

Abstract Submitted  
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**Time resolved photoluminescence study of CdSe/CdMnS/CdS core/shell/shell nanoplatelets heterostructures**<sup>1</sup> THOMAS SCRACE, State Univ of NY - Buffalo, SAVAS DELIKANLI, MEHMET ZAFER AKGUL, Bilkent University, JOSEPH MURPHY, TIM THOMAY, PEIYAO ZHANG, TENZIN NORDEN, ALEXANDER CARTWRIGHT, ATHOS PETROU, State Univ of NY - Buffalo, HILMI VOLKAN DEMIR, Bilkent University — We have recorded the time evolution of the photoluminescence (PL) for CdSe/CdMnS/CdS core/shell/shell solution-processed nanoplatelets (NP) using ultrafast pulses at 400 nm and 514 nm. Our NPs consist of a core with 5 monolayers (1.5 nm) of CdSe and average lateral dimensions of 55 x 10 nm<sup>2</sup>. Using 400 nm pulses we excite electron-hole pairs above the CdS shell bandgap; with 514 nm pulses we excite only in the CdSe core. The holes are primarily localized in the CdSe core, while the electrons are delocalized. Our measurements show that at  $\Delta t = 0$ , the peak PL energy for both kind of excitations is the same. As a function of time, both types of excitations result in a red-shift. The red shift of with 400 nm excitation is 60 meV and is described by two time scales:  $\tau_1 = 270$  ps and  $\tau_2 = 2.5$  ns. The red shift with the 514 nm excitation is 30 meV and is described by a single time scale:  $\tau_2 = 2.5$  ns. These results are discussed in terms of dipole layer formation[1]. [1] Gu, Y et. al. Phys. Rev. B 71 045340 (2005).

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