-inducing surface defects and contribute to the overall intensity. It can also be noted that, if the three MQW samples are compared, the intensity of SF_6 and the control sample are similar, while the H_3PO_4 is almost four times more intense.

Emissive coverage of the MQW samples after SF_6 and H_3PO_4 treatments. Finally, in order to attest the relevance of our approach for the realization of LED devices based on core-shell nanostructures, we extracted the CL spectra of SF_6 and H_3PO_4 MQW samples at the top; bottom

and middle of the NW (figure 6). It is noteworthy that the luminescence signal is still clearly visible until the NW base where the intensity is comparable to that of the yellow band. Note that the yellow band intensity remains constant all along the nanowire. This result is significantly better than the ones found in the literature^{30,31,40}, where the luminescence coming from the quantum well usually collapses near the middle of the nanowire. It demonstrates that our approach allows us to obtain a full shell luminescence on doped nanowire grown by continuous MOVPE. Both H₃PO₄ wet etching and SF₆ dry etching thus appear to be efficient and easy processes to completely etch the SiGaN_x layer.

Conclusion

This paper explores strategies to etch away the SiGaN_x layer covering the lateral facets of heavily Si-doped GaN NWs grown by MOVPE in order to achieve a full InGaN shell emissive coverage along the entire NW length and to improve its optical quality. Except for the BOE treatment, all samples showed an improved intensity of the *m*-plane QW luminescence as compared to the untreated fully doped NW. Among all the treatment performed, SF₆ dry etching and H₃PO₄ chemical etching appear to be the most promising ones in order to efficiently remove the SiGaN_x layer and thus, to extend the InGaN shell luminescence along the entire NW length. In addition, through the comparative study of different InGaN shell structures, we suggest that the H₃PO₄ treatment not only remove the SiGaN_x layer but also some GaN core surface defects. Finally, the study of the SF₆ and H₃PO₄ treated samples' luminescence along their whole length highlights a dramatic increase of the MQW's emissive coverage, confirming that fully covered core shell NWs

can be obtained through SiGaNx etching processes. This significant increase of the luminescence signal suggests that SiGaNx removal by etching is a very promising method for improving the efficiency of core-shell nanowire LEDs.

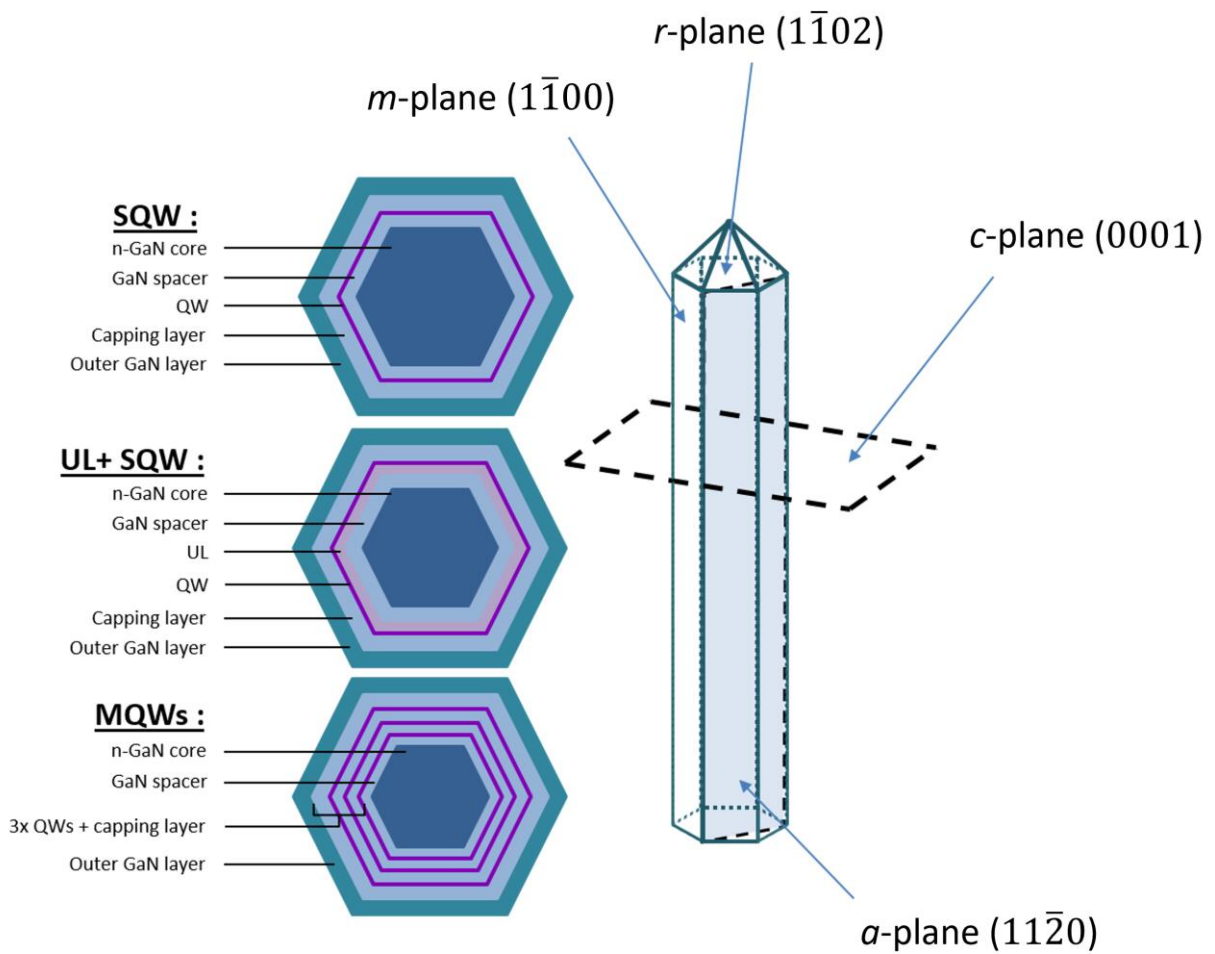


Figure 1. Structure of the three different shells and critical crystallographic planes in GaN nanowires

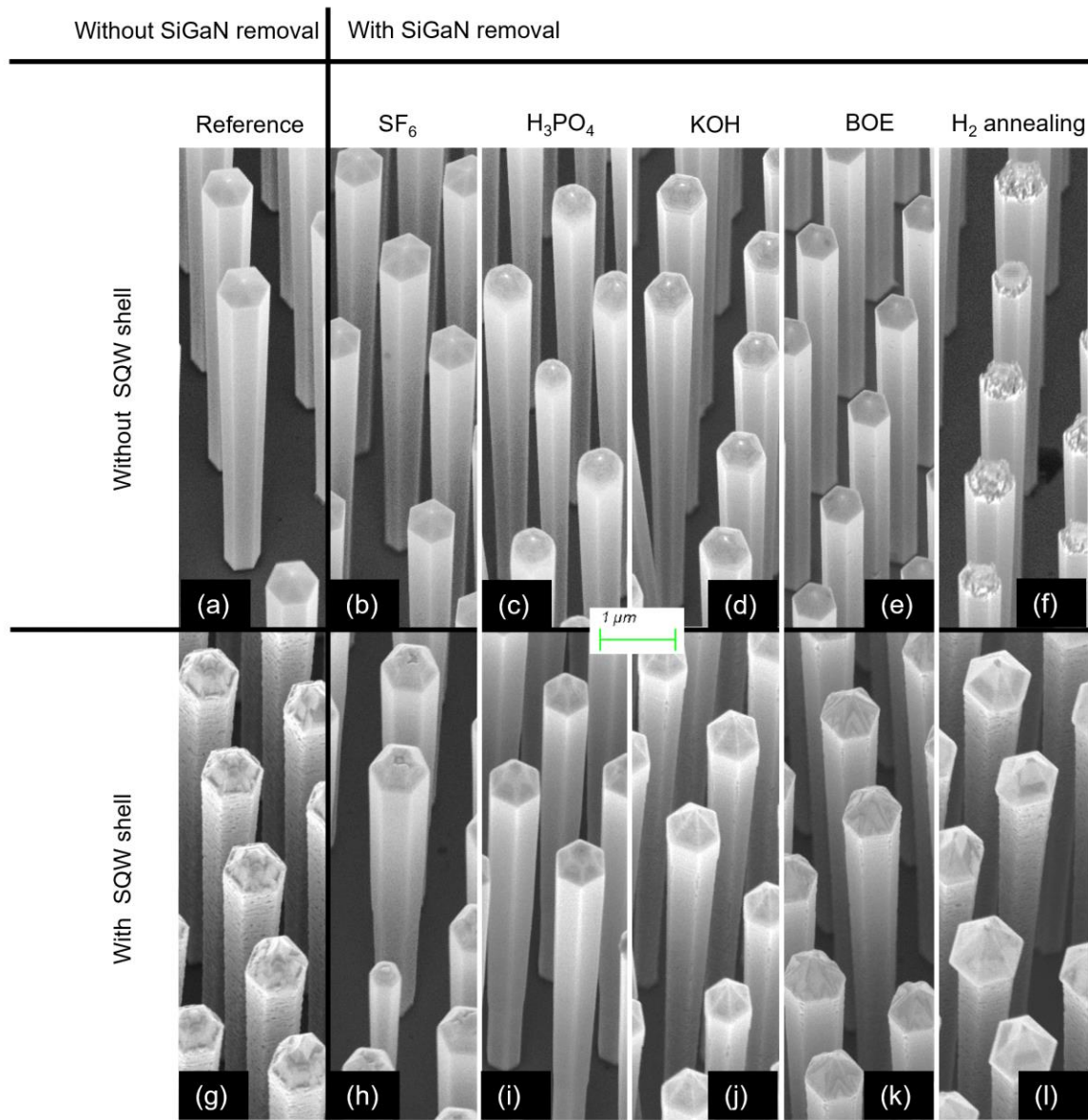


Figure 2. SEM images with 25° tilted view of: (a) to (f), doped core NW after surface treatment, and (g) to (l), the same samples after a subsequent shell growth

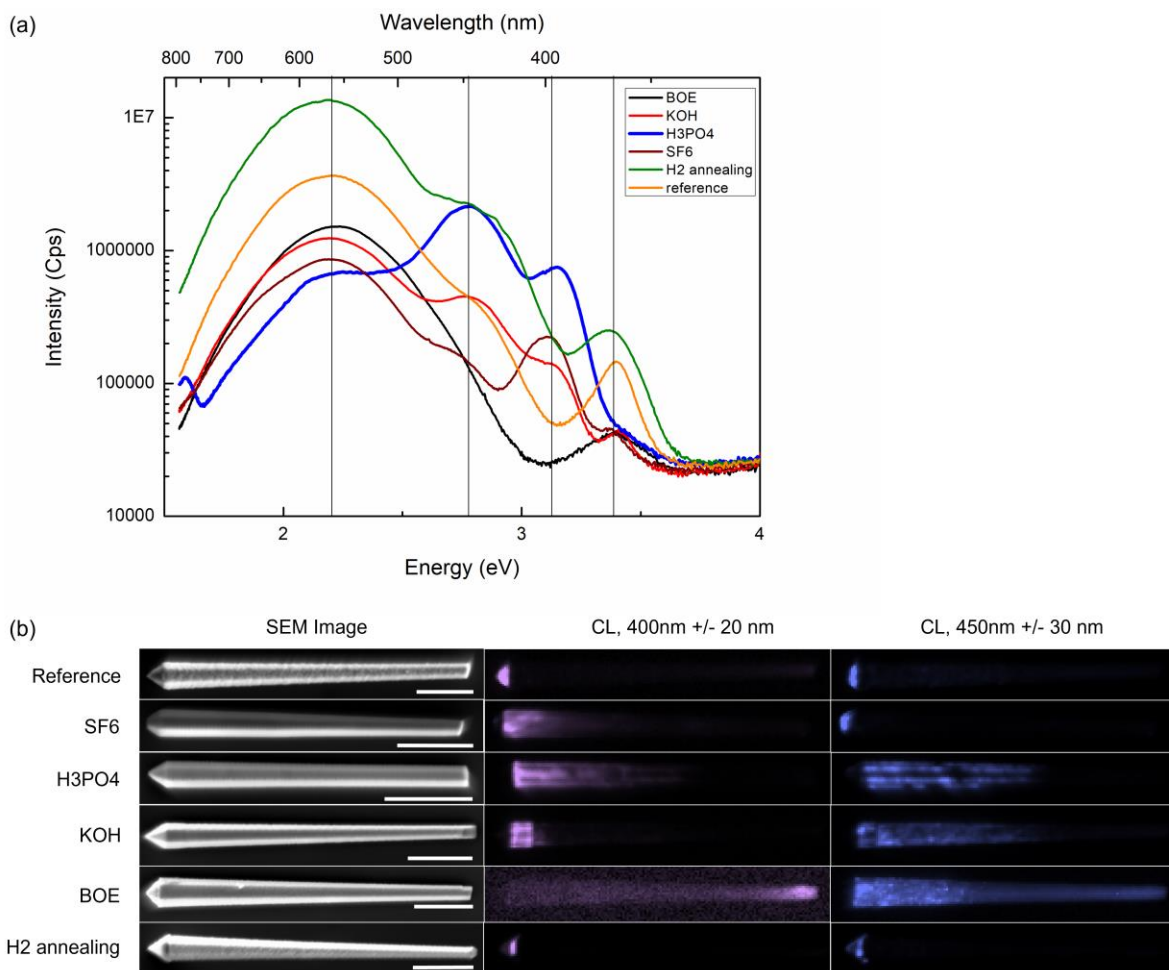


Figure 3. (a) CL spectra of SQW samples for treated and untreated samples. CL spectra were generated from the hyperspectral data set by adding the spectra of each pixel of the map, thus corresponding to the overall emission of the nanowire from top to bottom. The curve for H₃PO₄ has been highlighted as it is the best performing sample in this test. (b) SEM images of the NWs and corresponding normalized CL map of the luminescence intensity at 400 +/- 20 nm (c) and 450nm +/- 30nm (d). The scale bars are 2 μ m.

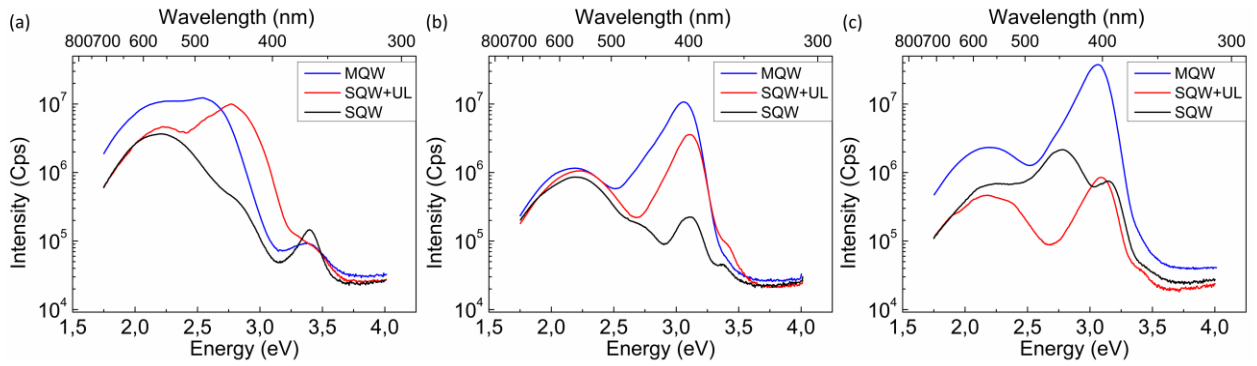


Figure 4. CL spectra for untreated (a), treated with SF_6 (b) and treated with H_3PO_4 (c) samples.

CL spectra were generated from the hyperspectral data set by adding the spectra of each pixel of the map, thus corresponding to the overall emission of the nanowire from top to bottom.

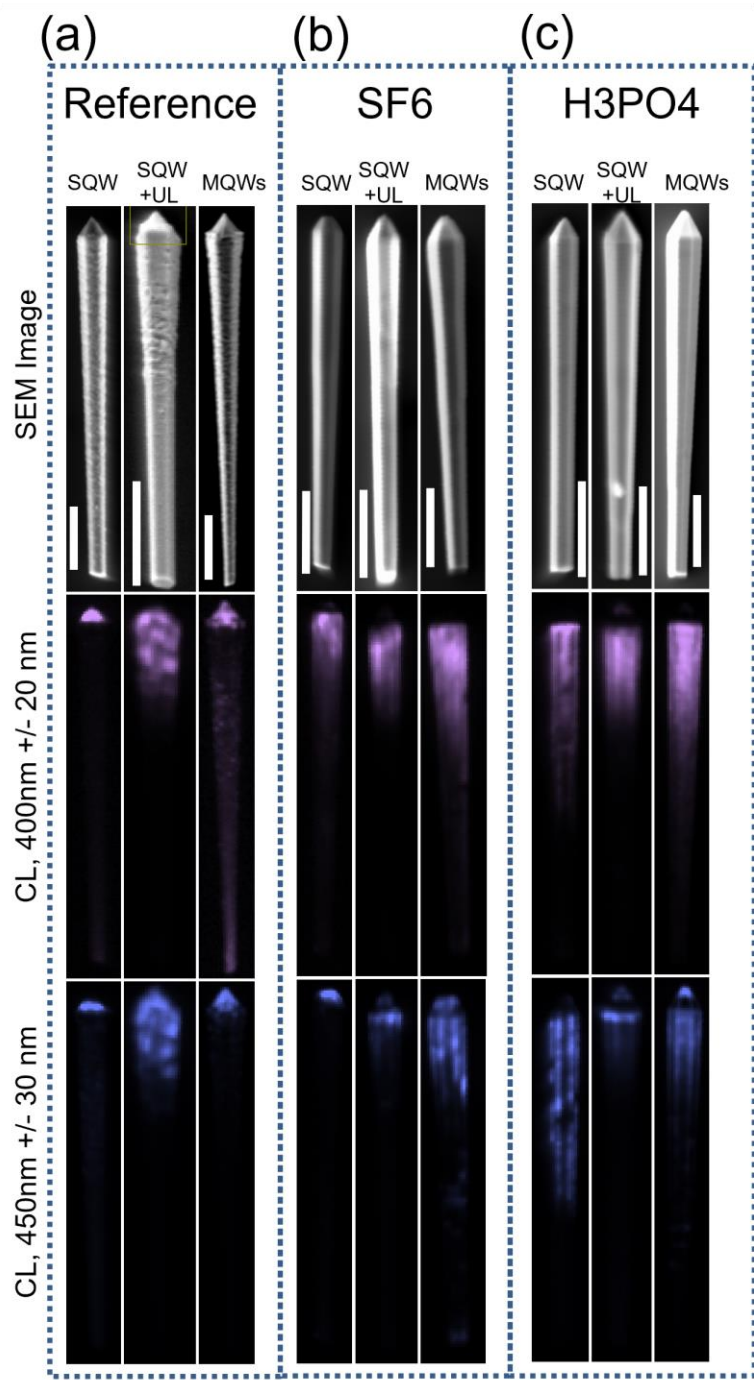


Figure 5. SEM image and CL maps at 400 and 450 nm for the reference sample (a), the SF₆ treated one (b) and the H₃PO₄ treated one (c). The scale bars are 2μm.

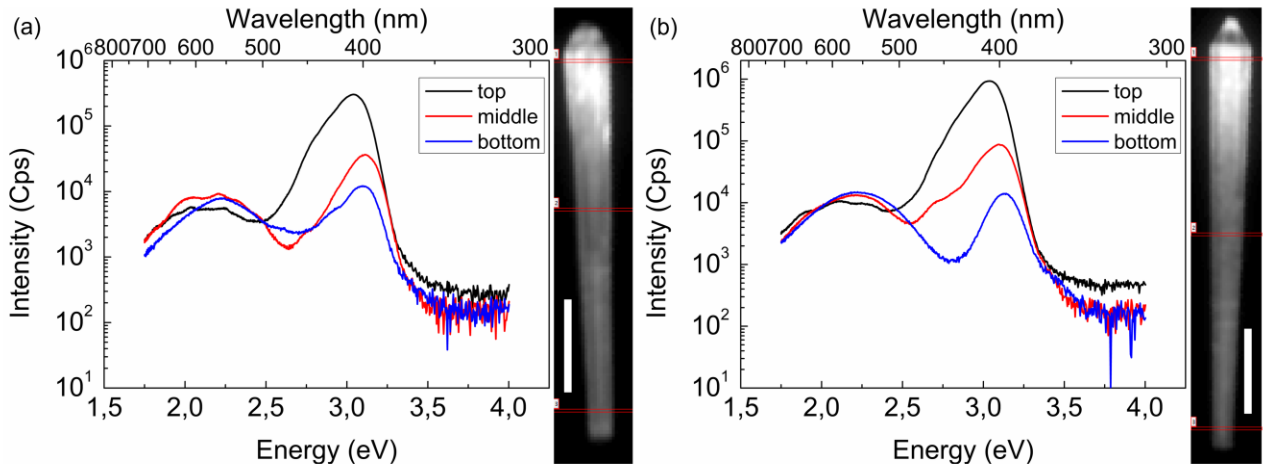


Figure 6. Room temperature CL spectra corresponding to the top, middle and bottom part of a SF₆ treated MQW sample (a) and a H₃PO₄ treated MQW sample (b). The scale bars are 2µm.

Type of shell		SQW	SQW+UL	MQW
Treatments	No treatment	Reference	Reference	Reference
	Chemical etching	BOE		
		KOH		
		H3PO4	H3PO4	H3PO4
	Dry etching	SF6	SF6	SF6
In situ thermal etching	H2 annealing			

Table 1. Summary of the sample used for each process

ASSOCIATED CONTENT

SEM image of doped GaN nanowire etched in H₃PO₄ at 125°C for 15 minutes, showing a faster etching at the junctions between *m*-planes, additional SEM image of doped GaN nanowires annealed under H₂ at 1175°C for 15s at x15 000 and x50 000 magnification, and supplementary panchromatic, SEM, and color reconstitution of the SQW+UL reference sample with CL spectrum for three indicated points.

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Author Contributions

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ABBREVIATIONS

LED, Light Emitting Diode; n-nid, doped/non intentionally doped; MOVPE, MetalOrganic Vapor Phase Epitaxy; QW, Quantum Well; QCSE, Quantum Confined Stark Effect; SAG,

Selective Area Growth; PECVD, Plasma Enhanced Vapor Phase Epitaxy; TMGa, Trimethylgallium; TMI_n, Trimethylindium; BOE, Buffered Oxide Etch; ECR-RIE, Electron Cyclotron Resonance Reactive Ion Etching; SQW, Single Quantum Well; SQW+UL, Single Quantum Well + Underlayer; MQWs, Multi Quantum Well; UL, Underlayer; SEM, Scanning Electronic Microscope; CL, Cathodoluminescence; YB, Yellow Band

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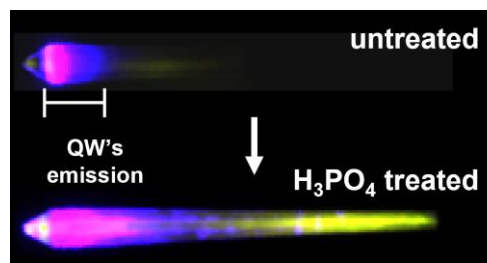
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Manuscript title: Etching of the SiGaN_x passivation layer for full emissive lateral facet coverage in InGaN/GaN core-shell nanowires by MOVPE

Authors: Julien Bosch, Pierre-Marie Coulon, Sébastien Chenot, Marc Portail, Christophe Durand, Maria Tchernycheva, Philip A Shields, Jesús Zúñiga-Pérez, Blandine Alloing



Cathodoluminescence mapping showing the evolution of the true color emission of an InGaN structure with no treatment between core and shell growth, and with an H₃PO₄ treatment