

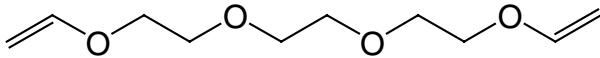
Supporting Information

General Procedures. All starting materials were commercially available and used after distillation or recrystallization. GLC analysis was performed with a flame ionization detector using a 0.2 mm x 25 m capillary column (OV-1). ^1H - and ^{13}C -NMR were measured at 400 MHz and 100 MHz, respectively, in CDCl_3 with Me_4Si as the internal standard. Infrared (IR) spectra were measured using NaCl plates.

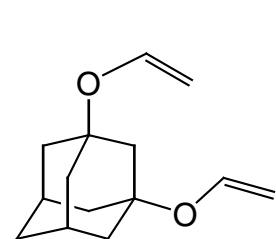
General Procedure for the Reduction of Alcohol with Vinyl Acetate. A typical reaction procedure is as follows: To a toluene solution (1.0 mL) of $[\text{IrCl}(\text{cod})]_2$ (0.01 mmol) and Na_2CO_3 (0.6 mmol) were added alcohol (1 mmol) and vinyl acetate (2 mmol) under Ar. The reaction mixture was stirred at 100 °C for 2 h. After quenching with wet ether, the product was isolated by column chromatography (230-400 mesh silica gel, *n*-hexane). After the reaction, the GC and GC-MS analyses were performed. The conversions and yields of products were estimated from the peak areas based on the internal standard technique using GC. The products **4**,¹ **6**,² **8**,³ **9**,⁴ **13**,⁵ **15**,⁶ **17**,⁷ **21**,⁸ **23**,⁹ **27**¹⁰ and **28**¹¹ were reported previously.

Spectral Data

{2-[2-(2-Vinyloxyethoxy)ethoxy]ethoxy}ethene (**11**):

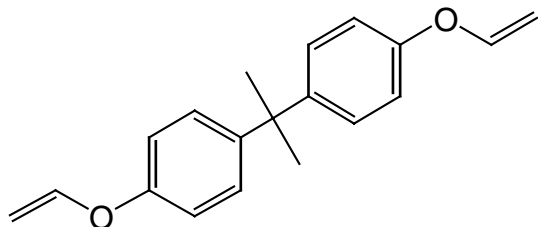
 ^1H NMR 6.50 (dd, $J = 6.8$ and 14.2 Hz, 1H), 4.19 (dd, $J = 2.0$ and 14.2 Hz, 1H), 4.01 (dd, $J = 2.0$ and 6.8 Hz, 1H), 3.83 (t, $J = 4.9$ Hz, 1H), 3.85 (d, $J = 2.9$ Hz, 2H), 3.73 (t, $J = 4.9$ Hz, 1H), 3.75 (d, $J = 2.9$ Hz, 1H), 3.69 (s, 2H); ^{13}C NMR 151.6, 86.5, 70.7, 69.6, 67.2; IR (neat) 2876, 1619, 1322, 1203, 1129 cm^{-1} .

1,3-Bis(vinyloxy)adamantane (**19**):



^1H NMR 6.45 (dd, $J = 6.3$ and 13.6 Hz, 1H), 4.45 (dd, $J = 1.0$ and 13.6 Hz, 1H), 4.07 (dd, $J = 1.0$ and 6.3 Hz, 1H), 2.38 (t, $J = 2.9$ Hz, 1H), 1.90 (s, 1H), 1.77 (s, 4H), 1.53 (t, $J = 2.9$ Hz, 1H); ^{13}C NMR 144.5, 91.5, 76.9, 46.1, 40.6, 34.8, 30.8; IR (neat) 2913, 1628, 1151, 1109, 1038 cm^{-1} .
Anal. Calcd for $\text{C}_{14}\text{H}_{20}\text{O}_2$: C, 76.33; H, 9.15. Found: C, 76.34; H, 9.27.

Bisphenol-A divinyl ether (25):



^1H NMR 7.18-7.14 (m, 3H), 6.91-6.88 (m, 2H), 6.62 (dd, J = 6.3 and 13.7 Hz, 1H), 4.73 (dd, J = 1.5 and 13.7 Hz, 1H), 4.39 (dd, J = 1.5 and 6.3 Hz, 1H), 1.65 (s, 3H); ^{13}C NMR 154.6, 148.2, 145.3, 127.8, 116.4, 94.7, 42.1, 31.0; IR (neat) 3036, 1644, 1504, 1385, 964, 836, 552 cm^{-1} . Anal. Calcd for $\text{C}_{14}\text{H}_{20}\text{O}_2$: C, 81.40; H, 7.19. Found: C, 81.29; H, 7.19.

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