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## The 2020 UV emitter roadmap

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# The 2020 UV Emitter Roadmap

Hiroshi Amano <sup>1, 2</sup>, Ramón Collazo <sup>3</sup>, Carlo De Santi <sup>4</sup>, Sven Einfeldt <sup>5</sup>, Mitsuru Funato <sup>6</sup>, Johannes Glaab <sup>5</sup>, Sylvia Hagedorn <sup>5</sup>, Akira Hirano <sup>7</sup>, Hideki Hirayama <sup>8</sup>, Ryota Ishii <sup>6</sup>, Yukio Kashima <sup>8, 9</sup>, Yoichi Kawakami <sup>6</sup>, Ronny Kirste <sup>10</sup>, Michael Kneissl <sup>11, 5</sup>, Robert Martin <sup>12</sup>, Frank Mehnke <sup>13</sup>, Matteo Meneghini <sup>4</sup>, Abdallah Ougazzaden <sup>14</sup>, Peter J. Parbrook <sup>15</sup>, Siddharth Rajan <sup>16, 17</sup>, Pramod Reddy <sup>18</sup>, Friedhard Römer <sup>19</sup>, Jan Ruschel <sup>5</sup>, Biplab Sarkar <sup>20, 3</sup>, Ferdinand Scholz <sup>21</sup>, Leo J. Schowalter <sup>22, 23</sup>, Philip Shields <sup>24</sup>, Zlatko Sitar <sup>10</sup>, Luca Sulmoni <sup>11</sup>, Tao Wang <sup>25</sup>, Tim Wernicke <sup>11</sup>, Markus Weyers <sup>5</sup>, Bernd Witzigmann <sup>19</sup>, Yuh-Renn Wu <sup>26</sup>, Thomas Wunderer <sup>27</sup>, Yuewei Zhang <sup>16</sup>

1 IMaSS, Nagoya Univ, Nagoya 464-8603, Japan

2 Nagoya Univ., Nagoya 464-8601, Japan

3 Department of Materials Science and Engineering, North Carolina State University, Raleigh, North Carolina 27695, USA

4 Department of Information Engineering, University of Padova, via Gradenigo 6/b, 35131, Padova, Italy

5 Ferdinand-Braun-Institut, Leibniz-Institut für Höchstfrequenztechnik, Gustav-Kirchhoff-Str. 4, 12489, Berlin, Germany

6 Department of Electronic Science and Engineering, Kyoto University, Katsura Campus, Nishikyo-ku, Kyoto 615-8510, Japan

7 UV Craftory Co., Ltd., Nagoya 464-0015, Japan

8 RIKEN, 2-1 Hirosawa Wako, Saitama, 351-0198 Japan

9 Marubun Corporation, 8-1 Oodenma-cho, Nihonbashi, Chuo Ward, Tokyo, 109-8577 Japan

10 Department of Materials Science and Engineering, North Carolina State University, Raleigh, NC 27695-7919, USA

11 Institut für Festkörperphysik, Technische Universität Berlin, Berlin, D-10623, Germany

12 Department of Physics, SUPA, University of Strathclyde, Glasgow, G4 0NG, UK

13 Georgia Institute of Technology, School of Electrical and Computer Engineering, 777 Atlantic Drive, Atlanta, GA 30332, USA

14 Georgia Institute of Technology, School of Electrical and Computer Engineering, Georgia Tech-CNRS, UMI 2958, 57070 Metz, France

15 Tyndall National Institute and School of Engineering, University College Cork, Cork, Ireland.

16 Department of Electrical and Computer Engineering, The Ohio State University, Columbus, Ohio, 43210, USA

17 Department of Materials Science and Engineering, The Ohio State University, Columbus, Ohio, 43210, USA

18 Adroit Materials, Inc. 2054 Kildaire Farm Rd, Suite 205, Cary, North Carolina 27518, USA 19 University of Kassel, Dept. of Electrical Engineering/Computer Science and CINSaT, Wilhelmshoeher Allee 71, D-34121 Kassel, Germany

20 Department of Electronics and Communication Engineering, Indian Institute of Technology Roorkee, Uttarakhand, India 247667

21 Institute of Functional Nanosystems, Ulm University, 89069 Ulm, Germany

22 Crystal IS Inc., 70 Cohoes Ave., Green Island, NY 12183 USA

23 Asahi Kasei Corporation, Fuji 416-8501, Japan

24 Department Electrical & Electronic Engineering, University of Bath, BAth, BA2 7AY, UK

25 Department of Electronic and Electrical Engineering, University of Sheffield, Sheffield, S1 3JD, UK

26 National Taiwan University, Institute of Photonics and Optoelectronics and Department of Electrical Engineering

27 Electronics Materials & Devices Laboratory, PARC, a Xerox company, 3333 Coyote Hill Road, Palo Alto, CA 94304

#### Abstract

Solid state UV emitters have many advantages over conventional UV sources. The (Al,In,Ga)N material system is best suited to produce LEDs and laser diodes from 400 nm down to 210 nm – due to its large and tuneable direct band gap, n- and p-doping capability up to the largest bandgap material AlN and a growth and fabrication technology compatible with the current visible InGaN-based LED production. However AlGaN based UV-emitters still suffer from numerous challenges compared to their visible counterparts that become most obvious by consideration of their light output power, operation voltage and long term stability. Most of these challenges are related to the large bandgap of the materials. However, the development since the first realization of UV electroluminescence in 1970s shows that an improvement in understanding and technology allows the performance of UV emitters to be pushed far beyond the current state. One example is the very recent realization of edge emitting laser diodes emitting in the UVC at 271.8 nm and in the UVB spectral range at 298 nm. This roadmap summarizes the current state of the art for the most important aspects of UV emitters, their challenges and provides an outlook of future developments.

#### List of terms and acronyms

AFM – atomic force microscopy – defined in sect. 3

CL - cathodoluminescence - def. in sect. 4

C-DLTS and O-DLTS – capacitance- / optical- deep level transient spectroscopy - def. in sect. 16

- DBR distributed Bragg reflector defined in sect. 7
- DFB Distributed Feedback defined in sect. 19
- DFT Density functional theory defined in sect. 5
- DUV Deep ultra violet defined in sect. 4
- EQE External quantum efficiency defined in sect. 1
- EBL electron blocking layer def. in sect. 10
- ELO epitaxial lateral overgrowth defined in sect. 2
- $\eta_{rad}$  radiative recombination efficiency defined in sect. 1
- $\eta_{ini}$  current injection efficiency defined in sect. 1
- $\eta_{\text{el}}$  electrical efficiency defined in sect. 1

FC – flip chip – def. in sect. 13

FDTD – finite-difference time-domain – def. in sect. 13

- LEE  $\eta_{ext}$  light extraction efficiency defined in sect. 1
- IQE  $\eta_{IQE}$  internal quantum efficiency defined in sect. 1
- HR-PhC highly reflective photonic crystal defined in sect. 13
- HTA high temperature annealing (of AIN) defined in sect. 2
- HVPE Hydride vapour phase epitaxy defined in sect. 3
- $I_f$  forward current defined in sect. 13
- LED light emitting diode defined in sect. 1
- MBE Molecular beam epitaxy defined in sect. 7
- MOVPE Metal-organic vapour phase epitaxy, same method as MOCVD defined in sect. 2
- MQW multi quantum well def. in sect. 12
- NRC nonradiative recombination centers def. in sect. 4
- PL Photoluminescence def. in sect. 4
- pss patterned sapphire substrate defined in sect. 2
- PVT physical vapour transport (growth of AIN) defined in sect. 3
- PDD point defect density defined in sect. 11
- QW quantum well defined in sect. 4
- RMS roughness root mean square roughness defined in sect. 3
- SIMS secondary ion mass spectrometry defined in sect. 5
- SNOM scanning near-field optical microscopy def. in sect. 4
- TDD threading dislocation density defined in sect. 2
- TEM transmission electron microscopy defined in sect 7
- TJ tunnel junction defined in section 7
- TE transverse electric defined in sect. 12
- TM transverse magnetic defined in sect. 12
- UV ultra violet
- UVA 315 nm 400 nm, according to ISO 21348

UVB - 280 nm - 315 nm, according to ISO 21348

- UVC 100 nm 280 nm, according to ISO 21348
- VCSEL Vertical cavity surface emitting laser def. in sect. 7
- V<sub>f</sub> forward Voltage def. in sect. 13
- Wall plug efficiency (WPE) defined in sect. 1
- XRD X-ray diffraction- defined in section 7

#### 1 – UV-LEDs: State-of-the-Art and Applications

Michael Kneissl<sup>1, 2</sup>

1 Institute of Solid State Physics, Technische Universität Berlin

2 Ferdinand-Braun-Institut, Leibniz Institut für Höchstfrequenztechnik

#### Status

Driven by the wide range of applications the development of group III-nitride based ultraviolet emitters has gained considerable momentum since the early 2000's [1,2]. In the UVA spectral band (400 nm - 315 nm) key applications include UV curing of polymers, inks, coatings, resins, and adhesives [3]. UVA emitters are also employed for sensing applications, e.g. the detection of blankophores and fluorescent labels as well as blood gas analysis and light therapy. Phototherapy, in particular the treatment of skin diseases like vitiligo and psoriasis, is one of the key applications for LEDs in the UVB wavelength range (315 nm - 280 nm). UVB-LEDs are also of great interest for the curing of surfaces since the penetration depth of UVB light in polymers is much smaller than that of UVA emitters. In addition, plant growth lighting constitutes a large scale application for UVB-LEDs. Here UVB-LEDs are employed to enhance the production of secondary metabolites in vegetables, which carry a number of health benefits for the consumer or to control the morphology and shapes of flowers [3]. Disinfection of drinking water and wastewater treatment are certainly the highest volume applications for LEDs emitting in the UVC spectral band (100 nm to 280 nm). In addition, sterilization of medical equipment, home appliances (e.g. refrigerators, washing machines), and various surfaces (e.g. in the food industry), would also benefit from UVC-LEDs. Sensing applications require specific emission wavelengths, depending on the absorption band of the gases or biomolecules of interest, and therefore cover the entire UV spectral range. Fig. 1 presents some of the key applications for UV-LEDs. As can be seen, the wavelength and power requirements vary greatly for these applications. Whereas sensing applications entail low power but spectrally pure LEDs, for UV curing and disinfection LED modules delivering many Watts of UV light are crucial.

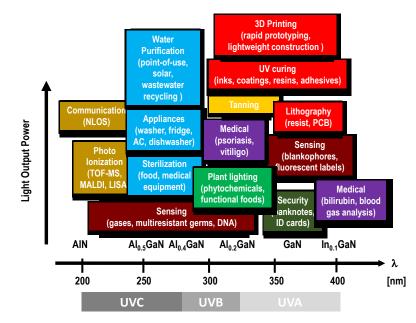


Figure 1. Key applications for UVA, UVB, and UVC LEDs. Figure reproduced from Ref. 3 with permission of Springer Nature.

Since UVA-LEDs with emission wavelength above 315 nm have already reached excellent performance levels with output powers of several watts per LED chip, UV curing has evolved as the first large scale application for UV-LEDs. Compared to conventional UV sources, like low and medium pressure mercury lamps, UV-LEDs provide a number of advantages. Besides a freely selectable single wavelength emission, UV-LEDs require no warm-up time, are electronically dimmable, exhibit no forward heat radiation, are highly temperature stable, and exhibit long lifetimes. Due to their small form factors, UV-LEDs provide a much greater design flexibility for UV modules and are also environmentally friendly, since they contain no mercury or produce ozone. Even though the output power levels of UVB- and UVC-LEDs are currently only in the 1 - 100 mW range, these shorter wavelength UV-LEDs have already found entry in first applications, especially in areas were solution with conventional UV sources are impossible or significantly more complex.

#### **Current and Future Challenges**

As can be seen from Fig. 2 the external quantum efficiencies (EQE) of UV-LEDs vary greatly with the emission wavelength. Whereas UVA-LEDs with emission longer than 315 nm exhibit EQEs exceeding 50%, LEDs in the UVB and UVC band lag significantly behind. Although record EQEs of more than 20% for LEDs emitting near 275 nm have been reported [4], the performance levels of commercial devices in these wavelength bands are still in the single digit percentage range.

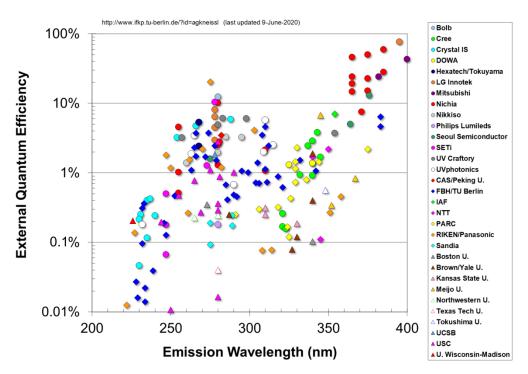


Figure 2. External quantum efficiencies of UV-LEDs in the spectral range between 200 nm and 400 nm. Figure adapted from Ref. 3 with permission of Springer Nature.

In order to analyse the shortcomings it is useful to take a look at the key parameters that describe the performance of UV-LEDs, i.e. the external quantum efficiency (EQE) and wall-plug efficiency (WPE). Both are related through the following equation

$$WPE = \frac{P_{out}}{I \cdot V} = EQE \frac{\hbar\omega}{e \cdot V} = EQE \cdot \eta_{el}$$

where  $P_{out}$ , I, and V represent the output power, operating current, and operating voltage of the UV-LED, respectively,  $\hbar\omega$  the emitted photon energy, and  $\eta_{el}$  the electrical efficiency [3]. The EQE in turn is dependent on the radiative recombination efficiency  $\eta_{rad}$ , the carrier injection efficiency  $\eta_{inj}$ , and the light extraction efficiency  $\eta_{ext}$ . Accordingly, the EQE can be described as

$$EQE = \eta_{rad} \cdot \eta_{inj} \cdot \eta_{ext} = \eta_{IQE} \cdot \eta_{ext}$$

where the product of the radiative recombination efficiency  $\eta_{rad}$  and the carrier injection efficiency  $\eta_{inj}$  is typically referred to as the internal quantum efficiency  $\eta_{IQE}$ . For example, record EQE and WPE for commercially available 280 nm LEDs are currently around 6.4% and 4.1% [5,6]. Although it is difficult to precisely determine the different contributions to the overall efficiencies, from the description of the material properties and device structure one would estimate the radiative efficiency to be around 40%, an extraction efficiency around 20%, and an electrical efficiency of 64%. Consequently, progress in all of the contributing factors will be required in order to improve the EQE, wallplug efficiency, and output power of UV-LEDs.

#### Advances in Science and Technology to Meet Challenges

There are a number of different possible solutions in order to improve each of the parameters that govern the overall UV-LED performance i.e.  $\eta_{rad}$ ,  $\eta_{inj}$ , and  $\eta_{ext}$ . For example, the extraction efficiency  $\eta_{ext}$  can be greatly enhanced by encapsulating the LED chips with a UV-transparent high refractive index material. A key challenge here is to find suitable materials, that are UV-transparent and also do not degrade under UV exposure. This is not a trivial task, but first candidates have emerged, e.g. fluoropolymers like CYTOP<sup>®</sup> [7]. Other approaches to enhance light extraction include the utilization of UV-reflective metal contacts together with a UV-transparent p-AlGaN heterostructure. However, finding suitable metal layers that are highly reflective in the UVB and UVC spectral range and at the same time enable low resistance ohmic contacts is not straightforward. Therefore, in many cases a trade-off between enhancing light extraction and degrading other parameters like operating voltages and electrical efficiency has to be made. Another key parameter is the radiative recombination efficiency  $\eta_{rad}$  which is strongly affected by the defect density in the semiconductor materials [8]. As most UV-LEDs are grown on sapphire substrates a large number of threading dislocations are generated at the AIN/sapphire interface. Even with advanced growth technologies in order to reduce the defect density, the threading dislocation density (TDD) in AlGaN heterostructures is typically between 10<sup>10</sup> cm<sup>-2</sup> and 10<sup>8</sup> cm<sup>-2</sup> [9,10], which corresponds to IQEs ranging from a few percent to over 60% [8]. Therefore, reducing the TDD is paramount to improving the radiative recombination efficiency and IQE of UV-LEDs. We have just provided these examples to illustrate the issues and refer to the discussion in the following articles describing in more detail the recent developments in materials and device technologies.

#### **Concluding Remarks**

Even though the large scale application of UV-LEDs will require substantial advances in the performance levels of UVB- and UVC-LEDs, it is clear that in the not too distant future group IIInitride based UV emitters will replace most conventional UV sources like mercury lamps. UV-LED and blue LED technologies share the basis of III-nitride material and device technologies. Although the technological solutions are very different there seem to be no fundamental roadblocks that would inhibit a steady improvement in UV-LED performance. How quickly these advances will be realized, will also depend on the magnitude of the research efforts in the field, but based on the trajectory of the performance improvements of the past decade one can anticipate that UVB and UVC LEDs with WPE of more than 10% should become commercially available by 2022 [2].

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#### 2 – AIN on sapphire

### Sylvia Hagedorn, Markus Weyers Ferdinand-Braun-Institut, Leibniz-Institut für Höchstfrequenztechnik

#### Status

UV LEDs are nearly exclusively bottom emitters, since p-GaN and contact materials are not transparent in the UV. Sapphire (or rather the pure corundum) is transparent and mass-produced for GaN LEDs. For UV transparency the semiconductor layer stacks have to start with AIN, which creates much bigger challenges in MOVPE (metalorganic vapour phase epitaxy) growth than GaN. The target is to obtain a low density of threading dislocations (TDD). Such dislocations form at the sapphire – AIN hetero-interface by coalescence of small nuclei and are non-radiative recombination pathways. A low TDD thus is prerequisite for high internal quantum efficiency [<sup>i</sup>]. While growth of thick layers is effective in reducing the TDD by mutual annihilation of dislocations, the high strain resulting from lattice mismatch and mismatch in the thermal expansion coefficients leads to strong wafer bow or even cracking of thicker layers. Strong bow makes processing of the wafers difficult, crack formation renders them useless for devices. Total internal reflection at the AIN-sapphire interface limits light extraction, which is especially low at short wavelengths. Patterned sapphire substrates (pss) are used for blue LEDs to enhance light extraction and to reduce TDD (or the layer thickness at which a low TDD is obtained). For UV LEDs this is still being developed. While UV LEDs have made tremendous progress in recent years, there is still plenty of room to improve their performance towards that achieved by blue LEDs which already approach the theoretical limits. The AIN-sapphire buffer layers are the basis for this further development. As yet, there is no completely satisfying solution that is proven by device performance and widely adopted.

#### **Current and Future Challenges**

Due to the high bond strength of AIN, high growth temperatures are preferred. On the other hand, high temperatures in an MOVPE reactor may result in surface reactions between the sapphire substrate and the gas phase. This does not only contain the injected process gases but also species stemming from parasitic deposits within the reactor. As one result of such pre-reactions, islands with N-polarity may form, which later are buried by the matrix with Al-polarity aided by the formation of AlON [<sup>ii</sup>]. So mastering the growth start is a challenge with strong impact on TDD. After coalescence of the initial nuclei, the TDD needs to be further reduced. A rough growth front with a high density of steps can help in this (by bending dislocations away from the +c surface normal). However, for the growth of the AlGaN device layers, high steps are detrimental since they can result in compositional modulation by a higher Ga incorporation rate at such steps ["]. To reproducibly manage the transition to a rough growth front and back to a smooth final AIN surface is a challenge. The pss approach for enhanced light extraction is hampered by the low surface mobility of Al. Thus the patterns (holes in the c-plane surface or pillars with c-plane top facet) need to be on the sub-micron scale (nano-pss). Also, AIN tends to nucleate in misoriented crystallites on non-c-plane sidewalls of such patterns. The c-oriented grains are difficult to coalesce if they are bounded by m-plane facets. So it remains a challenge to obtain smooth, fully coalesced AIN buffers on such nano-pss. Wafer bow results in an inhomogeneous temperature over the wafer making it a challenge to obtain homogeneous layer properties, especially when wafer size is being scaled up.

#### Advances in Science and Technology to Meet Challenges

There are a number of approaches towards reducing TDD. These include specific growth schemes for example by multistep growth with variation of the growth temperature, the ratio of TMAI and  $NH_3$ 

flow, the total pressure or by pulsed supply of the precursors. Although different growth recipes have been published as being successful, discussion with many groups active in the field indicates that reproducibility and homogeneity are issues that are being faced (but usually not discussed in scientific papers). A better understanding of the reasons for these limitations is required to optimize the production of AIN - sapphire templates.

A relatively new approach is the reduction of the TDD by annealing of thin AlN layers at high temperature (HTA) [<sup>iv</sup>]. This approach allows TDD in the range of  $5x10^8$  cm<sup>-2</sup> already for layer thicknesses of less than 1 µm reducing problems with wafer bow and cracks. It also works if sputtered layers of initially low quality (high TDD) are subjected to HTA (Fig. 1). In addition to being a relatively cheap process, sputtering also removes the issue of chemical attack on sapphire during heating up in an MOVPE environment. While a proof of principle has been given [<sup>v</sup>], further studies of process reproducibility and resulting device lifetime are necessary before this approach can be considered as validated.

To address the issue of light extraction, growth on nano-pss needs to be developed further [<sup>vi</sup>]. This includes finding the right patterns (which are different from those used for GaN not only in pitch but also in shape). Also the patterning of sapphire on this nanoscale needs to be optimized towards homogeneous and reproducible shape. The right growth parameters to obtain smooth and fully coalesced layers without misoriented crystallites need to be found. While the better understanding being developed in studies of planar growth provides guidelines, the presence of different orientations of sapphire and AlN will require modifications to the growth processes on nano-pss. While work currently is mainly on 2 inch sapphire substrates, higher volumes will require the

transition to larger diameters. To cope with wafer bow this will probably require thicker substrates which as a consequence may require another round of growth process optimization.

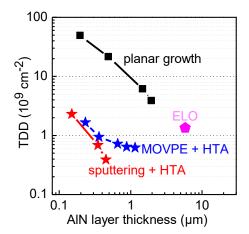


Figure 1. Reduction of threading dislocation density with grown AlN layer thickness. For planar growth, cracking limits the thickness that can be achieved. Epitaxial Lateral Overgrowth (ELO) patterning allows thicker layers and thus lower TDD but is not very economic [<sup>Vii</sup>]. High temperature annealing (HTA) achieves low TDD already for much thinner layers.

(S. Hagedorn et al., Ferdinand-Braun-Institut, unpublished results)

#### **Concluding Remarks**

While a low TDD and a smooth surface and not too big wafer bow are necessary requirements for AIN templates for UV LEDs, their suitability needs to be verified by growing device layer sequences, processing devices and studying their performance including reliability. Thus this topic will remain on the agenda for the foreseeable future. A combination of nano-patterning with defect reduction by HTA promises high crystalline perfection, acceptable wafer bow and improved light extraction and is worthwhile being studied more in detail [<sup>viii</sup>, <sup>ix</sup>].

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## 3 - The growth of bulk AIN and fabrication of AIN wafers

Ronny Kirste, Zlatko Sitar

North Carolina State University

#### Status

For most semiconductor technologies, like Si, Ge, III-arsenides, single crystalline substrates obtained from bulk growth provide for highest epitaxial layer quality and devices with optimum efficiency and lifetime. With the progress in hydride vapour phase epitaxy (HVPE) and ammonothermal growth of GaN crystals, these substrates are also entering InGaN and GaN device development. For Al-rich AlGaN epitaxial layers desired for UV laser diodes, LEDs, next generation power diodes, and sensors, single crystal AIN substrates are fabricated either via HVPE or sublimation growth/ physical vapor transport (PVT)[1-4]. In PVT growth, an AIN-powder source is sublimed at temperatures exceeding 2200°C in a nitrogen atmosphere. On the opposite side of the crucible, a seed is positioned and heated to temperatures slightly lower than the source. Due to the thermal gradient, the sublimed source molecules diffuse to the seed where they form AIN via a reverse reaction. The structural quality of the grown AIN crystals depends strongly on the type and quality of seed crystals, which are needed to provide structural information for the condensing Al and N atoms. Next to the control of thermal fields in PVT growth the use of native AIN seeds is essential to achieve high-quality crystalline boules and wafers. These seeds are typically obtained by spontaneous nucleation and growth of Lely-like, c-oriented platelets close to the equilibrium conditions. The first results of bulk growth of AIN on native seeds were presented in 2002; this process later became a basis for the development of an iterative boule expansion process and increase of the diameter of the native seeds over time.

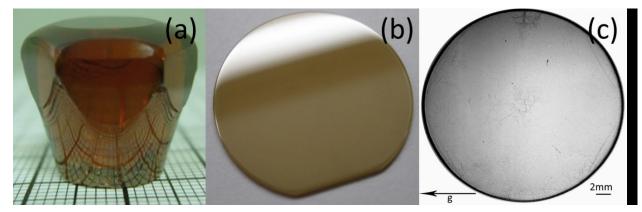


Figure 1: (a) Photograph of an AIN boule grown via the PVT method (mm-grid); (b) image of a 1" AIN wafer after mechanical and chemo-mechanical polishing; (c) x-ray topograph confirming that most of the wafer area is dislocation free.

Over the past two decades, PVT growth of AIN has been pursued by various laboratories around the world. [1, 2, 5, 6] Despite all these efforts, commercial availability of these substrates is still limited. Available PVT grown substrates typically have a dislocation density  $< 10^3$  cm<sup>-2</sup> and are available in diameters up to 2". Figure 1 shows a typical AIN boule (a) and a wafer cut from such a boule (b); an

X-ray topograph (c) from a high quality AIN wafer confirms the low dislocation density that can be achieved.

AFM imaging of polished AIN wafers reveals c/2 steps and an RMS roughness <0.1 nm. Using these AIN wafers, epitaxial growth via metalorganic vapour phase epitaxy (MOVPE) has been developed and doped and undoped AlGaN layers with up to 40% Ga-content can be grown pseudomorphically.[7, 8] Optically pumped UV lasers on AIN substrates have been demonstrated by various groups and the lowest threshold of 6 kW/cm<sup>2</sup> was achieved.[4, 9, 10]

#### **Challenges & Future Development**

While the crystal quality of PVT grown AIN substrates and epitaxial films grown on them is excellent, a few challenges need to be addresses in the near future before a widespread commercial adoption can be anticipated.

First, availability of 2" AlN wafers was limited up until recently. This hindered a possible entry even for fundamental epitaxy and device growth for many research facilities since many fabrication lines are designed for at least 2" wafers, with many foundries even moving further to 4". Although there is no fundamental limit for large-size AlN boules, the expansion process is relatively slow and will require significant long-term investment to achieve production of 4" or even larger boules.

Second, for UV light emitting diode (LED) applications, AIN substrates need to be transparent in the targeted emission window. PVT-grown AIN contains relatively high concentrations of C, Si, and O, which results in reduced thermal conductivity and various absorption centers that negatively impact the performance of any LEDs grown on them by absorbing the emitted UV light.[11] While the impact on UV laser diodes is not as significant, any leakage of the optical mode into the substrate will negatively impact the laser diode threshold. The most impactful absorption center for AIN is located at 4.7 eV or 265 nm, which is at the desired UVC LED emission wavelength for disinfection applications. Absorption coefficients exceeding 1000 cm<sup>-1</sup> are commonly observed at this wavelength. This absorption band has been associated with the luminescence emissions around 2.7 eV and 3.9 eV. It has recently been discussed that the absorption is related to the  $C_N^-$  point defect and the emission is related to the transition between the  $V_N$  and  $C_N$  point defects.[12] Thus, controlling these two point defects is key to reducing the absorption coefficient. Indeed, AIN epiready wafers have been deployed that have reasonable absorption coefficients <100 cm<sup>-1</sup>, which was achieved by compensation with Si (Figure 2 (a)).[11] While this approach significantly reduces the 265 nm absorption, the growth conditions to achieve the desired transparency may result in the formation of low angle grain boundaries in the PVT AIN crystals, which reduces the quality of the substrates up to the point that epitaxial layers show signs of relaxation and surface deterioration. Therefore, more efforts need to be made to reduce carbon and other point defects in PVT-grown AIN.[13] Alternatively, either the substrate could be removed during the LED fabrication or HVPE could be used to grow transparent AIN on PVT crystals to retain crystalline quality and obtain low point defect concentrations. Since the incorporation of C in HVPE and MOVPE can be controlled more efficiently, such substrates are excellent for UV LED and laser applications when the PVT seed is removed (Figure 2 (b)).

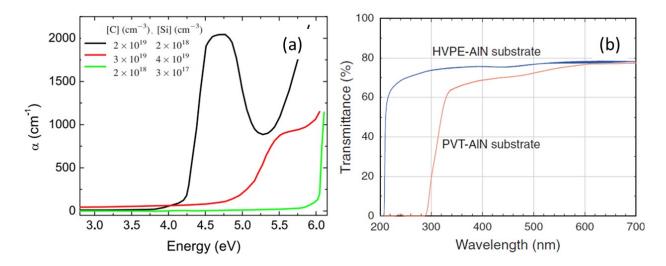


Figure 2: (a) 265 nm absorption can be reduced by orders of magnitude by controlling the carbon concentration in AIN; (b) using PVT substrates as seeds for HVPE and subsequent PVT seed removal results in transparency down to the fundamental bandedge.

Third, conductive AIN substrates would be beneficial for UV laser diodes as they would allow for vertical devices with more homogenous carrier injection, simplified laser design, and simpler fabrication. In addition, AIN and Al-rich AlGaN could potentially be used for high power Schottky diodes due to their high breakdown fields. AIN has a figure of merit that is nearly 200 times higher than that of SiC and 35 times higher than that of GaN. The main obstacle in using AIN substrates for power devices is their lack of conductivity – all AIN substrates are highly insulating. While the most promising n-dopant in AlGaN is Si, it forms DX-center and suffers from high activation energy and compensation for Al-content >85%.[14] Therefore, significant efforts are needed to better understand the effect of compensation and DX-center formation and increase doping efficiency.[7, 15, 16]

Fourth, the AIN surface and epitaxy of AIGaN films on AIN substrates needs to be addressed beyond simple trial-and-error approaches. This includes surface preparation, growth initiation, control of the surface morphology during growth, as well as engineering of possible relaxation schemes for thick AIGaN films. So far, optically pumped laser structures and fully fabricated laser diodes for electrical injection have been successfully grown on AIN substrates by a few groups.[4, 9] While significant efforts have been made to explain the epitaxial growth of AIGaN and the impact of various process parameters on the growth in general terms to allow for process transfer between different reactors, the reported results are not always optimal; in the worst case, epitaxy of AIGaN on AIN substrates.

Lastly, it should be mentioned that, despite high current prices, substrate cost is not a major challenge for future application of AIN substrates. The price per device will significantly drop for larger substrate areas (2", 4"...) and, even at the current price levels, the substrate contributes only 10-20% of the overall cost of a UV laser diode or LED. In addition, since the input cost of AIN crystal growth is not significantly different to that of SiC, the AIN substrate prices are expected to follow those of SiC in high volume production.

#### **Concluding remarks**

Device growth on native substrates is always more desirable than the growth on non-native substrates, since it allows for better crystal quality, resulting in more efficient and more robust devices. Although commercially available AlN substrates have improved greatly in quality and size over the years, a widespread penetration has not been achieved. This is mostly due to the challenges discussed above and lack of a demonstration of a vastly superior device grown on AlN. However, in order to advance AlN PVT and HVPE crystal growth, and consequently also AlN-based device technology, significant investments need to be made not only in further research and development of the AlN crystal growth, but also in AlGaN epitaxy, property control, and development of associated devices. Only the development of all these components in unison will ultimately justify the AlN-based technology.

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## 4 – Radiative recombination efficiency of AlGaN quantum wells; Do we estimate it accurately in a proper way?

Yoichi Kawakami, Mitsuru Funato, Ryota Ishii Kyoto University,

#### Status

Recent progress on the research of nitride semiconductors has led to the realization of highly efficient blue LEDs, active layers of which are composed of InGaN-based quantum wells (QWs). Currently, the improvement of efficiencies in Al-rich AlGaN-QWs based LEDs is a challenging subject for the development of deep ultraviolet (DUV) emitters. The external quantum efficiency (EQE) is expressed by the product of three terms, carrier injection efficiency, radiative recombination efficiency ( $\eta_{rad}$ ) and light extraction efficiency. Although EQEs are directly measurable values, it is difficult to resolve them accurately into the three terms.

The  $\eta_{rad}$  has generally been estimated by the temperature dependence of photoluminescence (PL) intensities. However, it should be noted that these values are overestimated in many cases if they were based on the assumption that the  $\eta_{rad}$  is 100 % at cryogenic temperatures. Precise PL measurements dependent on photo-excitation intensity [1], and rate equation analysis on time-resolved PL data [2], have revealed the failure of such an assumption. Moreover, one should be aware that  $\eta_{rad}$  is dependent on carrier density in the active layers. Carrier densities can be estimated precisely if the selective photo-excitation is made in the PL measurement, but it is not the case for the non-selective excitation.

In recent years, various experimental techniques have been reported for the estimation of  $\eta_{rad}$ . One approach is to detect the absolute emission intensities by means of the omnidirectional PL measurements using an integration sphere [3]. The second approach is to detect the heat generation induced by the non-radiative recombination processes, by employing techniques such as transient lens spectroscopy [4] or photo-acoustic spectroscopy [5]. In the former approach, estimation not only of light extraction efficiency but also of photon-recycling-effect is the key for the exact estimation of  $\eta_{rad}$ . In the latter approach, the quantification of the heat signal is the subject for the accuracy in determining  $\eta_{rad}$ .

#### **Current and Future Challenges**

If we aim at the improvement of  $\eta_{rad}$ , it is very important to understand the carrier recombination processes as initially discussed in the Shockley-Read-Hall model. The carrier recombination rate (*R*) can be divided into (1) nonradiative recombination rates due to defects and/or dislocations ( $R_{nr}$ ), (2) radiative recombination rate ( $R_r$ ) and (3) nonradiative recombination rate due to Auger process ( $R_{au}$ ), so that  $R=R_{nr}+R_r+R_{au}$ . In the so-called ABC model, the term of  $R_{nr}$ ,  $R_r$  and  $R_{au}$  is assumed to be proportional to injected carrier density (*n*), to the square of *n*, and to the cube of *n*, respectively if the background carrier density in the active layers is negligibly small. Consequently, *R* can be expressed as  $R=An+Bn^2+Cn^3$ , and we can derive  $\eta_{rad}=R_r/R=Bn^2/(An+Bn^2+Cn^3)$ . This model gives us apparently good results on the fitting of optical powers as a function of injected currents, by choosing proper parameters of *A*, *B* and *C*. Therefore, a lot of research groups have reported their fitting results based on this model. However, there has been no direct verification of such power laws in the ABC model, and there exists the risk not only for misunderstanding the physics of recombination processes but also for misinterpreting the  $\eta_{rad}$  estimation.

We have performed photoluminescence (PL) measurements on Al-rich AlGaN QWs in a wide range of photo-excited carrier densities, and carrier recombination processes have been analysed with a model based on rate equations. This has led to the quantitative estimation of  $\eta_{rad}$  values. Our finding was that radiative recombination processes are exciton recombination under relatively small carrier densities which correspond to the operating conditions of LEDs [2]. Since exciton recombination has to be treated as mono-molecular process, its radiative recombination should show single exponential decay which is independent on exciton density. Moreover, the probability of carriers/excitons to be captured by nonradiative recombination centres (NRCs) is not constant but decreases with increasing carrier/exciton densities because of the effect of saturation induced by the filling of NRCs. Figure 1 shows the PL lifetimes of an AlGaN/AIN QW emitting at 245 nm at room temperature which are increased with increasing photo-excited carrier densities due probably to the saturation of NRCs. These results indicate the failure of the conventional ABC model.

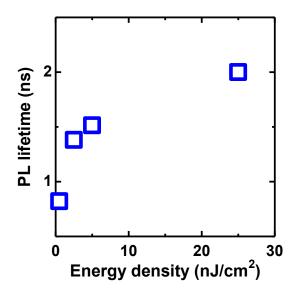


Figure 1. PL lifetimes of an AlGaN/AlN QW emitting at 245 nm under various photo-excitation energy densities which are taken at room temperature.

#### Advances in Science and Technology to Meet Challenges

The visualization of radiative and nonradiative recombination centers is the key to reveal the recombination mechanisms in semiconductors. Cathodoluminescence (CL) is a useful technique to map CL intensities in AlGaN-based DUV emission. However, the spatial resolution is affected not only by the spread of the incident electron beam in the sample but also by the diffusion length of the excitons/carriers. Scanning near-field optical microscopy (SNOM) has various merits for assessing the detailed recombination dynamics. We can choose the optical mode in the PL mapping with SNOM. In the illumination collection mode (IC-mode), both photo-excitation and the PL detection are performed through an optical fiber probe with small aperture. In the IC-mode, the spatial resolution is only limited by the diameter of aperture. The best spatial resolution achieved is down to about 10 nm in the visible spectral detection [6]. In the illumination mode (I-mode), the optical excitation is made through an optical fiber probe while the PL detection is made with a lens in the far-field configuration. In this mode the spatial resolution may be limited by the diffusion length like in the case of CL mapping. In the collection mode (C-mode), photo-excitation or electrical injection is done on large areas while the detection is through an optical fiber probe. Careful comparison of PL

mapping images taken at the same area in between IC-mode, I mode and C-mode (we named it at as multi-mode SNOM) gives us useful information on the carrier/exciton diffusion processes [7].

However, there have been existing technological difficulties in developing the SNOM in the DUV region. The shortest wavelength reported was 227 nm in the excitation, and longer than 240 nm in the detection with a spatial resolution of about 100 nm [8]. We have recently achieved an excitation with 206 nm, and the detection down to 220 nm with a spatial resolution better than 100 nm [9, 10]. As illustrated in Fig. 2, one target is to reach the detection wavelength down to 200 nm with an improved spatial resolution.

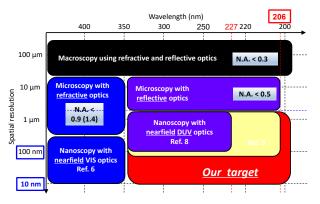


Figure 2. Current status and future prospect of PL-mapping technology.

#### **Concluding Remarks**

In this contribution, we have reviewed the issue in estimating  $\eta_{rad}$  of UV-emitting AlGaN QWs. The role of excitons in the emission mechanism has already been reported for InGaN QWs [11]. Since the exciton binding energies of AlGaN QWs are much larger than those of InGaN QWs, one should bear in mind the importance of excitons in discussing the recombination model. As for the origins of NRCs in Al-rich AlGaN QWs, point defects play more important role than in InGaN based QWs [12]. Development of characterization techniques such as DUV SNOM that visualizes the capture process of excitons or carriers to such NRCs and establishment of estimation scheme of  $\eta_{rad}$  may contribute to drastic improvement of EQE of UV LEDs.

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# 5 - Point defects Pramod Reddy<sup>1</sup>, Ramón Collazo<sup>2</sup> 1 Adroit Materials Inc. 2 North Carolina State University

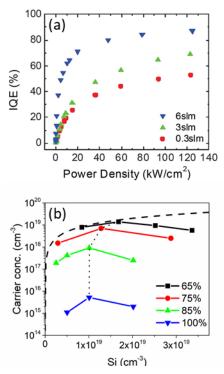
#### Status

There exists a "point defect problem" in Al rich  $Al_xGa_{1-x}N$  (x > 0.5) that has presented a major challenge to implementing AIGaN-based deep-UV optoelectronics. An increased incorporation of impurities and generation of native defects such as vacancies has been observed with increase in AI content (necessary for a technological push into deep-UV regimes) and is likely a consequence of the dependence of the defect formation energy on the Fermi energy whose range increases from GaN (3.4 eV) to AIN (6.1 eV). The incorporated point defects vary with the growth method, growth facet, doping type and concentration, and the growth environment, i.e., temperature, metal/N chemical potentials, pressure etc., and accordingly impact various optical and electronic properties of different regions in the optoelectronic device and hence the technological feasibility of ultra-wide bandgap-based deep UV emitters. Consequently, there are many point defect phenomena that are of high technological interest and we mention a few in the following: in metalorganic vapour phase epitaxy (MOVPE) grown AlGaN, when doped with Si (n-type) at low to moderate levels (<1x10<sup>19</sup> cm<sup>-3</sup>), and in unintentionally doped regions, carbon and vacancyoxygen complexes are reported to be the major compensating impurities and are a major constraint in designing active regions in optoelectronics including quantum wells.<sup>1,2</sup> At higher concentrations. Si itself forms an acceptor complex with an Al/Ga-vacancy resulting in a selfcompensation mechanism.<sup>3</sup> Further, at AI content>80%, Si relaxes from a donor type IIIsubstitutional site to an acceptor type DX configuration and results in a monotonically increasing activation energy from 20-50 meV for x<0.8 to around 300 meV for x=1.<sup>4</sup> In p-type AlGaN, doping itself is less understood with reports of both very low and very high dopant (Mg) activation energies.<sup>5,6</sup> Nevertheless, the hydrogen passivation and self-compensation by nitrogen vacancies have also been reported in Al-rich AlGaN similar to GaN. With recent availability of single crystal AIN substrates and very low dislocation density ( $<10^3$  cm<sup>-2</sup>) AlGaN epitaxy on them<sup>7</sup>, interesting phenomena have been observed. They include: the crucial balance between carbon, silicon and oxygen impurities in PVT-grown AIN single crystals whose ratios control the optical properties of the substrate<sup>8</sup>, and the observed dependence of the compensating point defect (vacancy-silicon complex) formation and hence electronic performance on the extended defect (threading dislocation) density.<sup>7</sup>

#### **Current and Future Challenges**

The various point defect phenomena mentioned in the previous section manifest very important technological challenges to the implementation of Al-rich AlGaN-based optoelectronics: The strong dependence of internal quantum efficiency (IQE) and associated non-radiative recombination coefficient in the quantum wells and active regions of UV emitters on the impurity concentrations demonstrates that point defects act as recombination centers thereby greatly reducing device performance.<sup>9,10</sup> Hence, low doped or "intrinsic" regions such as quantum wells require low point defect density for high IQEs. This is evidenced by the low IQE shown in Figure 1(a) for AlGaN/AIN multi-quantum well structures at low ammonia flows where incorporation of impurities such as carbon and oxygen is energetically favorable. Further, generation of native defects ( $V_{III}$  and  $V_N$ ) and formation of defect complexes have also been demonstrated to be energetically favorable. The point

defects have been identified by density functional theory (DFT) analysis in corroboration with thermodynamic analysis and electrical, secondary ion mass spectrometry and photoluminescence characterizations in III-nitrides especially at high AI content.<sup>1,2,11</sup> Further, at higher doping concentrations (n- or p-type) required in charge injection layers, conductivity exhibits a "knee behavior" due to self-compensation (vacancy-silicon complexes in n-type and nitrogen vacancies in p-type).<sup>7</sup> Hence, an upper doping limit exists that lowers the maximum conductivity resulting in increased ohmic losses and large contact resistances. Finally, the relaxation of Si from donor configuration to a DX center configuration has precluded AIN and Al<sub>x</sub>Ga<sub>1-x</sub>N (x > 0.8) from being employed in deep-UV optoelectronics.<sup>4</sup> Consequently, DX control is one of the most important technological challenges for implementing LEDs and lasers operating at wavelengths <240 nm. A summary of the Si behavior in AIGaN is shown in Figure 1(b) showing the "knee behavior" due to selfcompensation and the abrupt decrease in carrier concentrations for AI content >80%. So far, we have mentioned the point defect issues within the active regions of the device structure. However, point defects have also played a major role in substrate suitability. Specifically, in the development of physical vapor transport (PVT) AIN single crystals with dislocation densities  $< 10^3$  cm<sup>-2</sup> which promised highly efficient AlGaN-based optoelectronics, excessive carbon has led to absorption in the deep-UV region and has posed the most significant challenge to implementing UV emitters directly on PVT AIN substrates.<sup>12</sup>



**Figure 1.** (a) The IQE for AIGaN/AIN MQW structures on single crystal AIN substrates as a function of excitation power density shown for different point defect concentrations (here, higher ammonia flow corresponds to lower non-radiative recombination defect centre concentrations). (b) The dependence of carrier concentration as a function of doping concentration for different AI compositions in the alloy AIGaN. The black dashed line assumes the carrier concentration to be

#### Advances in Science and Technology to Meet Challenges

The difficulty in overcoming the point defect challenge arises from the dependence of the type and concentration of these defects on a large number of factors including growth technique, the precursors, growth mode, temperature, doping, etc. Developing specific experimental optimizations for each point defect may not be feasible due to the high iterative costs and the likely conflicting requirements of different point defects and other aspects of

crystal quality that affect device performance. A general solution would be a predictive point defect control scheme that may be employed to determine the optimal growth conditions within predefined constraints determined by other factors including growth rates, crystal quality and system limitations. Such a point defect scheme would employ (a) DFT analysis in corroboration with characterization techniques such as secondary ion mass spectroscopy (SIMS) and photoluminescence (PL) to identify the point defect responsible for a particular challenge and (b) dependence of thermodynamic and reaction kinetics describing the growth process on practical growth knobs and (c) finally understanding the relationship between the point defects and the thermodynamics and reaction kinetics describing the growth process. This would require a highly synergistic collaboration requiring the development and extension of DFT from typical zero kelvin and standard states to experimentally employed growth temperatures and supersaturated states determined from the developed thermodynamic and reaction models. In this direction, recent developments of a point defect control model directly providing a quantitative relationship between point defect formation energies and growth knobs has allowed for quantitative predictions of carbon as a function of growth conditions.<sup>13</sup> Finally, using the dependence of point defects on the chemical potentials (metal rich vs. N rich), and employing single crystal AIN substrates, record high IQEs (>80%) and low lasing thresholds (<10 kWcm<sup>-2</sup>) have been achieved.<sup>9,14</sup> An illustration is seen in Figure 1(a) where a shift towards N rich chemical potential via increased ammonia flow results in lower non-radiative recombination centre concentrations and a significant improvement in IQE. However, significant research efforts are necessary to generalize the scheme to different materials and growth techniques by including reaction kinetics and transients into the theoretical models so that for a given growth technique and practical constraints, the optimal growth conditions for a given application can be determined theoretically without the requirements for expensive experimental iterations. Further, point defects form when they are energetically favourable at equilibrium. Hence non-equilibrium doping techniques such as implantation need to be investigated in Al-rich AlGaN and AlN. Another solution would be to develop a point defect control scheme that targets a class of point defects having a common character where the exact identification is unnecessary. In this direction, manipulation of the Fermi level or electron chemical potential is another useful tool to control the defect formation energy of a particular class of defects i.e. charged compensating defects. Hence, defect quasi Fermi level (dQFL) control-based point defect reduction has been employed to improve the performance in different material systems such as nitrides and arsenides, with both n-type and p-type doping, with very different defect configurations, but with one common characteristic i.e. all the defects were of compensating type and charged.<sup>15,16</sup> Here, a significant advantage would be applying the point defect control scheme by external mechanism (in this case, an above bandgap light source) without altering the growth conditions. A summary of the developed point defect control frameworks is shown in Figure 2.



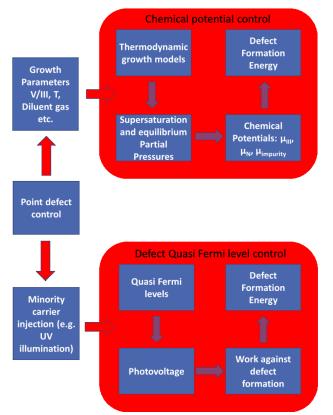


Figure 2: A schematic of targeted and predictable point defect control via chemical potential control and defect quasi Fermi level control.

#### **Concluding Remarks**

In conclusion, the "point defect problem" in Al-rich  $Al_xGa_{1-x}N$  (x > 0.5) requires a comprehensive solution for high performance deep-UV optoelectronics. Although experimental iterative approach towards low defect densities may be feasible in the short term, a long-term solution requires an understanding of point defect incorporation and "known and controlled" growth environments. In this direction, the energy of formation of the defects may present a general solution where the reaction constituent chemical potentials and electron chemical potential may be appropriately tuned within predefined constraints to obtain minimum defect incorporation and maximum efficiencies. This approach requires extending the DFT analysis beyond the current 0K and standard state calculations towards practical growth environment to understand point defect incorporation and developing and utilizing developed thermodynamic and reaction kinetic models that theoretically describe the growth environment which can then be controlled by internal knobs (e.g. gas flows, temperature etc.) and external knobs (e.g. illumination).

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#### 6 - Toward ohmic n-contacts on n-AlGaN with high Al mole fraction

#### L. Sulmoni

#### Institute of Solid State Physics, Technische Universität Berlin

#### Status

Ohmic contact formation on n-type III-nitrides is well established, at least for n-GaN. The basic mechanism responsible for ohmic behaviour involves N extraction by the contact metal upon thermal annealing and the formation of a heavily n-doped thin layer in the proximity of the interface [1,2]. In fact, N vacancies act as donors in GaN thus enhancing the probability of tunnelling for carriers through the thinner Schottky barrier [3,4]. Typically, the metal stack consists of four metals divided in 2 bilayers [2]. The first bilayer is responsible for the contact formation whereas the second one, usually referred to as capping layer, prevents the oxidation and the outdiffusion ("ball up") of the underlying metal stack. The most widely employed contact scheme is Ti/Al/Ni/Au. In many publications, the onset of the ohmic behaviour was attributed to the formation of a crystalline TiN layer (~5-20 nm thick) at the metal/semiconductor interface, indicating the reaction between Ti and GaN [1,2,4,5]. While investigating the n-contact formation for higher Al mole fractions, the most common approach has been to transfer the already established metallization scheme for n-GaN to the n-AlGaN material system. Unfortunately, Ti-based electrodes have shown to form ohmic contacts with n-AlGaN only up to an AlN molar fraction of 0.58 [6]. Typically, the best reported values of contact resistivities are lower than  $10^{-6} \Omega$  cm<sup>2</sup> on n-GaN. However, due to the need of a transparent current spreading layer, UV light emitters utilize n-Al<sub>x</sub>Ga<sub>1-x</sub>N [7]. Up to an Al mole fraction of x = 0.7, contact resistivities on  $n-Al_xGa_{1-x}N$  are similar to n-GaN but increase drastically to 1  $\Omega$  cm<sup>2</sup> on n-AIN (Fig. 1) [8], which goes hand in hand with the observed increase in n-sheet resistance [9]. In terms of wall-plug efficiency (WPE), devices emitting in the UV still exhibit significantly lower efficiencies than blue LEDs: >80% at 450 nm, >4% at 280 nm and >0.01% at 230 nm [7,10,11]. As discussed in the Sections 10 and 11, the reduced external quantum efficiency (EQE) at shorter wavelength is responsible to a significant extent for this efficiency drop. Nevertheless, minimizing the contact resistivities of p- and n-layers in deep UV LEDs and lasers is essential for improving WPE, reducing operating voltages and avoiding resistive Joule heating.

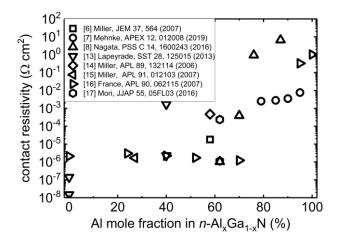


Figure 1. Contact resistivity as a function of the AI mole fraction for n-contacts on n-AI<sub>x</sub>Ga<sub>1-x</sub>N in the entire alloy composition.

#### **Current and Future Challenges**

The causes for the non-ohmic characteristics of Ti-based contacts on n-AlGaN with high Al mole fractions are twofold. Firstly, the Schottky barrier increases due to the lower electron affinity of AlGaN with increasing Al mole fraction as well as the lack of appropriate low work-function metals [1]. The lack of metals with work-functions lower than 4 eV is a fundamental physical limitation. Therefore, the main strategy in order to achieve ohmic contacts on n-AlGaN is to reduce the Schottky barrier width and therefore enhance tunnelling probabilities. Unfortunately, N extraction by Ti will become less energetically favourable as the AlN bond in the n-AlGaN layer is stronger than for GaN (lower formation enthalpy) [2]. During the annealing process, Ti substitutionally replaces Ga in the alloy leading to the formation of voids, highly defective Al+Ti+N phases and TiN protrusion islands, preferentially formed along dislocations, at the metal/nitride interface thus hindering the development of an uniform contact area [2,5]. Due to the higher resistivity of n-AlGaN layers and of n-contacts in comparison to n-GaN, an interdigitated n-electrode is usually implemented during the microfabrication process in order to improve the current uniformity in the emitting area.

Nonetheless, a careful adjustment of the metal stack and the thermal annealing conditions is not the only challenge in order to form low resistance and ohmic n-contacts to n-AlGaN at high Al mole fractions. Other important parameters play a critical role, such as the doping concentration in the underlying n-AlGaN layer and the dry etching conditions. Since UV LEDs are fabricated on a non-conductive substrate such as sapphire, the buried n-AlGaN contact layer must be exposed using plasma etching. Plasma-induced damage often results in an etched surface of poor quality thus undermining significantly the optical and electronic properties of n-AlGaN. Cao et al. reported that when increasing the Al mole fraction the plasma-induced defects act as deep-level centers, pinning the Fermi level and increasing the Schottky barrier height [12]. It is thus mandatory to carefully adjust the dry etching conditions and to introduce an adequate surface treatment in order to minimize the plasma damages and to assist the n-contact formation.

#### Advances in Science and Technology to Meet Challenges

In order to reduce the aggressive Ti-GaN reaction occurring in Ti-based n-contacts, different strategies have been suggested such as an Al layer with an adequate thickness on top of Ti (optimal Ti/Al ratio), the implementation of diffusion barriers between Ti and AlGaN (e.g., Cr, Zr, Nb,...) or the substitution of Ti by V. In the latter case, the generation of a heavily n-doped thin layer at the AlGaN interface is driven by the Al in the metal scheme while V acts as an indiffusion barrier [13]. Circumstantial evidence reported by several authors of the N extraction from the n-AlGaN layer and the consequent low n-contact resistance was the formation of a continuous thin AlN layer (~2-5 nm thick) at the interface [6,13]. Nevertheless, the driving force for the reaction metal/AlGaN weakens as AlGaN becomes more energetically stable with the increase in Al mole fraction: N atoms create a more stable compound with Al than with Ga due to the lower formation enthalpy of AlN (-318.1 kJ/mol) in comparison to GaN (-110.9 kJ/mol) [2,5]. In fact, it was possible to obtain ohmic V-based n-contacts only up to an AlN mole fraction of 0.7 [8]. The challenge is to find or develop a material (preferably highly conductive as a metal) able to form more stable nitrides than TiN or AlN on n-AlGaN layers with high Al content.

The high crystal quality with TDD lower than 10<sup>4</sup> cm<sup>-2</sup> of AIN bulk substrates is expected to improve the IQE of deep UV devices (see Section 3). Unfortunately, due to the high ionization energy of donors in n-AIGaN layers with very high AI mole fraction, conductive AIN bulk substrates are not expected to be utilized in near future. In fact, the practical use of such substrates would be beneficial especially for UV lasers to circumvent the need of exposing the buried n-AIGaN with plasma etching as mentioned before (current fabrication techniques typically involve substrate removal followed by a flip-chip process) thus allowing for a more homogeneous vertical carrier injection.

#### **Concluding Remarks**

Ohmic n-contacts on wide-bandgap semiconductors such as n-AlGaN with high Al content are challenging given several material limitations. First of all, the self-compensation of the Si dopant sets an upper limit on the amount of available donors in the n-AlGaN layer thus frustrating any effort in thinning the depletion region of the Schottky metal/semiconductor contact. Secondly, metals with sufficiently low work-function are not available. Finally, the N vacancy formation is hindered as the AlN bond in n-AlGaN becomes more and more energetically stable with decreasing Ga content and obstructs the formation of a heavily n-doped layer at the interface upon electrode annealing. Therefore, breakthroughs in terms of n-conductivity are expected to occur by new material concepts.

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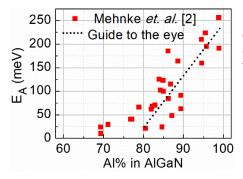
# 7 – Doping of AlGaN Biplab Sarkar<sup>1,2</sup>, Ramón Collazo<sup>2</sup> 1 Indian Institute of Technology Roorkee 2 North Carolina State University

#### Status:

A high electrical conductivity in n and p-type AlGaN is necessary to minimize the resistive losses in the epitaxial layers of UV light emitters. Furthermore, a high free carrier density also leads to low-resistance contacts. Thus, effective doping of AlGaN has been under serious investigation for more than two decades. N-type dopants (Si, Ge etc.) can readily be incorporated in GaN and Al-rich AlGaN epitaxial layers during growth. These dopants provide a relatively low ionization energy ( $E_A$ ) and high free electron density (> 10<sup>19</sup> cm<sup>-3</sup>) in AlGaN films. However, Mg is the only dopant that so far can be considered viable for p-type GaN and AlGaN epitaxial films. Mg shows a high  $E_A$  (> 120 meV) in metalorganic vapour phase epitaxy (MOVPE) grown p-GaN films, resulting in a free hole density that is only a fraction of the doping density. This becomes worse as the Al mole fraction increases in the AlGaN films, as  $E_A$  is expected to increase monotonically to > 500 meV for AlN films. As mentioned in the previous chapter, increasing the Al-mole fraction in AlGaN results in several point defects and charge compensators. Accordingly, several methods have been developed recently to achieve a high electrical conductivity in both n-type and p-type AlGaN epitaxial layers. This chapter is dedicated to highlight the advances in the doping of AlGaN to achieve a low  $E_A$  and high electrical conductivity.

#### n-type doping of (Al)GaN

A low  $E_A$  of Si donors in (Al)GaN yields a free electron density in the range of ~10<sup>19</sup> cm<sup>-3</sup> and conductivity of < 10 m $\Omega$ .cm in n-Al<sub>x</sub>Ga<sub>1-x</sub>N films (x < 0.8).<sup>1,2</sup> Availability of native substrates is a bonus for achieving a higher conductivity in Al-rich AlGaN films.<sup>3</sup> However, the donor  $E_A$  undergoes a steady rise in Al-rich AlGaN films when the Al mole fraction goes above 80%, as shown in Figure 1. This sharp increase in  $E_A$  is attributed to the formation of Si-DX centers acting as acceptor-type compensating point defects. Furthermore, the Si doping in AlGaN is also responsible for a "kneebehaviour" in resistivity with increase in the doping density.<sup>1,2</sup> While the low Si doping density is responsible for the formation of carbon and vacancy-oxygen complexes, a high Si doping leads to self-compensation and vacancy related complexes (e.g.  $V_{III}+Si$ ).<sup>4,5</sup> All these acceptor type point defects compensate the free electron density in Al-rich AlGaN films.



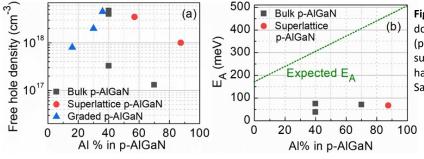
**Figure 1.** Donor  $E_A$  of Si in Al-rich AlGaN epitaxial films.<sup>2</sup> Si donor  $E_A$  in n-GaN is ~15 meV, which increases to ~50 meV in n-Al<sub>x</sub>Ga<sub>1-x</sub>N (x < 0.8) epitaxial films.<sup>1</sup> Thus, nearly full ionization of free carriers at room temperature is typically observed in n-Al<sub>x</sub>Ga<sub>1-x</sub>N (x < 0.8) epitaxial films.

Apart from Si, germanium and oxygen have also been attempted to dope n-AlGaN films. Ge typically shows a donor  $E_A$  of ~ 30 meV in GaN, and is predicted to allow a doping density beyond the Mott transition.<sup>6</sup> However, the onset of DX center formation is predicted to arise at an Al mole fraction of 52%, much lower than the Si counterpart. Therefore, a very high doping density can be achieved in Al<sub>x</sub>Ga<sub>1-x</sub>N films (x < 0.52) using Ge doping. Similarly, oxygen can also be incorporated in AlGaN epitaxial films, and gets incorporated during the growth due to the presence of residual oxygen in the growth chamber. However, oxygen donors show a much higher  $E_A$  than Si, and the onset of DX

centers also occurs at a lower Al mole fraction (~61% Al mole fraction) in AlGaN.<sup>7</sup> All these issues make Si the preferred choice for n-type doping of AlGaN epitaxial films.

#### p-type doping of AlGaN

Achieving a free hole density of the order of  $10^{18}$  cm<sup>-3</sup> in GaN and AlGaN films has been a major challenge for decades. Mg is the only known viable acceptor doping source that can occupy a substitutional site in (Al)GaN epitaxial films. A free hole density in the order of  $10^{17}$  cm<sup>-3</sup> is achievable in p-GaN films grown on foreign substrates using high temperature growth techniques (e.g. MOVPE). Although relatively lower temperature growth (e.g. molecular beam epitaxy) results in a higher free hole density due to lower hydrogen incorporation, the subsequent device processing incurs very high temperature process steps raising several reliability issues. Native AlN substrates offer a suitable alternative for high temperature p-GaN growth having free hole density nearly an order of magnitude higher than the foreign substrates.<sup>8</sup> The lower dislocation density of single crystal AlN substrates minimizes the incorporation of donor-type nitrogen vacancies in p-GaN. An  $E_A < 100$  meV is thus achievable in p-GaN films grown on single crystal AlN substrates using MOVPE.



**Figure 2.** Recent reports on Mgdoping of AlGaN; (a) free hole density (p), and (b) acceptor  $E_A$  for bulk and superlattice p-AlGaN films. Images have been reproduced from B. Sarkar, *et. al.* [3].

For AlGaN, the Mg acceptor  $E_A$  is believed to increase monotonically from 120-200 meV in GaN to 500 meV in AIN.9 Alongside, compensating nitrogen vacancies are also expected to exhibit a significant reduction in formation energy in p-AlGaN w.r.t. increase in the Al mole fraction.<sup>10</sup> However, recent studies reported a significantly lower acceptor  $E_A$  in Al-rich p-AlGaN films, as shown in Figure 2. Kinoshita et. al. has reported an  $E_A$  of < 72 meV in bulk p-Al<sub>0.7</sub>Ga<sub>0.3</sub>N films grown on sapphire substrate.<sup>9</sup> Similarly, Chen et. al. reported a free hole density of ~4.75 x 10<sup>18</sup> cm<sup>-3</sup> in p-Al<sub>0.4</sub>Ga<sub>0.6</sub>N by using In-surfactants and a Mg delta-doping scheme.<sup>11</sup> Along with improvement in bulk doping, alternative approaches such as superlattice structures and polarization doping schemes have also been developed to achieve a high free hole density in p-AlGaN (Figure 2). Short-period superlattice structures consisting of several thin p-AlGaN films with alternating Al mole fractions allow periodic oscillations of the valence band of p-AlGaN layers. This technique result in regions where the Mg energy level is much closer to the Fermi-level. The effective acceptor  $E_A$  is thus reduced, and a free hole density in the order of 10<sup>18</sup> cm<sup>-3</sup> can be achieved in p-AlGaN superlattice structures.<sup>12</sup> Similarly, a graded p-AlGaN film (polarization doping scheme) results in the presence of polarization induced 3D charges, and hence the film is degenerate of free carriers. A free hole density of the order of 10<sup>18</sup> cm<sup>-3</sup> has also been reported from polarization doped p-AlGaN films grown on foreign as well as native AIN substrates.<sup>13</sup>

#### **Current and future challenges:**

Doping all III-nitride epitaxial films will remain a major focus of the community for the coming years. Si doping of AlGaN up to 70% Al mole fraction can be considered to be matured enough. However, several scattering mechanisms limit the carrier mobility thereby increasing the sheet resistance of the epitaxial films. Thus, a major effort will be required to increase not only the free carrier density, but also the carrier mobility in Al-rich AlGaN. Crystal growth technique has to be optimized to minimize the point defect incorporation and produce smooth high quality epitaxial films. A lower dislocation density in the epitaxial films minimizes the point defect incorporation, and provides a better internal quantum efficiency. Thus, AlGaN growth optimization on low dislocation density

templates will require a major effort. Recent reports on AlGaN growth on native substrates and high temperature annealed templates are promising.<sup>3,14</sup> Moreover, contact mechanism in n-type Al-rich AlGaN films is still unknown for Al-rich AlGaN films. Unlike GaN and Ga-rich AlGaN films, obtaining a low resistance ohmic contact is a challenge. Ti/Al based contacts typically show a non-linear current-voltage characteristic when applied to Al-rich n-AlGaN films. Moreover, current conduction in Al-rich AlGaN films is dominated by different tunnelling mechanisms such as trap-assisted tunnelling. A major effort will thus be required to understand the contact formation mechanisms to enhance the contact performance and mitigate the reliability issues of future III-nitride deep-UV emitters.

Similarly, Mg doping of AlGaN will remain a major focus for the coming years. The role of dislocation density, point defects, vapour supersaturation, etc. are yet to be well understood for p-(AI)GaN films. Alternative methods such as polarization doping and superlattice structures will be required to attain maturity before these technologies can be commercialized. Apart from free carriers, contacting the p-side of the III-nitride light emitters remains a major challenge. Standard Ni/Au contact metallization scheme to p-GaN results in highly resistive ohmic contacts with a contact resistance of > mid  $10^{-3} \Omega$ .cm<sup>2</sup> in most of the reported literature. Alternatively, the tunnel junction concept has been developed to achieve low resistance ohmic contacts to the p-side of the III-nitride light emitters. In this scheme, the contact is applied to a highly doped n-GaN film grown on p-GaN. When bias is applied, electrons are drawn from the valence band of p-GaN into the n-GaN due to the inter-band tunneling. Similarly, free holes form the valence band of p-GaN are injected into the light emitter's active region. Thus, low resistance ohmic contacts can be formed in the p-side of the IIInitride light emitter by contacting the highly doped n-GaN instead of the p-GaN film. Note that an additional applied voltage (hence power loss) is required for the tunnel junction to allow holes to be injected into the light emitter's active region. Moreover, growth of subsequent n-GaN films on Mgdoped GaN at high temperatures result in "memory effect" where Mg incorporates into the n-GaN. Thus, excellent doping control is mandatory to ensure a sharp and thin depletion region across the highly doped tunnel junctions. All these open challenges simply indicate the quantum of efforts required to fully explore the potential of all III-nitride based devices.

#### Advances in Science and Technology to Meet Challenges:

Recent advances in growth, characterization and measurement techniques for III-nitride emitters has been a remarkable achievement. Several research groups have developed point defect control schemes (such as defect quasi fermi-level control, chemical potential control, vapour supersaturation, high quality templates etc.) resulting in very high free carrier density in Al-rich AlGaN epitaxial films. Formation of DX centers in Si-doped Al-rich AlGaN films during growth is a physics limited issue. However, recent reports indicate that high energy implantation (a non-equilibrium process) of Si into Al-rich AlGaN and AlN films may provide pathway to mitigate the DX center formation. A significantly lower  $E_A$  in ion-implanted n-AlN film grown on AlN substrate has been demonstrated recently.<sup>15</sup> Therefore, a paradigm shift of the doping scheme can be looked into to mitigate some of the critical doping issues in Al-rich AlGaN films.

Talking about the active region of the III-nitride light emitters, nearly defect-free quantum wells providing > 80% internal quantum efficiency have been developed for both foreign and native substrates. Similarly, for laser diodes, doped waveguides and reflecting mirrors has been optimized to provide excellent wave confinement. However, to achieve lasing in deep-UV spectrum, resistive losses in the p-type cladding region must be minimized. Bulk Mg-doping and alternate doping schemes seems promising to provide low resistance p-AlGaN cladding regions. So far, most of the IIInitride emitters have been designed having p-GaN as the contact layer. However, p-GaN absorbs the deep-UV light emitted by the active region of the emitter. Thus, recent reports on low  $E_A$  and high free hole density in Al-rich p-AlGaN seem promising to directly apply contact metallization to the p-AlGaN cladding layer. Moreover, standard Ni/Au contacts used in the p-side of laser diode are also known to absorb the deep-UV light. To overcome this absorption, reflective contact metallization schemes to replace the Ni/Au contact have been developed. Takano *et. al.* has even demonstrated improved light extraction efficiency in deep-UV flip-chip LEDs (275 nm) by using transparent p-AlGaN contact layer and Rh mirror contact.<sup>16</sup> All these reports indicate the importance of developing cutting-edge epitaxial film growth and contact metallization schemes to realize highly efficient deep-UV emitters.

#### **Concluding Remarks:**

This chapter provides a brief review of current status of n and p-type doping in Al-rich AlGaN films for future highly efficient III-nitride light emitters. Identification of different point defects is necessary to understand the charge compensators responsible for compensating the free carriers in Al-rich AlGaN films. Epitaxial films grown on low threading dislocation density templates seems promising in many aspects. Recent reports on Mg doping in Al-rich p-AlGaN films is very promising for deep-UV emitters. With advancement in growth techniques for both foreign and native substrates, high performance Al-rich AlGaN based devices like LEDs and laser diodes can be realized in the near future.

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#### 8 – Boron-containing (Al, Ga)N heterostructures

Ferdinand Scholz<sup>1</sup>, Abdallah Ougazzaden<sup>2</sup>,

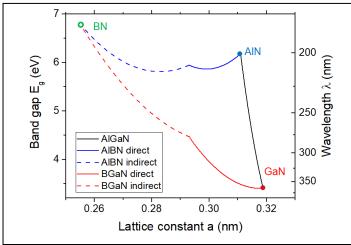
1 Institute of Functional Nanosystems, Ulm University

2 Georgia Institute of Technology, Georgia Tech-CNRS

#### Status

Boron (B) is the topmost and thus smallest element of the  $3^{rd}$  column of the periodic system of elements, being placed right above aluminum (AI) and gallium (Ga). Hence, one can expect a substantially smaller lattice constant and a larger band gap of B-containing nitrides (e.g.  $AI_{1-x}B_xN$ ) as compared to their B-free equivalents (i.e. AIN in this example). Typically, the lattice constant of such alloys follows Vegard's law being a linear relation between the composition x and the lattice constants of the binary end members (here AIN and BN), if both binaries crystallize in the same lattice type. However, while AIN and its Ga- and In-containing alloys crystallize in the hexagonal wurtzite structure, the thermodynamically most stable structure of BN is a hexagonal layered structure similar to graphite, whereas a wurtzitic phase only can be realized under specific boundary conditions. Anyway, by considering the data which are known for wurtzitic BN (w-BN), a variation of the lattice constants of  $AI_{1-x}B_xN$  by about 10 % when varying x can be expected with B incorporation into AIN leading to smaller lattice constants, while mixing Ga into AIN would lead to larger lattice constants. Therefore, B is a prospective candidate to manage lattice mismatch and strain in AlBGaN heterostructures.

The bandgap of such alloys typically follows a quadratic correlation described by a bowing parameter b besides the bandgaps of the binary end members. The band gap of w-BN is still heavily discussed (see, e.g., [1,2]); it was calculated to be about 6.8 eV with indirect character (Fig. 1), while the direct band gap is estimated to be around 10 eV [1]. Similarly, only sparse data are known about the bowing parameters. Calculated bandgap data for AIBN presented in [1] may be best described with a bowing parameter of 7 eV for the direct bandgap and of 4 eV for the indirect transition.



**Figure 1.** Bandgap versus lattice constant of the ternary alloys BGaN and AlBN. Notice the indirect band structure of w-BN. The connecting lines are plotted as a best compromise of unfortunately not yet fully consistent data from several sources. The direct band gap of w-BN is taken as 10.2 eV [1], whereas bowing parameters of 7 and 4 eV are assumed for the direct (full lines) and indirect transitions (broken lines), respectively, for both B-containing ternaries. For simplicity we assumed Vegard's law for the lattice constant. Any slight deviation (as sometimes discussed) would not change the major message of this diagram.

Of course, also essentially all other material characteristics are functions of the respective ternary or quaternary composition. Particularly the variation of the refractive index when mixing B to Al(Ga)N

alloys seems very promising for the realization of Bragg mirrors made of virtually strain-free AlGaN-AlBN superlattices [3]. Moreover, alloying with B creates new degrees of freedom for managing both, the spontaneous and the piezo-electric polarization in respective heterostructures [3, 4, 5]. Therefore, optical and electronic devices could benefit from the development of this material system.

#### **Current and Future Challenges**

These simple considerations lead to very promising properties of B containing group III nitride heterostructures. However, a closer look reveals a lot of challenges. The dominant problem is the controllable and reproducible synthesis of such alloys in high crystalline quality – being a major reason why their basic properties are still heavily debated. Calculations of their thermodynamic phase diagrams indicate that the AlBGaN material system exhibits a substantial miscibility gap at reasonable temperatures [6]. At temperatures around 1000°C, only few percent of B are expected to be incorporated into Al(Ga)N, whereas full miscibility is only predicted for temperatures exceeding 9000 K [7]. On the other hand, particularly for strain management, only small amounts of B are needed: 6% of B lead to lattice matching between AlBGaN quantum wells and AlN barriers for an emission wavelength of 270 nm, whereas only 2% are required for a strain-free Al<sub>x</sub>Ga<sub>1-x</sub>N-Al<sub>y</sub>B<sub>z</sub>Ga<sub>1-y-z</sub>N multi quantum well structure with the same target wavelength and reasonable barrier height (x ~ 0.55, y ~ 0.7).

Epitaxial growth results seem to confirm and even emphasize these problems. Very high B contents, mostly in the range of 10 – 15%, have been achieved by few groups by metalorganic vapor phase epitaxial growth at comparably low temperature of about 900 – 1000 °C (see, e.g., [8, 9]). Such layers and heterostructures exhibit fair X-ray diffraction results and have been studied with respect to their refractive index data in AlBN-AIN Bragg mirrors. Best reflectivities of only about 80% at 311 nm have been measured [3], but no PL is reported from such structures which may be taken as a confirmation of their limited crystalline quality. When directly compared to GaN, BGaN layers show reasonable luminescence only for a B content below 1%, whereas larger B concentrations dramatically reduce the emitted light intensity (see, e.g., [10]), although higher intensities are expected in strain-optimized structures [11]. By careful optimization of the growth procedure, strong PL intensities in AlBGaN layers containing 1-2 % of B similar as in respective B-free AlGaN layers (Fig. 2) have been recently reported [12]. Transmission electron microscopy (TEM) studies confirm the limited crystalline quality all AlBGaN layers with boron concentrations larger than some tenths of a percent. In consequence, many material properties of B containing nitrides are still questionable strongly limiting the consistency of device simulation studies.

#### Advances in Science and Technology to Meet Challenges

The major challenge for benefitting from the above mentioned promising properties of B containing AlGaN structures is the suppression of defect formation typically accompanied with the incorporation of major B amounts, e.g. by adopting nanoselective area growth [13]. To this end, a better understanding of the characteristics of B on the growing semiconductor surface is mandatory. Only scarce, if any data are available about its surface diffusion properties. Owing to the strong B-N bond, very high growth temperatures are recommended to enhance the B surface diffusion length. However, this may lead to strong parasitic gas phase reactions between the B precursor (typically triethyl-boron TEB) and ammonia (NH<sub>3</sub>). The mobility of B on the growing surface can also be increased by modulating the precursor flows [9, 14], i.e. supply the group III and group V precursors

sequentially, which also minimizes the risk of gas phase pre-reactions and hence may allow to use substantially higher growth temperatures. This approach, currently studied by many groups, needs further optimization. High resolution TEM may help to get a better understanding about the interaction between boron incorporation and defect generation. Moreover, some groups report about the incorporation of parasitic impurities, in particular oxygen and carbon, together with boron [15]. The reason for this behaviour is not yet well understood. The fairly strong chemical binding of the ethyl groups to boron in TEB may be responsible for the latter. Hence other B precursor molecules need to be investigated. Few studies have been done with diborane (B<sub>2</sub>H<sub>6</sub>) and borazine (B<sub>3</sub>N<sub>3</sub>H<sub>6</sub>) as well with tri-isopropyl-boron (TiPB), but here more detailed studies are needed to evaluate their usefulness for the growth of AlBGaN layers, particularly regarding the impurity problem. Such improved layer quality would also enable more elaborate determination of their structural and optoelectronic properties.

Obviously, the field is still at its infancy compared to the much more developed AlGaN and GalnN fields. For instance, successful p- or n-doping of BAIN and BGaN has not yet been reported. Similarly, high performance optical or electronic devices comprising AlBN and BGaN have yet to be demonstrated.

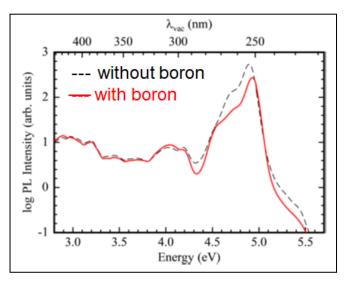


Figure 2. Low-temperature PL (10 K) of a thin AlGaN layer and a similar AlBGaN layer with about 1 % B. Notice the comparable intensity of both samples. From [12] with permission from John Wiley and Sons.

#### **Concluding Remarks**

The addition of boron to the group III nitride alloy components seems to open very promising solutions regarding strain, polarization and refractive index management in Al(B)GaN heterostructures. They may find applications in both optoelectronic (deep UV LEDs, UV detectors, sensors etc.) as well as in novel electronic devices (high electron mobility transistors, Schottky diodes etc.). However, we are obviously faced with very heavy problems regarding the epitaxial growth of B containing layers as a consequence of the extraordinary chemical properties of boron as a member of the second row of the periodic system. Very thorough optimization of the epitaxial growth procedure may lead to substantially better AlBGaN layer qualities as compared to today's situation, but it will remain a major challenge to achieve device performance data for, e.g., UV emitters better than obtainable by relying on B-free AlGaN structures.

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## 9 – Development of UV-A LEDs

# Peter J. Parbrook<sup>1</sup>, Tao Wang<sup>2</sup>

1 Tyndall National Institute and School of Engineering, University College Cork

### 2 University of Sheffield

#### Status

UVA covers the wavelength range 315-400 nm. Sources in this region have application in UV curing, photolithography, phototherapy (including melanoma treatment), security (banknote verification), dentistry, crime scene investigation and as a pump source for general illumination.<sup>1</sup> As a result of the wide range of applications, the UV market is dominated by UVA, along with the growing UVB/UVC area mainly for water purification.<sup>2</sup> The GaN bandgap (~362 nm) is in the centre of this range, with longer/shorter wavelengths achieved via InGaN, AlGaN (or InAlGaN) alloys. From a technological perspective the bandgap of GaN splits UVA into two distinct parts. For longer (>365 nm) wavelengths, devices can use adapted technology from that used in visible LEDs. For shorter wavelengths (<365 nm) there is a performance collapse, which can be seen clearly in the device external quantum efficiency (EQE) against wavelength plot shown in Figure 2 in Section 1 of this roadmap<sup>3</sup>. For most applications, the key parameter is raw optical power produced at the target wavelength. This correlates strongly to efficiency, due to the limits on the current density and heating that a device can tolerate, and hence the cost of ownership, running cost of the UV emission system. Improvements will lead to improved take up in systems using sources across this range.

Early reports of UVA-LEDs include: a 370 nm LED based on an AlGaN/GaN double heterostructure (DH) in 1994<sup>4</sup>; 350 nm UV-LEDs, using GaN/AlGaN quantum wells (QWs) for the active region (output power of 13  $\mu$ W at 20 mA) in 1998<sup>5</sup>; and the use of InGaN QWs for 380 nm was demonstrated by Mukai and Nakamura<sup>6</sup>. A 1 mW 348 nm UV-LED using AlInGaN quaternary as an active region at 50 mA was reported in 2002<sup>7</sup>. Using GaN as a substrate, the crystal quality can be greatly improved, leading to 352 nm UV-LEDs producing 0.55 mW at 20 mA (estimated internal quantum efficiency (IQE)>80%) in 2001<sup>8</sup>. These results are based on conventional GaN-on-sapphire technology, or GaN substrates. More recently, there have been significant improvements in the EQE of near UV-LEDs (NUV-LEDs), namely, the UVA-LEDs specifically in the spectral range from 365 to 400 nm, such as 30% EQE for 365 nm UVA-LEDs and 50% EQE for 385 nm UVA-LEDs. Properly packaged 365 nm UVA-LEDs with an output power of up to 12W have been reported.<sup>1</sup> For 380 nm LEDs such technology can now provide commercial devices with powers exceeding 1 W in a suitably packaged envelope. However, this is not the case for shorter wavelength devices.

#### **Current and Future Challenges**

The major challenge in all LEDs is to maximise wall plug efficiency (light output over electrical input). This can be defined in terms of the product of IQE, Light Extraction Efficiency (LEE) – giving EQE – and the electrical loss. As one targets shorter wavelengths these elements all become more challenging. LEE is the primary cause of the efficiency drop for wavelengths <365 nm. In all LED design a primary goal is to minimise all possible absorption losses in the device. For LEDs grown on GaN operating >365 nm there is minimal optical absorption as GaN is transparent. However, for wavelengths <365 nm optical absorption is harder to control as one generally is required to use a GaN or AIN based template layer to ensure high structural quality to ensure a high IQE. If devices are

grown on GaN many of the photons produced do not escape the diode, and LEE values of a few percent are typical. The alternative is to apply a "GaN-free technology" to prepare such LEDs on AIN templates. AIN, however, has a ~3% mismatch to GaN and UVA materials require high GaN content alloys, leading to significant strain control issues. Furthermore, convincing technologies to make a low resistance ohmic p-contact directly to AlGaN (as opposed to GaN) means that the vast majority of <360 nm LEDs still have absorption (hence LEE) issues.

IQE is affected by a number of factors in this region. For <360 nm point defects may become progressively more important at shorter wavelength as shown in **Figure 1**<sup>9</sup>. Typically, the V/III ratio for AlGaN growth is much less than (In)GaN to enhance atomic diffusion, potentially leading to significant increase in vacancy point defect density. A further issue is the control of the polarisation fields across the QWs. At longer wavelengths the need to include AlGaN to prevent carrier overflow, and the strain consequences to an otherwise GaN-based structure can lead to an IQE drop. Finally for <365 nm LEDs n-AlGaN must be used, resulting in alloy scattering and consequently a large mobility drop<sup>10</sup>, which coupled with the challenges in making n-AlGaN thick (due to strain and roughening) leads to additional voltage losses. This is in addition to further challenges in optimising the p- and n- electrical contact design.

#### Advances in Science and Technology to Meet Challenges

For UVA LEDs >365 nm package design is likely to have the biggest future impact on device performance, utilising best practice from those used in visible LEDs. This includes optimisation of light extraction and heat from the chip. Further, the low In content in the QWs for these devices means that carriers can potentially move to extended defects more easily than in their visible counterparts, so the optimisation of devices on lower defect GaN templates will be beneficial. This needs to be achieved either through obtaining best crystal quality cost effectively on sapphire (or Si with a substrate removal step and care to preventing cracking), or in finding routes to reuse expensive high quality templates with device lift off.

For devices operating <360 nm routes to improve the LEE are critical. Ideally this means a GaN-free growth process, or route to removing GaN material in post growth fabrication. An ideal would be use of thick, high quality AlGaN layers, which is an area but this is not easily achievable as the strain leads to cracking (GaN) or dislocation climb and layer roughening (AIN). There is developing research using such approaches (both for UVA and UVB LEDs) with improvements in emission quality and a more controlled roughening if a crystallographic miscut is introduced. However, this does lead to a step- bunched morphology with composition variations in the AlGaN, and additionally quantum well thickness variations. The impacts on this on device performance still requires further work.<sup>11</sup> Currently AlGaN on AlN is more promising, despite the issues of strain for the n-AlGaN buffer and

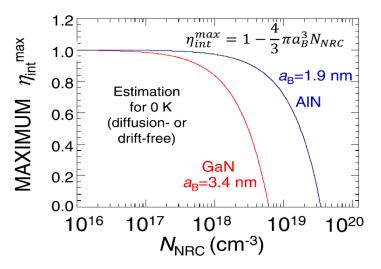


Figure 1. Maximum IQE as a function of point defect density.<sup>7</sup>

high Ga content AlGaN QWs (with polarisation field control issues) that this presents. Improved routes to allow smooth, thick AlGaN buffer growth of high crystal quality, for example using patterning to allow strain control, would revolutionise the short wavelength UVA.

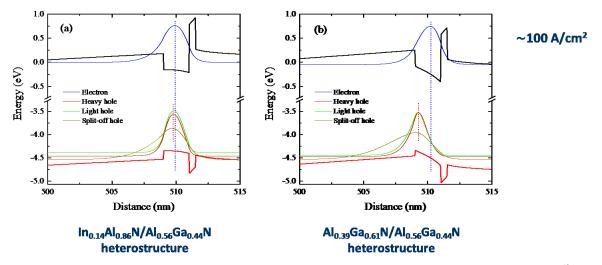


Figure 2. Improved electron-hole wavefunction overlap for InAIN-AlGaN vs AlGaN-AlGaN quantum well UVA LEDs (~340 nm)<sup>13</sup> [Copyright (2019) The Japan Society of Applied Physics].

As for UVB and UVC LEDs, making direct contact to AlGaN through doping and contact metal optimisation is also required. Such routes have demonstrated promise, but need work to retain the benefits to LEE while minimising electrical loss<sup>12</sup>. The high spontaneous polarisation in AlGaN means that polarisation fields are always an issue in UV LEDs. Thus the potential of using semi- or non-polar materials, which might allow thicker AlGaN as there is an easier route to strain relaxation, is worth investigating. In theory InAIN-AlGaN QWs can create polarisation-free active regions on c-plane for UVA, leading to improved electron-hole wavefuction overlap, as shown in Figure 2. However, while LED operation has been demonstrated the challenges in material growth are high<sup>13</sup>. Small quantities of In in InAlGaN QWs may have benefit for higher IQE, particularly in the 340-365 nm range.

#### **Concluding Remarks**

In conclusion, UVA LEDs can be split into two distinct bands. At longer wavelengths devices show high efficiency and can generally use adapted forms of the mature technologies using to make stateof-the-art blue LEDs. Devices with high powers are commercially available and improvements in package and defect reduction may lead to improved future performance. For shorter wavelengths (320-365 nm) where GaN absorption is a major problem compromising LEE in particular, development of a true GaN-free technology for devices is critical if UV LEDs to be fully exploited as sources in this range.

#### Acknowledgements

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#### 10 – UVB-LEDs

#### Tim Wernicke

Technische Universität Berlin, Institute of Solid State Physics

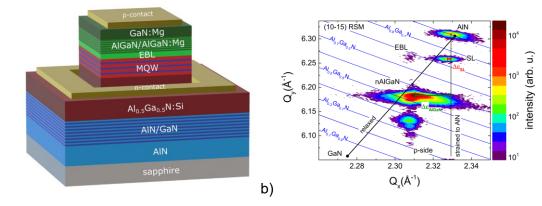
#### Status

For the UVB spectral range, 280 nm to 315 nm, the external quantum efficiency (EQE) values achieved so far are lower than those record efficiencies of LEDs emitting around 275 nm and UVA LEDs (see section 1), i.e., there is a "UVB gap" also extending into the UVA region especially in the 305 – 330 nm range with lower performance in comparison to shorter and longer emission wavelengths. Nevertheless UVB LEDs emitting at 305 nm are commercially available with an emission power of 100 mW at a current of 350 mA (EQE 6%) with an operation voltage of 7 V [1]. This EQE is similar to currently available LEDs emitting around 280 nm and 340 nm [1]. This disparity in reported record values and available products has technological and economic reasons. Research efforts and number of publications for UVB-LEDs is much smaller compared to the UVA and UVC spectral range, most likely because there is currently no high volume application (see Section 1) and combing all possible approaches to increase the light extraction efficiency (see section 13) have not been attempted yet. These novel approaches have a high potential for increasing the EQE of UVB LEDs as well but need to be tested for their impact on degradation behaviour before a general application. Nevertheless, UVB-LEDs have numerous applications, e.g., phototherapy, plant growth lighting and polymer curing (see section 1). Especially the first two require a peak wavelength at about 310 nm as light below 300 nm is harmful for these applications. From a technological point of view, the UVB-LEDs profit from most technological advances for UVC-LEDs but suffer from a lack of suitable substrates as they cannot be grown on defect reduced AIN – the large compressive strain leads to a formation of defects in UVB emitting quantum wells (QWs) [13, 16]. This can be mitigated by growing a relaxed  $Al_{0.5}Ga_{0.5}N$  buffer on AIN/sapphire template [6]. The relaxation process can be facilitated by introducing an AIN/GaN superlattice as indicated in Fig. 1 [6] and is only possible with a sufficiently high threading dislocation density (TDD) >  $3 \times 10^9$  cm<sup>-2</sup> as the compressive strain is released by tilt of edge type dislocations [6]. However, in this way a further reduction of the TDD in UVB LEDs is not easily possible leading to a lower internal quantum efficiency (IQE).

The electrical efficiency  $\eta_{el}$  of UVB-LEDs, which describes the losses by Joule heating at contacts and in the semiconductor layers, still gives room for improvement: Even comparably low operation voltages of 6 V at 100 mA [2] or 7 V at 350 mA [1] are several volts higher than the physical limit being defined by the bandgap in the quantum wells, i.e. close to 4 V (see also section 1). In the UVA these issues are solve with 365 nm LEDs exhibiting a very high electrical efficiency and low voltage of 3.49 V at 20 mA [14] and at 700 mA [1]. For UVB-LEDs ohmic n-contacts – Ti/Al-based as well as V/Albased - with resistivities of 10<sup>-5</sup>  $\Omega$ cm<sup>2</sup> (see section 7) and AlGaN layers with low resistivity have been realized. Therefore whole process needs to be optimized for lower voltages. Additionally, the operation voltage is increased if transparent p-doped AlGaN layers are employed [3,4].

The lifetime of UVB-LEDs can reach several thousand hours ([2], and see section 16) and even though their visible counterparts are much more stable, these values are sufficient for first applications.

In order to develop high efficiency and high power UVB-LEDs with EQEs exceeding 20% it is necessary to solve a number of challenges.



**Figure 1.** a) Schematic of a typical UVB-LED heterostructure. For coherent growth of the MQW emitting at 305 nm a relaxed  $Al_{0.5}Ga_{0.5}N$  layer is grown (from Susilo et al., pssa). b) XRD reciprocal space map around the AlN 105 reflection of a full UVB LED illustrating the strain relaxation process. The AlN/GaN superlattice is grown coherently strained on the AlN/sapphire template (nearly same  $Q_x$  value). The  $Al_{0.5}Ga_{0.5}N$  starts to grow first strained but then relaxes during further overgrowth ( $Q_x$  shifting to lower values) thus forming a metamorphic quasi-substrate that allows to grow EBL and buffer layer with the new lattice constant (from [5]).

#### **Current and Future Challenges**

a)

One of the main challenges to achieve UVB-LEDs with a higher light output power is reducing the threading dislocation density to allow for higher IQE values. Even though threading dislocations can be reduced by annihilation in a thick  $AI_{0.5}Ga_{0.5}N$  buffer layer the lowest threading dislocation density that could be achieved is still > 10<sup>9</sup> cm<sup>-2</sup>. To further increase the internal quantum efficiency of UVB emitting quantum wells, the relaxation process needs to be decoupled from threading dislocations and/or the annihilation of dislocations has to be enhanced. One approach is the structuring of the AIN surface before AIGaN overgrowth leading to relaxation and effective dislocation annihilation and TDD of 2 – 5 x 10<sup>8</sup> cm<sup>-3</sup> depending on the aluminium content in the layer [7]. A similar effect is achieved by introducing a 3D growth of  $AI_{0.55}Ga_{0.45}N$  and subsequent coalescence achieving TDD of 7 – 9 x 10<sup>8</sup> cm<sup>-3</sup> [16]. Another way to mitigate the effect of threading dislocations is growing UV-LEDs on highly miscut sapphire substrates of 1° with a step bunched surface leading to UVB-LEDs emitting at 300 nm with a record high EQE of 6% [2].

There are only few publications discussing the carrier injection efficiency  $\eta_{inj}$  mainly by a variation of the electron blocking layer (EBL) or by introducing an electron blocking heterostructure [9, 17] and the exact value of the injection efficiency is not clear, i.e., how much improvement of the external quantum efficiency can be achieved by further optimizing these layers. However, high efficiency LEDs require an almost perfect injection efficiency and the design of the electron blocking layer seems to have a crucial effect on the lifetime of the LEDs (see section 16).

The LEE has the highest potential for an EQE increase as current UVB LEDs do not incorporate techniques for increasing the LEE (see section 13). Successfully reducing the absorption in layers [17] and at interfaces as well as reducing the total internal reflection are the main tasks.

For the degradation of UVB LEDs the root cause seems to be deep levels at the pn-junction and within the quantum wells introduced by point defects leading to a reduction in output power (see section 16). The main challenge is to reduce these point defects by optimizing the growth parameters which will not only lead to longer lifetimes but also higher IQE and CIE.

#### Advances in Science and Technology to Meet Challenges

To further increase the IQE, a further reduction of the TDD down to 2x10<sup>8</sup> cm<sup>-2</sup> would be desirable [15]. Here approaches involving lateral overgrowth or maybe even AlGaN quasisubstrates prepared by hydride vapour phase epitaxy (HVPE) or achieving high quality AlGaN by high temperature annealing (see section 2) might be a necessary technological development. For the material of the QW novel indium or boron containing alloys might allow for a boost in IQE. Indium containing quaternary InAlGaN or even ternary InAlN might allow for an increased IQE by introducing localization and maybe also a reduced point defect density, however recent UVB LEDs with record high quantum efficiency employ pure AlGaN quantum wells [2,4] suggesting that TDD and point defect density are most important. Boron containing quaternary AlBGaN MQWs would allow to reduce the compressive strain (see section 8) and grow coherently strained on AlN making use of optimized AlN buffer layers or bulk substrates with much lower threading dislocation densities. However, the prerequisite for a successful implementation of such materials will be mastering their growth.

The LEE will by strongly increased by increasing the aluminium content in the p-doped AlGaN layers so the emitted photons are not absorbed in this layer [3,4], see also section 13. In comparison to UVC-LEDs, the necessary aluminium content to achieve transparency is much lower leading to a smaller increase of the operation voltage. However, very high LEEs will only be achieved in combination with reflective contacts leading to a significant improvement of LEE and EQE. Here the challenge is to find a highly reflective material that forms an ohmic contacts to p-doped AlGaN or realize photonic crystals (see section 13). The alternative approach to realize transparent LEDs is employing tunnel junctions as demonstrated for 326 nm LEDs (see section 17). To reduce internal reflections the substrate removal by laser lift off [19] or electrochemical etching [18] and/or roughening [12] will increase the light extraction efficiency. In parallel the encapsulation of UVB-LEDs will positively affect the LEE for transparent as well as non-transparent LEDs (see Section 11) and 13).

#### **Concluding Remarks**

UVB LEDs will also in future exhibit lower external quantum efficiencies than UVA and UVC LEDs due to the lack in suitable substrates with a low threading dislocation density. In future several approaches might allow for an effective reduction in threading dislocation density even though most of them require considerable development effort. A strong increase in LEE can be expected for the near future as increasing the aluminium content on the p-side and incorporating tunnel junctions lead to a lower additional voltage in comparison to UVC-LEDs due to the smaller bandgap of the material allowing for a high efficiency operation as soon as these techniques can be implemented.

#### Acknowledgements

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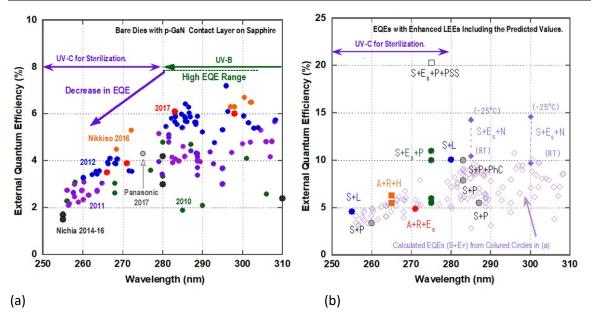
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# 11 – UVC LEDs Akira Hirano<sup>1</sup>, Hiroshi Amano<sup>2, 3</sup> 1 UV Craftory Co., Ltd. 2 IMaSS, Nagoya Univ 3 Nagoya Univ

#### Status

UVC LEDs with wavelengths between 250 and 280 nm are expected to be used as an alternative to low-pressure mercury lamps. In this section, UVC refers to this limited wavelength range. The main application of UVC light sources is sterilization, and the wavelength giving the peak disinfection efficiency is about 265 nm [1]. Low-pressure mercury lamps with an intense emission at the sterilization line (253.7 nm) have a high WPE of about 25% and a lifetime of 6,000 h. Although AlGaN-based deep UV-LEDs ( $\lambda$ <300 nm) in UVB have a potentiality of WPE of about 10% and the lifetime over 10,000h [1], WPE of 265 nm LEDs is significantly less than 10%. The LED lifetime decreases as the wavelength decreases from 280 nm [1,2]. With this background, at a wavelength of 275 nm, which is slightly shorter than 280 nm, higher initial EQEs over 10% were reported to be achievable by using a p-reflective contact and unknown encapsulation [3] and by integrating a preflective contact, patterned sapphire substrate, and lens [4], respectively. However, reliability tests have not yet been reported for the case of using a p-AlGaN contact, and the devices with a preflective electrode also exhibited an increased forward voltage (V<sub>f</sub>). In the case of using a p-GaN contact layer, EQEs of 4.9% at 270 nm with encapsulation [5] and 5.5-6.0% at 265 nm with distinctive roughening [6] were obtained using an AIN bulk substrate. Also, the results of reliability tests have been reported for 265 nm UVC LEDs with a p-GaN contact layer on sapphire [1,7]. Using a p-GaN layer, encapsulations using a fluoro-polymer [8] and a sapphire lens [2] were demonstrated to enhance the LEE 2.3-fold and 1.5-fold in UVC, respectively, which do not adversely affect the lifetime for UVC LEDs without increasing  $V_f$ . Figures 1(a) and (b) show the EQEs for bare dies of DUV-LEDs with a p-GaN layer grown on sapphire and the reported EQEs incorporating the techniques of enhancing LEE, respectively. Figure 1(b) includes the EQEs achievable from the results shown in Figure 1(a).



**Figure 1. (a)** EQEs from various groups for dies with p-GaN contact layer on sapphire. Colored data are the results obtained at UV Craftory. **(b)** Initial EQEs of UV-LEDs with roughening (+R), with encapsulation (+E), with a sapphire lens (+L), with a p-reflective contact (+P), with an n-reflective electrode (+N), and with the use of a PSS (+PSS). The subscripts 'S' and 'F' after '+E' indicate unknown/or silicone and fluoro-resin, respectively. 'A' and 'S' denote sapphire and AIN bulk substrates, respectively. The regrowth of an AIN layer with UVC transmittance by HVPE on an AIN substrate followed by removing the AIN bulk substrate is indicated as (+H). The data in **(b)** are limited to those published later than 2013. Also, the calculated achievable EQEs are shown by hollow symbols. Our calculated data for the case of fluoro-resin encapsulation from the data in **(a)** are indicated by hollow violet, and the solid violet is based on the experiment. In the graphs, data for 280-310 nm UVB-LEDs show the lower EQEs for UVC (250-280 nm). The EQE in UVC is considered to be suppressed by the decreases in IQE and CIE. [(a) and (b) are reproduced from Appl. Sci. [1]. (Copyright belongs to the authors)]

#### **Current and Future Challenges**

Generally, the near-future tasks are considered to be best discussed under the premise of using a p-GaN layer because the LED lifetime is expected to be longer than that of a low-pressure mercury lamp. An improvement in IQE by reducing the TDD and point defect density (PDD) and an improvement in LEE are considered to be priorities.

To reduce the TDD to near  $10^8$  /cm<sup>2</sup>, a typical approach is to use bulk AIN substrates with a TDD of less than  $10^6$  /cm<sup>2</sup>. Also, the TDD can be reduced to  $5 \times 10^8$  /cm<sup>2</sup> by using a  $1.0^\circ$ -miscut sapphire substrate [7]. Figure 2(a) indicates the further improvement of TDD by using  $1.5^\circ$ -miscut sapphire [1], showing the possible improvement in IQE by reducing the TDD.

The possibility of increasing the IQE by reducing the PDD has been reported [9,10]. A moderately Sidoped quantum well (QW) exhibited improved IQEs for AlGaN QWs with a TDD of greater than  $10^9$  /cm<sup>2</sup> near 250 nm [9]. Also, control of the PDD was shown by increasing the V/III ratio when growing 258 nm QWs [10] using an AlN bulk substrate. These results indicate that the IQE of UVC LEDs can be improved by reducing the PDD.

For the LEE, it is necessary to discuss the packaging techniques for UVC LEDs ( $\lambda$ <250nm). The molecular structure of the fluoro-resin used for encapsulating UVC-LEDs (265 nm) has been clarified to be polymerized perfluoro(4-vinyloxy-1-butene) with end groups of  $-CF_3$  [8]. However, the low refractive index of 1.35 limits the improvement in LEE. The refractive index of fluoro-resin is difficult to increase. Thus, consideration of the shape effect is important for increasing LEE. It is desirable to fabricate hemispherical lenses on surface-mounted devices (SMDs) by forming a lens array on a large AIN sheet (e.g., 4x4 inch<sup>2</sup>) followed by isolation to make small SMDs (e.g., 3.5x3.5 mm<sup>2</sup>). Another successful approach to improve LEE is to use sapphire lenses with a refractive index of 1.8 [2].

#### Advances in Science and Technology to Meet Challenges

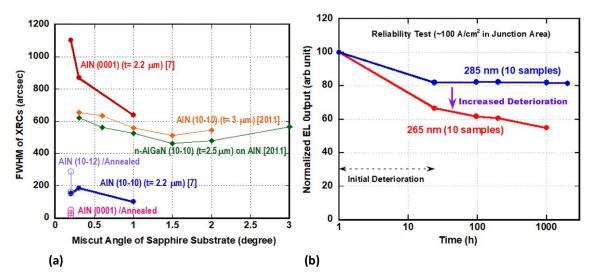
The utilization of EL emission from QWs penetrating p-contact layers is currently difficult in practice owing to the lack of both a low-resistivity p-contact and a conductive p-AlGaN cladding layer. Aluminium (Al) is the only metal with reflectivity in UVC. Furthermore, when an Al-based contact on AlGaN is annealed, the Al-based electrode loses its reflectivity in UVC. Thus, the development of an effective reflective structure above EB layers is considered to be a long-term challenge, which will be described in sections 13 and 17.

To date, mass-produced DUV-LEDs on sapphire have been grown on AIN with TDD of  $2-5 \times 10^8$  /cm<sup>2</sup> [3,7] or greater. To reduce the TDD to near  $10^8$  /cm<sup>2</sup>, studies on the behavior of AIN at around and above 1400°C will also be beneficial as well as fabricating and evaluating LEDs. Also, AIN templates with a low TDD is considered to be useful for determining the possible improvement in IQE because the required TDD for fabricating DUV-LEDs are considered to be between  $10^8$  /cm<sup>2</sup> and  $10^9$  /cm<sup>2</sup> [7]. The behavior of AIN is expected to be similar to that of GaN when the growth temperature is increased.

A detailed study on the decrease in EQE during reliability tests will be meaningful for achieving devices with comparable lifetime to those of low-pressure mercury lamps. In Figure 2(b), the increased deterioration of the output of UVC-LEDs compared with that of UVB-LEDs is shown. The reliability test for 265 nm LEDs suggests two main mechanisms for the initial and subsequent decrease in the EL output. The decrease in EL output for the UVC-LEDs was much larger than that for the UVB-LEDs, although fatal device faults resulting in no light or a short-circuit did not occur.

To improve the current injection efficiency (CIE), (AI)BN is considered to be a promising material. The possibility of using (AI)BN should be investigated, which is described in section 8.

The other approaches to improving EQE are the use of r-(10-12) and m-(10-10) planes, which have the possibility of improving CIE and IQE.



**Figure 2. (a)** Full width at half maximum (FWHM) of XRCs for AIN template and AlGaN layer on AIN template and **(b)** result of the reliability test for 265 nm LEDs. In **(a)**, the improvement in XRCs using sapphire substrates with high miscut angles is shown. Recently reported FWHMs of XRCs obtained using a sputtered AIN film on sapphire followed by annealing have also been added (hollow symbols), showing the possible improvement in IQEs, which is considered to be useful for examining the possible improvement. **[(a)** Reproduced from Proc. SPIE. 9926, 9926C and Jpn. J. Appl. Phys. [7] (Copyright ©Japan Society of Applied Physics)]

#### **Concluding Remarks**

The initial EQEs of UVC-LEDs (265 nm) have been reported to be about 2/3 of those for UVB-LEDs (280-300 nm). Further reduction of the PDD and TDD is expected to increase the IQEs of UVC-LEDs. Also, the improvement in LEE by encapsulating dies was demonstrated. The utilization of light towards the p-side without increasing V<sub>f</sub> and decreasing the lifetime of UVC-LEDs is also useful. The UVC-LEDs' lifetime is expected to be much longer than 6,000 h. To date, a decrease in EL output of about 50% for 265 nm LEDs during the reliability test was observed; thus, the deterioration of UVC-LEDs must be reduced.

The production of low-pressure mercury lamps is similar to that of fluorescent room lamps. To satisfy the Minamata Convention on Mercury by replacing low-pressure mercury lamps, an EL output of 100 mW for 2–3 US dollars must be targeted, keeping the cost-feasibility including the packaging in mind.

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# 12 – UVC LEDs with emission below 250nm

Frank Mehnke<sup>1</sup>, Leo J. Schowalter<sup>2,3</sup>, Tim Wernicke<sup>4</sup>

1 Georgia Institute of Technology, School of Electrical and Computer Engineering

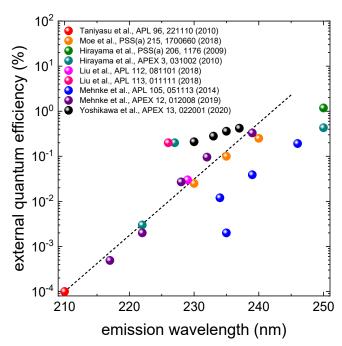
2 Crystal IS Inc.

3 Asahi Kasei Corporation

4 Technische Universität Berlin, Institute of Solid State Physics

### Status

Ultraviolet (UV) light emitting diodes (LEDs) in the UVC spectral region below 250 nm are of great interest as they, for example, enable in-situ sensing applications of diverse gases and liquids in industrial, medical, and automotive sectors. This includes the monitoring of nitrates in water,  $NO_x$  and  $SO_x$  in gas emissions, DNA purity analysis, and high-performance liquid chromatography.





**Figure 1 left.** External quantum efficiency of state-of-the-art UVC LEDs indicating the exponential reduction for emission wavelengths below 250 nm [1-9]. The wall plug efficiency of these short wavelength devices fall off even more sharply.

**Figure 1 right.** Image of Optan<sup>TM</sup> 235 which is in limited release from Crystal IS [10]. The diode is  $3.5 \times 3.5$ mm<sup>2</sup> in size with a  $0.8 \times 0.8$ mm<sup>2</sup> size chip and is packaged in standard Optan<sup>TM</sup> SMD package. It is designed to be run at 20 mA at a forward voltage of 6 V with powers binned from 50  $\mu$ W to greater than 500  $\mu$ W. These devices are rated to have a lifetime over 1,000 hours.

Currently, such short wavelength LEDs are, in design and technology, similar to LEDs emitting around 270 nm grown on c-oriented AIN templates or substrates (see Sections 2 and 3) but contain much higher aluminum mole fractions in the AlGaN/AlGaN multiple quantum wells (MQW) and n-side current spreading layers. The understanding of the n-type doping mechanisms of these high aluminum mole fraction layers enabled short wavelength LED emission down to the physical limit of AIN at 210 nm [1]. However, due to several material and heterostructure design challenges the external quantum efficiency (EQE) and wall plug efficiency (WPE) of such short emitting devices is exponentially dropping with decreasing emission wavelengths. As shown in figure 1 (left) one order of magnitude in EQE is lost for every 8 nm in reduced emission wavelength, i.e.,  $10^{-2}$  at 242 nm,  $10^{-3}$  at 234 nm,  $10^{-4}$  at 226 nm,  $10^{-5}$  at 218 nm, and  $10^{-6}$  at 210 nm [1-9]. This is attributed to a strongly

decreasing radiative recombination efficiency ( $\eta_{rad}$ ), carrier injection efficiency ( $\eta_{inj}$ ), and light extraction efficiency ( $\eta_{ext}$ ) with decreasing emission wavelength and directly translates in reduced emission powers. In addition, the WPE is further reduced at short wavelengths since making contacts to high aluminum mole fraction AlGaN is more difficult (see Section 6). Another issue with these short wavelength devices is that their lifetime is often much shorter than their longer wavelength counterparts. A successful commercial product will need to address all of these issues. Up to now there are only a few research reports on LEDs emitting below 250 nm [1-9] and only a limited-release of LEDs at 235 nm is commercially available (see figure 1 right) [10]. In order to open the market for LEDs emitting below 250 nm which are considered to become much more costeffective, small, robust, and persistent in comparison to discharge lamps both the efficiency and the emission power need to be improved. This will allow for applications such as gas sensing and enable new applications.

#### **Current and Future Challenges**

The main limiting mechanisms prohibiting higher emission powers and efficiencies of AlGaN-based UV LEDs with emission below 250 nm are related to the low  $\eta_{rad}$ ,  $\eta_{ini}$ , and the low  $\eta_{ext}$  and to high operation voltages (leading to a limitation of the driving current and the WPE) - all of which need to be addressed in future research. Firstly,  $\eta_{rad}$  within the MQW active region is strongly connected to the threading dislocation density (TDD) favoring LEDs on low TDD monocrystalline AIN substrates. However,  $\eta_{rad}$  might also be reduced by an increasing point defect incorporation (see Section 5) at these high aluminum mole fractions which is yet to be proven by measurements. Another potential issue is that  $\eta_{ini}$  into the MQWs decreases with decreasing emission wavelength as electron blocking becomes inefficient due to the smaller conduction band offset from the MQW to the AIN electron blocking layer (EBL). Furthermore, not only suppression of electron leakage but also supporting hole injection into the MQWs is a major challenge to the heterostructure design and the doping profile of the UV LEDs [7]. Finally,  $\eta_{ext}$  is reduced with decreasing wavelength due to the transition of the optical polarization of the emitted light from dominant transverse electric (TE) to dominant transverse magnetic (TM) [11, 12]. This is caused by valence band switching at an emission wavelength of around 240 nm and results in an emission pattern where photons tend to propagate parallel to the c-oriented surface. Consequently,  $\eta_{ext}$  through the substrate backside is reduced. Additional challenges arise by using monocrystalline AIN substrates as these may exhibit subbandgap absorption due to impurity incorporation during growth (see Section 3) depending on the particular growth method employed [13]. Furthermore, the poor p-type doping of high aluminum mole fraction AlGaN necessitates the usage of absorbing p-layers and p-contacts, which strongly reduces  $\eta_{ext}$ .

In order to further improve the WPE of UV LEDs with emission below 250 nm lower operation voltages are needed. In particular, a lower AlGaN:Si layer resistivity and n-contact resistivity need to be achieved for high aluminum mole fraction layers. Both exhibit physical limitations due to the large bandgap of the material. The AlGaN:Si layer resistivity is increasing with increasing aluminum mole fraction due to increasing donor ionization energy, DX-center formation, and incorporation of compensating point defects (see Section 7). The n-contact resistivity is increasing with increasing aluminum mole fraction due to a transition from Ohmic to Schottky behavior due to the low electron affinity, a very stable oxide, a high probability of forming deep levels during plasma etching, and the difficulty in forming shallow donors during the contact formation process (see Section 6).

Initial lifetime tests on sub-250 nm LEDs have suggested severely reduced lifetimes in comparison to longer wavelength UVC LEDs. In order to improve the lifetime of these device, the underlying mechanisms will need to be understood. However, an analysis of the degradation causes as well as a variation of growth and fabrication processes parameters has only recently been initiated [14].

#### Advances in Science and Technology to Meet Challenges

In the future, higher emission powers with better WPE and lifetime are desired. Longer lifetimes can be expected with future heuristic and analytical studies as well as an improved point defect control. Additionally, lower TDD may also lead to longer lifetimes. However, there is only a limited amount of data currently available comparing devices grown on AlN templates with devices grown on monocrystalline AlN substrates. A reduction of the forward voltage requires further development of highly conductive n-AlGaN layers by an improved control of the point defect density as well as low contact resistivity n- and p-electrodes especially by improved doping, novel contact metal stacks and improved surface treatments.

Future analysis of the LEDs will give insight to which portion the drop in EQE is related to  $\eta_{rad}$ ,  $\eta_{ini}$ , and  $\eta_{ext}$ , respectively, leading to a focus in the respective LED development. In order to improve  $\eta_{rad}$ , point defects resulting in nonradiative recombination in the active region can still result from the epitaxial growth conditions and further improvement is needed in this area. This measure will only be effective for growth on low TDD AIN, e.g., by growing LEDs pseudomorphically on monocrystalline AlN substrates with a TDD in the  $10^4 \text{ cm}^{-2}$  range [2]. The improvement of  $\eta_{inj}$  by a reduction of electron leakage needs to be obtained by clever heterostructure design, as the height of the EBL cannot be increased beyond the AIN bandgap and no larger bandgap semiconductor material is currently known. This may be addressed by multiple quantum barrier EBLs [4] which can provide a larger conduction band offset in comparison to AIN and allow for hole tunneling due to quantum interference. Advanced device simulations can provide the required design information, however, these calculations rely on several material parameters which are not well known and need to be further investigated. Furthermore, supporting the hole generation and injection needs to be mastered by heterostructure design and doping profile, e.g. the development of p-type polarization doping using aluminum mole fraction graded layers. It is also possible that new alloys such as AIBN (see Section 8) may offer an advantageous band alignment allowing for improved nini. In any case, clever device designs will be needed to maximize both  $\eta_{rad}$  and  $\eta_{inj}$  simultaneously since improvement in one can also be made at the expense of the other. next may be improved by shifting the optical polarization towards TE by tailoring the design of the MQW active region. The use of thin QWs and high aluminum mole fraction barriers has been demonstrated to increase TE emission at a given wavelength [11, 12]. Alternatively, growth of semi- and nonpolar MQWs might boost  $\eta_{ext}$  as the rotation of the crystal axis also leads to a rotation of the emission profiles [1], although currently the growth technology of such devices is much more immature compared to c-oriented ones. Nevertheless, the light extraction through the substrate can be improved for TE but especially for TM polarized light by one or multiple of the following approaches: a) substrate removal, b) substrate or bottom layer roughening, c) pre- or post-epitaxial patterning of the buffer layers i.e. epitaxial lateral overgrowth, patterned substrates, or photonic crystals (see Sections 13 and 14), as well as d) encapsulation. A big boost in  $\eta_{ext}$  could be gained by the use of transparent p-doped short period superlattices combined with reflective p-electrodes especially if the penalty of higher forward voltage could be avoided. Also, new approaches such as p-Si nanomembranes [6] acting as reflective contact yielded excellent results and might push  $\eta_{ext}$  and  $\eta_{ini}$  to higher levels. Another promising approach to overcome the transparency-conductivity dilemma is the implementation of p-(i)-n tunnel junctions (see Section 17). These offer the opportunity of using highly conductive and transparent AlGaN:Si layers and n-electrodes, which overcomes several but not all challenges of p-AlGaN.

#### **Concluding Remarks**

Improving the WPE, EQE and emission power of UV LEDs with emission below 250 nm cannot be performed by small single changes in the heterostructure design but requires manifold optimization and tradeoffs. These LEDs will benefit from technological improvements for longer wavelength LEDs, however, they will need additional development effort to solve the specific challenges, additionally hampered by the smaller expected market size. The exponential decrease of the EQE with decreasing emission wavelength is fundamental and will persist in spite of strong improvements for each wavelength. Additionally, applications based on these LEDs can only be realized when the lifetimes of the devices can be guaranteed.

Major breakthroughs in device performance of UV LEDs emitting below 250 nm may be expected to occur by new design concepts. For example, the development of tunnel junctions or p-Si nanomembranes can resolve the transparency-conductivity dilemma, pushing the device performance of UV LEDs with emission below 250 nm to higher levels.

#### Acknowledgements

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# **13 - Light Extraction Efficiency of UVC LEDs** Hideki Hirayama<sup>1</sup> and Yukio Kashima<sup>1,2</sup> 1 RIKEN

2 Marubun Company

#### Status

High output power 265-280 nm commercially available UVC LEDs have flip-chip (FC) geometry, and UVC light is extracted through the sapphire or AlN single crystal substrate. The performance of a UVC LED with wavelength at 280 nm with 50 mW output power at a driving forward voltage (Vf) of 5.5-7 V and a driving forward current (I<sub>f</sub>) of 350 mA exhibits an electrical power-to-light conversion efficiency (wall-plug efficiency; WPE) of about 2-3 %. The detailed breakdown of the total WPE [1] is estimated to be approximately 60% for internal quantum efficiency (IQE) including injection efficiency ( $\eta_{inj}$ ), 80% for electrical efficiency ( $\eta_{el}$ ), and about 6-8% for light extraction efficiency (LEE) [1, 2].

For a deep-UV (DUV) LED, the fact that the LEE is particularly small compared to that of a blue LED is a very big problem. The main reasons for low LEE in a DUV LED are that DUV light is heavily absorbed by the p-GaN contact layer, and total internal reflection occurs at the interface between the LED body and air. Therefore, improving LEE is the most important subject and greatly contributes to the improvement of output power and WPE of DUV LEDs.

The LEE of a current DUV LED was calculated by the combination of the ray-tracing method and the finite-difference time-domain (FDTD) method [2]. Figure 1 shows the calculation models of flip-chip (FC) DUV-LEDs with AI reflector mounted on ceramic package for (a) a usual LED and (b) an LED with lens. The assumed layer structure for a current LED on the sapphire substrate consists of a 4-µm-thick AIN, a 2 µm thick n-AlGaN buffer layer with emitting and electron blocking layers, a 1 µm thick p-GaN contact layer and Ni/Au p-type electrode with a reflectivity of 30%. The polarization of the emitted light determining the radiation profile and LEE is influenced by the aluminium contents of well and barriers and the strain state of the material and dominated by transverse electric polarization for emission wavelengths > 240 nm [3]. The reflectivity of the Al-coated reflector of the package is 90%. Using these conditions, the calculation result of LEE for a 280 nm UVC LED is 6.3% [2], which is in good agreement with the above estimated value.

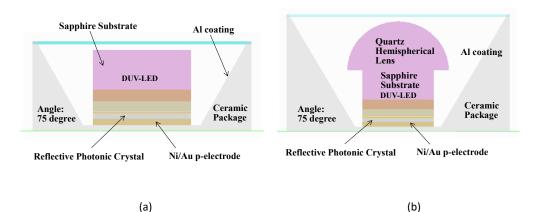


Figure 1. Calculation models of flip-chip (FC) DUV LEDs with AI reflector mounted on ceramic package for (a) a usual LED and (b) a LED with lens used for the analysis by the combination of ray-tracing and finite-difference time-domain (FDTD) methods.

#### **Current and Future Challenges**

In order to achieve a significant increase of LEE, it is effective to introduce a p-AlGaN transparent contact layer which is completely transparent to a wavelength of 270 nm instead of a p-GaN contact layer [2, 4, 5, 6]. It is also important to introduce a highly-reflective p-type electrode [4, 5], e.g., p-type silicon [7], Pd(1nm)/Al [8], Ni/Mg or rhodium [9] electrodes with reflectivity of >70%, instead of a Ni/Au electrode with reflectivity of 30% [2]. By introducing them, LEE improves more than twice and a calculated value of 13.1% was obtained [2]. However, since the hole concentration in the p-AlGaN contact layer is 1 x  $10^{14}$ cm<sup>-3</sup> or less and forward voltage (V<sub>f</sub>) rises by several volts [4], WPE cannot be greatly improved.

In order to improve LEE while maintaining WPE, thin p-GaN layers can be introduced. However, the LEE enhancement drops quickly already for thin GaN layers of 10 – 20 nm [10]. Therefore we propose forming a highly reflective photonic crystal (HR-PhC) in the p-GaN contact layer [2]. Figure 2 shows (a) a cross section of the schematic structure and (b) corresponding electric-field (E-field) mappings calculated by FDTD analysis and (c) the calculated results of LEE as a function of the distance from the quantum well (QW) emitting layer to a hole array PhC for 280 nm AlGaN UVC LEDs with HR-PhC [2]. In addition, as a comparison, we also conducted the LEE enhancement simulation with a p-GaN-free structure, i.e., using a model of p-AlGaN transparent contact layer with highly-reflective electrode [2]. In this case the radiation from the QW emitting layer does not penetrate into the PhC layer and reflected by the PhC, as confirmed from the cross-sectional E-field mappings shown in Fig. 2(b). In Fig. 2(c), we can confirm that with and without p-GaN the LEE is strongly enhanced especially when the vertical field resonant condition is satisfied. We found that the maximum increases in LEE enhanced by introducing the HR-PhC are by 2.8 and 1.8 times, respectively, for the p-GaN and p-AlGaN contact layer UVC LEDs [2]. The maximum LEE of the p-GaN and p-AlGaN contact layer 280 nm UVC LED with PhC were 18.0 and 23.8 %, respectively.

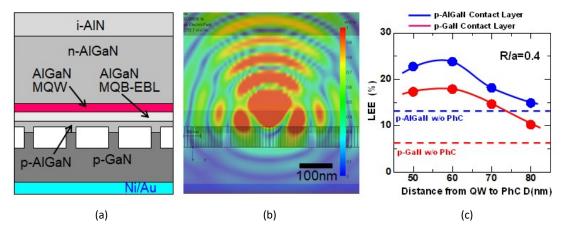


Figure 2. (a) Cross section of schematic structure and (b) corresponding electric-field (E-field) mappings calculated by FDTD analysis and (c) the calculated results of LEE as a function of the distance from the quantum well (QW) emitting layer to the PhC for 280 nm AlGaN UVC LEDs with HR-PhC

#### Advances in Science and Technology to Meet Challenges

As described above, when DUV light is emitted from the LED body into air, the DUV light is totally internally reflected at interfaces, and absorbed by a contact layer. Therefore, in order to increase the LEE of a DUV LED, it is quite important to use a low-absorption structure, e.g., by using a HR-PhC p-layer. When the absorption is low, the LEE is further enhanced by redirecting light into the light extraction cone within the semiconductor by p-side PhC [2], nanopatterned sapphire [4], structured

AlN/sapphire [3, 11]. The total internal reflection at the substrate/air interface can be alleviated by directly bonding a hemispherical lens [12], encapsulation [4, 13] or hybrid nanostructuring [14]. In the model shown in Fig. 1(b), a quartz hemispherical lens is assumed to be directly bonded to the back surface of the sapphire substrate. The calculation results of LEEs of the 280 nm UVC LEDs with quartz hemispherical lenses are 13.1 and 28.3 %, respectively, for the p-GaN and p-AlGaN contact layer cases which is a strong enhancement compared to 6.3% and 13.1%, respectively. Indeed, a record high EQE of 20% was demonstrated for a 275 nm LED with transparent p-AlGaN layer, reflective Rh contact and hemispherical encapsulation [4]. The main advancements which will lead to a widespread implementation of these light extraction techniques are reduction of the additional Voltage by improving the conductivity of p-AlGaN (see chapter 7), demonstrating the reliability of such LEDs (methods described in section 16) and implementing low cost fabrication techniques.

#### **Concluding Remarks**

To date, the efficiency and the output-power of an AlGaN DUV-LEDs are significantly limited by a low LEE, which is reduced by a strong absorption of a p-GaN contact layer. The LEE would be dramatically increased in near future by introducing an absorption-free contact layer, a highly-reflective p-type electrode, a reflective PhC fabricated on p-contact layer, by bonding lenses, by fabricating a vertical LED structure, and/or by combining these effects.

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# 14 – Nanostructuring for UV Emitters Philip Shields<sup>1</sup>, Robert Martin<sup>2</sup> 1 University of Bath 2 University of Strathclyde

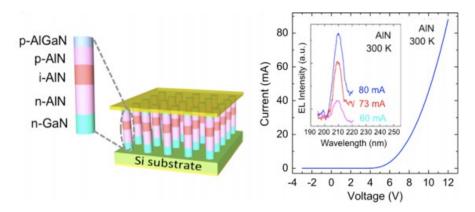
#### Status

Reports of using nanostructures in UV-emitting devices have increased sharply over the past 5 years as they can potentially solve the existing obstacles for obtaining efficient nitride-based UV-LEDs [1-8]. They follow a longer and more substantial effort in the use of nanostructures to aid visible LEDs. We describe below some examples of how existing uses of nanostructures have overcome the following key issues: the influence of defects on the internal quantum efficiency (IQE); the poor light extraction efficiency (LEE) in planar UV-devices due to the emission of predominantly TM-polarised light from AIN-rich AlGaN and the high absorption in conventionally-used layers; and the difficulty in achieving efficient p-doping in AlGaN materials.

Firstly, annealing AIN nanostuctures at high temperatures has been shown to improve the IQE in subsequent c-plane QWs by dramatically reducing the number of edge dislocations [2]. Laterally growing from these or other nano-patterned templates, such as sapphire or silicon, alongside the use of maskless techniques, permits the use of traditional epitaxial lateral overgrowth defect reduction techniques whilst coping with the poor mobility of AI species in the growth environment that traditionally leads to delayed coalescence and poor quality surfaces [3,4].

Secondly, the presence of scattering interfaces in nano-ELOG structures also improves the LEE, which is especially important for AIN-rich AlGaN materials for which the valence band ordering enhances TM-polarisation [5]. The nanoscale features can also promote strain relaxation, allowing strain engineering of the valence band to influence the dominant emission polarisation as well as a potential improvement of the crystal quality

Thirdly, the ability to manipulate strain and polarization in nanostructures has been employed to significantly increase the incorporation of the Mg p-dopant and enhance hole mobilities compared with Al(Ga)N planar films. For example, Fig. 1 shows nitrogen polar AlN nanowires grown directly on Si, which demonstrated increases in IQE and reductions in turn on voltage [6]. Highly conductive p-AlGaN can avoid the use of absorbing p-GaN, as in conventional UV LEDs, and increase both the electrical efficiency and reduce optical losses.



**Fig. 1:** Electrical and electroluminescence results from an AlN nanowire LED. Reprinted by permission from Springer Nature, Scientific Reports, Ref. [6], Copyright 2015.

Finally, the nanostructuring of additional materials such as AI can lead to enhanced UV emission through the coupling of surface plasmons with the emitting dipoles [7]. Photonic crystal effects have also been incorporated into the nanopatterning of AIN surfaces to increase LEE in deep-UV LEDs, using nanoimprint lithography [8].

#### **Current and Future Challenges**

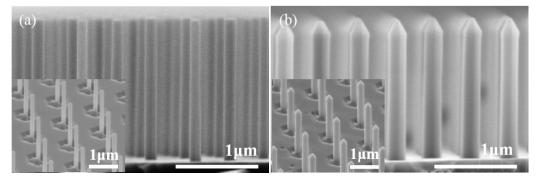
These examples show some different ways that nanostructuring can improve UV-emitting devices. However, simultaneously improving the IQE, LEE and electrical efficiency without introducing further detrimental effects remains a challenge. Following work in the visible regime, an ultimate goal might be the use of a regular and organised array of dislocation-free nanostructures as a scaffold for highaspect-ratio core-shell nanorods arranged to ensure maximum light scattering from both TE and TM emission. Enhanced IQE due to the high crystal quality would be combined with strain-relaxed material for improved dopant incorporation.

Achieving this requires solving a number of challenges, with the bottom-up growth of AlGaN materials difficult due to the low mobility of Al species. Core-shell structures have traditionally required MOVPE growth, whereas most reports of AlGaN nanostructure growth have relied on MBE, using UV-absorbing GaN pedestals as nucleation seeds. Controlling the AlGaN alloy composition and dopant incorporation across the various crystal facets is an additional challenge. Furthermore, most lighting applications require a large number of parallel-driven nanorods and need a 3D contacting architecture. Any filling materials need to be non-absorbing, which becomes more challenging for emission further into the UV, as does achieving sufficient heat extraction and conductivity. Whilst the maturity of UV LEDs lags behind visible LEDs, they are sufficiently well-developed to be commercially available. 3D nanoscale LEDs will need to demonstrate unequivocally that their advantages can justify their more complex creation to have a chance to compete.

Alternatively, nanostructuring could be directed simply at improving IQE or LEE in planar devices. However, recovering a planar surface from nanostructured material requires carefully controlled coalescence to ensure a net reduction of extended defects and complete surface recovery to stepflow growth since AlGaN QWs are sensitive to any underlying roughness. A further challenge will be to simultaneously optimise both IQE and LEE as the ideal pattern for one is likely to compromise the other. One way of disentangling the effects is to improve LEE by employing nanostructuring after growing the active layers. Again, the benefits must outweigh the increased complexity of fabrication.

#### Advances in Science and Technology to Meet Challenges

To fully realise the potential benefits of nanostructuring for Al(Ga)N UV-emitters, various advances are needed. These include further steps towards achieving defect-free, bottom-up growth of AlGaN nanowires with controlled size, spacing and height, without relying on GaN seeding. As well as improving MBE approaches in this respect, a key advance will be expansion out to MOVPE, which will also allow easier scale-up. Further work should build on the use of polarisation and strain relaxation to enhance doping. Hybrid approaches may be significant, for example building on the idea of the lateral re-growth of AlGaN on etched AlN nanorods reported by Coulon et al. ([9], Fig. 2). In this case, advances are required to increase the parameter space to grow high-quality non-polar material and improve precursor efficiency. Device designs and process flows optimised for performance and manufacture are required taking into account the specific vulnerabilities of electrical leakage and thermal management.



**Fig. 2:** Cross-section SEM images of AIN nanorod arrays (a) after etching and (b) after subsequent MOVPE regrowth. Reprinted by permission from ACS Appl. Mater. Interfaces, Ref. [9], Copyright 2018.

Where nanopatterned layers are to be used to create planar templates with reduced dislocation densities or for device structures, it is important to be able to balance dislocation bending with dislocation creation during coalescence of the neighbouring growth fronts. A deep understanding of the dislocation reduction process is required to achieve the best material quality for optimal LEE, additionally without having to resort to long growth runs.

Exploiting gratings or photonic crystal effects could help master the specific challenges in UV emitters but require the development of reliable techniques to manufacture the small nanostructure periodicities and the low edge roughnesses needed for the ultrashort wavelengths. Use of electron beam lithography is unlikely to be scalable, so large-area nanopatterning techniques such as nanoimprint and displacement Talbot lithography will likely become significant [10,11].

Use of flip-chip processing to minimise optical absorption and improve thermal efficiency can be beneficially combined with use of nanopatterned material or arrays of nanowires grown on large area Si for commercial scale-up. Other promising directions are extending nano-patterning to substrates beyond sapphire or growing Al(Ga)N on layered transition metal dichalcogenides, such as MoS<sub>2</sub>, on sapphire with a view to overcoming challenges associated with lattice and thermal expansion mismatches [1].

Effort should be applied to further exploit plasmonic effects as they have been shown to be effective in enhancing light emission. Advances in modelling and fabrication are required to optimise the positions of metal nanoparticles to enhance TE-emission and LEE, even to positions within the active regions themselves. Whilst Al is typically used in the UV, advantage could be obtained by using other non-oxidising materials such as Rh.

There will also be challenges to overcome for advanced characterisation methods, such as mapping of dopant incorporation in core-shell nanorods, perhaps using optical signatures to aid their rapid development. Developments with techniques such as cathodoluminesence and atom probe tomography for nanopatterned UV materials will accelerate progress.

#### **Concluding Remarks**

There are a number of reports that have signalled the promise of nanostructuring to potentially solve longstanding issues in planar UV emitters. Whilst the field is in an early stage, it has accelerated at a phenomenal rate and it will not be a surprise if many of the challenges are surmounted in the next few years. Particular game changers would be: the successful bottom-up growth of regular arrays of non-absorbing and dislocation-free AlGaN nanostructures, a detailed

understanding of the effect of polarity and strain on enhanced Mg incorporation for different AlGaN compositions, the convergence on an optimised 3D device architecture, and a routine capability to pattern materials with sub-100 nm resolution. Following that, the question will be whether nanostructured materials can be made in the multi-wafer growth reactors for commercial manufacture with sufficient yield to challenge conventional planar devices. This may be the toughest challenge of them all.

#### Acknowledgements

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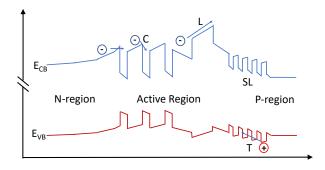
# 15 – Simulation of UV-Light Emitting Diodes and Lasers Bernd Witzigmann<sup>1</sup>, Friedhard Römer<sup>1</sup> and Yuh-Renn Wu<sup>2</sup> 1 University of Kassel 2 National Taiwan University

#### Status

Optoelectronic device simulation is a multi-scale, multi-physics task. Optical, electronic, and thermal properties, and their interaction govern the characteristics. Nano-scale regions require ideally atomistic methods, but electrical contacts, thermal reservoirs or optical resonators are often on the macroscopic scale. For lasers and LEDs in the III-nitride system, numerical simulations are routinely done nowadays as a guideline to design the device specifications, which starts from the epitaxial layers for optimum waveguiding and carrier injection (see Fig. 1), and carrier-photon interaction, up to the chip processing for placement of contacts, or light outcouplers (such as facet coatings or antireflection structures). A comprehensive, but probably still incomplete overview of the large variety of numerical codes can be found in [1], and the main approaches are as follows. The most widely used models are based on the drift-diffusion currents in combination with continuity equations for electrons and holes, and the electrostatic potential being evaluated by a Poisson equation. This coupled nonlinear system can discretized in up to 3 dimensions, and solved efficiently with a Newton type iterative method. Besides the current-voltage characteristics, one can study local current densities, recombination rates or carrier mobilities. For quantized regions, a single particle Schrödinger equation can be coupled to this system, also with decent convergence behaviour. Recent works on solving Schrödinger equations with the localization landscape theory to obtain the effective quantum potential provide an efficient tool to combine the semi-classical theory with quantum effects [9]. For non-equilibrium carrier transport, the Monte-Carlo method has been applied to LEDs, solving the semi-classical Boltzmann equation. While including non-equilibrium transport, it requires massive computational resources, and lacks a consistent inclusion of quantum effects. A full non-equilibrium quantum mechanical simulation has been realized with the nonequilibrium Greens function method, however again with large computational cost, and no model for non-radiative recombination [7].

The optical part requires numerical methods for solving Maxwells equations, such as finite element, finite difference methods or ray tracing approaches. Here, the challenge is to include the interaction between the electromagnetics and the electronics, which occurs via stimulated and spontaneous emission, or absorption.

For application to device design and analysis, active research on the models, equation frameworks and material parameters goes along with the advancement of material synthesis and technology capabilities. This is currently the case for III-nitride based light lasers and LEDs emitting at ultraviolet wavelengths. For their InGaN based counterparts, simulations, in combination with numerous experiments, have contributed significantly to pushing the electro-optical efficiencies and wavelength coverage in the visible close to optimum values. As example, numerical simulations of the quantum confined Stark effect have clarified the impact of polarization charges at interfaces. Also, the characteristics of devices on semi- and non-polar crystal orientations have been studied in great detail by microscopic simulations. With this knowledge, the potential performance gains compared to polar devices could be studied. The efficiency droop has been studied using various models, which have helped identifying the role of leakage currents and Auger recombination [8].



**Figure 1.** Schematic of conduction and valence band in a UV-LED. Carrier injection is influenced by tunneling (T), capture (C) and leakage (L). On the p-side, a superlattice (SL) supports dopant activation in AlGaN alloys. The active region consists of 3 quantum wells, alloy fluctuations will modulate the band edges, but are not shown for clarity here.

#### **Current and Future Challenges**

Experimental data for III-nitride based UV LEDs currently exhibit low quantum efficiencies, high operation voltages and low optical extraction efficiencies. UV light emitters in the III-nitride system differ from their counterparts emitting in the visible range mostly by the extensive use of AlGaN and AIN layers. This introduces alloy fluctuations in the entire device, and a different set of impurities and defects. The peculiarities of AlGaN with high aluminum mole fractions have not been studied in detail, and the existing models calculating the laser or LED performance have not been tested to large extent. The major challenge for the simulation models will be to quantitatively explain the efficiency characteristics, and come up with proposals for high performance laser and LED structures. For strain, continuum models might not be able to represent the local strain due to alloy fluctuations properly. Impurity modeling in III-nitrides is another challenging field, as these materials show high luminescence in the presence of defects. Most of the AlGaN related studies have been done for high electron mobility transistors, and while models for bulk and interface impurities are available in drift-diffusion based simulations, it is not clear how valid they are for UV light emitters. For silicon devices, the simulation of the process technology has resulted in a more accurate representation of the structural properties of the devices as input to the device simulation [6]. LED and laser simulation in the III-nitride system would benefit greatly from a process simulation, but is far from being established.

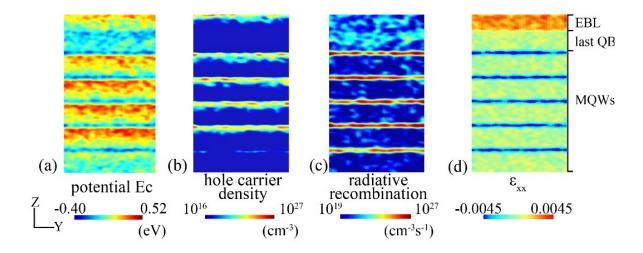


Figure 2. (a) is the calculated fluctuated potential Ec of UVC-LEDs by considering the random alloy fluctuation in EBL, QBs, QWs, and even the injection layer. (b) and (c) are the hole density and radiative recombination, which is clearly affected by random alloy fluctuation. (d) shows the calculated strain  $\epsilon_{xx}$  in the QW. Due to the fluctuated composition and local strain relaxation, the compressive strain of QW has been relaxed at the interface, which may not be good for TE emission [10].

#### Advances in Science and Technology to Meet Challenges

AlGaN crystallizes in the anisotropic wurtzite lattice and is a direct semiconductor for all compositions. Calibrated sets of parameters for modelling the electronic band structure and the influence of strain with the kp-Schrödinger method exist [3]. The luminescence polarization changes from perpendicular to parallel to the c-lattice direction with increasing aluminium content. In thin film structures grown along the c-lattice direction TM-polarized radiation dominates above 5.2eV. AlGaN exhibits spontaneous and piezoelectric polarization in the c-direction leading to sheet charges at hetero interfaces. The growth conditions of AlGaN material lead to alloy fluctuations. Ternary AlGaN layers build not only the active region, but also the carrier injection regions. The influence of local fluctuation on UVCLEDs has been analyzed in Ref. [10] with 3D Poisson, drift-diffusion and localized landscape model [9] to account the effect of quantum potential. Fig. 2 shows the cross section view of band profile, hole density, radiative recombination, and strain distribution. The preliminary results show that IQE is strongly affected by disorder as the blue LEDs. The poor hole injection due to low activated dopant and weaker blocking ability of the EBL caused by the fluctuating potentials may lead to strong droop effect. Furthermore, as shown in Fig. 2(b) and (2d), Carriers are less confined in the QW and extend into the QBs due to the alloy potential fluctuations. The hole wave function extension into the QBs will enhances TM emission as shown from a k.p. simulation of wave-functions admixture, which should then lead to poor light extraction. These preliminary results show that future simulation model should takes this effect into account for the whole structure, which leads an even higher computation burden if a full atomistic model is applied. Carrier injection in principle suffers from the low hole mobility in p-doped AlGaN alloys, and the high ionization energy in excess of 0.5eV for Mg acceptor atoms. Superlattices made of alternating thin layers with different band gap have been devised for enhancing the hole density and injection. Carriers in the superlattice are subject to quantization and minibands evolve due to the periodicity. The minibands and the polarization induced potential reduce the ionization energy and thus increase the hole density. The modelling and design of superlattices requires a quantum mechanical

approach. Miniband dispersion relations have been calculated with the Kronig-Penney model [4]. The non-equilibrium Green's function method enables superlattice transport modelling.

#### **Concluding Remarks**

While the basic equations for carrier transport and electromagnetics have been established, accurate material parameters and the impact of impurities still need improvement for UV light emitters. Parameters can be determined either by experiment or by ab initio simulations. A close collaboration between the communities in materials characterization and materials theory is beneficial in order to verify parameters and establish a general understanding of the device physics. Ab initio simulations for determining material and model parameters have given many valuable data for the device simulation community, such as Auger coefficients for GaN [2], or dopant activation in AlGaN. In the future, advanced methods beyond pure bulk computations will help understanding interfaces and quantum wells. In the field of materials characterization, techniques such as atomic probe tomography can resolve the structural and recombination properties of the device on an atomic level [5]. This will lead to much improved understanding of the optoelectronic properties.

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### 16 – Reliability of UV LEDs

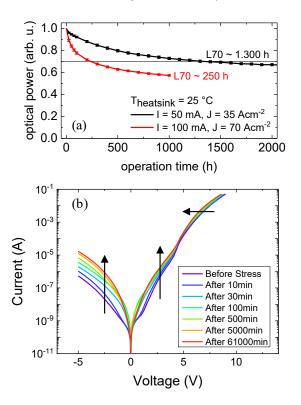
Carlo De Santi<sup>1</sup>, Matteo Meneghini<sup>1</sup>, Johannes Glaab<sup>2</sup>, Jan Ruschel<sup>2</sup>, Sven Einfeldt<sup>2</sup>

1 University of Padova

2 Ferdinand-Braun-Institut, Leibniz-Institut für Höchstfrequenztechnik

#### Status

In order to enable a wide market acceptance of UV LEDs, these devices need to be stable over typically several thousand hours of operation. For the time being, the lifetime of UV LEDs cannot yet keep up with their visible counterparts. Nevertheless UVB and near UVC LEDs with L70 lifetimes (time until 70% of the initial optical power is reached) in the order of 10.000 h have been demonstrated on commercially available devices. On the other hand the lifetime of state-of-the-art deep UVC LEDs with emission below 250 nm is still short [1]. Usually, during constant current operation, several degradation effects can be observed in parallel, including (i) a reduction in optical power, which is rapid within the first hundred hours and slower for longer operation times (see Fig. 1a); (ii) a variation of the operating voltage (see Fig. 1b for voltages > 4 V); (iii) an increase in the leakage current (see Fig. 1b for voltages < 4 V); (iv) changes in the spectral purity [2]; (v) catastrophic failure (devices suddenly stop working) [3]. Furthermore, it was found that typically the degradation is accelerated by the operation current [4, 5] (Fig. 1a) and temperature [4].



**Figure 1.** (a) Optical power (normalized to the value at 0 h) as a function of operation time of UVB LEDs operated at 50 mA and 100 mA, respectively, and 25 °C. The L70 values indicate the time at which the optical power of the LEDs reached 70 % of its initial value. (b) Current-voltage characteristics of UVB LEDs measured after different operation times over 1.000 h of operation at 100 mA and 20 °C.

So far, the main focus of most of the degradation studies has been on semiconductor issues of the UV LED chip. Here, the discussed possible physical causes of the degradation include:

(i) The generation of defects driven by current and temperature; such defects form deep

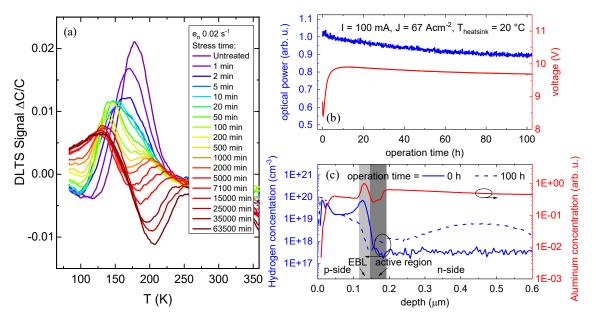
levels and behave as centers for non-radiative Shockley-Read-Hall recombination. For example, the degradation was correlated to an increased density of midgap states in 308 nm UVB LEDs [6].

- (ii) The migration of defects through the heterostructure [7]. Some authors reported on the migration of hydrogen [8] (see Fig. 2b and c) or aluminum [9]. Dislocations or V-pits and the electric field were proposed to enhance the migration.
- (iii) The pile-up of charges at heterointerfaces due to the generation and/or diffusion of defects that can affect the injection or escape efficiency of charge carriers [10].
- (iv) Changes in the charge distribution within the doped regions due to activation/compensation processes [1, 11].

#### **Current and Future Challenges**

To narrow down critical issues in the fabrication chain and to localize the degradation effects in the device are important aspects of current and future research on the degradation of UV LEDs. Therefore, empirical studies of the impact of heterostructure design (e.g. design of the electron blocking layer), epitaxial growth (e.g. variation of substrates), chip design (e.g. different insulator materials) and of chip fabrication and mounting on degradation are ongoing with the goal of enhancing device lifetimes.

Another essential aspect is to analyze the degradation mechanisms and stress-induced changes in the devices in order to understand the physics behind. Here the identification and localization of defects causing deep levels in the semiconductor are the key issue (See Section 5). Important techniques to study the density and the energy of these levels are the capacitance and optical deep level transient spectroscopy (C-DLTS and O-DLTS, see Fig. 2a), deep level optical spectroscopy, photocurrent spectroscopy and analysis of the low frequency noise. The structure of emerging defects, their changes and motion can be studied by techniques as transmission electron microscopy, secondary ion mass spectrometry (SIMS), cathodoluminescence, or electron beam induced current.



**Figure 2**. (a) Variation in C-DLTS signal in a 308 nm LED stressed at 100 mA. (b) Optical power and drive voltage as a function of operation time of one InAlGaN-based UVB LED operated at 100 mA and 20 °C. (c) Corresponding depth profiles of the LEDs shown in b) of the H-concentration (blue curves) at 0 h and 100 h of operation as determined by SIMS. The Al concentration profile (red curve, normalized to the max. value) is for the calibration of the depth.

Considering the future use of UV LEDs, new aspects might be important. For example for

applications which require the LED wavelength to stay in a certain spectral window, operationinduced parasitic emission bands lowering the spectral purity might be unacceptable, even if other electro-optical parameters stay constant. Also, it is worth studying approaches to make the devices less sensitive to temperature or to switching. Certain applications may require studying the influence of the environmental gas or of ionising radiation. For the use of UV LEDs in a harsh environment, packaging issues will be critical. Therefore, the impact of the UV radiation and the large amount of generated heat on the stability of encapsulation materials and optical reflectors used in the package have to be considered. Recently, a photon-induced generation of point defects that takes place even without any applied bias was reported for GaN-based devices [12] and could turn out to be relevant also in AlGaN-based UV LEDs.

Future UV LEDs may make use of new device concepts such as tunnel junctions (See Section 17) or polarization p-type doping which could result in unforeseen reliability issues.

#### Advances in Science and Technology to Meet Challenges

To improve device reliability, advances in clarifying the nature of the relevant defects and in understanding how they form and change during device operation are needed. Despite first studies showing that the defects form deep level acceptors [6], only little is known about the defect structure, i.e. whether it is an isolated vacancy or impurity, a complex of point defects and whether dislocations are involved. The charge state and the density of the defects are unknown as well. Therefore, spatially resolved measurement techniques, particularly luminescence studies, are needed at different stages of the degradation process. As the generation or activation of defects requires the supply of energy [8], hot carriers could be involved [5]. The generation of hot carriers in blue LEDs has already been verified and attributed to Auger recombination by Iveland *et al.* [12]. A similar study on UV LEDs is missing.

Moreover, additional work is needed to reduce the initial concentration of point defects and to suppress their migration, which could for instance be achieved by specific epitaxy techniques or conditions and by optimizing post-growth processes such as the activation of p-type conductivity. The introduction of additional layers into the heterostructure acting as migration barriers could be of interest as well.

Degradation could possibly be reduced through specific device concepts. For example the temperature and current density could be lowered and made more uniform by increasing the size of the active area. Additionally, an enhanced hole injection efficiency could reduce the excess of electrons with respect to holes in the active region, which may trigger the formation of defects if Auger recombination is involved.

Finally, it is mandatory to develop and improve the mathematical models that describe the performance of UV LEDs [1, 5, 14]. For example, De Santi *et al.* proposed a simple model for the temperature dependence of the emission spectrum of UVB LEDs [14]. Such models can be used for the analysis of the variation in performance at increasing stress time, and can eventually be extended to predict the LED lifetimes under different operation conditions.

#### **Concluding Remarks**

Before LEDs become competitive in the market of UV light sources, the origin of their degradation should be understood to improve their lifetime. The reduction in optical power in current UV LEDs is most likely dominated by a generation of deep-level acceptor states in the active region of the semiconductor heterostructure, which form non-radiative recombination centers. This process may be connected with a migration of point defects within the junction. Further studies are needed to

understand the involved physical mechanisms. Both an empirical optimization of the device reliability and an analysis of the involved physical processes are required.

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# **17 – Tunnel junction-based UV LEDs**Siddharth Rajan, Yuewei ZhangThe Ohio State University

#### Status

The major factors limiting UV LED efficiency are the poor light extraction efficiency and low carrier injection efficiency. Both of them are closely related to the low p-type AlGaN conductivity and high p-type contact resistance. Heavily doped p-GaN layers typically grown on top of p-AlGaN to enable hole injection lead to significant absorption losses, and poor light extraction efficiency. Alternate strategies involving p-type metal contacts to p-AlGaN [1] lead to significant improvement in the light extraction efficiency, but cause high electrical losses due to substantial increase in the operation voltage. Therefore, simultaneous improvement in the electrical efficiency and light extraction efficiency is intrinsically limited for the conventional device structures, and has remained a major challenge for the UV LED community. Tunnel-injected UV LED structures can enable non-equilibrium hole injection through transparent AlGaN layers, and enable better light extraction, thus simultaneously enabling low absorption and electrical loss.

Realizing efficient tunnel junctions (TJs) becomes more challenging as the bandgap of the semiconductor is increased. Recent work [2, 3, 4] has shown that efficient interband tunneling hole injection is feasible using tunnel junction structures where the spontaneous and piezoelectric polarization charges are used to create extreme electric fields [5]. These polarization sheet charges can create band-bending across nanometer-scale lengths, thereby reducing the tunneling barrier and increasing tunneling probability (Fig. 1(a)). This, and other innovations in semiconductor heterostructure design [6] have enabled high tunneling conductivity to be achieved for bandgaps as high as 5.4 eV, with demonstrations of ultra violet LEDs for wavelengths as low as 257 nm (Fig. 1(b)) in planar films [7], and 242 nm in nanowires [8]. The availability of such tunnel junctions is especially important for shorter wavelength (high Al-content films) since the thermally activated hole concentration decreases dramatically with increasing Al content in the p-AlGaN layer.

Interband tunnel junctions have the potential to enable ultra violet LEDs with power density and efficiency greatly exceeding state-of-art devices today. They could enable more efficient light extraction strategies for AlGaN-based UV LEDs by taking advantage of the transparent top and bottom n-AlGaN contacts, and the use of UV-reflective n-type contacts. Furthermore, the low sheet resistance of n-AlGaN could enable new light extraction strategies, similar to those used for visible LEDs. Finally, electrically injected lasers have been a long-standing challenge [9]. While lasing at wavelength as short as 271.8 nm under pulsed current was recently demonstrated [10], the devices showed high threshold current and poor emission power due to the deficit in hole injection. There are several potential applications for tunnel junctions in such devices, since they could enable efficient hole injection as well as a much thinner p-AlGaN layer, thereby lowering p-type conduction resistance, and reducing free carrier losses. However, each of these areas require a concerted effort to improve understanding of growth physics, transport, doping, and optical characteristics.

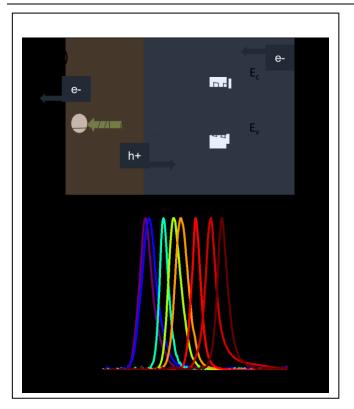


Figure 1. (a) Schematic energy band diagram of a tunnel-injected UV LED structure under forward bias. (b) Summary of the emission spectra achieved in tunnel-injected UV LEDs.

#### **Current and Future Challenges**

Even though tunnelling hole injection has been demonstrated to be feasible, several challenges remain to be addressed before we can realize the potential of tunnel-injected UV LEDs. While metalorganic vapour phase epitaxy (MOVPE) is the most technologically mature growth technique for UV LEDs, it brings some unique challenges for the growth and fabrication of tunnel-injected UV LED structures. The existence of hydrogen atoms in the MOVPE chamber leads to passivation of the Mg acceptors. The n-type top contact layer in the tunnel-injected UV LED structure impedes hydrogen diffusion, making it difficult to achieve Mg activation. To circumvent this problem, lateral Mg activation from etched sidewalls was developed for MOVPE-grown tunnel-injected blue LEDs [11]. However, these tunnel junction based visible LEDs have shown higher turn-on voltage and differential resistance when compared to standard LEDs with metal ohmic contacts.

Another challenge comes with the growth of the tunnel junction layer using MOVPE. The large difference in the growth temperature between AlGaN and the thin InGaN layer adopted in the tunnel junction structure could lead to decomposition and intermixing of the InGaN layer, resulting in poor tunnel junction performance. In comparison, molecular beam epitaxy (MBE) growth does not require p-AlGaN activation after growth, and provides much lower growth temperature difference between AlGaN and InGaN layers. However, recent work [12] demonstrates promising results from all-MOVPE based LEDs that could suggest a future pathway for UV emitters based on tunnel injection.

Low tunnel junction resistance values below  $2 \times 10^{-3} \Omega$  cm<sup>2</sup> have been achieved for ultra-wide bandgap AlGaN with Al content up to 70%. However, a substantial increase in the excess voltage drop is observed as the AlGaN bandgap increases in the tunnel junction structure as shown in Fig. 2. Even though polarization engineering has been utilized to shrink the tunnel barrier through the insertion of an ultra-thin InGaN layer between p+ and n+-AlGaN layers, the large conduction band and valence band offsets at the hetero-interfaces lead to extended depletion barriers for interband tunnelling [6]. Increasing the Al content in the AlGaN layers leads to further increase in depletion barriers and reduced tunneling probability. Therefore, a detailed understanding of the mechanisms in the tunnel junction layer is necessary to reduce the excess voltage drop. Furthermore, although tunnelling injection potentially reduces internal light absorption loss, there remains a severe limitation on the light extraction efficiency due to the lack of good encapsulants and reflective metals in the UV wavelength range.

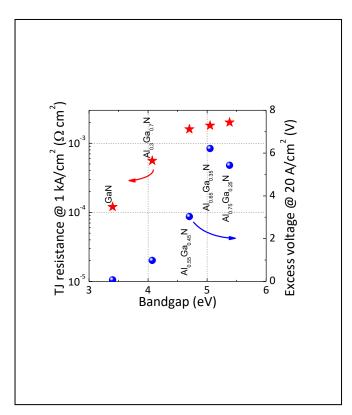


Figure 2. Summary of AlGaN tunnel junction (TJ) resistance and the excess voltage drop measured for tunnel-injected UV LEDs.

#### Advances in Science and Technology to Meet Challenges

To fully exploit the advantages of tunnel-injected UV LEDs, fundamental scientific and engineering studies related to growth science, carrier transport, and optics are needed. One promising solution could be a hybrid growth method employing both MOVPE and MBE growth techniques. The epitaxial nucleation, thick n-AlGaN bottom contact layer and the active region can be grown using MOVPE, while growth of the tunnel junction layer and the n-AlGaN top contact layer can be done using MBE to exploit the high doping concentrations and the sharp heterointerfaces. This method would combine the high internal quantum efficiency of MOVPE-grown active regions with high electrical injection efficiency of the tunnel junctions grown by MBE. Alternately, further investigation of MOVPE-based tunnel junction design and growth, and study of p-AlGaN activation could lead to solutions that take advantage of the relative maturity and performance of MOVPE growth.

In either case, it is critical to develop a better understanding of interband and intra-band tunnelling phenomena so that future designs can minimize voltage and resistance loss in tunnel-injected UV LEDs. For example, the introduction of high density polarization charges to reduce the depletion barrier was shown to reduce resistance due to conduction and valence band mismatch in p-

AlGaN/InGaN/n-AlGaN tunnel junction structures [6]. Background compensating charges, such as unintentional impurities or native defects, were found to be responsible for the extended depletion barriers in the tunnel junction layer. Therefore, growth optimization to reduce the compensating charge density could be critical for highly efficient interband tunnel junctions. Further improvement in the tunneling probability could be achieved by exploring the doping limit in AlGaN layers and by shrinking the interband tunnelling barrier.

The tunnel-injected UV LED structure minimizes the internal light absorption loss, making it possible to substantially improve the light extraction efficiency. Due to the high photon energy emitted from the UV LEDs, it is challenging to find transparent encapsulation materials with long operation lifetime. The tunnel-injected UV LED structure does provide flexibility to explore novel designs for light extraction and device packaging that are free from the widely used packaging materials. For example, tunnel junctions could allow for introduction of high reflectivity metal contact layers and specifically designed mesa structures to enable enhanced light extraction efficiency for both TE and TM polarized light.

#### **Concluding Remarks**

Tunnel-injected UV LEDs employing interband tunnel junctions for non-equilibrium hole injection hold great promise for high performance UV emitters. AlGaN tunnel junctions have been demonstrated for emission in AlGaN-based LEDs in wavelengths down to the UVC spectrum. While tunnel-injected UV LEDs emitting across the UV spectrum have been demonstrated, several opportunities still exist for improving the performance of the tunnel junctions, understanding the fundamental semiconductor physics in these structures, and for enhancing light extraction efficiency. A better understanding of these aspects could enable disruptive advances in solid state ultraviolet sources, and have significant impact on downstream applications.

#### Acknowledgements

We acknowledge funding from the National Science Foundation (Nos. ECCS-1408416 and PFI AIR-TT 1640700) and the OSU TCO Accelerator Award.

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# **18 – E-Beam Pumped Emitters**

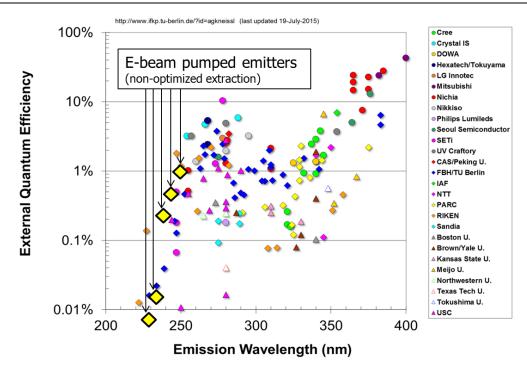
Thomas Wunderer Palo Alto Research Center, Inc., Palo Alto, CA

## Status

As we have seen in the previous sections, the challenges of realizing useful p-type doping in the ultra-wide bandgap AlGaN materials limit the performance of UV emitters. Using high-energy electron beam excitation as an alternative pump strategy for the generation of electron-hole pairs can circumvent many of the issues encountered in conventional p-n-junction configurations. When using e-beam excitation, electron-hole pairs are created through a sequence of scattering events within the semiconductor material that obviate the need for p-type doping. Thus, resistive electrical losses from contact and sheet resistances become negligible. Also, efficient and homogeneous carrier injection can be achieved even with a wide active zone. There is no need for an electron blocking layer or other means to address the asymmetry in the electrical properties between electrons and holes that influence the effectiveness of carrier injection. Furthermore, high light extraction efficiency can be realized, with low optical absorption materials within the entire device heterostructure, because no low bandgap materials are needed to improve the electrical properties. All of these aspects make electron beam excitation an attractive platform technology, in particular for emitters in the mid-UV where p-type doping otherwise limits device performance.

An e-beam pumped AlGaN chip producing spontaneous emission in the mid-UV was reported by Oto et al. in 2010 [1]. Although their claimed optical output power efficiency may have been obscured by X-ray radiation, the proof-of-concept was clearly demonstrated. Other teams followed showing pure optical output powers as high as 230 mW at  $\lambda$ =246 nm [2,3] and demonstrated the feasibility of implementing a compact form factor [4]. Record high optical intra-chip powers of about 10 Watts at 246 nm were shown with liquid-nitrogen cooled chips [5]. E-beam pumped spontaneous emitter sources compare favourably with state-of-the art efficiencies from conventional UV-LEDs (see Fig. 1), and can deliver high optical output powers at the same time.

Using e-beam excitation for lasers is also particularly intriguing for the same reasons discussed above. Despite the fact that laser emission in the deep-UV has still to be demonstrated, longer-wavelength III-N lasers reveal the great benefits of this compelling platform technology (Fig. 2) [6,7,8,9].



**Figure 1.** Comparison of e-beam pumped spontaneous emission device efficiencies with state-of-the art conventional LED performance. Collection of LED performance from M. Kneissl at http://www.ifkp.tu-berlin.de/?id=agkneissl.

#### **Current and Future Challenges**

E-beam pumped light emitters based on semiconductor materials share some common aspects with conventional LED manufacturing, including epitaxial growth and device fabrication processes. Various device design features follow identical optimization schemes as known for LED development. These involve improving the well-known efficiency contributions through optimization of material quality, heterostructure design and device architecture. The key difference to consider between e-beam pumped light emitters and p-/n-junction devices is how electrons and holes are created and injected into the active zone for photon generation.

E-beam pumping involves the bombardment of the target sample with high energy electrons (i.e., 5 – 30 keV) through which electron-hole pairs are generated in an extended volume that is dependent on the pump spot diameter and the chosen electron energy as well as the target materials. This process has an intrinsic energy efficiency limit of about 1/3. The carriers subsequently diffuse into the QWs where they recombine with the generation of photons. Proper electrical grounding removes net excess charge. Excess heating and possible hazards from soft x-ray generation require careful design and mounting schemes.

The operation of the electron beam requires a vacuum enclosure, dictating materials and manufacturing choices that are compatible with the mechanical, optical, electrical, and thermal needs.

Challenges for laser operation are not only related to the semiconductor gain chip, but are also convoluted with the actual e-beam properties. Both vertical and edge type laser configurations are feasible. Whereas it is straight-forward to realize a circular e-beam pump spot, as is convenient for vertical emitters, the short gain length makes essential the need for highly reflective mirrors. However, as known for electrically driven devices, epitaxial III-nitride DBRs are challenging to fabricate. Dielectric DBRs on the other hand might fulfil the optical requirements, but their robustness against extreme e-beam bombardment might be compromised. Also, in the case of vertical lasers, substrate removal might be required to minimize absorption losses in the optical

cavity. Edge emitting laser configurations are easier to fabricate. However, realizing a narrow and long continuous e-beam excitation spot is nontrivial, and its precise alignment with respect to the resonator is important.

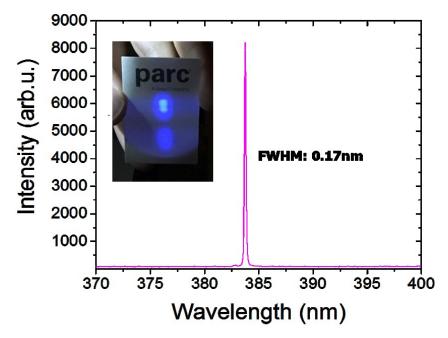


Figure 2. Laser spectrum from electron beam pumped GaN-based edge type laser. Inset: Far-field pattern of the laser emission made visible by focusing onto a fluorescing screen. The spot with lower intensity originates from reflections of the light on the wafer.

#### Advances in Science and Technology to Meet Challenges

In recent years much progress has been made to develop high quality AIN substrates and templates that provide the base for the growth of the device heterostructures. The structural material quality of today's UV emitters has approached a satisfying level with radiative efficiencies exceeding 90% for the best materials. These advancements have also allowed the successful demonstration of low threshold optically pumped lasers all the way down to emission wavelength of 237 nm. Challenges related to efficient carrier injection even at high carrier densities as seen in conventional p/n-junction devices are addressed by means of e-beam pumping. Light extraction is one of the factors that still limits UV emitter performance today, especially for wavelength shorter than 240 nm. This is where light polarization switches from TE to TM and photons are trapped within the chip. Structuring the chip surface and relying on scattering effects have showed promise for enhanced light extraction. However, the effective useable active area is compromised. An alternative approach to overcome the challenges of photon extraction is to fabricate devices on non- or semipolar AIN crystal orientations. For example, first MQW heterostructures grown on (2-201) AIN revealed very promising results [10] and further advances could enable efficient sub-240 nm emitters.

With respect to e-beam pumped edge type lasers further development of a compact line-shaped ebeam pump spot with high power density could significantly improve laser device performance. A longer excited resonator length can provide more gain per photon roundtrip and consequently reduce the laser threshold. Also implementing mirror concepts and laser cavities (for vertical emitting lasers) that do not degrade with high-energy electron bombardment would be of interest. For example, an e-beam pumped surface emitting laser with external cavity (VECSEL) could provide high optical output power with excellent beam properties and would circumvent the issues discussed above. Further advances on thermal management and implementation of ultra-compact components, including the electron gun, vacuum housing, beam shaping, and drive electronics are greatly desired. It is believed that these technical challenges can be mastered and that the final product could be as compact as a conventional light bulb.

### **Concluding Remarks**

Electron-beam pumping of semiconductor materials can be considered an enabling alternative excitation strategy for UV optical emitters. High pump powers of up to 100 Watts and high pump-power densities (e.g., ~1 MW/cm<sup>2</sup>) are accessible to achieve laser operation in both edge type and vertical emitting laser configurations. Especially for the ultra-wide bandgap AlGaN materials e-beam excitation is particularly appealing, as no p-type doping is required in the device heterostructure. As a result, device efficiencies of e-beam pumped spontaneous UV emitters have been shown to meet or exceed state-of-the-art performance levels from conventional LEDs, while delivering high optical output powers. Further advances in miniaturization of various device components will realize a platform technology for compact and efficient emitter systems that can access a wide range of emission wavelengths.

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# 19 – UV laser diodes

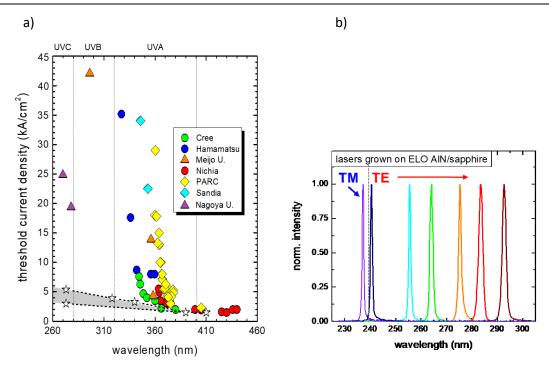
Tim Wernicke<sup>1</sup>, Michael Kneissl<sup>1,2</sup>

- 1 Institute of Solid State Physics, Technische Universität Berlin
- 2 Ferdinand-Braun-Institut, Leibniz Institut für Höchstfrequenztechnik

# Status

Although AlGaN-based ultraviolet light emitting diodes (UV-LEDs) with emission wavelength as short as 211 nm have been realized with emitters covering the entire composition range (see section 12) the situation is quite different for current-injection UV laser diodes (LDs). Due to the greater complexity and increased requirements commercially available UV laser diodes are limited to emission wavelengths above 370 nm [1] and the shortest emission wavelength for a currentinjection LD was for a long time 326 nm [1]. However recently edge emitting laser diodes emitting in the UVC at 271.8 nm [2], and UVB spectral region at 298 nm [3] were demonstrated. As shown in Fig. 1, lasers emitting below 370 nm exhibit an increase in threshold current density with shorter emission wavelength [4]. Laser emitting around 400 nm exhibit threshold current densities of 1-2 kA/cm<sup>2</sup>, while the thresholds of lasers emitting below 340 nm exceed 15 kA/cm<sup>2</sup> and can be only operated under pulsed current conditions [1]. The AlGaN quantum well (QW) based currentinjection lasers in the UVB and UVC wavelength region exhibit a laser threshold of 41.2 kA/cm<sup>2</sup> [3], 25 kA/cm<sup>2</sup> [2], and 19.6 kA/cm<sup>2</sup> [5].

Optically pumped lasing from AlGaN heterostructures with emission wavelengths from 310 nm down to 237 nm [6, 7, 8, 9] has been demonstrated and over the past decade the threshold power densities have been significantly reduced and are now reaching <10 kW/cm<sup>2</sup> for AlGaN multiple quantum well (MQW) structures grown on low defect density bulk AlN substrates [6]. Investigations of the optical gain in such AlGaN MQW heterostructures show a high material gain [7], moderate optical waveguide losses even in the presence of Mg-doped layers [10], and fairly large confinement factors [7]. Advanced laser concepts such as vertical cavity surface emitting lasers (VCSELs) [11], distributed feedback (DFB) [12], and tapered lasers are currently being explored for emitters in the wavelength



**Figure 1.** a) Previously published threshold current densities of current injection laser diodes [2] with data added from [15,16, 17]. b) Spectra of optically pumped lasers in the range of 237 nm to 292 nm [7].

range from 400 nm to 369 nm. For emitters in the UVB and UVC spectral range research activities are predominantly focused on edge-emitting lasers due to significant challenges in materials development (e.g. Mg-doping of AlGaN alloys) and device fabrication technologies (e.g. ohmic contacts).

## **Current and Future Challenges**

The challenges for UV lasers are tremendous but depend strongly on the emission wavelength and composition. In the spectral range above 370 nm, low threshold density GaN-based laser diodes operating in continuous-wave mode with long lifetimes are already commercially available. Lasers emitting at shorter wavelength require AlGaN or InAlGaN active regions and confinement heterostructures leading to increasing tensile strain with increasing aluminium mole fraction when grown on low defect density bulk GaN substrates [1]. In order to avoid the formation of cracks in the laser heterostructure the development of special strain relief techniques is required. Furthermore, the performance characteristic of these lasers also degrades much faster due to the increased operating voltages and currents leading to Joule heating as well as the higher defect densities in these materials. For UVB lasers one of the biggest challenges is the lack of high quality AlGaN substrates with a lattice constant corresponding to the medium AlGaN composition range. For example, growth on bulk AIN substrates is not ideal since the strong compressive strain in the AIGaN quantum well heterostructures leads to relaxation and defect formation during growth [13] and consequently a degradation of the optical properties. This might be the main reason for higher lasing threshold of the demonstrated UVB laser diode in comparison for UVC laser diodes grown on bulk AIN [2, 3]. On the upside, highly conductive n-AlGaN layers and low resistance Ohmic n-contacts have been demonstrated for emitters in this aluminium alloy composition range as also described in section 6 and 7. Furthermore, due to the only moderately higher Mg acceptor ionization energies, low resistivity Mg-doped p-AlGaN layers are also easier to realize [7]. UVC lasers with high structural quality can be realised on low defect density bulk AIN [2, 5, 6, 8]. Since these high aluminium mole

fraction AlGaN heterostructures can be grown pseudomorphically strained on bulk AlN [13], they exhibit excellent optical properties as described in the previous paragraph. However, efficient carrier injection and confinement as well as low resistivity p-AlGaN cladding layers with an aluminium content exceeding 70% pose very difficult challenges for the realization of AlGaN QW lasers emitting in the UVC. As described in section 5, the ionization energy for Mg acceptors in AlGaN alloys steadily increases with the aluminium content leading to very low room temperature conductivities. Here polarization doped p-AlGaN layers [14] without Mg employed in all UVB and UVC laser diodes so far [2, 3] allow moderate Voltages of 13.8 V at the threshold current density of 25 kA/cm<sup>2</sup> [2].

Finally, designing current injection structures that allow high current density operation [2, 3] is a crucial challenge to achieve lasing and find a suitable way for optimizing the structures

#### Advances in Science and Technology to Meet Challenges

For laser diodes in the UVA spectral range emitting below 370 nm recent research developments including reducing point defects in AlGaN alloys (section 5), improved doping schemes (section 6) as well as low resistance Ohmic contacts to AlGaN layers (section 7) should lead to a reduction in operation voltages, cw operation, and improved laser diode lifetimes. As it is often the case, laser diode technologies will also profit from the progress made in the development of UV-LEDs (section 9). However, the root causes for the strong increase in threshold current density at shorter wavelengths is not yet well-understood. Since the increased strain in the AlGaN layers is certainly playing a role different approaches for the growth of low defect density AlGaN heterostructures are critical. This could be either achieved by strain relaxation which, however, needs to be attained without introducing additional threading dislocations or by realizing low defect density AlGaN templates allowing for the coherent growth of a laser diode heterostructure with AlGaN cladding layers exceeding an Al mole fraction of 30%. Another option is the incorporation of InAIN cladding layers for which n- and p-doping was successfully demonstrated [15] and which can be grown latticematched onto low defect density bulk GaN substrates. For lasers in the UVB spectral range the creation of high quality heterostructures by using metamorphic AlGaN buffer layers that promote strain relaxation (see section 10 and [3, 9]) are effective to reduce the TDD. In the longer term Al<sub>0.5</sub>Ga<sub>0.5</sub>N quasi substrates grown, e.g., by hydride vapour phase epitaxy (HVPE), could provide near lattice-matched templates with low threading dislocation densities, smooth surfaces and vertical current-injection through a conducting substrate.

Now, shortly after the first realization of UVC laser diodes it is not clear which factors contribute to the lasing threshold. However, carrier injection and optical confinement under the condition that the diode can sustain high current densities to reach the threshold are probably the most crucial contributions. Part of this is optimizing the polarization doping for efficient hole injection transport [14].

Novel concepts, especially tunnel junctions will also be investigated as these would allow the incorporation of low resistance Si-doped AlGaN cladding layers on both sides of the active region with only a thin Mg-doped tunnel heterojunction for hole injection. Here the key challenge is to minimize the excess voltage drop at the AlGaN tunnel-junction, e.g., by using low-bandgap interlayers (see section 17). The best growth technologies for deep UV tunnel-junction laser diodes either by MOVPE or hybrid MOVPE/MBE as well as the optimum design for a low optical loss of the lasing mode at the tunnel-heterojunction are currently being investigated. Another approach for carrier injection that is currently investigated is the pumping by a high energy electron beam with acceleration voltages of 5 - 10 kV (see section 18). Using this approach, electron-hole pairs are

excited in the laser heterostructure through e-beam irradiation and thus completely avoiding all problems connected to Mg-doping and current-injection. However, such a device requires a very different laser heterostructure design as well as a detailed understanding of the carrier dynamics in order to efficiently collect the charge carriers in the AlGaN quantum wells. Another concern is the long term stability under high density electron beam irradiation as well as the effects of Joule heating, since only 1/3 of the energy is converted to generate electron-hole pairs and the rest is dissipated via various other channels.

Advanced laser concepts will be a topic of intense research as soon as technological building blocks are present and they will be first realized in the UVA region. For DFB, DBR and tapered lasers the technological prerequisites exist and advances will be achieved by optimizing the fabrication technology. Future breakthroughs for VCSELS in the UVA region will be achieved by improving the fabrication of high reflectivity mirrors, e.g., based electro chemical etching [11], and reducing the optical losses in p-contact layers. An extension towards shorter emission wavelength will then be possible by improved epitaxial growth and fabrication technology in combination with tunnel junctions or e-beam pumping.

#### **Concluding Remarks**

Laser diodes emitting in the UV spectral range have so far proven to be a much bigger challenge than UV LEDs. For the recent realization of deep UV current-injection laser diodes significant advances in material quality and device fabrication were required. With the recent improvements in materials growth (e.g., bulk AIN substrates, polarization doping, minimizing point defects) and fabrication technologies (e.g., Ohmic contact formation) initiated by the development of deep UV-LEDs significant reduction of the lasing threshold of UVB and UVC laser diodes can be expected.

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