Aggregation-enhanced emission (AEE)-active polyamides with methylsulfonyltriphenylamine units for electrofluorochromic applications

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A R T I C L E   I N F O

Article history:
Received 16 December 2016
Received in revised form 22 January 2017
Accepted 23 January 2017
Available online 24 February 2017

Keywords:
Triphenylamine
Sulfone
Polyamide
Electrochromism
Electrofluorochromism

A B S T R A C T

In this work, a series of AEE-active polyamides containing methylsulfonyltriphenylamine units were prepared from a newly synthesized diamine "4,4'-diamine-4 '-methylsulfonyltriphenylamine" and three dicarboxylic acids, which were highly soluble in various organic solvents and exhibited outstanding thermostability. The resulting polymer films showed one reversible redox couple along with apparent color changing from colorless to purple. The polymer solutions revealed relatively weak fluorescence with quantum yields in the range of 2.2-26.2%, which could be enhanced by induced aggregation in poor solvents. Furthermore, the bright fluorescence of the solid polymer film could be reversibly tuned by direct electrochemical redox of triphenylamine with a high contrast ratio ($I_{on}/I_{off}$) of 234. Overall, this comprehensive investigation of their interesting electrochromic and electrofluorochromic bifunctional properties not only supplies a deep understanding of the optical essence upon electrical stimuli but also paves the way for their future intelligent applications.

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1. Introduction

Recently, stimuli-responsive fluorescent materials have attracted intense interest due to their potential applications in sensors and displays [1–4]. In particular, a promising approach to achieve reversible fluorescence switching is based on the conversion of the redox states driven by electrical stimuli, which is known as electrofluorochromism [5–7]. In view of the fascinating advantages and wide application foreground, several efforts have been made to exploit various electrofluorochromic materials, mainly including dyads and electroactive fluorophores [8–13]. Specially, the latter ones behave more attractive since their fluorescence could be switched by electrochemical modulation directly and readily. To further meet the requirement of commercial application, some parameters such as rapid response time, high contrast ratio and long term stability are crucial for high performance electrofluorochromic materials. However, most of the existing materials could not achieve all these demands at the same time, especially lack of high contrast due to their poor fluorescence essence. To better satisfy the future optoelectronic applications, more versatile materials are urgent to be further exploited.

Triphenylamine (TPA) with special starburst structure has been widely applied in various optoelectronic materials such as organic light emitting diodes (OLEDs), dye-sensitized solar cells (DSSCs) and polymer memory devices, owing to its strong electron-donating and transporting capability [14–19]. Moreover, the TPA derivatives always yield obvious changes in color during the oxidation process, and thus have been developed as attractive anodic electrochromic materials [20–25]. In addition, para-protective TPA has also been demonstrated an ideal electro-switching modulator for its stable electroactivity by effectively preventing coupling reaction [26,27]; several reported fluorescent TPA derivatives exhibited interesting electrochemical fluorescence switching behaviors [28–30]. However, the aggregation-caused quenching (ACQ) essence of most of these materials resulted in
weak fluorescence in the solid states, which greatly decreased the fluorescence “on/off” sensitivity. To address this problem, the opposite of ACQ, aggregation-induced emission (AIE) effect which was first reported by Tang [31] paves a facile way for the design and synthesis of efficient solid-state materials. Remarkably, sulfone-containing materials are receiving increasing attention due to their effectively enhanced photoluminescence quantum yields [32,33]. Adachi and coworkers have reported amount of TPA-sulfone derivatives with interesting thermal activated delayed fluorescence (TADF) properties, ascribed to the small energy gap between the lowest singlet (S₁) and the lowest triplet (T₁) states resulting in strong reverse intersystem crossing (RISC) process [34–37]. Lee et al. reported several donor-acceptor molecules with high-efficiency blue light-emitting based on sulfone as acceptor [38,39]. In view of the propeller-like structure and electron-donating characteristic of TPA as well as the enhanced fluorescence and electron- withdrawing properties of sulfone unit, TPA-sulfone derivatives may readily act as electro and AIE-active modulators in electro-switching system, which will greatly promote the development of electrochromic and electrofluorochromic materials.

In this work, we herein report the synthesis of a new diamine monomer, “4,4’-diamic4-4’-methylsulfonyltriphenylamine”, and its derived electroactive polyamides. Their properties such as organic solubility, thermal stability, AEE fluorescence, redox, electrochromic and electrofluorochromic performances are also disclosed and investigated in detail.

2. Experimental section

2.1. Materials

4-Methylsulfonylaniline (TCI), 4-fluoronitrobenzene (Acros), cesium fluoride (CsF, Acros), 10% palladium on charcoal (Pd/C, TCI), hydrazine monohydrate (TCI), triphenyl phosphite (TPP, Acros) were used as received. The model compound with aliphatic structure was synthesized and characterized as described in the supporting information. Calcium chloride was dried under vacuum at 180 °C for 20 h prior to use. N-methyl-2-pyrrolidinone (NMP) and dimethyl sulfoxide (DMSO) were dried using 4 Å molecular sieves prior to use. Tetrabutylammonium perchlorate (TBAP, Acros) was recrystallized twice from ethanol under nitrogen atmosphere and then dried under vacuum at 40 °C before use. Commercially available dicarboxylic acids that include trans-1,4-Cyclohexanedicarboxylic acid (3a, TCI), 4,4’-dicarboxyldiphenylether (3b, TCI) and 2,2-bis(4-carboxy-phenyl)hexafluoropropane (3c, TCI) were dried under vacuum at 100 °C prior to use. Other commercially available chemicals and solvents were used without further purification.

2.2. Synthesis of 4,4’-dinitro-4’methylsulfonyltriphenylamine

In a 250 mL round-bottom flask equipped with a stirring bar, a mixture of 12.0 g (70.2 mmol) of 4-Methylsulfonylaniline, 20.8 g (147.4 mmol) of 4-fluoronitrobenzene and 22.4 g (147.4 mmol) of CsF in 130 mL of DMSO was heated under nitrogen atmosphere at 165 °C for 24 h. After cooling to room temperature, the reaction solution was poured into 500 mL of ethanol to form yellow crude which was subsequently washed thoroughly by hot water and ethanol. Then the crude was recrystallized from DMF/ethanol, yielding 24.1 g of the desired dinitro compound in 83.1% with a melting point of 271 °C. FT-IR (KBr): 1307 cm⁻¹, 1578 cm⁻¹ (–NO₂ stretch). 1H NMR (300 MHz, DMSO-d₆, δ ppm): 8.24 (d, J = 9.2 Hz, 4H), 7.97 (d, J = 8.9 Hz, 2H), 7.43 (d, J = 8.9 Hz, 2H), 7.31 (d, J = 9.2 Hz, 4H), 3.27 (s, 3H).

2.3. Synthesis of 4,4’-diamino-4’methylsulfonyltriphenylamine

In a 500 mL round-bottom flask, 15 g (8.8 mmol) of the dinitro compound 1 and 1.5 g of Pd/C were dispersed in 70 mL of ethanol. After heating the mixture to reflux, 8.4 g hydrazine monohydrate was added slowly into the react system. Then the mixture was stirred at the reflux temperature for 8 h. The Pd/C was removed by filtration and the resulting filtrate was cooled under a nitrogen flow to grow light-yellow crystals. Then the crystals were collected and dried under vacuum (11.0 g, 86.1% of yield, mp = 235 °C). 1H NMR (300 MHz, DMSO-d₆, δ ppm): 7.54 (d, J = 9.0 Hz, 2H, Ha), 6.93 (d, J = 8.6 Hz, 4H, Hb), 6.65–6.48 (m, 6H, Hz + Hc), 5.17 (s, 4H, –NH₂), 3.05 (s, 3H, –CH₃). 13C NMR (75 MHz, DMSO-d₆, δ ppm): 153.58, 146.97, 133.86, 128.32, 128.08, 127.29, 114.90, 113.70, 44.35.

2.4. Synthesis of polyamides

A series of polyamides were prepared via polycondensation (as shown in Scheme 2) [40]. The prepared polyamides were abbreviated to 4a, 4b and 4c, respectively. In a typical procedure, a mixture of 0.353 g (1 mmol) of the diamine monomer 2, 0.172 g (1 mmol) of 3a, 0.12 g of calcium chloride, 1 mL of triphenyl phosphite and 2.5 mL of NMP was heated with stirring at 120 °C for 3 h. After cooling to room temperature, the obtained viscous polymer solution was poured slowly into 250 mL of methanol, producing a stringy, fiber-like precipitate. The resulting polymer was washed thoroughly with hot water and methanol, and then precipitated by DMAc/ethanol twice for further purification. The 1H NMR data of these polyamides are listed as follows.

Polymide 4a: 1H NMR (300 MHz, DMSO-d₆, δ ppm): 9.98 (s, 2H), 8.02–7.45 (m, 6H), 7.14 (d, J = 8.4 Hz, 4H), 6.83 (d, J = 8.6 Hz, 2H), 3.11 (s, 3H), 2.44–2.24 (m, 2H), 2.02–1.81 (m, 4H), 1.60–1.38 (m, 4H).

Polymide 4b: 1H NMR (300 MHz, DMSO-d₆, δ ppm): 10.34 (s, 2H), 8.06 (d, J = 8.4 Hz, 4H), 7.84 (d, J = 8.7 Hz, 4H), 7.70 (d, J = 8.9 Hz, 2H), 7.31–7.12 (d, J = 8.4 Hz, 6H), 6.91 (d, J = 8.7 Hz, 2H), 3.12 (s, 3H).

Polymide 4c: 1H NMR (300 MHz, DMSO-d₆, δ ppm): 10.53 (s, 2H), 8.05 (d, J = 8.4 Hz, 4H), 7.83 (d, J = 8.6 Hz, 4H), 7.70 (d, J = 8.8 Hz, 2H), 7.54 (d, J = 7.9 Hz, 4H), 7.22 (d, J = 8.6 Hz, 4H), 6.92 (d, J = 8.8 Hz, 2H), 3.12 (s, 3H).

The polyamide films were prepared via solution casting. 0.3 g of sample was dissolved in 6 mL of DMAc to form the homogeneous solution, which was then poured into a 6 cm glass Petri dish and placed in the oven at 90 °C for 12 h to release the solvent slowly. After that, the damp-dry film was further dried at 160 °C for 12 h under vacuum. These as-prepared films were about 30–40 μm in thickness and were subsequently used for solubility tests and thermal analyses.

2.5. Measurements

Fourier transform infrared (FT-IR) spectra were recorded through a Bruker Vector 22 spectrometer at a resolution of 4 cm⁻¹ in the range of 400–4000 cm⁻¹. Nuclear magnetic resonance (NMR) spectra were determined on a BRUKER-300 spectrometer at 300 MHz for 1H NMR and 75 MHz for 13C NMR in deuterated DMSO. Inherent viscosities (ηinh) were measured through an Ubbelohde viscometer with a 0.5 g/dL of DMAc solution at 25 °C. Weight-average molecular weight (Mw) and number-average molecular weights (Mn) were obtained via gel permeation chromatographic (GPC) analysis on the basis of polystyrene calibration on a PL-2000 instrument with DMF as an eluent at a flow rate of 1.0 mL/min. Differential scanning calorimetric (DSC) analysis was
performed on a TA instrument DSC Q100 at a scanning rate of 10 °C/min in a nitrogen flow of 50 mL/min. Thermogravimetric analysis (TGA) was conducted with the TA 2050, with a heating rate of 10 °C/min under nitrogen atmosphere. Electrochemistry measurements were carried out on a CHI 660e electrochemical analyzer. Cyclic voltammetry (CV) was conducted by using a three-electrode cell. The working electrode was prepared by drop-coating of the polymeric solutions onto an ITO glass substrate (the area of polymer films was about 0.5 cm×2 cm). A platinum wire was used as an auxiliary electrode and a homemade Ag/AgCl, KCl (sat.) was used as a reference electrode. Ultraviolet-visible (UV–vis) spectra were measured using a Shimadzu UV 3101-PC spectrophotometer. Photoluminescence (PL) spectra and fluorescent quantum yield were determined with an Edinburgh FLS920 fluorescence spectrophotometer. Fluorescence quantum yield (ΦF) was determined using a calibrated integrating sphere.

3. Results and discussion

3.1. Monomers and polyamides synthesis

As shown in Scheme 1, the new diamine 2 with methyl-sulfonyltriphenylamine unit was synthesized via a two-step route; the diamine was successfully synthesized by Pd/C-catalyzed reduction of the dinitro 1 which was resulted from the nucleophilic displacement reaction of 4-methyl fluoronitrobenzene with 4-fluoronitrobenzene. The structures of the dinitro and diamine compounds were analyzed and confirmed by FT-IR and NMR techniques. The characteristic bands of nitro groups at around 1578 and 1307 cm⁻¹ disappeared after the reduction reaction. Meanwhile, typical N–H stretching absorptions pairs corresponding to the amino group appeared in the region of 3300–3500 cm⁻¹, indicating the formation of the new diamine (Fig. S1). Furthermore, the ¹H NMR spectrum of the diamine monomer 2 with the assignments for proton signals was illustrated in Fig. 1a, which further supports that the target diamine monomer with methyl-sulfonyltriphenylamine unit has been synthesized successfully.

A series of polyamides (4a–4c) were prepared from the diamine 2 with three different dicarboxylic acids through the phosphorylation technique. All the polymerization processes were homogeneously and afforded highly viscous polymer solutions after 3 h. The inherent viscosities of these polyamides ranged from 0.87 to 1.25 dL/g and the GPC measurement showed the number-average molecular weights (Mn) of 41200–69600 (Table S1). All these high-molecular-weight polyamides could be solution-casted into free-standing films. The typical FTIR spectrum of polyamide 4a is illustrated in Fig. S2, which exhibited the characteristic absorption bands of 1666 and 3321 cm⁻¹. The proposed structures of these polyamides were also confirmed by ¹H NMR spectra with amide resonance peaks appearing at around 9.9–10.6 ppm (Fig. 1b).

3.2. Solubility and thermal properties

The solubility tests of these polyamides were studied by dissolving 10 mg sample in 1 mL organic solvents (Table S1). The series of polyamides exhibited excellent solubility in polar solvents, which is greatly beneficial for the fabrication of large-scale thin film devices. The thermal properties of these polyamides were investigated by TGA and DSC measurements, with the results summarized in Table 1. During the decomposition processes in the TGA curves (Fig. S3), the 10% weight loss temperatures in nitrogen atmospheres were recorded in the range of 443–470 °C, and the carbonized residue was over 44% at 800 °C. The glass transition temperatures were observed at 306–334 °C depending on the stiffness of polymer structure. Thus, the excellent thermal properties of these polyamides will certainly contribute to extending their service time in optoelectronic devices.

3.3. Optical properties

The optical properties of these polyamides were examined with the detail spectroscopic data summarized in Table 2. The absorption maxima of these polyamides were located at 309–338 nm in NMP...
solutions and 312–344 nm in solid films, respectively, due to the π−π* transition of methylsulfonyltriphenylamine moieties. The dilute NMP solutions of these polyamides showed fluorescence maxima at 485–488 nm with quantum yields of 2.2–26.2% (Fig. 2a). Compared with the polyamide 4b and 4c, the high fluorescence of polyamide 4a could be ascribed to its reduced charge-transfer effect between TPA donor and benzamide acceptor [28]. Moreover, these polyamide films revealed enhanced fluorescence with quantum yields increasing to 2.7–32.1%. To further confirm the AEE effect, water, as the poor solvent for the polyamides, was added into the NMP solutions of polyamide 4a and different percentages of NMP-water mixtures were examined by fluorescence spectroscopy (Fig. 2b). In pure NMP solution, the polyamide 4a exhibited yellowish-green emission with a maximum peak at around 490 nm. With increasing water fractions from 0 to 50%, during which the polyamide aggregated to particles, the emission spectra were blue-shifted and the fluorescence intensity enhanced. The restriction of intramolecular rotations resulted in the higher fluorescence intensity [41]. By further increasing water fractions, however, the fluorescence intensity decreased gradually with a slight red-shift, which could be explained by the fact that the more increasing aggregation led to the enhanced intermolecular stack. These fluorescence spectra changes could be explained by the competition of ACQ and AEE effects. In addition, it is well known that the materials containing donor-acceptor structures are expected to show solvatochromic behaviors. To better demonstrate the solvatochromatic properties, the model compound of polyamide 4a which has better solubility was investigated in various solvents with different polarity. Fig. 2c and d exhibited the normalized UV−vis and PL spectra of the model compound in different solvents. All the absorption spectra were similar without obvious shift in the peak maximum, indicating the little effect of the solvent polarity on the ground-state electronic transition. However, the emission spectra revealed strong dependence of solvent polarity; in low-polar toluene, the model compound emitted strong blue light, while in the high-polar solvent DMSO, the emission color became yellow. A dramatic bathochromic shift of the emission peak from 445 to 501 nm was recorded with the changes of solvent polarity and this phenomenon could be ascribed to the fast inter-conversion process from the emissive excited state to the low emissive state [42].

### 3.4. Electrochemical and electrochromic properties

The redox behavior of these polyamides was investigated by CV measurement in a three-electrode cell with a polyamide-coated ITO electrode as the working electrode, a Pt wire as the counter electrode and an Ag/AgCl electrode as the reference electrode. As shown in Fig. 3, the polyamide 4a–4c all exhibited one reversible redox couple with onset potentials of the oxidation process ranging from 0.88 to 0.91 V, which indicates that the structures of dicarboxylic acid have a weak influence on the oxidation of triphenylamine core. On the basis of their onset oxidation potentials and ferrocene/ferrocenium reference, the highest occupied molecular orbital (HOMO) levels of these polyamides were estimated to be 5.26–5.29 eV. The bandgaps (Eg), as determined from the onsets of the polyamide films, lay between 3.11 and 3.20 eV. Thus, the corresponding lowest unoccupied molecular orbital (LUMO) levels were calculated to be 2.06–2.18 eV (LUMO = HOMO - Eg). All the details of electrochemical data as well as HOMO and LUMO levels are summarized in Table 2.

Considering the phenomenon that the polymer films coated on ITO substrates were observed to switch their appearance colors between colorless and purple reversibly, spectroelectrochemical measurements were performed to compare the optical properties of polyamide 4a at different applied potentials (Fig. 4A). Upon the electro-oxidation steadily arising from 0.0 V to 1.2 V, the intensity of the characteristic absorption peak at 312 nm for the neutral form of the polyamide decreased gradually, while three new peaks at 388, 545 and 854 nm grew up. The spectra changes were ascribed to the formation of a monocation radical, which caused the TPA structure to flatten off and absorb long wave lights.

Electrochromic switching studies were also performed through square-wave potential stepping experiments with the transmittance change at the given wavelength monitored as a function of time. The typical switching behavior of polyamide 4a with a cycle time of 10 s was depicted in Fig. 4B. After 50 cycles, no obvious spectral decay was observed, indicating the highly stability as electrochromic device. The switching time plays an imperative role in electrochromic applications, and here, it was calculated as 90% of the time taken for the full switching state. The polyamide 4a revealed a satisfactory switching time of 4.1/1.0 s for coloring/bleaching process. In addition, coloration efficiency (CE) is also an important parameter of electrochromic materials, and it was defined as: CE = ΔA/ΔQd (ΔA: optical absorbance change; ΔQd: injected/ejected charge per unit sample area during a redox step). The CE value of polyamide 4a ranged from 121 cm2C−1 for the first
Fig. 2. (a) UV–vis absorption and PL spectra of the polyamides 4a–4c in NMP solutions ($1 \times 10^{-3}$ M). (b) PL spectra of polyamide 4a in NMP-water with different water fractions (the inset photographs were taken under illumination of a 365 nm UV light). Absorption (c) and PL behavior (d) of the model compound in various solvents with different polarity (solution concentration was $1 \times 10^{-3}$ M and the inset photographs were taken under illumination of a 365 nm UV light.

Fig. 3. Cyclic voltammetric diagrams of the polymers (a) 4a, (b) 4b, (c) 4c, (d) ferrocene in CH$_3$CN containing 0.1 M TBAP at a scan rate of 100 mV s$^{-1}$. 
cycle to 113 cm\(^2\) C\(^{-1}\) for the fiftieth cycle.

### 3.5. Electrofluorochromic performance

The polyamide 4a with methylsulfonfyltriphenylamine exhibited shining fluorescence, especially in the solid state, but TPA\(^+\) is known as an effective fluorescence quencher [13]. Therefore, the fluorescence changes under a series of positive potentials were recorded to demonstrate its electrofluorochromic switching properties. Upon excitation at 365 nm, the neutral polyamide film exhibited vivid cyan fluorescence, while the strong fluorescence vanished to dark and the intensity of the fluorescence peak at 474 nm decreased under positive potentials. When applying reverse potentials, during which the TPA\(^+\) was reduced to the neutral state, the fluorescence recovered, and notably, the electrofluorochromism offered a high contrast ratio (\(I_{\text{off}}/I_{\text{on}}\)) of 234. It is worth noting that no shift of spectral band was observed during the changes of the fluorescence intensity, which indicates that the fluorescence quenching originated from the electrochemical oxidizing of the polyamide 4a to its cation radical form without other side reactions. Furthermore, the dynamic response behavior was also examined by repeating oxidation and reduction steps between 0.0 V and 1.2 V (Fig. 5b), which revealed excellent reversibility during the fluorescence “off/on” process. The contrast ratio exhibited dependent behavior on the switching time, changing from 234 to 9 by reducing the switching time from 360 to 10 s. In addition, the long-term durability of the electrofluorochromic polyamide was also studied. No obvious decay in either “on” or “off” state was observed in 50 continuous switches when the switching time was set as 10 s. Considering the easy processability and excellent electrochromic/electrofluorochromatic properties, the polyamide 4a holds great potential for practical intelligent applications in the future.

### 4. Conclusion

In summary, we have designed and synthesized a series of AEE-active polyamides with methylsulfonfyltriphenylamine units, which exhibited excellent solubility and thermal stability. These polyamides exhibited reversible electroactive and favorable electrochromic properties with color changing from colorless to purple. Notably, the polyamide 4a displayed an AEE feature with a high solid-state fluorescent quantum yield up to 32.1%, and the strong fluorescence could be effectively modulated by electrochemical switching with a high contrast of 234. Thus, the durable and high-contrast electrochromic/electrofluorochromatic polyamide 4a will provide useful information for developing bifunctional displays or

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**Fig. 4.** (A) Spectral changes of 4a film coated on an ITO-coated glass substrate (in 0.1 M TBAP/CH\(_3\)CN) at different applied potentials (0.0–1.2 V). The insets show the color changes. (B) Optical switching between 0.0 and 1.2 V of 4a film (a cycle time of 10 s). (a) Absorbance changes and (b) current consumptions monitored at 545 nm. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

**Fig. 5.** (a) Fluorescence spectra changes of 4a film upon different applied potentials from 0.0 to 1.2 V. (b) Fluorescence switching response under applied step potential between 0.0 and 1.2 V with a step duration time for 360, 60, 40, 20, 10 s at each potential.
other electro-switching devices.

Appendix A. Supplementary data

Supplementary data related to this article can be found at http://dx.doi.org/10.1016/j.dyepig.2017.01.078.

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