Abstract Submitted for the MAR12 Meeting of The American Physical Society

Ultrafast Photo Physics of P3HT/PCBM blends for Organic Photovoltaic applications BILL PANDIT, SANJEEV SINGH, ZEEV VALY VAR-DENY, Department of Physics and Astronomy, University of Utah, Salt lake City, Utah, 84112 — We studied the ultrafast dynamics of photoexcitations in pristine polymer films of regio-regular polythiophene, regio-random polythiophene, and their blends with the fullerene derivative C_{61} -PCBM using the pump-probe photomodulation (PM) spectroscopy with ~ 150 fs time resolution. Our transient PM spectrum covers the broad spectral range of 0.25 - 2.4 eV using two different laser systems; which allows us to simultaneously monitor the dynamics of various photoinduced absorption bands such as intrachain excitons, charge transfer excitons, and polaronpairs. Surprisingly, we have been able to monitor the decay of intrachain exciton on the polymer chains in films of polymer/fullerene blends, but unable to detect the subsequent generation of polarons in the donor (D) and acceptor (A) materials up to ~ 1 ns. We explain this finding considering that the excitons in the polymer chains form charge transfer excitons upon reaching the D-A interface, rather than undergo a more direct dissociation on the D-A materials. The understanding of charge separation at the D-A interface is crucial for improving the power conversion efficiency of organic solar cell devices. Supported in part by the DOE grant No. DE-FG02-04ER46109.

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Date submitted: 13 Nov 2011

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