Controlling wafer-scale epitaxial growth of transition metal dichalcogenides using MOCVD

Tanushree H. Choudhury,^{1*} Mikhail Chubarov¹, Danielle R Hickey², Saiphaneendra Bachu², Tianyi Zhang², Mauricio Terrones^{2, 4, 5}, Nasim Alem^{1, 2}, and Joan M. Redwing^{1, 2}

¹2D Crystal Consortium-Materials Innovation Platform (2DCC-MIP), Materials Research Institute, The Pennsylvania State University, University Park, PA 16802, USA

²Department of Materials Science and Engineering, The Pennsylvania State University, University Park, PA 16802, USA

⁴Department of Chemistry, The Pennsylvania State University, University Park, PA, USA

⁵Department of Physics, Center for 2-Dimensional and Layered Materials, The Pennsylvania State University, University Park, PA, USA

*Contact@E-mail: tuc21@psu.edu

Abstract

Our research is focused on the development of an epitaxial growth technology for layered dichalcogenides MX_2 (M=Mo, W and X= S, Se), based on metal organic chemical vapor deposition (MOCVD). This process uses metal hexacarbonyl and hydride chalcogen precursors to deposit monolayers on 2" sapphire wafers in a cold-wall reactor. A multi-step precursor modulation growth method is used to independently control nucleation density and the lateral growth rate of monolayer domains on the sapphire substrate [1]. Using this approach, uniform, coalesced monolayer TMD films were obtained at growth rates on the order of ~1 monolayer/30 min. In-plane X-ray diffraction demonstrates that the films are epitaxially oriented with respect to the sapphire [2]. Controlling the growth temperature and chalcogen flux was crucial in establishing an epitaxial relation. For example, dark-field transmission electron microscopy (DF-TEM) of transferred WS₂ monolayers show ~95% single orientation coverage with minimal bilayer and inversion domains (Figure 1). These singlecrystal transferred films also show narrow exciton linewidths (~31 meV) and negligible defect-related emission at 80 K (Figure 1). The key features observed during the growth of MoS₂, WS₂, MoSe₂ and WSe₂ will be compared and discussed. The authors acknowledge financial support of the U.S. National Science under NSF cooperative agreement DMR-1539916 and EFRI 2-DARE Grant EFRI-1433378.

References

- X. Zhang, T.H. Choudhury, M. Chubarov, Y. Xiang, B. Jariwala, F. Zhang, N. Alem, G.C. Wang, J.A. Robinson and J.M. Redwing, Nano Lett. 18, (2018)1049.
- [2] M. Chubarov, T.H. Choudhury, X. Zhang and J.M. Redwing, Nanotechnology 29 (2017)055706.

Figures



Figure 1: (a) Photograph of 2" WS_2 film on c-sapphire, (b) AFM micrograph of WS_2 film with < 1% bilayer, (c) Composite dark-field TEM map of the film corresponding to (b) and (d) Temperature-dependent photoluminescence (PL) of WS_2 monolayer

Graphene2020