Scalable methodology for the direct synthesis of atomically thin WS₂ films

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Abstract

Tungsten disulfide (WS₂) in the monolayer form is a promising 2D material for the next generation of optoelectronic devices due to its direct bandgap in the visible [1]. However, the development of such technologies needs scalable synthesis processes able to provide WS₂ deposition with full control in terms of structural quality, thickness and, hence, optical properties.

In this contribution, we show the flexibility of a pure chemical methodology for depositing mono and few layer WS₂ films with defined layer structures on different substrates including SiO₂, Si, Al₂O₃, CVD- and epitaxial-graphene (van der Waals heterostructures). This CVD technique uses tungsten hexacarbonyl (W(CO)₆) and sulfur (S) as precursors and is based on a single step chemical mechanism (W(CO)₆ + S₂ \rightarrow WS₂ + 6CO) which allows a good control in the material growth (especially on the large area) compared with more traditional two step processes based on the metal oxide physical deposition and subsequent sulfurization [2]. A full chemical (Raman, EDS), structural (SEM, AFM) and optical characterization (photoluminescence and spectroscopic ellipsometry) of deposited WS₂ films is provided to give further insight to the optical response of WS₂ exciton system and its dependence on WS₂ films structural quality and interface nature, e.g. presence of an epitaxial relationship with the substrate or occurrence of charge transfer processes.

References

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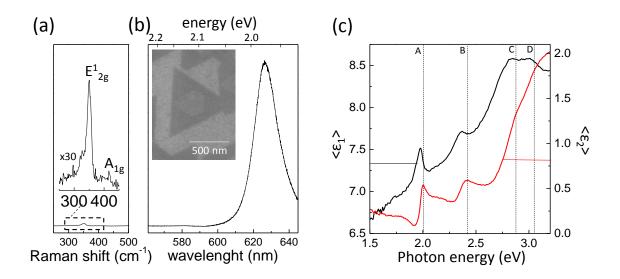


Figure. (a) Raman and (b) photoluminescence spectra (in the same intensity scale) of WS₂ film deposited on SiO₂ substrate (SEM image in the inset). (c) Ellipsometric spectra of the imaginary (< ϵ_1 >) and real part (< ϵ_2 >) of the pseudodielectric function ($\epsilon = <\epsilon_1 > + <i\epsilon_2 >$) of WS₂ film on epitaxial graphene