

Synthesis and characterization of polyfluorene-based photoelectric materials: the effect of coil segment on the spectral stability†

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The origin of the low-energy emission of fluorene-based rod-coil block copolymers still remains controversial. In this work, a series of polyfluorene-based rod-coil block copolymers having different coil segments, *i.e.*, poly[2,7-(9,9-dihexylfluorene)]-*block*-poly(2,2,3,3,4,4,4-heptafluorobutyl methacrylate, (PF-*b*-PHFBMA), PF-*b*-poly(butylmethacrylate) (PF-*b*-PBMA), PF-*b*-poly(2-hydroxyethyl methacrylate) (PF-*b*-PHEMA) and PF-*b*-poly(acrylic acid) (PF-*b*-PAA), were synthesized using the ATRP technique. The optical and surface properties and thermal behaviors of these copolymers were systematically investigated. In particular, different thermal treatment conditions, including annealing temperature, annealing time and annealing atmosphere were introduced to study the effect of coil segment on the copolymer spectral stability. The incorporation of PBMA, PHEMA and PAA segments to PF could indeed improve the copolymer spectral stability, while the PHFBMA block brought undesirable low-energy emission. In addition, water contact angle (WCA) measurements of the copolymer films before and after annealing further demonstrated that the low-energy emission of PF-based rod-coil block copolymers was attributed to the molecular aggregation rather than the formation of fluorenone defects.

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Introduction

Fluorene-based polymers (PFs) have been intensively studied as blue-light-emitting materials in polymer light-emitting diodes (PLEDs) due to high photoluminescence efficiency, good photostability and the emission of polarized blue light.^{1–5} The poor spectral stability under thermal treatment or passage of current caused the appearance of a low-energy emission band at 520–530 nm, significantly limiting their application in blue PLEDs.^{4,5} Three models, *i.e.*, the creation of keto defects in the backbone of PFs,^{6–8} the formation of aggregates or intermolecular excimer emission^{9–11} as well as the combination of the two mechanisms,^{12,13} have been suggested to clarify this phenomenon. The exact origin of this low-energy emission band remains unclear.¹⁴ Nevertheless various efforts have been made to address this disadvantage, including using either dendrimers or bulky substituents in the 9-position of the fluorene moiety,^{15,16} copolymerization techniques,^{17,18} purification of the fluorene monomers before polymerization,¹⁹ or blends of PFs with some nonconjugated polymers such as polystyrene (PS) or

poly(vinylidenequinoline),²⁰ while the undesirable emission band was not eliminated.²¹

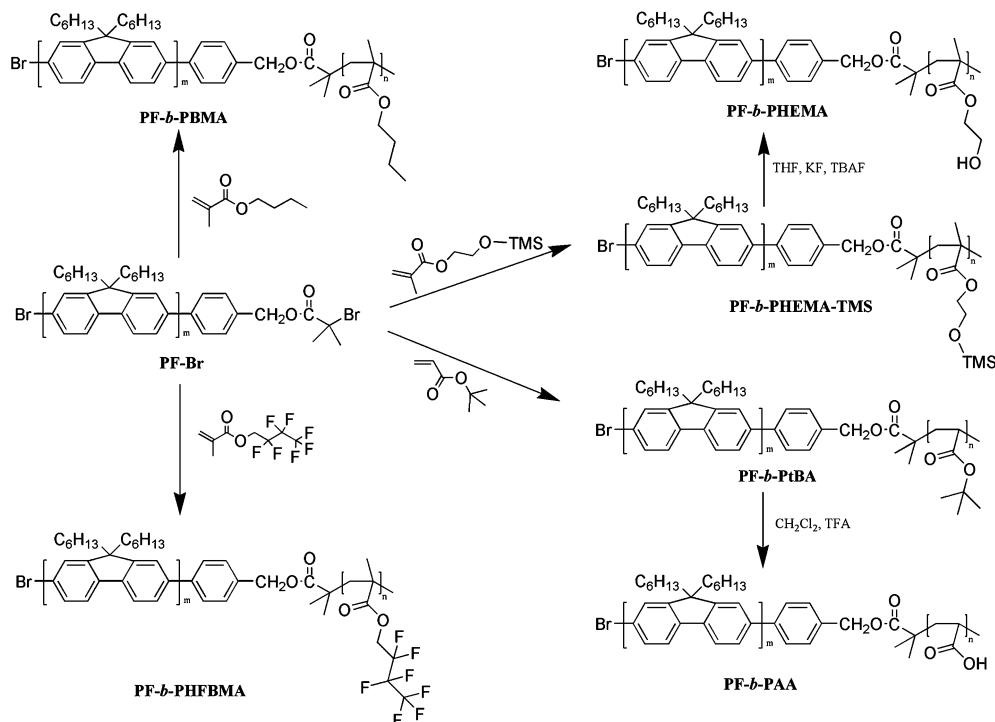
In recent studies, coil-like blocks nonconjugated and hole/electron transporting molecules were incorporated into polyfluorene backbones to suppress the aforementioned low-energy emission.^{21–24} So far, many coil segments have been used to form PF-based rod-coil block copolymers, such as poly(methyl methacrylate) (PMMA), polystyrene (PS), poly(2-hydroxyethyl methacrylate) (PHEMA), poly(2-(dimethylamino)ethyl methacrylate) (PDMAEMA) and poly(ethylene oxide) (PEO).^{25–30} The aggregation and microphase separation behavior of these block copolymers caused to the incompatibility of rod and coil segments can lead to the formation of unusual nanoscale morphologies and photophysical properties in both solution and solid state.^{14,31} These functions, which can be adjusted flexibly through introducing different coil segments, have attracted great attention for optoelectronic or sensory applications.^{32,33} However, the influence of coil segments related to the microphase separation behavior on the photophysical properties of PF-based rod-coil block copolymers has not been fully explored.

Recently, we have synthesized a series of PF-based rod-coil block copolymers with PHFBMA as the coil segment by ATRP and reported that the micelle structure formed by different selective solvents has a dramatic influence on their optical properties.³⁴ The previous study was achieved in solution, however the influence of separation behavior on the optical properties of these block copolymers in solid state has not ever

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Scheme 1 Synthesis route of polyfluorene-based rod-coil block copolymers PF-*b*-PHFBMA, PF-*b*-PBMA, PF-*b*-PHEMA, PF-*b*-PAA.

been studied. Based on this idea, four coil segments having different polarity and solubility are selected to form PF-based rod-coil block copolymers PF-*b*-PHFBMA, PF-*b*-PBMA, PF-*b*-PHEMA and PF-*b*-PAA in this study. These copolymers with a similar polymerization degree are well synthesized through ATRP technique. Scheme 1 shows the synthetic routes and corresponding synthetic conditions. Particular attention is focused on the effect of coil segment on the copolymer spectral stability under different thermal treatment conditions. In addition, water contact angle (WCA) measurements about the surface properties and thermal stability of resulting polymers were also carried out.

Experimental section

Materials

α -{4-[2-(2-Bromo-2-methylpropoxy)methyl]phenyl}- ω -bromopoly[2,7-(9,9-dihexyl-fluorene)] (PF-Br) was synthesized using a similar procedure, which has been reported by Chen *et al.*²⁹ and us.³⁴ 1,1,4,7,7-Pentamethyldiethylenetriamine (PMDETA, 98%) was purchased from Aldrich. 2-Hydroxyethyl methacrylate (HEMA, 95%) and tetrabutylammonium fluoride (TBAF, 1 M in tetrahydrofuran) were obtained from TCI (Shanghai) Development. HEMA was purified by washing an aqueous solution of monomer with hexane to remove ethylene glycol dimethacrylate, salting the monomer out of the aqueous phase by addition of NaCl, drying over MgSO₄, and distilling under reduced pressure. Potassium fluoride (KF, 99%), trifluoroacetic acid (TFA, 99%), copper(I) chloride (CuCl, 98%), butyl methacrylate (BMA, AR) were obtained from Sinopharm Chemical Reagent

(SCRC). 2,2,3,3,4,4,4-Heptafluorobutyl methacrylate (HFBMA, 98%) was obtained from A Better Choice for Research Chemicals GmbH & Co. KG. (ABCR). *tert*-Butyl acrylate (*t*BA, 99%) was purchased from Aladdin. BMA, HFBMA and *t*BA were washed with 5 wt% aqueous NaOH solution to remove inhibitor before polymerization. All other reagents and solvents were obtained from SCRC and used without further purification.

Synthesis of PF-*b*-PHFBMA

100 mg (0.017 mmol) PF-Br first dissolved in 2.5 mL of cyclohexanone, 14.2 μ L PMDETA and 3.4 mg CuCl were added to the solution. The mixture was degassed and then filled with nitrogen three times at 0 °C, followed by addition of 339 μ L (1.7 mmol) HFBMA. Finally, the flask was immersed into an oil bath preheated at 80 °C for 24 h. The polymerization was terminated by opening the flask and exposing the mixture to air. After cooling to room temperature, the copolymer solution was diluted with THF and passed through an alumina column to remove the catalyst. Finally, the resulting solution was concentrated and poured into methanol to precipitate the product. The product was further purified twice by redissolving/precipitating with THF/methanol and dried in vacuum for 24 h to obtain PF-*b*-PHFBMA.

Synthesis of PF-*b*-PBMA

The synthesis process was the same as the synthesis process of PF-*b*-PHFBMA except replacing HFBMA by BMA (270 μ L, 1.7 mmol).

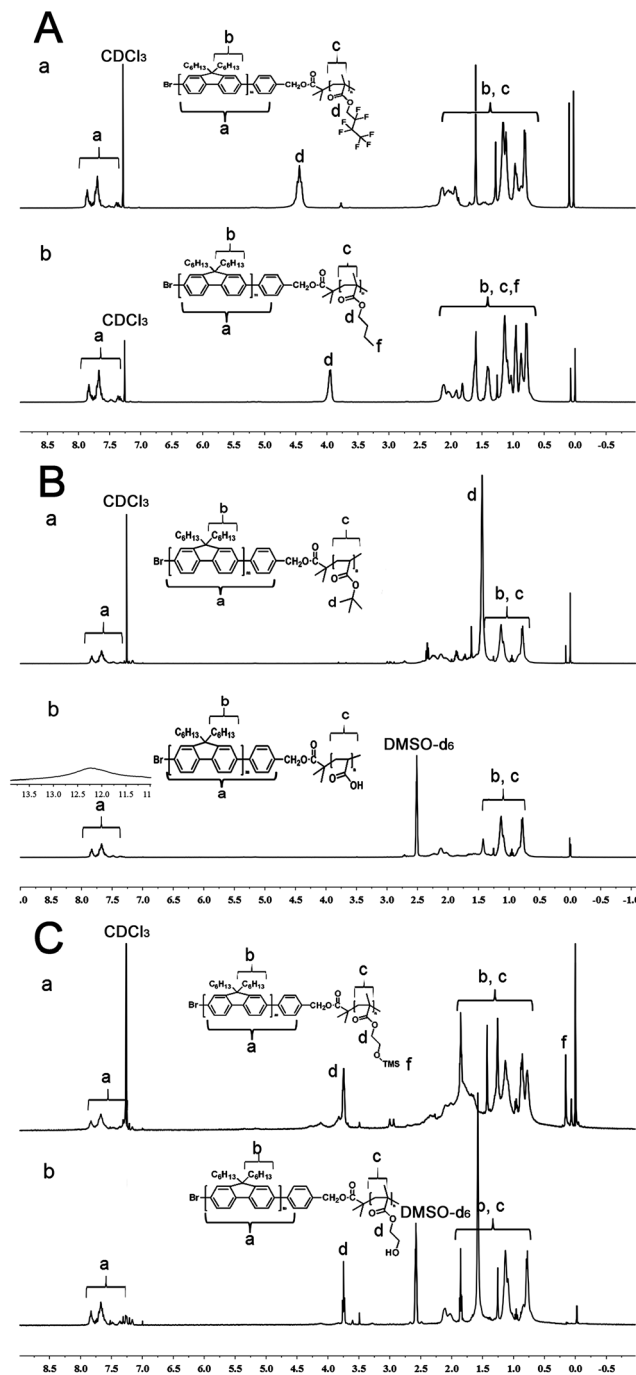


Fig. 1 ^1H NMR spectra of polyfluorene-based rod-coil block copolymers (A-a) PF-*b*-PHFbMA and (A-b) PF-*b*-PBMA in CDCl_3 ; (B-a) PF-*b*-PtBA in CDCl_3 and (B-b) PF-*b*-PAA in $\text{DMSO}-d_6$; (C-a) PF-*b*-P(HEMA-TMS) in CDCl_3 and (C-b) PF-*b*-PHEMA in $\text{DMSO}-d_6$.

Synthesis of PF-*b*-PAA

PF-*b*-PtBA was first prepared using PF-Br as macroinitiator and tBA as monomer. The synthesis process was the same as the synthesis process of PF-*b*-PHFbMA except replacing HFBMA by tBA (370 μL , 1.7 mmol). Then the obtained copolymer PF-*b*-PtBA (60 mg) was dissolved in 15 mL dry dichloromethane. 6.5 μL TFA was added and the solution was stirred at room

temperature for 24 h. A precipitate was obtained after pouring the concentrated solution into *n*-hexane. The result copolymer PF-*b*-PAA was dried under vacuum for 24 h.

Synthesis of PF-*b*-PHEMA

2-(Trimethylsilyl)ethyl methacrylate (HEMA-TMS) instead of HEMA is usually used to synthesize block copolymers due to its high solubility in organic media.³⁵ PF-*b*-P(HEMA-TMS) was first prepared using PF-Br as macroinitiator and HEMA-TMS as monomer. The synthesis process was the same as the synthesis process of PF-*b*-PHFbMA except replacing HFBMA by HEMA-TMS (354 μL , 1.7 mmol). Then the obtained PF-*b*-P(HEMA-TMS) (70 mg) was dissolved in 15 mL dry THF, KF (5 mg) and TBAF (5 μL) were added to the polymer solution and stirred for 24 h at room temperature. The polymer solution was concentrated under reduced pressure and poured into water. The target copolymer PF-*b*-PHEMA was separated by filtration, and dried under vacuum for 24 h.

Measurements

^1H NMR spectra were measured on a BrukerAvance 400 instrument with deuterated chloroform and dimethyl sulfoxide- d_6 as the solvent.

Fourier-transform infrared (FTIR) spectra were obtained on an Avatar 360 FTIR spectrophotometer by dispersing samples in KBr disks.

The molecular weight (M_n) and molecular weight distribution (M_w/M_n , PDI) of the polymers were determined at 40 $^\circ\text{C}$ by GPC equipped with a waters 1515 isocratic HPLC pump, three styragel columns (Waters HT4, HT5E, and HT6) and a waters 2414 refractive index detector (set at 30 $^\circ\text{C}$), using THF as the eluent at the flow rate of 1.0 mL min^{-1} . A series of polymethyl methacrylate narrow standards were used to generate a conventional calibration curve.

Thermogravimetric analysis (TGA) was conducted on a SDT Q600 under a heating rate of 10 $^\circ\text{C min}^{-1}$ from room temperature to 800 $^\circ\text{C}$ in nitrogen gas atmosphere.

The fluorescence measurement was carried out on a Hitachi 7000 spectrofluorometer with a xenon lamp as a light source.

Water contact angle (WCA) measurements were carried out on a Contact Angle Measuring Instrument (KRUSS, DSA30). The wetting liquid used was water. For each angle reported, at least five sample readings from different surface locations were averaged.

The films used for fluorescence and WCA measurements were prepared as follows. Glass slides were first cleaned successive using acetone, ethyl alcohol and deionized water. The polymer solution (10 mg mL^{-1} in THF) was then spin-cast onto clean glass slide at 2000 rpm for 30 s, and dried naturally for 24 h.

Results and discussion

Characterization of polymers structure

According to the relevant literature²⁹ and our previous work,³⁴ PF-OH ($M_n = 5947 \text{ g mol}^{-1}$, $M_w/M_n = 1.56$) was first prepared

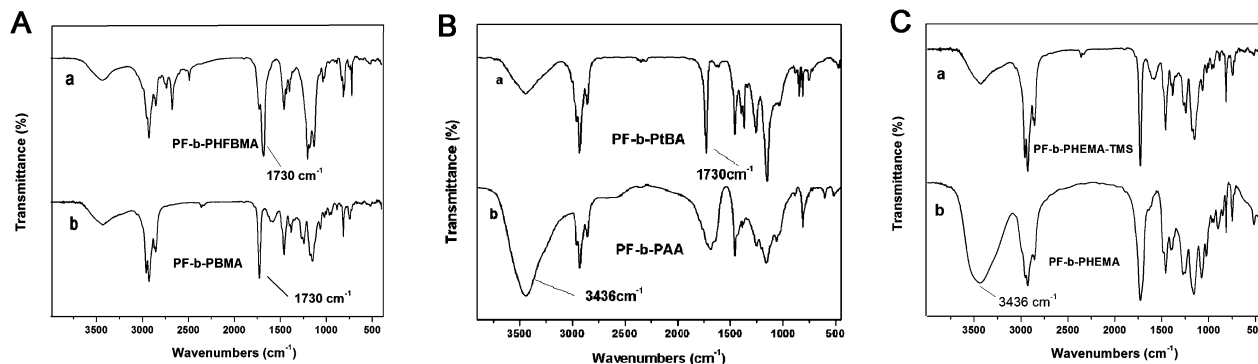


Fig. 2 FTIR spectra of polyfluorene-based rod-coil block copolymers (A-a) PF-*b*-PHFBMA and (A-b) PF-*b*-PBMA; (B-a) PF-*b*-PtBA and (B-b) PF-*b*-PAA; (C-a) PF-*b*-P(HEMA-TMS) and (C-b) PF-*b*-PHEMA.

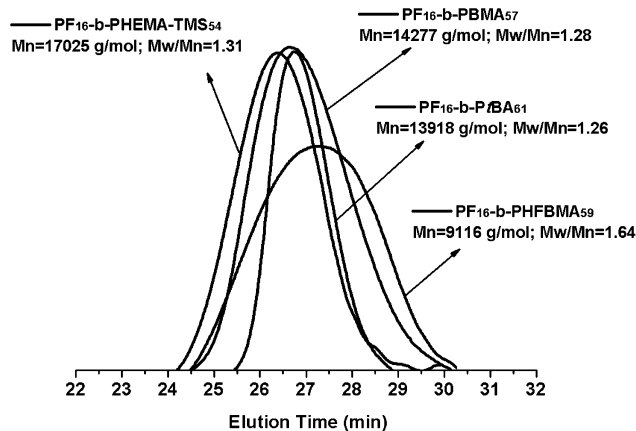


Fig. 3 GPC traces of polyfluorene-based rod-coil block copolymers PF-*b*-PHFBMA, PF-*b*-PBMA, PF-*b*-P(HEMA-TMS), PF-*b*-PtBA.

through a modified Suzuki coupling reaction with 2-bromo-9,9-di-*n*-hexylfluoreneboronic acid and a 4-bromobenzyl alcohol end-capper. After esterification of PF-OH with 2-bromoisobutryl bromide, bromo-ended polyfluorene PF-Br was obtained. $M_n = 6067 \text{ g mol}^{-1}$, $M_w/M_n = 1.48$. The successful synthesis of PF-OH and PF-Br can be confirmed by ^1H NMR and FTIR (Fig. S1 and S2 of the ESI†).

All polyfluorene-based rod-coil block copolymers were synthesized *via* ATRP method with the bromo-ended polyfluorene as macroinitiator (shown in Scheme 1). And their chemical structures were confirmed by ^1H NMR and FTIR. Fig. 1A-a and A-b show the ^1H NMR spectra of PF-*b*-PHFBMA and PF-*b*-PBMA respectively. The appearance of peak at 4.42 ppm in Fig. 1A-a and peak at 3.95 ppm in Fig. 1A-b, corresponding to the protons of the $-\text{O}-\text{CH}_2-\text{CF}_2-$ group of HFBMA and the $-\text{O}-\text{CH}_2-$ group of BMA, respectively, indicate the successful synthesis of PF-*b*-PHFBMA and PF-*b*-PBMA. The FTIR spectra of PF-*b*-PHFBMA (Fig. 2A-a) and PF-*b*-PBMA (Fig. 2A-b)

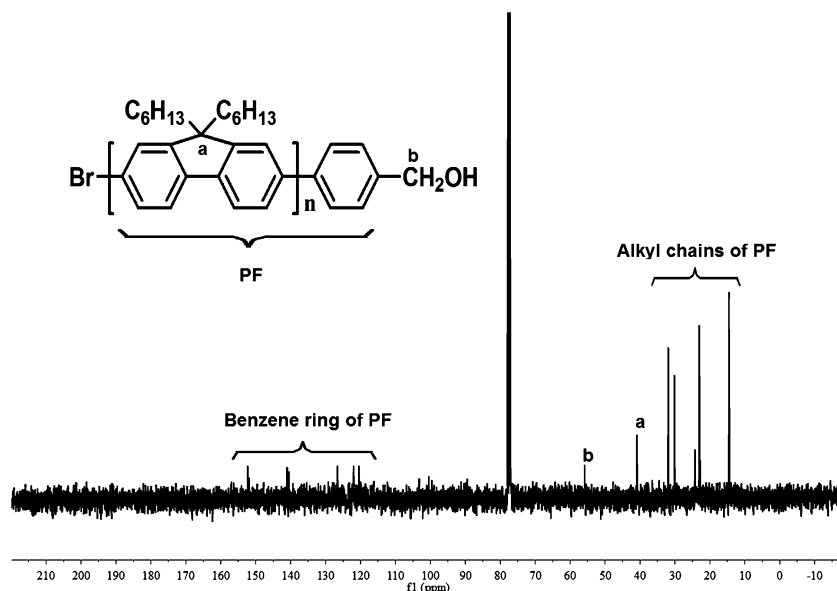


Fig. 4 ^{13}C NMR spectrum of PF-OH in CDCl_3 .

both exhibit more stronger absorption peak at 1730 cm^{-1} which is assigned to the stretching vibrations of $\text{C}=\text{O}$ group of PHFBMA and PBMA. These results also confirm the successful synthesis of PF-*b*-PHFBMA and PF-*b*-PBMA (the original ^1H NMR and FTIR spectra of PF-Br for comparison are respectively shown in Fig. S1-b and S2-b of the ESI†).

Fig. 1B-a and 2B-a respectively show the ^1H NMR and FTIR spectra of PF-*b*-PtBA. The appearance of peak at 1.44 ppm corresponding to the protons of *tert*-butyl group (Fig. 1B-a) and the more stronger absorption peak at 1730 cm^{-1} which is assigned to the stretching vibrations of $\text{C}=\text{O}$ group of PtBA (Fig. 2B-a) both indicate the successful synthesis of PF-*b*-PtBA. After hydrolysis, the ^1H NMR spectrum of PF-*b*-PAA in DMSO (Fig. 1B-b) show the obviously decreases of the *tert*-butyl resonance at 1.44 ppm and the appearance of the peak at 12.25 ppm assigned to the protons of carboxyl groups. In the FTIR spectrum of PF-*b*-PAA (Fig. 2B-b), a new adsorption peak characteristic of carboxylic acid group appears at 3436 cm^{-1} and the signal at 1370 cm^{-1} attributed to the symmetric bending vibration of the *tert*-butyl group disappears. These results reveal the successful preparation of PF-*b*-PAA (the original ^1H NMR and FTIR spectra of PF-Br for comparison are respectively shown in Fig. S1-b and S2-b of the ESI†).

The ^1H NMR spectra of PF-*b*-PHEMA-TMS and PF-*b*-PHEMA are shown in Fig. 1C. In Fig. 1C-a, the appearance of peak at 3.75 ppm corresponding to the protons of the $-\text{CH}_2-$ group of $-\text{CH}_2-\text{O}-\text{Si}(\text{CH}_3)_3$, and the appearance of peak at 0.16 ppm corresponding to the protons of $-\text{CH}_3$ groups of $-\text{Si}(\text{CH}_3)_3$, both indicate the successful incorporation of HEMA-TMS to PF segment. After the deprotection reaction using KF/TBAF as the catalyst, the disappearance of peak at 0.16 ppm (Fig. 1C-b) indicate that the $-\text{TMS}$ groups on PHEMA are successful removed. The successful synthesis of PF-*b*-PHEMA also can be confirmed by the appearance of the adsorption peak at 3436 cm^{-1} in Fig. 2C-b, which is assigned to the stretching vibrations of the $-\text{OH}$ group of PHEMA (the original ^1H NMR and FTIR spectra of PF-Br for comparison are respectively shown in Fig. S1-b and S2-b of the ESI†).

Fig. 3 shows the GPC traces for the four as-prepared copolymers. The relatively narrow polydispersity indicates the good controllability of polymerization. However, the result of PF-*b*-PHFBMA seems to be unusual, which might be due to the collapsed coil structure of PHFBMA block in THF.³⁴ Additionally, the molecular weights of PF-*b*-PHEMA ($M_n = 13\,137\text{ g mol}^{-1}$) and PF-*b*-PAA ($M_n = 10\,502\text{ g mol}^{-1}$) can be calculated by the results of PF-*b*-PHEMA-TMS and PF-*b*-PtBA, since PF-*b*-

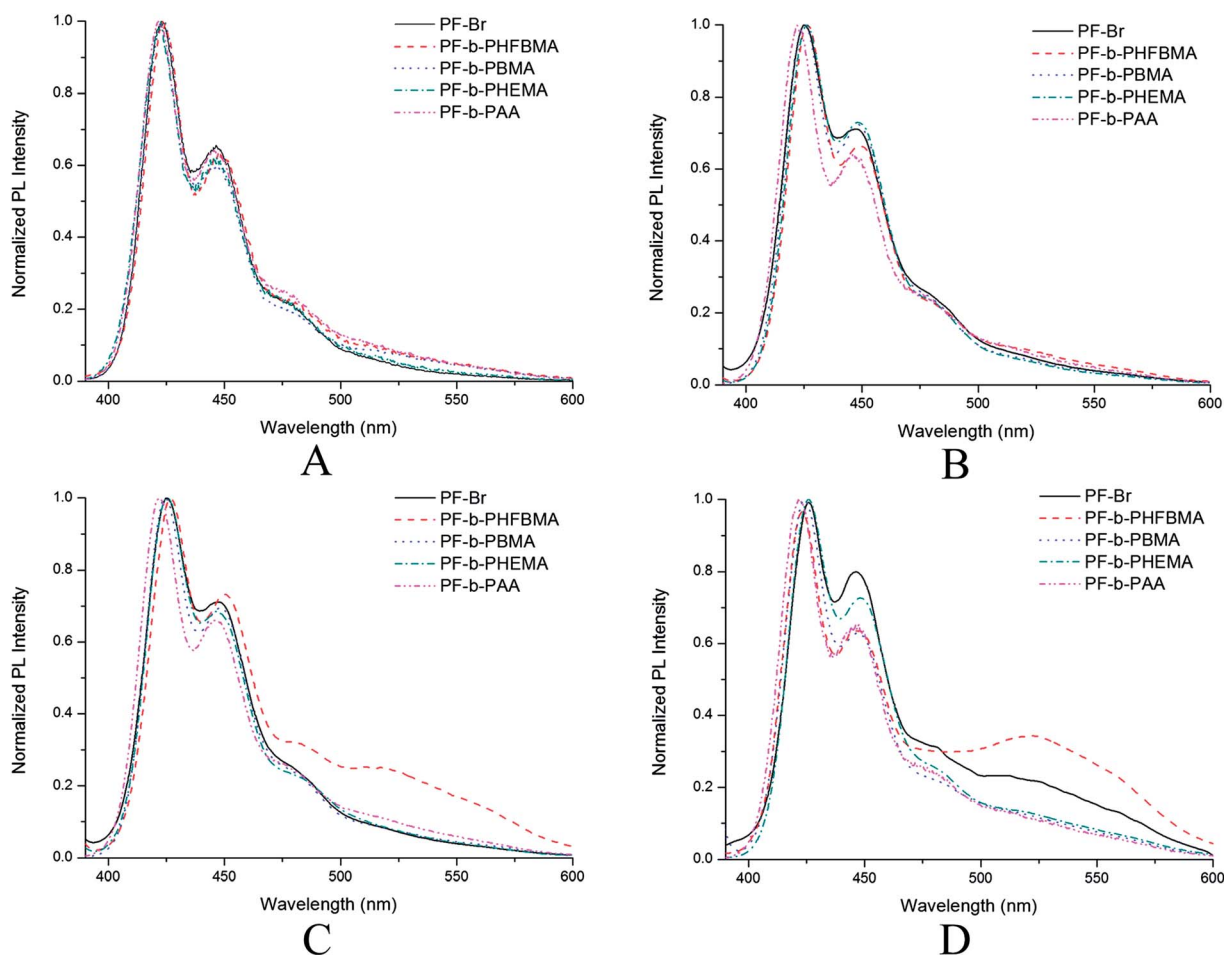


Fig. 5 Fluorescence spectra of PF-Br, PF-*b*-PHFBMA, PF-*b*-PBMA, PF-*b*-PHEMA and PF-*b*-PAA films (A) annealed 1 h at $50\text{ }^\circ\text{C}$ in air, (B) annealed 1 h at $100\text{ }^\circ\text{C}$ in air, (C) annealed 1 h at $150\text{ }^\circ\text{C}$ in air and (D) annealed 1 h at $200\text{ }^\circ\text{C}$ in air.

PHEMA-TMS and PF-*b*-PtBA exhibit better solubility in THF eluent.

The influence of annealing on the spectral stability of polymers

It is well known that the formation of fluorenone defect structure might affect the spectral stability of polymers during the polymerization of fluorene monomer. Therefore, the hydroxyl-ended polyfluorene was prepared in the dark. The ^{13}C NMR spectrum of PF-OH in Fig. 4 shows that the characteristic peak at $\delta = 194$ ppm corresponding to carbonyl is absence, which confirms that fluorenone defects are prevented successfully during the reaction.³⁶

After excluding the impact of fluorenone defects, the effect of incorporating different coil segments on the spectral stability of polyfluorene-based copolymers have been investigated under different thermal treatment conditions, including annealing temperature, annealing time and annealing atmosphere. Green index ϕ , defined as intensity ratio of green emission (~ 530 nm) to the strongest peak of the blue emission ($I_{\text{green}}/I_{\text{blue}}$),^{37,38} was introduced to depict the low-energy emission. The larger the value of ϕ is, the stronger the low-energy emission is, which indicates the poorer the spectral stability.

First, the spectral stability of polyfluorene-based polymers under different annealing temperature for 1 h was investigated.

From the results shown in Fig. 5(A)–(D), no new emission band appears in the spectra of PF-Br, PF-*b*-PHFBMA, PF-*b*-PBMA, PF-*b*-PHEMA and PF-*b*-PAA when the annealing temperature is 50 °C or 100 °C. As the temperature increases to 150 °C, emission band at 525 nm only appears in the spectrum of PF-*b*-PHFBMA, the green index $\phi = 0.25$. When the temperature continues increasing to 200 °C, this emission band is also observed in the case of PF-Br, the green index $\phi = 0.23$, simultaneously, the intensity of this emission band in the spectrum of PF-*b*-PHFBMA increases significantly, the green index $\phi = 0.34$. As a whole, the annealing temperature has influence on the spectral stability to some extent.

Fig. 6(A)–(D) show the spectra of polyfluorene-based polymers PF-Br, PF-*b*-PHFBMA, PF-*b*-PBMA, PF-*b*-PHEMA and PF-*b*-PAA under different annealing time at 150 °C in air. When the annealing time is 1 h, the emission band at 525 nm only appears in the spectrum of PF-*b*-PHFBMA, the green index $\phi = 0.25$ (Fig. 6(B)). As the annealing time go on, the spectra of PF-*b*-PBMA, PF-*b*-PHEMA and PF-*b*-PAA still maintain steady while the emission band at 525 nm is observed in the spectrum of PF-Br at 6 h, the green index $\phi = 0.22$ (Fig. 6(D)). The intensity of the emission band at 525 nm appeared in the spectrum of PF-*b*-PHFBMA increases significantly ($\phi = 0.69$) as the annealing time increase to 6 h (Fig. 6(D)). The above results suggest that polyfluorene can maintain the spectral stability in a short

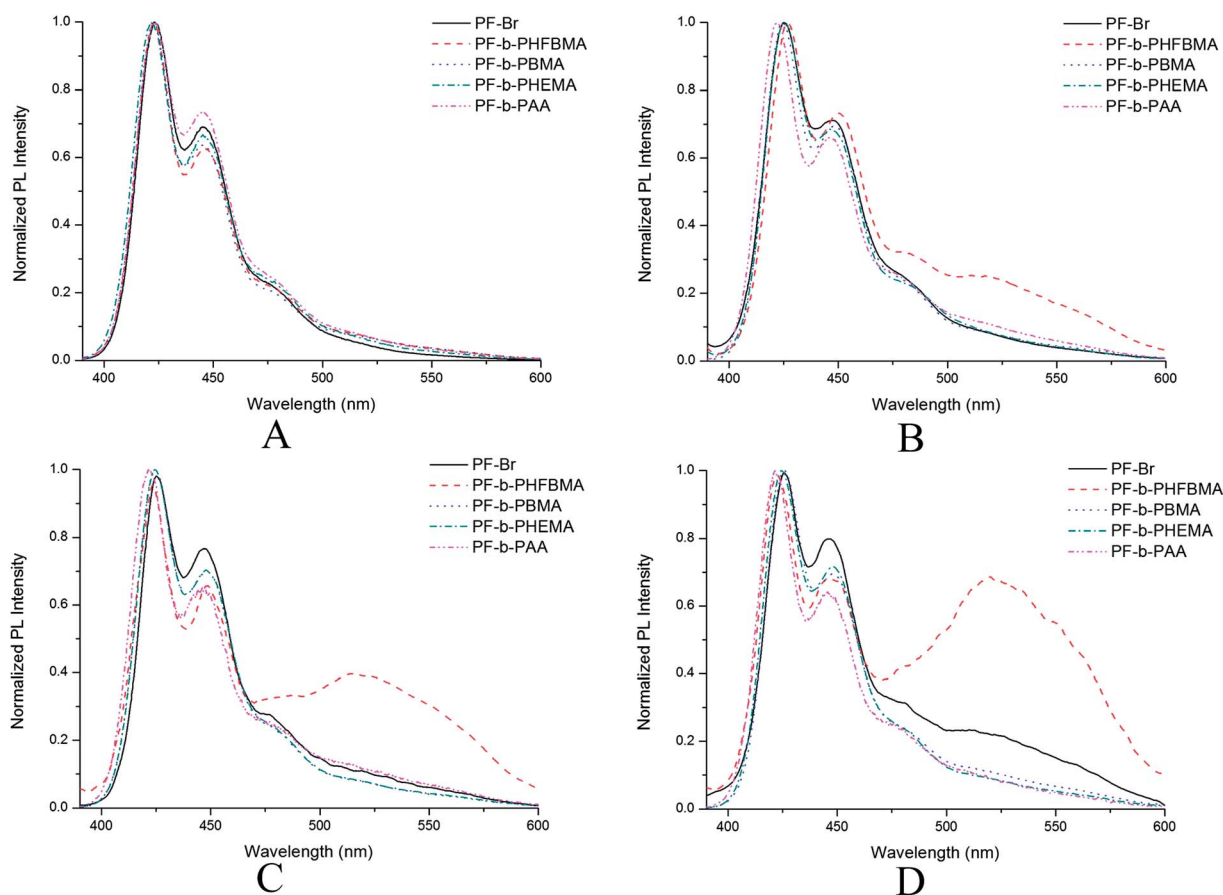


Fig. 6 Fluorescence spectra of PF-Br, PF-*b*-PHFBMA, PF-*b*-PBMA, PF-*b*-PHEMA and PF-*b*-PAA films (A) pristine spin-cast, (B) annealed 1 h, (C) annealed 3 h and (D) annealed 6 h at 150 °C in air.

period annealing time. However, when the annealing time exceeds a certain value, low-energy emission begin to appear. In a word, introducing the coil blocks PBMA, PHEMA and PAA can improve the spectral stability of polyfluorene while incorporating the coil block PHFBMA has the opposite effect.

To confirm whether the fluorenone defect structure was formed due to annealing in air, the spectral stability of polyfluorene-based polymers annealing in different atmosphere at 150 °C for 1 h was investigated. The corresponding emission spectra are depicted in Fig. 7. When compared to the pristine spin-casted film, no low-energy emission is present in the spectra of PF-Br, PF-*b*-PBMA, PF-*b*-PHEMA and PF-*b*-PAA either in vacuum or in air. In contrast, PF-*b*-PHFBMA exhibits the 525 nm emission band in air and in vacuum. From the results above, we suppose that the fluorenone defect structure might not form in air due to the limited oxidation in solid state, and the low-energy emission of PF-*b*-PHFBMA film might be resulted from molecular aggregation.

In combination with the results discussed above, both PF-Br and PF-*b*-PHFBMA exhibit the 525 nm low-energy emission

band after annealing in air at 200 °C for 1 h (Fig. 5(D)) or at 150 °C for 6 h (Fig. 6(D)). The ¹³C NMR spectrum of PF-OH (Fig. 4) and the spectra of rod-coil copolymer films annealed in vacuum (Fig. 7(C)) excluding the impact of fluorenone defects on the spectral stability of polyfluorene-based polymers. Therefore, we tentatively suggest that introducing the coil blocks PBMA, PHEMA and PAA can inhibit the low-energy emission of polyfluorene while incorporating the coil block PHFBMA has the opposite effect, and aggregation behavior between fluorene chain is the cause of the low-energy emission band.

To PF-*b*-PHFBMA, the unfavorable enthalpic interactions between the fluorinated segment and the PF block can drive the segregation of copolymer. During the segregation process, PHFBMA is oriented to gather on the air-polymer surface due to its low surface free energy and self-aggregated property. If the annealing temperature is constant, the amount of fluorinated segments on the surface will increase gradually with the increase of annealing time, and finally reached a maximal value.^{39,40} The accompanied aggregation of PF blocks on the

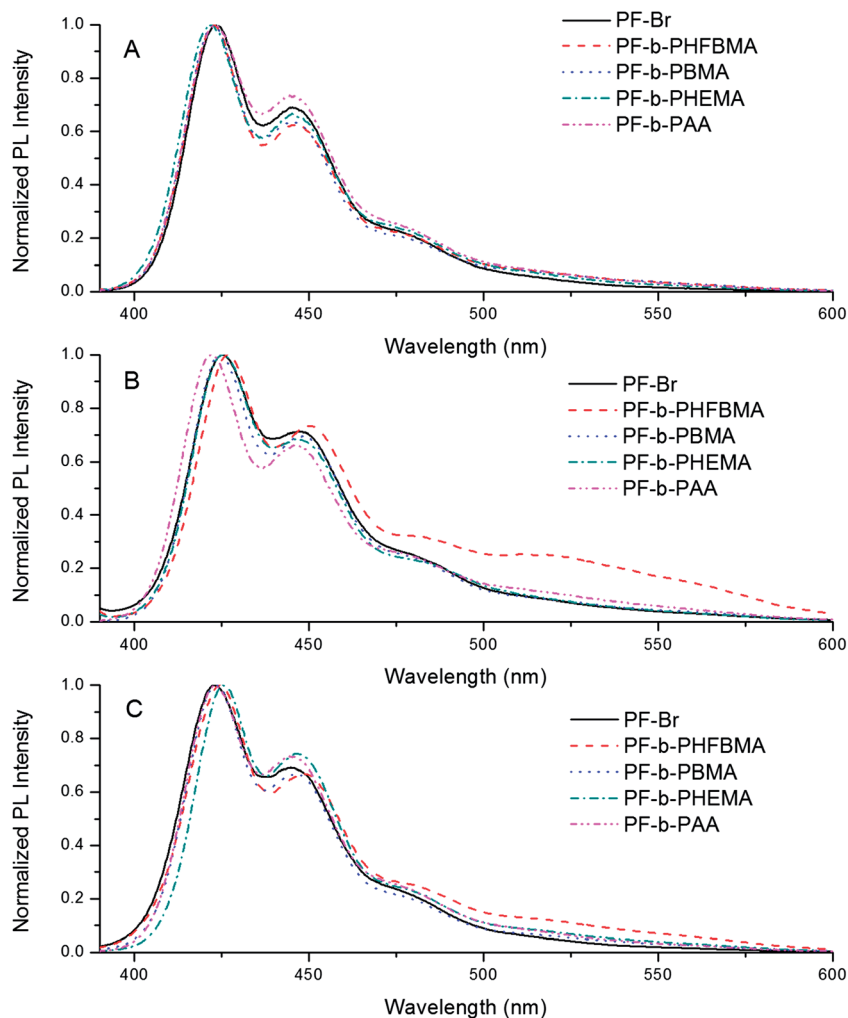


Fig. 7 Fluorescence spectra of PF-Br, PF-*b*-PHFBMA, PF-*b*-PBMA, PF-*b*-PHEMA and PF-*b*-PAA films (A) pristine spin-cast, (B) annealed 1 h at 150 °C in air, (C) films annealed 1 h at 150 °C in vacuum.

other side causes the low-energy emission band, and the intensity of the emission gradually increase as the annealing time go on (Fig. 6). The increase of the annealing temperature can promote the moving of PHFBMA segments and the aforementioned maximal value can be reached in a shorter period time when the temperature is higher.^{39,40} It is a reasonable explanation about the appearance of low-energy emission band and the enhanced intensity of the emission as the increase of the annealing temperature (Fig. 5). Compared with PHFBMA block, PBMA, PHEMA and PAA blocks have a better compatibility with PF, the separation trend of copolymers PF-*b*-PBMA, PF-*b*-PHEMA and PF-*b*-PAA are relatively weaker. Aggregation of PF blocks is less likely to occur even at a relative high temperature. Therefore, increasing annealing temperature or extending annealing time, a low-energy emission band never appear in the spectra of the polyfluorene-based copolymers with PBMA, PHEMA and PAA as coil blocks (Fig. 5 and 6).

Surface properties

The further investigation about the surface properties of as-prepared films is to gain some supporting information for the above conclusion. By comparison of the results obtained before and after annealing at 150 °C in air for 1 h, one can find that the WCA of PF-*b*-PHFBMA film increases from 107° to 113° and the other as-prepared films have no distinct difference in Fig. 8A and B. The phenomenon confirms the fact that fluorinated blocks prefer to migrate onto the surface of film, and the accompanied aggregation of PF blocks on the other side causes

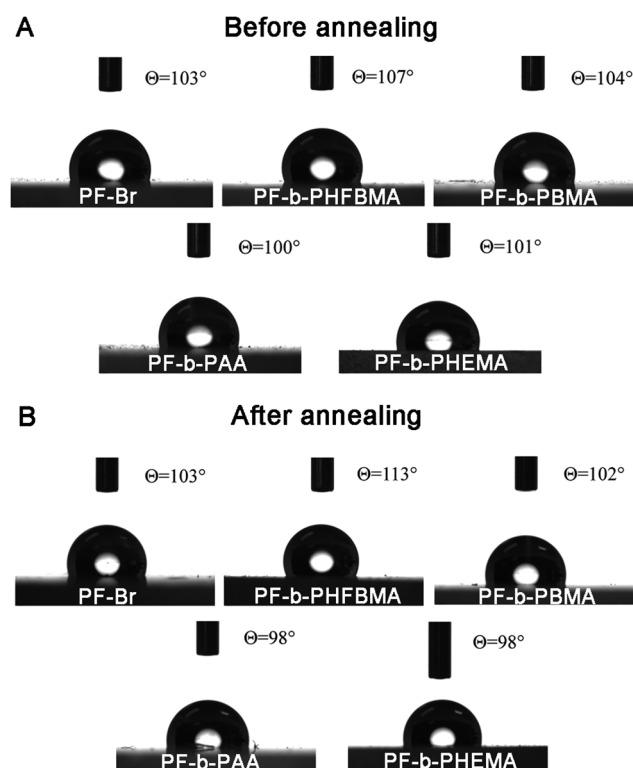


Fig. 8 The static WCA images of films (A) before annealing and (B) after annealing at 150 °C in air for 1 h.

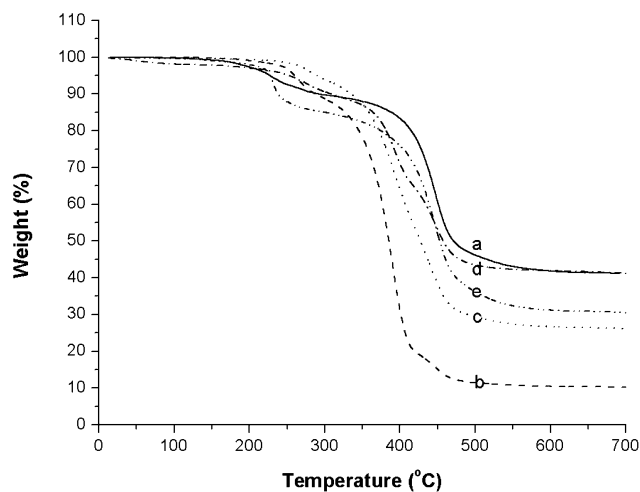


Fig. 9 TGA curves of (a) PF-Br, (b) PF-*b*-PHFBMA, (c) PF-*b*-PBMA, (d) PF-*b*-PHEMA and (e) PF-*b*-PAA.

the low-energy emission band. These provide the supporting information for the spectral instability of PF-*b*-PHFBMA film.

Thermal properties

The thermal behavior of the initiator PF-Br and the polyfluorene-based rod-coil block copolymers were studied using TGA (Fig. 9). It is evident from Fig. 9 that PF-Br presents two thermal degradation temperatures at 225 °C and 425 °C, which are attributed to the decomposition of the micromolecules and polyfluorene, respectively. The rest weight percentage of PF at 700 °C is 41.4%. Block copolymers PF-*b*-PHFBMA, PF-*b*-PBMA, PF-*b*-PHEMA and PF-*b*-PAA undergo two-step decompositions, the flexible-coil blocks first decomposition under a lower temperature (225–330 °C). Subsequently, the PF rod segments begin to decompose from 350 °C to 500 °C. Their rest weight percentage at 700 °C are 10.2%, 26.1%, 41.1% and 30.5%, respectively. The thermal degradation temperatures of polyfluorene-based rod-coil block copolymers decrease slightly compared to PF-Br, indicating that the incorporation of coil blocks with polyfluorene have a negative effect on the thermal stability of the polymers. However, these rod-coil block copolymers still are thought to be good thermal stability within the slightly reduction.

Conclusion

A series of polyfluorene-based rod-coil block copolymers, including PF-*b*-PHFBMA, PF-*b*-PBMA, PF-*b*-PHEMA and PF-*b*-PAA, were synthesized *via* ATRP technique using PF-Br as macroinitiator. The macroinitiator was prepared through modified Suzuki coupling reaction and esterification. The structures of resulted copolymers were confirmed by the ¹H NMR and FTIR spectra. The investigations about spectral stability of polyfluorene-based rod-coil block copolymers under different thermal treatment operations demonstrate that incorporating the coil blocks PBMA, PHEMA and PAA can

prevent the low-energy band emission while the coil block PHFBMA produce the opposite effect on the spectral stability. A proper coil block is essential for the stabilization of the emission spectra of polyfluorene-based copolymers. Simultaneously, after excluding the impact of fluorenone defects, aggregation behavior between fluorene chains is considered to be the cause of the low-energy emission band. Furthermore, surface properties of copolymers are used to further explore the supporting information for the spectral instability of PF-*b*-PHFBMA film.

The thermal behaviors of all polymers investigated by TGA indicate that incorporation of coil blocks with polyfluorene does not much affect the thermal properties of polymers, which can be used to prepare the electronic device with inhibited low-energy emission.

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