

A redox conjugated polymer-based all-solid-state reference electrode

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Supplementary Materials

1. The synthesis scheme and ¹H, ¹³C NMR spectra of the synthesized compounds.

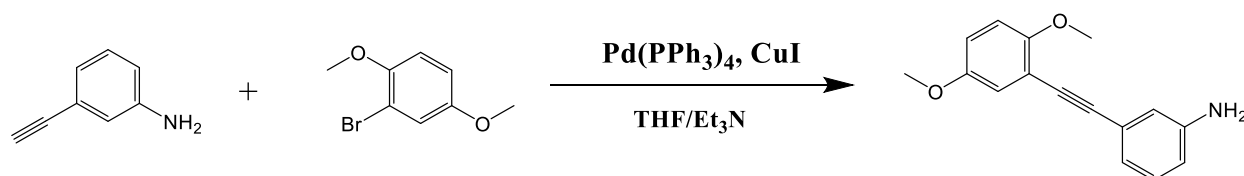


Figure S1. The synthesis scheme

To a N₂ protected 50ml round-bottom flask, Pd(PPh₃)₄ (148 mg, 0.128 mmol, 5% eq), CuI (49 mg, 0.256 mmol, 10% eq), 3-Ethynyl aniline (300 mg, 2.56 mmol), 2-bromo-1,4-dimethoxybenzene (612 mg, 2.8 mmol) were dissolved in 12 mL THF/Et₃N(3/1), then the reaction was heated to 60°C and stirred overnight. The solvents were removed via vacuum, the residue was extracted by ethyl acetate from water, dried over anhydrous sodium sulfate, finally purified by silica gel (R_f=0.35) chromatography (hexane/EA=6/1) to obtain the targeted compound. ¹H NMR (400 MHz, CDCl₃) δ 7.10 (t, *J* = 7.8 Hz, 1H), 7.01 (d, *J* = 2.4 Hz, 1H), 6.95 (d, *J* = 7.6 Hz, 1H), 6.88-6.84 (m, 1H), 6.83-6.79(m, 2H), 6.63 (d, *J* = 12.0 Hz, 1H), 3.85 (s, 3H), 3.76 (s, 3H), 3.66 (br, 2H); ¹³C NMR (100 MHz, CDCl₃) δ 154.7, 153.5, 146.4, 129.4, 124.3, 122.4, 118.3, 118.2, 115.9, 115.5, 113.4, 112.5, 94.6, 93.9, 56.8, 56.0.

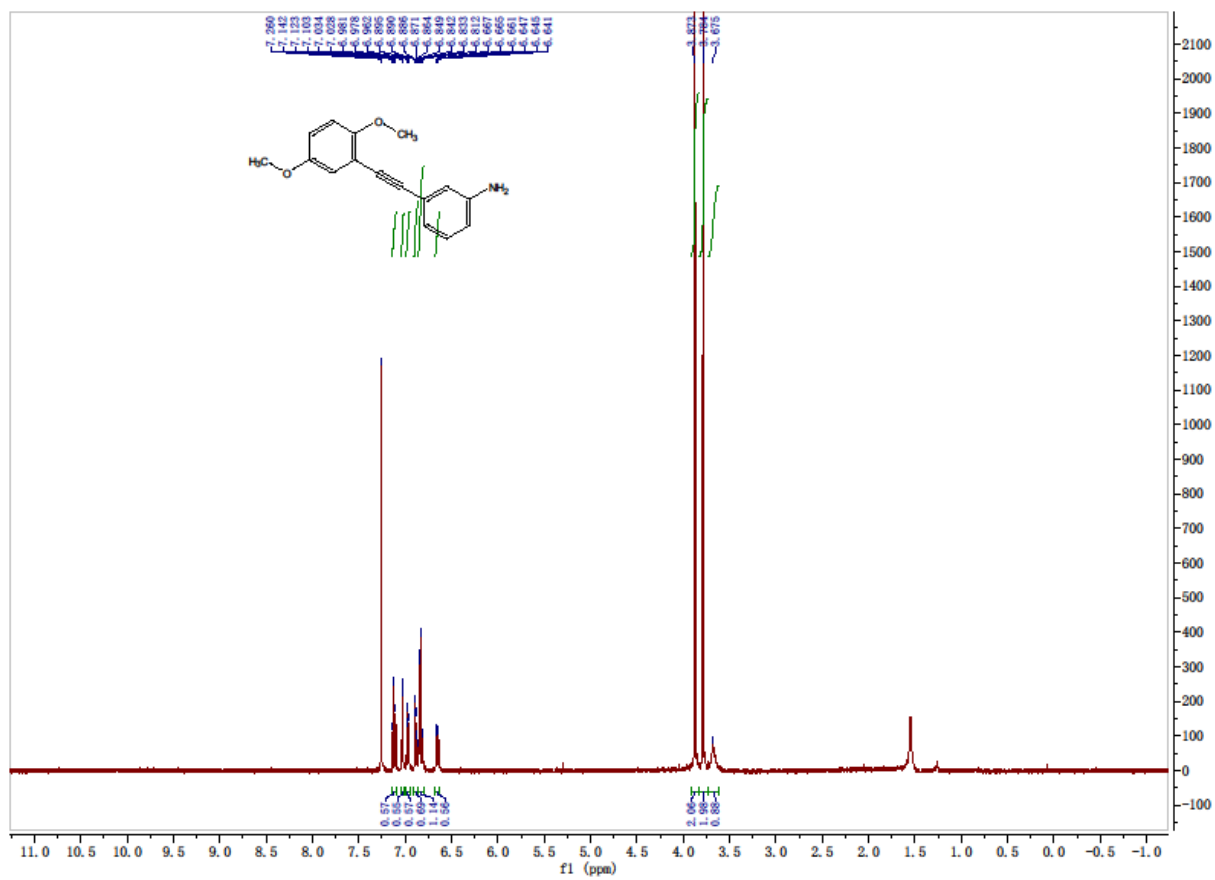


Figure S2. ^1H NMR spectrum

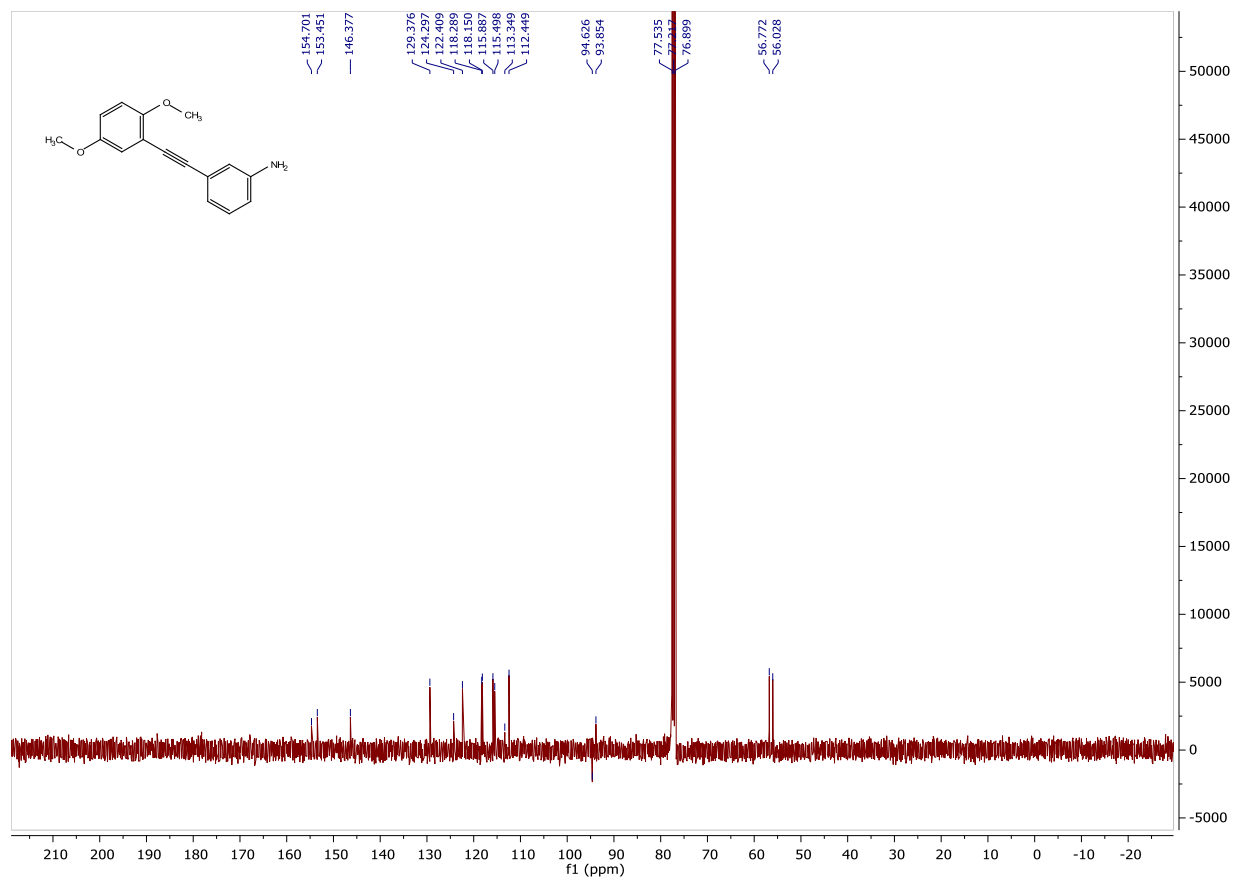


Figure S3. ^{13}C NMR spectrum

FTL-14
 [Elemental Composition]
 Data : 16Feb02_E02-002 Date : 02-Feb-2016 16:15 Page: 1
 Sample: FTL-14
 Note : 70eV
 Inlet : Direct Ion Mode : EI+
 RT : 0.29 min Scan#: (4,5)
 Elements : C 400/0, H 800/0, O 2/2, N 1/1
 Mass Tolerance : 10mmu
 Unsaturation (U.S.) : -0.5 - 10000.0

Observed m/z	Int%	Err [ppm / mmu]	U.S.	Composition
253.1098	100.0	-1.7 / -0.4	10.0	C 16 H 15 O 2 N = 253.1103

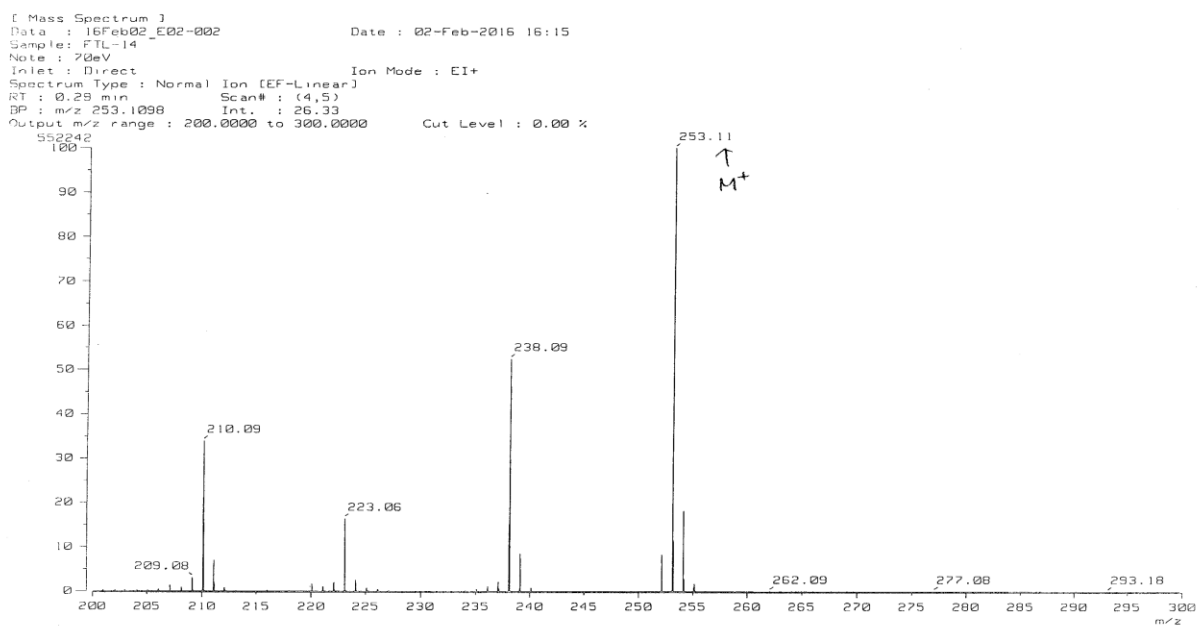
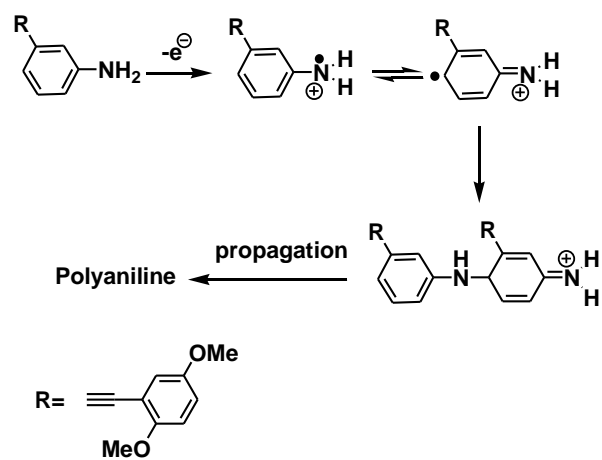


Figure S4. Mass spectrum

Aniline oxidation:



De-methylation by BBr₃

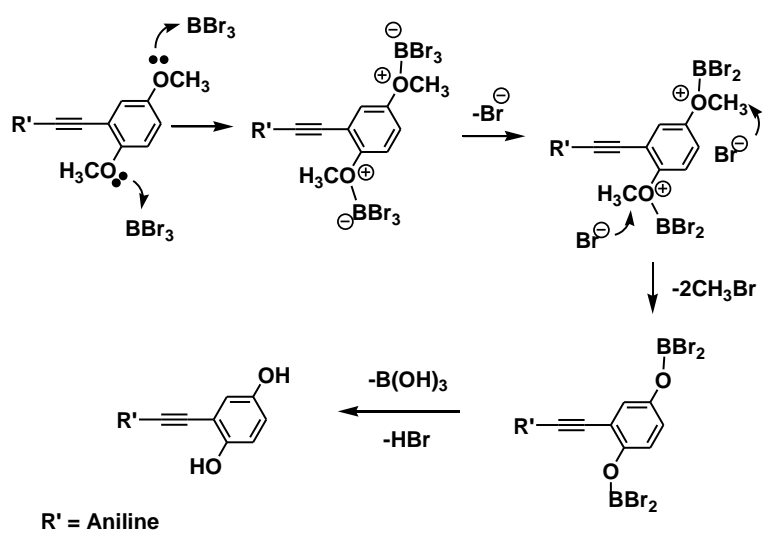


Figure S5. Mechanism of Aniline oxidation and de-methylation by BBr₃

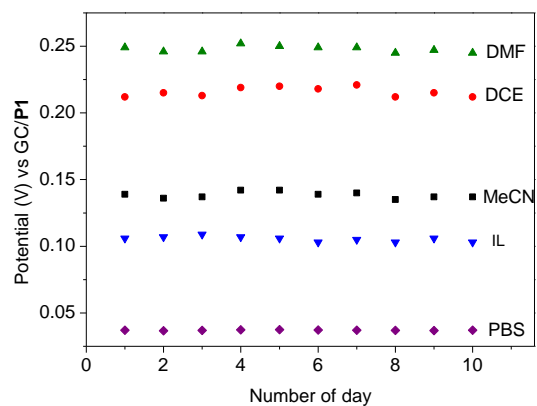


Figure S6. Stability of GC/P1 as RE in DMF, DCE, MeCN, [Bmim][NTf₂] ionic liquid and PBS, with FcMeOH as redox probe and its average potentials plotted.