Comparative Study of Alternating Low-band-Gap Benzothiadiazole Co-oligomers

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Received: 21 June 2010 / Accepted: 10 January 2011 / Published online: 29 January 2011
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Abstract The benzothiadiazole – arylene alternating conjugated oligomers have been designed and synthesized via Suzuki coupling reaction. The structures and properties of the conjugated oligomers were characterized by $^1$H NMR, $^{13}$C NMR, UV–vis absorption spectroscopy, photoluminescence (PL) spectroscopy. The luminescent measurements demonstrate that polybenzothiadiazoles are good chromophores able to form thin films by Langmuir-Blodgett (LB) technique, making them suitable for further applications. Also the electrical properties of obtained films confirm the good potential of these novel aryl-based $\pi$-conjugated polymers for the development of various electrical and electrochemical solid-state devices.

Keywords Benzothiadiazole · Copolymers · Suzuki coupling · Electroconductivity · Luminescence · Atomic force microscopy

Introduction

During the past decade, organic light-emitting diodes (OLED) have been one of the most intensively explored topics in chemistry and applied physics owing to their wide applications in full-color displays with color filters, backlighting for liquid crystal displays, and general lighting sources [1–3]. While a large number of organic materials exhibit high fluorescence quantum efficiencies and bright in the visible region, there are some drawbacks such as low stability, high driving voltage associated with these materials. Therefore, they are not suitable for use in large area flat-panel displays. To overcome these problems and improve properties of electroluminescence (EL) devices, it is highly desirable to design and synthesize new dipolar and multifunctional materials, which could maintain well the carrier balance. $\Pi$-conjugated polymers/oligomers due to their combined desirable properties, such as fluorescence quantum yield, and good film-forming and hole-transporting properties have emerged as a very promising candidate for OLEDs. Color tuning in these polymers can be achieved by incorporating an electron-deficient monomer, benzothiadiazole (BT), into the polymer backbone [4, 5]. By incorporating BT into the alkylphenothiazine or alkyl phenoxazine polymer, the resulting copolymers possessed high electron affinity and preferential electron-transporting properties. An electron-rich and electron-deficient heterocycles composing the polymers possess a dual band of absorption in the visible spectrum consisting of short- and long-wavelength absorption bands. Moreover, benzothiadiazole derivatives normally are efficient fluorophores.

Many of desirable properties of conjugated organic polymers are dependent on the material’s band gap ($E_{\text{gap}}$), which is energetic separation between the material’s filled valence and empty conduction bands. Because of this, the significant effort has been applied to control the polymer $E_{\text{gap}}$ and to produce low $E_{\text{gap}}$ useful in technological fields novel materials.
Phenothiazine, as well as phenoxazine, is well known very strong electron donor and has high HOMO energy level because of sulfur atom. In addition, the HOMO energy level of phenothiazine copolymers is dominated by contribution of phenothiazine unit. Therefore, it is expected phenothiazine ring to be an excellent building block for lowering the ionization potential of conjugated polymers. Moreover, efficient electrogenerated chemiluminescence was observed in co-oligomers of methylphenothiazine and interesting redox properties were found in other oligophenothiazines [6]. Phenothiazines as well as phenoxazines are highly nonplanar. The possible consequence of the non-planarity of these rings for the photophysics, light-emitting properties, charge transport of π-conjugated polymers are intriguing and motivated. However, the device based on poly(alkylphenothiazine) are not enough to make efficient device since it has unbalanced charge transporting property. In order to improve electron affinity and transporting properties of material the benzothiadiazole was introduced to the oligomer backbone.

The research presented here deals with the synthesis, characterization and optical properties of novel π-conjugated oligomers based on 2,1,3-benzothiadiazole. Reported here palladium promoted coupling reaction of Suzuki is effective method of obtaining this class of highly luminescent π-extended benzothiadiazole derivatives. It is expected that these polymers could be used as air-stable conducting materials (through base doping for various electrical and electrochemical solid-states).

**Experimental**

**Materials and Instruments**

1H NMR and 13C NMR spectra were recorded using a Bruker 300 spectrometer. Chemical shifts are denoted in 1 unit (ppm) and were referenced to internal tetramethylsilane (0.0 ppm). The splitting patterns are designated as follows: s (singlet), d (doublet), t (triplet), quin (quintet), and m (multiplet). Preparative column chromatography was carried out on glass columns of different sizes packed with silica gel Merck 60 (40–63 μm). MS spectra were taken on a Bruker microOTOF-Q, FWHM-17500, 20 Hz. Chemical reagents were obtained from Aldrich and used as received. Tetraakis (triphenyl)phosphine palladium were obtained from Lancaster and used as received. Anhydrous THF was purified by vacuum distillation before use.

Atomic force microscopy (AFM) studies of Langmuir-Blodgett films were carried out using the AFM Dimension V Veeco.

Gel Permeation Chromatography (GPC) analysis of the polymer was performed on a Waters gel permeation chromatograph with Shodex gel columns and Waters model 150 C refractive index detectors using THF as eluent and polystyrene standards as reference.

Thermogravimetric Analysis (TGA) were conducted with a TA instrument Q50 TGA at a heating rate of 20 °C/min under nitrogen gas flow.

UV–vis absorption spectra were recorded on UV–VIS HP 8452A diode array spectrophotometer in either 1 or 4 cm quartz cuvettes depending on the sample concentration (10−4–10−6 mol dm−1). Emission spectra were measured at 295 K in 1 cm long-neck sealed quartz cuvettes using 90° geometry on Hitachi F–2500 fluorescence spectrophotometer. All solutions were purged with N₂ for 20 min prior measurement. All emission spectra were corrected using correction data obtained with rhodamine and methylene blue as quantum counters for wavelength out to 720 nm, beyond which the response was estimated from the manufacturer’s photomultiplier response data.

The luminance-current–voltage characteristics were recorded by combining the spectrometer through a Keithly model 2400 programmable voltage-current source. All the measurements were carried out at room temperature under ambient conditions.

The calculations were performed applying the DFT TD/B3LYP method and the standard 6–31 G* atomic basis set. The results reported here were obtained by utilizing the GAUSSIAN 03 code.

**Supplementary Materials—synthesis**

**4,7-Dibromobenzothiadiazole (2)** To a 250 ml two–necked round bottom flask were added 5.00 g (36.72 mmol) benzothiadiazole (1) and 75 ml of HBr (48%). A solution containing 17.47 g Br₂ (109.19 mmol) in 50 ml of HBr was added dropwise very slowly (slow addition is essential!). After the total addition of Br₂, the solution was heated at reflux for 6 h. Precipitation of a dark orange solid was noted. The mixture was cooled to room temperature, and a sufficient amount of a saturated solution of NaHSO₃ was added to completely consume any excess of Br₂. The mixture was filtered under vacuum and washed exhaustively with water. The solid was then washed once with cold Et₂O and dried under vacuum for ca. 20 h [7–9].

**Data of 4,7-Dibromobenzothiadiazole 2** Y=90% (9.71 g, 33.03 mmol), orange crystals, mp. 190 °C, 1H NMR (CDCl₃) δ: 7.72 (s, 2 H), 13C NMR (CDCl₃) δ: 152.9, 132.3, 113.9.

**3,7-Bis(4,4,5,5-Tetramethyl-2,1,3-Dioxaborolan-2-yl)-N,N-Diethylphenoxazine (4a)** To a solution of 1.00 g 3,7-dibromo-N-nonylphenoxazine (3a) (2.14 mmol) in dry THF (30 ml) at −78 °C was added, with a syringe,
0.44 ml (4.71 mmol) of n-butyllithium (2.5 M in hexane), the mixture was stirred at −78 °C, then warmed to 0 °C for 15 min, and cooled again to −78 °C for 15 min. 2-Isoproxy-4,4,5,5-tetramethyl-1,3-dioxaborolane (1.07 ml (5.25 mmol)) was added rapidly to the solution, warmed to room temperature, and stirred for 24 h. The mixture was poured into water and extracted with diethyl ether. The organic layers were washed with brine, and dried over magnesium sulfate. The solvent was removed, and the residue was purified by column chromatography with hexane:ethyl acetate (1:1) as eluent [10, 11].

Data of 3,7-Bis(4,4,5,5-Tetramethyl-2,1,3-Dioxaborolan-2-yl)-N-Nonylphenoxazine (5a) Y=45% (0.05 g, 0.16 mmol), dark brown crystals, 1H NMR (CDCl3) δ: 7.64 (d, J=8.3 Hz, 2 H, arom. H), 6.52 (d, J=3.2 Hz, 2 H, arom. H), 6.44 (dd, J=3.2 Hz, J=1.7 Hz, 2 H, arom. H), 3.85 (t, J=7.1 Hz, 2 H, CH2), 1.81−1.79 (m, 2 H, CH2), 1.35−1.27 (m, 24 H, CH3), 1.25−1.23 (m, 10 H, CH3), 0.83 (t, J=6.3 Hz, 3 H, CH3). 13C NMR (CDCl3) δ: 153.2, 143.9, 127.80 (2 C linked to boron, broad peak). 125.6, 124.5, 122.9, 122.7, 115.3, 111.8, 111.5, 103.8, 47.6, 31.8, 29.5, 28.0, 27.6, 26.8, 22.6, 14.0.

Poly[(N-Nonylphenoxazine-3,7-Diyl-alt-(2,1,3-Benzothiadiazole)] (5a) The following general Suzuki polymerization procedure was used for the preparation of all the copolymers. To a three-necked flask 0.20 g 3,7-bis(4,4,5,5-tetramethyl-2,1,3-dioxaborolan-2-yl)-N-nonylphenoxazine (4a) (0.35 mmol), 0.10 g 4,7-dibromobenzothiadiazole (2) (0.35 mmol), 0.01 g Pd(PPh3)4 (4.80 μmol), and 0.57 g potassium carbonate (0.04 mmol) was added. A mixture of water (0.21 ml) and toluene (15 ml) was added to the flask, and the reaction vessel was degassed. The mixture was heated at 90 °C for 24 h under nitrogen and then precipitated into methanol. The polymer was filtered and washed with methanol and acetone and then dried under vacuum overnight to afford the neutral oligomer [12, 13].

Results and Discussion

The synthesis of dibromobenzothiadiazole is outlined in Scheme 1. The reaction of 1 with molecular bromine (added
dropwise very slowly) in hydrobromic acid exclusively affords 4,7-dibromobenzothiadiazole (2) as a major product in 95% yield (Scheme 1) [7].

The synthesis of oligomers 5a–b was provided as shown in Scheme 2. Conversion of N-alkyl-3,7-dibromo-N-alkyl-phenoxazine (3a), -phenothiazine (3b) to the boronic ester (4a–b) was achieved (according to literature general procedure [8]) under reaction conditions in presence 2-isopropoxy-4,4,5,5-tetramethyl-1,3,2-dioxaborolane. Compounds 4a–b underwent Suzuki palladium-catalyzed polycondensation. Alternated DA oligomers 5a–b were obtained in 50% yield after purification by silica column chromatography. The molar ratio of phenoxazine moiety in the oligomers was controlled by adjusting the molar ratio between dibromobenzothiadiazole and monomers 4a–b while a 1/1 molar ratio between the dibromides and the bisborylated compounds was maintained.

The structures of the obtained co-oligomers were confirmed by 1HNMR, 13CNMR as well as MS spectroscopy. All structures were found to be highly soluble in most of organic solvents such as methylene chloride, chloroform, toluene, and o-dichlorobenzene.

GPC analysis was used to estimate the molecular weight of each sample. The molecular weight data are summarized in Table 1. It was determined that the oligomers were all of rather low molecular weight. Both oligomers gave number-average molecular weights of about 3,500 g/mol. This value corresponds to a degree of polymerization of seven to eight, meaning that the polymers synthesized here have an average of about 30 aromatic rings per chain. The molecular weights obtained were adequate for processing and film formation, and thus, further optimization of the polymerization was not attempted. The average molecular weight (Mn5a–b) of the copolymers was determined by gel permeation chromatography against polystyrene standards in THF (polydispersity in every case near 1.6).

Thin Films Formation

Thin films of these compounds were also easily fabricated from their solutions in chloroform by Langmuir-Blodgett (LB) technique.

Synthesized oligomers 5a–b dissolved in organic solvents (chloroform) were spread on the water, then organic layers were deposited by LB technique onto ITO (50 Ω) or a set of eight interdigital, buried Au electrodes (200 μm wide and spaced) photolithographically fixed on SiO2 thermally coated silicon substrates. Etching canals in silicone substrates were then filling by mixture of gold and trace amount of wolfram. This configuration provides flat, polished (measured by interferometric method) electrodes, ready to LB deposition. All initially non measured under water films were deposited at the velocity lower than the draining rate of film of carboxylic acids i.e. 1.3 mm/min. After deposition, films were stored in vacuum dessicator prior to use.

Langmuir monomolecular films of 5a–b were spread from CHCl3 monolayer solution on high purity water at room temperature. Langmuir-Blodgett deposition was carried

| Polymer | Mn [g/mol] | Mw [g/mol] | Tg [°C] | PDI |
|---------|-----------|-----------|---------|-----|
| 5a      | 3,500     | 5,200     | 130     | 1.57|
| 5b      | 3,600     | 5,400     | 80      | 1.62|

Table 1 Summary of GPC data (polystyrene standard) for benzothiadiazole co-oligomers
out with a KSV System 5000 LB through at a surface pressure of around 20–25 mN/m.

The sample processing was carried out at ca. 22 °C as well as conductivity measurements. The transference of LB film was Y–type in first deposition. The relationship between absorbance and number of layers and constant transfer ratio during the deposition indicate on constant architecture of LB film layers.

The LB film morphology was examined using atomic force microscopy. Experimental results on the topography (Fig. 1) show that the LB film is compact and highly ordered. On Fig. 1, for comparison, is also shown the topography of clean substrate (glass slide, Fig. 1b) before oligomer deposition. The high degree of homogeneity is observed for the film 5a. The surface roughness is rather smooth, it is due the reorganization of oligomer molecules during the LB deposition. The value of average roughness equals 9.8 nm. No clear crystalline fibers were observed in the AFM images.

The major result of investigations was pointed on the electrical conductivity of Langmuir-Blodgett (LB) films consisting several layers of synthesized compound (5a) as an effective conducting material.

Figure 2 shows typical current–voltage (I-V) curves of fabricated LB films at room temperature. The I-V curves are asymmetrical and non-linear. The forward currents follow approximately an exponential trend. This I-V behaviour is connected with diode-like type conductivity. The conductivity found for oligomer 5a LB film equals $\sigma = 1.2 \times 10^{-4} \text{ S/m}$.

From current–voltage characteristics obtained at higher temperatures (60 °C, 80 °C), it was found that the characteristics of the I-V maintain its asymmetrical shape and character. The forward currents increase strongly with increasing voltage.

Therefore, these results indicate that obtained films behave as p-type semiconductors. Moreover, good environmental stabilities observed for this material makes it a promising candidate for sensing elements.

Figure 3 displays the voltage-luminance results of compound 5a. The maximum luminance brightness is 6,857 cd m$^{-2}$ at 11 V. The brightness of 5a is improved largely, which denotes that compound has good hole-transporting ability and luminance property.

Figure 4 illustrated the luminance efficiency at different voltages. As the voltages increased gradually, the maximum luminescent efficiency increased to 3.10 cd A$^{-1}$. This phenomenon is a consequence of the formation of excitons required to emit light.

Optical Properties

While the origin of the dual-band absorption sometimes encountered in DA-type semiconducting polymers remains a source of debate, two mainstream rationales have been frequently proposed. A first assumption attributes the lower-energy optical transition to the presence of intermolecular charge-transfer excitons occurring on the presence of covalently bound DA segments along the backbone. A second assumption considers the presence of low-lying unoccupied energy levels, strictly localized on the electron-deficient heterocycles, yet forming a discrete “band” of easily accessed energy states within the bandgap of the conjugated system in its ground state. In both cases, the higher-energy transitions appear localized on the most-electron-rich building units incorporated along the polymer backbone with a clear dependence on their relative concentration to the electron-deficient heterocycles.

This distinct contribution from the electron-rich segments on the short-wavelength absorption band in DA backbones as well, the long-wavelength absorption band (red shifted) is illustrated in Figure 5.

The photophysical characteristic of the copolymers (5a–b) were investigated by ultraviolet–visible (UV–vis)
absorption and photoluminescence (PL) in diluted chloroform solution and in solid state. The optical data summarized are also listed in Table 2. A plot of extinction coefficient versus wavelength for the compound as measure in chloroform is depicted in Figure 5. As expected, 5a–b exhibit a dual band of absorption, in this case peaking at 280, 425 nm and 280, 400 nm for 5a–b, respectively. 2,1,3-Benzothiadiazole adsorbs at 306 nm [14]. It should be noted that the absorption bands characteristic for phenoxazine ring are localized at about 400 nm. However, the extent of conjugation between neighbouring benzothiadiazole units and phenoxazine ones will largely depend on the twist angle. The above data clearly suggest that the corresponding polymer has potential as low-band-gap material. The UV–vis spectrum of 5b is very similar to spectrum of 5a, but there is very characteristic broad absorption at 490–510 nm in solid state which was found also in another compounds based on benzothiadiazole and N-alkyphenothiazine (poly(3,7-divinylene-N-octyphenothiazine-alt-benzothiadiazole)) [15]. Similar donor-acceptor interaction between the electron-poor benzothiadiazole unit and the relatively electron-rich unit $\lambda_{\text{max}}$ at 460 nm, leading to an orange colour was found also in case of carbazole – benzothiadiazole copolymers [16].

In general, the absorption range of solid of 5a and 5b is slightly red-shifted than those of solution because of the formation of aggregates (Fig. 5). However, the absorption range of polyphenothiazine and 5b are not much different, this is due to that phenothiazine impede formation of aggregates in the solid state [17].

The photoluminescent properties of the synthesized oligomers (5a–b) were investigated by fluorescence spectroscopy in dilute chloroform solution (2 μM). The fluorescence quantum yield of the oligomers 5a–b were rather good (5a-$\varphi_F$=0.42, 5b-$\varphi_F$=0.48). The fluorescence quantum yield of benzothiadiazole is very low ($\varphi_F$=0.006). Molecules 5a and 5b have rather large Stokes’ shifts, 162 (61 728 cm$^{-1}$) and 145 nm (68 965 cm$^{-1}$) respectively.
The emission spectra of 5a and 5b solution were recorded at different excitation wavelengths in the range 310–390 nm, but the emission values were almost the same. The phenoxazine/phenothiazine – benzothiadiazole based oligomer emits in the blue region. However, when excited diluted solution of 5a at 389 nm the emission was centred at 476 nm (2.61 eV, Fig. 6, Table 2). In case of 5b the emission maximum was found at 480 nm (2.58 eV, Fig. 7). The observed Stokes shift according to phenoxazine emission suggests that there is a large twist between benzothiadiazole and neighbouring aromatic units.

The luminescence emission spectrum of 5a has well resolved vibronic structures in dilute chloroform solution. The emission spectrum of 5a has a vibronic progression with the 0–0 transition at 476 nm. Incorporation of benzothiadiazole moieties into polyphenoxazine results in a slight red shift in the 0–0 in the emission band from 458 nm (polyphenoxazine) [12] to 476 nm, both polyphenoxazine and polyphenoxazine with incorporated benzothiadiazole emits blue light in diluted solutions.

2,1,3-Benzothiadiazole – based oligomers and polymers have been widely studied in recent years as active materials in various optoelectronic devices because of the heterocyclic group and the observed low-band-gap in polymers containing it [18]. Copolymerization of benzothiadiazole with phenoxazine, phenothiazine, fluorene [4] and carbazole [16] or other suitable arylenes can be used as a means to tune the HOMO – LUMO levels in the resulting polymers. The HOMO and LUMO energy level of π-conjugated polymer are important for understanding charge injection processes in the luminescent devices.

By the theoretical study the luminescence of benzothiadiazole – phenoxazine oligomer (5a) was found as 506 nm (Table 2), in case of 5b the value is 482 nm. The above data, similar to experimental (476 nm—5a, 480 nm—5b), clearly suggests that the corresponding copolymers has potential as low-band-gap materials. The low optical band gaps of the oligomers should result from the alternating between the electron-rich unit of alkylphenothiazine/alkylphenoxazine and the strong electron-deficient unit of

| Table 2 Summary of physical properties of benzothiadiazole co-oligomers |
|----------------------------------|---------------------------------|-----------------|-----------------|
| Polymer | In solution | In solid state | Egap$^b$ [eV] | Egap$^c$ [eV] |
|         | abs $\lambda_{max}$ [nm] | PL $\lambda_{max}$ $^a$ [nm] | Stokes shift [cm$^{-1}$] | abs $\lambda_{max}$ [nm] |
| 5a | 280 | 476 | 61 728 | 0.42 | 355 | 2.45 | 2.61 |
| 5b | 284 | 480 | 68 965 | 0.48 | 360, 500 | 2.57 | 2.58 |

$^a$ polymer in chloroform solution was excited at 389 nm

$^b$ theoretical data

$^c$ figured out from the absorption edge from the PL spectrum
benzothiadiazole. The application of these oligomers are currently being investigated.

Conclusions

In summary, according to the donor-acceptor-alternating strategy, this paper described the synthesis of novel π-conjugated co-oligomers based on benzothiadiazole structure through well-known palladium catalyzed Suzuki polycondensation. The ability of connection of electron-donating and electron-withdrawing groups in benzothiadiazole co-oligomers, has been applied to developing methods for controlling the polymer E_g and producing technologically useful low-band-gap conjugated materials. The use of electron-withdrawing groups, however, provides reduced HOMO-LUMO energies and lower E_g values. The application of such alternating co-oligomers employing electron-withdrawing groups appears to be a promising new approach for the production of low E_g materials.

The obtained semiconducting oligomers - bifunctional materials as viable luminance and high hole-transporting one, exhibit excellent solubility in common organic solvents, thermal stability and luminescence in blue region and can be cast into uniform films. They possess good fluorescence quantum. The band gap values of synthesized compounds are in the adequate range (2.54–2.75 eV) for testing as OLEDs. Thin, ordered film fabricated of these compounds were found as semiconducting material for diode application.

Furthermore, it must be emphasized that benzothiadiazole-based copolymers have not only good processability due to the backbone, in which alkyl groups are incorporated, but also bright and uniform luminescence in the visible region. Since stable, bright blue emission is still a limiting factor in high performance LED devices. Polymers with these unique
backbones will be able to be introduced in many applications for low cost electronics, which needs good reproducibility of device performance and moderate field-effect mobility.

Acknowledgments Authors are acknowledged to Wrocław University of Technology and to the Polish Ministry of Science and Higher Education (Grant No. NN 204 244934) for financial support.

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