Band alignment at GaN/single-layer WSe₂ interface

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ABSTRACT: We study the band discontinuity at the GaN/single-layer (SL) WSe₂ heterointerface. The GaN thin layer is epitaxially grown by molecular beam epitaxy on chemically vapor deposited SL-WSe₂/c-sapphire. We confirm that the WSe₂ was formed as an SL from structural and optical analyses using atomic force microscopy, scanning transmission electron microscopy, micro-Raman, absorbance, and micro-photoluminescence spectra. The determination of band offset parameters at the GaN/SL-WSe₂ heterojunction is obtained by high-resolution x-ray photoelectron spectroscopy, electron affinities and the electronic bandgap values of SL-WSe₂ and GaN. The valence band and conduction band offset values are determined to be 2.25 ± 0.15 and 0.80 ± 0.15 eV respectively, with type II band alignment. The band alignment parameters determined here provide a route towards the integration of group III nitride semiconducting materials with transition metal dichalcogenides (TMDs) for designing and modeling of their heterojunction based electronic and optoelectronic devices.

KEYWORDS: GaN, single layer WSe₂, 3D/2D heterojunction, HRXPS, band alignment, molecular beam epitaxy

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1. INTRODUCTION

Group-III nitrides, GaN as a primary material, are well demonstrated with enormous applications in high efficiency electronic and optoelectronics devices such as high electron mobility transistors, light emitting diodes, and laser diodes.¹⁻⁴ Recently, group VI transition metal dichalcogenides (TMDs) in the form of MX_2 has emerged as a novel atomic layered material system with challenging thermoelectric, electronic and optoelectronic properties.⁵⁻⁸ Among the TMDs, tungsten diselenide (WSe_2) in a single layer form is of potential interest for such devices owing to its direct bandgap of ≈ 1.65 eV and prominent transport properties.^{9,10} GaN has small lattice mismatch of $\approx 3.2\%$, $\approx 1.0\%$ and $\approx 0.8\%$ with majorly studied TMDs such as WSe_2 , tungsten disulfide (WS_2) and molybdenum disulfide (MoS_2)¹¹, respectively, in comparison to commonly used substrates such as sapphire, SiC and Si.^{12,13} Moreover, recent advances in the integration of 2D-layered materials with wide band gap group III nitride semiconductors is exciting due to their variety of applications in high current tunnel diodes.^{14,15} Several efforts were made to grow GaN on closely lattice matched TMDs. Yamada *et al.* presented the growth of GaN on bulk MoS_2 by plasma-enhanced molecular beam epitaxy¹⁶. There were recent attempts to grow GaN on large area MoS_2 and WS_2 layers,^{11,17} layered MoS_2 on GaN epilayers¹⁸ by chemical vapor deposition (CVD) growth techniques and layer transferred p- MoS_2 on GaN¹⁴. Though the deposition of large area single-layer WSe_2 on sapphire exhibiting direct bandgap has been demonstrated,^{9,19} the growth of GaN on such large area monolayered WSe_2 has not yet been explored.

So far, the band offset parameters were determined for either group III nitrides on various other semiconducting materials or solely TMDs based dissimilar heterostructures using X-ray photoemission spectroscopic core-levels evaluated with respect to the valence band maximum and bandgap studies. Though, the band offset parameters (junction type: valence band offset- ΔE_v & conduction band offset- ΔE_c) are measured for various heterojunctions in the literature.²⁰⁻²⁵ Recently we have reported the growth of GaN on SL- MoS_2 and the band alignment parameters (Type-II: 1.86 ± 0.08 & 0.56 ± 0.10 eV) for GaN/SL- MoS_2 hetero-interface.²⁶ The band offset parameters (valence band offset (VBO)- ΔE_v and conduction band offset (CBO)- ΔE_c) and type of junction by HRXPS for epitaxially formed GaN/SL- WSe_2 hetero-interface is yet to be experimentally investigated. The determination of band offset parameters is required to study the group III nitride/TMDs heterojunction based devices. These hybrid heterojunctions, exhibiting type I and II band alignments, can be utilized as TMD quantum well based light emitting devices

and the tunnel diodes, respectively.^{6,15} Thus, the growth of GaN/SL-WSe₂ and the determination of band offsets are necessary to provide a route towards the integration of group III nitrides with TMDs for designing the electronic and optoelectronic devices.

In this report, in order to study the band discontinuity at the GaN/SL-WSe₂ heterointerface, thin GaN layers were epitaxially grown on a CVD prepared SL-WSe₂. The sustainability of SL-WSe₂ during GaN growth is confirmed by the atomic force microscopy (AFM), micro-photoluminescence (μ PL) and Raman spectroscopies. In addition, the band offset parameters for GaN/SL-WSe₂ heterojunction were determined using high-resolution x-ray photoelectron spectroscopy (HRXPS) and electronic band gap values of respective constituent layers in the heterojunction.

2. EXPERIMENTAL SECTION

The growth experiments of GaN on WSe₂/c-sapphire substrates were carried out by Veeco 930 Gen plasma assisted molecular beam epitaxy (PAMBE) system at substrate temperature of 500 °C. The large area WSe₂ SLs were prepared on c-sapphire substrates using CVD and the details can be found elsewhere¹⁹. The ion and a cryo pumps were employed to attain a base pressure of 3×10^{-11} Torr and oxygen partial pressure $< 10^{-11}$ Torr, as obtained by a residual gas analyzer (RGA). The substrates were thermally outgassed in introduction chamber at 200 °C for 30 mins, further cleaning was carried out in preparation chamber at 300 °C for 60 mins and in the growth chamber at 400 °C for 30 mins. For GaN growth, the nitrogen plasma source was operated with the flow rate of 1 standard cubic centimeter per minute (sccm) and RF power of 300 W and Ga metal was evaporated by standard Knudsen cell with beam equivalent pressure (BEP) value of 2.10×10^{-8} Torr. Prior to this, thin AlN buffer layer was grown. The corresponding chamber pressure was $\approx 2.5 \times 10^{-5}$ Torr. The thickness of the GaN epilayer (sample C) is measured to be ≈ 500 nm. The thickness of GaN layer in sample B was estimated to be 6 ± 1 nm from growth rate calibrations. The Agilent technologies 5400 atomic force microscopy was used in tapping mode to acquire the surface morphology of the samples. Structural quality of GaN epilayer was investigated by CuK α High Resolution X-ray Diffraction (HRXRD) having four-bounce Ge(022) monochromator. High-angle annular dark field scanning transmission electron microscopy (HAADF-STEM) was utilized by operating a probe-corrected FEI Titan at an acceleration voltage of 300 kV. A cross-sectional TEM specimen of sample B was prepared by using FEI's Helios Dual Beam focused ion beam

(FIB)/SEM equipped with an Omniprobe. Electron energy loss spectroscopy (EELS) acquisition was performed by using Gatan's GIF Quantum of Model 966. Using Horiba Aramis room temperature (RT) μ PL, the emission of GaN and WSe₂ layers was deduced with excitation lines of He-Cd laser of 325 nm and He-Ne laser of 633 nm, respectively. Absorbance spectra were acquired with a Shimadzu UV3600 spectrophotometer equipped with integrating sphere. The high-resolution XPS measurements were carried out using a Kratos Axis Ultra DLD spectrometer equipped with a monochromatic Al K α X-ray source ($h\nu = 1486.6$ eV) operating at 150 W, a multi-channel plate and delay line detector under a vacuum of $\sim 10^{-9}$ mbar. The high-resolution spectra were collected within the limits of spatial resolution at a fixed analyzer pass energy of 20 eV. In order to eliminate the shifts in the HRXPS spectra associated with the surface charging effect, the measurements were acquired both with and without electron beam charge compensation. For both cases, no changes were observed in the determined band alignment. In absence of electron beam charge, the surface of samples was electrically connected with a clean copper (Cu) foil. The remnant binding energy shifts were referenced to the adventitious carbon (C 1s) signal.^{27–29} In this study, CVD grown SL-WSe₂ (sample A), MBE grown GaN on SL-WSe₂ (sample B) and GaN epilayer (sample C) were used to determine the band offsets at GaN/SL-WSe₂ hetero-interface.

3. RESULTS AND DISCUSSION

To investigate the surface morphology, root mean square (rms) roughness of GaN/WSe₂ (B) and GaN epilayer (C), and thickness of WSe₂ layer (A), AFM scans were carried out in tapping mode. Figures 1(a-c) show the AFM surface topography scans collected on samples C, A and B, respectively. The AFM images were obtained at different scales to view the fine features according to surface smoothness of the formed films. Color bars on top of the respective images in Figure 1 show height contrast of the features. From Figure 1(a), the root mean square (rms) surface roughness for GaN film is ≈ 2 nm. Moreover, AFM is one of the most commonly used technique to determine the number of monolayers of TMDs samples. The thickness of the large area WSe₂ layer is found to be ≈ 0.76 nm from the line profile as shown in Figure 1(b) for sample A, which is in agreement with the thickness of Se-W-Se single-layer.¹⁹ The inset to Figure 1(b) displays the optical microscopy image of large area WSe₂ layer. Figure 1(c) shows the AFM surface morphology for sample B exhibiting surface rms roughness of ≈ 4 nm which is higher than that of relaxed GaN epilayer (sample C) that results from the lattice mismatch between thin GaN and

WSe₂. Thus, AFM study confirms that the CVD deposited large area WSe₂ is in the form of a single layer. Further, HRXRD measurements were performed to investigate the crystalline structure and quality of the GaN epilayer (sample C). Figure 1(d) presents the 2 θ - ω scans, which were obtained on the symmetric (on-axis) and asymmetric (off-axis) planes of sample C. Asymmetric planes were studied in skew symmetric geometry.⁴ Figure 1(d) displays c-oriented peaks: at GaN(0002) at 34.52°, GaN(0004) at 73.20° along with the buffer layer AlN(0002) at 35.89° and Al₂O₃ substrate (0006) at 41.45°, respectively. The asymmetric reflection shows GaN(10 $\bar{1}$ 1) peak at 36.81° GaN(20 $\bar{2}$ 2) at 78.77° along with the buffer layer peak AlN(10 $\bar{1}$ 1) at 38.50° and substrate peak Al₂O₃(11 $\bar{2}$ 3) at 43.08°, respectively. The respective inset shows the ω scans for on-axis GaN(0002) and off-axis GaN(10 $\bar{1}$ 1) reflections, with broadening values of $\approx$ 350 and $\approx$ 980 arc-sec, respectively. Thus, from HRXRD analysis, sample C possess single crystalline wurtzite structure grown along c-axis exhibiting moderate crystal quality.

HAADF-STEM studies were performed on sample B (GaN/WSe₂/c-sapphire) to confirm the existence and the thickness of GaN and WSe₂ layers. Figure 2(a) shows the cross sectional HAADF-STEM image recorded on sample B which is prepared by FIB. Prior to this, the Ti metal layer having thickness of $\approx$ 500 nm was deposited on sample B by e-beam evaporation method to avoid any physical damage of thin GaN and WSe₂ layers at the heterojunction caused by focused ion beams while the sample preparation was done for STEM studies. This reveals that the thickness of GaN/WSe₂ heterojunction is $\approx$ 7 nm. Figure 2(b) shows the magnified STEM image of Figure 2(a) obtained across the GaN/WSe₂/c-sapphire heterojunction (sample B). STEM-EELS imaging data sets were acquired to generate the elemental maps of W, Se, Ga and Al elements by employing W-M (1809 eV), Se-L (1436 eV), Ga-L (1117 eV), and Al-K (1560 eV) EELS edges, respectively. The elements W, Se, Ga and Al across the junction are, respectively, represented by Blue, Green, Red, and Yellow colored pixels. Figure 2(c) depicts the elemental mapping of heterojunction showing the existence of both W and Se atoms at the interface. A STEM-EDS line profile was generated to re-confirm the presence of SL-WSe₂ that is sandwiched between sapphire substrate and GaN thin film. Figure 2(d) represents the EDS line profiles obtained for elements Al, Ga, W and Se across the junction. The intensity of line profiles for W and Se atoms increases in the region of SL-WSe₂ whereas the line profiles for Ga and Al elements exhibit high intensity in the regions of overgrown GaN and underlying sapphire substrates, respectively, as shown in Figure 2(d). Therefore, it is

noted that the line profiles in Figure 2(d) were in accordance with the respective elemental color mapping as depicted in Figure 2(c). Thus, Figures 2 (b-d) depict the existence of WSe₂ in the form of single layer having thickness of ≈ 0.78 nm which is in corroboration with AFM analysis. Overall, the HAADF-STEM analysis is a clear evidence of the formation of GaN/SL-WSe₂ heterojunction.

Micro-Raman spectroscopy was used to investigate the optical quality as well as to ensure that the CVD grown WSe₂ is an SL. Figure 3(a-d) shows the Raman spectra for WSe₂/c-sapphire, GaN/WSe₂/c-sapphire and GaN/c-sapphire. In order to obtain the enhanced Raman signal from GaN and WSe₂ layers, the measurements were carried out with excitation lines of 325 and 633 nm by which signal can overcome the reduced resonant excitation effect.^{26,30} Thus, both the excitation sources are required to collect the high intensity phonon modes from the individual layer in GaN/SL-WSe₂ heterojunction as described in Figure 3(b and c). The two characteristic peaks in Figure 3(a and b) observed at 248 cm⁻¹ and 259 cm⁻¹ are attributed to E_{2g}^1 and A_{1g} phonon modes.^{10,19} These modes correspond to the in plane vibration of W and Se atoms (E_{2g}^1) and out of plane vibration of Se atoms (A_{1g}) in WSe₂. The intensity ratio (I_A/I_E) of these modes is higher than the reported values which can be attributed to the variation in doping levels.³¹ The observed peaks at 222, 241, 377 and 398 cm⁻¹ stem from the WSe₂ layer which are consistent with the previous studies.^{10,32} The peaks represented by asterisks are from the sapphire substrate. The absence of a peak at ≈ 307 cm⁻¹ confirms that the formed WSe₂ is a single layer which is corroborated by AFM studies. These results also show the sustainability of WSe₂ layer during the growth of GaN in PAMBE. In consequence of the reduced resonant excitation effect²⁶, in Figure 3(b), E_2^H mode of GaN exhibits lower intensity than the E_{2g}^1 and A_{1g} phonon modes of WSe₂ while the sample B is excited with 633 nm line. In contrast, in Figure 3(c), the intensity of E_2^H mode for GaN is higher than that of the phonon modes (E_{2g}^1 and A_{1g}) of WSe₂ with 325 nm excitation. Further, Figure 3(c) shows a low intensity single peak for both E_{2g}^1 and A_{1g} phonon modes which is in good agreement with the literature.³³⁻³⁵ In Figure 3(b and c), the E_2^H peak of thin GaN grown on WSe₂ is significantly blue shifted from that of the bulk GaN (sample C) having the peak at 568.5 ± 0.5 cm⁻¹ which results from the strain accompanied by the pseudomorphic growth of GaN on SL-WSe₂/c-sapphire. Thus, AFM, STEM and micro-Raman measurements confirm that the existence of SL-WSe₂ during the growth of GaN.

In order to investigate the optical quality of both WSe₂, bulk like GaN epilayer and GaN/WSe₂ (samples A-C) room temperature μ PL and absorbance measurements were performed. Figure 4 (a and b) show the RT μ PL and optical absorbance spectra acquired on GaN epilayer and WSe₂. The optical band edge emissions (E_{opt}^{GaN} and $E_{opt}^{SL-WSe_2}$) for GaN and WSe₂ are observed at ≈ 3.43 and ≈ 1.63 eV, respectively. The low intensity PL for sample B could be due to the presence of octahedral phase of WSe₂, which will be discussed later. The WSe₂ (A) and GaN/WSe₂ (B) exhibits the direct optical bandgap which is matching with the value of single layer²³. As represented in Figure 4(b) the observed states X_A , X_B , $X_{A'}$ and $X_{B'}$ for sample A at ≈ 1.63 , ≈ 2.08 , ≈ 2.44 and ≈ 2.91 eV in the absorbance spectrum are the exciton resonances corresponding to the transitions from broken inversion symmetry induced two spin-split valence sub-bands to the conduction band and vice versa^{36–38}. Thus, the spin splitting of WSe₂ is observed to be ≈ 0.45 eV which is smaller than that of MoSe₂ exhibiting the largest value of ≈ 0.55 eV among MX₂ family^{39,40}. Hence, this observation confirms that the CVD grown WSe₂ sample is in the form of SL, as corroborated by earlier AFM, STEM and micro-Raman measurements. However, these are not an exact band to band transitions due to the involvement of excitons. The electronic band gap is measure of actual gap of a material which is the summation of optical bandgap and exciton (electron-hole) binding energy ($E_g = E_{opt} + E_b$).⁴⁰ Hence, these observed optical bandgap values differ from reported electronic bandgap values of $E_g^{SL-WSe_2} = 2.08$ eV and $E_g^{GaN} = 3.45$ eV by their respective exciton binding energy (E_b) values of ≈ 0.450 and ≈ 0.023 eV as reported in the literature.^{23,41,42}

High-resolution XPS measurements were extensively used to measure the valence band offset (VBO) of a heterointerface. In order to evaluate VBO at GaN/SL-WSe₂ interface, the energy difference between the Ga and W core levels from the GaN/SL- WSe₂ heterojunction sample and the energy of core levels relative to the respective valence band maximum (VBM) of the GaN epilayer and SL- WSe₂ samples need to be acquired. As the escape depth of photo emitted electrons in HRXPS is significantly low, over grown GaN layer of heterojunction sample has to be thin enough so that the electrons removed from both thin overgrown GaN and underlying SL- WSe₂ layers can be easily examined.⁴³ However, the surface area of SL-WSe₂ is large enough to carry out the XPS measurements, due to the non-continuity of the WSe₂ film on sapphire substrate, the region of interest on WSe₂/sapphire and GaN/WSe₂ samples was selectively chosen within the

spatial resolution of HRXPS measurements.^{23,44} This is executed by comparing the intensity of Ga 2p, W 4f and Al 2p core-levels which allowed us to collect the photoemission signal from solely SL-WSe₂/sapphire and GaN/SL-WSe₂ heterostructures for samples A and B, respectively. Since LMM auger line of Ga at ≈ 397 eV overlaps with N 1s core level for Al K α anode as x-ray source in HRXPS measurements, the N 1s core levels are not considered for the analysis of VBO determination. Furthermore, Ga 3d is also excluded from the analysis as it exists close to the valence band states.⁴⁵ Thus, in order to determine the VBO, the core levels W 4f, Se 3d and Ga 2p were used for the analysis. The valence band offset (VBO) for GaN/SL-WSe₂ heterojunction, can be calculated by the method provided by Kraut et al.⁴⁶, expressed as

$$\Delta E_v = \left(E_{W4f_{7/2}}^{WSe_2} - E_{VBM}^{WSe_2} \right) + \left(E_{Ga2p_{3/2}}^{GaN/WSe_2} - E_{W4f_{7/2}}^{GaN/WSe_2} \right) - \left(E_{Ga2p_{3/2}}^{GaN} - E_{VBM}^{GaN} \right) \quad (1)$$

From Figure 5a (i and ii), the first term of equation (1) deduced by taking into account the position of W 4f_{7/2} core-level referenced with respect to the VBM, which is measured to be 31.56 $\pm$ 0.10 eV. The values of the valence band maxima (VBM) are determined by extrapolating the linear fit to the leading edge of the valence band to the base level.^{47,48} Prior to this, W 4f core-level is deconvoluted so as to access the accurate value that contributed solely from W-Se bonding in SL-WSe₂. In this peak deconvolution process, we use Voigt (mixed Lorentzian-Gaussian) line shapes for fitting the W-Se (trigonal prismatic -2H) and W-O chemical states, respectively, centered at 32.52 $\pm$ 0.10 and 36.15 $\pm$ 0.10 eV. The later state with low intensity results from WO₃ precursor or excess W metal bonding with the oxygen of c-Al₂O₃ at the interface of WSe₂/c-sapphire during high temperature CVD growth.

In order to evaluate the subsequent term, GaN/SL-WSe₂ sample is used for XPS measurements where we consider the difference between Ga 2p_{3/2} and W 4f_{7/2} core-levels. Figure 5b (i and ii) depicts the Ga 2p and W 4f core-levels which are acquired on GaN/SL-WSe₂ heterointerface. Figure 5(b-i) represents the Ga 2p core-level which is deconvoluted with Ga-N bonding and any other states are not observed. The Figure 5(b-ii) shows the fitting of W 4f core-level equipped with three W 4f_{7/2} chemical states at 30.81 $\pm$ 0.10, 31.94 $\pm$ 0.10 and 36.17 $\pm$ 0.10 eV, ascribed to the W-Se bonding in 1T-WSe₂ (octahedral), W-Se bonding in 2H-WSe₂ (trigonal prismatic), and W-O bonding in WO₃, respectively.⁴⁹ The W 4f_{5/2} core-level has similar deconvolutions at higher binding energy values differed by ≈ 2.16 eV from W 4f_{7/2} bonding states. The octahedral phase could be due to unintentional N-plasma intercalation of WSe₂ single layer during GaN growth which is as similar as lithium intercalation reported for the case of MoS₂³⁷. The presence of

octahedral phase results in the low intensity PL signal for sample B as presented in Figure 4(b). The W-Se bonding peak infers that the sustainability of SL-WSe₂ at GaN growth temperature under UHV oxygen free environment. The absence of any other chemical state associated with W or Se in Ga 2p core-level spectrum is a clear evidence of Vander Waal epitaxy. Thus, the energy difference between W 4f_{7/2} and Ga 2p_{3/2} core-levels is observed to be 1085.92±0.10 eV as described in Figure 5(b).

The last term indicates the core-level energy 1115.23±0.10 eV of Ga 2p_{3/2} measured relative to the respective VBM of 2.27±0.15 eV as shown in Figure 5(c). This VBM measured with respect to the Fermi level for nearly intrinsic GaN which was utilized in the present study is lower than that of the n-type GaN in the literature (2.7 eV).⁴⁷ In Figure 5c(i), Ga 2p_{3/2} and 2p_{1/2} core-levels are deconvoluted with Ga-N bonding state at 1117.50±0.10 and 1144.40±0.10 eV, respectively. The Fermi level position with respect to the VBM as shown in Figure 5(a-ii and c-ii) infers that GaN epilayer and SL-WSe₂ are nearly intrinsic materials. Thus, the VBO from equation (1) is determined to be ≈2.25±0.15 eV. The VBO is also confirmed by using VB spectrum, Se 3d and Ga 2p core levels of samples A and B (see supporting information).

As a result, the substitution of VBO (ΔE_v) obtained from HRXPS analysis (Figure 5) and electronic bandgap (E_g) values of SL-WSe₂ and GaN epilayer in equation (2), allows to measure the conduction band offset (CBO) ΔE_c for GaN/SL-WSe₂ heterostructure.

$$\Delta E_c = E_g^{WSe_2} + \Delta E_v - E_g^{GaN} \quad (2)$$

Thus, the CBO (ΔE_c) is determined to be 0.88±0.10 eV. Furthermore, the measured CBO value is verified by the Anderson's affinity rule which is defined as the difference between electron affinity values of constituent semiconductors of heterojunction.⁵⁰ The electron affinity (the energy separation between vacuum level and CBM) values χ^{GaN} , χ^{SL-WSe_2} of GaN and SL-WSe₂ were ≈4.1 and ≈3.3 eV, respectively.^{51,52} UPS measurements were performed to measure the electron affinity of GaN. Here, electron affinity (χ) of SL-WSe₂ is extracted using the formula $\chi = \phi - (E_g - E_F)$ from our earlier reported work function value deduced by ultraviolet photoelectron spectroscopic (UPS) measurements⁵², where ϕ is the work function (3.61 eV) and E_g is the electronic bandgap (2.08 eV) and E_F is the Fermi level position with respect to VBM (1.8 eV) of an SL-WSe₂. For more details see the supporting information. It should be noted that

electron affinity is material specific and independent of the Fermi level position. Hence, from Anderson's rule, calculated CBO for GaN/SL-WSe₂ heterojunction is 0.8 eV. The determined CBO value of 0.88 ± 0.15 eV from HRXPS studies in the present study is consistent with the value (0.8 eV) obtained by affinity rule, which facilitates in understanding the electron transport properties across the heterojunction.

Thereby, the experimentally measured offset parameters from this study are presented as a schematic of band discontinuity diagram in Figure 6 which pertains to type-II heterostructure.

4. CONCLUSIONS

In conclusion, GaN epitaxial thin layers were deposited on SL-WSe₂/c-sapphire substrates by PAMBE to study the band discontinuity at GaN/SL-WSe₂ heterointerface. We confirm that the CVD deposited WSe₂ is a single-layer by complementary characterization tools such as AFM, HAADF-STEM, micro-Raman modes, absorbance and μ PL studies. The determination of band offset parameters at GaN/SL-WSe₂ heterostructure was carried out by utilizing the HRXPS, electron affinities and electronic band gap values. We determine the valence band and conduction band offset values, respectively, of 2.25 ± 0.15 eV and 0.8 ± 0.15 eV with type II band alignment at GaN/SL-WSe₂ heterostructure. This demonstration of unprecedented band offset parameters renders a route towards the integration of 3D group III nitride materials with 2D transition metal dichalcogenide layers.

ASSOCIATED CONTENT

Supporting Information: Confirmation of valence band offset by using Ga 2p and Se 3d HRXPS core levels. Determination of electron affinity for GaN and SL-WSe₂ by ultraviolet photoelectron spectroscopy (UPS).

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Notes The authors declare no competing financial interest.

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Figures and captions.

Figure 1 shows the AFM scans for (a) GaN epilayer, (b) WSe₂ and (c) GaN/WSe₂ samples, respectively. The line profile of 1(b) represents the thickness of WSe₂ layer. The inset to Figure 1(b) is optical microscopy image of WSe₂ sample. Figure 1(d) shows the HRXRD 2θ-ω scans of on-axis and off-axis reflections and the inset shows the respective ω-scans acquired on GaN epilayer.

Figure 2 (a and b) show the low and high magnification cross-sectional images of high-angle annular dark field - scanning transmission electron microscopy (HAADF-STEM) for sample B. (c) shows the STEM-EELS elemental color maps (W-Blue, Se-Green, Ga-Red, and Al-Yellow). Figure 2(d) represents the STEM-EDS elemental line profiles acquired across the GaN/WSe₂/c-sapphire heterojunction.

Figure 3(a-d) show the micro-Raman spectra of samples A-C collected on WSe₂, GaN/WSe₂ and bulk like GaN samples, respectively. Here, He-Cd (325 nm) and He-Ne (633 nm) excitation sources ($E_{exc} > E_g$) were used to enhance the Raman signal based on the band gap of each material.

Figure 4(a and b) show the micro-photoluminescence (right y-axis) and absorbance spectra (left y-axis) of GaN and WSe₂ samples. The green colored PL peak in Figure 4(b) is for GaN/WSe₂ sample.

Figure 5a (i and ii) present the W 4f core-level and valence band spectra for SL- WSe_2 . b(i and ii) show the spectra of Ga 2p and W 4f core-levels for GaN/SL-WSe_2 . c(i and ii) show Ga 2p core-level and valence band spectra acquired on GaN epilayer. The peak positions of core-levels are shown in parentheses.

Figure 6 shows the schematic representation of type II band alignment at GaN/SL-WSe₂ heterointerface.

TOC