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Synthesis of polymethylene-*b*-poly(vinyl acetate) block copolymer *via* visible light induced radical polymerization and its application†

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New well-defined amphiphilic polymethylene-*b*-poly(vinyl acetate) (PM-*b*-PVAc) diblock copolymers were synthesized *via* a tandem strategy combining polyhomologation of ylides, chain-end functionalization and a visible light induced degenerative iodine transfer polymerization using decacarbonyldimanganese ($Mn_2(CO)_{10}$). Firstly, an iodo terminated polymethylene (PM-I) was prepared by polyhomologation of ylides initiated by a new organic borane based on catecholborane followed by functional group transformation using iodine as the end-capping reagent. Then, a series of amphiphilic PM-*b*-PVAc diblock copolymers with controlled molecular weight ($M_n = 5530-17\ 020\ g\ mol^{-1}$) and relatively narrow molecular weight distribution ($D = 1.31-1.45$) were obtained efficiently through a visible light induced degenerative iodine transfer polymerization of vinyl acetate (VAc) using PM-I as macroinitiator in the presence of $Mn_2(CO)_{10}$ under weak visible light irradiation at 40 °C. The micelles of such an amphiphilic diblock copolymer formed in tetrahydrofuran were observed by transmission electron microscopy (TEM). Moreover, the fabrication of porous films using PM-*b*-PVAc diblock copolymers *via* a static breath-figure process was investigated.

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Introduction

The low cost, excellent physical properties, easy processability and recyclability of polyolefins have led to their commercial success.¹ However, the poor compatibility of polyolefins with other polar materials limits their applications. To overcome this deficiency, the development of polyolefin-based (co)polymeric materials with novel architectures and components is needed.² To explore the highly active and efficient strategies targeting the functionalization of non-polar polyolefins at their main-chain ends or branches was of active interest. By virtue of such functional groups or their derivatives formed by chemical transformation, novel functionalized polyolefin copolymers were synthesized *via* a variety of strategies and had a wide range of applications.¹⁻⁶

Recently, Shea and coworkers⁷ developed polyhomologation of ylides which can be used to build up a carbon backbone polymers one carbon at a time giving rise to a polyethylene

analog named polymethylene. Various main-chain end functionalized polymethylene can be prepared by polyhomologation of ylides using different boranes followed by functional group transformation. Recently, a variety of polymethylene-based copolymers with different architectures and chemical components have been synthesized *via* a combination of polyhomologation of ylides with nitroxide-mediated radical polymerization (NMP),⁸ atom transfer radical polymerization (ATRP),⁹ reversible addition-fragmentation chain transfer polymerization (RAFT),¹⁰ ring-opening polymerization (ROP)¹¹ ring-opening metathesis polymerization (ROMP),¹² living cationic polymerization,¹³ Diels-Alder reaction¹⁴ and “click” chemistry.¹⁵

Light induced polymerization has been widely employed in conventional radical polymerization as a key technique in various applications, such as coatings, adhesives, medicine, optics and microelectronics.¹⁶ The use of light in controlled/living radical polymerization also brings several distinct advantages, including temporal and spatial control over chain growth, easy preparation of well-defined polymers at room temperature and minimization of possible higher temperature side reactions.^{17,18} Although extensive efforts have been made in this area, it is still crucial to explore an effective light induced polymerization in synthetic block polymers.

Among monomers for constructing block copolymer, vinyl acetate (VAc) has gained extensive attention due to the wide applications of poly(vinyl acetate) (PVAc) in different areas, such

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as paints, adhesives, additives to pharmaceuticals and so on.¹⁹ Furthermore, the largest volume water-soluble polymer poly(vinyl alcohol) is also made commercially available by the hydrolysis of PVAc. In contrast to a large variety of the polymerization systems for conjugated vinyl monomers, the controlled/living radical copolymerizations of VAc are difficult because of its low copolymerizability²⁰ and the lack of effective systems for both monomers. Although the reversible addition-fragmentation chain transfer (RAFT) polymerization or macromolecular design *via* interchange of xanthate systems are effective for various monomers including VAc and other conjugated monomers by designing the dithiocarbonyl compounds, there are no effective agents for controlling the copolymerizations.²¹

Koumura and coworkers²⁰ reported the controlled/living radical polymerization of VAc and its copolymerization with methyl acrylate in bulk or fluoroalcohols using dimanganese decacarbonyl [Mn₂(CO)₁₀] in conjunction with an alkyl iodide as an initiator under weak visible light. The dinuclear complex itself is stable and inactive in the dark; in conjunction with halogen containing compounds, it is capable of generating free radicals and initiating polymerization under visible light. In addition, since alkyl halides are also used as coiniciators to generate initiating radicals, so it provides molecular design flexibility in macromolecular synthesis.

Iodo functionalized polyethylene, prepared by the addition of iodine after catalyzed polyethylene chain growth on magnesium, acted as an efficient macroinitiator for the thermal-induced controlled radical polymerization of styrene and methyl methacrylate using Mn₂(CO)₁₀ at 80 °C was reported by Ciftci and coworkers.²² The proposed mechanism involved metal-catalyzed homolysis of the C-I bond of the initiator and dormant polymer chain end, resulting in polymers with controlled molecular weights.

Herein, we'd like to report a combined synthetic strategy targeting the PM-*b*-PVAc diblock copolymers with relatively narrow molecular weight distribution. Firstly, iodo terminated polymethylene (PM-I) was prepared by polyhomologation of dimethylsulfoxonium methylide using a new organic borane based on catecholborane followed by the functional group transformation using iodine as the end-capping reagent. Then, a visible light induced degenerative iodine transfer polymerization of VAc was performed using PM-I as macroinitiator in the presence of Mn₂(CO)₁₀ under weak visible light at 40 °C, obtaining amphiphilic PM-*b*-PVAc diblock copolymers with relatively low molecular weight distribution. The properties of such diblock copolymer such as the micellization and the formation of ordered porous films were also investigated.

Experimental

Materials

All manipulations involving air- and/or moisture-sensitive compounds were carried out in a N₂ filled dry box or using Schlenk techniques. Mn₂(CO)₁₀ (99%, Aldrich) was purified by sublimation and stored in a refrigerator in the dark. Toluene (Sinopharm Chemical Reagent Co., Ltd., ≥99.5%) was refluxed

over sodium and distilled under nitrogen before use. Sodium hydride (Beidouxing Company, Tianjin, 95%) was stirred in *n*-hexane for 6 h and standing for 1 h, then *n*-hexane drew off by a needle, in this way washed three times with *n*-hexane, dried in a vacuum, and stored under nitrogen. Trimethylsulfoxonium iodide (Acros Organics, 98%), benzyltributylammonium chloride (TCI, >98%), dichloromethane (Sinopharm Chemical Reagent Co., Ltd., ≥99.5%) was refluxed over sodium and distilled under nitrogen before use. Styrene (Lingfeng Company, Shanghai, ≥99%) was degassed with nitrogen prior to use. Vinyl acetate (VAc, 99%, Acros) was distilled under reduced pressure and stored under nitrogen. Catecholborane (Aldrich, 1.0 M in THF), iodine (Sinopharm Chemical Reagent Co., Ltd., >99.8%), tetrahydrofuran (Tianlian Company, Shanghai, ≥99%) was refluxed over sodium and distilled under nitrogen before use, chloroform (Shenbo Company, Shanghai, ≥99%).

Synthesis of PM-I

In a typical procedure to obtain a new organic borane, this round-bottom flask was then capped with a rubber septum and deoxygenated by degassing and backfilling with nitrogen for three times, styrene (1.5 equiv., 1.5 mmol) was freeze-dried three times in liquid nitrogen, and then dried toluene (10 mL) were introduced into a nitrogen-purged, 500 mL, round-bottom flask equipped with a stirring bar. Then the solution was dried with nitrogen for 0.5 h. The solution was cooled to 0 °C, adding dropwise of catecholborane solution (1.1 equiv., 1.1 mmol), then was stirred at 80 °C for 3 h. Such reaction mixture was added *via* syringe under nitrogen protection to the solution of dimethylsulfoxonium methylide (72 mmol, synthesized in a similar procedure described in literature²³) to operate the polyhomologation in a similar procedure as described in our previous work²⁴ at 80 °C for 3 h. Cooling to room temperature, an iodine solution (20 mL saturated solution of iodine in dried toluene) was added and the resulting mixture was stirred under the condition of dark for 3 h. The polymer was precipitated in methanol under nitrogen atmosphere and collected by filtration, washed with methanol for several times, and dried under vacuum. PM-I, ¹H NMR (400 MHz, C₆D₄Cl₂, 50 °C) δ: 3.09 (2H, -CH₂-I), 2.60 (2H, C₆H₅-CH₂-), 1.06-1.53 (94H, -CH₂-), 0.67-1.05 (5H, -CH₃) ppm. FT-IR (KBr): 3082, 3059, 3025 (ν, C-H on phenyl), 2849-2918 (ν, -CH₂-), 560 (ν, C-I) cm⁻¹.

Synthesis of block copolymers using VAc monomer with polymethylene macroinitiators

In a typical polymerization, PM-I (0.1 g, 0.1 mmol) and Mn₂(CO)₁₀ (0.02 g, 1.282 mmol) was added to a 100 mL round-bottom flask equipped with a stirring bar. This flask was then capped with a rubber septum and deoxygenated by degassing and backfilling with nitrogen for three times. Then, VAc was placed in another flask equipped with a stirring bar. This flask was capped with a rubber septum, purged with nitrogen for about 3 times, and charged with deoxygenated VAc (5 mL, 54.3 mmol) *via* syringe. The VAc was purged with nitrogen at room temperature for about 30 min, then transfer it into round-



bottom flask with a syringe under a nitrogen atmosphere. The formulation was stirred and exposed to light continuously at 40 °C for several minutes under a blue LED, in which the sample was surrounded by silver paper. At the end of the photo irradiation, the reaction was terminated with liquid nitrogen and the resultant polymer was washed with hexane for three times under nitrogen atmosphere and dried under vacuum then analyzed by ¹H NMR, GPC and FT-IR. ¹H NMR (400 MHz, C₆D₄Cl₂, 50 °C) δ: 5.11 (88H, -CH₂-CH-), 2.06 (264H, -OCCH₃), 1.33 (94H, -CH₂-), 0.93 (5H, -CH₃) ppm. FT-IR (KBr): 3082, 3059, 3025 (ν, C-H on phenyl), 2849–2918 (ν, -CH₂), 1740 (ν, C=O), 1148 (ν, -C(O)-O-C) cm⁻¹.

Formation of honeycomb films *via* the breath figure method

The porous polymer films were fabricated by a static breath-figure process in 50 mL wide-mouth glass bottle with a cap. The glass slides (1.0 × 1.0 cm²) were cleaned sequentially in an ultrasonic bath of acetone, ethanol and deionized water, and dried overnight at 40 °C *in vacuo* finally. Relative humidity (R. H.) in the glass vessel were achieved by adding saturated aqueous solutions of deionized water (R. H. = 95%). A piece of glass slide was placed into the glass vessel, which was 1 cm higher than the liquid level. The PM-*b*-PVAc (run 10 in Table 1) were used to prepare the films. Polymer solutions with different concentrations were prepared by dissolving PM-*b*-PVAc in THF, CHCl₃ and CH₂Cl₂, respectively. A certain amount of solution (10 μL) was dropped and spread on the substrate with a microsyringe in a static humid condition, which was controlled at 20 °C, respectively. The polymer films were dried at room temperature after complete evaporation of solvents.

Instruments and measurements

Blue LED (light-emitting diode, Shenzhen Tengyi, 220 V, 40 W). The ¹H NMR analysis was performed on a Bruker Avance III 400 MHz NMR spectrometer using C₆D₄Cl₂ as the solvent. GPC measurements: molecular weights (*M_w* and *M_n*) and molecular

weight distributions (*D*) were determined using GPC equipped with a Waters 1515 Isocratic HPLC pump, a Waters 2414 refractive index detector, and a set of Waters Styragel columns (HR3, HR4, and HR5, 7.8 × 300 mm²). GPC measurements of polymers were run at 35 °C using THF as eluent with a flow rate of 1.0 mL min⁻¹. The data were calibrated with linear polystyrene standards. FT-IR spectra were recorded on a Nicolet AVATAR-360 FT-IR spectrophotometer with a resolution of 4 cm⁻¹. Transmission Electron Microscope (TEM) observations were performed on a TEM (JEOL JEM-1230) instrument operated at 80 kV. Scanning electron microscopy (SEM) was performed on a SEM (JEOL JSM-6390LV) instrument operated at 10 kV.

Results and discussion

The PM-*b*-PVAc copolymer was synthesized according to a procedure illustrated in Scheme 1. Firstly, PM-I was prepared *via* the strategy of polyhomologation of ylides combining with chain-end functionalization. Then, a series of well-defined PM-*b*-PVAc copolymers were synthesized *via* visible light induced degenerative iodine transfer polymerization of VAc in toluene at 40 °C using Mn₂(CO)₁₀ as initiator and PM-I as a macromolecular chain transfer agent.

Synthesis and characterization of iodo terminated polymethylene (PM-I)

The PM-I were successfully synthesized by polyhomologation of dimethylsulfoxonium methylide following the functional group transformation using iodine as the end-capping reagent. The chain structure of PM-I was confirmed by ¹H NMR, ¹³C NMR and FT-IR spectra.

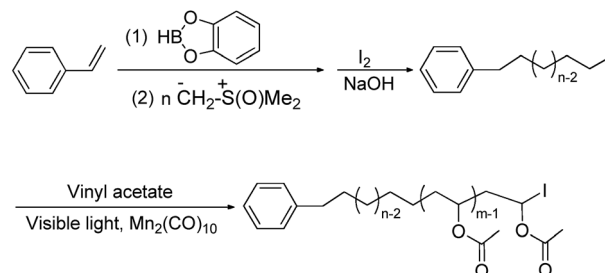
Fig. 1 showed the ¹H NMR spectrum of PM-I. The characteristic resonances of aryl unit in the PM-I can be observed at 7.22–7.08 ppm. The resonance of the methylene group adjacent to the iodine group appeared at 3.09 ppm. And the triplet peak at 2.60 ppm was assigned to methylene protons next to the aryl group, indicating that the halogenated compound based on polymethylene was formed. The signal at 0.93 ppm assigns to the side methyl groups for the existence of dimethylsulfoxonium ethylide impurity in the dimethylsulfoxonium methylide. The integral ratio of peak (a) to (e) is 2 : 2 and the ratios (aromatic protons : peak (a)) is 5 : 2 indicating the formation of

Table 1 Results of PM-*b*-PVAc diblock copolymers prepared *via* visible light induced degenerative iodine transfer polymerization^a

Run	Time [min]	Conv. [%]	<i>M_n</i> , _{theory} ^b [g mol ⁻¹]	<i>M_n</i> , _{NMR} ^c [g mol ⁻¹]	<i>M_n</i> , _{GPC} [g mol ⁻¹]	<i>D</i>
1	10	12.0	5180	4540	5530	1.31
2	20	17.4	6590	6600	6790	1.32
3	30	22.4	8410	8500	8520	1.45
4	35	24.7	9400	9500	9490	1.38
5	40	26.6	10 200	10 100	10 600	1.39
6	50	31.6	11 950	12 000	12 060	1.41
7	60	35.9	13 350	13 600	13 690	1.45
8	90	41.5	15 400	15 700	15 780	1.45
9	150	42.2	16 390	16 000	16 780	1.43
10	240	44.7	16 900	17 000	17 020	1.41

^a Polymerization conditions: PM-I/[VAc]₀/[Mn₂(CO)₁₀] = 1/470/1, *V*_{toluene}/*V*_{VAc} = 1/1.5, samples were irradiated by blue LED light at 40 °C.

^b *M_n*,_{theory} = [VAc]₀/[PM-I]₀ × *M*_{VAc} × conv. ^c The *M_n*,_{NMR} was calculated by ¹H NMR *M_n*,_{NMR} = (integral area of -CH₂-/integral area of -CH₂-I) × *M*_{-CH₂-} + *M*_{-C₆H₅} + *M*_{-I}.



Scheme 1 Synthetic routes for the preparation of well-defined PM-*b*-PVAc diblock copolymers.



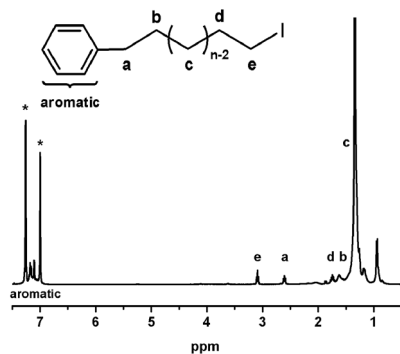


Fig. 1 ^1H NMR spectrum of PM-I.

PM-I. Meanwhile, the integral ratio of peak (d) to peak (c) was 2 : 120, which is agreement with GPC information. All of the information shown in ^1H NMR spectrum of PM-I proved that the functionalization proceeded quantitatively. Furthermore, by ^{13}C NMR analysis, the chemical shifts at 128.32 and 6.33 ppm indicated the presence of carbon atoms of aryl and next to the iodine. It was note worthy that the signals at 142.67 and 33.77 ppm prove the aryl connect to the chain of methylene evidently. In the FT-IR spectrum of PM-I, the peak of C-I stretching vibration absorption had formed at 560 cm^{-1} and the peaks at $3000\text{--}3150\text{ cm}^{-1}$ proved the existence of the benzene ring.

The readily precipitated PM-I chains are simply recovered by filtration and dried. Well-defined PM-I ($\bar{D} = 1.10$) of relatively low molar mass (M_n , GPC = 1000 g mol^{-1}) with high functionality ($\sim 100\%$) was employed as macroinitiator in the following polymerization of VAc.

Synthesis and characterization of PM-*b*-PVAc diblock copolymer

A series of amphiphilic PM-*b*-PVAc diblock copolymers were obtained *via* a visible light induced degenerative iodine transfer polymerization of VAc using PM-I/ $\text{Mn}_2(\text{CO})_{10}$ system under weak visible light at $40\text{ }^\circ\text{C}$.

For the PM-I can solve in the VAc completely, we first have a test for bulk polymerization, the polymerization showed typical radical polymerization behaviour, uncontrolled molecular weight ($M_n = 62\,000\text{ g mol}^{-1}$) and a wide molecular weight distribution ($\bar{D} = 1.75$). The conversion of VAc reached 88.7% after 30 min under visible light at $40\text{ }^\circ\text{C}$.

The polymerization could not take place within 24 h at $40\text{ }^\circ\text{C}$ under light irradiation without $\text{Mn}_2(\text{CO})_{10}$, which could initiate the quick polymerization of VAc. When the polymerization was conducted in toluene with both PM-I and $\text{Mn}_2(\text{CO})_{10}$ under visible light, the obtained polymers showed controlled molecular weight and a narrow molecular weight distribution, which are shown in Table 1.

The GPC traces ($\bar{D} = 1.31\text{--}1.45$) of PM-*b*-PVAc diblock copolymers moved to higher molecular weight in comparison with PM-I indicating the successful chain extension and the formation of PM-*b*-PVAc diblock copolymers. There are no

shoulder and no weight tails in GPC. According to the comparison of the elution time of PM-I and diblock polymers, no self coupling of PM-I was observed.

High reaction temperatures are undesirable for visible light induced degenerative iodine transfer polymerization with alkyl iodides due to the possibility of decomposition of the chain end of iodo-terminated polymers. Because alkyl-I/ $\text{Mn}_2(\text{CO})_{10}$ can decompose at low temperature ($t = 40\text{ }^\circ\text{C}$)²⁵ under visible light, so the polymerization of vinyl acetate can be performed at $40\text{ }^\circ\text{C}$, which should also result in the reduced decomposition rate of the iodo-terminated poly(vinyl acetate).

The system is sensitive to moisture, because traces of water could result in the hydrolysis of C-I end groups,²⁶ leading to the decomposition of the iodo-terminated polymer with the formation of aldehyde end groups and acetyl iodide.²⁷ Increased polarity of the reaction medium may accelerate this decomposition reaction, so toluene was selected as the medium solvent. Because alkyl iodides are UV and light sensitive, all the polymerization reactions were performed in the dark.

Fig. 2 shows the GPC curves of the PM-*b*-PVAc diblock copolymers (a) and the dependence of M_n and \bar{D} on conversion (b) in visible light induced degenerative iodine transfer polymerization of VAc with PM-I/ $\text{Mn}_2(\text{CO})_{10}$ at $40\text{ }^\circ\text{C}$. The M_n increased in direct proportion to the monomer conversion and agreed with the calculated values assuming that one initiator molecule generates one polymer chain. As the polymerization proceeded, the GPC curves shifted to high molecular weights though the \bar{D} s became broader. One of the main reasons for the broadening of the \bar{D} s in the later stages of the polymerization would be the accumulation of the less reactive primary C-I

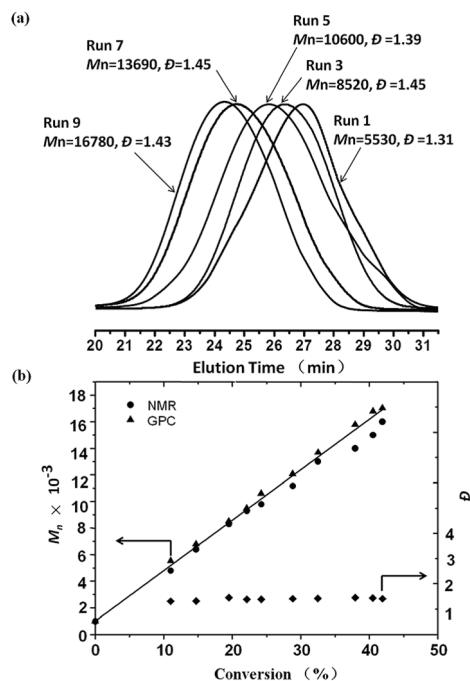


Fig. 2 GPC curves of PM-*b*-PVAc diblock copolymers (a) and dependence of M_n and \bar{D} on conversion (b) in visible light induced degenerative iodine transfer polymerization of VAc.



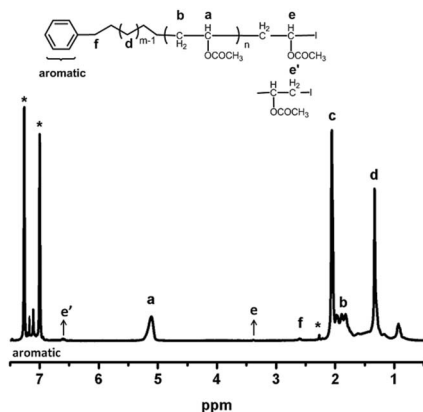


Fig. 3 ^1H NMR spectrum of PM-*b*-PVAc diblock copolymer (run 3 in Table 1).

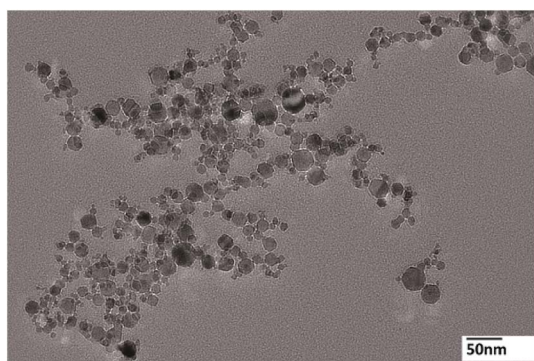


Fig. 4 TEM image of the precipitated aggregates of PM-*b*-PVAc.

terminal originating from the head-to-head addition, which is inherent in VAc radical polymerization.^{25,28,29} These results indicate that $\text{Mn}_2(\text{CO})_{10}$ is effective for the fast and controlled

radical polymerization of VAc under weak light irradiation, which induces slow and continuous generation of the highly active manganese pentacarbonyl radical species [$\text{Mn}(\text{CO})_5$]. To the best of our knowledge, this is the first example of visible light induced degenerative iodine transfer polymerization of VAc at 40 °C with an iodo-terminated polymethylene as the initiator in the presence of $\text{Mn}_2(\text{CO})_{10}$.

As shown in Fig. 3, the main absorption signals of poly(vinyl acetate) were assigned as shown. The methine proton (a) in the polymer backbone absorbs at 5.11 ppm, and the methylene (b) group appears at 1.77–1.93 ppm. Methyl groups in polymer (c) absorb at 2.06 ppm. The signals (e and e') at 3.38 ppm and 6.71 ppm indicates the internal connectivity of the end VAc with head to head and head to tail two different styles. Chain end fidelity is an important issue not only to show the controlled character of the polymerization but also further use of the obtained polymers for chain extension and preparation of block copolymers. The PM-*b*-PVAc was purified and analyzed by ^1H NMR using 1,2-dichlorobenzene- D_4 as the deuterium reagent. The iodide chain-end functionality (82.4%) is calculated on the basis of the ratio of (f) (next to benzene ring) to (e) and (e') (the iodide chain-end functionality = $(I_e + I_{e'}/2)/(I_f/2) \times 100\%$).

Micellization of PM-*b*-PVAc diblock copolymer

The amphiphilic copolymers in a selective solvent can form aggregates with different morphologies such as spheres, vesicles, rod like, ribbon like, and so forth.^{30–32} The amphiphilicity of the resultant PM-*b*-PVAc diblock copolymer was investigated through its micellization in aqueous media. The polymer (run 3) was dissolved in THF to prepare 1 mg mL^{-1} solutions. A solution of PM-*b*-PVAc/THF (0.5 mL), then 5 mL of membrane-filtered water was added with stirring at a flow rate of 0.5 mL min^{-1} . THF was then evaporated for 2 days at 25 °C in

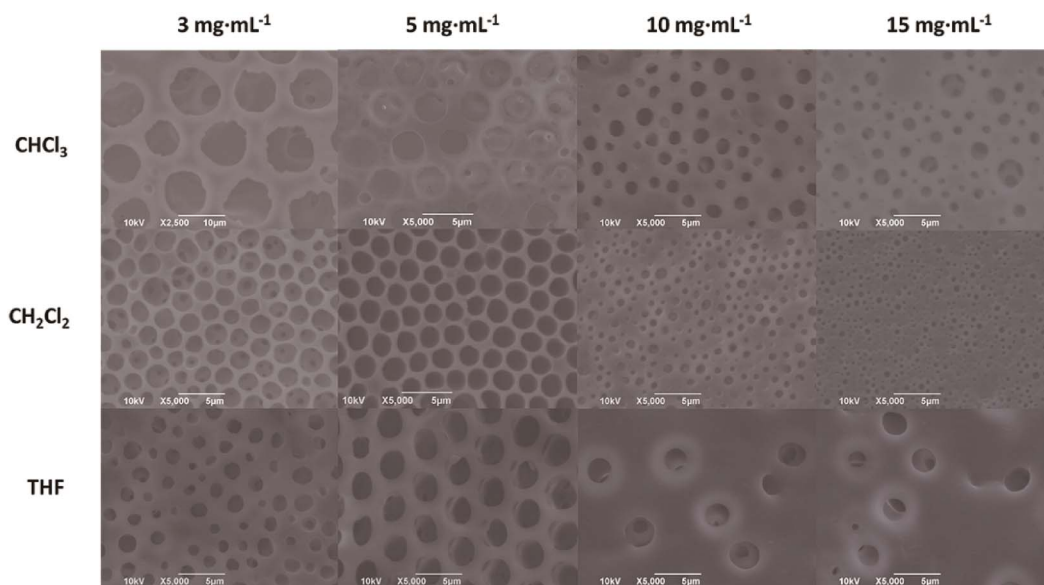


Fig. 5 SEM images of porous films fabricated from PM-*b*-PVAc (run 10 in Table 1) solutions in various solvents at different concentrations.



a water bath. Opaque dispersions of particles were obtained after evaporation.

Then, the aggregates of diblock copolymers deposited on a carbon-coated copper grid were observed by TEM after the complete evaporation of water at room temperature and atmosphere.

As shown in Fig. 4, the amphiphilic PM-*b*-PVAc diblock copolymer aggregated into spheres with diameters of 10–30 nm. Consulted from the study of the model of diblock copolymer micelles,^{33,34} such tendency may be explained by the increasing PVAc corona layer thickness because of the longer PVAc segment in PM-*b*-PVAc.

Fabrication of honeycomb porous PM-*b*-PVAc films

Ordered honeycomb structured films can be prepared by a simple bottom-up process *via* the breath-figure (BF) technique, which takes advantage of the condensation of water droplets on a cold surface.³⁵

The regularity of pores on the surface of film is influenced by various factors in BF process, including polymer structure, polymer molecular weight, polymer concentration, solvent, temperature and humidity. In this work, the effects of solvent and polymer concentration on the surface morphology of PM-*b*-PVAc (run 10, Table 1) film were investigated in the fabricating porous films *via* a static BF process developed by Li and co-workers.³⁶ It was found that PM-*b*-PVAc diblock copolymer can be readily dissolved in chloroform (CHCl₃), dichloromethane (CH₂Cl₂) and tetrahydrofuran (THF). So, the polymer solutions in such three solvents were employed to fabricate porous polymer films in a static BF process at 20 °C under a relative humidity of 95%. All of the PM-*b*-PVAc solutions in THF, CH₂Cl₂ and CHCl₃ can form regular porous films at the concentration of 5 mg mL⁻¹ using the static BF process.

Polymer concentration is an important influencing factor.^{35c,37} The porous films of PM-*b*-PVAc diblock copolymer (run 10 in Table 1) were fabricated at 20 °C under R. H. of 95% with different concentration of 3, 5, 10 and 15 mg mL⁻¹, respectively. SEM images of such films were shown in Fig. 5. The formation of disordered porous film was probably due to the combined effects of lower solution viscosity and lower concentration in 3 mg mL⁻¹ polymer solution, which cannot inhibit the coalescence of water droplets efficiently, resulting in larger pores ($D =$ average pore size, $D_{\text{CHCl}_3} = 9.66 \mu\text{m}$, $D_{\text{CH}_2\text{Cl}_2} = 1.99 \mu\text{m}$, $D_{\text{THF}} = 1.69 \mu\text{m}$) and less regularity. With the polymer concentration rising to 5 mg mL⁻¹, the porous films with ordered pore structures were observed ($D_{\text{CHCl}_3} = 2.79 \mu\text{m}$, $D_{\text{CH}_2\text{Cl}_2} = 2.07 \mu\text{m}$, $D_{\text{THF}} = 2.15 \mu\text{m}$). The porous films with irregular pore structures and smaller pore sizes were formed with the increasing of polymer concentration to 10 mg mL⁻¹ or 15 mg mL⁻¹, respectively. Such results are probably attributed to the weak Marangoni convection and the increased difficulty of water-droplets sinking into the polymer solution with higher viscosity. At the same time, the average pore sizes of films fabricated from the polymer solution in CHCl₃ and CH₂Cl₂ are becoming smaller as $D_{\text{CHCl}_3} = 1.39/1.02 \mu\text{m}$ and $D_{\text{CH}_2\text{Cl}_2} = 0.77/0.41 \mu\text{m}$ at the concentration of 10 mg mL⁻¹/15 mg mL⁻¹, respectively. Being

a better candidate, CH₂Cl₂ will be employed as solvent in the forthcoming systematic investigation on the fabrication of highly ordered porous PM-*b*-VAc films.

Conclusions

New organic borane was prepared by the hydroboration of styrene using catecholborane and initiated *in situ* the polyhomologation of ylides. Subsequently, the post functionalization was carried out directly in the reaction system to achieve terminal -I group forming the iodo terminated polymethylene. The polymerization of vinyl acetate was successfully carried out in toluene using PM-I as macroinitiator in the presence of Mn₂(CO)₁₀ under weak visible light at 40 °C. The amphiphilic PM-*b*-PVAc diblock copolymers with a series of molecular weights and narrow molecular weight distributions were synthesized successfully. The capabilities of such amphiphilic diblock copolymer in the formation of micelles and highly ordered porous films were presented in this work. It is worthy for PM-*b*-PVAc diblock copolymers to be investigated further in the application fields of functional micelles and highly ordered porous films.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- 1 T. C. Chung, *Prog. Polym. Sci.*, 2002, **27**, 39–85.
- 2 M. J. Yanjarappa and S. Sivaram, *Prog. Polym. Sci.*, 2002, **27**, 1347–1398.
- 3 J. Y. Dong and Y. Hu, *Coord. Chem. Rev.*, 2006, **250**, 47–65.
- 4 R. Gody-Lopez, F. D'Agosto and C. Boisson, *Prog. Polym. Sci.*, 2007, **32**, 419–454.
- 5 J. Mazzolini, E. Espinosa, F. D'Agosto and C. Boisson, *Polym. Chem.*, 2010, **1**, 793–800.
- 6 Y. Zhao, L. Wang, A. Xiao and H. Yu, *Prog. Polym. Sci.*, 2010, **35**, 1195–1216.
- 7 (a) K. J. Shea, J. W. Walker, H. Zhu, M. Paz and J. Greaves, *J. Am. Chem. Soc.*, 1997, **119**, 9049–9050; (b) K. J. Shea, *Chem.-Eur. J.*, 2000, **6**, 1113–1119; (c) J. Luo and K. J. Shea, *Acc. Chem. Res.*, 2010, **43**, 1420–1433.
- 8 (a) X. Z. Zhou and K. J. Shea, *Macromolecules*, 2001, **34**, 3111–3114; (b) N. Alkayal, H. Durmaz, U. Tunca and N. Hadjichristidis, *Polym. Chem.*, 2016, **7**, 2986–2991.
- 9 (a) J. Z. Chen, K. Cui, S. Y. Zhang, P. Xie, Q. L. Zhao, J. Huang, L. P. Shi, G. Y. Li and Z. Ma, *Macromol. Rapid Commun.*, 2009, **30**, 532–538; (b) H. C. Lu, Y. Xue, Q. L. Zhao, J. Huang, S. G. Xu, S. K. Cao and Z. Ma, *J. Polym. Sci., Part A: Polym. Chem.*, 2012, **50**, 3641–3647; (c) Y. Xue, H. C. Lu, Q. L. Zhao, J. Huang, S. G. Xu, S. K. Cao and Z. Ma, *Polym.*



- Chem.*, 2013, **4**, 307–312; (d) Z. Zhang, H. Zhang, Y. Gnanou and N. Hadjichristidis, *Chem. Commun.*, 2015, **51**, 9936–9938; (e) Z. Zhang, M. Altaher, H. Zhang, D. Wang and N. Hadjichristidis, *Macromolecules*, 2016, **49**, 2630–2638.
- 10 X. Wang, J. P. Gao, Q. L. Zhao, J. Huang, G. L. Mao, W. Wu, Y. N. Ning and Z. Ma, *J. Polym. Sci., Part A: Polym. Chem.*, 2013, **51**, 2892–2899.
- 11 (a) C. Yuan, H. C. Lu, Q. Z. Li, S. Yang, Q. L. Zhao, J. Huang, L. H. Wei and Z. Ma, *J. Polym. Sci., Part A: Polym. Chem.*, 2012, **50**, 2398–2405; (b) Q. Z. Li, G. Y. Zhang, J. Z. Chen, Q. L. Zhao, H. C. Lu, J. Huang, L. H. Wei, F. D'Agosto, C. Boisson and Z. Ma, *J. Polym. Sci., Part A: Polym. Chem.*, 2011, **49**, 511–517.
- 12 (a) H. Zhang, Y. Gnanou and N. Hadjichristidis, *Polym. Chem.*, 2014, **5**, 6431–6434; (b) H. Zhang, Z. Zhang, Y. Gnanou and N. Hadjichristidis, *Macromolecules*, 2015, **48**, 3556–3562; (c) D. Zhang, H. Zhang, N. Hadjichristidis, Y. Gnanou and X. Feng, *Macromolecules*, 2016, **49**, 2484–2492.
- 13 H. Zhang, S. Banerjee, R. Faust and N. Hadjichristidis, *Polym. Chem.*, 2016, **7**, 1217–1220.
- 14 N. Alkayal and N. Hadjichristidis, *Polym. Chem.*, 2015, **6**, 4921–4926.
- 15 N. Alkayal, H. Durmaz, U. Tunca and N. Hadjichristidis, *Polym. Chem.*, 2016, **7**, 2986–2991.
- 16 (a) B. Aydogan, Y. Yagci, L. Toppare, S. Jockusch and N. J. Turro, *Macromolecules*, 2016, **45**, 7829–7834; (b) S. Dadashi-Silab, S. Doran and Y. Yagci, *Chem. Rev.*, 2016, **116**, 10212–10275; (c) X. Pan, M. A. Tasdelen, J. Laun, T. Junkers, Y. Yagci and K. Matyjaszewski, *Prog. Polym. Sci.*, 2016, **62**, 73–125; (d) M. Chen, M. Zhong and J. A. Johnson, *Chem. Rev.*, 2016, **116**, 10167–10211.
- 17 S. Yamago and Y. Nakamura, *Polymer*, 2013, **54**, 981–994.
- 18 K. Matyjaszewski and N. V. Tsarevsky, *J. Am. Chem. Soc.*, 2014, **136**, 6513–6533.
- 19 (a) K. E. Uhrich, S. M. Cannizzaro, R. S. Langer and K. M. Shakesheff, *Chem. Rev.*, 1999, **99**, 3181–3198; (b) K. Y. Lee and D. J. Mooney, *Chem. Rev.*, 2001, **101**, 1869–1880; (c) H. Lee, R. Mensire, R. E. Cohen and M. F. Rubner, *Macromolecules*, 2012, **45**, 347–355.
- 20 (a) K. Koumura, K. Satoh and M. Kamigaito, *Macromolecules*, 2008, **41**, 7359–7367; (b) K. Koumura, K. Satoh and M. Kamigaito, *Macromolecules*, 2009, **42**, 2497–2504; (c) K. Koumura, K. Satoh and M. Kamigaito, *J. Polym. Sci., Part A: Polym. Chem.*, 2009, **47**, 1343–1353.
- 21 (a) G. Moad, E. Rizzardo and S. H. Thang, *Acc. Chem. Res.*, 2008, **41**, 1133–1142; (b) G. Moad, E. Rizzardo and S. H. Thang, *Polymer*, 2008, **49**, 1079–1131.
- 22 M. Ciftci, S. Norsic, C. Boisson, F. D'Agosto and Y. Yagci, *Macromol. Chem. Phys.*, 2015, **216**, 958–963.
- 23 B. B. Busch, M. M. Paz, K. J. Shea, C. L. Staiger, J. M. Stoddard, J. R. Walker, X. Z. Zhou and H. Zhu, *J. Am. Chem. Soc.*, 2002, **124**, 3636–3646.
- 24 F. Xu, P. Dong, K. Cui, S. Z. Bu, J. Huang, G. Y. Li, T. Jiang and Z. Ma, *RSC Adv.*, 2016, **6**, 69828–69835.
- 25 M. C. Iovu and K. Matyjaszewski, *Macromolecules*, 2003, **36**, 9346–9354.
- 26 B. Boutevin, J. P. Hugon, Y. Pietrasanta and A. Sideris, *Eur. Polym. J.*, 1978, **14**, 353–356.
- 27 E. K. Euranto, A. Noponen and T. Kujanpaa, *Acta Chem. Scand.*, 1966, **20**, 1273–1280.
- 28 M. Wakioka, K. Y. Baek, T. Ando, M. Kamigaito and M. Sawamoto, *Macromolecules*, 2002, **35**, 330–333.
- 29 K. Koumura, K. Satoh, M. Kamigaito and Y. Okamoto, *Macromolecules*, 2006, **39**, 4054–4061.
- 30 I. Astafieva, X. F. Zhong and A. Eisenberg, *Macromolecules*, 1993, **26**, 7339–7352.
- 31 Y. Yu, L. Zhang and A. Eisenberg, *Langmuir*, 1997, **13**, 2578–2581.
- 32 G. Riess, *Prog. Polym. Sci.*, 2003, **28**, 1107–1170.
- 33 G. S. Kwona and K. Kataoka, *Adv. Drug Delivery Rev.*, 1995, **16**, 295–309.
- 34 J. Z. Chen, Q. L. Zhao, H. C. Lu, J. Huang, S. K. Cao and Z. Ma, *J. Polym. Sci., Part A: Polym. Chem.*, 2010, **48**, 1894–1900.
- 35 (a) G. Widawski, M. Rawiso and B. Francois, *Nature*, 1994, **369**, 387–389; (b) U. H. F. Bunz, *Adv. Mater.*, 2006, **18**, 973–989; (c) M. H. Stenzel, C. Barner-Kowollik and T. P. Davis, *J. Polym. Sci., Part A: Polym. Chem.*, 2006, **44**, 2363–2375.
- 36 (a) L. Li, C. K. Chen, A. J. Zhang, X. Y. Liu, K. Cui, J. Huang, Z. Ma and Z. H. Han, *J. Colloid Interface Sci.*, 2009, **331**, 446–452; (b) A. J. Zhang, H. Bai and L. Li, *Chem. Rev.*, 2015, **115**, 9801–9868.
- 37 (a) L. W. Zhu, B. H. Wu, L. S. Wan and Z. K. Xu, *Polym. Chem.*, 2014, **5**, 4311–4320; (b) X. Y. Li, Q. L. Zhao, T. T. Xu, J. Huang, L. H. Wei and Z. Ma, *Eur. Polym. J.*, 2014, **50**, 135–141; (c) J. P. Gao, W. Wu, L. Rong, G. L. Mao, Y. N. Ning, Q. L. Zhao, J. Huang and Z. Ma, *Eur. Polym. J.*, 2014, **59**, 171–179.

