Polymer-Derived Ceramic Composite Fibers with

Aligned Pristine Multiwalled Carbon Nanotubes

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Supporting Information

- **1. Synthesis of Polyaluminasilazane.** 40.008 g CerasetTM VL20 (purchased from Kion Corporation, Charlotte, NC) and Al sec-butoxide (purchased from Sigma-Aldrich, St Louis, MO) was reacted in a 100 mL conical flask at 125°C for 36 hours under argon (Ar) atmosphere with magnetic stirring. The mixture was cooled for overnight and a solid polymer was obtained.
- 2. Synthesis of the block copolymer poly(3-hexylthiophene)-*b*-poly (poly (ethylene glycol) methyl ether acrylate) (P3HT-b-PPEGA). P3HT macroinitiator (180 mg, 0.028 mmol), CuBr (8.0 mg, 0.056 mmol), PMDETA (0.056 mmol) and PEGA (2.514 g, 5.537 mmol) were dissolved in 4 mL toluene. After degassed by three cycles of freeze-pump-thaw, the polymerization was carried out at 100°C for 20 hours. After getting rid of the copper by passing through a short column, the block copolymer was precipitated in cold diethyl ether for 3 times and then dried under vacuum at 60°C. The molecular weight (M_n) of P3HT-PPEGA was determined by gel permeation chromatography (GPC) to be 16900 with a polydispersity index M_w/M_n of 1.5. The molar ratio of the 3-hexylthiophene unit to PEGA was determined to be 1: 2.8 by comparing the peaks of proton a and b as shown in Figure S3. ¹H NMR (CDCl₃): 6.98 (s, 4 proton of thiophene), 4.17 (br s, CH-COO-CH₂), 3.64 (t, -CH₂-CH₂-OCH₃), 3.39 (s, -CH₂-OCH₃), 2.79 (t, α-CH₂).

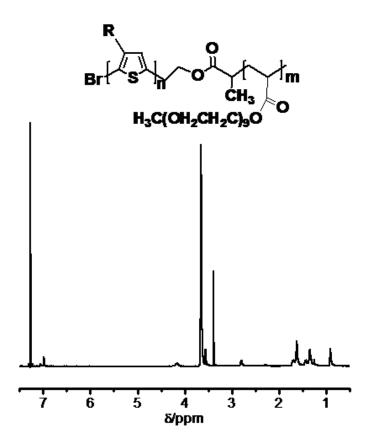


Figure S1. ¹H NMR Spectrum of P3HT-b-PPEGA block copolymer.

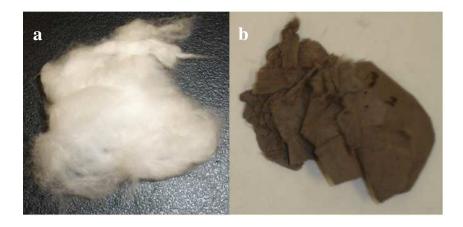


Figure S2. Nonwoven fiber mats of electrospun fibers of (a) pure polyaluminasilazane and (b) polyaluminasilazane/MWCNT (0.3 wt %).

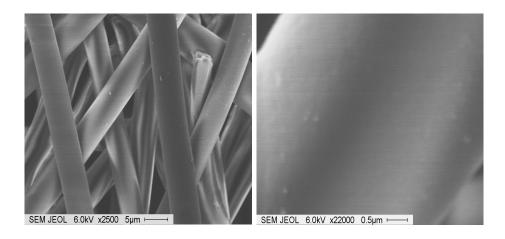


Figure S3. SEM images of polyaluminasilazane fibers electrospun from 20~% polyaluminasilazane chloroform solutions.