# Electronic Supplementary Information for

# Electron Transport in a Sequentially Doped Naphthalene Diimide Polymer Khaled Al Kurdi,<sup>1</sup> Shawn A. Gregory,<sup>2</sup> Samik Julkhi,<sup>1</sup> Maxwell Conte, <sup>1</sup> Stephen Barlow, <sup>1</sup> Shannon K. Yee, <sup>3</sup> Seth R. Marder<sup>1,2\*</sup>

<sup>1</sup>School of Chemistry and Biochemistry, Georgia Institute of Technology, Atlanta, GA <sup>2</sup>School of Materials Science and Engineering, Georgia Institute of Technology, Atlanta, GA <sup>3</sup>George W. Woodruff School of Mechanical Engineering, Georgia Institute of Technology, Atlanta, GA

\* Corresponding author E-mail: seth.marder@chemistry.gatech.edu

Contents:

I.	Synthesis and Characterization of PNBS	pS2
II.	Additional UV-visNIR Data	pS3
III.	Additional XRD Data	pS5
IV.	AFM Data	pS5

## I. Synthesis and Characterization of PNBS



Figure S1. Synthesis of PNBS.

**Table S1.** Comparison of molecular weight characteristics of PNBS synthesized in this work and reported in the literature.

This work <sup>a</sup>	Literature
$M_n$ = 42.8 kDa	$M_n = 39.7 \text{ kDa}$
$M_w$ = 86.1 kDa	$M_w = 147.6 \text{ kDa}$
PDI=2.10	PDI = 3.72.

<sup>a</sup> Molecular weight of the synthesized polymer was analyzed via Gel Permeation Chromatography (GPC) using chloroform as an eluent. GPC sample was prepared by preparing 5 g L<sup>-1</sup> solution of the polymer in 0.25% triethyl amine in chloroform (HPLC grade) and then filtered through a 20  $\mu$ m PTFE filter before running the analysis.



**Figure S2.** Characterization of PNBS by (a) thermogravimetric analysis and (b) differential scanning calorimetry (second heating/cooling cycle), acquired using Mettler Toledo instruments at a heating/cooling rate of  $10 \,^{\circ}\text{C}$  / min.

#### II. Additional UV-vis.-NIR Data

To compare the absorption of PNBS upon doping with N-DMBI-H to (N-DMBI)<sub>2</sub> samples prepared in a glovebox in Schlenk cuvettes were doped with different amounts of N-DMBI-H. The samples were then measured without heating. The samples were then heated for various times and measured.



**Figure S3.** UV-vis.-NIR spectra showing reactivity of PNBS with N-DMBI-H in chlorobenzene. The samples were prepared in a nitrogen filled glovebox in schlenk cuvettes to avoid exposure to oxygen and moisture. (a) doping of PNBS with different concentrations of DMBI-H. (b) PNBS pristine solution absorption evolution upon doping with 25% DMBIH with and without heating of the sample. (c) Doped PNBS with DMBIH solutions in chlorobenzene were heated at 100 °C for 10 minutes. (d) PNBS pristine solution absorption evolution upon doping with 50% DMBIH with and without heating of the sample.



Figure S4. UV-Vis-NIR spectra of PNBS films sequentially doped with (RuCp\*mes)<sub>2</sub>.



**Figure S5.** UV-Vis-NIR spectra of a PNBS film sequentially doped with (N-DMBI)<sub>2</sub> (5 mM) showing the effect of exposure to air for various times.

# **III. Additional XRD Data**



Figure S6. GIXRD of PNBS films sequentially doped with (RuCp\*mes)<sub>2</sub>.



### IV. AFM Data

**Figure S7.** AFM images, using a Bruker AFM instrument in tapping mode, where the scan frequency was fixed to 0.5 Hz, measured for the pristine, lightly doped and highest conductivity doped (N-DMBI)<sub>2</sub> doped PNBS films. (a) Pristine PNBS, (b) 2.5 mM doped PNBS, (c) 7.5 mM doped PNBS.