

Cite this: *RSC Adv.*, 2017, 7, 12269

Hyperbranched conjugated polymers containing 1,3-butadiene units: metal-free catalyzed synthesis and selective chemosensors for Fe³⁺ ions†

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Hyperbranched conjugated polymers (P1 and P2) containing 1,3-butadiene repeating units appended with carboxylic ester groups were synthesized in good yield by metal-free catalyzed polymerization of diethyl 4,4'-(5-formyl-1,3-phenylene)(2*E*,2'*E*)-bis(but-2-enoate) (M1) or dimethyl 4,4'-(5-formyl-1,3-phenylene)(2*E*,2'*E*)-bis(but-2-enoate) (M2). The corresponding P1 and P2 polymers were structurally characterized by NMR and IR spectroscopy. The polymers are highly soluble in common organic solvents. The UV-vis absorption bands of P2 along with its emission spectrum was red-shifted compared to that of P1, indicating the extension of conjugated length. The hyperbranched conjugated polymers were used as ratiometric fluorescence chemosensors for the detection of the biologically important metal ion Fe³⁺. The corresponding Stern–Volmer constants (*K*_{sv}) of P1 and P2 are $2.90 \times 10^4 \text{ M}^{-1}$ and $3.83 \times 10^4 \text{ M}^{-1}$, respectively. Detection limits of $6.99 \times 10^{-7} \text{ M}$ for P1 and $6.69 \times 10^{-7} \text{ M}$ for P2 were achieved. The neutral polymer P1 or P2 was hydrolyzed to obtain water-soluble hyperbranched conjugated polymer P3. The ability of P3 to detect different Fe³⁺ ions in aqueous solution was investigated. The results demonstrate that these polymers can serve as sensitive and selective sensors for Fe³⁺ ions in organic and aqueous solutions.

Received 13th January 2017
Accepted 6th February 2017

DOI: 10.1039/c7ra00540g

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Introduction

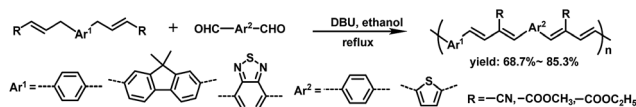
Hyperbranched conjugated polymers have attracted much attention because of their unique properties, such as good solubility, low viscosity, tunable electrical properties and processability, and also because of their potential applications in organic electronics and optoelectronics, including organic electroluminescence,^{1–5} ion detection,^{6–8} explosive detection,^{9–14} disease diagnosis,¹⁵ and biological identification.^{16,17} Various transition-metal-catalyzed polymerization methods have been used for the synthesis of hyperbranched conjugated polymers, such as Suzuki coupling polymerization,^{18–20} click polymerization,^{21,22} and Sonogashira coupling polymerization.^{23,24} These well-established polymerization routes have played an indispensable role in the synthesis of hyperbranched conjugated polymers. However, these methods have drawbacks, such as harsh reaction conditions, tedious synthetic steps, the use of environmentally harmful transition-metal catalysts and solvents. In addition, the trace residues of transition-metal catalysts in polymeric products can seriously affect the efficiency and the life of the resulting devices.²⁵

Recently, various linear conjugated polymers were synthesized with transition-metal-free catalyzed methods.^{26–35} However, these methods, including Wittig reaction,^{4,36} Knoevenagel condensation reaction,¹¹ arylimino-deoxy-trisubstituted reaction,³⁷ and acid-promoted cyclotrimerization reactions,³⁸ were rarely explored to prepare hyperbranched conjugated polymers. In addition, these transition-metal-free catalyzed methods also require harsh reaction conditions and complex synthetic steps. Therefore, in order to expand the application of hyperbranched conjugated polymers, it is desirable to develop efficient polymerization method and new structure under mild reaction conditions by using transition-metal-free catalyst.

In our previous work,³⁹ we have established an efficient transition-metal-free catalyzed polymerization method. Reactions of α,β -unsaturated compounds containing γ -H with aldehydes in ethanol in the presence of 1,8-diazabicyclo[5.4.0]undecene-7-ene (DBU) as the basic catalyst gave a series of linear π -conjugated polymers (Scheme 1), which have alternating arylene and 1,3-butadiene units in the main chain. This method has many advantages, including fewer reaction steps, mild reaction conditions, environmentally friendly solvents (*i.e.*, in ethanol even without purity and protection from oxygen), and no toxic metal byproducts. On the basis of these results, herein we report the synthesis of new hyperbranched conjugated polymers containing carboxylic ester groups appended to repeating 1,3-butadiene units and their uses as sensitive and selective sensors for Fe³⁺ ions in organic and aqueous solutions. Iron is the second most abundant metal

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† Electronic supplementary information (ESI) available. See DOI: 10.1039/c7ra00540g



Scheme 1 Synthesis of π -conjugated polymers containing 1,3-butadiene units in the main chain.

element in nature, and plays very important roles in the whole ecosystem.⁴⁰ Recently, fluorescence sensors based on conjugated polymers for the detection of Fe^{3+} have received much attention owing to their signal amplification properties.^{41–48}

This special macromolecular structure shows several advantages in Fe^{3+} detection. First, the carboxylic ester group has been reported to act as chelating group for metal ions, especially for Fe^{3+} ions. Therefore, neutral polymers P1 and P2 are expected to coordinate with Fe^{3+} in organic solutions, and the strong association interaction between carboxylic ester groups and Fe^{3+} ions will cause the fluorescence quenching.⁴⁹ Second, the carboxylic ester group could be hydrolyzed to obtain water-soluble hyperbranched conjugated polymers. We expected that our synthesized hyperbranched conjugated polymers could be used in the field of selective detection of Fe^{3+} from environmental and biological samples.

Experimental

Materials and instruments

Chemical reagents and different metal salts were purchased from Energy chemical, Merck or Aladdin and used as received. THF and 1,4-dioxane were distilled from sodium benzophenone ketyl under dry nitrogen immediately prior to the use. ^1H and ^{13}C NMR spectra were recorded on a Bruker 600 MHz spectrophotometer in CDCl_3 using tetramethylsilane (TMS, $\delta = 0$) as an internal reference. IR spectra were taken on a FT-IR Bruker Tensor 27 spectrometer at a nominal resolution of 2 cm^{-1} using KBr pellet. UV-vis spectra were collected on Hitachi U-3900 spectrophotometer (Tokyo, Japan). Photoluminescence (PL) spectra were recorded on a Hitachi F-7000 spectrofluorometer (Tokyo, Japan). Thermogravimetric analysis (TGA) measurements were carried out under dry nitrogen on Q1000DSC + LNCS + FACSQ600SDT instruments at $20\text{ }^\circ\text{C min}^{-1}$. The average molecular weight and polydispersity index (PDI) of the polymers were determined using Viscotek gel permeation chromatography (GPC) analysis with THF as the eluent and polystyrene as the standard. Cyclic voltammetry (CV) was conducted on an electrochemistry workstation (CHI 660A, Chenhua Shanghai) with the polymer film on Pt plate as the working electrode, Pt slice as the counter electrode, and saturated calomel electrode (SCE) as a reference electrode in a 0.1 M tetra-*n*-butylammonium hexafluorophosphate acetonitrile solution at a scan rate of 50 mV s^{-1} . The pH value was measured by using of a MRFS: TOSHNIWAL pH meter.

Monomer synthesis

3,5-Bis(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)benzaldehyde (1). Compound **1** was synthesized according to published

procedures with a yield of 71%.⁵⁰ ^1H NMR (400 MHz, CDCl_3) δ (ppm): 10.06 (s, 1H), 8.49 (s, 1H), 8.41 (s, 2H), 1.36 (s, 24H). Elemental analysis for **1** ($\text{C}_{19}\text{H}_{28}\text{B}_2\text{O}_5$): C, 63.74; H, 7.88; B, 6.04; O, 22.34. Found: C, 63.68; H, 7.84; B, 6.12; O, 22.39.

Diethyl 4,4'-(5-formyl-1,3-phenylene)(2*E*,2'*E*)-bis(but-2-enoate) (M1). Compound **1** (1 g, 2.79 mmol), ethyl (*E*)-4-bromobut-2-enoate (1.6 g, 8.37 mmol) and KF (1.62 g, 27.9 mmol) were dissolved in anhydrous 1,4-dioxane (30.00 mL). The solution was bubbled under nitrogen at room temperature for 10 min, and then, $\text{Pd}(\text{OAc})_2$ (14.7 mg, 0.15 mmol) was added. The reaction mixture was stirred for 72 hours at room temperature. The reaction solution was then washed with saturated NaCl water and extracted with Et_2O (100 mL \times 3). The combined organic layer was dried with magnesium sulfate, and the solvent was removed by rotary evaporation. The crude product M1 was purified on silica gel chromatography using petroleum ether/ethyl acetate (5/1, v/v) as the eluent to afford a light yellow oily liquid (1.1 g, 60%). ^1H NMR (400 MHz, CDCl_3) δ (ppm): 9.99 (s, 1H), 7.59 (s, 1H), 7.27 (s, 1H), 7.10–7.01 (m, 1H), 5.86–5.79 (m, 1H), 4.19 (d, $J = 6.8\text{ Hz}$, 2H), 3.59 (d, $J = 6.0\text{ Hz}$, 2H), 1.29 (d, $J = 6.8\text{ Hz}$, 3H). ^{13}C NMR (100 MHz, CDCl_3) δ (ppm): 191.8, 166.1, 145.7, 139.3, 137.2, 135.2, 128.3, 123.1, 77.3, 77.0, 76.6, 60.3, 37.8, 14.1. Elemental analysis for M1 ($\text{C}_{19}\text{H}_{22}\text{O}_3$): C, 76.48; H, 7.43; O, 16.09. Found: C, 76.39; H, 7.35; O, 16.13.

Dimethyl 4,4'-(5-formyl-1,3-phenylene)(2*E*,2'*E*)-bis(but-2-enoate) (M2). The same procedure for M1 was applied for the synthesis of M2, using methyl (*E*)-4-bromonut-2-enoate instead of ethyl (*E*)-4-bromonut-2-enoate. The crude product M2 was purified on silica gel chromatography using petroleum ether/ethyl acetate (4/1, v/v) as the eluent to afford a light yellow oily liquid (1.16 g, 63%). ^1H NMR (400 MHz, CDCl_3) δ (ppm): 9.98 (s, 1H), 7.58 (s, 2H), 7.26 (s, 1H), 7.12–7.05 (m, 2H), 5.85 (d, $J = 15.5\text{ Hz}$, 2H), 3.73 (s, 6H), 3.59 (d, $J = 6.4\text{ Hz}$, 4H). ^{13}C NMR (100 MHz, CDCl_3) δ (ppm): 191.8, 166.5, 146.0, 139.3, 137.2, 135.1, 128.3, 122.7, 77.2, 77.0, 76.7, 51.5, 37.8. Elemental analysis for M2 ($\text{C}_{17}\text{H}_{18}\text{O}_3$): C, 75.53; H, 6.71; O, 17.76. Found: C, 75.45; H, 6.67; O, 17.82.

General polymerization

To a flask charged with monomer in ethanol solution, DBU was added. The reaction mixture was stirred under refluxed for 72 h. The reaction mixture was then heated at high temperature ($110\text{ }^\circ\text{C}$) for 8 h. The solvent was removed under reduced pressure, and the residue was extracted with CH_2Cl_2 and washed with HCl solution (2 M, 100 mL \times 3) and water. The solvent was removed under reduced pressure.

Synthesis of polymer P1. M1 (0.3 g, $9.08 \times 10^{-4}\text{ mol}$) and DBU (62.16 mg, $4.09 \times 10^{-4}\text{ mol}$) were used. P1 was obtained as yellow solid (yield: 79%). ^1H NMR (400 MHz, CDCl_3) δ (ppm) 7.82–7.27 (8H, ArH, C=CH), 6.63–6.62 (d, $J = 5.6\text{ Hz}$, 2H, CH=CH), 6.52–6.50 (d, $J = 7.2\text{ Hz}$, 2H, CH=CH), 4.12–4.08 (m, 4H, CH_2), 1.23–1.19 (t, 6H, CH_3). IR (KBr pellet, cm^{-1}): 3030, 2914, 1740, 1600, 1450, 1350.

Synthesis of polymer P2. M2 (0.3 g, $9.92 \times 10^{-4}\text{ mol}$) and DBU (62.16 mg, $4.09 \times 10^{-4}\text{ mol}$) were used. P2 was obtained as



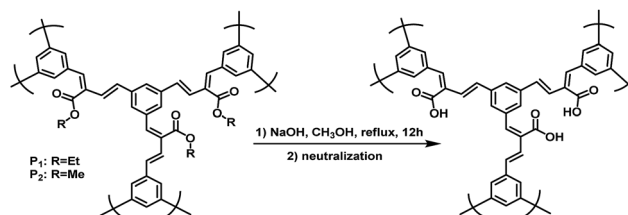
yellow solid (yield: 80%). ^1H NMR (400 MHz, CDCl_3) δ (ppm) 7.62–7.52 (9H, ArH, C=CH), 6.61–6.59 (d, $J = 9.2$ Hz, 2H, CH=CH), 6.50–6.47 (d, $J = 12.8$ Hz, 2H, CH=CH), 4.09 (s, 6H, CH_3). IR (KBr pellet, cm^{-1}): 3004, 2908, 1745, 1650, 1480, 1380.

Hydrolysis of P1 or P2 to form P3. Neutral carboxylic-ester-containing polymer (P1 or P2) was added to a solution of sodium hydroxide in methanol; the mixture was refluxed for 12 h.⁶ After cooling to room temperature, the mixture was neutralization with dilute hydrochloric acid, the solvent was removed by rotary evaporation and the obtained mixture was dialyzed (MW cutoff, 4 kDa) against water for 3 days to remove small molecules. FT-IR (KBr, cm^{-1}): 3450, 1800, 1650, 1400, 700.

Results and discussion

Synthesis and structure characterization

The polymer structures and synthetic routes are shown in Scheme 2. Compound **1** was synthesized according to the published procedures.⁵⁰ 3,5-Bis(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl) benzaldehyde (**1**), diethyl 4,4'-(5-formyl-1,3-phenylene)(2*E*,2'*E*)-bis(but-2-enoate) (**M1**) and dimethyl 4,4'-(5-formyl-1,3-phenylene)(2*E*,2'*E*)-bis(but-2-enoate) (**M2**) were prepared by Pd-catalyzed Suzuki cross-coupling reaction under argon atmosphere. In our previous work, we have established a transition-metal-free polymerization method and successfully synthesized a series of linear conjugated polymers.³⁹ In this work, the hyperbranched conjugated polymers P1 and P2 were synthesized by the simple AB₂ protocol based on our reported polymerization method. The polycondensation was carried out in the same way by refluxing in ethanol for 72 h with 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU) as the catalyst. The yield of P1 and P2 were 79% and 80%, respectively. This result indicates that the molecular weight is very less affected by the side group. It is worth noting that the carboxylic ester-functional group can be readily obtained without special reaction for the monomer during polymerization. All the polymers displayed excellent solubility in common organic solvents such as chloroform, tetrahydrofuran, dichloromethane and chlorobenzene. The water-soluble polymer P3 was obtained by the hydrolysis of P1 or P2 in a solution of sodium hydroxide in methanol with a yield of 80% (Scheme 3). Table 1 summarized



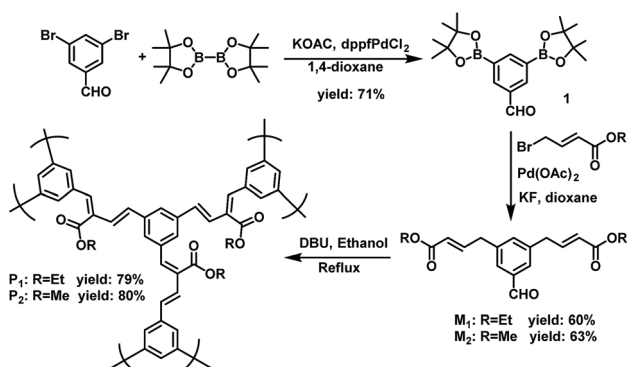
Scheme 3 Synthetic routes of P3.

the polymerization results and thermal properties of the polymers. The number-average molecular weight (M_n) of P1 and P2 are 6141 and 6026 with polydispersity index (PDI) of 2.575 and 2.465, respectively. It can be seen that in general, the polymerization afforded conjugated polymers with relatively high molecular weight.

All the monomers were carefully purified and characterized by ^1H and ^{13}C NMR (see Experimental section and ESI† for details). The structures of polymer P1 and P2 were verified by ^1H NMR and FT-IR. In the ^1H NMR spectrum of P1, the peaks attributable to the aromatic protons were observed at 7.82–7.27 ppm. Another set of olefinic protons were observed at 6.63–6.62 and 6.52–6.50 ppm. The proton signals of ester methylene and methyl for P1 appear at 4.12–4.08 and 1.23–1.19 ppm, respectively (Fig. S6†). In the ^1H NMR spectrum of P2, the peak at 4.09 ppm is assigned to the signal from $-\text{COOCH}_3$ of P2 (Fig. S7†). Examples of the IR spectra of P1 and P2 are given in Fig. S9.† P1 and P2 shows absorption bands at 3030 and 3004 cm^{-1} due to their CH=CH stretching vibrations. The strong peaks at 1740 cm^{-1} for P1 and 1745 cm^{-1} for P2 are associated with their C=O groups. As for P3, we see a broad band of strong absorption peaks at 3450 cm^{-1} due to its OH stretching vibrations of COOH groups. IR spectrum indicated successful synthesis of the target polymers P1, P2 and P3. Because of the poor resolution of related resonances in the ^1H NMR spectra of P1 and P2, the important parameter of degree of branching is hardly assessed.

Thermal properties

The thermal properties of the target polymers were obtained by thermogravimetric analysis (TGA) under a nitrogen atmosphere at a heating rate of 20 $^\circ\text{C min}^{-1}$. The TGA diagram of P1 (shown in Fig. S10†) showed a mass loss of 5% starting at 240 $^\circ\text{C}$ as the threshold of the thermal decomposition, and a loss of about 50% at 500 $^\circ\text{C}$. On the other hand, the TGA trace of P2 showed



Scheme 2 Synthetic routes of monomers and polymers.

Table 1 Molecular weights and thermal properties of polymers

Polymer	Yield (%)	M_n^a	M_w^a	PDI ^a	T_d^b [$^\circ\text{C}$]
P1	72%	6141	15 812	2.575	240
P2	75%	6026	14 854	2.465	200

^a Determined by GPC at using THF as an eluent against polystyrene standards. ^b 5% weight loss temperature measured by TGA under a nitrogen atmosphere.



that polymer decomposition started at 200 °C and the weight loss was about 55% at 600 °C.

Photophysical properties

Fig. 1a and b depict the UV-vis spectra of polymer P1 and P2, respectively, measured in different organic solvents, including CH₂Cl₂, DMSO, CH₃CH₂OH, THF, and in solid-state films. P1 and P2 exhibited no significant absorption peaks in most tested organic solvents, while they showed significant absorption peaks at 325 and 330 nm in THF, respectively (Fig. 1). The different absorption features of P1 and P2 in these solvents are due to the different polarity of solvents. It is reported that molecular aggregation in solution exhibited distinct change in the absorption band and can possibly be influenced by the polarity of solvents.¹⁶ The polymer P1 in THF solution showed an absorption maximum at 325 nm attributable to phenyl moiety. The solid film of P1 also showed a similar absorption behaviour with an absorption maximum at 327 nm. This result indicates the low aggregation degree of P1 hyperbranched chains in the thin film state. On the other hand, polymer P2 in THF solution showed an absorption peak at 330 nm corresponding to phenyl moiety. The absorption peaks were red-shifted compared to those of P1, suggesting an extension of the conjugated length. It is likely that the substituent has a steric influence on the electronic structure, possibly due to the fact that bulky groups can interrupt the linear π -system, leading to the reduction in the π -conjugated length. The solid film of P2 showed a similar absorption behavior with an absorption maximum at 350 nm. The absorption peaks are red-shifted compared to their solution absorption spectra, which was mainly attributed to the enhanced intermolecular interactions between neighboring molecules in the film state.

The photoluminescent (PLs) spectra of the hyperbranched conjugated polymers in solution excited at the absorption maxima are shown in Fig. 2a and b. As shown in Fig. 2a and b, P1 and P2 exhibited low fluorescence intensity in most organic solvents, while they showed relatively high fluorescence intensity in THF: P1 displayed an emission peak at 410 nm (450 nm in the solid state); P2 showed a peak maximum at 420 nm (460 nm in the solid state). The emission peaks of both P1 and P2 in the solid state are broaden and red-shifted compared with those in THF solution. This result suggests the formation of aggregates in the solid state. The fluorescence quantum yields (Φ_F) of the polymers in THF were measured in comparison with

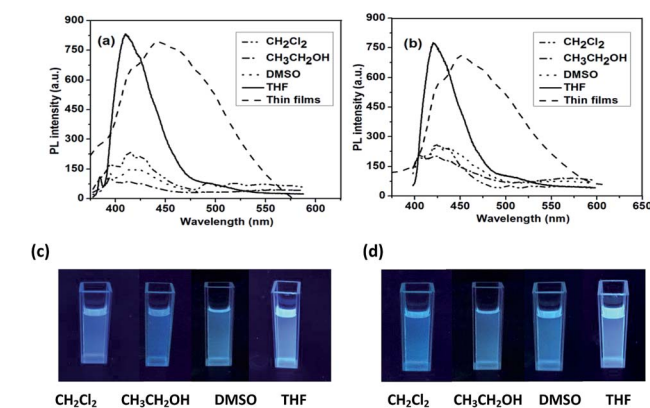


Fig. 2 Fluorescence spectra of (a) P1 and (b) P2. Emission images observed for (c) P1 and (d) P2 measured in different solutions.

quinine sulfate as a standard (in 0.1 M H₂SO₄ solution, $\Phi_F = 0.55$).⁵¹ The results are 32% for P1 and 34% for P2. The relatively low quantum yields of P1 and P2 might be caused by imperfect structures presented in the hyperbranched polymers with higher contents of branch units and carboxylic ester groups attached to the main chain, which reduced the conjugated length.

Electrochemical properties

The electrochemical behaviors of polymers P1 and P2 were investigated by cyclic voltammetry (CV) as shown in Fig. 3 (the results are summarized in Table 2). CV was performed in acetonitrile (0.1 M Bu₄NPF₆) at a scan rate of 50 mV s⁻¹ at room temperature using an SCE as the reference electrode (all potentials reported are *versus* SCE). A platinum electrode coated with a thin polymer film was used as the working electrode. The onset potentials for oxidation (E_{ox}) are observed to be 0.55 V and 0.54 V for P1 and P2, respectively, and the reduction (E_{red}) of P1 and P2 were found to be -0.75 V and -0.74 V, respectively. With the formula $E_{HOMO} = -e(E_{ox} + 4.29)$ eV and $E_{LUMO} = -e(E_{red} + 4.29)$ eV, we estimated the highest occupied molecular orbital (HOMO) and the lowest unoccupied molecular orbital (LUMO) energy levels of the polymer P1 and P2 to be -4.80 eV, -4.79 eV and -3.54 eV, -3.55 eV, respectively. There are no obvious differences in the HOMO and LUMO levels between polymers

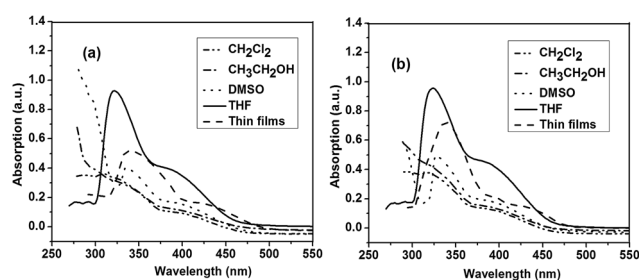


Fig. 1 UV-vis spectra of (a) P1 and (b) P2.

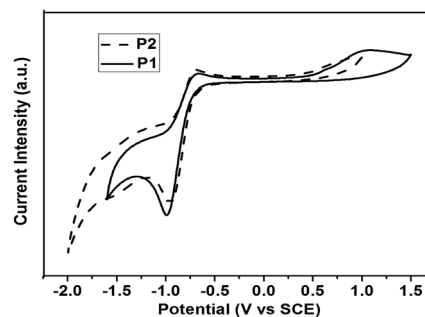


Fig. 3 Cyclic voltammograms of the polymer in 0.1 M Bu₄NPF₆ acetonitrile solution.



Table 2 Optical and electrical properties of the polymers

Polymer	Abs. (nm) ^a	PL (nm) ^a	HOMO ^b (eV)	LUMO ^b (eV)	E _g ^c (eV)
P1	325, 380	410	−4.80	−3.54	1.26
P2	330, 385	420	−4.79	−3.55	1.21

^a THF was used as an the synthesis solvent. ^b $E(\text{HOMO}) = -e(E_{\text{ox}} + 4.29)$ eV, $E(\text{LUMO}) = -e(E_{\text{red}} + 4.29)$ eV, observed in the cyclic voltammetric analyses. ^c $E_g = E(\text{LUMO}) - E(\text{HOMO})$ (eV).

P1 and P2 due to the same main chain structures and the same end groups.

Ion sensing properties

Due to the same main chain of P1 and P2, we chose polymer P1 as a model to investigate its ion sensing property. The results of P2 about ion sensing property were shown in ESI.† The ion sensing properties of polymer P1 was investigated for various metal ions. Variations in the UV-vis absorption and fluorescence spectra of P1 were caused by various competitive metal ions, including Mg^{2+} , Al^{3+} , K^+ , Fe^{3+} , Cu^{2+} , Ba^{2+} , Ni^{2+} , Zn^{2+} , Mn^{2+} , Pb^{2+} , Ag^+ , Hg^{2+} , Ca^{2+} , Li^+ and Cd^{2+} . The measured concentration of P1 in THF was 1.0×10^{-5} M. The influence of different metal ions ($[\text{metal ion}] = 1.0 \times 10^{-4}$ M) on the UV-vis absorption spectra and fluorescence intensity of P1 are shown in Fig. 4a and b. From the UV-vis absorption spectra (Fig. 4a), we can see that P1 showed a significant absorption band centered at about 325 nm upon addition of Fe^{3+} ions, whereas weaker absorption bands were observed with other ions. Emission quench was observed for P1 upon addition of Fe^{3+} and Cu^{2+} , and the quench with Fe^{3+} ions is much more significant compared to that with Cu^{2+} . No fluorescence reduce was observed upon treatment with other metal ions. This result indicates the good selectivity of P1 toward the sensing of Fe^{3+} . According to Swager's group reported, if the absorption peak of UV-vis spectra was red-shifted, the mechanism of quenching is consistent with the static quenching by a charge transfer complex.⁴⁹ For P1 the addition of 1.0×10^{-4} M Fe^{3+} shifts the absorption onset from 325 nm to 334 nm (Fig. 4a). For P2 shows the similar phenomenon from 330 nm to 340 nm (Fig. S12†). Therefore, these results indicate that the fluorescence quenching induced by Fe^{3+} occurs mainly *via* static quenching mechanism owing to the strong association interaction between the Fe^{3+} ions and the receptors,

carboxylic ester groups (the proposal mechanism was shown in Fig. S11†).⁵²

In order to quantitatively measure the effect of metal cations on fluorescence intensity, we calculated the relative fluorescence intensity (I/I_0), in which I_0 is the fluorescence intensity without addition of quencher and I is the fluorescence intensity after the addition of metal cations. The I/I_0 values of P1 and P2 for Mg^{2+} , Al^{3+} , K^+ , Ba^{2+} , Ni^{2+} , Zn^{2+} , Mn^{2+} , Pb^{2+} , Ag^+ , Hg^{2+} , Ca^{2+} , Li^+ and Cd^{2+} are almost 1. However, this value of P1 is 0.72 and 0.36 for Cu^{2+} and Fe^{3+} , respectively; this value of P2 is 0.72 and 0.41 for Cu^{2+} and Fe^{3+} , respectively. Conspicuously, the highest fluorescence quenching effect of both P1 and P2 was observed in the presence of Fe^{3+} ions.

Fig. 5 shows the fluorescence change of P1 upon titration with Fe^{3+} ion ($[\text{Fe}^{3+}] = 0$ to 120×10^{-6} M). With addition of Fe^{3+} ions, the fluorescence intensity of P1 at 425 nm continuously reduced. The corresponding Stern-Volmer constants (K_{sv}) were calculated from the Stern-Volmer relationships (eqn (1)),⁵³ where $[Q]$ is the concentration of the quencher (*i.e.*, Fe^{3+}).

$$I_0/I = 1 + K_{\text{sv}}[Q] \quad (1)$$

At low concentration (0 to 20×10^{-6} M), I_0/I increases linearly with the concentration of quencher. The corresponding Stern-Volmer constant (K_{sv}) were calculated to be $2.90 \times 10^4 \text{ M}^{-1}$ for P1 and $3.83 \times 10^4 \text{ M}^{-1}$ for P2 (shown in ESI S13†). On the basis of these relationships and $3 \times \delta_{\text{blank}}/k$ (where δ_{blank} is the standard deviation of the blank solution and k is the slope of the calibration plot), the detection limit was determined by blank standard method and was calculated to be 6.99×10^{-7} M for P1 and 6.69×10^{-7} M for P2. These numbers indicated that both P1 and P2 were highly sensitive to Fe^{3+} ions.

It is well known that large side groups attached to the polymer backbone will reduce the inter polymer aggregation. We investigated the fluorescent quenching of P1 and P2 in THF solution upon the addition of Fe^{3+} ions. The absorption and emission spectra, and the quantum yield of P1 and P2 in THF solution are very similar, which implied the change of side groups did not have much influence on the main chain of the polymer. However, Fe^{3+} -induced quenching is much less efficient for P1 ($K_{\text{sv}} = 2.90 \times 10^4 \text{ M}^{-1}$) than P2 ($K_{\text{sv}} = 3.83 \times 10^4 \text{ M}^{-1}$), which indicates the bulky side groups indeed stifle interpolymer aggregation and lower the response to metal ions as expected.⁵⁴

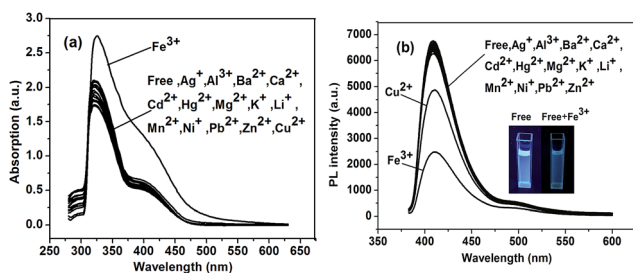


Fig. 4 (a) UV-vis absorption spectra of P1; (b) fluorescence spectra and images of P1.

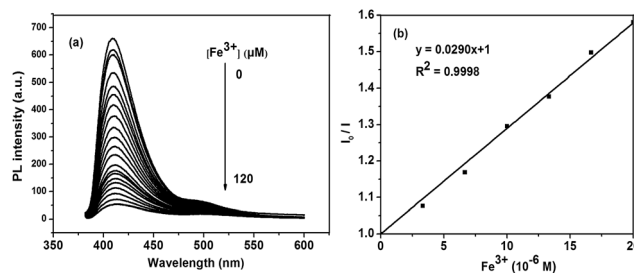


Fig. 5 (a) Fluorescence response of P1 to Fe^{3+} in THF solution; (b) the concentration line chart of P1.



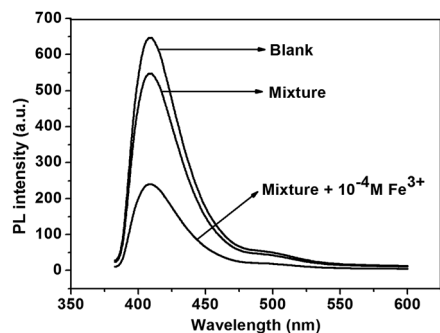


Fig. 6 Fluorescence change of P1 upon titration of metal ion mixtures (concentration of each metal ion was 10^{-4} M) with or without Fe^{3+} in THF solution.

It is reported that the interference from other metal ions will influence the fluorescence quenching assay for a specific metal ion when the sensor is used in complex media. The selectivity of polymers for Fe^{3+} in the presence of excesses various metal ions was investigated (Fig. 6, and S14† for P2). Upon addition of a mixture of metal ions including Mg^{2+} , Al^{3+} , K^+ , Cu^{2+} , Ba^{2+} , Ni^{2+} , Zn^{2+} , Mn^{2+} , Pb^{2+} , Ag^+ , Hg^{2+} , Ca^{2+} , Li^+ and Cd^{2+} (concentration of each metal ion was 1.0×10^{-4} M), the fluorescence intensity of P1 has negligible change. However, when Fe^{3+} (1.0×10^{-4} M) was added to the mixture solution, the fluorescence intensity could be quenched up to 62% of P1 and 57% for P2.

Ion sensing properties of water-soluble polymer P3

Water-soluble hyperbranched conjugated polymer P3 was obtained by hydrolysis of neutral polymer P1 or P2. The structure of P3 was determined by IR (Fig. S9†), showing a strong broad absorption band at 3450 cm^{-1} attributed to the OH stretching vibration of $-\text{COOH}$ group in P3.

Polymer P3 was also examined as a sensor for metal cations. As shown in Fig. 7, dramatic quenching of the emission by Fe^{3+} was observed, while almost no change could be found in the presence of other cations (a small quenching was observed with Cu^{2+}), indicating the good selectivity of P3 for Fe^{3+} in aqueous solution.

Fig. S15† showed the titration of P3 with Fe^{3+} in aqueous solution. The fluorescence intensity at 415 nm exhibited a good linear relationship with the concentration of Fe^{3+} ($0\text{--}40\text{ }\mu\text{M}$).

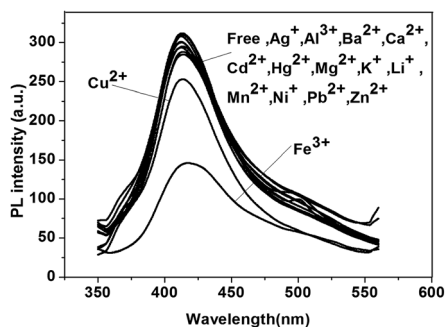


Fig. 7 The fluorescence spectra of P3.

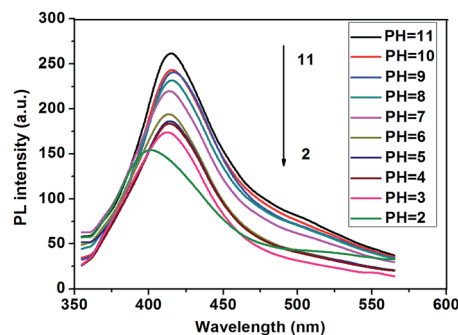


Fig. 8 pH-Dependent emission spectra of P3.

The corresponding Stern–Volmer constants (K_{sv}) was calculated to be $1.17 \times 10^4\text{ M}^{-1}$, and the detection limit was determined to be $5.86 \times 10^{-6}\text{ M}$. According to the results, water-soluble P3 could be considered as a selective detector of Fe^{3+} ion in aqueous environment.

In order to investigate the pH effect on the fluorescent properties of P3, we measured the fluorescence in 10 mM $\text{NaH}_2\text{PO}_4\text{--Na}_2\text{HPO}_4$ buffer solutions with a pH range of 2 to 11. As show in Fig. 8, at basic level ($\text{pH} \geq 7$), P3 exhibits similar emission behavior with the emission maximum located approximately at 415 nm, which is nearly identical to its neutral polymer P1 and P2 in THF. This result indicates that the polymer isolates well in basic solutions. As the pH decreases to 2, a break point is observed where the absorption maximum shifts from 415 to 400 nm. A further decrease in the pH results in significant reduction of the emission intensity of P3. This pH-dependent fluorescence property indicates that P3 may have two different forms under different pH conditions.⁶ At high pH, the majority of the carboxylic acid groups are deprotonated (the anionic form). The electrostatic repulsions between anionic carboxylate groups surrounding the polymer backbones reduce the unfavorable aggregation of the polymer chains. While at low pH, the carboxylate groups can be partially neutralized, leading to decreased charge density and thus a decrease in the inter chain electrostatic repulsion. On one hand, P3 may aggregate through hydrogen bonding between carboxylic acid groups and aromatic $\pi\text{--}\pi$ stacking of hydrophobic backbones. On the other hand, the carboxylic acids make the conjugated main chain twisted and thus decrease the conjugated length. Therefore, at low pH, the emission maxima show blue shift.

Conclusion

In summary, we have synthesized functional hyperbranched polymers containing carboxylic ester groups attached to 1,3-butadiene repeating units. Synthesized polymers P1 and P2 are soluble in common organic solvents and are thermally stable. The neutral hyperbranched conjugated polymers can be hydrolyzed to obtain water-soluble hyperbranched conjugated polymer P3. The fluorescence intensity of these hyperbranched polymers had a good linear response with Fe^{3+} ions over other ions, indicating their potential uses as ratiometric fluorescence chemosensor for Fe^{3+} ions in organic and aqueous solutions.



Acknowledgements

This work was supported by National Science Basic Research Plan in Shaanxi Province of China (2016JM2019) and the National Key Research Project MOST (2016YFA0202400).

Notes and references

- 1 X.-Y. Cao, X.-H. Zhou, H. Zi and J. Pei, *Macromolecules*, 2004, **37**, 8874–8882.
- 2 Y. Xin, G.-A. Wen, W.-J. Zeng, L. Zhao, X.-R. Zhu, Q.-L. Fan, J.-C. Feng, L.-H. Wang, W. Wei, B. Peng, Y. Cao and W. Huang, *Macromolecules*, 2005, **38**, 6755–6758.
- 3 J. Sun, J. Yang, C. Zhang, H. Wang, J. Li, S. Su, H. X. Xu, T. Zhang, Y. Wu, W.-Y. Wong and B. Xu, *New J. Chem.*, 2015, **39**, 5180–5188.
- 4 K. R. Prashanth Kumar, M. G. Murali and D. Udayakumar, *Des. Monomers Polym.*, 2014, **17**, 7–18.
- 5 W. Wu, R. Tang, Q. Li and Z. Li, *Chem. Soc. Rev.*, 2015, **44**, 3997–4022.
- 6 B. Bao, X. Zhan and L. Wang, *J. Polym. Sci., Part A: Polym. Chem.*, 2010, **48**, 3431–3439.
- 7 W. Ding, H. Zhang, J. Xu, Y. Wen, J. Zhang, H. Liu, Y. Yao and Z. Zhang, *J. Polym. Sci., Part A: Polym. Chem.*, 2016, **54**(23), 3694–3700.
- 8 H. Li, X. Wu, Y. Xu, H. Tong and L. Wang, *Polym. Chem.*, 2014, **5**, 5949–5956.
- 9 L. Chen, Y. Gao, Y. Fu, D. Zhu, Q. He, H. Cao and J. Cheng, *RSC Adv.*, 2015, **5**, 29624–29630.
- 10 Y. Xu, X. Wu, Y. Chen, H. Hang, H. Tong and L. Wang, *Polym. Chem.*, 2016, **7**, 4542–4548.
- 11 X.-S. Ma, Y.-Z. Cui, Y.-Q. Ding, F.-R. Tao, B. Zheng, R.-H. Yu and W. Huang, *Sens. Actuators, B*, 2017, **238**, 48–57.
- 12 B. Zheng, Y. Li, F. Tao, Y. Cui and T. Li, *Sens. Actuators, B*, 2017, **241**, 357–363.
- 13 R. Hu, J. W. Y. Lam, J. Liu, H. H. Y. Sung, I. D. Williams, Z. Yue, K. S. Wong, M. M. F. Yuend and B. Z. Tang, *Polym. Chem.*, 2012, **3**, 1481–1489.
- 14 W. Z. Yuan, R. Hu, J. W. Y. Lam, N. Xie, C. K. W. Jim and B. Z. Tang, *Chem.–Eur. J.*, 2012, **18**, 2847–2856.
- 15 D. Ding, G. Wang, J. Liu, K. Li, K.-Y. Pu, Y. Hu, J. C. Y. Ng, B. Z. Tang and B. Liu, *Small*, 2012, **8**(22), 3523–3530.
- 16 H. Ejima, H. Matsuno and T. Serizawa, *Langmuir*, 2010, **26**, 17278–17285.
- 17 D. Wang, T. Zhao, X. Zhu, D. Yan and W. Wang, *Chem. Soc. Rev.*, 2015, **44**, 4023–4071.
- 18 X. Wu, H. Li, Y. Xu, H. Tong and L. Wang, *Polym. Chem.*, 2014, **6**, 2305–2311.
- 19 J. Sun, J. Yang, C. Zhang, H. Wang, J. Li, S. Su, H. Xu, T. Zhang, Y. Wu, W.-Y. Wong and B. Xu, *New J. Chem.*, 2015, **39**, 5180–5188.
- 20 H. T. Nguyen, L.-T. T. Nguyen, T. T. Nguyen, A. T. Luu and T. V. Le, *J. Polym. Res.*, 2014, **21**(552), 2–11.
- 21 D. Li, X. Wang, Y. Jia, A. Wang and Y. Wu, *Chin. J. Chem.*, 2012, **30**, 861–868.
- 22 J. Wang, J. Mei, E. Zhao, Z. Song, A. Qin, J. Z. Sun and B. Z. Tang, *Macromolecules*, 2012, **45**, 7692–7703.
- 23 C. P. Sen, V. D. Goud, R. G. Shrestha, L. K. Shrestha, K. Ariga and S. Valiyaveetil, *Polym. Chem.*, 2016, **7**, 4213–4225.
- 24 J. Qu, M. Shiotsuki, N. Kobayashi, F. Sanda and T. Masuda, *Polymer*, 2007, **48**, 6481–6490.
- 25 J. Oriou, F. Ng, G. Hadzioannou, G. Garbay, M. Bousquet, L. Vignau, E. Cloutet and C. Brochon, *Polym. Chem.*, 2014, **5**, 7100–7108.
- 26 M. R. Raj and S. Anandan, *RSC Adv.*, 2013, **3**, 14595–14608.
- 27 S. Chhatre, S. Agarkar, V. Dhas, S. Nagane, V. Thakare, S. Ogale and P. Wadgaonkar, *J. Mater. Chem.*, 2012, **22**, 23267–23271.
- 28 M. Yoon, H. W. Yoon, A. J. Kim, R. S. Ree, Y.-S. Gal, J. W. Lee, J. H. Lee and S.-H. Jin, *Mol. Cryst. Liq. Cryst.*, 2011, **550**, 304–310.
- 29 B. J. Laughlin and R. C. Smith, *Macromolecules*, 2010, **43**, 3744–3749.
- 30 S. Barik, T. Bletzacker and W. G. Skene, *Macromolecules*, 2012, **45**, 1165–1173.
- 31 S. Barik and W. G. Skene, *Polym. Chem.*, 2011, **2**, 1091–1097.
- 32 S. Dufresne and W. G. Skene, *J. Phys. Org. Chem.*, 2012, **25**, 211–221.
- 33 A. Upadhyay and S. Karpagam, *Polym. Bull.*, 2016, **73**, 2741–2760.
- 34 M. S. Maji, T. Pfeifer and A. Studer, *Chem.–Eur. J.*, 2010, **16**, 5872–5875.
- 35 M. Bourdeaux and W. G. Skene, *Macromolecules*, 2007, **40**, 1792–1795.
- 36 Q. He, F. Bai, J. Yang, H. Lin, H. Huang, G. Yu and Y. Li, *Thin Solid Films*, 2002, **417**, 183–187.
- 37 Y. Sun, Y. Sun, Q. Pan, G. Li, B. Han, D. Zeng, Y. Zhang and H. Cheng, *Chem. Commun.*, 2016, **52**, 3000–3002.
- 38 X.-Y. Cao, W.-B. Zhang, J.-L. Wang, X.-H. Zhou, H. Lu and J. Pei, *J. Am. Chem. Soc.*, 2003, **125**, 12430–12431.
- 39 X. Cai, Y. Liu, T. Lu, R. Yang, C. Luo, Q. Zhang and Y. Chai, *Macromol. Rapid Commun.*, 2016, **37**, 2005–2010.
- 40 J. Y. Cabon, P. Giamarchi and A. L. Bihan, *Anal. Chim. Acta*, 2010, **664**, 114–120.
- 41 B. Muthuraj, S. Hussain and P. K. Iyer, *Polym. Chem.*, 2013, **4**, 5096–5107.
- 42 G. Saikia and P. K. Iyer, *Macromolecules*, 2011, **44**, 3753–3758.
- 43 J. Y. Xu, Y. Zhou, G. F. Cheng, S. X. Liu, M. T. Dong and C. B. Huang, *Luminescence*, 2015, **30**, 451–456.
- 44 C. J. Qin, Y. X. Cheng, L. X. Wang, X. B. Jing and F. S. Wang, *Macromolecules*, 2008, **41**, 7798–7804.
- 45 P. Li, Y. Zhao, L. Yao, H. Nie and M. Zhang, *Sens. Actuators, B*, 2014, **191**, 332–336.
- 46 W. C. Ding, J. K. Xu, Y. P. Wen, H. Zhang and J. J. Zhang, *J. Photochem. Photobiol., A*, 2016, **314**, 22–28.
- 47 X. F. Wu, B. W. Xu, H. Tong and L. X. Wang, *Macromolecules*, 2010, **43**, 8917–8923.
- 48 H. Zhang, G. Zhang, J. K. Xu, Y. P. Wen, B. Y. Lu, J. Zhang and D. W. Ding, *Sens. Actuators, B*, 2016, **230**, 123–129.
- 49 Q. Zhou and T. M. Swager, *J. Am. Chem. Soc.*, 1995, **117**, 7017–7018.
- 50 K. Ono, R. Aizawa, T. Yamano, S. Ito, N. Yasuda, K. Johmoto, H. Uekusa and N. Iwasawa, *Chem. Commun.*, 2014, **50**, 13683–13686.



- 51 D. Nagaraja, R. M. Melavanki, N. R. Patil, H. S. Geethanjali and R. A. Kusanur, *Luminescence*, 2015, **30**, 495–502.
- 52 G. Zhou, Y. Cheng, L. Wang, X. Jing and F. Wang, *Macromolecules*, 2005, **38**, 2148–2153.
- 53 J. R. Lakowicz, *Principles of Fluorescence Spectroscopy*; Kluwer Academic/Plenum Publishers: New York, 1999.
- 54 H. Tong, L. Wang, X. Jing and F. Wang, *Macromolecules*, 2002, **35**, 7169–7171.

