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**Supporting Information** 

## High performance functionalized anthracene organic supercapacitors

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Figure S1 SEM micrographs for A. Pristine *tert*-butyl-ethylene-ketone-ANT, B. PANI, and C-F. Pristine *tert*-butyl-ethylene-ketone-ANT/PANI composite with varying weight ratios.



Figure S2 N<sub>2</sub> adsorption desorption curve for tert-butyl-ethylene-ketone-Anthracene (Surface area: 16.9 m<sup>2</sup>/g), polyaniline (Surface area: 36.5 m<sup>2</sup>/g) and tert-butyl-ethylene-ketone-Anthracene/polyaniline composite (3:7) (Surface area: 43.6 m<sup>2</sup>/g).



Figure S3 A. CV curves for electrode prepared using RGO/AC composite (1:1 weight ratio). B. Corresponding specific capacitance calculated form each of the CV at various scan rate.



Figure S4. Cycling a fresh device to 3000 cycles followed by a 72 hrs resting and then a final 1000 cycles. After resting only, a 0.6% drop in capacitance is observed.



Figure **S5 Equivalent series circuit to fit the EIS measured for aqueous device.** 

NMR spectra were recorded on Bruker DPX 400 instruments. The Chemical shifts, given in ppm, are relative to the residual solvent peaks.

## Spectral data for the anthracene derivatives.

**ketone-ANT**: Following the general procedure **1**, the product **methyl-ethylene-ketone-ANT** was obtained as a pale yellow solid and the structure of the compound was confirmed by <sup>1</sup>H and <sup>13</sup>CNMR data with those reported in previous literature.<sup>1</sup>

<sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>): δ 8.49 (d, *J* = 15.6 Hz, 1H), 8.47 (s, 1H), 8.21 (dd, *J* = 7.6, 3.8 Hz, 2H),

8.03 (dd, J = 7.6, 3.8 Hz, 2H), 7.54 – 7.47 (m, 4H), 6.72 (d, J = 15.6 Hz, 1H), 2.57 (s, 3H).

<sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>): δ 197. 8, 140.6, 136.0, 131.4, 129.5, 129.1, 128.6, 126.6, 125.6, 125.2, 28.2.

**HRMS (ESI)**: calc. for [(C<sub>18</sub>H<sub>14</sub>O)H] (M+H) 247.1123, measured 247.1113.

*tert*-butyl-ethylene-ketone-ANT: Following the general procedure **1**, the product *tert*-butylethylene-ketone-ANT was obtained as a pale yellow solid and the structure of the compound was confirmed by <sup>1</sup>H and <sup>13</sup>CNMR data with those reported in previous literature.<sup>1</sup>

<sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>):  $\delta$  8.63 (d, J = 15.6 Hz, 1H), 8.46 (s, 1H), 8.22 (dd, J = 7.6, 3.8 Hz, 2H),

8.06 (dd, J = 7.6, 3.8 Hz, 2H), 7.52 – 7.49 (m, 4H), 7.10 (d, J = 15.6 Hz, 1H), 1.30 (s, 9H).

HRMS (ESI): calc. for [(C<sub>21</sub>H<sub>20</sub>O)H] (M+H) 288.1514, measured 288.1509.

**cyano-ethylene-ANT**: Following the general procedure **2**, the product **cyano-ethylene-ANT** was obtained as an orange solid and the structure of the compound was confirmed by <sup>1</sup>H and <sup>13</sup>CNMR data with those reported in previous literature.<sup>1</sup>

<sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>): δ 8.96 (s, 1H), 8.66 (s, 1H), 8.10 (d, *J* = 15.6 Hz, 2H), 7.94 (d, *J* = 15.6 Hz, 2H) (dd, *J* = 7.6, 3.8 Hz, 2H), 7.70 − 7.66 (m, 2H), 7.61 − 7.57 (m, 2H).

<sup>13</sup>C NMR (100 MHz, CDCl<sub>3</sub>): δ 160.7, 132.6, 131.0, 129.7, 129.2, 128.5, 126.2, 124.0, 113.0,
111.3.

HRMS (ESI): calc. for [(C<sub>18</sub>H<sub>11</sub>N<sub>2</sub>)H] (M+2H) 256.1000, measured 256.2632.



Figure S6: <sup>1</sup>H and <sup>13</sup>CNMR Spectra of methyl-ethylene-ketone-ANT

Chemical shift (ppm)



Figure S7: <sup>1</sup>H and <sup>13</sup>CNMRSpectra of *tert*-butyl-ethylene-ketone-ANT

Chemical shift (ppm)



## Figure S8: <sup>1</sup>H and <sup>13</sup>C NMR spectra of cyano-ANT



Chemical shift (ppm)



Figure S9. PL spectra of deposited anthracene derivatives on glass slides using ethanol (insets showing glass coated with material under UV). Fluorescence spectroscopy measurements were carried out using a Fluorolog spectrophotometer (HORIBA Scientific, Irvine, CA) (Shown in Figure S7).



Figure S10. SEM micrograph of Polyaniline.



Figure S11. SEM micrograph of reduced graphene oxide.



Figure S12. IV characteristic of composites with varying concentration of PANI and slop indicate with PANI, conductivity increases.

Conductivity of the *tert*-butyl-ethylene-ketone-ANT/PANI thin films (2 cm length).

ANT with PANI	Pure ANT	30% PANI	50% PANI	70% PANI	90% PANI
Material conductivity $(\Omega^{-1} \text{ cm}) \ge 10^{-3}$	3.56	5.50	5.58	6.00	6.46



**Figure S13. Long term stability analysis. A.** Self-discharge behavior of asymmetric nonaqueous device at two different charging voltage after 100 cycles. **B**. Effect of floating at different voltage values.