Supporting Information

Scalable Fabrication of Electrochromic Films Using a Cobalt-Complexed Poly(*N*-vinylcarbazole) Metallopolymer

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Synthesis of 3-([2,2':6',2''-terpyridin]-4'-yl)-9-vinyl-9H-carbazole (3)

Compound 2 0.5 g (1.75 mmol) was dissolved in 10 ml of ethanol in a two neck round bottom flask and then added 2-actylpyridine 0.63 mL (10.5 mmol) and potassium hydroxide 0.34 g (6.1 mmol). The reaction mixture was heated to 60 °C for 1 hour and then added ammonium hydroxide NH₄OH 5 ml and then continued for 10 hr. The precipitated product was collected by filtration and washed with ethanol. The product was dried under vacuum and obtained as a white powder (0.35 g; yield 42%). The product was confirmed by ¹H-NMR: ¹H NMR (400 MHz, CDCl₃) δ 8.86 (s, 2H), 8.80 – 8.68 (m, 4H), 8.63 (d, J = 1.5 Hz, 1H), 8.21 (d, J = 7.6 Hz, 1H), 8.05 (dd, J = 8.6, 1.8 Hz, 1H), 7.90 (td, J = 7.8, 1.7 Hz, 2H), 7.73 (dd, J = 29.0, 8.4 Hz, 2H), 7.57 – 7.47 (m, 1H), 7.41 – 7.30 (m, 4H), 5.62 (dd, J = 15.9, 0.8 Hz, 1H), 5.25 (d, J = 9.1 Hz, 1H); ¹³C NMR (126 MHz, CDCl₃) δ 156.50, 155.80, 150.79, 149.11, 139.91, 136.87, 130.99, 129.41, 126.62, 125.83, 124.61, 123.93, 123.69, 121.41, 120.96, 120.57, 119.22, 118.82, 110.77, 110.63, 102.92.

Synthesis of PVK-3-tpy

0.1 g (0.23 mmol) of compound 3 was dissolved in 5 mL of dry tetrahydrofuran (THF) under argon atmosphere and added azobisisobutyronitrile (AIBN) 0.01 g and then heated to 60 $^{\circ}$ C and continued for 12 h. After that the reaction mixture was concentrated by evaporating the excess solvent and then precipitated by adding dropwise into methanol. The precipitated solid was collected by filtration and dried under vacuum. The ¹H-NMR study confirms that the absence of vinyl bond in the polymer product indicated the polymerized product was obtained.

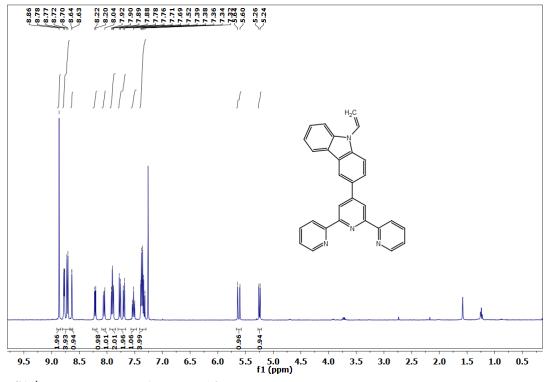


Figure S1. ¹H-NMR spectrum of compound **3**.

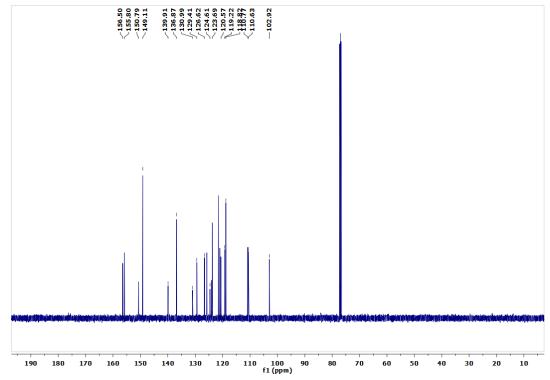


Figure S2. ¹³C-NMR spectrum of compound 3.