Conjugated Poly(phenylacetylene) Films Cross-Linked with Electropolymerized Polycarbazole Precursors

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ABSTRACT: In this work, we describe the synthesis and electropolymerization of conjugated substituted polyacetylenes, poly(N-alkoxy-(3-ethynylphenyl)carbazole) or PPA-Cz-CN with electropolymerizable carbazole side groups to form conjugated polymer network (CPN) films. The phenylacetylene monomer was functionalized with a carbazole group separated by an alkylene spacer. Polymerization of the monomer in solution is accomplished using a Rh catalyst to form a “precursor polymer”. The electrochemical behavior and cross-linking of the carbazole side group was then investigated by cyclic voltammetry (CV) and spectroelectrochemistry. A trend in the redox electrochemical behavior was observed with varying spacer length between the poly(phenylacetylene) backbone and carbazole side-group. The resulting film combines the electrooptical properties of the conjugated poly(phenylacetylene) polymer with polycarbazole units in a cross-linked electropolymerized film as evidenced by the CV and spectroelectrochemical behavior. This study emphasizes the preparation of polymer materials with mixed π-conjugated species arising from the electrochemical cross-linking of a designed precursor polymer.

Introduction

Polyacetylene is one of the first reported conductive polymers with both high conductivity and interesting electrooptical properties. They have been synthesized using a variety of catalysts and polymerization conditions, often under inert atmosphere. Substituted polyacetylenes derivatives have also been extensively studied and are readily synthesized by a variety of metal carbene and metal halide catalysts. A key advantage of substituted polyacetylenes is their stability and solubility compared to polyacetylenes at ambient conditions. Using metathesis polymerization techniques with group V and VI metal catalysts, their configuration and conformation (cis-trans and cisoidal-transoidal) can be controlled toward higher or lower π-conjugation lengths. On the other hand, Rh catalysts are known to control the formation of cis and trans configurations on the polymer backbone primarily through an insertion mechanism. The polymerization of poly(phenylacetylene)s (PPA) is well-known and has been thoroughly investigated. While substituted polyacetylenes have been investigated for various electrooptical and nonlinear optical applications, they have been less studied as materials for electroluminescent devices and organic semiconductors. However, several reports by our group and others have shown electroluminescence properties based on fluorophore incorporation and active charge-carrier transport property optimization for these type of polymers.

Poly(N-vinylcarbazole) (PVK) materials have been extensively studied. They exhibit interesting electrical and optical properties as light emitting diode materials, solar cells, and applications in various electrochromic devices. Additionally, PVK shows good hole-transport properties which have important applications for improving the performance of organic electroluminescent devices. Others have reported applications in amperometric chemical sensors. PVK exhibits these interesting properties primarily due to the carbazole group. On the other hand, an all 3,6-position linkage of the carbazole units will lead to the formation of conjugated poly(3,6-N-vinylcarbazole). The synthesis of polycarbazoles (PCz) is also of great interest for electrical conductivity and electrochromic device applications. Several groups have investigated its electropolymerization on Pt and Au substrates.

Recently, several groups including our group have reported the conversion of “precursor polymers” to form conjugated polymer network (CPN) films on conducting surfaces. The approach consists of synthesizing a precursor polymer, which by design contains pendant electroactive monomer units. Electropolymerization or chemical oxidation results in a conjugated polymer network having both intra- and intermolecular cross-linkages between the pendant monomer units. The films formed are characterized by high optical quality (transparency), uniform coverage, good adhesion, smoothness in morphology, and controlled ion permeability. Moreover, by controlling the amount of conjugated species and doping, it should be possible to control electrical conductivity. Thus, the process is interesting for depositing insoluble cross-linked ultrathin films of conjugated polymers for practical electrooptical applications. A variety of combinations should be possible for the design of a precursor polymer backbone and the “electroactive monomer” side group to form the CPN film. It is also possible to copolymerize with small molecule electroactive monomers with varying compositions in order to control the degree of cross-linking and linear polymer formation. We have applied these materials to a number of possible applications toward electroluminescent devices. We have also previously reported the surface grafting of carbazole-functionalized polyfluorenes on carbazole self-assembled monolayer (SAM) modified indium tin oxide (ITO) surfaces.

With the interest in combining the electrooptical properties of PPA and PCz in a CPN film, we have synthesized a series of substituted poly(phenylacetylene)s containing carbazole unit side groups. As previously mentioned, the stereochemical
troluminescence properties of soluble poly(phenylacetylene)s previously reported the synthesis, characterization, and elec-

tromicrostructure of poly(phenylacetylene) can be controlled using polymer and a cross-linked unit; (b) electrografting of the polymer to 

Electropolymerization and cross-linking scheme for carbazole Figure 1.

![Figure 1. Electropolymerization and cross-linking scheme for carbazole substituted polyacetylene: (a) the chemical structure of the precursor polymer and a cross-linked unit; (b) electrografting of the polymer to a SAM-modified ITO electrode surface.](image)

microstructure of poly(phenylacetylene) can be controlled using metal catalysts. Rh catalysts can preferentially give cis-rich polymers while W catalysts give trans-rich polymers. We have previously reported the synthesis, characterization, and electroluminescence properties of soluble poly(phenylacetylene)s with pendant carbazole groups. The effective conjugation length of the PPA backbone played an important role in photo(luminescence (PL) quantum efficiency and photoconduct-

Experimental Section

Materials. All the reagents for monomer synthesis were commercially available and used without further purification. The toluene solvent used in the polymerization was distilled and purified over CaH₂. The Rh catalyst, [nbd][RhCl]₂, was purchased from Aldrich and used as received. For the electrochemical studies, tetrabutylammonium hexafluorophosphate (TBAH), also from Aldrich) was dried in vacuo at 40 °C for a day. Predistilled THF was dried by refluxing over a benzophenone-sodium complex for a half day.

Monomer Synthesis for PPA-Cz-Cn series. Carbazole substituted phenylacetylene monomers with various alkyl chain lengths, e.g., C3, C6, and C8 (abbreviated as Cz-C3, Cz-C6 and Cz-C8, respectively) were synthesized. Monomer Cz-C3 was synthesized according to Scheme 1. The total yield of Cz-C3 was 58%. The synthetic route to monomer Cz-C6 and Cz-C8 is shown in Scheme 2. The yields of Cz-C6 and Cz-C8 were 39% and 49%, respectively. The structure and purity of the monomers were confirmed by NMR, IR, and elemental analysis and is described as follows:

\[ \text{N-Ethylp-ethynylbenzoxylcarbazole, Cz-C3. Yellowish crystal. Yield: 58%. Mp: 94.0 °C. H NMR (400 MHz, CDCl₃): } \delta \text{ (ppm) 3.04 (s, 1H), 3.85 (t, 2H, J = 5.6 Hz), 4.37 (s, 2H), 4.51 (t, 2H, J = 5.6 Hz), 7.05 (d, 2H, J = 8.8 Hz), 7.20–7.45 (m, 8H), 8.09 (d, 2H, J = 8.0 Hz). IR (KBr, cm}^{-1}: 3274, 2888, 2865, 1626, 1597, 1485, 1452, 1327, 1152, 1021, 745, 725. Anal. Calcd For C_{28}H_{29}NO: C, 85.02; H, 7.75; N, 3.75; O, 4.15. Found: C, 84.80; H, 7.43; N, 3.27; O, 4.15.}

\[ \text{N-Hexoyp-ethynylphenylcarbazole, Cz-C6. White crystal. Yield: 39%. Mp: 81.6 °C. H NMR (400 MHz, CDCl₃): } \delta \text{ (ppm) 1.40–1.55 (m, 4H), 1.74 (m, 2H), 1.93 (m, 2H), 3.02 (s, 1H), 3.89 (t, 2H, J = 6.0 Hz), 4.34 (t, 2H, J = 6.8 Hz), 6.77 (d, 2H, J = 8.8 Hz), 7.20–7.48 (m, 8H), 8.10 (d, 2H, J = 8.0 Hz). IR (KBr, cm}^{-1}: 3256, 3052, 2940, 1607, 1509, 1438, 1426, 1453, 1327, 1154, 1021, 750, 725. Anal. Calcd For C_{30}H_{28}NO: C, 84.98; H, 6.86; N, 3.81; O, 4.35. Found: C, 84.76; H, 6.83; N, 3.57; O, 4.13.}

\[ \text{N-Octoyp-ethynylphenylcarbazole, Cz-C8. White crystal. Yield: 49%. Mp: 80–81 °C. H NMR (400 MHz, CDCl₃): } \delta \text{ (ppm) 1.10–1.50 (m, 8H), 1.73 (m, 2H), 1.86 (m, 2H), 2.98 (s, 1H), 3.89 (t, 2H, J = 6.4 Hz), 4.28 (t, 2H, J = 6.8 Hz), 6.78 (d, 2H, J = 8.8 Hz), 7.20–7.48 (m, 8H, J = 8.0 Hz), 8.09 (d, 2H). IR (KBr, cm}^{-1}: 3256, 2855, 1607, 1509, 1438, 1463, 1452, 1327, 1154, 1021, 1015, 750, 727. Anal. Calcd For C_{32}H_{30}NO: C, 85.02; H, 7.39; N, 3.54; O, 4.04. Found: C, 84.80; H, 7.43; N, 3.27; O, 4.15.}

Polymization To Form PPA-Cz-Cn. Poly(phenylacetylene)s were prepared by metal-catalyzed polymerization using a Rh catalyst according to the literature. Polymizations were performed in a Schlenk tube equipped with a three-way stopcock under nitrogen atmosphere. Unless otherwise specified, the polymerization conditions were as follows: In toluene, 30 °C, 24 h; [Mₙ] = 200 mM, [Cat] = 1.0 mM, and [Cocat] = 2 mM. Polymers were isolated by precipitation in a large excess of methanol, separated by filtration, and the yields of the polymers determined by
for 3 days. The slides were then rinsed successively with toluene, K804, K805, and J806 with CHCl3 as eluent at a flow rate of 1.0 mL/min for chromatography (GPC) on a Jasco Gulliver system (PU-980, CO-Cz-C6, and triethoxy(11-carbazoylundecyl)silane solution in toluene at 60 °C).

**Results and Discussion**

**Synthesis of Triethoxy(11-carbazoylun-decyl)silane SAM.** The synthesis of triethoxy (11-carbazoylundecyl) silane was carried out as previously reported.23 The 9-undec-10-enyl-9H-carbazole was synthesized by the reaction between carbazole and 11-bromoundec-l-ene with potassium hydroxide in 20 mL DMF at room temperature. 9-Undec-10-enyl-9H-carbazole, triethoxysilane, and H2PtCl6 were charged in a one-neck flask under nitrogen. The flask was sealed and put into a sonication bath to yield the triethoxy(11-carbazoylundecyl)silane.26

**SAM Modification of the ITO.** The triethoxy(11-carbazoylundecyl)silane (Figure 1) was used to modify all the ITO surfaces by SAM. ITO was cleaned using standard sonication solutions (2% glutathione solution) followed by copious water rinsing and last by argon ion plasma treatment. The ITO was treated by the 1% triethoxy(11-carbazoylundecyl)silane solution in toluene at 60 °C for 3 days. The slides were then rinsed successively with toluene and THF then dried by a dry nitrogen flow.

**Instrumentation and Electropolymerization Setup.** The molecular weights of the polymers were determined by gel permeation chromatography (GPC) on a Jasco Gulliver system (PU-980, CO-965, RI-930, and UV-1570). Elution was done on Shodex columns K804, K805, and J806 with CHCl3 as eluent at a flow rate of 1.0 mL/min and calibrated with polystyrene (PSi) standards. FT-IR spectra, UV−vis spectra, and NMR spectra were recorded on a Shimadzu FT-IR-8100 spectrophotometer, a Jasco V-550 spectrophotometer, and a JEOL EX-400 spectrometer, respectively. Thermogravimetric analysis (TGA) was conducted in air on a Perkin-Elmer TGA7 thermal analyzer.

The cyclic voltammetry (CV) experiments were carried out on a Princeton Applied Research Parstat 2263 with a modified ITO substrate as the working electrode coupled with a Pt plate counter and Ag/AgCl reference electrode. Cyclic voltammetry was utilized to prepare the cross-linked films from a 0.1 wt % of the precursor polymer solution of 0.1 M TBAH/CH2 Cl2, where TBAH = tetrabutylammonium hexafluorophosphate. Copious washing with CH2 Cl2 solvent and dry nitrogen drying was done prior to any film analysis.

Electrochemical-quartz crystal microbalance (QCM) measurements were performed on a R-QCM (Maxtek Inc.) with an Amel 2049 potentiostat/galvanostat and Power lab system utilizing the QCM crystal as the working electrode and a Pt plate as counter. The cyclic voltammetry (CV) experiments were carried out on a R-QCM (Maxtek Inc.) with an Amel 2049 potentiostat/galvanostat and Power lab system utilizing the QCM crystal as the working electrode and a Pt plate as counter. The cyclic voltammetry (CV) experiments were carried out on a R-QCM (Maxtek Inc.) with an Amel 2049 potentiostat/galvanostat and Power lab system utilizing the QCM crystal as the working electrode and a Pt plate as counter. The cyclic voltammetry (CV) experiments were carried out on a R-QCM (Maxtek Inc.) with an Amel 2049 potentiostat/galvanostat and Power lab system utilizing the QCM crystal as the working electrode and a Pt plate as counter. The cyclic voltammetry (CV) experiments were carried out on a R-QCM (Maxtek Inc.) with an Amel 2049 potentiostat/galvanostat and Power lab system utilizing the QCM crystal as the working electrode and a Pt plate as counter.

**Characterization of the polymers** was also done by IR and NMR. The IR spectra of the polymers exhibited no absorption band around 3300 cm−1, usually attributed to the stretching vibration of C=C bond, which was seen in the monomers. In the 1H NMR spectra of the polymers, a signal at around 3.0 ppm assigned to the acetylenic proton of monomers was not observed. All of these results clearly point to the reaction of the triple bonds of the monomers to form polymer main chains composed of alternating single-double bonds. It is known that Rh catalysts provide poly(phenylacetylenes) with cis−transoidal structures, showing a signal due to the olefinic proton of the main chain around 6 ppm.23 In fact, with the 1H NMR spectra of the polymers, a resonance around 5.8 ppm was indeed observed, indicating the cis−transoidal structure of the backbone. From the integrated intensities of the olefinic proton and two carbazolyl protons (ca. 8.1 ppm), the cis contents were determined to be 74%, 64% and 93% for PPA-Cz-C3, PPA-Cz-C6, and PPA-Cz-C8, respectively. According to the TGA trace, the weight loss of the polymers in air occurred above 370 °C, whereas that of poly(phenylacetylene) usually starts at 250 °C. These results suggest that these polymers have superior thermal stability compared to typical poly(phenylacetylene).1−4

The normalized UV−vis spectra of the neutral PPA-Cz-C6 and PPA-Cz-C8 are shown in Figure 2. The absorbance spectra of the neutral PPA-Cz-C6 and PPA-Cz-C8 are consistent with the fact that carbazoyl moieties were incorporated as polymer side groups. The absorption band, tailing from 350 to 450 nm with an onset at 500 nm is due to the absorption (π→π∗) of the poly(phenylacetylene) main chain. This onset is similar for both, indicating little conformational change attributed to the addition of the carbazoyl groups. The other peaks below 350 nm were attributed to the absorption by the π→π∗, n→π∗, and benzenoid transitions of carbazole and phenyl groups. The conjugation length of these polymers is typical of most PPAs. The electrooptical band gap, ΔEg, of the un-crosslinked polymer can be calculated from the onset of absorbance and is equivalent to 2.46 eV, which is typical of poly(phenylacetylene).2,3

### Table 1. Summary of Polymerization of Cz-C3, Cz-C6, and Cz-C8 with [nbdRhCl2]a

<table>
<thead>
<tr>
<th>monomer</th>
<th>[M]/[Cat]</th>
<th>yield, %</th>
<th>Mw/Mn</th>
<th>solubilitya</th>
</tr>
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<tbody>
<tr>
<td>Cz-C3</td>
<td>200</td>
<td>60</td>
<td>148.0</td>
<td>insoluble</td>
</tr>
<tr>
<td></td>
<td>100</td>
<td>60</td>
<td>148.0</td>
<td>insoluble</td>
</tr>
<tr>
<td>Cz-C6</td>
<td>200</td>
<td>60</td>
<td>109.6</td>
<td>partly soluble</td>
</tr>
<tr>
<td>Cz-C8</td>
<td>200</td>
<td>60</td>
<td>93.9</td>
<td>soluble</td>
</tr>
</tbody>
</table>

*In toluene, at 30 °C. b By GPC (eluuent CHCl3, PSi standard). □ In THF, CHCl3, DMF etc.
the deposited films which appears at about 0.7 V on the first cycle. Similar to PVK, the first cycle always showed an oxidation onset at 0.9 V and the appearance of the oxidation−doping peak in the 0.6−0.7 V range with subsequent cycles. This means that the appearance of the 0.7 V peak is indeed due to the presence of the cross-linked carbazole units, regardless of the polymer backbone. All the CVs were reversible in the oxidation−reduction cycles at scan rates within the 20−100 mV/s range. Like other polyacetylenes, these polymers exhibited p-doping behavior. Thus, these observations point to a well-controlled electrochemical and cross-linking behavior which should allow work-function tuning when applied to actual LED devices using ITO substrates. Recently, extensive electrochemical studies on a class of polyacetylene ionomers have been investigated in which the importance of internal ion compensation to control various oxidation states has been emphasized.

The average values of the redox couple were calculated for PPA-Cz-C3, PPA-Cz-C6, and PPA-Cz-C8 and were found to be 0.461, 0.441, and 0.427 V, respectively. This was calculated from the anodic and cathodic peaks taken as the maximum of both peaks in a precursor polymer-free scan on the substrates after 10 cycles at 50 mV/s in 0.5 mg/mL polymer/CH2Cl2 in 0.1 M TBAH. From these data, it can be observed that the redox potential decreased as the length of the alkyl chain increased. Longer alkyl side chains tend to weaken any electron withdrawing effect from the PPA polymer backbone and decrease the redox potential. This may also be attributed to a change in the conformation of the polymer backbone influenced by steric effects of the side group. Nevertheless, such property may allow the possibility of band-gap tuning of the HOMO level for these polymers.
polymers with respect to device optimization. It should be noted that continuous potential cycling up to 10 cycles below 1.1 V for all these polymers, resulted in a linear area increase of the charge per cycle, corresponding to the linear deposition of polymer films. This observation of linear increase is consistent with what we have observed with other precursor polymers and the electropolymerization of PVK. Furthermore, these films are transparent and are insoluble to organic solvents indicating the electropolymerization of PVK.18-22 Furthermore, these films are transparent and are insoluble to organic solvents indicating a cross-linked film. For precursor polymers in general, we have observed that a “certain free volume” is necessary for effective cross-linking and film formation.22 In this case, a longer alkyl spacer or decoupling of the electropolymerizable carbazole monomer units results in a more homogeneous film formation.

**Electrochemical—Quartz Crystal Microbalance measurements.** Electrochemical—quartz crystal microbalance measurements were performed under much the same manner as the corresponding aforementioned CV. The difference from the previous electrochemistry setup is that the working electrode was changed from the Cz modified ITO slide to an unmodified Au coated quartz crystal with a 1.327 cm² working electrode area. The mass deposition per cycle was studied during the electrochemical polymerization (Figure 4). While it is true that viscoelastic effects do play a role in frequency changes when characterizing films by EQCM, it has been shown that EQCM is an effective tool for characterizing sufficiently thin films.32 For thin films (where the film thickness is much less than the wavelength of the piezoelectrically launched shear waves) a simplified form of the mass frequency relation can be used.

From the well-known Sauerbrey equation, the frequency shift \( \Delta F \) is a function of several parameters in the QCM setup.33 As can be seen in Figure 4b, the trace of the mass gain corresponds to that of the amount of charge transferred to electrode. This indicates that the deposition is due to the oxidation of the pendant carbazole units starting from about 0.9 V in the anodic scan of the CV cycle. The decrease of mass and amount of charge corresponds to the anion dedoping of the supporting electrolyte, which confirms the electroactivity of the deposited film. The \( \Delta F \) vs potential plot shows a regular increase in mass deposited with each cycle upon the cathodic scan. As the scan number increases, a dedoping of supporting electrolyte can also be seen. The plot of mass gain vs time has an induction period at the beginning of the experiment, but after the second cycle, the growth takes on a linear increase, giving a homogeneous film growth. This may be due to the absence of the carbazole SAM “adhesion” layer utilized with the ITO electrodes. However, this induction behavior is typical of most electropolymerization experiments.32 Likewise, E-QCM measurements of several cycles under precursor polymer-free conditions (see Supporting Information) showed no increase in deposited mass (or frequency shift) following each cycle. This is consistent with the reversible CV oxidation—reduction cycle as shown in Figure 3b. Thus, these polymers are well-behaved in terms of their film deposition properties and ion/mass transport.

**Spectroelectrochemistry.** To observe the presence of both the cross-linked carbazole and polyacetylene backbone species on the deposited film, spectroelectrochemistry measurements were made. UV—vis absorption spectra were measured in situ together with CV film deposition on PPA-Cz-C6 in 0.1 M TBAH/CH₂Cl₂ electrolyte (WE, modified ITO; CE, Pt wire; RE, Ag/AgCl) at a scan rate of 20 mV s⁻¹. Potentiodynamic measurements were then made on this deposited film to show doping/dedoping dependence with potential.

From the CV method, the peak at 550 nm was tracked and showed a linear increase with increasing CV cycle indicating deposition of the polymer (Figure 5a). After completion of the CV deposition, the two peaks at 400 and 850 nm were monitored during a series of potentiostatic measurements between 0 and 1.1 V (Figure 5b). Both peaks increased with higher potential but disappeared at lower and zero potentials. Therefore, the absorbance spectra in Figure 5b shows the doped (oxidized) and dedoped (reduced) states of the deposited films. Since the UV—vis spectra was monitored from a previously deposited film (no electropolymization taking place), the peaks at 400 and 850 nm can be assigned to the polaronic band originating form the contribution of poly(carbazole) species and their complex ion couple with the hexafluorophosphate ions. The assignments are polaron bonding level to the \( \pi^* \) conduction band for the 400 nm and bonding level to the antibonding state for the 850 nm.34 With active CV deposition, this is not easily observed. Thus, from Figure 5a, the 550 nm peak can be assigned to the \( \pi \rightarrow \pi^* \) transition of the polyacetylene, which increases linearly with film deposition during CV cycling.

From these measurements, the band-gap, \( \Delta E_g \), could be calculated and compared between the different electroactive species present on the film. The calculated \( \Delta E_g \) for the cross-linked polycarbazole species was estimated to be 3.13 eV based on extrapolation of the onset of absorbance in the UV—vis for the polycarbazole peak. The \( \Delta E_g \) for the cross-linked polyacetylene species was calculated from the onset of reduction and oxidation through CV methods and was found to be 2.45 eV. Some of the references for polyacetylene have given \( \Delta E_g \),
values of up to 1.88 eV. Thus, the results from potentiostatic spectroelectrochemistry methods confirmed the attributes of the PPA double—single bond polymer backbone in the presence of cross-linked polycarbazole units. However, their exact quantitative contribution to the cross-linked structure in the film is not easily distinguished by spectroelectrochemical methods alone.

AFM Imaging. Figure 6 shows the AFM image of the electropolymerized PPA-C6-Cz electropolymerized at 80 mV/s in 0.1 M TBAH/CH2Cl2 electrolyte (WE, modified ITO; CE, Pt wire; RE, Ag/AgCl wire). Electrochemical polymerization was carried out under CV conditions, for 10 cycles at a range of -50 to +1100 mV at a scan rate of 80 mV s-1 on the triethoxy(11-carbazoylundecyl)silane modified ITO plate. The films were optically smooth and uniform both by visual inspection and optical microscopy. The AFM image was made under ambient and dry conditions using MAC mode (noncontact). The morphology consisted of globular features on the order of 100–200 nm diameter which are regular in size and from relatively from large domains. The morphology is similar to that of our previously electrodeposited precursor polymers, e.g., the layer-by-layer electrodeposition of poly(methyhydrosiloxane)-functionalyzed polythiophenes under CV conditions in which smooth films were formed with each cycle.19,32

Conclusion

This work has shown that carbazole substituted poly(phenylacetylene)s (PPA) can be electrochemically cross-linked and deposited on triethoxy(11-carbazoylundecyl)silane-modified ITO resulting in the formation of conjugated polycarbazole networks without significantly decomposing the backbone. The two redox peaks observed at 0.6 V vs Ag/AgCl and above 1.0 V vs Ag/AgCl as confirmed by CV measurements were due to the carbazole and PPA unit charge complex couples, respectively. Regular and stepwise growth of the films was confirmed through in situ spectroelectrochemistry and electrochemical-QCM. These results emphasize that the following PPA-PCz-Cn polymer design is feasible for the electropolymerization of carbazole without affecting the PPA conjugated backbone significantly. The deposition method resulted in a combination of PPA and the PCz properties in one film. This was confirmed on the basis of the electrochemical species being identified (doping and dedoping) and followed using spectroelectrochemical measurements. CV trends with spacer chain length differences and a well-behaved electrochemical deposition and cross-linking behavior should allow HOMO-level and work-function tuning with these materials. Measurements are underway to determine their compatibility in polymer light emitting diode (PLED) applications as hole transport materials.

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Supporting Information Available: Figures showing CV scan of PPA control, five cycle CV for electrochemical crosslinking/deposition, and monomer free E-QCM study. This material is available free of charge via the Internet at http://pubs.acs.org.

References and Notes


