Synthesis and Characterization of \(-\)-Conjugated Multi-branched Molecules Bearing Carbazole and Phenothiazine Peripheral Groups

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Synthesis and Characterization of π-Conjugated Multi-branched Molecules Bearing Carbazole and Phenothiazine Peripheral Groups

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A novel class of conjugated multi-branched molecules bearing N-hexyl substituted carbazoles and phenothiazine as peripheral groups and [2,4,5-Tris-(diethoxyphosphorylmethyl)benzyl]phosphonic acid diethyl ester core has been synthesized through Horner-Emmons reaction in moderate to good yield. 8HCz-Den and 8HPTz-Den molecules have absorption maxima at 349 and 381 nm in solution states and maximum PL intensities were observed at 434 and 506 nm, respectively. They exhibited good self-film forming properties by solution casting and spin coating. In particular, by doping [2,6-bis(2-(5-(dibutylamino)phenyl)vinyl)-4H-pyran-4-ylidene]propanedinitrile, red emission could be achieved through complete Förster energy transfer process from the light harvesting host to the guest molecule.

Keywords: absorption; electroluminescence; energy transfer; multi-branched molecule; photoluminescence

INTRODUCTION

Multi-branched and conjugated polymers have attracted considerable attention because of their various applications in electronics and optoelectronics such as organic light-emitting diodes (OLEDs) [1–4]

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and photovoltaic cells [5]. However, it is difficult to control their molecular weight and purify these polymers. Dendrimers, which are perfect monodispersed macromolecules with a regular and highly branched three-dimensional architecture, represent a key stage in the ongoing evolution of polymer chemistry [6–9]. Along with flexible backbone dendrimers, conjugated dendrimers with rigid structures, such as phenylacetylene [10–12], phenylene vinylene [13–15], and polyphenylene [16] dendrimers have also been developed in recent years.

Carbazole and phenothiazine based multi-branched molecules can also be employed for carrier transporting materials and as luminescent host materials in EL devices [17–19]. We herein report the synthesis and photophysical characterization of a novel class of conjugated multi-branched molecules bearing carbazoles and phenothiazines as peripheral groups. In addition, we fabricate the composite film after doping \[2,6\text{-bis}(2\text{-}(5\text{-}(\text{dibutylamino})\text{phenyl})\text{vinyl})\text{-}4H\text{-pyran}\text{-}4\text{-ylidene})\text{propanedinitrile (DADB) into two multi-branched molecules with the concentration. We investigated the possibility to show red emission through complete or partial Förster resonance energy transfer process.}

**EXPERIMENTAL**

**Instrumental Analysis**

$^1$H NMR spectra were recorded on a Varian Mercury NMR 300 Hz spectrometer using deuterated chloroform (CDCl$_3$-d) purchased from Cambridge Isotope Laboratories, Inc. Elemental analysis was performed by using an EA1112 (Thermo Electron Corp.) elemental analyzer. Time-of-flight mass spectrometry (MALDI-TOF) was performed using a Voyager-DE STR MADI-TOF (matrix; DHB) mass spectrometer.

Thermal properties were studied under a nitrogen atmosphere on a Mettler DSC 821° instrument. Thermal gravimetric analysis (TGA) was conducted on a Mettler TGA50 thermal analysis system under a heating rate of 10°C/min.

**Absorption and Photoluminescence Spectroscopy**

Studying absorption and PL spectral behavior, thin films of multi-branched molecules were fabricated on quartz substrates as follows. The solution (2 wt%) of each molecule in monochlorobenzene was filtered through an acrodisc syringe filter (Millipore 0.2 μm) and
subsequently spin-cast on the quartz glass. The films were dried overnight at 80°C for 24 hours under vacuum. Absorption spectra of film samples and chloroform solution (conc. 1.5 x 10^{-6} mole/L) were obtained using a UV-vis spectrometer (HP 8453, PDA type) in the wavelength range of 190–1100 nm. PL spectra were recorded with an AMINCO-Bowman series-2 luminescence spectrometer.

Synthesis

We synthesized the intermediate compounds, 2, 3 by the literature method [20,21].

Synthesis of Compound 1

The same procedure was employed to prepare the compound 2: Instead of phenothiazine, we used carbazole. 1H NMR (400 MHz, CDCl3) δ 10.10 (s, 1H), 8.29 (s, 2H), 8.18 (d, J = 8.0 Hz, 2H), 7.93 (s, 3H), 7.74 (d, J = 8.0 Hz, 2H), 7.51 (t, J = 8.0 Hz, 2H), 7.50 (d, J = 16.0 Hz, 2H), 7.45 (d, J = 8.0 Hz, 2H), 7.44 (d, J = 8.0 Hz, 2H), 7.29 (t, J = 8.0 Hz, 2H), 7.24 (d, J = 16.0 Hz, 2H), 4.31 (t, 4H), 1.86–1.94 (m, 4H), 1.39–1.46 (m, 4H), 1.29–1.37 (m, 8H), 0.90 (t, 6H).

Anal. Calcd for C_{47}H_{48}N_{2}O C, 85.93; H, 7.37; N, 4.26. Found: C, 85.90; H, 7.53; N, 4.07.

Synthesis of 8HCz-Den

Compound 1 (820 mg, 1.25 mmol) and 3 (212 mg, 0.312 mmol) were dissolved in 30 ml of dry THF. The resulting solution was added drop-wise slowly to THF solution (20 mL) of potassium tert-butoxide (140 mg, 1.25 mmol) under N₂ for 12 h at room temperature. The reaction was quenched by adding a small amount of HCl solution and extracted with dichloromethane. The organic phase was dried over MgSO₄. Chromatography on silica gel (hexane/chloroform 1:4) gave 612 mg (yield: 73%) of green powder. T.g 96°C. 1H NMR (300 MHz, CDCl3) δ 8.12 (s, 8H), 8.02 (d, J = 8.4 Hz, 8H), 8.00 (s, 2H), 7.77 (d, J = 15.3 Hz, 4H), 7.75 (s, 8H), 7.71 (s, 4H), 7.57 (d, J = 8.4 Hz, 8H), 7.44 (t, J = 8.4 Hz, 8H), 7.41 (d, J = 15.3 Hz, 8H), 7.34 (d, J = 15.3 Hz, 4H), 7.30 (d, J = 8.4 Hz, 8H), 7.16 ~ 7.22 (m, 16H), 7.11 (d, J = 8.4 Hz, 8H), 4.00 (t, 16H), 1.66 ~ 1.75 (m, 16H), 1.18 ~ 1.29 (m, 48H), 0.84 (t, 24H); MALDI-TOF MS m/z calcd C_{198}H_{198}N_{8} (M + H)^+ 2688.5812, Found 2689.8746.

Synthesis of 8HPTz-Den

Compound 2 (1.12 mg, 1.55 mmol) and 3 (263 mg, 0.387 mmol) were dissolved in 30 ml of dry THF. The resulting solution was added
dropwise slowly to THF solution (20 mL) of potassium tert-butoxide (174 mg, 1.55 mmol) under N₂ for 12 h at room temperature. The reaction was quenched by adding a small amount of HCl solution and extracted with dichloromethane. The organic phase was dried over MgSO₄. Chromatography on silica gel (hexane/chloroform 1:4) gave 810 mg (yield: 71%) of orange powder. T_g 80°C; ¹H NMR (300 MHz, CDCl₃) δ 7.76 (s, 2H), 7.57 (d, J = 15.3 Hz, 4H), 7.50 (s, 8H), 7.46 (s, 4H), 7.10 ~ 7.19 (m, 40H), 6.81 ~ 6.97 (m, 20H), 6.78 (d, J = 8.4 Hz 8H), 6.61 (m, J = 8.4 Hz 8H), 3.71 (t, 16H), 1.69 ~ 1.78 (m, 16H), 1.34 ~ 1.42 (m, 16H), 1.25 ~ 1.33 (m, 32H), 0.87 (t, 24H); MALDI-TOF MS m/z calcd C₁₉₈H₁₉₉N₈ S₈ (M+H)⁺ 2944.3578, Found 2945.3600.

**RESULTS AND DISCUSSION**

We herein report the synthesis and photophysical characterization of two new π-conjugated multi-branched molecules bearing carbazoles and phenothiazines as peripheral groups.

In order to prepare 8HCz-Den and 8HPTz-Den, compounds 1 and 2 were prepared by the Heck coupling reaction of 3-vinyl-9H-hexylcarbazole/3-vinyl-10H-hexylphenothiazine and 3,5-dibromobenzaldehyde [20].

The convergent syntheses of the multi-branched molecules were conducted based on the typical Horner-Emmons condensation of 1/2 and [2,4,5-tris(diethoxy-phosphorylmethyl)benzyl]phosphonic acid diethyl ester core (Scheme 1). The reaction yield of this coupling reaction is fairly high to be around 71–73%.

The identity and purity of the synthetic materials were confirmed by ¹H NMR, MALDI-TOF mass spectrometry, and elemental analysis. These synthetic materials were found to have a good self-film-forming property and showed good solubility in various organic solvents such as chloroform, xylene, chlorobenzene, and tetrahydrofuran (THF).

The thermal properties of the multi-branched molecules were characterized by differential scanning calorimetry (DSC) and thermogravimetric analysis (TGA). 8HCz-Den and 8HPTz-Den exhibit no discernible crystalline-isotropic transitions observed in DSC thermograms (see Table 1). The glass transition temperatures (T_g) of the two molecules are 96 and 80°C for 8HCz-Den and 8HPTz-Den, respectively. This indicates that all molecules can exhibit an amorphous morphology in solid films. When comparing two glass transition temperatures, we found the fact that they are strongly dependent on the molecular planarity of carbazole units. It implies that a carbazole moiety has more planar than the phenothiazine moiety. Non-polar van
der Waals interaction between carbazole moieties is an important molecular specific interaction which can affect the physical properties significantly. TGA measurements at a heating rate of 10°C/min under nitrogen revealed good thermal stability. Two multi-branched molecules are thermally stable up to 392–445°C. (see Table 1) Higher thermal stability of 8HCz-Den can be attributed to the reason we described above.

**TABLE 1** Thermal and Photophysical Properties of 8HCz-Den and 8HPTz-Den

<table>
<thead>
<tr>
<th>Compound</th>
<th>T_g (°C)</th>
<th>T_d (°C)</th>
<th>ε (M⁻¹cm⁻¹)</th>
<th>λ_{max} (nm)</th>
<th>λ_{ex} (nm)</th>
<th>λ_{em} (nm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>8HCz-Den</td>
<td>96</td>
<td>445</td>
<td>356,000</td>
<td>349</td>
<td>350</td>
<td>345</td>
</tr>
<tr>
<td>8HPTz-Den</td>
<td>80</td>
<td>392</td>
<td>175,000</td>
<td>381</td>
<td>384</td>
<td>370</td>
</tr>
</tbody>
</table>
Figure 1 displays the absorption and PL spectra of the synthesized compounds such as $4$ and $5$ in dilute chloroform solutions and films. The absorption $\lambda_{\text{max}}$ of 8HCz-Den and 8HPTz-Den appeared 349 and 381 nm in solution, respectively. In the ground state, two branched large molecules do not exhibit high molecular interaction.

The PL spectra of two molecules in solutions and films are featureless and are almost a mirror-image of the low-energy absorption band. The PL spectra of the synthesized molecules in films exhibit a red-shift and become broader in comparison to the spectra in chloroform. These features are usually observed for organic and polymer conjugated materials mainly due to the different polarity of the environments. When we compare the PL spectra of two compounds in film state, 8HCz-Den showed larger spectral shift ($\Delta \lambda_{\text{emission}} = 51$ nm) than that of 8HPTz-Den ($\Delta \lambda_{\text{emission}} = 7$ nm). Therefore, in an excited state, the carbazole moiety induces higher molecular interaction to induce this effect.

In order to examine the functions of these compounds as host material in OLEDs, [2,6-bis(2-(5-(dibutylamino)phenyl)vinyl)-4H-pyran-4-ylidene]propanedinitrile (DADB) [22] was mixed into 8HCz-Den and 8HPTz-Den at six different concentrations in monochlorobenzene. Subsequently, thin films were fabricated on a quartz substrate for obtaining the PL spectra with excitation at 370 nm (8HCz-Den) and 365 nm (8HPTz-Den). Figure 2 shows the variation of the PL spectra with the DADB concentration (conc. $= 1$–$11$ wt.%); (dashed line) absorption spectrum of polystyrene doped with DADB. 8HCz-Den and 8HPTz-Den exhibited greenish blue and green emission predominantly; the maximum intensity of emission appeared at

**FIGURE 1** Absorption and PL spectra of the multi-branched molecules. (a) Solution in chloroform ($1.5 \times 10^{-6}$ M), (b) Film (thickness: 80 nm).
around 485 and 513 nm, respectively. With an increase in the concentration of DADB, the intensity at 485 and 513 nm decreases drastically. Further, red emission at around 620 nm increases significantly. DADB is capable of electronic excitation via Förster energy transfer from the host material. Photoluminescence (PL) studies of the doped matrix demonstrated that a significantly high energy transfer from 8HCz-Den and 8HPTz-Den to DADB is achieved. From Figure 2, it is clearly evident that the emission spectra of 4 and 5 and absorption spectrum of DADB dye overlap favorably, which enables an efficient energy transfer. In particular, we speculated that efficient energy transfer from a singlet and triplet state of the carbazole moiety to the DCM derivatives can enhance red emission in the DADB doped matrix. When doped with 5 wt% DADB into 8HCz-Den, we were able to obtain the highest PL intensity at 613 nm. The concentration of DADB could be optimized to 5 wt% in this experiment. Therefore, if we investigate the optimum concentration for complete or incomplete energy transfer from the host to the dopant, red or white emission behavior can be observed and it can be utilized for white light electroluminescence in our future study.

CONCLUSION

We have synthesized novel multi-branched molecules for host materials for red emitting EL devices. The HOMO and LUMO energies of the synthesized molecules are suitable for hole transporting materials in a multilayer EL device. The absorption and PL spectra indicate that the

![FIGURE 2 PL spectra of the multi-branched molecules doped with DADB in film states. (a) 8HCz-Den doped with DADB, (b) 8HPTz-Den doped with DADB (thickness: 30 nm).](image)
molecules bearing carbazoles or phenothiazines as peripheral groups can exhibit highly isolated photophysical properties. It can be considered that these carbazole-labeled multi-branched molecules can potentially be used for white organic light-emitting devices optimizing the concentration of DADB.

REFERENCES