

Demonstration of Ultraviolet μ LED Array With Novel Electrical Contact Etch Mask

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Abstract—We report on the realization of a uniform array of electrically driven, individually addressable micro light-emitting diodes (μ LEDs) with diameters of $2.5\ \mu\text{m}$ emitting at $372\ \text{nm}$. This highly ordered array was fabricated via a top-down fabrication approach using a novel Ni/Au/Ni structure which combines both a Ni/Au low-resistance electrical contact and durable Ni etch mask. The use of this novel structure allows individual μ LEDs to be addressed with promising electroluminescence intensity and narrow linewidth at ultraviolet-A (UV-A) band. This improved control over μ LED output power and emission location makes our approach an important step towards the development of high efficiency UV-A μ LEDs for a range of applications.

Index Terms—Electrical contact, GaN, micropillar, ultraviolet LED.

I. INTRODUCTION

ULTRAVIOLET (UV) light has numerous applications in sterilization and disinfection, [1], [2] curing of polymers, resins, inks for 3D printing, [3], [4] spectroscopy and non-destructive testing, [5] and shows promise for high-speed optical communication [6] and displays [7]. Still further applications have been identified as materials and fabrication advances enable emission of shorter wavelengths, [8] higher efficiencies, [9], [10] and smaller devices [11]. Within the UV spectrum, UV-A light, with a wavelength between $315\text{--}400\ \text{nm}$, is of especially high interest for photolithography [12] and as a photocatalyst for air and water treatment [7], [8]. Conventional UV-A light emitting diodes (LEDs) based on mesa structures typically have a low overall efficiency, caused in part by especially low light extraction efficiency, typically between 10% - 40% [3]. The poor light extraction efficiency can be improved by shrinking the commonly used mesa structures to micropillars and nanowires, [15] thus μ LEDs are of great interest.

Micropillar and nanowire gallium nitride (GaN) structures can be fabricated either through bottom-up or top-down processes. Selective area epitaxy (SAE) can be used to grow micropillar and nanowire structures from a patterned template in a bottom-up fabrication process [16]. However, this

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approach only offers limited control over the growth rate, diameter, and length of the resulting structures [17]. As an alternative, structures can be instead formed through a top-down approach. In this case, crystal growth is performed at the full wafer scale and structures are formed through photolithography and reactive ion etching (RIE) [18]. This fabrication approach uses well established processes produces highly uniform structures with precise control over their placement and spacing.

Though our previous works have demonstrated successful top-down fabrication of micropillar and nanowire GaN LEDs [18], [19], [20], [21], [22], [23], another limiting factor for high-efficiency UV μ LEDs is poor p-type doping of GaN. Magnesium can be incorporated as a p-type dopant during epitaxial growth, but it forms electrically-inactive complexes with hydrogen [24] and must be activated after growth, typically through high temperature annealing [25]. This activation process is highly inefficient, with only up to 5.25% activation [26] due to magnesium's high activation energy [27]. This poor doping efficiency leads to an extremely high p-GaN work function, predicted to be 6.6 eV, higher than that of all pure metals [28]. The issue of high p-GaN work function has led to the development and optimization of alloyed or multi-layer metal contacts, such as commonly used nickel (Ni)/gold (Au) in an attempt to minimize the Schottky barrier height at the metal-semiconductor interface and form a low-resistance, ohmic electrical contact [29].

The use of nanowire and micropillar structures has been shown to improve both the quantum efficiency and optical output power of the device [30]. As research investigates ever smaller devices, existing fabrication processes become steadily less applicable. Fabrication of GaN LEDs typically consists of deposition and patterning of an etch mask, RIE to form the mesa structure, post-etch removal of the etch mask, and finally, deposition of p-type electrical contacts [21], [31]. However, this well demonstrated process becomes increasingly challenging to implement for ever shrinking devices. The extremely precise lithographic alignment necessary to accurately produce these electrical contacts post-etching presents significant challenges and necessitates a new approach. Previous works [21], [32], [33], [34] use a single, common electrical contact to avoid this challenge. However, this prevents the possibility of individual control of single μ LEDs.

In this work, we propose a novel solution to these challenges and demonstrate the successful fabrication of an array of individually addressable $2.5\ \mu\text{m}$ diameter UV μ LEDs emitting at $372\ \text{nm}$. Our proposed $20\ \text{nm}$ Ni/ $20\ \text{nm}$ Au/ $200\ \text{nm}$ Ni

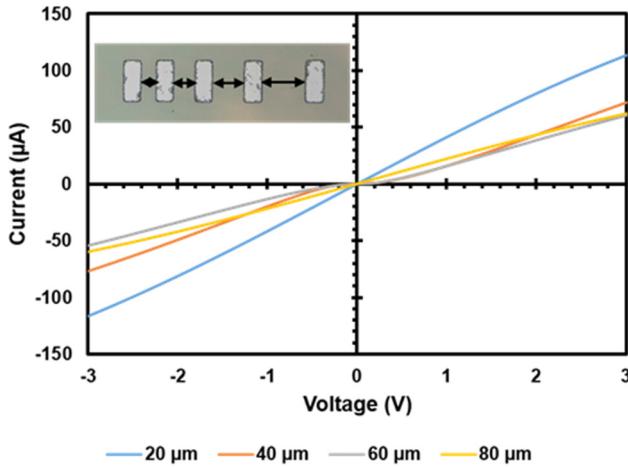


Fig. 1. I-V characteristics from 20 nm Ni/20 nm Au/200 nm Ni TLM measurements after annealing with spacings of 20 μm, 40 μm, 60 μm, and 80 μm between pads. Inset: microscope image of TLM structures used for testing, with spacing indicated by double arrows.

metal structure serves to both act as the durable plasma etch mask necessary for the fabrication of high aspect ratio devices and as a low resistance electrical contact for fabricated devices. By pre-depositing the electrical contacts for each device, the need for a universal, common electrical contact is eliminated and individual UV μ LEDs within the array can be driven independently of the rest of the array, offering greater control over the emitters. This novel fabrication approach offers several advantages over existing processes, most notably enabling the fabrication of individually-addressable micropillar and nanowire LED devices and arrays.

II. CONTACT STRUCTURE OPTIMIZATION AND DEVICE FABRICATION

Our initial optimization of the p-type metal contact stack for our UV-A μ LEDs was performed through a series of transfer length method (TLM) experiments. These initial experiments used a contact metal stack consisting of 20 nm of Ni and an Au interlayer of 20 nm, 40 nm, 60 nm, or 80 nm, to serve as the ohmic contact, followed by a 200 nm of Ni to serve as the durable etch mask for the chlorine-based plasma etch. All metals were deposited via thermal evaporation onto the surface of the top p-GaN layer, doped to a level of $\sim 10^{18}/\text{cm}^3$ with Mg, forming a set of 30 μm x 100 μm contact pads spaced 20 μm, 40 μm, 60 μm, and 80 μm apart, as shown in the inset of Figure 1. Following metal evaporation, samples were transferred to a rapid thermal annealing (RTA) tool and were annealed for 5 minutes at 500°C in an oxygen environment. A second set of similar TLM contacts, consisting of a single 20 nm Ni layer deposited onto p-GaN, were also deposited to serve as a reference, and were also annealed for 5 minutes at 500°C in an oxygen environment. TLM measurements were then performed and the electrical performance of the Ni/Au/Ni contacts was compared to that of the single layer Ni control contact structures. As shown in Figure 2, the use of a Ni/Au/Ni contact structure resulted in a reduction of contact resistance of up to 75% when compared to contacts

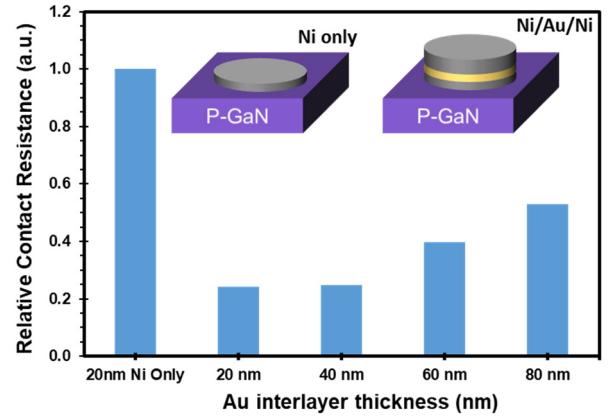


Fig. 2. Relative contact resistance comparison of annealed single layer Ni contact and three-layer Ni/Au/Ni contacts with varying Au interlayer thicknesses.

composed solely of Ni. For a simple Ni/Au bilayer contact, annealing in an oxygen-containing atmosphere causes outdiffusion and oxidation of Ni. This results in the formation of a discontinuous Au layer, interrupted by NiO islands at the GaN surface. The presence of NiO, a p-type semiconductor itself, at the GaN-metal interface serves to reduce the overall contact resistance. Additionally, further interdiffusion at the surface leads to the formation of an Au-Ga solid solution and thus acceptor-like Ga vacancies at the GaN-metal interface, further reducing contact resistance. This process of interdiffusion and oxidation progresses differently with the addition of a Ni capping layer, as with our novel contact structure. Exposed Ni metal quickly oxidizes during annealing, as with the Ni/Au contact. However, unlike the bilayer Ni/Au contact, the thick Ni capping layer and increasing Au interlayer thickness slows the formation or transport of NiO to the GaN-metal interface. The presence of un-oxidized Ni in contact with the p-GaN surface, detrimental to the formation of a low-resistance, ohmic contact. This can be seen in Figure 2 as an increase in measured contact resistance with increasing Au interlayer thickness [35], [36], [37]. Based on this initial optimization, an Au interlayer thickness of 20-40 nm showed the lowest contact resistance. After annealing, these Ni/Au/Ni electrical contacts showed nearly linear current-voltage (I-V) characteristics for all different spacing lengths, as shown in Figure 1. Based on these TLM results, an Au thickness of 20 nm was selected for subsequent μ LED fabrications.

Fabrication of the UV-A μ LED array was performed on a c-plane epitaxial stack similar to that used by Huang, [38] which consisted of a 20 nm AlN buffer layer, a 2 μm undoped GaN layer, a 2.5 μm n-AlGaN lower cladding layer, an InGaN/AlGaN active region, a 15 nm p-AlGaN upper cladding layer, and a 200 nm p-GaN contact layer, as shown in Figure 3(a). The active region consisted of 10 periods of 3 nm $\text{In}_{0.03}\text{Ga}_{0.97}\text{N}$ wells and 11 nm $\text{Al}_{0.02}\text{Ga}_{0.98}\text{N}$ barriers. The epitaxial structure was grown on a sapphire substrate using metalorganic chemical vapor deposition (MOCVD). As shown in Figure 3(b), to fabricate devices, lift-off resist (LOR) and photoresist were deposited and patterned using 405 nm direct write lithography using a Heidelberg DWL 66+,

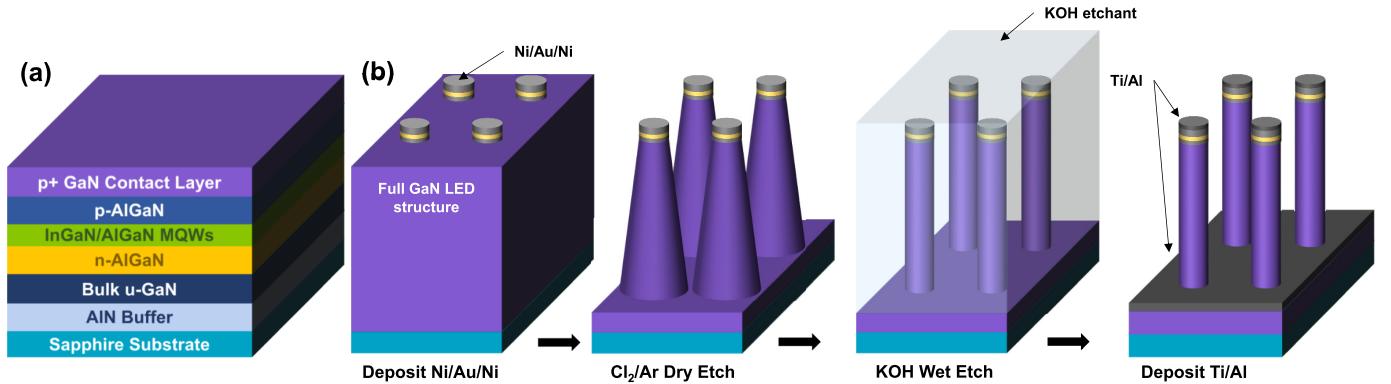


Fig. 3. (a) GaN UV LED epitaxial structure. (b) Device fabrication process showing deposition of Ni/Au/Ni contact and etch mask, Cl₂/Ar dry etching, wet etching in KOH solution, and final deposition of Ti/Al electrical contacts.

and the metal stack was deposited using highly anisotropic thermal evaporation. Based on prior TLM results, Ni/Au/Ni thicknesses of 20/20/200 nm were utilized. After metal lift-off, dry etch was performed using a Cl₂/Ar plasma (40/10 sccms Cl₂/Ar, 500 W inductively coupled plasma (ICP) power, 225 W forward power, 440 nm/min etch rate, 192 second duration) in a PlasmaTherm ICP-RIE to form a large array of 2.5 μ m diameter, 1.4 μ m tall micropillars with a 10 μ m pitch, as shown in Figure 4(a). The sidewalls of these micropillars are left roughened, as shown in Figure 4(b), indicative of surface damage typically caused by the ICP-RIE etch process [39]. Hydroxyl-based chemistries, such as KOH, are commonly used on GaN structures to passivate and to remove surface damage caused by dry etching [22]. The wafer was submerged in a solution of 40% AZ400K (a photoresist developer containing 2 wt% KOH) heated to 80°C for a total of 15 minutes. To ensure that this etch duration was sufficient to produce highly vertical sidewalls, the wafer was examined and imaged using a scanning electron microscope (SEM), shown in Figure 4(c). Following this inspection, the p-type metal was annealed at 450°C for 40 seconds under nitrogen. A universal n-type contact was then deposited using thermal evaporation. The deposition of the n-contact was self-aligned in that the micropillars themselves were used to mask deposition of the n-metal on the sidewalls of the micropillars. The sample was carefully aligned directly above the evaporation source, and the highly anisotropic nature of thermal evaporation allowed metal deposition to occur on the tops of the micropillars and on the n-AlGaN at the bases of the micropillars, but not on the micropillar sidewalls. This n-metal stack consisted of 40/200 nm Ti/Al. After this deposition, the LED array was completed and the individually addressable UV-A μ LEDs could be tested and characterized.

III. ELECTRICAL AND OPTICAL CHARACTERIZATIONS OF μ LEDs

Electrical testing of individual μ LEDs was performed by manually probing the n- and p-contacts and applying a forward bias. These tests show a turn on voltage of approximately 5 V, as shown in Figure 5, above this threshold voltage, individual μ LEDs illuminate as shown in the inset of Figure 5.

By placing an optical fiber near the probed device, electroluminescence (EL) spectra can be collected. These spectra show a peak emission wavelength of 372 nm, as shown in Figure 6. Emission peaks show a consistent FWHM of approximately 11 nm, similar to what is reported in other works [40], [41], [42]. As drive current is increased from 60 μ A to 150 μ A, we observe both an increase in EL intensity and a slight redshift of approximately 2 nm in peak emission wavelength. This redshift is expected and is likely due to an increase in junction temperature with increasing drive current [42].

We have also analyzed light extraction efficiency from UV-A μ LEDs with three-dimensional (3D) finite-difference time-domain (FDTD) simulations using commercially available Synopsys Fullwave software which is specifically designed to model light propagation in a range of structures. These simulations model the propagation of electromagnetic waves through a defined structure by finding direct solutions for Maxwell's curl equations. Simulations were conducted using methods similar to those used in Ref. [20].

Simulation setup begins with modelling of the structure of interest, a 2.5 μ m diameter, 1.4 μ m tall μ LED. Material types and properties can be defined for the different layers within the structure and our modeled μ LED uses a similar layer structure as our fabricated μ LEDs, shown in Figure 3(a). Three dipole sources were placed at the center of the MQW region, aligned with the x, y, and z-axes and a 20 nm cubic monitor was defined encompassing these dipole sources to capture the full luminous flux. This flux is compared against the luminous flux measured by another set of monitoring planes placed outside the simulated μ LED. These exterior planes form a rectangular prism, extending 500 nm beyond the sides and top of the μ LED structure and extending 200 nm below the bottom nAlGaN surface. The external monitor planes act as perfect absorbers, preventing unintended wave reflections off the simulation boundaries. Overall light extraction efficiency (LEE) can then be calculated as the ratio of flux measured by the external monitor planes around the sidewalls and top μ LED surface and the total flux produced by the combined dipole sources in the MQW region.

Two sets of simulations were conducted, one with a simple, single 20 nm layer of Ni covering the top surface of the

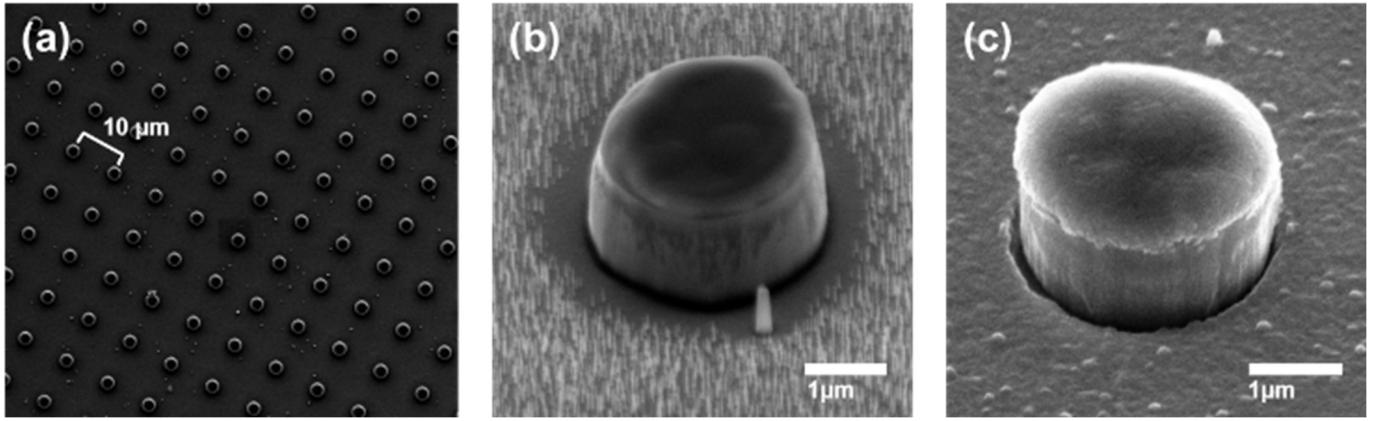


Fig. 4. SEM images of (a) fabricated ordered UV-A μ LED array, (b) micropillar with flared base after dry etching, (c) μ LED after follow-up wet etching and n-metal deposition.

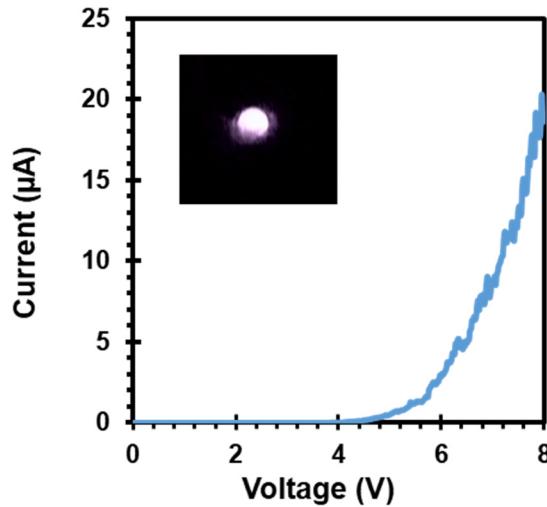


Fig. 5. I-V characteristics from a single UV-A μ LED. Inset: microscope image of one illuminated UV-A μ LED.

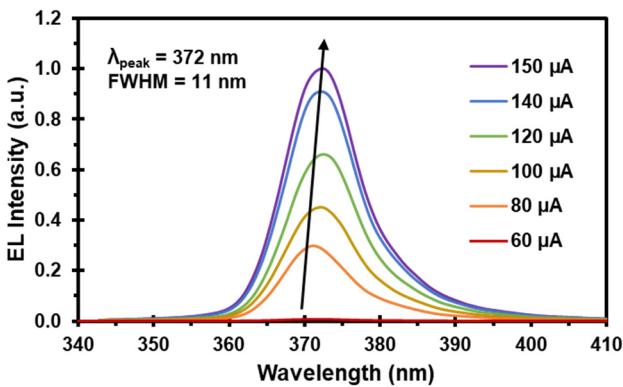


Fig. 6. EL spectra recorded from a single μ LED under different drive currents.

μ LED that represents common p-contact cases, and another using our novel 20 nm Ni/20 nm Au/200 nm Ni contact structure covering the μ LED top surface. Simulated electric field intensity within the two μ LED structures can be seen in Figure 7. These simulation results indicate that even in the case of a thin 20 nm layer of Ni covering the top surface,

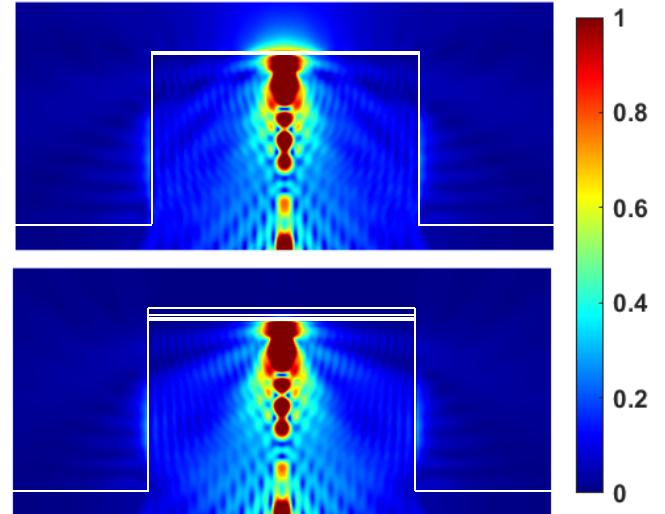


Fig. 7. Simulated electric field intensity within μ LEDs, (top) with a 20 nm Ni contact deposited on the top surface, and (bottom) with a 20 nm Ni/20 nm Au/200 nm Ni contact deposited on the top surface.

extraction efficiency is only 1.6% through the top surface with the majority of escaping light being emitted instead through the sidewalls. When our novel Ni/Au/Ni structure is simulated, the additional metal layers have a negligible effect on total light extraction from the whole μ LED structure, calculated to be 25.8% for the Ni top contact and 24.2% for the Ni/Au/Ni contact. Overall, these results suggest that the use of a Ni/Au/Ni contact structure offers several advantages in terms of fabrication and electrical contact resistance without significantly affecting total light extraction from the device.

IV. CONCLUSION

In conclusion, we have successfully demonstrated an array of individually addressable UV-A GaN μ LEDs emitting at 372 nm. These μ LEDs were fabricated using a novel Ni/Au/Ni structure to form highly uniform 2.5 μ m diameter, 1.4 μ m tall μ LEDs with a 10 μ m pitch. This novel Ni/Au/Ni structure serves as both a durable dry etch mask during device fabrication and as a low resistance electrical contact during

device operation, allowing individual μ LEDs to be probed and tested. Electrical testing of these μ LEDs shows a consistent, narrow linewidth of 11 nm and a turn on voltage of 5 V. This novel approach enables an extremely high level of control over the output of μ LED arrays as individual devices can be driven rather than the whole array. Additional study of this process could further improve the effectiveness of this Ni/Au/Ni structure by minimizing the capping Ni layer thickness based on the desired etch depth, potentially lowering device turn on voltages. Overall device efficiency could also be improved through additional optimization of the annealing process to further reduce contact resistance. By controlling individual μ LEDs, total emission power and the location of light generation can be easily varied making this a promising approach for numerous applications including photolithography and photocatalytic processes. This approach can also be applied to μ LEDs emitting at other wavelengths, including visible light, which can open the door for higher resolution displays.

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