



Graphene oxide and poly(3-hexylthiophene) nanoscale interface interactions probed by KPFM

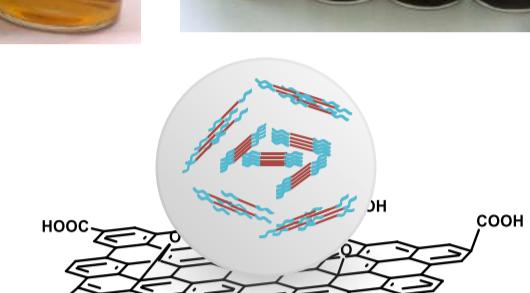
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Introduction

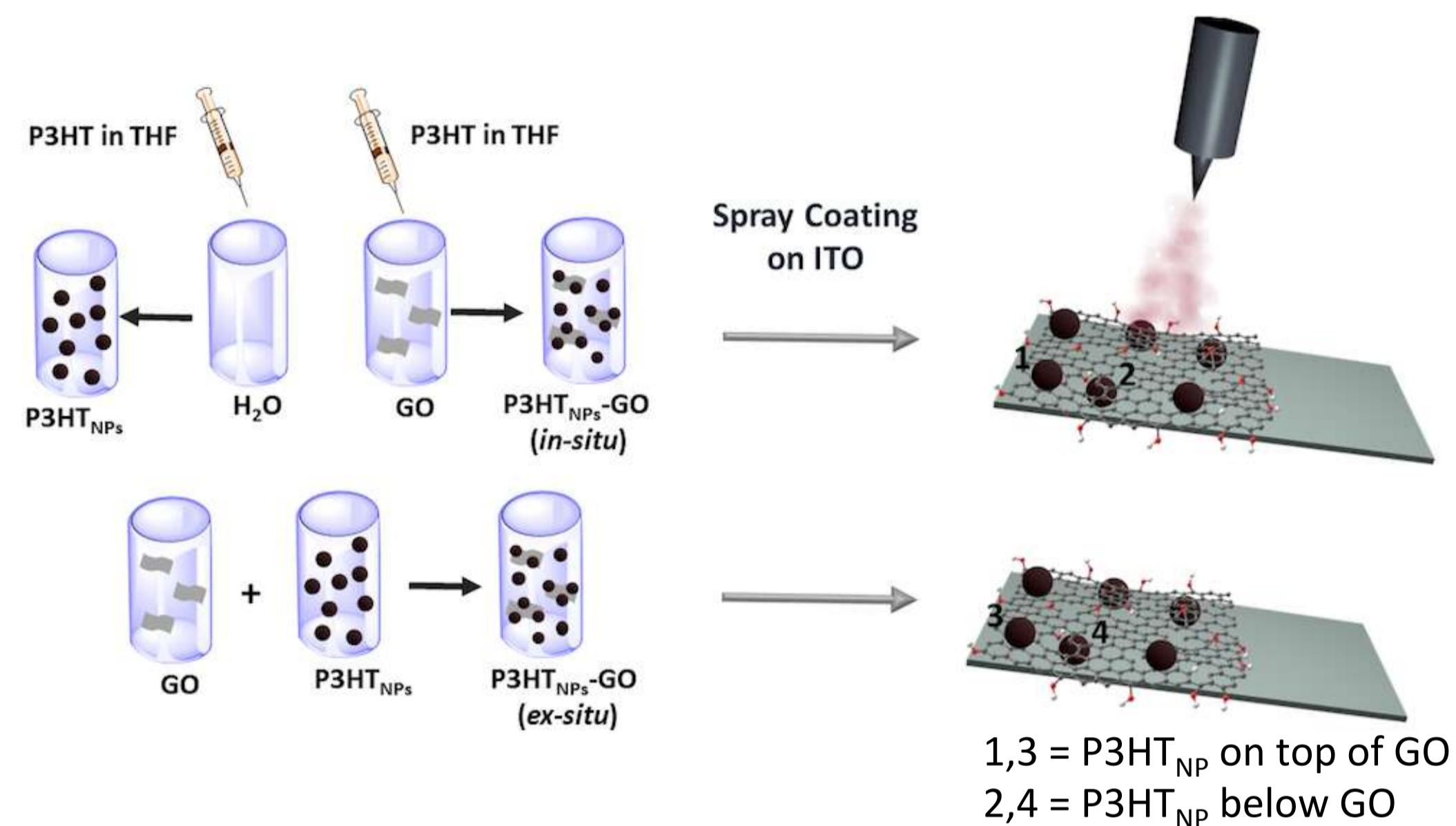


Graphene oxide (GO) is a chemically modified form of graphene containing many different types of oxygen functional groups on its basal plane and edges. Their presence imparts hydrophilicity and allows for ease of processing from water dispersions. We recently have shown that sheets of graphene oxide critically affect the aggregate structure of water dispersible nanoparticles P3HT_{NPs} and thus the performance of optoelectronic thin film devices.¹

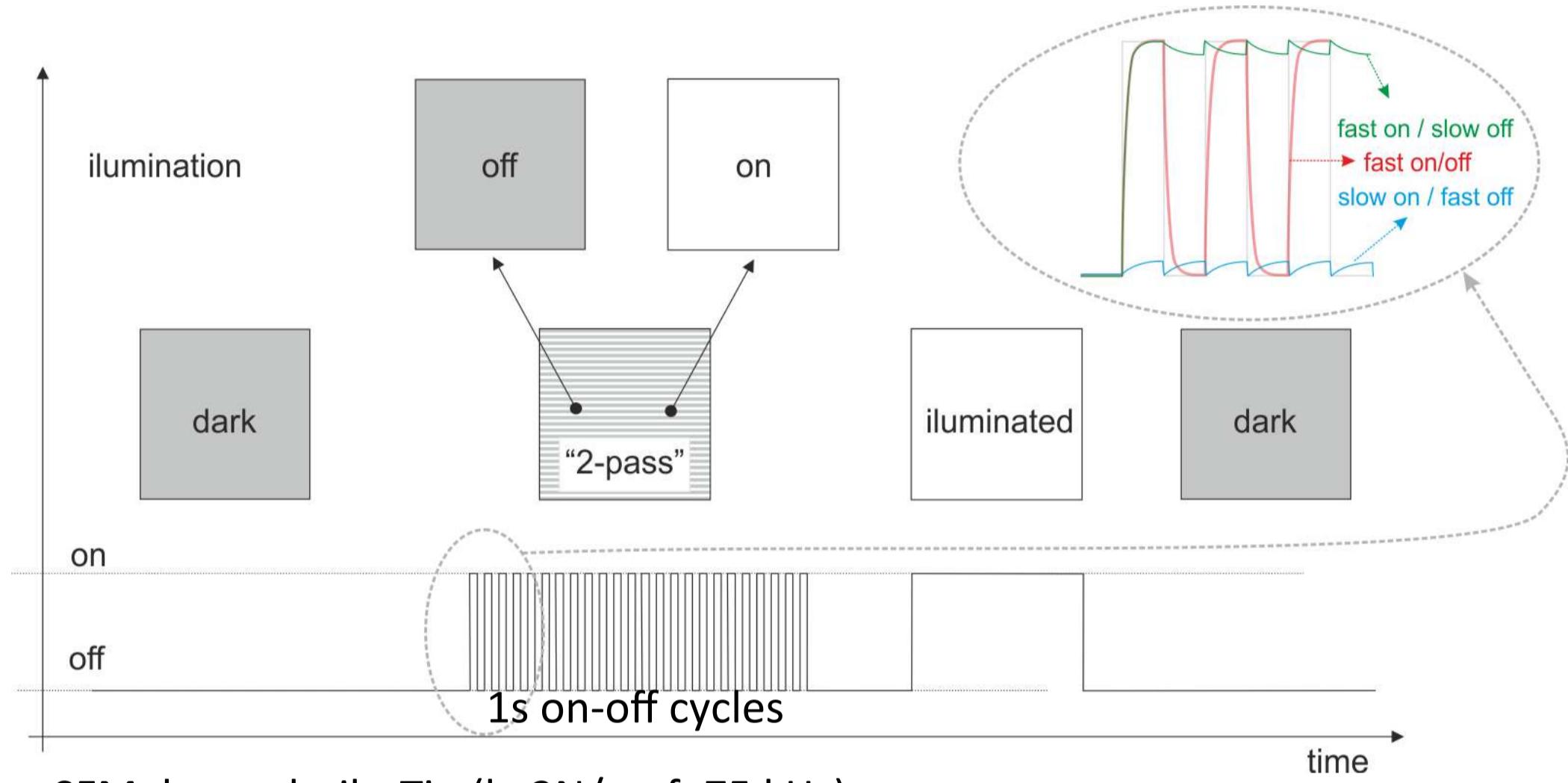
In this work we elucidate the photogenerated charge transfer dynamics of (P3HT_{NPs}-GO) ensembles with defined aggregate structure across their interface by Kelvin Probe Force microscopy (KPFM).^{2,3}

¹E. Istif et al, Adv. Funct. Mater 2018, 1707548; ²E. Palacios-Lidon, Nanotechnology 2009, 20, 085707; ³E. Palacios-Lidon, Nanoscale 2019, 11, 11202

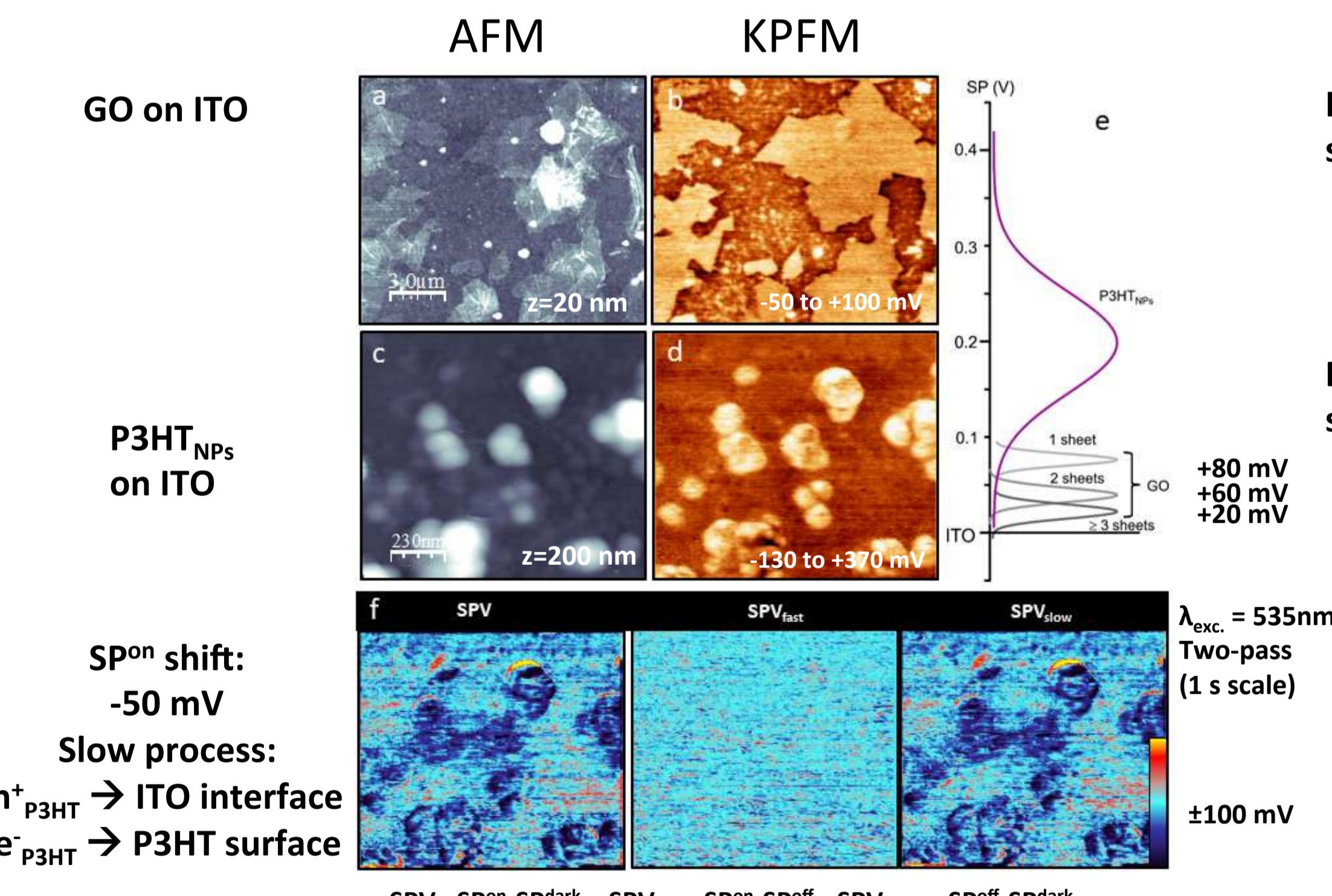
Preparation of P3HT_{NPs}-GO ensembles



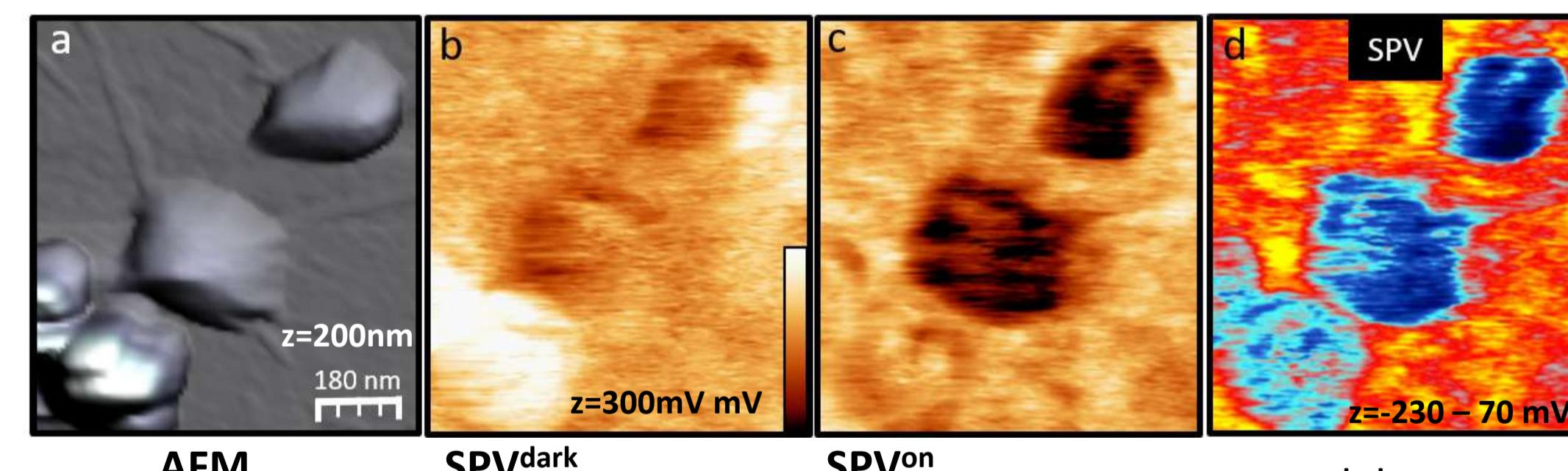
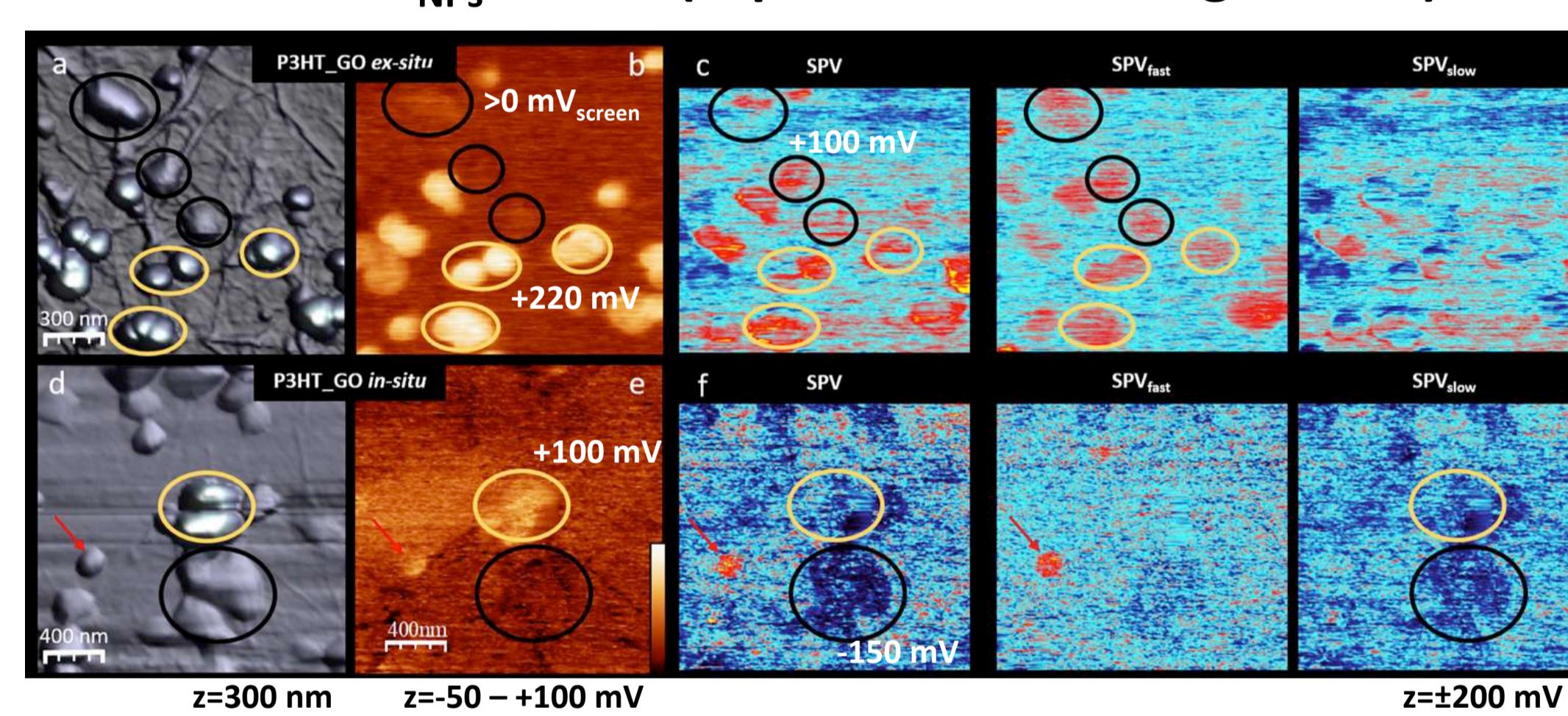
KPFM: Two-pass technique



KPFM STUDIES: Surface potential (SP) and Surface photovoltage (SPV)

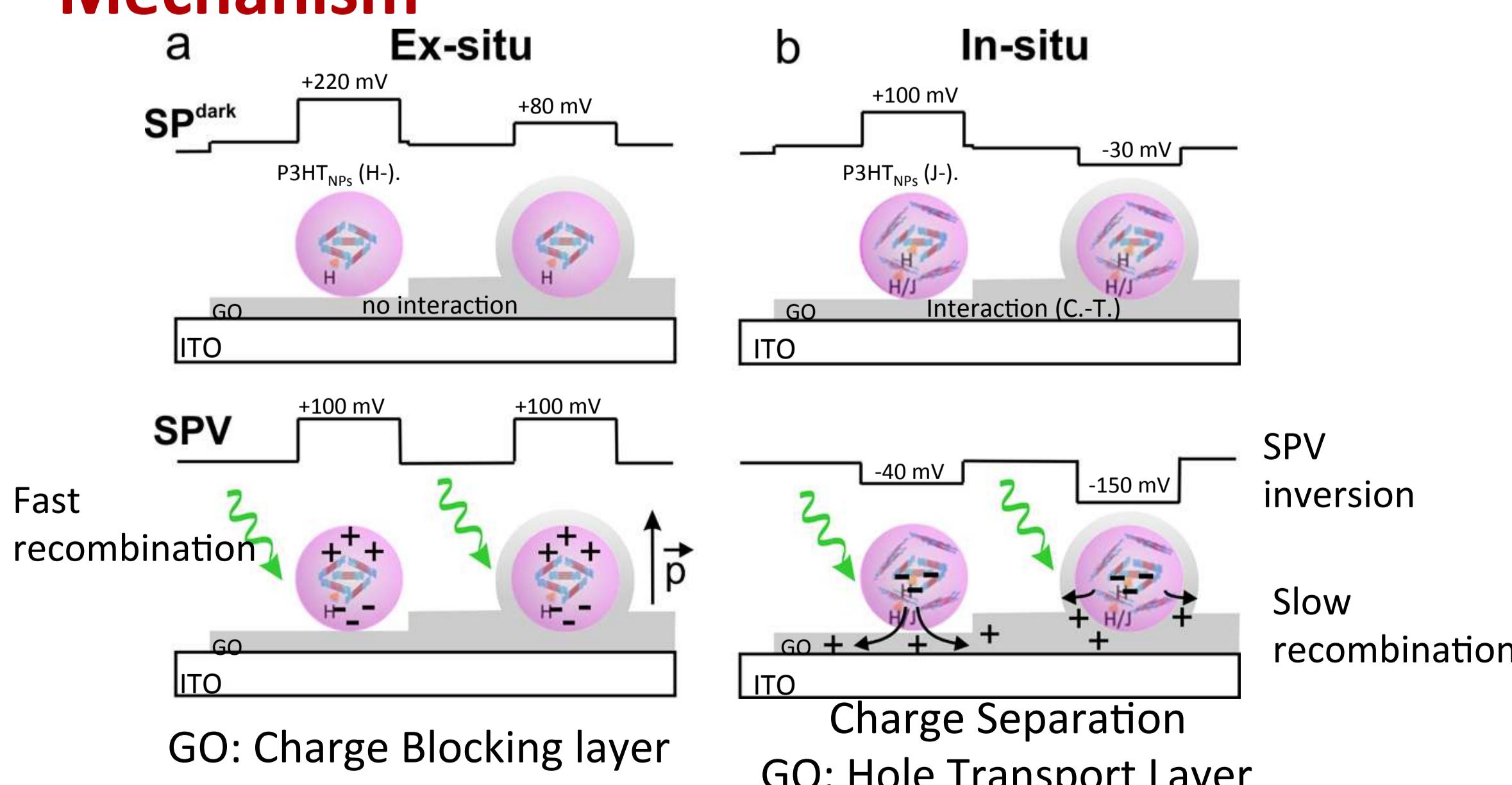


P3HT_{NPs} on GO (top and down configuration)



P3HT_{NPs}: SP substructure
Low SP on GO: C.-T. P3HT_{NPs} → GO (modified Ef) → D-A C.-T.
SPV: P3HT more negative; GO positive patches (10 nm scale)
h⁺ transferred from P3HT to GO (10 nm scale at 1 s time scale)

Mechanism



Conclusions

- GO acts as HTL
P3HT_{NPs}-GO (in-situ): Photoactive layered system
- GO acts as CBL
P3HT_{NPs} – GO (ex-situ) non-interacting system

