

Time Resolved photoluminescence spectra of WS₂ and MoS₂ at high excitation fluence

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Semiconducting two-dimensional Transition Metal Dichalcogenides (TMDs), such as MoS₂ and WS₂, exhibit the indirect-to-direct bandgap transition in the monolayer limit and a large exciton binding energy (> 0.1 eV) [1], allowing to investigate the physics of excitons at room temperature by means of PhotoLuminescence (PL) experiments. Despite since 2010 many PL investigations have been performed, both in the steady state and out of equilibrium, a complete study of the time evolution of the PL spectrum after a pulsed excitation is still lacking, especially at high exciton density. In this regime, the generation of an electron hole plasma (EHP) and of an electron hole liquid (EHL) have been reported for freestanding monolayer MoS₂, in steady state PL experiments [2]. In this seminary we report novel Time Resolved PhotoLuminescence (TR-PL) data for mechanically exfoliated monolayers of WS₂ and MoS₂ on SiO₂/Si substrates, for exciton densities ranging from 10⁹ cm⁻² to 10¹³ cm⁻². Our data allow to investigate the time evolution of the PL spectrum after an ultra-short light pulse (t=300 fs, hv=2.4 eV) excitation, with a time resolution of 50 ps, showing how the density of excitons strongly affects both the spectrum and the time dynamics of the PL light.

[1] Thomas Mueller et al, Exciton physics and device application of two-dimensional transition metal dichalcogenide semiconductors, npj 29 (2018)

[2] Yiling Yu et al, Room-Temperature Electron–Hole Liquid in Monolayer MoS₂, ACS Nano 2019, 13, 9, 10351–10358