

S/TEM Characterization of Vertical Heterostructures Formed by Mono- to Multi-layer Graphene and WSe₂

Saiphaneendra Bachu¹, Benjamin Huet¹, Danielle Reifsnyder Hickey², Chenhao Qian¹, Joan Redwing¹ and Nasim Alem³

¹Pennsylvania State University, United States, ²Pennsylvania State University, Pennsylvania, United States, ³Pennsylvania State University, Washington, District of Columbia, United States

Two-dimensional (2D) transition metal dichalcogenides (TMD) are emerging semiconductor materials in the field of nano optoelectronics owing to their desirable physical and electronic properties [1]. They are touted to replace silicon in the sub-5 nm regime where silicon suffers from quantum confinement effects [2]. Hence, there is a great interest in growing large area defect-free electronic grade TMD materials towards target optoelectronic applications. A recent development in the synthesis of TMD materials is the use of 2D growth substrates such as graphene and hBN as alternatives to traditional sapphire and silicon. Compared to the traditional substrates, 2D substrates offer potential advantages such as step edge free flat surfaces, absence of dangling bonds etc. [3]. Moreover, 2D substrates could enable true van der Waals epitaxy with the as-grown TMD with relaxed lattice matching conditions [4]. In addition, usage of 2D substrates could lead to the formation of 2D vertical heterostructures with enhanced optoelectronic properties and exciting physical phenomena [5, 6]. For example, H. Buch et al. have observed superlubricity in epitaxial graphene-WS₂ vertical heterostructures [6].

In this work, we investigate the nucleation of WSe₂ synthesized on mono- to multi-layer graphene via MOCVD process. The mono- to multi-layer graphene was first prepared using chemical vapor deposition (CVD) technique on Cu substrate and then transferred to sapphire substrate prior to the WSe₂ growth. As-grown heterostructures are characterized using scanning/transmission electron microscopy (S/TEM) imaging to understand the atomic structure of the heterostructures, epitaxy between graphene and WSe₂ layers and the role of defects in graphene as the underlying mechanism behind the nucleation of WSe₂ on Graphene.

Figure 1a shows a scanning electron microscope (SEM) image of WSe₂ triangles (~ 500 nm) synthesized on CVD grown multilayer graphene after the MOCVD step. Layer numbers of graphene are labeled in the image. It can be observed that the nucleation density of WSe₂ triangles is increasing with the graphene layer number. Selected area diffraction (SAD) pattern obtained from the as-grown heterostructures in monolayer graphene area is presented in Figures 1b. It is evident that WSe₂ triangles (green hexagon) maintain an epitaxial relationship with the graphene layer (blue hexagon). Figures 2 shows the atomic resolution images of graphene and WSe₂. While Figure 2a is a monochromated high-resolution TEM image taken from monolayer graphene region, Figure 2b is an atomic resolution high-angle annular dark-field (HAADF)-STEM image taken from a WSe₂ triangle. The presentation will further uncover the role of the substrate and epitaxy, graphene layer number, and substrate defects on the nucleation density of WSe₂ [7].

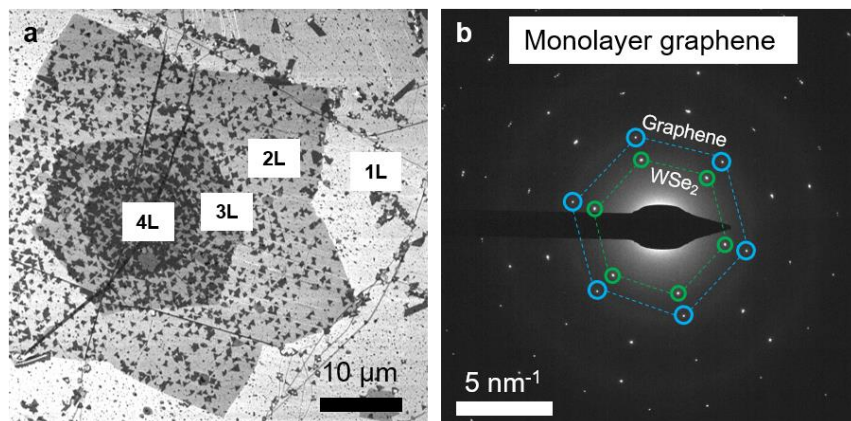


Figure 1. (a) SEM image of WSe_2 triangles grown on multilayer CVD graphene; the labels indicate the number of graphene layers and (b) SAD pattern obtained from heterostructures grown on the monolayer graphene area showing epitaxy between graphene and WSe_2 .

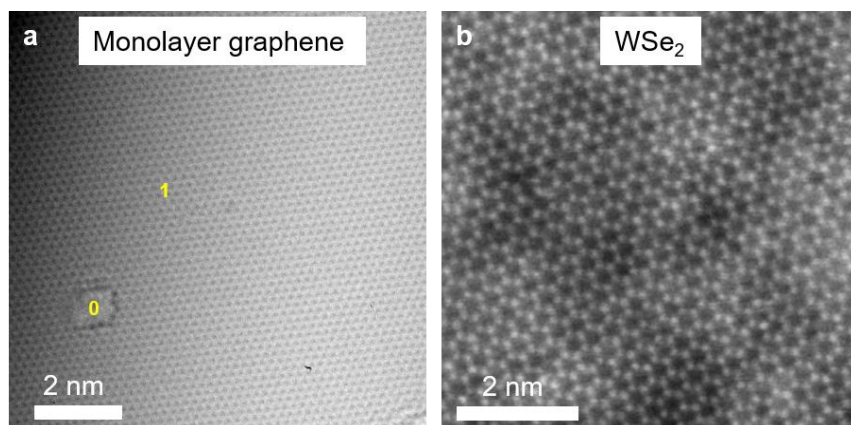


Figure 2. (a) Monochromated HRTEM image from monolayer graphene region; the numbers in the image indicate the number of graphene layers and (b) atomic resolution HAADF-STEM image of WSe_2 grown on the monolayer graphene region.

References

- [1] Jariwala, Deep, et al., *ACS nano* 8.2 (2014): 1102-1120.
- [2] Tsutsui, Gen, et al., *IEEE Transactions on nanotechnology* 4.3 (2005): 369-373.
- [3] Zhang, Xiaotian, et al., *ACS nano* 13.3 (2019): 3341-3352.
- [4] Bianco, G. V., et al., *RSC advances* 5.119 (2015): 98700-98708.
- [5] Lin, Yu-Chuan, et al., *Nano letters* 14.12 (2014): 6936-6941.
- [6] Büch, Holger, et al., *Nano Research* 11.11 (2018): 5946-5956.
- [7] This work was supported by the National Science Foundation (NSF), in part under the CAREER program (DMR-1654107), in part by the program EFRI 2-DARE: 2D Crystals by Activated Atomic Layer Deposition (EFRI-1433378), and in part by the Penn State 2D Crystal Consortium-Materials Innovation Platform (2DCC-MIP) under NSF cooperative agreement DMR-1539916.