

Abstract Submitted  
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**Fully conjugated donor-acceptor block copolymers as model systems for studies of charge transfer**<sup>1</sup> MELISSA APLAN, YOUNG-MIN LEE, CHRISTOPHER GRAY, THOMAS MALLOUK, ENRIQUE GOMEZ, Pennsylvania State University — Fully conjugated block copolymers, consisting of an electron donor and an electron acceptor block, can serve as the active layer in organic photovoltaic devices. Incorporating the donor-acceptor interface within the chemical structure enables model studies of charge transfer. We synthesized a series of block copolymers consisting of a P3HT electron donor and a push-pull polymer electron acceptor, yielding: poly(3-hexylthiophene)–block–poly-((9-(9-heptadecanyl)-9H-carbazole)-1,4-diyl-alt-[4,7-bis(3-hexylthiophen-5-yl)-2,1,3-benzothiadiazole]-2',2''-diyl) (P3HT-b-PCT6BT), poly(3-hexylthiophene)–block–poly-((9,9-dioctylfluorene)-2,7-diyl-alt-[4,7-bis(3-hexylthiophen-5-yl)-2,1,3-benzothiadiazole]-2',2''-diyl) (P3HT-b-PFT6BT), and poly(3-hexylthiophene)–block–poly-((2,5-dihexylphenylene)-1,4-diyl-alt-[4,7-bis(3-hexylthiophen-5-yl)-2,1,3-benzothiadiazole]-2',2''-diyl) (P3HT-b-PPT6BT). By altering the electron rich unit of the acceptor, we adjust the energy difference between donor and acceptor HOMOs. Using photoluminescence, we observe no evidence of exciton dissociation to a charge transfer state in P3HT-b-PCT6BT. In P3HT-b-PFT6BT and P3HT-b-PPT6BT we observe varying degrees of intrachain charge transfer. These results measure the critical driving force needed for charge transfer.

<sup>1</sup>Fully conjugated donor-acceptor block copolymers as model systems for studies of charge transfer

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