

Abstract Submitted  
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**Experimental Determination of Charge/Neutral Branching Ratio in  $\pi$ -Conjugated Polymers by Broad-band Ultrafast Spectroscopy<sup>1</sup>**

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— We demonstrate a reliable method of determining the branching ratio,  $\eta$  of photogenerated charge (polarons) to neutral (excitons) photoexcitations in various  $\pi$ -conjugated polymer films and solutions using femtosecond ultrafast spectroscopy with broad spectral range from 0.14 to 2.7 eV. We found that both excitons and polarons are instantaneously photogenerated, but  $\eta$  critically depends on the film nanomorphology, which, in turn controls the interchain coupling. In films,  $\eta$  varies between 1% for derivatives of poly(p-phenylene vinylene) casted from chloroform solution, to more than 30% for regio-regular poly-3-hexyl thiophene. Our results show that charge photogeneration quantum efficiency in these materials is an interchain process; and this has ramifications for their use in solar cell applications.

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