

The use of carbon paste electrode mixed with multiwall carbon nanotube/polyimide composites as electrode for sensing ascorbic acid

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Mass spectrum (m/e): calculated for $C_{40}H_{30}N_4O_7 = 678.6$. Found 677.2. 1H NMR (d_6 -DMSO): $\delta = 8.40$ (s, 1H, due to H_4), $\delta = 8.05$ - 8.03 (d, 1H, due to H_8), $\delta = 7.63$ - 7.60 (2H, due to H_7, H_9), $\delta = 7.29$ - 7.23 (t, 4H, due to H_2, H_6), $\delta = 7.17$ - 7.12 (t, 4H, due to H_3, H_5), $\delta = 6.87$ (t, 1H, due to H_1)

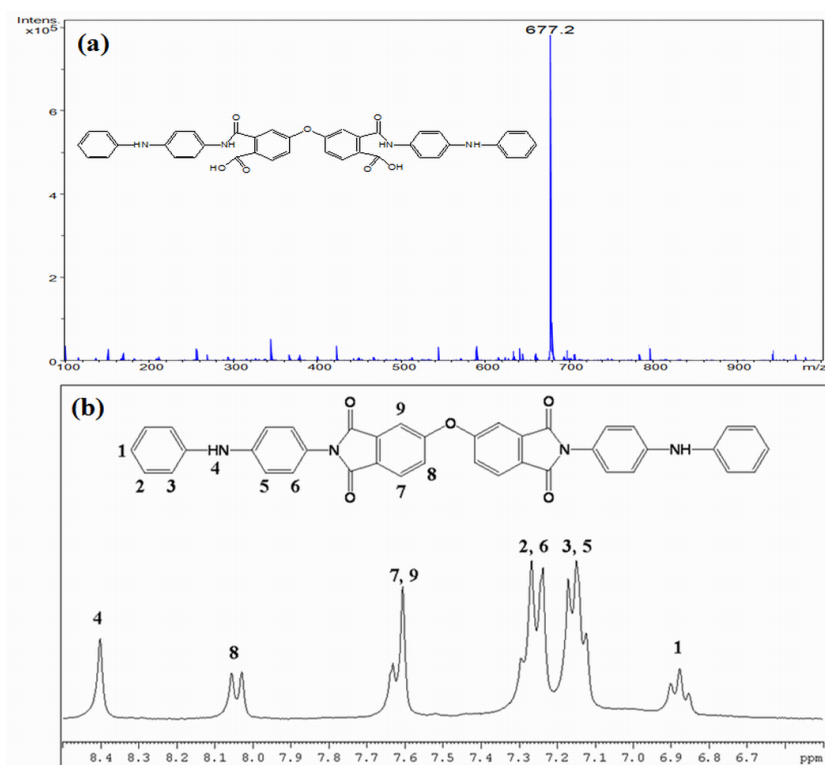


Fig. S1 (a) Mass of oligoaniline and (b) 1H NMR of imidic monomer of oligoaniline.

The detailed characterizations for imidic monomer of oligoaniline was listed as follows: FTIR (KBr, cm^{-1}): 3380 (s, ν_{NH}), 1774 (m, $\nu_{\text{C=O}}$ asymmetric stretching), 1708 (vs, $\nu_{\text{C=O}}$ symmetric stretching), 1592 (s, $\nu_{\text{C=C}}$ of benzenoid rings), 1521 (vs, $\nu_{\text{C=C}}$ of benzenoid rings), 1323 (s, $\nu_{\text{C-N}}$), 1102 (m, δ_{CH}), 890 (m, δ_{CH}), 740 (m, imide ring deformation).

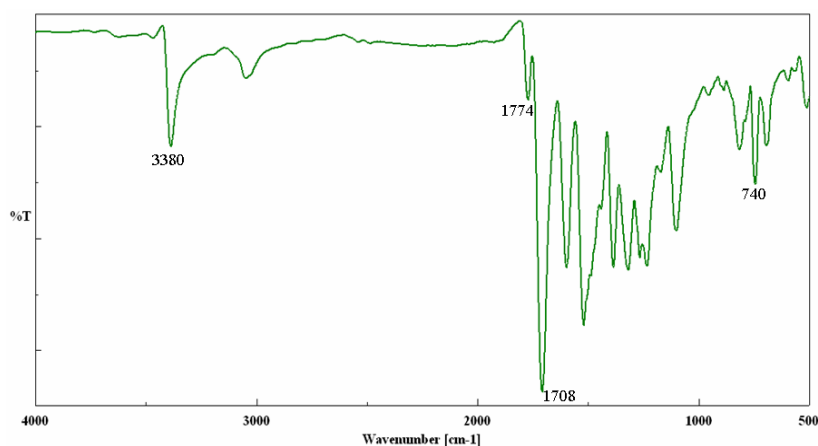


Fig. S2 FTIR spectra of the imidic oligoaniline

The detailed characterizations for EPAA was listed as follows: FTIR (KBr, cm^{-1}): 1711 cm^{-1} (vs, $\nu_{\text{C=O}}$ symmetric stretching), 1590 (s, $\nu_{\text{C=C}}$ of benzenoid rings), 1510 (vs, $\nu_{\text{C=C}}$ of benzenoid rings).

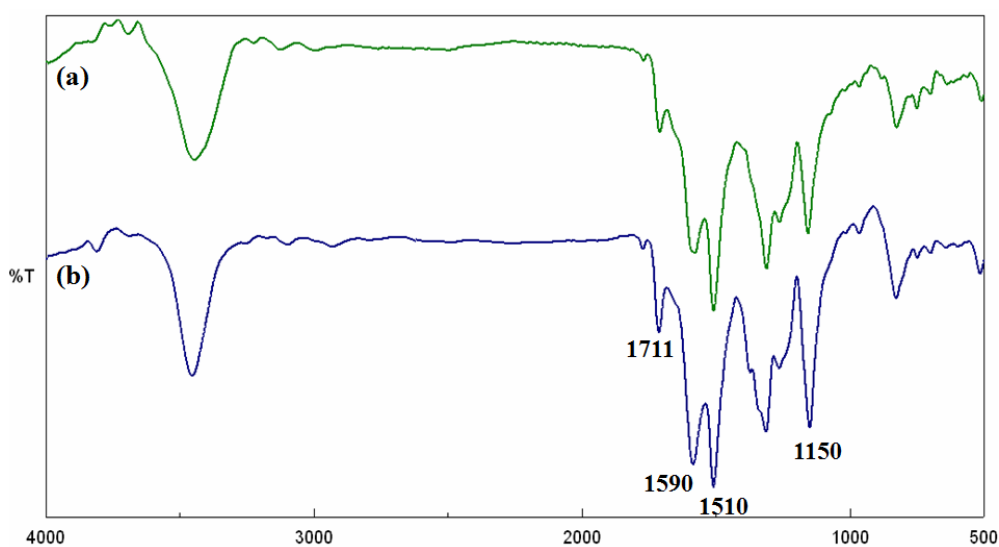


Fig. S3 FTIR spectra of (a) EPAA and (b) AF-MWCNT/EPAA composites

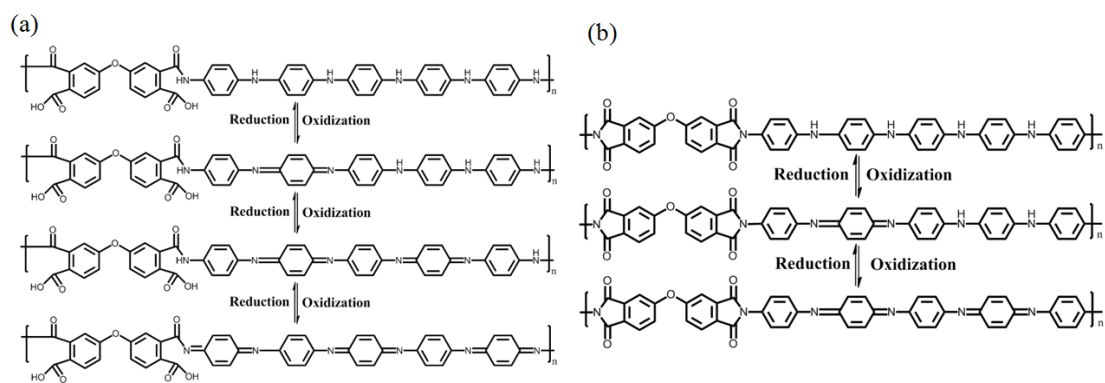


Fig. S4 Molecular structures of (a) EPAA and (b) EPI at various oxidation states

Table S1 The conductivity of AF-MWCNT/EPI

AF-MWCNTs Loading [wt%]	Conductivity (S/cm)
0	5.46×10^{-6}
1	1.32×10^{-5}
3	3.55×10^{-3}
5	3.31×10^{-1}
10	8.87×10^{-1}