GROWTH AND CHARACTERIZATION OF THIN MoS₂ LAYERS ON EPITAXIAL GRAPHENE ON SiC(0001)

A. Schütze^a, M. Zeißig^a, M. Wanke^a, F. Speck^a, and Th. Seyller^a

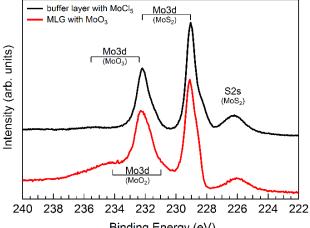
^aInstitut für Physik, TU Chemnitz, Reichenhainer Straße 70, 09126 Chemnitz, Germany

The unique property of layered metal dichalcogenides such as MoS₂ to transform from an indirect to a direct semiconductor when reducing the thickness to one monolayer offers new possibilities for electronic devices. For applications, however, the availability of scaleable production methods is a prerequisite.

In the present work, we investigate the chemical vapor deposition (CVD) of thin layers of MoS₂ using sulfur powder and two different molybdenum precursors, MoO₃

[1] and MoCl₅ [2]. MoS₂ was deposited graphene monolayer on (MLG) SiC(0001). epitaxially grown on Alternatively, the so-called buffer layer (BL), which is a graphene-like layer strongly bound to SiC(0001) [3], was used as substrate. Both BL and MLG were prepared by sublimation growth in argon at atmospheric pressure as described elsewhere [4].

Samples were characterized using X-ray photoelectron spectroscopy (XPS) for the chemical composition of the samples, atomic force microscopy Fig. 1. Mo3d and S2s core-level spectra of CVD-MoS₂



Binding Energy (eV)

and low-energy electron diffraction for films on buffer layer and monolayer graphene using the structure and crystallinity of the MoCl₅ and MoO₃ precursors, respectively.

deposited layers. For MoO₃ as precursor, a rather inhomogeneous MoS₂ growth is found which is accompanied by interface oxidation of the SiC due to oxygen intercalation. On the other hand, MoCl₅ as precursor results in an improved homogeneity of the deposited films and sulfur intercalation as suggested by XPS data. For the latter, a thickness of the deposited MoS₂ of approximately 1 to 3 monolayers is derived from analysis of core-level intensities.

Keywords: MoS₂; Graphene; CVD; XPS

References

- [1] S. Wang et al., Chem. Mater. 26 (2014) 6371.
- [2] Y. Yu et al., Sci. Rep. 3 (2013) 1866.
- [3] M. Ostler et al., Phys. Status Solidi B 247 (2010) 2924.
- [4] K. V. Emtsev et al., Phys. Rev. B 77 (2008) 155303.