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# Eu<sup>3+</sup> optical activation engineering in Al<sub>x</sub>Ga<sub>1-x</sub>N nanowires for red solid-state nano-emitters

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## ABSTRACT

In this work, Eu<sup>3+</sup>-implanted and annealed Al<sub>x</sub>Ga<sub>1-x</sub>N ( $0 \leq x \leq 1$ ) nanowires (NWs) grown on GaN NW template on Si (111) substrates by plasma-assisted molecular beam epitaxy are studied by  $\mu$ -Raman, cathodoluminescence (CL), nano-CL, and temperature-dependent steady-state photoluminescence. The preferential location of the Eu<sup>3+</sup>-implanted ions is found to be at the Al<sub>x</sub>Ga<sub>1-x</sub>N top-section. The recovery of the as-grown crystalline properties is achieved after rapid thermal annealing (RTA). After RTA, the red emission of the Eu<sup>3+</sup> ions is attained for all the samples with below and above bandgap excitation. The <sup>5</sup>D<sub>0</sub> → <sup>7</sup>F<sub>2</sub> transition is the most intense one, experiencing a redshift with increasing AlN nominal content ( $x$ ) from GaN to AlN NWs. Moreover, AlN nominal content and annealing temperature alters its spectral shape suggesting the presence of at least two distinct optically active Eu<sup>3+</sup> centers (Eu1 and Eu2). Thermal quenching of the Eu<sup>3+</sup> ion luminescence intensity,  $I$ , is found for all the samples from 14 K to 300 K, being the emission of Eu<sup>3+</sup>-implanted AlN NWs after RTA at 1200 °C the most stable ( $I_{300\text{ K}}/I_{14\text{ K}} \sim 80\%$ ). The GaN/AlN interface in this sample is also found to have a key role in the Eu<sup>3+</sup> optical activation.

## KEYWORDS

*Al<sub>x</sub>Ga<sub>1-x</sub>N, nanowires, europium, luminescence, red-emitters*

# 1. INTRODUCTION

The nanowire (NW) configuration in semiconductor systems such as oxides, nitrides, and arsenides is highly promising for the next generation nanoscale photonic, optoelectronic and photovoltaic devices thanks to the characteristic large area to volume ratio [1–3]. In particular, for light-emitting devices, it was demonstrated that group III-nitride (III-N) NWs can achieve higher light extraction compared to their 2D counterparts [4,5]. Furthermore, the efficient strain relaxation during NW growth allows the use of highly lattice-mismatched substrates (*e.g.*, silicon) with a reduced density of extended defects compared to thin-film epitaxy, facilitating their compatibility with current microtechnology. III-N nanostructures, with a direct and tunable bandgap energy from 6.2 eV (AlN) to 0.7 eV (InN), have shown unique properties suitable for light-emitting diodes (LEDs), micro-LED technology, and solid-state lasers operating in a wide range of the electromagnetic spectrum [6–12].

The implementation of micro-LED emissive displays requires a monolithic integration of independently controlled red-green-blue (RGB) LEDs to attain high emission efficiency and thermal stability [13]. Blue and green LEDs are obtained by  $\text{In}_x\text{Ga}_{1-x}\text{N}$  bandgap/nanostructure engineering. However, when further increasing InN content, necessary to achieve red emission, the efficiency is greatly reduced due to lower crystalline quality and increased piezoelectric/spontaneous polarization [14,15]. Thus, a practical and viable strategy is needed to improve the efficiency of III-N red-emitters for monolithic integration of RGB LEDs.

The incorporation of the rare-earth (RE) trivalent europium ions ( $\text{Eu}^{3+}$ ) into III-N layers [16–23] and nanostructures [24–32] was proved as an excellent strategy to obtain red emission characterized by the sharp and stable  $\text{Eu}^{3+}$  intra- $4f^6$  transitions. This approach was successfully implemented for *in situ*  $\text{Eu}^{3+}$ -doped GaN-based emitting devices [31,33–38], with  $\text{Eu}^{3+}$ -related luminescence external quantum efficiency (EQE) values reaching ~29% at room temperature (RT) and ~48% at 77 K [39]. In order to sustain the EQE value up to RT, it is important to overcome the RE thermal quenching. Indeed, as a possible solution, it was proposed that wider bandgap semiconductors show weaker RE thermal quenching [40,41], therefore paving the way to further explore the incorporation of RE in  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  and AlN hosts.

As an alternative to *in situ* RE incorporation, *ex-situ* ion implantation constitutes a suitable method to obtain a controlled doping profile with concentration even beyond solubility limits [18,25,26,32,42–47]. This technique requires post-implantation thermal annealing to recover the implantation-induced damage and to optically activate the RE ions. Nevertheless, previous reports indicate that the ion implantation-induced damage decreases with increasing AlN content in  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  layers [48–50], which is an additional argument to explore such materials. To the best of our knowledge, studies on ternary  $\text{Al}_x\text{Ga}_{1-x}\text{N}:\text{Eu}^{3+}$  over the entire range of AlN content were only performed in layers and revealed a reduced thermal quenching with enhanced  $\text{Eu}^{3+}$  luminescence efficiency for high AlN content hosts [18,51,52]. Besides,  $\text{Eu}^{3+}$ -implantation in III-N NWs can play a key role for improving  $\text{Eu}^{3+}$  luminescence efficiency [26–28]: on the one hand, NWs are expected to be less affected by residual implantation-induced damage compared to 2D structures [25]; and on the other hand,  $\text{Eu}^{3+}$ -implantation occurs not only on the upper surface (*c*-plane) but also on side facets. Although this increases the complexity of implantation damage build-up since crystallographic surface planes can lead to distinct damage in GaN during implantation [53,54], it provides an additional parameter to control the distribution of  $\text{Eu}^{3+}$  and defects within the NW via the implantation geometry [25]. Furthermore, it was shown that the

formation of extended defects during ion implantation is reduced for NWs as compared with thin films [25].

Therefore, exploring  $\text{Eu}^{3+}$ -implantation of  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NWs turns out to be a very promising strategy for efficient red light-emitting devices with expected lower implantation damage, lower thermal quenching, higher efficiency, and higher light extraction. In the present work, the  $\text{Eu}^{3+}$  luminescence, achieved after RTA, is studied with emphasis on the most intense  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  transition. Both excitations, above and below the hosts' bandgap energies, are explored in detail. At least two main  $\text{Eu}^{3+}$  centers whose optical activation depends on both AlN nominal content and annealing temperature are identified. The temperature dependence of this emission is evaluated in order to explore its thermal stability in these nanostructures. The contribution of the GaN/AlN interface to the optical activation of these centers is also highlighted.

## 2. EXPERIMENTAL SECTION

The  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NWs with different AlN nominal content,  $x$  ( $0 \leq x \leq 1$ ), were grown in nitrogen-rich conditions by plasma-assisted molecular beam epitaxy (PA-MBE) on Si (111) substrates. The substrate temperature was varied between 850-930 °C depending on  $x$ ; the substrate was continuously rotated to promote homogeneous NW growth [55]. A 3 nm-thick AlN buffer layer was deposited prior to NW growth to improve their vertical orientation [56]. The GaN NW template was grown on the AlN buffer layer, followed by the growth of  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NW sections [57]. The obtained NW structure (Fig. 1a), in terms of density and dimensions of each section, depends on  $x$ , as indicated in Table 1. The existence of fluctuations in the AlN molar fraction along the growth axis ( $c$ -axis) was demonstrated in these NWs, and it was found to dominate the structural and optical properties of  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NWs at the nanoscale [57–60].

$\text{Eu}^{3+}$ -implantation was realized at room temperature (RT) using 300 keV  $\text{Eu}^{3+}$  ions with a fluence of  $10^{14} \text{ cm}^{-2}$  at a tilt angle of 45° with respect to the  $c$ -axis (Fig. 1a). During implantation, the samples were rotated for a more homogeneous distribution of  $\text{Eu}^{3+}$  ions and implantation-induced defects [25,61]. Such implantation geometry favors that  $\text{Eu}^{3+}$  ions are mostly incorporated in the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  top-section, being their insertion in GaN NW template unlikely to occur due to shadowing effects caused by the high density of NWs and their longitudinal dimensions.

Monte Carlo simulations, obtained by SRIM 2013 code [62], predict a projected range of ~40 nm and ~55 nm, for GaN and AlN respectively (maximum  $\text{Eu}^{3+}$  concentration  $\sim 2.3\text{-}2.5 \times 10^{19} \text{ cm}^{-3}$ ). It is important to note that SRIM does not consider the three-dimensionality of the NWs and therefore the projected range may be slightly different. Indeed, unlike layered structures for which the ions only impinge on the top facet, for oblique incidence the ions can impinge on the sidewall facets of the NWs, causing a higher estimated doped thickness than predicted by SRIM.

Post-implantation rapid thermal annealing (RTA) was carried out for  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  ( $x > 0$ ) NWs during 30 seconds under nitrogen ( $\text{N}_2$ ) flow at 1000 °C and 1200 °C. For GaN NWs ( $x = 0$ ), RTA was performed at 1000 °C only, since higher annealing temperatures may induce severe damage in GaN due to nitrogen out-diffusion [63]. A common strategy to protect GaN for annealing temperatures above 1000 °C is the use of an AlN cap layer [64,65]. Thus, by annealing  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  ( $x > 0$ ) NWs at 1200 °C, it is not expected to strongly damage the GaN NW template, as it is protected by the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  top-section [57]. Similar annealing conditions were reported to successfully achieve the recovery of the lattice damage introduced by the implantation and the optical activation of  $\text{RE}^{3+}$  ions in III-N layers [66,67] and nanostructures [26–28].

Table 1 summarizes the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NWs studied in this work, and the used  $\text{Eu}^{3+}$ -implantation and RTA conditions. The samples' labels and their structural characteristics (NWs density and dimensions of each section) are also included.

**Table 1.** Structural characteristics of the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NWs studied in this work: typical length ( $h$  in nm), average diameter ( $d$  in nm), and NWs density ( $\text{cm}^{-2}$ ). (N/A: not applicable).

$\text{Al}_x\text{Ga}_{1-x}\text{N}$		$x = 0$	$x = 0.3$	$x = 0.5$	$x = 0.75$	$x = 1$
<b><math>\text{Al}_x\text{Ga}_{1-x}\text{N}</math>-asgr (as-grown)</b>						
$\text{Al}_x\text{Ga}_{1-x}\text{N}$ NWs top-section	$d$	N/A	~35	~40	~45	~175
	$h$		~225	~260	~225	~500
GaN NW template	$d$		~35	~40	~35	~75
	$h$		< 200	< 200	< 200	~1500
GaN NWs	$d$	~35	N/A			
	$h$	~750				
NWs density		$\sim 1.5 \times 10^{10}$	$\sim 2.5 \times 10^{10}$			$\sim 5 \times 10^9$
<b><math>\text{Al}_x\text{Ga}_{1-x}\text{N}</math>-asimp (as-implanted)</b>						
$\text{Eu}^{3+}$ fluence of $10^{14} \text{ cm}^{-2}$ , 300 keV, $45^\circ$ tilt vs $c$ -axis						
<b><math>\text{Al}_x\text{Ga}_{1-x}\text{N}</math>-1000</b>						
RTA at 1000 °C for 30 s in $\text{N}_2$ ambient						
<b><math>\text{Al}_x\text{Ga}_{1-x}\text{N}</math>-1200</b>						
<b>Rapid thermal annealing (RTA)</b>		N/A	RTA at 1200 °C for 30 s in $\text{N}_2$ ambient			

Scanning electron microscopy (SEM) of the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -asgr samples was performed using a Zeiss Ultra-55 system. Figs. 1b and 1c show representative SEM images of the  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$ -asgr NWs in side-view and top-view, respectively. In Fig. S1 (Supplementary Information), representative SEM images for the other NWs are depicted. Due to their density and dimensions, the vertical NWs exhibit partial coalescence caused by their bundling, as reported by Kaganer *et al.* for GaN NWs [68]. Radial growth of the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NWs top-section is observed for high AlN nominal contents (Fig. S1e for  $x = 0.75$ , and Fig. S1g for  $x = 1$  in Supplementary Information) due to limited diffusion of Al atoms along the sidewalls towards the top [69]. This effect also contributes to the coalescence of the NWs at the top-section. Moreover, substrate surface coverage (fill factor) is lower for  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NWs ( $0 < x < 1$ ) in comparison to AlN NWs (Figs. S1d, S1f, and S1h in Supplementary Information), which reduces the shadowing effects during  $\text{Eu}^{3+}$ -implantation and increases the probability of  $\text{Eu}^{3+}$  to reach the GaN NW template.

RT  $\mu$ -Raman measurements (Horiba Jobin-Yvon HR800) were performed on  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NW ensemble (beam spot  $\sim 1 \mu\text{m}^2$ ), under 441.6 nm excitation line (He-Cd laser) in backscattering geometry  $z(x, \cdot)\bar{z}$  to reduce the Si (111) substrate contribution. The beam was focused using a  $\times 100$  magnification objective ( $NA = 0.9, f = 0.1 \text{ cm}$ ).

PL measurements were carried out on  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NW ensemble (beam spot  $\sim 5 \text{ mm}^2$ ) using the 325 nm (3.8 eV) excitation line (He-Cd laser) with an excitation power density of about  $0.5 \text{ W/cm}^2$ .

Such energy allows above bandgap excitation for GaN NWs ( $x = 0$ ) and below bandgap excitation for  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NWs ( $x \geq 0.3$  in our samples). The PL signal was measured in a dispersive system SPEX 1704 monochromator (1 m, 1200 grooves/mm) coupled to a water-cooled Hamamatsu R928 photomultiplier tube. The samples were mounted in a cold finger of a closed-cycle helium cryostat and the temperature was controlled between 14 K and 300 K.

CL measurements were conducted on AlN-1000 and  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1200 ( $x > 0$ ) NW ensembles (beam spot  $\sim 2 \mu\text{m}^2$ ), at 5 K, under 5 kV acceleration voltage. The electron beam leads to the excitation of all the luminescence mechanisms present in the material, especially the creation of electron-hole pairs allowing above bandgap excitation.

It is important to mention that in CL, PL, and  $\mu$ -Raman measurements, the average optical response of a few hundreds of NWs is collected simultaneously. Furthermore, a 5 kV acceleration voltage used in CL measurements results in a lower penetration depth compared to PL. In other words, the CL response comes predominantly from the NWs top-section ( $\sim 70$ -190 nm), while the PL optical response is mainly from the GaN/ $\text{Al}_x\text{Ga}_{1-x}\text{N}$  interface and the GaN NWs template (up to  $\sim 80$  nm below the interface) due to the lower absorption of the used laser line in the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NWs top-section for  $x > 0$ .

Additionally, AlN-1200 NWs were dispersed on a foreign Si substrate, allowing nano-CL measurements on individual NWs, at 5 K, under 10 kV acceleration voltage.

### 3. RESULTS AND DISCUSSION

#### 3.1. Structural and vibrational properties

Wurtzite (WZ) III-N semiconductors belong to the  $C_{6v}^4$  space group with four atoms in the primitive cell, occupying  $C_{3v}$  sites. At the Brillouin zone (BZ) center,  $\mathbf{q} = 0$ , six optical modes are active in first-order Raman scattering processes  $A_1(\text{TO})$ ,  $A_1(\text{LO})$ ,  $E_1(\text{TO})$ ,  $E_1(\text{LO})$ ,  $E_2^{\text{H}}$ , and  $E_2^{\text{L}}$ . In the backscattering configuration  $z(x, \cdot)\bar{z}$ ,  $E_2^{\text{L}}$ ,  $E_2^{\text{H}}$ , and  $A_1(\text{LO})$  phonons are allowed according to the selection rules [70]. For ternary alloys such as  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ ,  $A_1(\text{LO})$  is characterized by a one-mode behavior (phonon frequency varies continuously from the  $A_1(\text{LO})$  of one end binary to that of the other one with the mode strength remaining almost constant), while  $E_2^{\text{H}}$  exhibits a two-mode behavior (two impurity modes:  $E_2^{\text{H}}$  (GaN-like) and  $E_2^{\text{H}}$  (AlN-like), close to the frequency of the respective binaries), which blueshifts with  $x$  [71–73]. Nevertheless, it is important to consider that their behavior could be influenced by strain and compositional effects, while  $E_2^{\text{H}}$  (GaN-like) could also be affected by the possible superposition with the alloy  $E_1(\text{TO})$  phonon [71–73].

Fig. 2a illustrates representative normalized  $\mu$ -Raman spectra of the  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$ -(asgr, asimp, 1000, and 1200) NWs. The calculated phonon density of states (DOS) for GaN and the disorder-induced Raman spectrum for erbium (Er)-implanted AlN layers are included for comparison [74]. Fig. S2 (Supplementary Information) depicts the  $\mu$ -Raman response of all the samples acquired in the same conditions. It should be mentioned that the allowed  $A_1(\text{LO})$  phonon was not detected in these experiments. Except for AlN-asgr presenting  $E_2^{\text{H}}$  (GaN) and  $E_2^{\text{H}}$  (AlN),  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -asgr exhibit an asymmetric response in the  $E_2^{\text{H}}$  (GaN) region (blue rectangle in Fig. 2a and Fig. S3a). The presence of this asymmetry (to higher frequencies) and its higher full width at half maximum (FWHM) when compared to  $E_2^{\text{H}}$  (GaN) of the binary, suggest more than one contribution. For  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$ -asgr, such response can be deconvoluted into two phonons (section S2.2. in Supplementary Information):  $E_2^{\text{H}}$  (GaN) at  $568 \text{ cm}^{-1}$  associated with the GaN NW

template and  $E_2^H$  (GaN like) at  $571\text{ cm}^{-1}$  related to the alloy top-section. The resulting active phonons for all the samples are summarized in Fig. 2b and Table S1 (Supplementary Information).

We found that all  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -asgr samples exhibit the  $E_2^H$  (GaN) phonon at  $567\text{-}568\text{ cm}^{-1}$  (experimental error of  $\pm 1\text{ cm}^{-1}$ ) indicating that both GaN-asgr NWs and GaN NW template are nearly-relaxed on Si (111) substrate (dotted line in Fig. 2b indicates the strain-free value of  $567.6\text{ cm}^{-1}$  [74]). For  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -asgr ( $x = 0.3, 0.5, \text{ and } 0.75$ ), the  $E_2^H$  (GaN-like) phonon (within the dashed rectangle in Fig. 2b) is composed of one or two components, indicating possible compositional fluctuations within the same NW or in different NWs, as discussed in other works [57,75].  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -asgr  $E_2^H$  (GaN-like) frequencies are found to be lower than the ones obtained for  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  layers [73], which can be due to *i*) the tensile strain introduced by the growth of  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  on the GaN NW template, and/or *ii*) the lower AlN content compared to the nominal value. For the AlN-asgr, the  $E_2^H$  (AlN) phonon located at  $655\text{ cm}^{-1}$ , is slightly redshifted with respect to the strain-free value ( $657.4\text{ cm}^{-1}$  [74]), evidencing a tensile strain in the AlN section as a result of their growth on GaN NW template with higher *in-plane* lattice parameter ( $a_{\text{GaN}} = 3.189\text{ \AA}$  and  $a_{\text{AlN}} = 3.112\text{ \AA}$  [76]).

After  $\text{Eu}^{3+}$ -implantation, the lattice damage induces a higher absorption in the affected region. Thus, the  $\mu$ -Raman signal (double absorption in backscattering geometry) and the signal-to-noise ratio (SNR) are strongly reduced. The  $\mu$ -Raman spectra of the  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$ -asimp (Fig. 2a) exhibits disorder-activated second-order Raman scattering processes involving phonons from the entire BZ. As a result, the feature located at the  $E_2^H$  phonon mode shows an asymmetry at the low-frequency side (orange rectangle in Fig. 2a), while the asymmetry is observed at the high-frequency side in  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$ -asgr. A thorough analysis in the  $E_2^H$  (GaN) region reveals the presence of a strain-free  $E_2^H$  (GaN) phonon originating from the GaN NW template region not affected by  $\text{Eu}^{3+}$ -implantation, and an additional contribution located at  $562\text{ cm}^{-1}$  from the GaN NW template region affected by  $\text{Eu}^{3+}$ -implantation (detailed in section S2.2. in Supplementary Information). The decrease of the  $E_2^H$  (GaN-like) contribution confirms that the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  top-section is the most affected by  $\text{Eu}^{3+}$ -implantation. One can notice that a similar behavior is obtained for  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -asimp (with  $0.3 < x < 1$ ). When it comes to AlN-asimp, the  $E_2^H$  (GaN) phonon originating from the GaN NW template keeps the same frequency, while the  $E_2^H$  (AlN) experiences a redshift (Fig. S2b in Supplementary Information), indicating that  $\text{Eu}^{3+}$ -implantation affects the AlN top-section. The obtained phonon redshift can be associated with hydrostatic tensile stress within the implanted volume, in good agreement with X-ray diffraction and molecular dynamics results on RE-implanted GaN NWs [25,77,78]. In GaN-asimp and AlN-asimp, we have estimated *quasi*-hydrostatic tensile stress ( $\sigma_{xx} \approx \sigma_{yy} \approx \sigma_{zz}$ ) of  $\sim 1\text{ GPa}$  (section S2.3. in Supplementary Information).

After annealing,  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$  NW ensemble shows a  $\mu$ -Raman response dominated by first-order scattering processes at the BZ center (Fig. 1a), similar to the one of the  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$ -asgr. This result, together with an improved SNR for the highest annealing temperature, is consistent with a recovery of the as-grown crystalline structure. Furthermore, we confirm that the implantation-induced tensile stress gradually decreases after annealing for the binaries (Fig. S4 in Supplementary Information).

## 3.2. Emission properties

### 3.2.1. Above bandgap excitation of $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1200 ( $x > 0$ )

While the  $\text{Eu}^{3+}$  free ion intra- $4f^6$  transitions, such as from  ${}^5\text{D}_J (J=0,1,2)$  to  ${}^7\text{F}_J (J=0-6)$  multiplets, are forbidden by electric dipole according to Laporte selection rules, they happen to be partially allowed when the ions are introduced in a host material, since the crystal-field perturbation leads to the admixture of states with different parities [79,80]. Besides, crystal-field perturbation together with  $4f$  electron spin-orbit coupling and electrostatic interaction contribute to the splitting of the energy levels in different Stark levels [81]. In III-N hosts,  $\text{Eu}^{3+}$  ions likely incorporate into cation substitutional sites ( $C_{3v}$  symmetry) [82], which leads to  ${}^7\text{F}_1$  and  ${}^7\text{F}_2$  multiplets splitting into two (one doublet and one singlet) and three (two doublets and one singlet) distinct levels, respectively. Slightly different  $\text{Eu}^{3+}$  environments (*e.g.*, substitutional cation site surrounded by cation/anion vacancies) can lower the symmetry and lift the degeneracy [44,81,83], reaching a maximum number of three and five levels for the  ${}^7\text{F}_1$  and  ${}^7\text{F}_2$  multiplets, respectively. The  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_0$  transition is characterized by its singlet character, *i.e.* no crystal field splitting is allowed for  ${}^5\text{D}_0$  and  ${}^7\text{F}_0$  levels, implying that the number of the observed  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_0$  transitions corresponds to the number of non-equivalent active sites if accidental overlaps are not considered [32,44,81,83,84].

Fig. 3a represents 5 K CL spectra of  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1200 NW ensembles ( $x > 0$ ) with well-resolved characteristic sharp luminescence lines of the  $\text{Eu}^{3+}$  intra- $4f^6$  shell transitions, for which the most intense emission corresponds to the  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  transition. The intensity of each spectrum was normalized to the corresponding  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  transition. The observation of intra- $4f^6$  shell transitions indicates that the implantation and annealing conditions successfully achieved the activation of the  $\text{Eu}^{3+}$  ions in these hosts. The CL spectra magnified around the intra- $4f$  transitions are shown in Figs. S5a-S5f (Supplementary Information). Table S2 (Supplementary Information) summarizes the  $\text{Eu}^{3+}$  peaks extracted and the corresponding assignments. Fig. 3b shows a magnification around the  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_0$  transition for the  $\text{AlN}$ -1200 sample and reveals that at least two non-equivalent active sites (590.2 nm – shoulder and 590.4 nm) are formed. The possible presence of a third  $\text{Eu}^{3+}$  peak (broad peak at 591.8 nm) cannot be discarded.

The evolution of the integrated CL intensity of the  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  transition as a function of  $\text{AlN}$  nominal content is represented in Fig. 3c. For  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1200 ( $0 < x < 1$ ), the intensity is similar independently on  $x$ , while for  $x = 1$ , it increases by an order of magnitude. This may be explained by the higher intensity ratio  $I_{{}^5\text{D}_0 \rightarrow {}^7\text{F}_2} / I_{{}^5\text{D}_1 \rightarrow {}^7\text{F}_1}$  found for  $\text{AlN}$ -1200, suggesting a higher radiative recombination probability from the excited state  ${}^5\text{D}_0$  instead of  ${}^5\text{D}_1$  for the  $\text{Eu}^{3+}$ -implanted  $\text{AlN}$  NWs. Since luminescence properties are affected by the host's crystalline quality, a higher  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  CL intensity for  $\text{AlN}$ -1200 might indicate a lower structural damage inflicted by ion implantation. Furthermore, when comparing the integrated intensity of the  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  transition for  $\text{AlN}$ -1200 and  $\text{AlN}$ -1000 (also included in Fig. 3c), the value is doubled for the highest annealing temperature. This result demonstrates that the annealing temperature is a crucial parameter for the optimization of the  $\text{Eu}^{3+}$  emission and the efficient recovery of the as-grown crystalline properties.

In order to localize the  $\text{Eu}^{3+}$  emission in the NWs with the highest CL intensity, nano-CL was performed on dispersed  $\text{AlN}$ -1200 NWs. Fig. 4a shows the SEM image of the dispersed NWs (Fig. 4a) consisting of two coalesced NWs at the  $\text{AlN}$  top-section. Nano-CL intensity maps integrated at distinct central energies 4.722 eV (262 nm, Fig. 4b), 3.404 eV (364 nm, Fig. 4c), and 1.989 eV (624 nm, Fig. 4d) are also shown. The choice of the central energy of 4.722 eV comes directly from the  $\text{AlN}$ -1200 CL spectrum (Fig. S6 in Supplementary Information), which cannot

be associated with AlN near band edge (NBE) expected to be  $> 6$  eV, but with defects emission from AlN [85,86]. In turn, 3.404 eV corresponds to the NBE emission from GaN NW template. The NW heterostructure consisting of an AlN NW top-section and GaN NW template is clearly resolved in Fig. 4. In this analysis, it is important to mention that the spatial localization of recombination processes is dependent on the carrier diffusion length, which is about 70 nm for electrons in GaN NWs [87]. Fig. 4d indicates that the distribution of optically active  $\text{Eu}^{3+}$  centers is preponderant in two regions: AlN top-section and GaN/AlN interface, with the latter revealing the most intense  $\text{Eu}^{3+}$  emission at 1.989 eV.

As mentioned above, when incorporated in III-N hosts,  $\text{Eu}^{3+}$  ions tend to occupy cation substitutional, or near-substitutional, lattice sites, which favor the achievement of the trivalent charge state  $\text{Eu}^{3+}$ . Depending on the surrounding environment, two dominant optically active  $\text{Eu}^{3+}$  centers are observed in GaN hosts and are usually identified as Eu1 and Eu2 [30,88]. Mitchell *et al.* ascribed Eu1 (at higher energy) and Eu2 (at lower energy) to complex defects involving substitutional Ga sites ( $\text{Eu}_{\text{Ga}}$ ) and different next-neighbors in the crystalline lattice: nitrogen vacancies ( $V_{\text{N}}$ ) and cation vacancies ( $V_{\text{Ga}}$ ), respectively [89]. Theoretical studies revealed that these complexes have a stable configuration [90]. Additionally, it was demonstrated that electrical excitation of GaN:Eu devices favors the excitation of the Eu2 center (also labeled as OMPVE7) [35,91]. In fact, ion implantation might induce a high concentration of cation and nitrogen vacancies ( $V_{\text{III}}$  and  $V_{\text{N}}$ ), leading to the incorporation of  $\text{Eu}^{3+}$  ions in cation sites and to the formation of complexes with the generated point defects, which can provide energetic levels within the host's bandgap able to assist  $\text{Eu}^{3+}$  excitation. Recently, two dominant optically active  $\text{Eu}^{3+}$  centers (Eu1 and Eu2) were identified by photoluminescence in  $\text{Eu}^{3+}$ -implanted AlN NWs, with similar spectral shape to the corresponding Eu1 and Eu2 centers in GaN [26].

Warnick *et al.* determined theoretically the migration barriers for  $V_{\text{III}}$  and  $V_{\text{N}}$  in GaN and  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$ . In strain-free GaN, the energetic barriers of  $V_{\text{N}}$  and  $V_{\text{Ga}}$  were found to be similar, while in  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$ , the migration energy was found higher for  $V_{\text{N}}$  and lower for  $V_{\text{III}}$  compared to GaN. When  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$  samples are subjected to strain, the migration barriers for  $V_{\text{III}}$  are reduced, while  $V_{\text{N}}$  barriers keep the same value. Since the diffusion rate increases exponentially with the decrease of the migration barrier energy, the mobility of  $V_{\text{III}}$  is higher in strained regions, as is the case of the GaN/AlN interface [92]. Moreover, the annealing process necessary to optically activate the  $\text{Eu}^{3+}$  ions leads to a reorganization of the crystalline lattice, allowing not only the a reduction of the density of defects introduced by implantation but also the diffusion to energetically favorable locations [92]. Therefore, as indicated in Fig. 4d, the regions corresponding to AlN NW top-section and GaN/AlN interface can affect the optical properties of the  $\text{Eu}^{3+}$  ions differently, because of the different surrounding environment, the lattice and defects dynamics after implantation and annealing.

### 3.2.2. Above and below bandgap excitation: the ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$ transition

Fig. 5a displays a magnification of the 5 K CL and 14 K high-resolution (HR) PL spectra for  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1200 ( $x > 0$ ) NW ensembles around the most intense transition,  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$ . For the annealing temperature of 1000 °C, HR PL spectra for  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1000 NW ensembles is shown in Fig. 5b. On the one hand, CL enables above bandgap excitation of  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  for which the  $\text{Eu}^{3+}$  emission is possible through all the excitation mechanisms. On the other hand, PL obtained using 3.8 eV laser excitation, *i. e.* below  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  ( $x \geq 0.3$  in our samples) bandgap energy,  $\text{Eu}^{3+}$  emission can be achieved through direct and indirect processes. In a direct process, the excitation

is resonant with an excited electronic state of the ion ( $4f$  or  $5d$ ), while indirect processes occur by energy-transfer (ET) from the host to the  $4f$  electrons subsystem through non-radiative Auger-type mechanisms. These mechanisms have been discussed by different authors for GaN-based layered structures [18,93–95]. Following Lozykowski *et al.* [95], the most probable excitation mechanism occurs through ET from excitons bound to isovalent traps to  $4f$  electrons created by RE ions. Fig. 5c describes the defect-trap mediated ET model for the excitation/emission of  $\text{Eu}^{3+}$  ions in  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NWs. For above bandgap excitation (GaN NWs), two mechanisms are possible: *i*) creation of excitons in the host followed by their localization at the defect-traps; and *ii*) localization of one of the carriers at the defect-traps, and subsequently the localization of the other. In both cases, a non-radiative recombination and ET leaves  $4f$  electrons in an excited state, resulting in intra- $4f$  emission. For below bandgap excitation ( $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NWs,  $x \geq 0.3$ ) only the second mechanism is allowed.

As mentioned above for AlN-1200, at least two  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_0$  transitions were observed (Fig. 3b), indicating that at least two optically active centers are involved, as also confirmed by the different relative intensities of the most intense  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  three peaks in PL and CL (Fig. 5a). Moreover, it is possible to observe that PL favors Eu2 in comparison to CL, *i.e.* the ratio  $I_{\text{Eu}2}/I_{\text{Eu}1}$  is higher in the PL spectrum. This agrees with the nano-CL results since the central energy of 1.989 eV corresponds to the Eu2 center and its optical contribution is higher at the interface GaN/AlN (Fig. 4d). We remind that the CL response of the NW ensemble comes predominantly from the NWs top-section, while the PL experiments provide information not only from the NWs top-section but also from the GaN/ $\text{Al}_x\text{Ga}_{1-x}\text{N}$  interface and the GaN NW template.

For  $\text{Al}_{0.5}\text{Ga}_{0.5}\text{N}$ -1200, CL and PL spectra show nearly identical luminescence response, while for  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1200 (with  $x = 0.3$  and  $0.75$ ), these spectra exhibit similar shape with slightly different peaks' resolution and energy, which can be associated with compositional fluctuations in the NWs and different resolution of both techniques. Alloy disorder and/or compositional fluctuations felt by the RE ions also contribute to the increase of the broadening of the  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  emission [18,51,67]. A tentative representation of the spectral shape evolution with the AlN nominal content is shown in Fig. 5d. Here, the lower (higher) energy center Eu1 (Eu2) contribution is represented with luminescence intensities of  $I_{\text{Eu}1}$  ( $I_{\text{Eu}2}$ ). When considering such representation, the ratio  $I_{\text{Eu}2}/I_{\text{Eu}1}$  is found to decrease when  $x$  increases. For the annealing temperature of 1200 °C, it is possible to tune the  $\text{Eu}^{3+}$  optical active center by changing the nominal AlN content, which verifies that the relative abundance of the  $\text{Eu}^{3+}$  centers is extremely sensitive to the surrounding environment.

The annealing temperature is a parameter of utmost importance for device development and performance because it can affect the contribution of the different  $\text{Eu}^{3+}$  centers [30], as defined here by the ratio  $I_{\text{Eu}2}/I_{\text{Eu}1}$ . Furthermore, it was demonstrated that the electrical properties of  $\text{Eu}^{3+}$ -implanted/annealed diode structure degrade when subjected to high temperature and high pressure annealing at 1400 °C [47]. For GaN: $\text{Eu}^{3+}$  layers capped with AlN, it was reported that the Eu2 center is predominant for annealing temperatures of 1300 °C [88], while an opposite behavior was recently reported in AlN: $\text{Eu}^{3+}$  NWs showing that Eu2 is favored for rapid thermal annealing treatments at 1000 °C [26]. Therefore, the evaluation of the impact of the annealing temperature in the  $\text{Eu}^{3+}$  emission is needed. To do so, the behavior of the  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  transition for the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1000 NW ensembles is explored as a function of  $x$  and its behavior compared to that of the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1200 samples. For the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1000 (Fig. 5b), the behavior of the  $\text{Eu}^{3+}$  active center, shows an opposite trend when compared to the one observed for  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1200 NWs. In fact, while GaN-1000 and  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$ -1000 show a higher contribution of the Eu1 center

(i.e.  $I_{\text{Eu}2} < I_{\text{Eu}1}$ ), the contribution of the Eu2 increases for AlN-1000, which indicates that the ratio  $I_{\text{Eu}2}/I_{\text{Eu}1}$  increases when  $x$  increases. Note that for  $\text{Al}_{0.5}\text{Ga}_{0.5}\text{N}$ -1000 and  $\text{Al}_{0.75}\text{Ga}_{0.25}\text{N}$ -1000 samples, the emission exhibits a shoulder at the high-energy side (grey rectangle in Fig. 5b) that could be associated with GaN NW template (detailed in section S3.3 in Supplementary Information).

Moreover, as seen in Fig. 6, a redshift of the  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  transition components is observed when  $x$  increases, in agreement with reports on  $\text{Al}_x\text{Ga}_{1-x}\text{N}:\text{Eu}^{3+}$  layers [18,29,51]. This shift is due to the higher expansion of the  $4f$ -orbitals in AlN caused by the larger electron affinity of Al compared to Ga [51].

### 3.2.3. Below bandgap excitation: temperature dependence of ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$ transition

Temperature-dependent HR PL measurements at 14 K and 300 K are shown for  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$ -(1000, 1200) and AlN-(1000, 1200) samples (Figs. 7a and 7b, respectively). The evolution for all the temperatures and all the implanted samples are shown in Fig. S8 (Supplementary Information). The optical center Eu1 (Eu2) is highlighted with violet (orange) dashed line. When increasing the temperature from 14 K to 300 K, the PL spectra become less resolved due to the phonon broadening, rendering the distinction of the optically active  $\text{Eu}^{3+}$  centers even more difficult. In addition, competitive thermally-induced non-radiative processes cause an overall decrease of the  $\text{Eu}^{3+}$  intensity (Figs. 7a, 7b and S8). Such thermal quenching results from the ET to either radiative and/or non-radiative defects in the vicinity of each  $\text{Eu}^{3+}$  center. Moreover, for AlN hosts, the ratio  $I_{\text{Eu}2}/I_{\text{Eu}1}$  decreases with temperature, indicating that Eu1 is thermally more stable. For the ternaries, the peak broadening renders the identification of such behavior more difficult. The different quenching behaviors of  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1000 and  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1200 NW ensembles is better seen in Fig. 7c which represents the evolution of the ratio of the  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  integrated PL intensity at 300 K and 14 K,  $I_{300\text{ K}}/I_{14\text{ K}}$ , versus  $x$ . For the annealing temperature of 1000 °C, we note that the thermal quenching is lower for the ternary alloys when compared to GaN-1000 NWs and characterized by  $I_{300\text{ K}}/I_{14\text{ K}} \sim 50\%$ . The fact that a strong quenching is present in the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1000 (with  $x > 0.5$ ) NWs can be tentatively associated with the predominance of the Eu2 center, which is highly sensitive to temperature in AlN hosts. However, the thermal quenching is reduced for  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1200 NWs when the AlN nominal content increases, reaching its maximum for AlN-1200 NWs and characterized by the highest  $I_{300\text{ K}}/I_{14\text{ K}}$  value of  $\sim 80\%$ . This effect can be explained by the higher contribution of the thermally stable Eu1 center with respect to the Eu2 center for the annealing temperature of 1200 °C.

## 4. CONCLUSIONS

To summarize and conclude, the incorporation of  $\text{Eu}^{3+}$  ions in implanted and annealed  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NWs was studied by  $\mu$ -Raman and nano-CL, being observed that the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  top-section is the most affected region. A partial recovery of the implantation-induced damage was obtained after RTA as confirmed by  $\mu$ -Raman. RTA also provided the optical activation of the  $\text{Eu}^{3+}$  ions, being possible to be excited with above (CL) and below (PL) bandgap energies. The  ${}^5\text{D}_0 \rightarrow {}^7\text{F}_2$  transition is the most intense and its energy redshifts from GaN to AlN NWs. It was found that its spectral shape depends on two effects: AlN nominal content and annealing temperature, indicating that such behavior is attributed to at least two optically active  $\text{Eu}^{3+}$  centers (Eu1 and Eu2). In this sense, the ratio  $I_{\text{Eu}2}/I_{\text{Eu}1}$  increases with  $x$  for the lower annealing temperature while the opposite trend is

verified for the higher annealing temperature. In addition, temperature-dependent PL evidenced a thermal quenching of the  $\text{Eu}^{3+}$  luminescence, for which AlN-1200 NWs show the lowest quenching ( $I_{300\text{ K}}/I_{14\text{ K}}$  value of  $\sim 80\%$ ) compared to the rest of the samples. Thus, the interplay between the AlN nominal content and the annealing temperature can be used to tune the optical contribution of  $\text{Eu}^{3+}$  centers in  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NWs. This significant result can help to optimize the optical activation of  $\text{Eu}^{3+}$  centers in III-N semiconductors since some of them are favored under electrical excitation for red light-emitting diodes. Therefore, it can be a viable route to improve the efficiency of these devices.

## **ASSOCIATED CONTENT**

Supplementary Material Complementary experiments including scanning electron microscopy images and  $\mu$ -Raman response of  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NWs, deconvolution procedure of  $\text{E}_2^{\text{H}}$  (GaN) and  $\text{E}_2^{\text{H}}$  (GaN-like) phonons, determination of hydrostatic strain in GaN and AlN NWs, CL and PL response of the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NWs, peak assignment of the  $\text{Eu}^{3+}$  intra- $4f^6$  transitions, and heterogeneity of the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  NWs and its impact on the  $\text{Eu}^{3+}$  emission.

## **CONFLICT OF INTEREST**

The authors declare that they have no known financial/commercial Conflict of Interest.

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## FIGURE CAPTIONS

**Fig. 1:** (a) Schematic representation of the  $\text{Eu}^{3+}$ -implantation geometry. Representative SEM images of the  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$ -asgr NWs in (b) side-view and (c) top-view. Vertical side arrows correspond to the different NW sections.

**Fig. 2:** (a) RT normalized  $\mu$ -Raman spectra of the  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$ -asgr,  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$ -asimp,  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$ -1000, and  $\text{Al}_{0.3}\text{Ga}_{0.7}\text{N}$ -1200 NW ensembles using laser line excitation of 441.6 nm. The spectra were normalized to the maximum of the  $\text{E}_{2\text{H}}$  (GaN) phonon intensity. The feature arising from the Si (111) substrate is indicated by (\*). The calculated phonon DOS for GaN and the disorder-induced Raman spectrum for Er-implanted AlN layers are included for comparison [74]. Two rectangles (blue and orange) is inserted to show the difference in the asymmetry for the as-grown and as-implanted samples. (b)  $\text{E}_{2\text{H}}$  (GaN) and  $\text{E}_{2\text{H}}$  (GaN-like) phonon frequency as a function of  $x$ . The dashed horizontal line corresponds to the relaxed value for the  $\text{E}_{2\text{H}}$  (GaN) phonon ( $567.6\text{ cm}^{-1}$  [74]). The evolution of the  $\text{E}_{2\text{H}}$  (GaN-like) with AlN content for  $\text{Al}_x\text{Ga}_{1-x}\text{N}$  layers, taken from Davydov *et al.*, is also included for comparison [73].

**Fig. 3:** (a) 5 K CL spectra of  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1200 NW ensembles, indicating the different  $\text{Eu}^{3+}$  intra- $4f_6$  transitions. (b) 5 K CL spectrum of AlN-1200 magnified around the  $5\text{D}_0 \rightarrow 7\text{F}_0$  transition. (c)  $5\text{D}_0 \rightarrow 7\text{F}_2$  integrated CL intensity (in logarithmic scale) as a function of the AlN nominal content,  $x$ . The integrated CL intensity of the AlN-1000 NWs is also included.

**Fig. 4:** 5 K nano-CL intensity maps of dispersed AlN-1200 NWs [corresponding SEM image in (a)] integrated at the central energies: (b) 4.722 eV (262 nm), (c) 3.404 eV (364 nm), and (d) 1.989 eV (624 nm).

**Fig. 5:** (a) 5 K CL and 14 K PL spectra around the most intense  $5\text{D}_0 \rightarrow 7\text{F}_2$  transition for the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1200 NW ensembles. (b) 14 K PL spectra around the most intense  $5\text{D}_0 \rightarrow 7\text{F}_2$  transition for the  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1000 NW ensembles. (c) Schematic representation of the  $\text{Eu}^{3+}$  excitation/emission model when using the 3.8 eV laser line: A - photoexcitation of electrons (holes) from the VB (CB) to the CB (VB); B - localization of the carriers at the defect-traps; C - recombination of the exciton formed at the defect-trap; D - energy-transfer (ET) to the  $\text{Eu}^{3+} 4f$  levels through Auger processes; E - excitation of the  $\text{Eu}^{3+}$  ion from the ground state ( $7\text{F}_0$ ) to an excited state ( $5\text{D}_1$ ), leading to  $\text{Eu}^{3+}$  emission. (d) Spectral shape representation of the  $5\text{D}_0 \rightarrow 7\text{F}_2$  transition when dominated by Eu1 (top) and Eu2 (bottom).

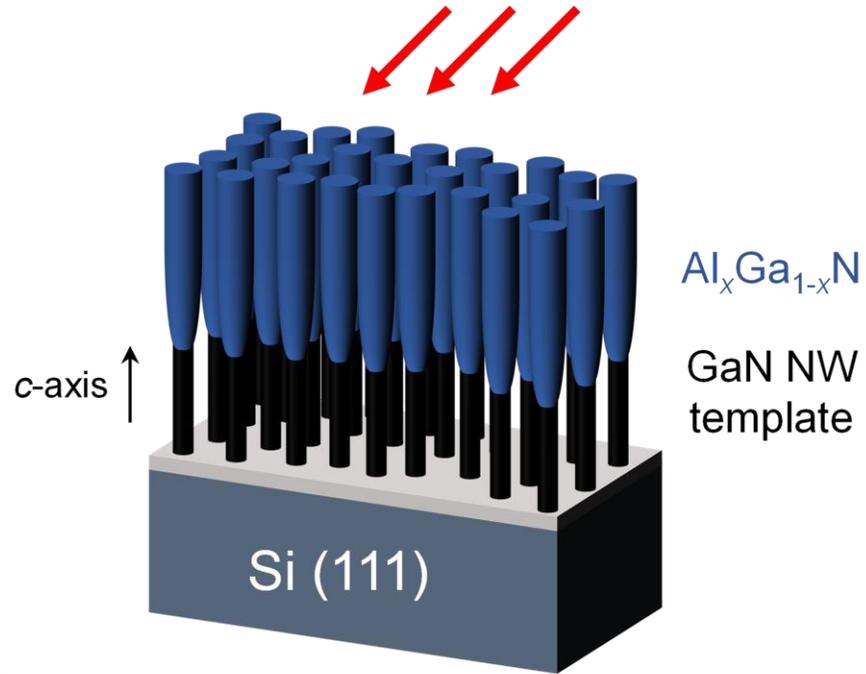
**Fig. 6:** Evolution of the  $5\text{D}_0 \rightarrow 7\text{F}_2$  resolved peaks as a function of  $x$ . For  $x = 0$ , the values were extracted from the 14 K PL spectrum of GaN-1000. Green stars correspond to the most intense  $5\text{D}_0 \rightarrow 7\text{F}_2$  peaks obtained by Gruber *et al.* for  $\text{Eu}^{3+}$ -doped  $w$ -AlN single crystal [17].

**Fig. 7:** 14 K and 300 K HR PL spectra around the  $5\text{D}_0 \rightarrow 7\text{F}_2$  transition for  $x = 0.3$  (a) and  $x = 1$  (b) for  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1000 NW ensembles (top figures) and  $\text{Al}_x\text{Ga}_{1-x}\text{N}$ -1000 NW ensembles (bottom figures). Vertical dashed lines indicate Eu1 (violet) and Eu2 (orange) centers. (c) Evolution of the  $I_{300\text{ K}}/I_{14\text{ K}}$  ratio with the Al nominal content for all the studied samples.

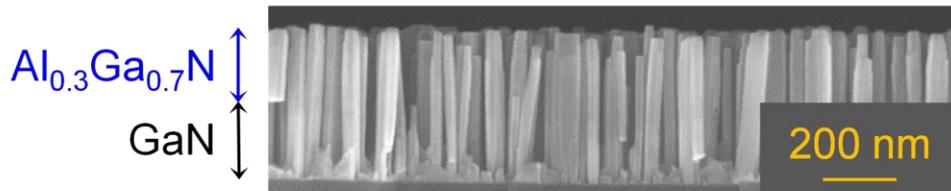
Figure 1

(a)

$\text{Eu}^{3+}$  ion beam at  $45^\circ$  tilt



(b)



(c)

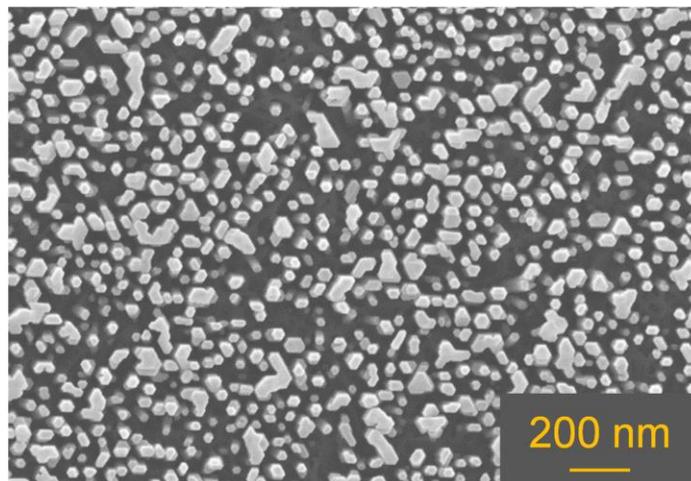
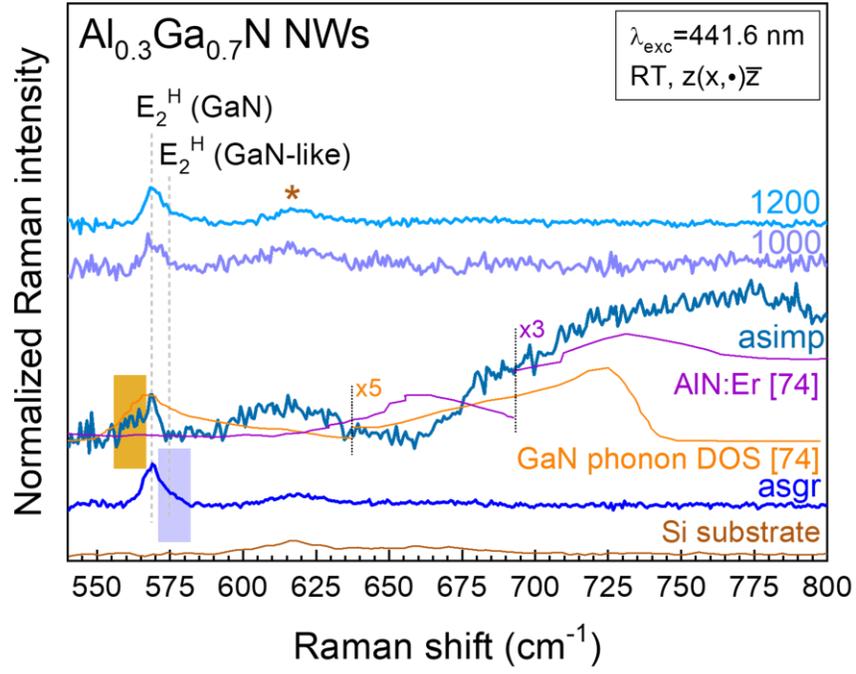


Figure 2

(a)



(b)

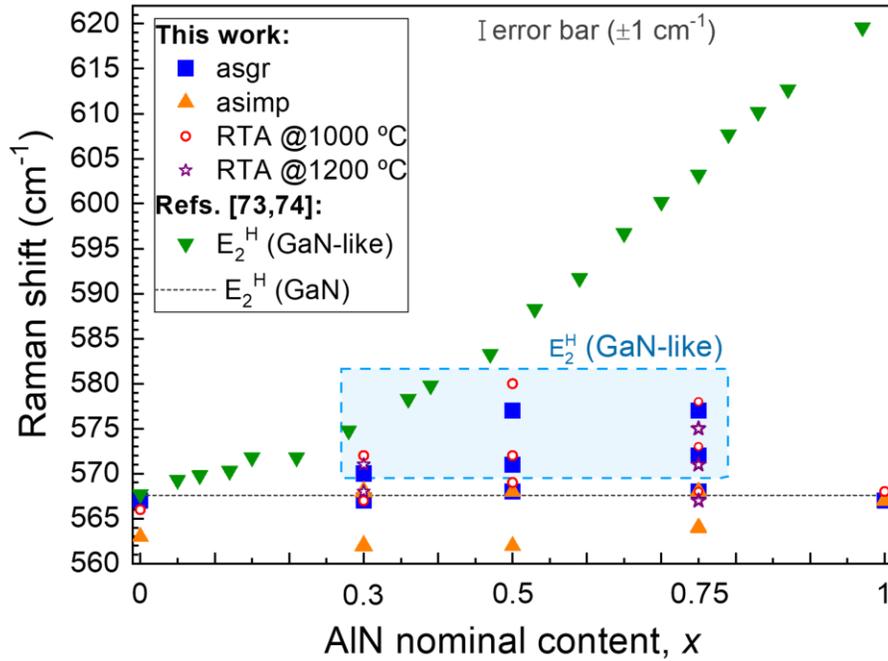


Figure 3

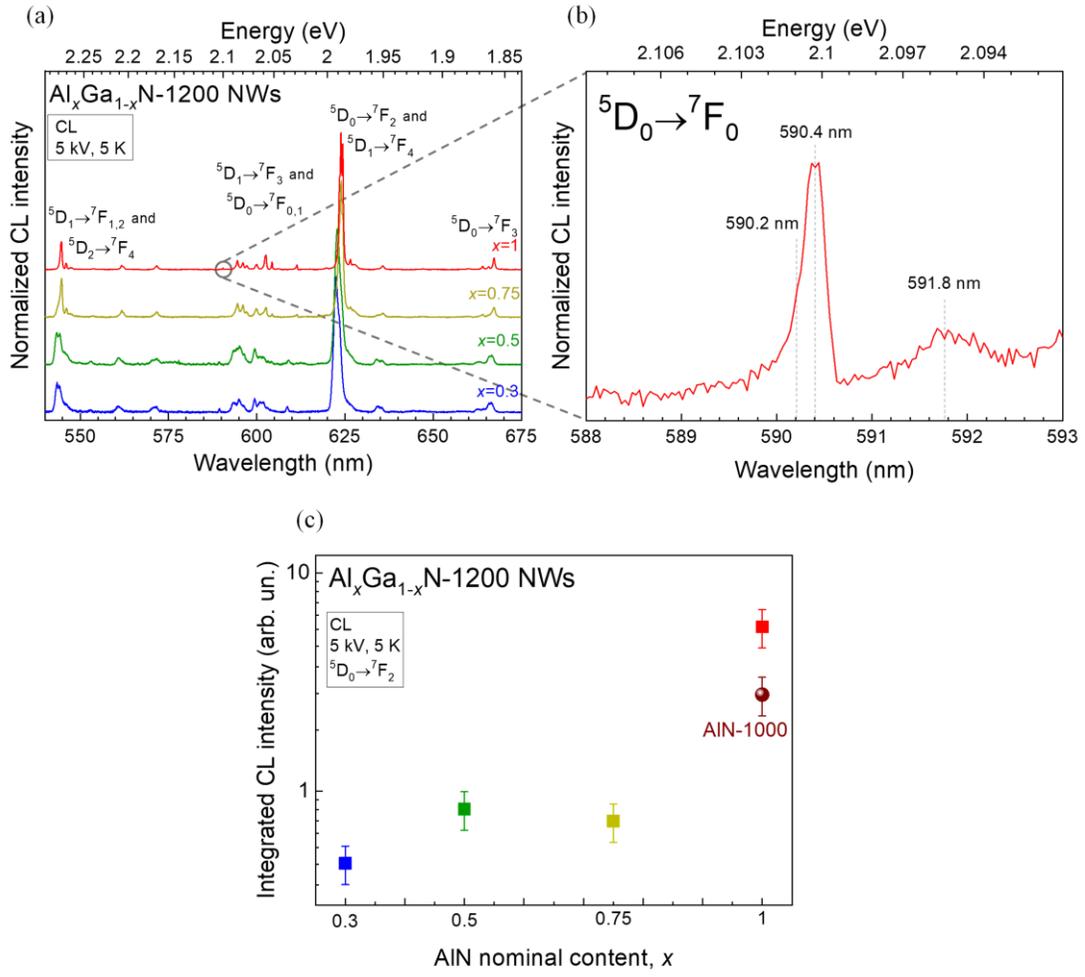


Figure 4

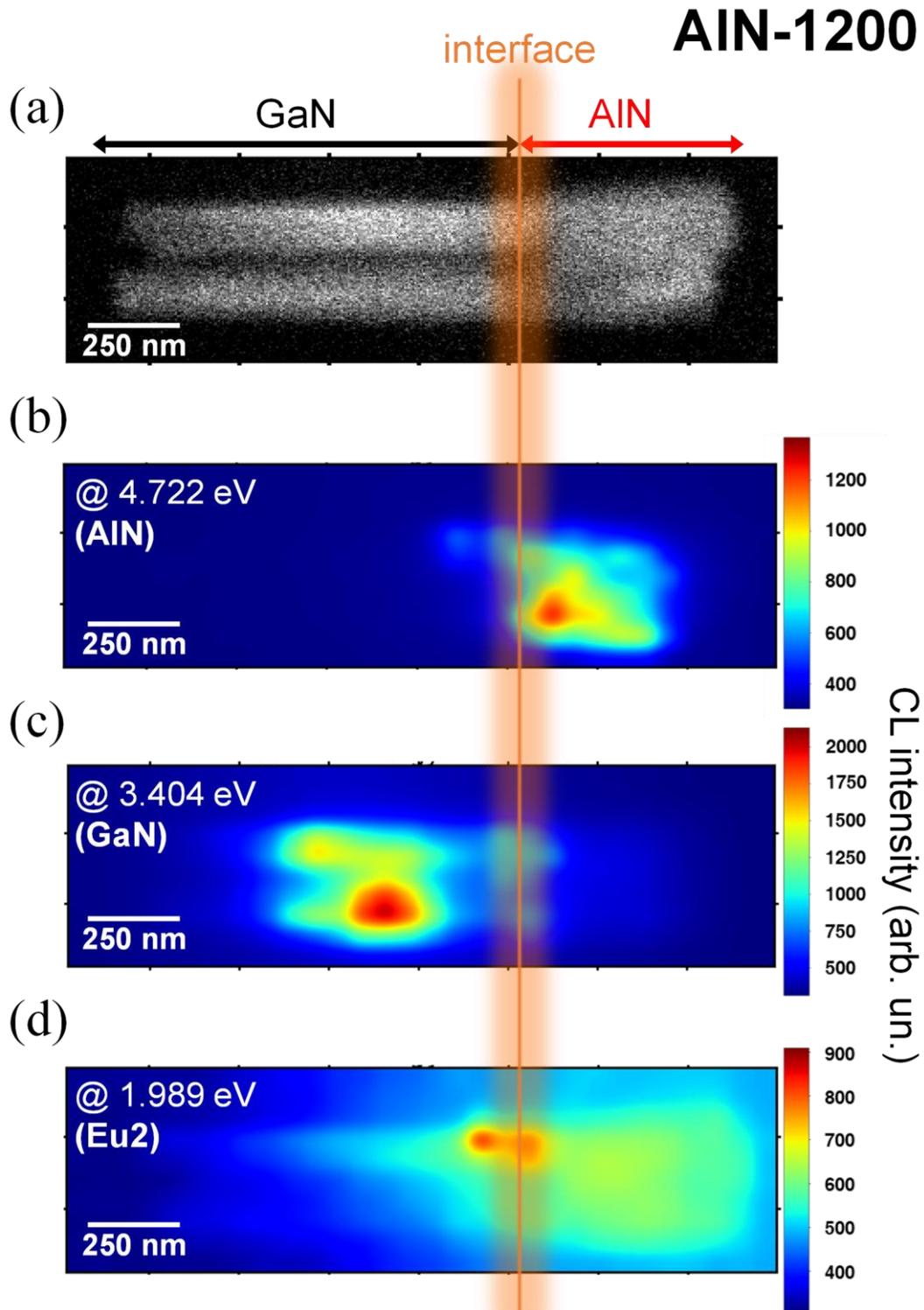


Figure 5

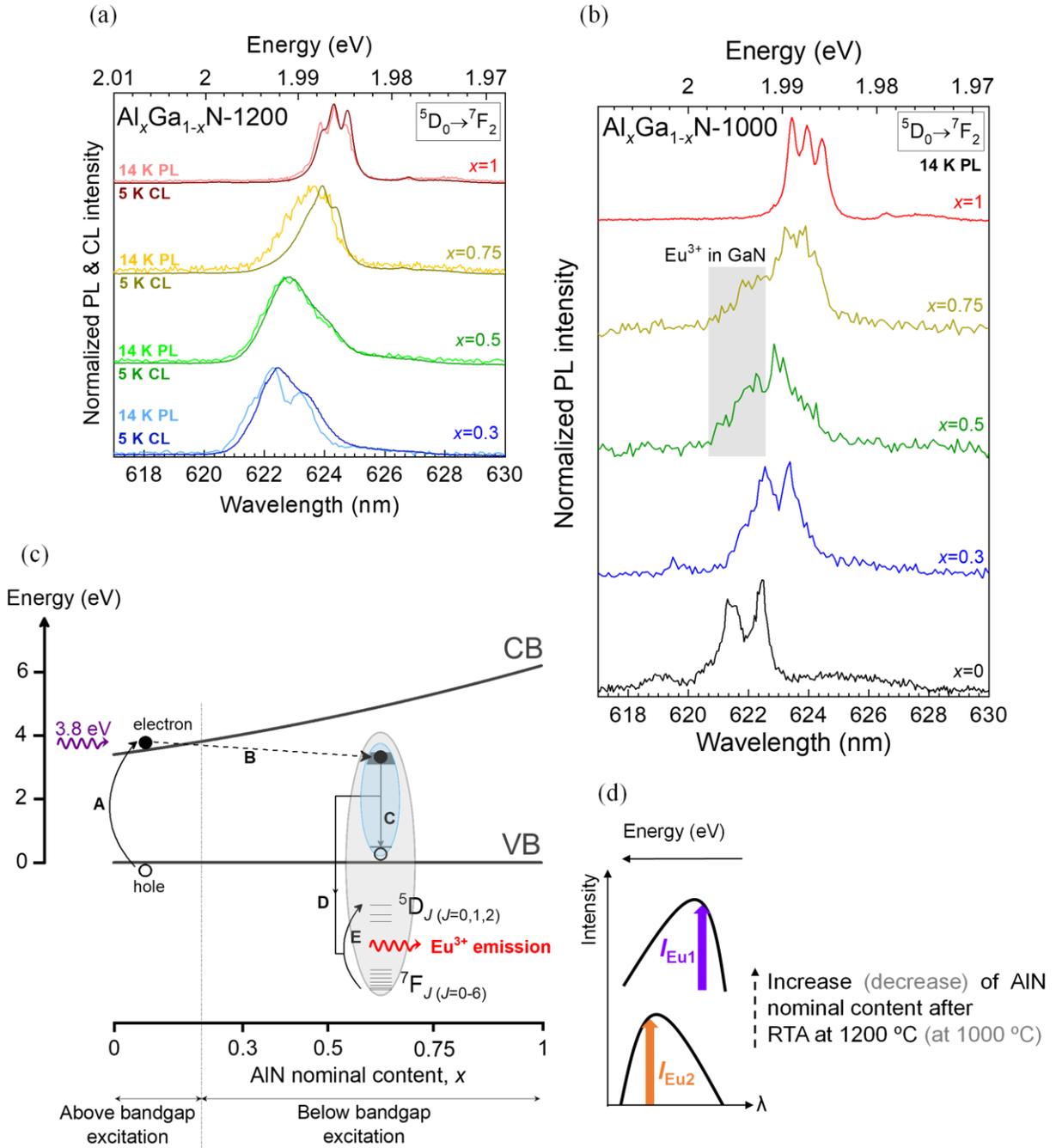


Figure 6

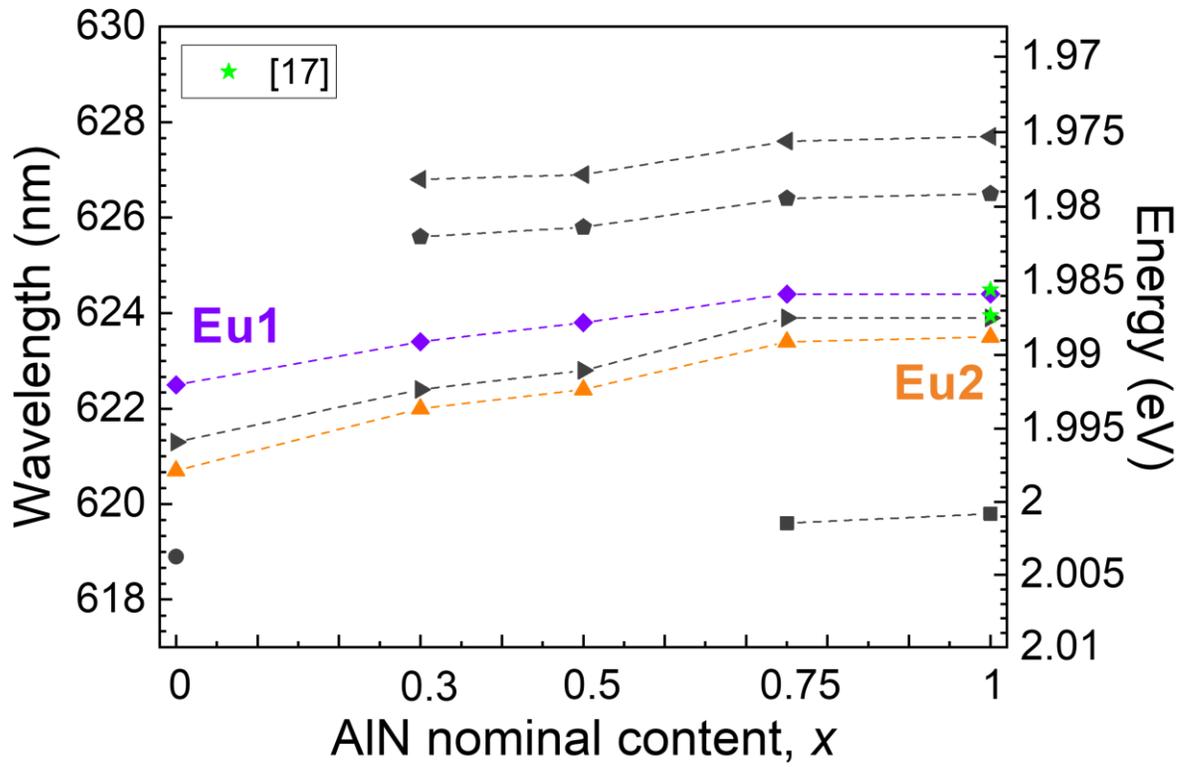


Figure 7

