Poly[ethynylene(3-n-butyl-2,5-thiophenediyl)-ethynylene]: A Soluble Polymer Containing Diacetylene Units and Its Conversion to a Highly Cross-Linked Organic Solid

We are engaged in a program directed at the synthesis and study of highly cross-linked organic solids. Our approach has been to prepare low molecular weight polymers and oligomers consisting of aromatic moieties linked by diacetylene units, to cross-link the diacetylene units thermally, and to examine the properties of the resulting solids. Our initial efforts were directed toward the preparation of monomers having as high a ratio of carbon to hydrogen as possible. Although this approach yielded materials with interesting properties (high Young's modulus, hardness, and thermal stability) the oligomeric/polymeric intermediates had poor solubility and high cure energies (in some cases exceeding 500 J/g); these values of cure energy made the preparation of large pieces difficult.

We chose to introduce an alkyl group into the aromatic portion of the monomer to increase the solubility and to lower the cure energy of the polymer derived from it. This paper reports our results of a study of poly[ethynylene-(3-n-butyl-2,5-thiophenediyl)ethynylene] and related oligomers and polymers: preparation of these diacetylenic oligomers and polymers in soluble form, conversion of the oligomers to hypercross-linked organic solids with high carbon content, and characterization of these solids. An accompanying paper by Rutherford and Stille describes related work.4 Our studies demonstrate that the properties of poly[ethynylene-(3-n-butyl-2,5-thiophenediyl)ethynylene] derived from the oxidative coupling of 3-n-butyl-2,5-diethynylthiophene (1) can be controlled by the addition of 2-ethynylthiophene (2) as an end-capping agent. We have also prepared 2,3,5-triethynylthiophene (3) and carried out the oxidative copolymerization of 1, 2, and 3 in a 10:1 molar ratio to investigate the properties of a diacetylene network polymer.

Scheme I outlines the syntheses of 1 and 3. The nickel-catalyzed6 coupling of 3-bromothiophene (4) with n-butylmagnesium bromide in ether, followed by treatment of the resulting 3-n-butylthiophene with bromine in acetic acid,9 gave 3-n-butyl-2,5-dibromoethynylene (5) in an overall yield of 87%. The palladium- and copper-catalyzed coupling of the dibromide (trimethylsilyl)acetylene followed by deprotection of 3-n-butyl-2,5-bis[(trimethylsilyl)ethynyl]thiophene with potassium hydroxide in methanol gave 1 in >99% yield.9 We prepared 2,3,5-triethynylthiophene (3) from 2,3,5-tribromoethynylene (6) in 74% overall yield using a procedure analogous to that used for the conversion of 5 to 1.

Polymerization of 1 by oxidative coupling of the acetylenes with dioxygen and catalytic amounts of cuprous chloride in o-dichlorobenzene for 10 min at 65 °C followed by cooling over 2 h to room temperature gave a completely homogeneous solution of the polymer represented by the structure 7 (Scheme II, eq 3).12 Copolymerization of 1 and 2 (added as an end-capping agent) in molar ratios of 1:2 = 3:1, 5:1, 10:1, 100:1, and 1000:1 all gave completely homogeneous solutions in o-dichlorobenzene of the polymers 9a–e (eq 4). The oxidative copolymerization of 1, 2, and 3 in molar ratios of 10:1:1, respectively, gave the network polymer 11 (eq 5), also soluble in o-dichlorobenzene.

Examination of 9a by 1H NMR spectroscopy clearly showed resonances assignable to the end-capping groups. At molar ratios of 1:2 > 10:1, resonances of the end-capping groups were no longer observed in the 1H NMR spectrum. Infrared spectral analysis did not show acetylenic C–H stretching vibrations in any of the polymers prepared.13 Analysis by GPC confirms that added 2 does indeed decrease the molecular weight.14 The number-average molecular weights (Mn) and polydispersities (Z) of 9a–e decrease with increasing amount of end capping: 9a (Mn = 3700; Z = 1.9); 9b (Mn = 4500; Z = 2.1); 9c (Mn = 6000; Z = 2.3); 9d (Mn = 13000; Z = 2.1); 9e (Mn = 16000; Z = 2.2). The addition of the triacetylenic monomer 3 for the preparation of 11 resulted in an increase in both molecular weight (Mn = 9000) and polydispersity (Z = 2.7) relative to 9c. The polymers 9a–e are soluble in solvents such as tetrahydrofuran and toluene. The lower molecular weight materials, 9a and 9b, are also soluble in methylene chloride. The solubilities of 7, 9a–e, and 11 are remarkable in comparison to that of the oligomer 13 derived from the oxidative polymerization of 2,5-diethynylthiophene (14).3 We found that 13 was soluble only at low molecular weights, when formed by coupling of a 1:1 molar ratio of 14 and 2.

The hypercross-linked solids 8, 10a–e, and 12 were prepared by molding the oligomers/polymers under pressure (10 000 lb/in.2) into disks (ca. 1 cm in diameter and 0.25 mm thick) and heating to 200 °C for 1 h. The cross-linking processes of the oligomers and polymers, 7 and 9a–e, are characterized (DSC, ramp rate = 5 °C/min) by a cure onset temperature of ~160 °C, a cure maximum temperature of ~180 °C, and a cure energy of ~300 J/g. As expected, 11 had a lower cure onset temperature of 110 °C, a lower cure maximum temperature of 137 °C, and a higher cure energy of 350 J/g reflecting the higher reactivity of the triacetylenic monomer unit. The thermal stabilities (TGA, ramp rate = 10 °C/min) of 8, 10a–e, and 12 were virtually indistinguishable: under argon, the characteristic temperature (T10%) at which they had lost...
10% of their weight was $T_{10\%} = 470$ °C; the corresponding temperature for 50% weight loss was $T_{50\%} = 950$ °C. Under air, all had a $T_{10\%} = 425$ °C and a $T_{50\%} = 525$ °C. These values are similar to those observed for analogous nonbutylated materials.\(^2\) For example, the organic solid derived from the thermal cross-linking of 13, poly(2,5-diethynylthiophene), derived from the oxidative polymerization of 14 and 2, exhibited, under argon ($T_{10\%} = 540$ °C, $T_{50\%} > 1100$ °C\(^{15}\)) and air ($T_{10\%} = 330$ °C, $T_{50\%} = 400$ °C), thermal stability similar to those of 8, 10a--e, and 12.\(^2\)

Although the disks that we prepared from 8, 10d, 10e, and 12 were extremely brittle and had surprisingly low densities: $\rho \approx 1.2$ g/cm\(^3\),\(^{16,17}\) those from 10a--c formed solids having good mechanical properties. The black, shiny disks formed from 10a--c had $\rho \approx 1.34$ g/cm\(^3\) and Young's moduli (~14000 MN/m\(^2\)) clearly indicative of a highly cross-linked organic solid.\(^{16,17}\) We have also found that smooth, noncracking, yellow films can be cast on glass from toluene or tetrahydrofuran solutions. When thermally treated at 200 °C the resultant adherent, shiny, black films cracked due to shrinkage.

The solubility, Young's modulus and thermal stability of poly[ethynylene(3-n-butyl-2,5-thiophenediyli)-ethynylene] and related oligomers and polymers provide an attractive system for the exploration of the properties of hypercross-linked solids. We are continuing to investigate these materials.

**Acknowledgment.** We are grateful to the National Science Foundation for support of this research (Grant CHE-85-08702). We thank Drs. Jim Barber, Howard Tennant, and Larry Truesdale of Hyperion Catalysis In-
ternational, Lexington, MA, and Dr. Owen Webster of E. I. du Pont deNemours and Co. for helpful discussions. We thank Dr. Timothy M. Miller (Massachusetts Institute of Technology) for assistance. We thank Professor Richard R. Schrock, Gui Bazan, and Steven A. Krouse (Massachusetts Institute of Technology) for allowing us to use the GPC equipment and computer programs in their laboratory to characterize these polymers. We also thank Professor John K. Stille (Colorado State University) for discussions concerning this class of polymers. Thermal analysis was carried out by using an apparatus at Hyperion Catalysis International.

Registry No. 1 (homopolymer), 116954-15-5.

References and Notes

(3) We refer to these materials—solids designed to maximize the volume densities of the bonds linking the chains of the polymeric precursors—as hypercross-linked solids.
(9) Spectral and elemental analyses of 3, the trimethylsilyl-protected ethynylated monomers [3-n-butyl-2,5-bis[trimethylsilyl]ethynyl]thiophene and 2,3,5-tris[(trimethylsilyl)ethynyl]thiophene, and the polymers (7, 9a-e, 11) were satisfactory. The deprotected monomers (1-3) have spectral analyses consistent with the assigned structures.
(10) Caution: Compound 3 was isolated only in small (milligram) amounts. Only limited quantities should be stored or manipulated as pure (undiluted) material. Although we have had few incidents with these substances, all deprotected ethynyl aromatics and oligomeric and polymeric materials mentioned in this paper should be treated as potentially explosive materials.
(13) We estimate the limits of detection by infrared analysis to be approximately one free acetylene per 125 3-n-butyl-2,5-di-ethynylthiophene units.
(14) Number-average molecular weights (Mn) and polydispersities of these polymers were measured in tetrahydrofuran and referenced to polystyrene standards by using a Polymer Laboratories PLgel 10 μm polystyrene mixed-bed resin column with a range of Mn of 500-4 x 10⁶. Because the reference standards are of polystyrene polymer, the absolute values of molecular weights reported here are probably less accurate than the relative values. The chromatograms were analyzed by using software developed in the laboratories of Professor Richard R. Schrock at the Massachusetts Institute of Technology.
(15) For this case, T₉₀% exceeded the upper temperature limit of the thermal analyzer (1100 °C). At 1100 °C, 70% of the mass of the organic solid derived from the polymer 13 was retained.
(16) Densities of typical organic polymers range from 0.94 g/cm³ (polyethylene) to ~1.33 g/cm³ (Thermid-600, a polyimide thermoset polymer) with Young's moduli from 2450 MN/m² (polyethylene) to 4650 MN/m² (Thermid-600).

Matthew R. Callstrom, Thomas X. Neenan, and George M. Whitesides*
Department of Chemistry, Harvard University Cambridge, Massachusetts 02138
Received May 23, 1988; Revised Manuscript Received September 13, 1988