

Gate tuneable ultrafast charge transfer in graphene/MoS₂ heterostructures

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We report ultrafast pump-probe measurements on a graphene/MoS₂ heterostructure and demonstrate sub picosecond exciton dissociation and charge transfer from MoS₂ to graphene, one order of magnitude faster than in type II two-dimensional heterostructures [1]. The process can be controlled by applying an external gate and shifting the Fermi level of the graphene layer. For pump-probe measurements we excite the gate controlled graphene/MoS₂ heterostructure at 400 nm, well above the MoS₂ bandgap [2], and probe the normalized differential transmission changes ($\Delta T/T$) of the MoS₂ first exciton (A exciton) at 660 nm with time resolution ~ 200 fs. In this configuration, MoS₂ acts as the absorbing material for visible wavelengths while graphene is the electron scavenger [3], as depicted in Fig. 1. We observe strong dependence of recombination dynamics in MoS₂ upon gate voltage biasing and graphene doping. Specifically, higher p-doping in graphene increases the built in potential difference at the graphene/MoS₂ interface (Fig. 1a), enhances the rate of electrons transfer from MoS₂ to graphene (Fig. 2) and as a result reduces the A excitons lifetime in MoS₂. On the other hand, if graphene is n-doped, the built in field is weakened resulting in slower electrons dynamics (Fig.

1b and 2). Charge transfer in layered heterostructures was previously reported [1, 4, 5]. Here we demonstrate that this process can be electrically controlled by external gating. This mechanism is key for applications such as photodetectors [3] and non-volatile memories [6].

References

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Figures

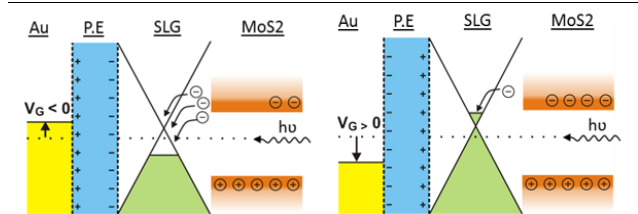


Figure 1: Sketch of the MoS₂ to graphene electron transfer as a function of graphene doping.

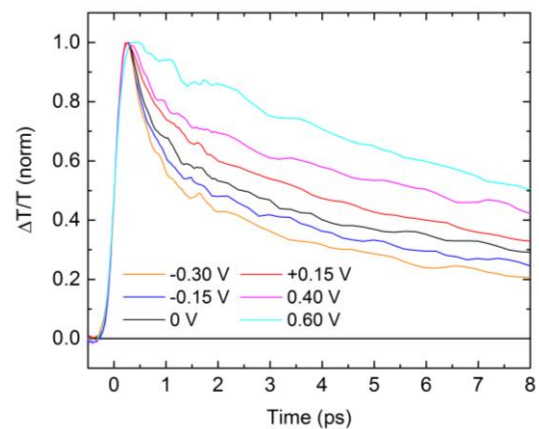


Figure 2: Pump-probe dynamics of graphene/MoS₂ heterostructure probed at 660 nm for different applied gate-voltages.