Supplemental to the Solvent-Free Functionalization of Carbon Nanotubes by Christopher A. Dyke and James M. Tour*

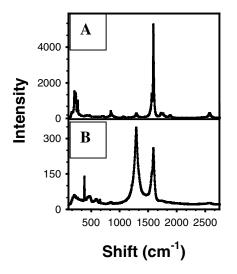
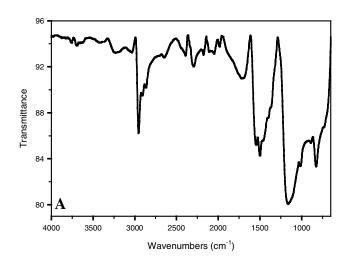
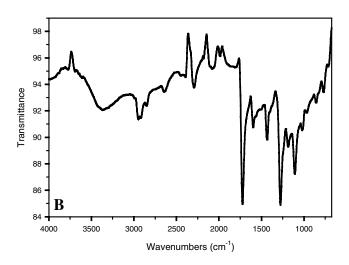


Figure A. Raman spectra (780.6 nm excitation) of (**A**) pristine SWNTs, and (**B**) the product of the diazonium salt of 4-chloroaniline and SWNTs in a ball-miller. Spectrum **B** displays an increased disorder mode compared to pristine SWNTs.

NMR analysis was attempted on the **3** and **4**. However, as is typical in the NMR analysis of SWNTs, we were unable to obtain meaningful ¹H NMR signals (500 MHz, inverse probe) due to any one or more of the following reasons: low concentration, the exceedingly short T₂ relaxation times for these large molecules with their slow tumbling rates, and traces of Fe impurities. CP-MAS NMR (200 MHz wide bore) was equally uninformative showing signals that were too broad to meaningfully interpret.





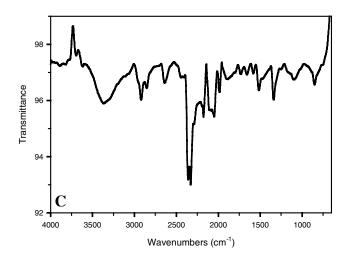


Figure B. ATR-IR spectra of (A) tert-butyl-substituted product 3, (B) aryl ester 4, and (C) aryl nitro 5.