

Emission properties of near band-gap excitons and exciton kinetics in Niobium-doped Rhenium Disulphide (ReS₂:Nb)

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Atomically thin Transition Metal Dichalcogenides (TMDs) are a new class of materials which exhibit variety of physical properties.[1] Some of these properties are quite similar to the bulk-like materials, however, other rely strongly on quantum confinement effects. Among TMDs one can find well known semiconductors like MoS₂, WSe₂, superconductors like NbSe₂, PdTe₂, or semimetals e.g. WTe₂, TiSe₂, however, there are still some TMD chemical compounds for which their basic physical parameters are unknown. The ReS₂ is one of those. In this work we examine emission properties of near band-gap excitons and exciton kinetics in the ReS₂:Nb.

We utilize a polarization-resolved, high spectral resolution photoluminescence experiment at T=10 K in order to obtain emission properties of a ReS₂:Nb thin layer. The PL spectrum of the ReS₂:Nb shows a doublet of linearly-polarized emission bands E₁, and E₂ centered at 1.553 eV and 1.585 eV, respectively. According to the latest theoretical studies [2] these bands can be formed due to radiative recombination process of excitons near the fundamental gap of rhenium disulphide at the Γ point of the Brillouin zone (direct gap). Both PL lines are orthogonally polarized. Although the physical picture lying behind this observation remains still unclear, it can be attributed to the existence of zigzag Re chains along one of the lattice vectors in the plane of a ReS₂:Nb layer. At the high energy range of the PL spectrum another two emission bands are observed: the E₃ band located at 1.644 eV, and E₄ one centered at ~1.67 eV, both being polarization insensitive. In contrast to previous observations existing in the literature the E₄ band reveals its fine structure. It consists of four equidistant PL peaks separated by ~5 meV each. The E₃ and E₄ bands are attributed to the recombination of excitons at higher energy states within the ReS₂ band structure, however exact origin of those states is ambiguous.

Exciton kinetics on the E₁ and E₂ energy levels is examined in a time-resolved, pump-probe differential reflection experiment at T=10 K. Decay traces of the differential reflection signal are typically composed of two components. The fast component of ~0.2 ps decay time, which is close to the setup resolution, is attributed to radiative decay of a free exciton within the fundamental gap of ReS₂. The slow component of ~1.2 ns decay time constant is associated to the recombination process of bound exciton states near the fundamental gap of the ReS₂. These time constant parameters were unavailable in the literature, up to now.

[1] M. Chhowalla, H. S. Shin, G. Eda, L.J. Li, K. P. Loh, and H. Zhang. *Nature Chem.* **5**, 263 (2013)

[2] S. Tongay, H. Sahin, Ch. Ko, A. Luce, W. Fan, K. Liu, J. Zhou, Y. S. Huang, C. H. Ho, J. Yan, D. F. Ogletree, S. Aloni, J. Ji, S. Li, J. Li, F. M. Peeters, and J. Wu *Nature Comm.* **5**, 3252 (2014).