



Cite this: *RSC Adv.*, 2019, 9, 12370

Synthesis of block cationic polyacrylamide precursors using an aqueous RAFT dispersion polymerization†

Bo Huang,^a Jie Jiang,^a Mutian Kang,^a Pingwei Liu,^{*ab} Hailong Sun,^{*c} Bo-Geng Li^a and Wen-Jun Wang^{id} ^{*ab}

Synthesis of cationic polyacrylamides (CPAMs) by introducing cationic polymer precursors followed by chain extension of acrylamide (AM) homopolymer blocks *via* RAFT polymerization is a promising approach for engineering high-performance CPAMs. However, the aqueous solution polymerization of AM usually leads to high viscosity, thus limiting the solid content in the polymerization system. Herein a novel approach is introduced that uses a random copolymer of AM and methacryloxyethyltrimethyl ammonium chloride (DMC) as a macro RAFT chain transfer agent (mCTA) and stabilizer for aqueous RAFT dispersion polymerization of AM. The AM/DMC random copolymers synthesized by RAFT solution polymerization, having narrow dispersities (D_s) at different molecular weights and cationic degrees (C_s), could serve as the mCTA, which was confirmed by mCTA chain extension in aqueous solution polymerization of AM under different C_s , solid contents, AM addition contents, extended PAM block lengths, and mCTA chain lengths. The block CPAMs had a D value of less than 1.2. A model was developed using the method of moments with consideration of the diffusion control effect, for further understanding the chain extension kinetics. Predicted polymerization kinetics provided an accurate fit of the experimental data. The AM/DMC random copolymers were further used for aqueous RAFT dispersion polymerization of AM under different polymerization temperatures, C_s , and mCTA chain lengths. The resulting products had a milky appearance, and the block copolymers had D_s of less than 1.3. Higher C_s and longer chain lengths on mCTAs were beneficial for stabilizing the polymerization systems and produced smaller particle sizes and less particle aggregation. The products remained stable at room temperature storage for more than a month. The results indicate that aqueous RAFT dispersion polymerization using random copolymers of AM and DMC at moderate cationic degrees as a stabilizer and mCTA is a suitable approach for synthesizing CPAM block precursors at an elevated solid content.

Received 11th April 2019
 Accepted 11th April 2019

DOI: 10.1039/c9ra02716e

rsc.li/rsc-advances

Introduction

Water-soluble cationic polyacrylamides, CPAMs, have been widely used in mining, oil recovery, paper manufacturing, dust control, and sewage treatment.¹ The cations in the CPAMs offer good electrostatic interaction properties with colloidal particles in water, which thus make them strong candidates for use as

flocculation aids.¹ Most commercial available CPAMs are random copolymers of acrylamide (AM) and cationic monomers,² for instance, dimethyl diallyl ammonium chloride, acryloxyethyl trimethyl ammonium chloride, and methacryloxyethyltrimethyl ammonium chloride (DMC). In order to achieve good performance, CPAMs usually have very high molecular weights,³ which provide polymer chain bridging among colloidal particles to enhance flocculation efficiency in addition to the charge density.¹ However, such a linear structure at high molecular weight restrains the number of the cations participating in electrostatic interaction with colloidal particles owing to random distribution of the cationic monomer, leading to inefficient use of cationic functionality,⁴⁻⁷ in addition to long dissolution time. Chain structure tuning makes it possible to improve the efficiency of utilizing cationic functionality in the CPAMs. It has been reported that flocculation performance can be enhanced *via* engineering a comb-branched CPAM possessing cationic homopolymer branches.⁴⁻⁷ Star structures^{8,9} have also been observed to promote flocculation. However, even with

^aState Key Lab of Chemical Engineering, College of Chemical and Biological Engineering, Zhejiang University, 38 Zheda Road, Hangzhou 310027, China. E-mail: wenjuncwang@zju.edu.cn; liupingwei@zju.edu.cn

^bInstitute of Zhejiang University – Quzhou, 78 Jiuhua Boulevard North, Quzhou, China 324000

^cState Key Laboratory of Hydraulics and Mountain River Engineering, Sichuan University, 24 South Section 1, Yihuan Road, Chengdu, China 610064. E-mail: sunhl@scu.edu.cn

† Electronic supplementary information (ESI) available: ¹H NMR spectra of AM/DMC random copolymer (M1), and block CPAMs respectively synthesized by aqueous RAFT solution polymerization (SA₂₈₁C₇₀-A₆₀₀-3) and dispersion polymerization (DA₁₇₀C₁₇₇-6-45). See DOI: 10.1039/c9ra02716e



such structures, there is still a fraction of cationic monomers not performing. High charge density in polymerization of cationic monomer leads to difficulty in producing polymers of high molecular weight owing to the positive charge rejection. A copolymer of AM with a moderate cationic monomer content is believed to have the similar electrostatic interaction efficiency to that of a cationic homopolymer, which increases the efficiency of cationic monomer utilization. This can be accomplished through an arm-first strategy^{12–16} by positioning cationic copolymer blocks at branch ends of a hyperbranched AM homopolymer with CPAM precursors followed by chain extension of AM homopolymer block as arm. The resulting CPAMs have high flocculation efficiencies.^{17,18} However, such structures are not possible with conventional radical polymerization techniques, but can be produced using the Reversible Addition–Fragmentation Chain Transfer (RAFT) Polymerization method.^{10,11}

A number of researchers have utilized RAFT polymerization in the generation of PAM. McCormick and co-workers^{19,20} investigated an aqueous RAFT polymerization of AM, while Schork *et al.*²¹ successfully synthesized PAM using an inverse mini-emulsion RAFT polymerization. AM could also be rendered to diblock or random hydrophilic/amphiphilic copolymers *via* RAFT copolymerization, such as block copolymers of AM with ionic liquid monomer²² or with monoacryloxyethyl phosphate,²³ and random copolymers of AM with acrylonitrile.²⁴ The RAFT copolymerization could synthesize multiblock copolymers likewise, such as a ABA triblock copolymer of AM and styrene as a supramolecular elastomer,²⁵ or a triblock of AM, acrylonitrile, and *N,N*-diethylamino ethyl methacrylate as a stimuli responsive material.²⁶ In a dispersed system,²⁷ AM/sodium 2-acrylamido-2-methylpropanesulfonate random copolymer was synthesized *via* RAFT copolymerization. An imprinted AM copolymer was also synthesized using RAFT precipitation polymerization.²⁸ Graft, star, or branch copolymers are also possible with RAFT polymerization.^{29–39} Shan *et al.*⁴⁰ used a multi-arm star RAFT chain transfer agent (CTA) to synthesize star PAM copolymers. A similar approach was used for preparation of hyperbranched copolymers with AM and acrylonitrile.⁴¹ Wang *et al.*^{14,15} synthesized hyperbranched PAMs using a semi-batch strategy by controlling crosslinker addition. The PAM-based nanogels were obtained *via* a RAFT dispersion polymerization.⁴²

Although AM polymers can be synthesized by aqueous RAFT solution polymerization, high viscosities of polymerization systems at high solid contents limits control of the polymerization, and low solid content sacrifices polymerization efficiencies. This low-solid issue can be overcome with aqueous RAFT dispersion polymerization to produce CPAMs.^{43–45} During the dispersion polymerizations, stabilizers including water-soluble polymers, like poly(acryloyloxyethyl trimethylammonium chloride)⁴⁴ and poly(ethylene glycol),⁴⁵ are used to stabilize the polymerization system. We speculate that the copolymerization of cationic monomer with AM at moderate charge densities should provide similar stabilization as a homopolymer of the cationic monomer. In addition, if a RAFT functionality is introduced into the polymer chain, the

copolymer could act as a macro RAFT chain transfer agent (mCTA) in addition to being a stabilizer. The macro RAFT chain transfer agent of cationic monomer with AM is believed to be facile for chain extension of AM, which is difficult to achieve when using the homopolymer of DMA having the terminal RAFT functionality.⁴⁶ An approach of using cationic monomer as a stabilizer and macro RAFT chain transfer agent (as well as a component of final product) for aqueous RAFT dispersion copolymerization is thus developed for synthesizing block CPAMs at elevated solid contents. These block CPAMs are the precursors for further producing star or hyperbranched CPAMs with cationic blocks situated at polymer chain ends.^{47–50}

Model-based polymer chain productions provide precise control over copolymer composition and chain topology distributions.^{51–53} Much attention has been paid to model developments for RAFT polymerization. Vana and co-workers⁵⁴ simulated the polymerization rate and dispersity (\bar{D}) during RAFT polymerization using Monte Carlo simulations, while Tobita^{55,56} and Luo *et al.*⁵⁷ modeled the RAFT mini-emulsion polymerization. Kinetic methods based on elementary reactions in RAFT processes are also commonly used for modeling. Barner-Kowollik and co-workers^{58–60} employed the commercial PREDICI software package to model a RAFT process. Vivaldo Lima *et al.*⁶¹ also used PREDICI to simulate RAFT dispersion polymerizations. By introducing the method of moments, Zhu *et al.*⁶² modeled RAFT polymerization and studied the influence of reaction rate constants. Vivaldo Lima *et al.*^{63–66} compared different RAFT polymerization mechanisms and investigated the effect of the fragmentation rate constants. Zhu and co-workers^{62,67} developed a semi-batch RAFT polymerization model in both homogeneous and heterogeneous systems for regulating the copolymer microstructure. Wang and co-workers^{16,68} prepared a series of hyperbranched polyacrylamide and poly(methyl methacrylate) by using semi-batch RAFT polymerization, and developed a comprehensive understanding of the branching mechanisms.^{16,59}

In this work, we synthesized random co-polymers of AM and DMC with moderate cationic contents *via* an aqueous RAFT solution polymerization. The resulting copolymers could serve as both stabilizers and macro RAFT chain transfer agents for aqueous dispersion polymerization of AM in the presence of ammonium sulfate. The block CPAMs are the precursors for further preparation of star or hyperbranched CPAMs having terminal cationic blocks. The synergy of CPAM and salt is beneficial for road dust suppression.⁶⁹ To develop a better understanding of the mCTA chain extension kinetics with AM, an aqueous RAFT solution polymerization of AM with mCTA was also conducted. A mathematic model was developed and used to predict and guide the synthesis of block CPAM precursors under different experimental conditions, including various extended PAM chain lengths, DMC composition and chain length in mCTAs, and AM addition content. Furthermore, the aqueous dispersion RAFT polymerizations of AM using the mCTA as stabilizer under different polymerization temperatures, charge densities (cationic degrees, C_s), and chain lengths of mCTA were investigated. The block CPAMs were synthesized at elevated solid contents using aqueous RAFT dispersion



polymerization with random copolymer of cationic monomer and AM at moderate charge density as both stabilizer and macro RAFT chain transfer agent.

Experimental

Materials

AM ($\geq 98.5\%$, Shanghai Lingfeng Chemical Reagent) was purified by recrystallization in acetone, and DMC (80% solution in water, Sigma-Aldrich) was washed and precipitated in acetone as well. Initiator 2,2'-azobis[2-(2-imidazolin-2-yl) propane] dihydrochloride (Va-044, $\geq 98\%$, J&K Chemical) and ammonium sulfate (AS, $\geq 99\%$, Sinopharm Chemical Reagent) were used as received. The RAFT chain transfer agent, 3-benzyltrithiocarbonyl propionic acid (BCPA), was synthesized following a procedure from the literature.⁷⁰ Other materials were used as received.

Synthesis of AM/DMC random copolymers

The AM/DMC random copolymers mCTAs were synthesized in acetate acid/sodium acetate buffer solution at pH = 5.0 and 50 °C. BCPA was used as CTA while Va-044 served as the initiator. We take Run M3 as an example. A 250 mL flask equipped with a mechanical stirring mixer was charged with 7.11 g AM (0.1 mol), 25.96 g 80% DMC (0.1 mol), 0.151 g BCPA (0.556 mmol), and 157.9 g buffer solution subsequently. After purging with N₂ for 1 h, 0.090 g Va-044 (0.278 mmol) was introduced to the flask. The polymerization was initiated and kept at 50 °C for 6 h under the N₂ protection. In order to eliminate the residual Va-044, the resulting solutions were kept at 80 °C in sealed bottles for an additional 6 h.

Synthesis of block CPAMs *via* aqueous RAFT solution polymerization

The aqueous RAFT solution polymerization of AM was conducted using AM/DMC random copolymers as mCTA. The block CPAMs were produced *via* chain extension. As an example consider the run designated SA₁₇₀C₁₇₇-A₆₀₀-3. This is a RAFT solution copolymerization run using M3 (PAM₁₇₀-*r*-PDMC₁₇₇) as mCTA, [AM]₀/[mCTA]₀ = 600/1, and 3.0 wt% AM content. A portion of the 0.1 mmol mCTA M3 solution (32.44 g), 4.27 g AM (0.06 mol), and 105.3 g acetate/acid acetate buffer solution were added into a 250 mL flask. After purging with N₂ for 1 h, 0.016 g Va-044 (0.05 mmol) was charged into the flask. The polymerization was carried out at 50 °C for 3 h. Aliquots were taken during polymerization and terminated by adding trace hydroquinone solution.

Synthesis of block CPAMs *via* aqueous RAFT dispersion polymerization

Dispersion polymerizations of AM were carried out using AM/DMC random copolymers as both stabilizer and mCTA in the present of ammonium sulfate AS. The addition of salt to the water could induce block CPAMs to form a polymer-rich phase and become a dispersion system due to the salting-out phenomenon. Consider DA₁₇₀C₁₇₇-6-50 as an example. A

250 mL flask equipped with a mechanical stirring mixer was charged with 31.14 g M3 solution, which contained 0.0934 mmol mCTA, 3.98 g AM (0.056 mol), 18.57 g AS, and 12.64 g deionized water. The polymerization system was purged with N₂ for 1 h, adjusted to pH = 5.5, initiated with 0.0151 g Va-044 (0.0467 mmol), and kept at 50 °C for 3 h under the N₂ protection (Scheme 1).

Characterization

Overall conversions were determined by the brominating titration method,^{71,72} while DMC conversions of mCTAs were determined with ¹H NMR spectra acquired on a Bruker Advance 400 spectrometer with D₂O as solvent. Molecular weights were measured with a Polymer Laboratory PL-GPC 50 gel permeation chromatograph (GPC) equipped with three columns (PL aquagel-OH 50, PL aquagel-OH 40, and PL aquagel-OH 20) and a RI detector. The eluent was 0.4 mol L⁻¹ sodium nitrate and 3 wt% acetonitrile/water solution at a flow rate of 0.8 mL min⁻¹ at 30 °C. The PEO standards ($M_{w1} = 1190 \text{ kg mol}^{-1}$, $D_1 = 1.21$; $M_{w2} = 885.5 \text{ kg mol}^{-1}$, $D_2 = 1.10$; $M_{w3} = 77.35 \text{ kg mol}^{-1}$, $D_3 = 1.05$, where M_w refers to weight-average molecular weight) were used for calibration. Particle sizes of CPAM dispersions were characterized by a dynamic light scattering analysis (DLS) using a Malvern Nanosizer ZS-90. Sample measurements were carried out at the polymerization temperatures. Samples were directly charged into polystyrene cuvettes and sonicated for 90 s to remove bubbles. Particle morphology in dispersions were characterized using a FEI Cryogenic Transmission Electron Microscope (Cryo-TEM, Talos F200c). Apparent viscosities of samples were measured with #27 rotor at 5 rpm and 25 °C using a rotational viscometer (LVDV, Shanghai Fangrui Instrument Co., Ltd).

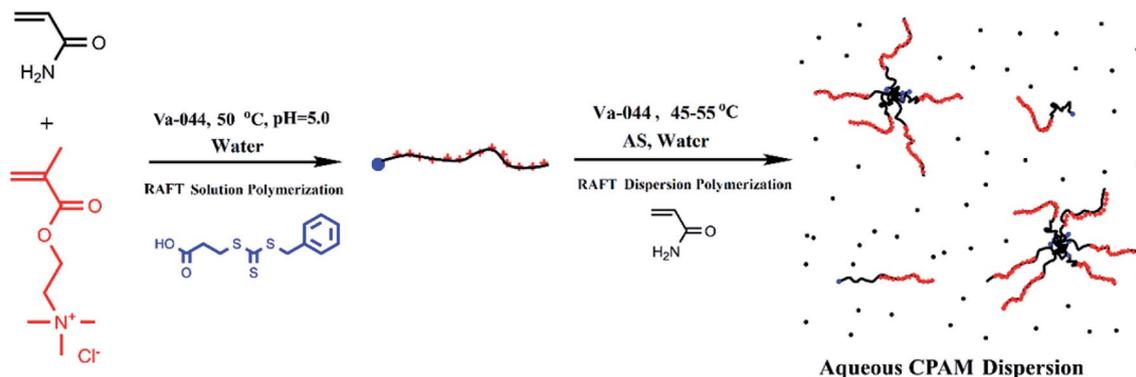
Results & discussion

Aqueous RAFT solution copolymerization of AM and DMC

Six AM/DMC random copolymers were synthesized *via* aqueous RAFT solution polymerization. The experimental conditions and characterization results are summarized in Table 1. It can be seen that all runs reached overall conversions of >95% and dispersities within 1.2 (see Fig. 1). ¹H NMR spectrum of Run M1 is given in Fig. S1 of the ESI. † DMC was more reactive than AM. Number-average molecular weights (M_n s) determined by GPC were close to those obtained from NMR measurements. Well-controlled AM/DMC random copolymer samples were produced.

Targeting the preparation of random copolymers having same chain length of $([AM]_0 + [DMC]_0)/[BCPA]_0 = 360$ and different cationic degrees, 4 runs having C of 20% (Run M1), 40% (Run M2), 50% (Run M3), and 60% (Run M4) were synthesized. Their GPC traces in Fig. 1A are overlapping. They move to slightly higher molecular weight with the increasing cationic degree, indicating similar chain lengths for these copolymers. The copolymers having same cationic degree of 50% and different chain lengths were also prepared. Run M5 had a M_n of 28.8 kg mol⁻¹, while M_n s for Run M3 and M6 were





Scheme 1 Synthesis of block CPAMs via aqueous RAFT dispersion polymerization.

48.8 and 69.2, respectively. The D_s for all three samples were within 1.2.

Aqueous RAFT solution polymerization of AM using mCTA

Ten RAFT solution polymerizations of AM using AM/DMC random copolymers as mCTA were carried out. The experimental conditions and characterization results are summarized in Table 2. ^1H NMR spectrum of Run SA₂₈₁C₇₀-A₆₀₀-3 is shown in Fig. S2 of the ESI.† Run SA₁₇₀C₁₇₇-A₆₀₀-3 was conducted using M3 as mCTA at $[\text{AM}]_0/[\text{M3}]_0/[\text{Va-044}]_0 = 1200/2/1$ and $C = 50\%$. The AM conversion, $-\ln(1 - X_{\text{AM}})$, and GPC trace versus polymerization time and M_w and D versus conversion are given in Fig. 2. The narrow distributed GPC traces progress with polymerization time, indicating successful chain extension of AM/DMC random copolymer with AM. Linear relationships between $-\ln(1 - X_{\text{AM}})$ and t as well as M_w with X_{AM} , and D under 1.1 suggest a controlled characteristic in the polymerization using AM/DMC random copolymer as mCTA.

Influence of mCTA cationic degree on kinetics and acquisition of model parameters

Aqueous RAFT solution polymerizations of AM were conducted under mediation of mCTA having different cationic degrees ranging from 19.9 to 61.7%. The conversions,

molecular weights and distributions are plotted in Fig. 3. It can be seen that the mCTAs with approximately the same M_n but different cationic degrees had little influence on kinetics. The same kinetic parameters can be applied to the RAFT polymerization of AM using mCTAs with different charge densities.

A kinetic model was developed (see Appendix) for the aqueous RAFT solution polymerization of AM under mediation of mCTA. Most of the model parameters can be found from the literature as listed in Table 9 except k_a , k_f , and k_{ct} , which were estimated from experimental data collected from Run SA₂₈₁C₇₀-A₆₀₀-3, SA₂₀₆C₁₄₃-A₆₀₀-3, SA₁₇₀C₁₇₇-A₆₀₀-3, and SA₁₃₃C₂₁₄-A₆₀₀-3.

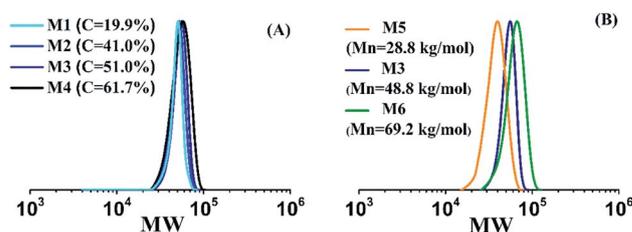


Fig. 1 GPC traces of AM/DMC random copolymer mCTAs having different cationic degrees (A) and chain lengths (B). Experimental conditions: $[\text{BCPA}]_0/[\text{Va-044}]_0 = 2/1$, solid content (SC) = 15.0 wt%, pH = 5.0 (sodium acetate/acid acetate buffer solution), and $T = 50\text{ }^\circ\text{C}$.

Table 1 Experimental conditions and results of synthesis of AM/DMC random copolymers^a

Run	$[\text{AM}]_0/[\text{DMC}]_0^a$	X^b (%)	x_{AM}^b (%)	x_{DMC}^b (%)	Chain structure ^c	C^d (%)	M_n^{NMR} (kg mol^{-1})	$M_n^{\text{GPC},e}$ (kg mol^{-1})	D^e
M1	288/72	97.4	97.6	96.6	PAM _{281-r} -PDMC ₇₀	19.9	34.4	41.1	1.1
M2	216/144	96.8	95.4	99.0	PAM _{206-r} -PDMC ₁₄₃	41.0	43.6	45.7	1.2
M3	180/180	96.5	94.5	98.5	PAM _{170-r} -PDMC ₁₇₇	51.0	48.8	48.6	1.2
M4	144/216	96.4	92.3	99.1	PAM _{133-r} -PDMC ₂₁₄	61.7	53.9	52.6	1.2
M5	108/108	95.4	94.5	96.3	PAM _{102-r} -PDMC ₁₀₄	50.5	28.8	34.1	1.2
M6	252/252	97.9	96.7	99.1	PAM _{244-r} -PDMC ₂₅₀	50.6	69.2	67.7	1.2

^a All runs had $[\text{BCPA}]_0/[\text{Va-044}]_0 = 2/1$, solid content SC = 15.0 wt%, pH = 5.0 (sodium acetate/acid acetate buffer solution), and $T = 50\text{ }^\circ\text{C}$. All $[\text{AM}]_0/[\text{DMC}]_0$ were based on $[\text{BCPA}]_0 = 1$. ^b Overall conversion (X), and AM (x_{AM}) and DMC conversions (x_{DMC}). X_s were determined using the brominate titration method, while x_{DMC} s were determined from ^1H NMR spectra, and x_{AM} s were estimated from both X_s and x_{DMC} s for the copolymerizations. ^c Resulting polymer chain structures are based on the AM and DMC conversions. ^d Cationic degree values were calculated from ^1H NMR spectra. ^e Number-average molecular weights and dispersities were determined by GPC.



Table 2 Experimental conditions and results of aqueous RAFT solution polymerization of AM using mCTAs^a

Run	mCTA	[AM] ₀ /[mCTA] ₀	[AM] ₀ (wt%)	X _{AM} ^b (%)	Chain structure ^c	Mn ^{The,d} (kg mol ⁻¹)	Mn ^{GPC,e} (kg mol ⁻¹)	D ^e
SA ₁₇₀ C ₁₇₇ -A ₃₀₀ -3 ^f	M3	300/1	3.0	55.1	(PAM ₁₇₀ - <i>r</i> -PDMC ₁₇₇)- <i>b</i> -PAM ₁₆₅	60.5	65.0	1.1
SA ₁₇₀ C ₁₇₇ -A ₆₀₀ -3	M3	600/1	3.0	82.5	(PAM ₁₇₀ - <i>r</i> -PDMC ₁₇₇)- <i>b</i> -PAM ₄₉₅	83.9	86.7	1.1
SA ₁₇₀ C ₁₇₇ -A ₁₂₀₀ -3	M3	1200/1	3.0	87.0	(PAM ₁₇₀ - <i>r</i> -PDMC ₁₇₇)- <i>b</i> -PAM ₁₀₄₄	122.9	119.2	1.1
SA ₂₈₁ C ₇₀ -A ₆₀₀ -3	M1	600/1	3.0	83.8	(PAM ₂₈₁ - <i>r</i> -PDMC ₇₀)- <i>b</i> -PAM ₅₀₃	70.1	73.5	1.1
SA ₂₀₆ C ₁₄₃ -A ₆₀₀ -3	M2	600/1	3.0	79.1	(PAM ₂₀₆ - <i>r</i> -PDMC ₁₄₃)- <i>b</i> -PAM ₄₇₅	77.3	73.1	1.1
SA ₁₃₃ C ₂₁₄ -A ₆₀₀ -3	M4	600/1	3.0	80.9	(PAM ₁₃₃ - <i>r</i> -PDMC ₂₁₄)- <i>b</i> -PAM ₄₈₅	88.3	86.9	1.1
SA ₁₀₂ C ₁₀₄ -A ₆₀₀ -3	M5	600/1	3.0	84.9	(PAM ₁₀₂ - <i>r</i> -PDMC ₁₀₄)- <i>b</i> -PAM ₅₀₉	64.9	71.9	1.2
SA ₂₄₄ C ₂₅₀ -A ₆₀₀ -3	M6	600/1	3.0	84.4	(PAM ₂₄₄ - <i>r</i> -PDMC ₂₅₀)- <i>b</i> -PAM ₅₀₆	105.1	102.6	1.2
SA ₁₇₀ C ₁₇₇ -A ₆₀₀ -1.5	M3	600/1	1.5	78.7	(PAM ₁₇₀ - <i>r</i> -PDMC ₁₇₇)- <i>b</i> -PAM ₄₇₂	82.3	89.7	1.1
SA ₁₇₀ C ₁₇₇ -A ₆₀₀ -4.5	M3	600/1	4.5	91.3	(PAM ₁₇₀ - <i>r</i> -PDMC ₁₇₇)- <i>b</i> -PAM ₅₄₈	87.7	93.9	1.1

^a All runs had [mCTA]₀/[Va-044]₀ = 2/1, pH = 5.0 (sodium acetate/acid acetate buffer solution), *T* = 50 °C, and *t* = 2 h. ^b AM conversions were determined by brominate titration method. ^c Resulting polymer chain structures are based on the AM conversions and mCTA chain structures.

^d Theoretical number-average molecular weights are estimated from number-average molecular weights of mCTAs calculated from ¹H NMR spectra and PAM block lengths estimated from AM conversions. ^e Number-average molecular weights and dispersities were determined by GPC.

^f Sample code SA₁₇₀C₁₇₇-A₃₀₀-3 represents a RAFT solution copolymerization run using PAM₁₇₀-*r*-PDMC₁₇₇ as mCTA, [AM]₀/[mCTA]₀ = 300/1, and AM adding content of 3.0 wt%.

Effect of solid content on kinetics and diffusion-controlled termination correction

The aqueous RAFT solution polymerizations of AM at different solid contents were conducted to investigate the diffusion-controlled effect on the kinetics as shown in Fig. 4. Three AM addition contents were used including 1.5 wt% (SA₁₇₀C₁₇₇-A₆₀₀-1.5), 3.0 wt% (SA₁₇₀C₁₇₇-A₆₀₀-3), and 4.5 wt% (SA₁₇₀C₁₇₇-A₆₀₀-4.5), corresponding to overall solid contents of 3.2, 6.4, and 9.6 wt%, respectively. The conversions increased slightly at the same polymerization times when changing the AM addition

contents from 1.5 to 3.0 wt%, while the conversions went up substantially with the further increment of the AM addition contents from 3.0 to 4.5 wt%, suggesting diffusion controlled reactions. RAFT polymerization of AM is quite rapid. Even at a low conversions, diffusion control occurs much earlier. This is a result of higher radical concentrations due to lower radical addition rates to CTA and reduced radical termination rates.⁷⁵ However, this effect had little influence on the polymerization control. Run SA₁₇₀C₁₇₇-A₆₀₀-4.5 samples still possessed narrow dispersities with linear increments of molecular weight *versus* polymerization times, which implies good control should be

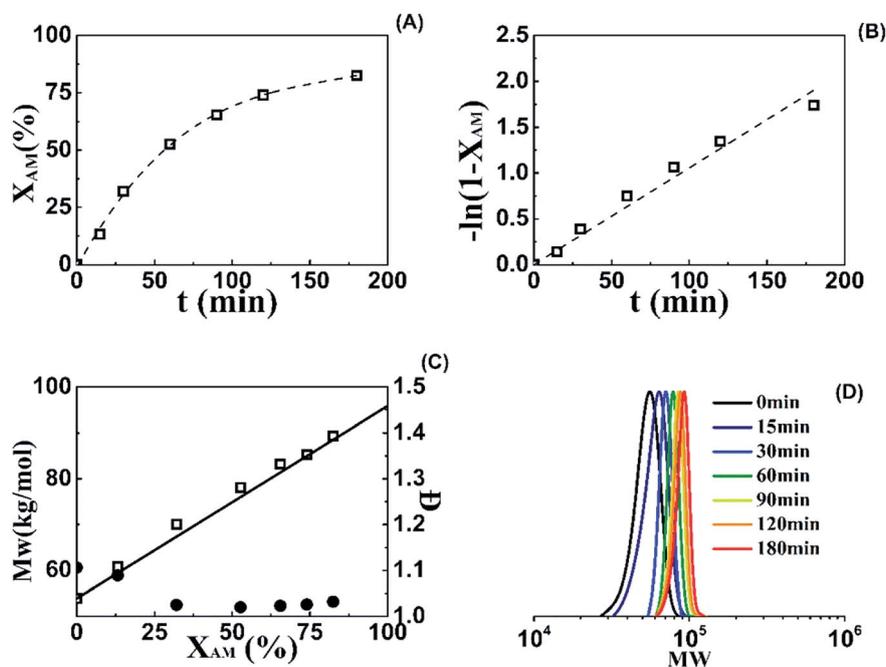


Fig. 2 Plots of characterizing the aqueous RAFT solution polymerization of AM using M3 as mCTA including (A) AM conversion *versus* polymerization time, (B) $-\ln(1 - X_{AM})$ *versus* polymerization time, (C) weight-average molecular weight and dispersity *versus* AM conversion, and (D) GPC trace of polymers at different polymerization times. Experimental conditions: [AM]₀/[M3]₀/[Va-044]₀ = 1200/2/1, [AM]₀ = 0.423 M, pH = 5.0, sodium acetate/acetate acid as buffer solution, and *t* = 50 °C.



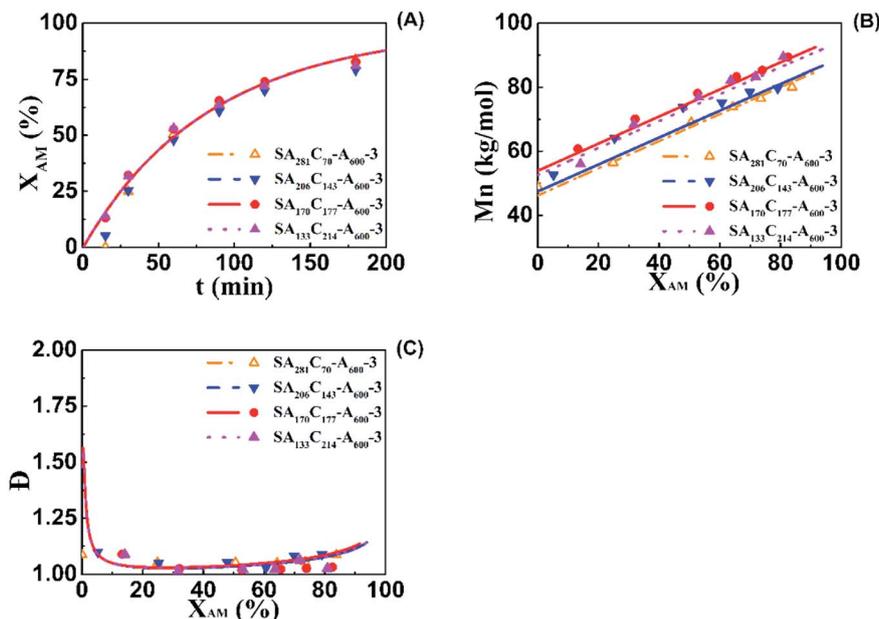


Fig. 3 Plots of characterizing the aqueous RAFT solution polymerization of AM with different mCTA cationic degrees including (A) AM conversion versus polymerization time, (B) number-average molecular weight versus AM conversion, and (C) dispersity versus AM conversion. Experimental conditions: $[AM]_0/[mCTA]_0/[Va-044]_0 = 1200/2/1$, $[AM]_0 = 0.423$ M, pH = 5.0, sodium acetate/acetate acid as buffer solution, and $t = 50$ °C. The points are experimental data while lines are model simulation results.

achievable in polymerization systems at high solid contents, such as aqueous dispersion systems.

To describe the diffusion-controlled effect, semi-empirical eqn (A5)–(A7) were used to quantify the termination constant

with the consideration of contribution of number-average chain length (\bar{r}_N) and free volume fraction (v_f).^{77–79} When the solid content increases, the free volume fraction v_f decreases due to the existence of more polymer chains in the same volume. The

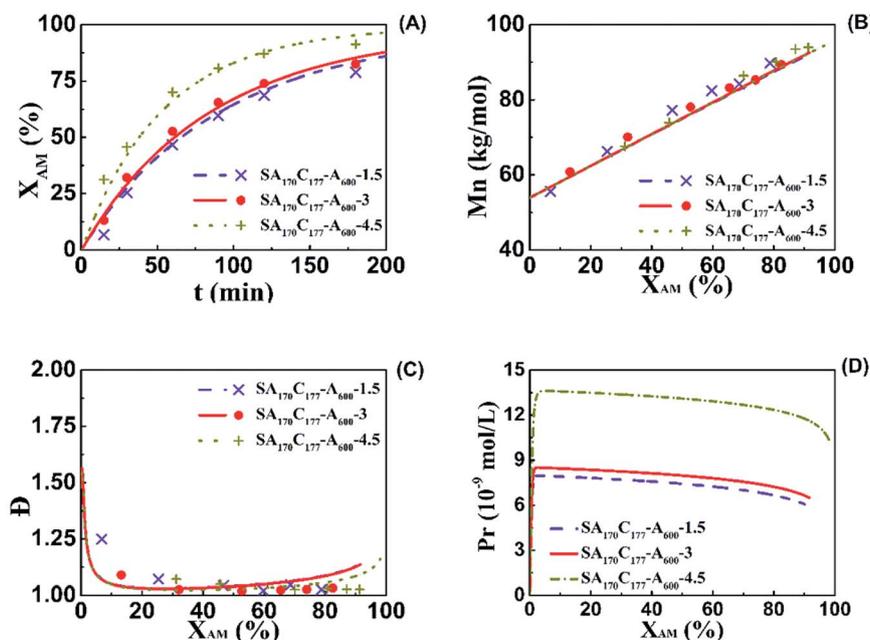


Fig. 4 Plots of characterizing the aqueous RAFT solution polymerization of AM with different AM addition contents including (A) AM conversion versus polymerization time, (B) number-average molecular weight versus AM conversion, (C) dispersity versus AM conversion, and (D) propagation radical (Pr) concentrations estimated by the model. Experimental conditions: $[AM]_0/[M3]_0/[Va-044]_0 = 1200/2/1$, pH = 5.0, sodium acetate/acetate acid as buffer solution, and $t = 50$ °C. The points are experimental data while lines are model simulation results.



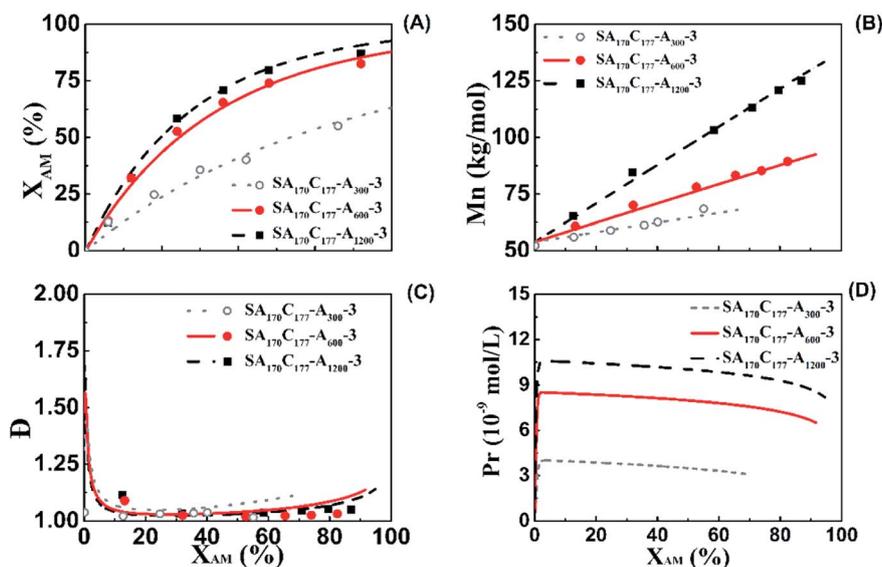


Fig. 5 Plots of summarizing model-predicted and experimental polymerization kinetics results for aqueous RAFT solution polymerization of AM at different extended PAM block lengths including (A) AM conversion versus polymerization time, (B) number-average molecular weight versus AM conversion, (C) dispersity versus AM conversion, and (D) propagation radical concentration (Pr) values estimated with the model. Experimental conditions: $[M3]_0/[Va-044]_0 = 2/1$, $[M3]_0 = 0.653$ mM, pH = 5.0, sodium acetate/acetate acid as buffer solution, and $t = 50$ °C. The lines are model-predicted results while the points are experimental data.

parameters of $k_{t,D}^0$, a , and b in eqn (A6) were estimated from the experimental data of Run $SA_{170}C_{177}A_{600}-1.5$, $SA_{170}C_{177}A_{600}-3$ and $SA_{170}C_{177}A_{600}-4.5$, and listed in Table 9.

Measured viscosities of CPAM samples at 30 min with solids contents ranging from 1.5 to 4.5 wt% were 5.0 mPa s ($X = 25.4\%$, $SA_{170}C_{177}A_{600}-1.5$), 14.2 mPa s ($X = 32.1\%$, $SA_{170}C_{177}A_{600}-3$), and 54.7 mPa s ($X = 45.7\%$, $SA_{170}C_{177}A_{600}-4.5$).

Corresponding propagation radical concentration (Pr) values estimated from the model were 7.76×10^{-9} , 8.24×10^{-9} , and 13.2×10^{-9} mol L $^{-1}$, respectively, as shown in Fig. 4D. Even for conversions as low as 32%, the viscosity and Pr values were 5.8, 14.2 and 24.6 mPa s and 7.69×10^{-9} , 8.24×10^{-9} and 13.4×10^{-9} mol L $^{-1}$, respectively. This indicates that the increase in

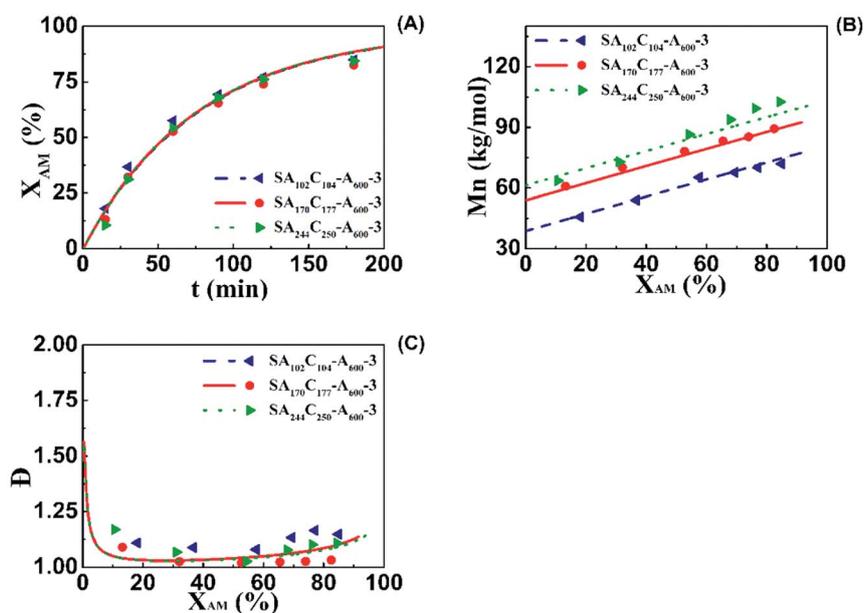


Fig. 6 Model-predicted and experimental polymerization kinetics for aqueous RAFT solution polymerization of AM using mCTAs of various chain lengths. Plots show (A) AM conversion versus polymerization time, (B) number-average molecular weight versus AM conversion, and (C) dispersity versus AM conversion. Experimental conditions: $[AM]_0/[mCTA]_0/[Va-044]_0 = 1200/2/1$, $[AM]_0 = 0.423$ mM, pH = 5.0, sodium acetate/acetate acid as buffer solution, and $t = 50$ °C. The lines are model-predicted results while the points are experimental data.



Table 3 Experimental conditions and results of aqueous RAFT dispersion polymerization of AM using mCTAs^a

Run	mCTA	T (°C)	[AM] ₀ (wt%)	X _{AM} ^b (%)	Chain structure ^c	Mn ^{The,d} (kg mol ⁻¹)	Mn ^{GPC,e} (kg mol ⁻¹)	D ^e
DA ₁₇₀ C ₁₇₇ -6-45 ^g	M3	45	6.0	57.4	(PAM ₁₇₀ - <i>r</i> -PDMC ₁₇₇)- <i>b</i> -PAM ₃₄₅	73.3	71.4	1.2
DA ₁₇₀ C ₁₇₇ -6-50	M3	50	6.0	94.0	(PAM ₁₇₀ - <i>r</i> -PDMC ₁₇₇)- <i>b</i> -PAM ₅₆₄	88.9	90.7	1.2
DA ₁₇₀ C ₁₇₇ -6-55	M3	55	6.0	96.5	(PAM ₁₇₀ - <i>r</i> -PDMC ₁₇₇)- <i>b</i> -PAM ₅₇₉	89.9	87.8	1.3
DA ₂₀₆ C ₁₄₃ -6-50	M2	50	6.0	91.6	(PAM ₂₀₆ - <i>r</i> -PDMC ₁₄₃)- <i>b</i> -PAM ₅₅₀	82.7	82.7	1.2
DA ₁₈₉ C ₁₅₈ -6-50	M7 ^f	50	6.0	92.8	(PAM ₁₈₉ - <i>r</i> -PDMC ₁₅₈)- <i>b</i> -PAM ₅₅₇	85.8	90.9	1.1
DA ₁₀₂ C ₁₀₄ -4.2-50	M5	50	4.2	68.6	(PAM ₁₀₂ - <i>r</i> -PDMC ₁₀₄)- <i>b</i> -PAM ₄₁₂	58.1	67.0	1.1
DA ₁₇₀ C ₁₇₇ -4.2-50	M3	50	4.2	85.0	(PAM ₁₇₀ - <i>r</i> -PDMC ₁₇₇)- <i>b</i> -PAM ₅₁₀	85.0	82.5	1.1
DA ₂₄₄ C ₂₅₀ -4.2-50	M6	50	4.2	94.2	(PAM ₂₄₄ - <i>r</i> -PDMC ₂₅₀)- <i>b</i> -PAM ₅₆₅	109.3	104.6	1.2

^a All runs had [AM]₀/[mCTA]₀/[Va-044]₀ = 1200/2/1, AS = 28 wt%, and *t* = 2 h. ^b AM conversions were determined by brominate titration method.

^c Resulting polymer chain structures based on the AM conversions and mCTA chain structures. ^d Theoretical number-average molecular weights estimated from number-average molecular weights of mCTAs calculated from ¹H NMR spectra and PAM block lengths estimated from AM conversions. ^e Number-average molecular weights and dispersities were determined by GPC. ^f An AM/DMC random copolymer with Mn^{GPC} = 46.3 kg mol⁻¹, D = 1.2, and C = 45.5% produced by aqueous RAFT solution copolymerization. ^g Sample code DA₁₇₀C₁₇₇-6-45 represents a RAFT dispersion copolymerization run using PAM₁₇₀-*r*-PDMC₁₇₇ as mCTA, 6.0 wt% AM content and *T* = 45 °C.

propagation radical concentrations resulted in greater AM conversion.

Model-predicted and experimental polymerization kinetics

The effect of the extended PAM block length on aqueous RAFT solution polymerization using M3 as mCTA was predicted using the model and validated with experimental data, as shown in Fig. 5. The ratios of [AM]₀/[M3]₀ were 300/1 (SA₁₇₀C₁₇₇-A₃₀₀-3), 600/1 (SA₁₇₀C₁₇₇-A₆₀₀-3), and 1200/1 (SA₁₇₀C₁₇₇-A₁₂₀₀-3). It can be seen that the model predictions are in good agreement with the experimental data. Run SA₁₇₀C₁₇₇-A₃₀₀-3 had the lowest

polymerization rate as a result of having the shortest CPAM length. With the increase of the extended PAM block length from Run SA₁₇₀C₁₇₇-A₃₀₀-3 to SA₁₇₀C₁₇₇-A₁₂₀₀-3, the polymerization rate increased while maintaining good control characteristics.

Viscosities of CPAM samples at 60 min having different extended PAM blocks ranging from 300 to 1200 units were 10.7 mPa s (*X* = 26.1%, SA₁₇₀C₁₇₇-A₃₀₀-3), 23.0 mPa s (*X* = 52.4%, SA₁₇₀C₁₇₇-A₆₀₀-3), and 61.7 mPa s (*X* = 59.7%, SA₁₇₀C₁₇₇-A₁₂₀₀-3). The corresponding Prs estimated from the model were 3.99 × 10⁻⁹, 7.95 × 10⁻⁹, and 9.87 × 10⁻⁹ mol L⁻¹, respectively.

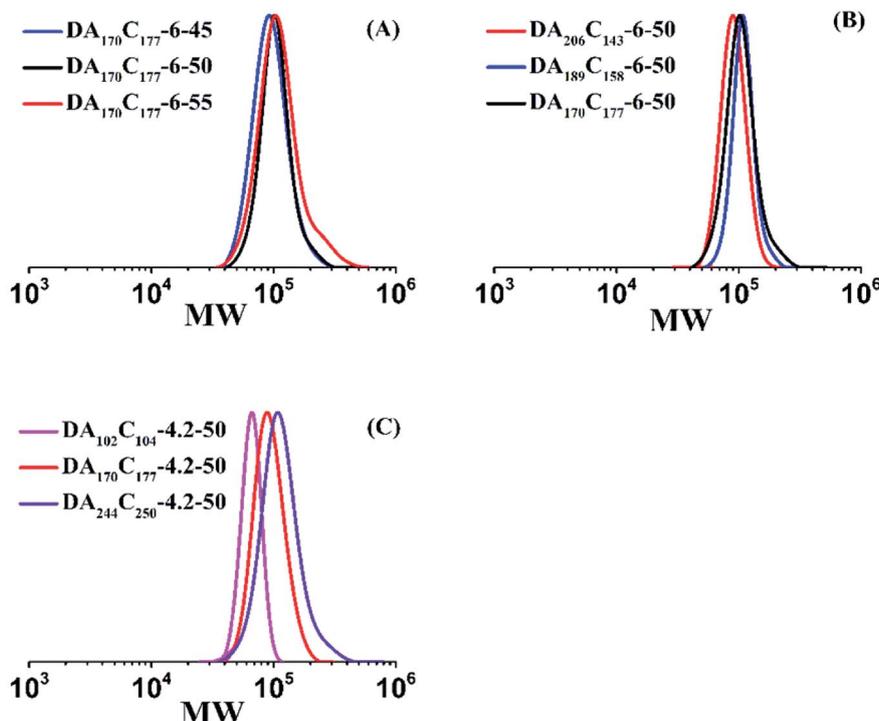


Fig. 7 GPC traces of block CPAM dispersion samples under various experimental conditions including (A) polymerization temperature, (B) cationic degree on mCTA having the same chain length, and (C) molecular weight of mCTA having the same *C* of 51%.



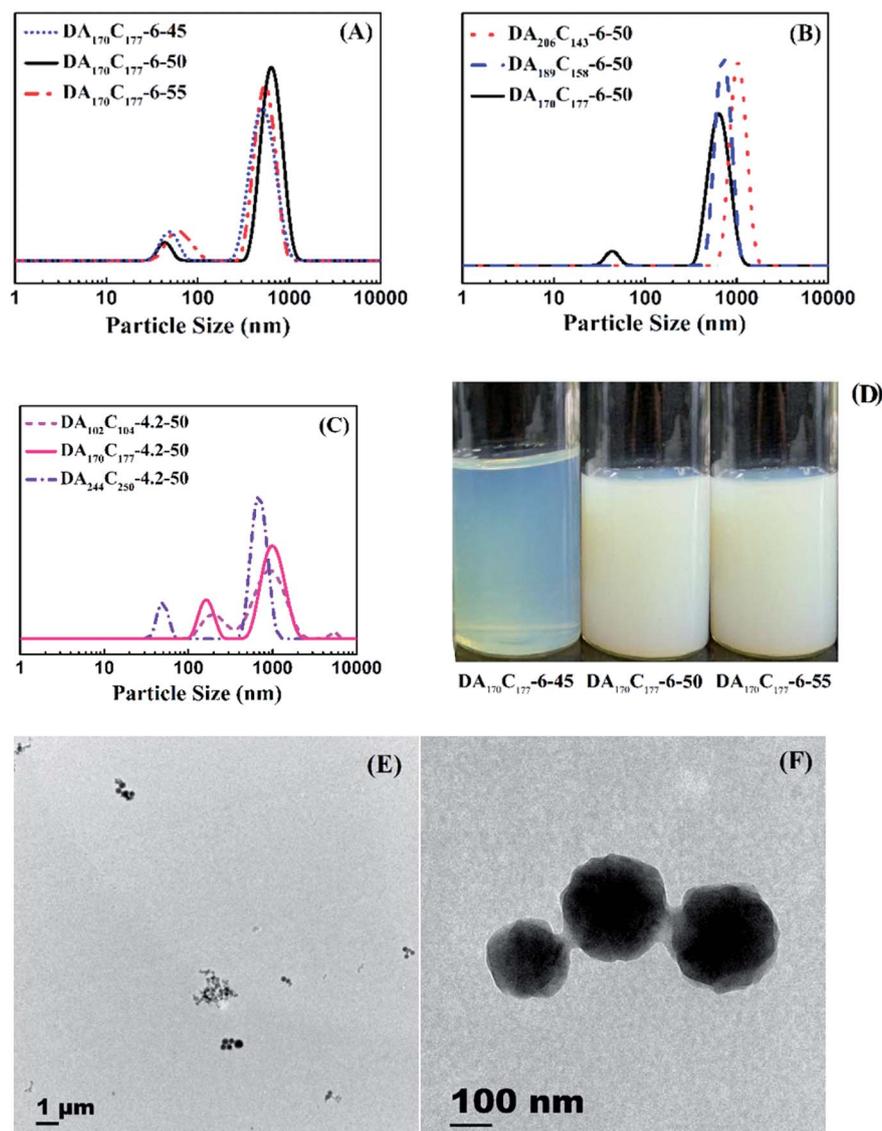


Fig. 8 Appearance of block CPAM dispersion samples and their intensity-based particle size distributions. The samples were synthesized using aqueous RAFT dispersion polymerization under various experimental conditions including a range of (A) polymerization temperatures, (B) cationic degrees on mCTA having the same chain lengths, and (C) molecular weights of mCTA having the same C_s of 51%. All the samples were measured at the same temperature as their polymerizations. Also shown are (D) images of Run $DA_{170}C_{177}$ -6-45, $DA_{170}C_{177}$ -6-50 and $DA_{170}C_{177}$ -6-55, and Cryo-TEM images of particles from Run $DA_{102}C_{104}$ -4.2-50 under (E) low and (F) high magnification.

Viscosities and Prs of the samples at 32% conversion were 12.7, 14.2, and 23.7 mPa s and 3.74×10^{-9} , 8.24×10^{-9} , and $10.3 \times 10^{-9} \text{ mol L}^{-1}$, respectively, indicating that longer CPAM chain lengths produce higher solution viscosity and enhanced the diffusion control, promoting propagation.

The influence of mCTA chain length on aqueous RAFT AM solution polymerization was also examined by model prediction and experiment. Three runs having the same C of 51% but different number-average molecular weights ranging from 28.8 ($SA_{102}C_{104}$ -A₆₀₀-3), 48.8 ($SA_{170}C_{177}$ -A₆₀₀-3) to 69.2 kg mol^{-1} ($SA_{244}C_{250}$ -A₆₀₀-3). Both model-predicted and experimental conversions, molecular weights and distributions are plotted in Fig. 6. Good agreement between the model prediction and experiment further confirm the reliability of the model. The

trend for conversions *versus* polymerization times in three runs is almost overlapping, suggesting the chain length of mCTA had little influence on the kinetics. The narrow dispersities for the three runs indicate good polymerization control for the mCTAs at different chain lengths.

Aqueous RAFT dispersion polymerization of AM using mCTA

The AM/DMC random copolymers were further used as mCTA and stabilizer for preparing block CPAMs *via* aqueous RAFT dispersion polymerization of AM. Eight runs were polymerized at a ratio of $[AM]_0/[mCTA]_0/[Va-044]_0 = 1200/2/1$ and 28 wt% of ammonium sulfate for 2 h. Various experimental conditions including polymerization temperatures, mCTA cationic degrees



and chain lengths were studied as shown in Table 3. ^1H NMR spectrum of Run DA₁₇₀C₁₇₇-6-45 is given in Fig. S3 of the ESI.† All runs had AM conversions between 57.4–96.5%, molecular weights close to the theoretical values, and dispersities within 1.3 (see GPC traces given in Fig. 7), suggesting good control of mCTA chain extension. During polymerizations, phase separations occurred, and dispersion systems were formed. The products had a milky appearance as shown in Fig. 8. The particle sizes of the samples were determined by DLS. Their Z-average particle size distributions are provided in Fig. 8. Most samples except Run DA₂₀₆C₁₄₃-6-50 and DA₁₈₉C₁₅₈-6-50 had two peaks with one peak at approximately 20–200 nm contributed by separated phase particles, and another peak approximately 1000 nm attributed to the aggregation of particles. Particle morphology of Run DA₁₀₂C₁₀₄-4.2-50 was further characterized with Cryo-TEM. TEM images are shown in Fig. 8E and F. The particles with approximately 100 nm and the particle aggregates having a size of approximately 1 μm can be observed, which is consistent with the DLS results and confirms particle aggregation.

The products were stable and could be stored at room temperature for more than one month. Moreover, the solid content in the dispersion system went up to 12.9 wt% (Run DA₁₇₀C₁₇₇-6-50) with an apparent viscosity of 23.4 mPa s, in comparison to the solution system having similar chain structure and length at a solid content of 6.4 wt% (Run SA₁₇₀C₁₇₇-A₆₀₀-3) with an apparent viscosity of 108.5 mPa s. We also synthesized a solution sample having the same solid content (12.9 wt%), chain structure and length as Run DA₁₇₀C₁₇₇-6-50. This sample had a viscosity of 750.2 mPa s, indicating that viscosity can be limited and suggesting that CPAM can be prepared at a high solids content using dispersion polymerization.

The aqueous RAFT dispersion polymerizations of AM using mCTA were carried out at 45 °C (DA₁₇₀C₁₇₇-6-45), 50 °C (DA₁₇₀C₁₇₇-6-50), and 55 °C (DA₁₇₀C₁₇₇-6-55). The higher temperature accelerated polymerizations. The AM conversion went up, from 57.4% at 45 °C to 96.5% at 55 °C, and \bar{D} rose slightly. However, polymerization temperature had little influence on the particle size distributions of the samples. All three samples had two peaks with one approximately 20–100 nm and another around 1000 nm.

Table 4 Elementary reactions in aqueous RAFT solution polymerization

Elementary reaction	Scheme
Initiation	$I \xrightarrow{f, k_{\text{td}}} 2P_0'$ $P_0' + M \xrightarrow{k_p} P_1'$
Propagation	$P_r' + M \xrightarrow{k_p} P_{r+1}' \quad r = 1, 2, \dots$
Pre-equilibrium	$P_r' + TP_0 \xrightleftharpoons{k_a/k_t} P_r \dot{T} P_0 \xrightleftharpoons{k_{a0}/k_{t0}} P_0' + TP_r$
Core-equilibrium	$P_r' + TP_s \xrightleftharpoons{k_a/k_t} P_r \dot{T} P_s \xrightleftharpoons{k_a/k_t} P_s' + TP_r$
Termination	$P_r' + P_s' \xrightarrow{k_{tc}} P_{s+r} P_r' + P_s' \xrightarrow{k_{td}} P_r + P_s P_r' + P_s \dot{T} P_r \xrightarrow{k_{ct}} P_{r+s+1}'$

Table 5 Definition of moments for each chain species

Type of chains	Definition of Moments
Propagating radical	$Y_m = \sum_{r=2}^{\infty} r^m [P_r']$
Dormant	$Q_m^T = \sum_{r=2}^{\infty} r^m [TP_r]$
Intermediate radical	$Y_m = \frac{1}{2} \sum_{r=2}^{\infty} r^m \sum_{s=2}^{r-2} [P_s \dot{T} P_{r-s}]$ $Y_{m,n}^T = \sum_{r=2}^{\infty} r^m s^n \sum_{s=2}^{r-2} [P_r \dot{T} P_s]$
Dead	$Q_m = \sum_{r=2}^{\infty} r^m [P_r]$

Three mCTAs having the same chain lengths of 348 but different C_s of 41.0% (DA₂₀₆C₁₄₃-6-50), 45.5% (DA₁₈₉C₁₅₈-6-50), and 51.0% (DA₁₇₀C₁₇₇-6-50) were used for the dispersion polymerization. In RAFT solution polymerization, the cationic degree did not affect the kinetics, which was also found in RAFT dispersion polymerizations of AM. However, the charge density in mCTA had a significant effect on phase separation since the positive charge of mCTA made the dispersion stable. When mCTA with lower C was used, the repulsive force between particles decreased. Run DA₂₀₆C₁₄₃-6-50 had Z-average particle size (D_z) of 1057 nm. With the increase of C on mCTA to 45.5%, the particle sizes of the dispersion phase became smaller with $D_z = 723.0$ nm. The increment of C on mCTA to 51.0% further modified the particle size distribution with it becoming bimodal with a D_z at 43.8 nm and another, presumably for the aggregates at 656.4 nm.

Also studied was the effect of mCTA chain length for a fixed C of 51% on RAFT dispersion polymerizations. The molecular weights of mCTAs ranged from 28.8 kg mol⁻¹ (DA₁₀₂C₁₀₄-4.2-50), 48.8 kg mol⁻¹ (DA₁₇₀C₁₇₇-4.2-50), to 69.2 kg mol⁻¹ (DA₂₄₄C₂₅₀-4.2-50). Longer mCTA chains increased the viscosity of the polymerization system, resulting in higher conversions (from 68.6 to 94.2%) and CPAM molecular weight (from 82.5 to 104.6 kg mol⁻¹) due to diffusion-controlled effect. The longer mCTA (DA₂₄₄C₂₅₀-4.2-50) was beneficial to stabilizing the polymerization system and produced particles having the smallest particles ($D_z = 49.73$ nm). When the mCTA became shorter, the formed particles became larger, D_z of 165.7 nm for Run DA₁₇₀C₁₇₇-4.2-50 and 210.5 nm for Run DA₁₀₂C₁₀₄-4.2-50, respectively, and more particle aggregation occurred.

Conclusions

Random copolymers of AM and DMC were synthesized *via* RAFT solution copolymerization at a pH of 5 with BCPA as CTA. The copolymers were used as macro RAFT chain transfer agent mCTA and stabilizer for chain extension in aqueous RAFT dispersion polymerization of AM. Random copolymers had C values ranging from 19.9 to 61.7%, M_n values ranging from 28.8 to 69.2 kg mol⁻¹, and \bar{D} values of less than 1.2. The aqueous



Table 6 Kinetic equations for chain species

Type of chains	Mass balance equation
Propagating radical chains	$\frac{d[P_r^*]}{dt} = k_p[P_{r-1}][M] - k_p[P_r^*][M] - k_a[P_r^*]\left([TP_0] + \sum_s [TP_s]\right) + \frac{1}{2}k_f\left([P_r\dot{TP}_0] + \frac{1}{2}\sum_s [P_r\dot{TP}_s]\right) - \sum_s (k_{tc} + k_{td})[P_r^*][P_s^*] - k_{ct}[P_r^*]\left(\sum_s [P_s\dot{TP}_0] - \sum_s \sum_t [P_s\dot{TP}_t]\right)$
Dormant chains	$\frac{d[TP_r]}{dt} = \frac{1}{2}\left(k_{f,0}[P_r\dot{TP}_0] + \frac{1}{2}\sum_s k_f[P_r\dot{TP}_s]\right) - \left(k_{a,0}[P_r] + \sum_s k_a[P_s^*]\right)[TP_r]$
Primary intermediate radical chains	$\frac{d[P_r\dot{TP}_0]}{dt} = k_{a,0}[P_0][TP_r] + k_a[P_r^*][TP_0] - \frac{1}{2}(k_{f,0} + k_f)[P_r\dot{TP}_0] - \sum_s k_{ct}[P_s^*][P_r\dot{TP}_0]$
Intermediate radical chains	$\frac{d[P_r\dot{TP}_s]}{dt} = k_a[P_r^*][TP_s] + k_a[P_s^*][TP_r] - k_f[P_r\dot{TP}_s] - \sum_t k_{ct}[P_t^*][P_r\dot{TP}_s]$
Dead chains	$\frac{d[P_r]}{dt} = \sum_t \sum_{s=0}^r k_{tc}[P_s^*][P_{r-s}^*] + \sum_s k_{td}[P_r^*][P_s^*] + \sum_{s=0}^r k_{ct}[P_s^*][P_{r-s}\dot{TP}_0] + \sum_{s=0}^r k_{ct}[P_{r-s}^*]\sum_{t=0}^s [P_t\dot{TP}_{s-t}]$

RAFT polymerizations of AM using random copolymers as mCTAs with $[mCTA]_0/[Va-044]_0 = 2/1$ were conducted. It was observed that the progress of narrow-distributed GPC traces of resulted block CPAMs ($\mathcal{D} \leq 1.2$) with polymerization times and existence of linear relationships between $-\ln(1 - X_{AM})$ and t , and M_w and X_{AM} , suggesting a controlled characteristic in AM block extension with AM/DMC random copolymers as mCTAs. The cationic degrees and chain lengths of mCTAs had little influence on the chain propagation of AM. When the AM addition content was more than 3.0% and the PAM block was extended from 300 to 1200 units, the polymerization rate increased rapidly due to diffusion control effect. In order to better understand the chain extension kinetics in aqueous solution, a model using the method of moments with the consideration of diffusion control effect was developed. Experimental data were well fit with model predictions. The AM/DMC random copolymers were used as stabilizer for aqueous RAFT dispersion polymerization of AM, to produce the block CPAM precursors at an elevated solid content. Stable milky CPAM dispersions were obtained with particle sizes from 10 to 200 nm and aggregates of approximately 1000 nm, which were stable on storing at room temperature for more than 1 month. The resulting block CPAMs had \mathcal{D} values of less than 1.3 with molecular weight ranging from 58.1 to 109.3 kg mol⁻¹. Effects of temperature (45 to 55 °C), mCTA cationic degree (41.0 to 51.0%) and molecular weights (28.8 to 69.2 kg mol⁻¹) on aqueous dispersion were investigated. High polymerization temperature promoted the polymerization but had little influence on particle size, while the charge density on mCTA had little effect on the reaction rate but had a great impact on phase separation. With the increase of charge density from 41.0 to 51.0% on mCTAs, the particle sizes of the dispersion phase became smaller and formed fewer aggregates. Longer mCTA chains resulted in higher conversions and produced the particles having a smaller particle size and less aggregation. The results indicate that the aqueous RAFT dispersion polymerization using random copolymers of AM and DMC at moderate

cationic degrees as stabilizer and mCTA is a suitable approach in synthesizing CPAM block precursors at an elevated solid content.

Appendix

Reaction mechanism of RAFT aqueous polymerization

A kinetic model of aqueous RAFT solution polymerization of AM mediated by mCTA was developed based on the elementary reactions listed in Table 4. Chain transfer reactions to monomer and solvent were not considered. The number-average chain length (\bar{r}_N), weight-average chain length (\bar{r}_W), \mathcal{D} , and M_n were calculated using eqn (A1)–(A4) using the method of moments.^{73,74} The moments of various chain species are defined in Table 5. The specific derivations for the balance of species and moments are shown in Tables 6 and 7.

$$\bar{r}_N = \frac{Y_1 + Y_1^T + Q_1 + Q_1^T}{Y_0 + Y_0^T + Q_0 + Q_0^T} + \bar{r}_{N,mCTA} \quad (A1)$$

$$\bar{r}_W = \frac{Y_2 + Y_2^T + Q_2 + Q_2^T}{Y_1 + Y_1^T + Q_1 + Q_1^T} + \bar{r}_{W,mCTA} \quad (A2)$$

$$\mathcal{D} = \frac{\bar{r}_W}{\bar{r}_N} \quad (A3)$$

$$M_n = (\bar{r}_N - \bar{r}_{N,mCTA})m_{AM} + M_{n,mCTA} \quad (A4)$$

where $\bar{r}_{N,mCTA}$ and $\bar{r}_{W,mCTA}$ represent the contribution made by mCTA, respectively. m_{AM} is the molecular weight of AM.

Diffusion-controlled termination model

With the polymerization proceeded, the amount and length of polymeric chains would increase, which lead to increased viscosity. As a result, reactions involved in long chains could become diffusion controlled, which is quite common in a radical polymerization process.^{75,76} Here we adopted a semi-empirical equation,^{77–79} only for diffusion-controlled



Table 7 Differential equations of moments

Zeroth-order moments	Mass balance equation
Propagating radical chains	$\frac{dY_0}{dt} = 2fk_d[I] + k_f Y_0^T - k_a Y_0 Q_0^T - k_{td} Y_0 Y_0 - k_{tc} Y_0 Y_0 - k_{ct} Y_0 Y_0^T$
Intermediate radical chains	$\frac{dY_1^T}{dt} = k_a Y_0 Q_0^T - k_f Y_0^T - k_{ct} Y_0 Y_1^T$
Dormant chains	$\frac{dQ_0^T}{dt} = k_f Y_0^T - k_a Y_0 Q_0^T$
Dead chains	$\frac{dQ_0}{dt} = k_{td} Y_0 Y_0 + \frac{1}{2} k_{tc} Y_0 Y_0 + k_{ct} Y_0 Y_0^T$
First-order moments	Mass balance equation
Propagating radical chains	$\frac{dY_1}{dt} = k_p Y_0 [M] + \frac{1}{2} k_f Y_1^T - k_a Y_1 Q_1^T - k_{td} Y_0 Y_1 - k_{tc} Y_0 Y_1 - k_{ct} Y_1 Y_0^T$
Intermediate radical chains	$\frac{dY_1^T}{dt} = k_a Y_1 Q_1^T + k_a Y_0 Q_1^T - k_f Y_1^T - k_{ct} Y_0 Y_1^T$
Dormant chains	$\frac{dQ_1^T}{dt} = \frac{1}{2} k_f Y_1^T - k_a Y_0 Q_1^T$
Dead chains	$\frac{dQ_1}{dt} = k_{td} Y_0 Y_1 + k_{tc} Y_0 Y_1 + k_{ct} Y_1 Y_0^T + k_{ct} Y_0 Y_1^T$
Second-order moments	Mass balance equation
Propagating radical chains	$\frac{dY_2}{dt} = k_p Y_0 [M] + 2k_p Y_1 [M] + \frac{1}{2} k_f Y_{2,0}^T - k_a Y_2 Q_2^T - k_{td} Y_0 Y_2 - k_{tc} Y_0 Y_2 - k_{ct} Y_2 Y_0^T$
Intermediate radical chains	$\frac{dY_2^T}{dt} = k_a Y_2 Q_2^T + 2k_a Y_1 Q_1^T + k_a Y_0 Q_2^T - k_f Y_2^T - k_{ct} Y_0 Y_2^T$
Dormant chains	$\frac{dY_{2,0}^T}{dt} = k_a Y_2 Q_2^T + k_a Y_0 Q_2^T - k_f Y_{2,0}^T - k_{ct} Y_0 Y_{2,0}^T$
Dormant chains	$\frac{dQ_2^T}{dt} = \frac{1}{2} k_f Y_{2,0}^T - k_a Y_0 Q_2^T$
Dead chains	$\frac{dQ_2}{dt} = k_{td} Y_0 Y_2 + k_{tc} Y_0 Y_2 + k_{ct} Y_1 Y_1 + k_{ct} Y_0 Y_2^T + 2k_{ct} Y_1 Y_1^T + k_{ct} Y_2 Y_0^T$
Mass balance equation	
Small molecule	$\frac{d[I]}{dt} = -k_d[I] \frac{d[M]}{dt} = -k_p [M] Y_0$

termination reaction because the termination rate constants are several order of magnitude higher than the other kinetic rate constants.

$$\frac{1}{k_t} = \frac{1}{k_{t,C}} + \frac{1}{k_{t,D}} \quad (\text{A5})$$

$$k_{t,D} = k_{t,D}^0 (\bar{r}_N)^{-a} \exp\left(\frac{-b}{v_f}\right) \quad (\text{A6})$$

$$v_f = \left[0.025 + \alpha_p (T - T_{gp})\right] \varphi_p + \left[0.025 + \alpha_m (T - T_{gm})\right] \varphi_m + \left[0.025 + \alpha_s (T - T_{gs})\right] \varphi_s \quad (\text{A7})$$

Table 8 Physical property parameters

Parameters	Values	References
α_p (K ⁻¹)	4.8×10^{-4}	16
α_m (K ⁻¹)	1×10^{-3}	80
α_s (K ⁻¹)	0.012	81
T_{gm} (K)	260	84
T_{gs} (K)	136	84
T_{gp} (K)	438	82

Table 9 Kinetic rate constants

Parameters	Values	Ref.
k_d (s ⁻¹)	4.02×10^{-5}	83
k_p (L mol ⁻¹ s ⁻¹)	$4.266 \times 10^7 \exp(-3909.61/T)$	83
$k_a = k_{a,0}$ (L mol ⁻¹ s ⁻¹)	4×10^4	This work
$k_f = k_{f,0}$ (s ⁻¹)	5×10^3	This work
k_{tc} (L mol ⁻¹ s ⁻¹)	$2.0 \times 10^{10} \exp(-1553.01/T)$	83
$k_{t,D}^0$	3×10^{12}	This work
a	0.3	This work
b	0.7	This work
k_{ct}	2×10^6	This work

where $k_{t,D}^0$, a , and b are adjustable parameters to correlate the experimental data. The other parameters are obtained from the literature with their physical meanings described in the Table 8.

Author contributions

The manuscript was written with contributions of all authors. All authors have given approval to the final version of the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work is financially supported by National Key Research and Development Program of China (2016YFC0206003), and National Natural Science Foundation of China (21420102008, 21536011, 21074116, and 20936006). The authors thank Mr Eryuan Fang for his assistance in performing GPC analyses at State Key Laboratory of Chemical Engineering (Zhejiang University).

Notes and references

- C. S. Lee, J. Robinson and M. F. Chong, *Process Saf. Environ. Prot.*, 2014, **92**, 489.
- K. E. Lee, N. Morad, T. T. Teng and B. T. Poh, *Chem. Eng. J.*, 2012, **203**, 370.
- D. A. Z. Wever, F. Picchioni and A. A. Broekhuis, *Prog. Polym. Sci.*, 2011, **36**, 1558.
- M. Ma and S. Zhu, *Colloid Polym. Sci.*, 1999, **277**, 115.



- 5 M. Ma and S. Zhu, *Colloid Polym. Sci.*, 1999, **277**, 123.
- 6 R. Subramanian, S. Zhu and R. H. Pelton, *Colloid Polym. Sci.*, 1999, **277**, 939.
- 7 D. Li, S. Zhu, R. H. Pelton and M. Spafford, *Colloid Polym. Sci.*, 1999, **277**, 108.
- 8 Á. Blanco, E. Fuente, M. C. Monte, N. Cortés and C. Negro, *Ind. Eng. Chem. Res.*, 2009, **48**, 4826.
- 9 F. Brouillette, D. Morneau, B. Chabot and C. Daneault, *Pulp Pap. Can.*, 2004, **105**, 31.
- 10 J. Chiefari, Y. K. Chong, F. Ercole, J. Krstina, J. Jeffery, T. P. T. Le, R. T. A. Mayadunne, G. F. Meijs, C. L. Moad, G. Moad, E. Rizzardo and S. H. Thang, *Macromolecules*, 1998, **31**, 5559.
- 11 Z. Wang, J. He, Y. Tao, L. Yang, H. Jiang and Y. Yang, *Macromolecules*, 2003, **36**, 7446.
- 12 H. T. Lord, J. F. Quinn, S. D. Angus, M. R. Whittaker, M. H. Stenzel and T. P. Davis, *J. Mater. Chem.*, 2003, **13**, 2819.
- 13 L. Zhang and Y. Chen, *Polymer*, 2006, **47**, 5259.
- 14 I. Capek and W. Funke, *Macromol. Chem.*, 1990, **191**, 121.
- 15 D. Wang, W.-J. Wang, B.-G. Li and S. Zhu, *AIChE J.*, 2013, **59**, 1322.
- 16 D. Wang, X. Li, W.-J. Wang, X. Gong, B.-G. Li and S. Zhu, *Macromolecules*, 2012, **45**, 28.
- 17 W.-J. Wang, D. Wang, B.-G. Li and S. Zhu, *CN Pat.*, 101693749B, 2009.
- 18 W.-J. Wang, D. Wang, B.-G. Li and S. Zhu, *CN Pat.*, 102153714B, 2010.
- 19 D. B. Thomas, A. J. Convertine, R. D. Hester, A. B. Lowe and C. L. McCormick, *Macromolecules*, 2004, **37**, 1735.
- 20 D. B. Thomas, A. J. Convertine, L. J. Myrick, C. W. Scales, A. E. Smith, A. B. Lowe, Y. A. Vasilieva, N. Ayres and C. L. McCormick, *Macromolecules*, 2004, **37**, 8941.
- 21 G. Qi, C. W. Jones and F. J. Schork, *Macromol. Rapid Commun.*, 2007, **28**, 1010.
- 22 K. Vijayakrishna, S. K. Jewrajka, A. Ruiz, R. Marcilla, J. A. Pomposo, D. Mecerreyes, D. Taton and Y. Gnanou, *Macromolecules*, 2008, **41**, 6299.
- 23 S. S. Eamegdool, M. W. Weible II, B. T. T. Pham, B. S. Hawckett, S. M. Grieve and T. Chan-ling, *Biomaterials*, 2014, **35**, 5549.
- 24 A. Asadujjaman, B. Kent and A. Bertin, *Soft Matter*, 2017, **13**, 658.
- 25 M. Hayashi, A. Noro and Y. Matsushita, *Macromol. Rapid Commun.*, 2016, **37**, 678.
- 26 S. Lin, J. Shang and P. Theato, *Polym. Chem.*, 2017, **8**, 2619.
- 27 S. Bai, Y. Wang, B. Liu, Y. Zhu and R. Guo, *Colloids Surf., A*, 2018, **553**, 446.
- 28 E. Abdollahi, A. Khalafi-Nezhad, A. Mohammadi, M. Abdouss and M. Salami-Kalajahi, *Polymer*, 2018, **143**, 245.
- 29 X.-M. Wu, L.-L. Wang, Y. Wang, J.-S. Gu and H.-Y. Yu, *J. Membr. Sci.*, 2012, **421–422**, 60.
- 30 R. Ranjan and W. J. Brittain, *Macromolecules*, 2007, **40**, 6217.
- 31 J. Huang, W.-J. Wang, B.-G. Li and S. Zhu, *Macromol. Mater. Eng.*, 2013, **298**, 391.
- 32 D. Yang, D.-P. Yu, X.-F. Dong, A.-J. Shen, G.-W. Jin, Z.-M. Guo, J.-Y. Yan, M.-Y. Liu and X.-M. Liang, *Chin. J. Anal. Chem.*, 2015, **43**, 1439.
- 33 L. Xiang, W.-J. Wang, B.-G. Li and S. Zhu, *Macromol. React. Eng.*, 2017, **11**, 1700023.
- 34 M. Takara, M. Toyoshima, H. Seto, Y. Hoshino and Y. Miura, *Polym. Chem.*, 2014, **5**, 931.
- 35 L. Xun, J. Qiao, L. Qi, J. Huang and H. Cai, *Anal. Methods*, 2015, **7**, 9906.
- 36 C. Gupta, M. J. Sverdløve and N. R. Washburn, *Soft Matter*, 2015, **11**, 2691.
- 37 K. S. Silmore, C. Gupta and N. R. Washburn, *J. Colloid Interface Sci.*, 2016, **466**, 91.
- 38 C. Gupta and N. R. Washburn, *Langmuir*, 2014, **30**, 9303.
- 39 L. Xiong, X. Qin, H. Liang, S. Huang and Z. Lian, *Polym. Compos.*, 2017, **38**, 27.
- 40 Y. Shan, Y. Fu and M. Qin, *Adv. Mater. Res.*, 2012, **476–478**, 2256.
- 41 M. Qi, K. Li, Y. Zheng, T. Rasheed and Y. Zhou, *Langmuir*, 2018, **34**, 3058.
- 42 K. Ma, Y. Xu and Z. An, *Macromol. Rapid Commun.*, 2015, **36**, 566.
- 43 I. Capek and W. Funke, *Macromol. Chem.*, 1990, **191**, 121.
- 44 M. S. Cho, K. J. Yoon and B. K. Song, *J. Appl. Polym. Sci.*, 2002, **83**, 1397.
- 45 S. Guorong and C. Zhihai, *J. Appl. Polym. Sci.*, 2009, **111**, 1409.
- 46 D. Wang, PhD thesis, Zhejiang University, 2012.
- 47 M. Semsarilar and S. Perrier, *Nat. Chem.*, 2010, **2**, 811.
- 48 L. Barner, T. P. Davis, M. H. Stenzel and C. Barner-Kowollik, *Macromol. Rapid Commun.*, 2007, **28**, 539.
- 49 B. Voit I and A. Lederer, *Chem. Rev.*, 2009, **109**, 5924.
- 50 A. B. Lowe and C. L. McCormick, *Prog. Polym. Sci.*, 2007, **32**, 283.
- 51 X. Li, W.-J. Wang, B.-G. Li and S. Zhu, *Macromol. React. Eng.*, 2014, **9**, 90.
- 52 X. Li, W.-J. Wang, F. Weng, B.-G. Li and S. Zhu, *Ind. Eng. Chem. Res.*, 2014, **53**, 7321.
- 53 X. Li, W.-J. Wang, B.-G. Li and S. Zhu, *Macromol. React. Eng.*, 2011, **5**, 467.
- 54 M. Drache, G. Schmidt-Naake, M. Buback and P. Vana, *Polymer*, 2005, **46**, 8483.
- 55 H. Tobita, *Macromol. Theory Simul.*, 2009, **18**, 108.
- 56 H. Tobita, *Macromol. Theory Simul.*, 2009, **18**, 120.
- 57 Y. Luo and B. Yu, *Polym.-Plast. Technol. Eng.*, 2004, **43**, 1299.
- 58 P. Vana, T. P. Davis and C. Barner-Kowollik, *Macromol. Theory Simul.*, 2002, **11**, 823.
- 59 A. Feldermann, A. A. Toy, H. Phan, M. H. Stenzel, T. P. Davis and C. Barner-Kowollik, *Polymer*, 2004, **45**, 3997.
- 60 C. Barner-Kowollik, J. F. Quinn, D. R. Morsley and T. P. Davis, *J. Polym. Sci., Part A: Polym. Chem.*, 2001, **39**, 1353.
- 61 G. Jaramillo-Soto, M. L. Castellanos-Cárdenas, P. R. García-Morán, E. Vivaldo-Lima, G. Luna-Bárcenas and A. Penlidis, *Macromol. Theory Simul.*, 2008, **17**, 280.
- 62 A. R. Wang and S. Zhu, *J. Polym. Sci., Part A: Polym. Chem.*, 2003, **41**, 1553.
- 63 J. J. Hernández-Meza, G. Jaramillo-Soto, P. R. García-Morán, J. Palacios-Alquisira and E. Vivaldo-Lima, *Macromol. React. Eng.*, 2009, **3**, 101.



- 64 J. C. Hernández-Ortiz, E. Vivaldo-Lima, M. A. Dubé and A. Penlidis, *Macromol. Theory Simul.*, 2014, **23**, 147.
- 65 P. López-Domínguez, J. C. Hernández-Ortiz, K. J. Barlow, E. Vivaldo-Lima and G. Moad, *Macromol. React. Eng.*, 2014, **8**, 706.
- 66 J. Pallares, G. Jaramillo-Soto, C. Flores-Cataño, E. Vivaldo-Lima, L. M. F. Lona and A. J. Penlidis, *J. Macromol. Sci., Part A: Pure Appl. Chem.*, 2006, **43**, 1293.
- 67 X. Sun, Y. Luo, R. Wang, B. G. Li, B. Liu and S. Zhu, *Macromolecules*, 2007, **40**, 849.
- 68 S. Liang, X. Li, W. J. Wang, B. G. Li and S. Zhu, *Macromolecules*, 2016, **49**, 752.
- 69 W.-J. Wang, M. Kang, B. Huang and B.-G. Li, *CN Pat.*, 109180859A, 2018.
- 70 M. Jesberger, L. Barner, M. H. Stenzel, E. Malmström, T. P. Davis and C. Barner-Kowollik, *J. Polym. Sci., Part A: Polym. Chem.*, 2003, **41**, 3847.
- 71 G. S. Misra and H. Narain, *Macromol. Chem. Phys.*, 1968, **113**, 85.
- 72 G. S. Misra and U. D. N. Bajpai, *J. Macromol. Sci., Part A: Pure Appl. Chem.*, 1979, **13**, 1135.
- 73 E. Mastan and S. Zhu, *Eur. Polym. J.*, 2015, **68**, 139.
- 74 Y.-N. Zhou and Z.-H. Luo, *Macromol. React. Eng.*, 2016, **10**, 516.
- 75 A. R. Wang and S. Zhu, *Macromol. Theory Simul.*, 2003, **12**, 196.
- 76 D. Achilias and C. Kiparissides, *J. Appl. Polym. Sci.*, 1988, **35**, 1303.
- 77 F. L. Marten and A. E. Hamielec, *J. Appl. Polym. Sci.*, 1982, **27**, 489.
- 78 P. Neogi, in *Diffusion in Polymers*, Marcel Dekker, New York, 1996. p. 143.
- 79 H. Suzuki and V. B. F. Mathot, *Macromolecules*, 1989, **22**, 1380.
- 80 M. A. Dube, K. Rilling and A. Penlidis, *J. Appl. Polym. Sci.*, 1991, **43**, 2137.
- 81 *Handbook of Chemistry and Physics*, ed. Y. Yao, T. Xie and Y. Gao, Shanghai Scientific and Technical Publishers, Shanghai, 1985.
- 82 K. M. Jones, D. Bhattacharya, J. L. Brash and A. E. Hamielec, *Polymer*, 1986, **27**, 602.
- 83 J. Brandrup, E. Immergut, E. Grulke, A. Abe, and D. Bloth, in *Polymer Handbook*, Wiley, New York, 4th edn, 1999.
- 84 R. F. Fedors, *J. Polym. Sci., Polym. Lett. Ed.*, 1979, **17**, 719.

