Metal-Organic Chemical Vapor Deposition of Wafer-Scale and Carbon- Free WS₂ Atomic Layers

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Abstract

The successful growth of Two-Dimensional (2D) TMDs materials is one of the prerequisites in order to take advantage of their remarkable electronic and optoelectronic properties in real applications. This requires the systematic study of the growth mechanism of such atomic films. Here, I will present the results on the growth of WS₂ and WSe₂, via the metal-organic chemical vapor deposition (MOCVD) approach. One of the major advantages of MOCVD is the use of volatile precursors for both, the metal and the chalcogen sources, in contrast to the more common metal oxide powder-based CVD approach[1], where the control over the precursor supply is limited. Although the MOCVD method results in large-scale growth, it might include intrinsic carbon contamination [2] arising from the volatile organic-based precursors themselves. This, together with the high rate of precursor decomposition at the growth teperatures, leads to the formation of nano-crystalline films.

Here, a Growth-Etch cycle technique is developed, in which the precursors are sequencially delivered while a small amount water vapor is introduced to the growth chamber. This causes the re-evaporation of small and defective domains as well as the carbon contaminants from the growth substrate, allowing the highly crystalline domains to expand, resulting in high-quality large domains or continuous atomic layers, depending on growth conditions[4]. This methodology could be further extended to other 2D materials in general and TMDs in particular. Finally, I will describe how the MOCVD approach can be used to study the vander Waals apitaxy of TMDs on different substrates.

References

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Figures

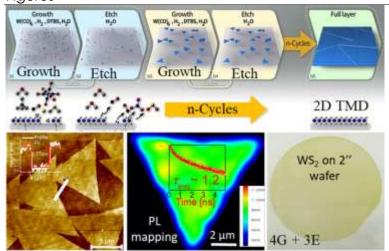


Figure 1: Schematic model of the growth-etch method, Topography map with the corresponding height profile of a monolayer WS $_2$, PL intensity map and TRPL decay measurement on a single crystal WS $_2$, a full coverage monolayer WS $_2$ on a 2" sapphire substrate