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Synthesis of poly(methyl methacrylate)-*b*-polyethylene (PMMA-*b*-PE) block copolymers via conventional emulsion polymerization†

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Poly(methyl methacrylate)-*b*-polyethylene (PMMA-*b*-PE) block copolymers were recently obtained simply by involving ω -unsaturated PMMA oligomers prepared by CCTP in the radical polymerization of ethylene performed in dimethylcarbonate (DMC) solution (F. Baffie, G. Patias, A. Shegiwal, F. Brunel, V. Monteil, L. Verrieux, L. Perrin, D. M. Haddleton and F. D'Agosto, *Angew. Chem., Int. Ed.*, 2021, **60**, 25356–25364). However, ω -unsaturated PMMA oligomers are more effectively obtained as a latex by emulsion CCTP. In the present paper, these resulting latexes were used as seeds in the radical emulsion polymerization of ethylene in water. The effects of the methacrylic oligomer molar mass and the initial solids content of the seeds were investigated. Cryo-TEM analysis enabled demonstration of the formation of stable latexes composed of faceted spherical particles. The increase of molar mass correlated to the consumption of unsaturated oligomers with the polymerization time indicated the formation of block copolymers. The present study successfully transposed the chemistry performed in DMC solution and PMMA-*b*-PE block copolymers were obtained by a simple post-treatment of a PMMA latex.

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Introduction

Due to its low production costs and excellent mechanical properties, polyethylene (PE) has long been the most widely produced polymer and indeed chemical globally. Its fully hydrocarbon-based backbone ensures excellent chemical and mechanical properties and stability. The production of macromolecular architectures containing ethylene and polar monomers is thus particularly relevant as such polymers will combine the properties of PE with the functionality of the polar units. However, the synthesis of such structures *e.g.*, by copolymerization of ethylene with polar monomers, is not trivial,^{1,2} and only few industrial solutions are available even though the potential applications are many and a great deal of research has been carried out over many decades. Reversible-deactivation radical polymerization (RDRP) represents today one of the most versatile tools to result in complex architectures combining polar and apolar monomers. Recently, ethylene-based copolymers were developed by our group through

the use of reversible addition–fragmentation chain transfer (RAFT),^{3–5} organotellurium mediated radical polymerization (TERP)⁶ and iodine transfer polymerization (ITP)^{7,8} under mild conditions (≤ 200 bar, ≤ 80 °C) in dimethylcarbonate (DMC), but also in dispersed media or *via* polymerization-induced self-assembly (PISA).^{9,10} These techniques, however, require the use of controlling agents, which is not always compatible with the cost of the final structures that has to remain very low, particularly when considering ethylene-based products.

Thus, using much more conventional radical polymerization techniques, complex architectures could be obtained simply by involving ω -unsaturated methacrylic oligomers in the radical polymerization of a wide range of monomers. These types of oligomers can be obtained by catalytic chain transfer polymerization (CCTP) as discovered by Smirnov and Marchenko.^{11,12} CCTP involves the use of [Co]^{II} or [Co]^{III} catalysts such as cobaloximes as catalytic chain transfer agents (CCTA) in the radical polymerization of methacrylates.^{13–19} CCTP has been carried out successfully in all processes used for free radical polymerization including solution, suspension and bulk as well as commercial processes using emulsion polymerization and indeed very recently to photopolymerization to 3D networks.²⁰ The use of CCTP in dispersed media is not just used efficiently commercially but has raised a lot of interest in academia.^{21–26} The transition of carrying out CCTP in solution to dispersed media is however not trivial. Indeed, the compartmentalization effects associ-

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ated to emulsion polymerization²⁷ affect the efficiency of the CCTA, which needs to partition between the aqueous phase and the polymer particles. In most cases, lower catalytic activity and thus higher molar mass are observed when CCTP is conducted in dispersed media due to mass transport limitations, partitioning and potential deactivation of catalysts in aqueous phase. It is also important for the polymerizations to be at less than 100% instantaneous conversion so the glass transition of the swollen polymer particle allows the catalyst to move in and out of the aqueous phase.^{11,21,23,28–31}

Low molar mass ω -unsaturated methacrylic oligomers have been used in radical polymerization for the synthesis of branched copolymers, amphiphilic dispersants or multi-block copolymers.^{32–36} In most of the cases, the propagating radical is added on the unsaturated chain end leading to the formation of an intermediate radical. Depending on the monomer used for the radical polymerization, and thus the corresponding propagation rate, two types of architectures can be obtained. For mostly styrene, acrylate, acrylonitrile the propagation rate is relatively high and propagation step occurs leading to grafted structures.^{37,38} However, in the case of acrylates, depending on the nature of the acrylate and its molecular weight, block copolymer structures can be obtained after termination steps of the intermediate radical.^{32,39–41} When methacrylates are used, the rate of propagation is lower than the rate of β -scission leading to an addition–fragmentation chain transfer mechanism and to the formation of block copolymers.^{34,42–45}

The use of ω -unsaturated oligomers synthesized by CCTP in emulsion has also been established. For instance, block copolymers with a poly(*n*-butyl methacrylate) block have been produced in emulsion polymerization from PMMA macromonomers.^{34,42,44,45} They can also be used as *in situ* stabilizer in order to conduct a surfactant-free emulsion polymerization of *n*-butyl acrylate (BA).^{42,46,47}

We recently showed that PMMA produced by CCTP (PMMA_n) can serve as efficient degradative chain transfer agent in the radical polymerization of ethylene conducted in DMC under relatively mild conditions (80 bar and 70 °C).⁴⁸ Detailed characterizations of the obtained products combined with DFT calculations showed that the propagating polyethylenyl radicals produced upon initiation of ethylene polymerization rapidly react with the ω -chain end of the PMMA_n to form an intermediate radical (Fig. 1). The fate of this intermediate radical was demonstrated to be termination by H-abstraction to lead to PMMA_n-*b*-PE for $n > 2$. Upon consumption of the double bond of PMMA_n, the molar mass of the PE segment increased consistently with a polymerization mechanism implying a degradative chain transfer reaction. Depending on the amount of ω -unsaturated oligomers, the molar mass of the PE block could be adjusted.

As mentioned above, CCTP of MMA can easily be carried out in emulsion using sodium dodecylsulfate (SDS) as surfactant to produce a stable latex incorporating ω -vinyl functional PMMA.^{34,42,44,45} Building on our results obtained in DMC, we now report application in aqueous dispersed media by the

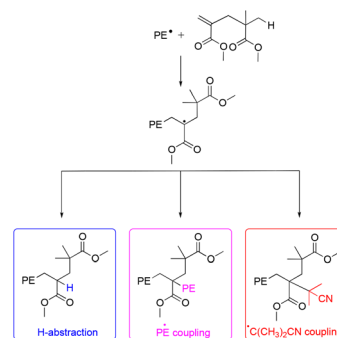


Fig. 1 Possible fate of the propagating radicals in ethylene polymerization in the presence of PMMA₂. Reproduced from ref. 48.

direct use of PMMA_n from CCTP latexes as seeds in ethylene polymerization. Indeed, this approach would be a powerful way to produce PMMA-*b*-PE copolymers directly in water, simply by post-treating the seed latex with initiator under ethylene pressure. The study explores the effect of the molar mass of PMMA oligomers and of the initial solids content of the seed latex on the polymer architecture. Morphologies of the produced particles are also investigated.

Materials and methods

Materials

Methyl methacrylate (MMA, Aldrich, ≥99%), 4,4'-azobis(4-cyanovaleric acid) (ACVA, Aldrich, ≥98%), sodium dodecylsulfate (SDS, MP Biomedicals, LLC), ammonium persulfate (APS, Aldrich, ≥98%) and sodium bicarbonate (NaHCO₃, Aldrich, Bioreagent) were used as received. Water was deionized before use (Purelab Classic UV, Elga LabWater). The catalyst bis[(difluoroboryl)dimethylglyoximate]cobalt(II) (CoBF) was synthesised as previously described.^{32,49} Deuterated solvents were supplied by Eurisotop.

Analytical techniques

Nuclear magnetic resonance (NMR). All NMR experiments were performed at 90 °C in a mixture of tetrachloroethylene and deuterated benzene (2 : 1 v : v) using a Bruker Avance 400 spectrometer. Proton NMR spectra were recorded with a 5 mm BBFO + probe with a z-gradient coil. Chemical shifts are given in parts per million (ppm) with the benzene deuterated solvent used as internal standard.

Dynamic light scattering (DLS). The intensity-weighted mean diameter (or Z-average diameter) (Z_{ave}) of the latex particles and the polydispersity index (PDI) were measured at 25 °C using a Zetasizer Nano Series (Nano ZS) from Malvern Instruments. Before measurements, the latex was diluted with deionized water. The mean particle diameter was averaged over three consecutive runs. The data were collected at a 173° scattering angle using the fully automatic mode of the Zetasizer system and fitted with the cumulant analysis.



Differential scanning calorimetry (DSC). DSC analyses were performed on a Mettler Toledo DSC 3. Around 10 mg of samples were precisely weighted in 40 μL alumina pans. Two heating steps separated by one cooling step were performed at a rate of 10 $^{\circ}\text{C min}^{-1}$ with temperatures ranging from to -40 $^{\circ}\text{C}$ to 150 $^{\circ}\text{C}$. Crystallization, melting and glass transition temperatures (T_c , T_m , T_g) were recorded on the second cycle.

Size exclusion chromatography in THF (SEC-THF). SEC-THF analyses were performed using a Viscotek system (Malvern Instruments) including a four-capillary differential viscometer, a differential refractive index (RI) detector and a UV detector. THF was used as the mobile phase at a flow rate of 1 mL min^{-1} at 35 $^{\circ}\text{C}$. All samples were injected at a concentration of 3 to 5 mg mL^{-1} after filtration through a 0.45 μm PTFE membrane. The separation was carried out on three Mixed C columns (SDVB, 5 μm , 300 \times 7.5 mm) from Agilent Technology and a guard column. The average molar masses (number-average molar mass, M_n , and weight-average molar mass, M_w) and the dispersity, $\bar{D} = M_w/M_n$, were calculated from the RI signal with a calibration curve based on narrow poly(methyl methacrylate) standards from Polymer Standards Service.

High temperature size exclusion chromatography (HT-SEC). HT-SEC analyses were performed using an Agilent Infinity II High Temperature GPC instrument equipped with differential refractive index (DRI), viscometry (VS) and dual angle light scattering (LS 90 + 15) detectors. The system was equipped with 2 \times PLgel Olexis columns (300 \times 7.5 mm) and a 10 μm guard column. 1,2,4-Trichlorobenzene (TCB) was used as mobile phase with 250 ppm of butylated hydroxytoluene (BHT) as antioxidant. Analyte samples were dissolved at 150 $^{\circ}\text{C}$ and hot filtered through a stainless-steel frit with 10 μm pore size before injection. Samples were run at 1 mL min^{-1} at 160 $^{\circ}\text{C}$. Polystyrene standards (Agilent EasiVials) were used to create a third-order calibration within a molar mass range of 6 570 000–580 g mol^{-1} . M_n and \bar{D} values were determined by conventional calibration using Agilent GPC/SEC software.

Cryogenic-transmission electron microscopy (cryo-TEM). The diluted latex samples were dropped onto 300 mesh holey carbon films (Quantifoil R2/1) and immediately quench-frozen in liquid ethane. The grid was then mounted on a precooled Gatan 626 specimen holder and transferred to a Philips CM120 microscope operating at an accelerating voltage of 120 kV (Centre Technologique des Microstructures (CTM), platform of the Université Claude Bernard Lyon 1, Villeurbanne, France). The average particle size of the particles was calculated out of 200 particles using ImageJ software. The number-average (D_n , nm) and the weight-average (D_w , nm) particle diameters and the particle-diameter dispersity (D_w/D_n) were calculated with the following equations:

$$D_n = \frac{\sum n_i D_i}{\sum n_i} \quad D_w = \frac{\sum n_i D_i^4}{\sum n_i D_i^3}$$

where n_i is the number of particles with diameter D_i .

Fourier-transform infrared spectroscopy (FT-IR). FT-IR measurements were performed on a iS50 + ATR FT-IR device from Thermo Fisher Scientific at room temperature. The

spectrometer is equipped with a diamond crystal ATR accessory, a deuterated triglycine sulfate (DTGS) detector and KBr optics. A small amount of dried latex was pressed directly on the diamond crystal. Background and sample were acquired using 32 scans at a spectral resolution of 4 cm^{-1} from 4000 to 400 cm^{-1} . The FTIR spectra were treated with Omnic software.

Synthetic procedures

General procedure for the preparation of ω -functional poly(methyl methacrylate) oligomer latexes via CCTP in emulsion. CoBF (22.43 mg, 0.0583 mmol) was placed in a 250 mL round bottom flask together with a stirring bar. Nitrogen was purged in the flask for at least 10 min. Subsequently, MMA (112 mL, 105.28 g, 1.05 mol) previously deoxygenated for 30 min using argon was added to the flask *via* a deoxygenated syringe. The mixture was vigorously stirred under inert atmosphere until total dissolution of the catalyst. Meanwhile, ACVA (1.36 g, 4.84 mmol), SDS (2.14 g, 7.43 mmol) and 250 mL of water were charged into a three-neck, 500 mL double jacketed reactor, equipped with a thermometer and an overhead stirrer. The mixture was purged with nitrogen and stirred at 325 rpm for at least 30 min. Subsequently, the mixture was heated under inert atmosphere. When the temperature in the reactor reached 70 $^{\circ}\text{C}$, the addition of the solution of MMA and CoBF started using a deoxygenated syringe and a syringe pump (feeding rate = 1.87 mL min^{-1} , feeding time = 60 min). When the addition was over, stirring continued for a further 60 min under the same conditions. Stable latex containing PMMA oligomers of different degrees of polymerization (n) were obtained and labelled PMMA $_n$.

General procedure for the ethylene seeded emulsion polymerization using PMMA $_n$ as seeds. The reaction was conducted in a stainless-steel reactor (160 mL, from Parr Instrument Co.) equipped with a thermometer, a pressure sensor, a mechanical stirring apparatus, and safety valves. In a typical polymerization procedure (Table 2), the seed latex of PMMA $_{16}$ was initially added in the autoclave with NaHCO_3 (pH = 8) and deoxygenated under argon for 30 min. APS (123 mg, 0.54 mmol) and NaHCO_3 (to reach pH = 8) were dissolved in water (the volume of water added depends on the initial solids content targeted and the total volume of the reactor $V_{\text{tot}} = 50$ mL) and this solution was introduced after degassing *via* an injecting chamber pressurized with 30 bar of ethylene into the preheated autoclave at 80 $^{\circ}\text{C}$. Ethylene gas was then introduced into the reactor until the desired pressure (100 bar) was reached. The reaction medium was stirred at 250 rpm. In order to manage the polymerization safely over 50 bar of ethylene, a 1.5 L intermediate tank was used. It was cooled at -20 $^{\circ}\text{C}$ during a filling step from the ethylene bottle. After isolation, it was then heated back and connected to the reactor to get the desired pressure. This tank was used to charge the reactor. After the desired polymerization time, the stirring speed was reduced, and the autoclave was cooled with iced water. When the temperature inside the autoclave dropped below 35 $^{\circ}\text{C}$, the remaining pressure was carefully released by venting to the atmosphere. The solids content (τ_s) was determined by gravimetric analysis and the consumption of the double bond carried



by PMMA_n was followed by ¹H NMR spectroscopy performed on dried samples. Copolymers were also analyzed by SEC and DSC after drying the latex sample. The latex was directly analyzed by DLS to measure the particle size. Some latexes were also observed by cryo-TEM.

Preparation of the PMMA₁₆/PE blend for DSC analysis. 1.5 mg of PMMA₁₆ dried extract was added in the alumina pan directly with 9.5 mg of PE obtained by radical emulsion polymerization with the same initiating and stabilization systems.

Results and discussion

Our previous study demonstrated that PMMA-*b*-PE copolymers can be formed by combining CCTP and radical polymerization of ethylene in DMC.⁴⁸ In that work, PMMA_n oligomers obtained by CCTP were solubilized in DMC in the presence of

AIBN. PMMA_n chain-end double bonds were quickly consumed, and the molar mass of the PE block increased with the reaction time. Herein, latexes of PMMA_n particles are used as seeds in the radical emulsion polymerization of ethylene to transpose the chemistry performed in DMC to aqueous media and thus produce particles of PMMA_n-*b*-PE block copolymers.

Seeded emulsion polymerization of ethylene with PMMA₁₆

Using SDS as surfactant and ACVA as initiator, two batches of latex were produced with two polymerization degrees (namely PMMA₁₆ and PMMA₃₆) adjusted by the amount of CoBF involved in the recipe (Table 1). Both latexes were stable with particle size of 147 and 101 nm, respectively. The two significantly different *T_g* values (34 °C for PMMA₁₆ and 93 °C for PMMA₃₆) are directly related to the difference of molar masses.

Seeded radical emulsion polymerization of ethylene was first carried out using the PMMA₁₆ latex. The seed latex was diluted to an initial solids content of 5 wt% and introduced into the reactor. Subsequently, an aqueous phase containing APS was injected in the reactor with NaHCO₃ used as buffer. APS was chosen as initiator in our system although persulfate initiators could react with the remaining CoBF³¹ in the PMMA₁₆ seed latex. To counterbalance this effect, the APS concentration was increased compared to the reference concentration used in more classical emulsion polymerization of ethylene previously performed in our group.⁵⁰

After 4 h of reaction under 100 bar of ethylene at 80 °C (run 1, Table 2), a stable latex was obtained. The particle size did not vary significantly (147 nm for the seed PMMA₁₆ and

Table 1 Characteristics of the PMMA_n latexes

PMMA _n	τ_s^a (%)	Z_{ave} (nm)/PDI ^b	$M_{n,NMR}^c$ (g mol ⁻¹)	$M_{n, SEC}$ (g mol ⁻¹)/ D^d	T_g^e (°C)
PMMA ₁₆	16.9	147 0.07	1 470	1200 1.8	34
PMMA ₃₆	19.9	101 0.09	3 630	3000 2.2	93

^a Determined by gravimetric analysis. ^b Determined by DLS.

^c Calculated by assuming that there is one vinyl per chain.

^d Determined by SEC-THF using conventional PMMA calibration.

^e Determined by DSC at 10 °C min⁻¹.

Table 2 PMMA₁₆ and PMMA₃₆ seeded emulsion polymerizations of ethylene and seeded emulsion polymerizations of ethylene with a PMMA latex obtained by FRP^a

Run	PMMA _n	<i>T</i> (h)	$\tau_{s,i}^b$ (%)	$\tau_{s,f}^b$ (%)	Z_{ave}^c (nm)	PDI ^c	PMMA conv. ^d (%)	DP _{PE} ^e	$M_{n,SEC}^f$ (g mol ⁻¹)	D^f	T_g^g (°C)	T_c^g (°C)
1 ^h	PMMA ₁₆	4	5.0	7.4	155	0.06	99	26	4400 505 200	6.1 1.4	—	42 56
2 ⁱ	PMMA ₁₆	4	5.0	5.4	146	0.05	27	0	1500	2.0	n.d.	n.d.
3-t ₁	PMMA ₁₆	0.5	5.0	5.4	142	0.04	49	3	1500	1.9	55	—
3-t ₂	PMMA ₁₆	1	5.0	4.9	151	0.02	72	2	2300	2.2	68	—
3-t ₃	PMMA ₁₆	2	5.0	5.5	154	0.05	93	5	2900	4.7	72	42
3-t ₄	PMMA ₁₆	3	5.0	6.1	159	0.04	99	13	4000 419 100	5.9 1.6	86	45
4-t ₁	PMMA ₃₆	0.5	10.0	9.4	113	0.09	72	3	3000	2.2	87	—
4-t ₂	PMMA ₃₆	1	10.0	9.8	100	0.07	96	11	8700	4.1	88	42
4-t ₃	PMMA ₃₆	2	10.0	12.1	110	0.02	95	48	6700 410 300	4.8 5.7	97	42
4-t ₄	PMMA ₃₆	4	10.0	14.6	120	0.07	92	105	5900 837 000	5.7 1.5	—	42 56
4-t ₅	PMMA ₃₆	6	10.0	15.6	117	0.03	92	123	7 300 1 016 000	5.2 1.5	—	42 56
5	PMMA ₁₆	4	10.0	11.6	155	0.03	98	3	5200	3.9	80	—
6	PMMA-FRP	4	5.0	14.8	68	0.03	n.d.	n.d.	7200	72.2	—	55

^a APS (0.54 mmol), NaHCO₃ (pH = 8) at 80 °C and 100 bar in water (*V*_{tot} = 50 mL). ^b The initial and final solids content determined by gravimetric analysis. ^c Determined by DLS. ^d Calculated by comparing the ¹H NMR signals intensities of the vinylic protons of PMMA₁₆ and residual vinylic protons at the end of the reaction. ^e Calculated by assuming that there is one methacrylic oligomer per PE chain, DP_{PE} is the degree of polymerization and is calculated according to the eqn (1) using NMR. ^f Determined by HT-SEC with a conventional calibration based on PS standards. Two separate entries per row corresponds to a bipopulated molar mass distribution. ^g Determined by DSC (“—” is used when glass transition or crystallization was not visible). ^h This experiment can also be considered as a run 3-t₅. ⁱ Experiment conducted without ethylene pressure.



155 nm for the obtained latex). The 2.4 wt% increase in solids content correlated with the appearance of an intense signal in the ^1H NMR spectrum of the dry extract at 1.3 ppm (Fig. 2), corresponding to the methylene protons of PE, are consistent with the formation of PE during the reaction. The almost total disappearance of the characteristic signals of the vinylic protons of PMMA_{16} , in the 5.6 to 6.5 ppm region, indicated that they were almost quantitatively consumed (99% conversion after 4 h). The polymerization degree of PE (DP_{PE}) was measured by NMR (eqn (1)) assuming that the obtained product is a $\text{PMMA-}b\text{-PE}$ block copolymer. The integral of the methoxy signal from PMMA at 3.5 ppm (I_{C}) was thus fixed at a value of 48.

$$\text{DP}_{\text{PE,NMR}} = \frac{I_{\text{PE}} - I_{\text{C}}'}{4 \cdot \text{CONV} \cdot \text{PMMA}_n} \quad (1)$$

HT-SEC chromatograms of the starting PMMA_{16} and the product obtained in run 1 are compared in Fig. 3. It shows the consumption of the starting PMMA_{16} and the formation of higher molar mass product at lower elution time. In addition, a second minor population at even lower elution time ($t_{\text{el}} = 12.4$ min), *i.e.*, higher molar masses, appeared. A similar behavior was observed in our previous study conducted in DMC, in which the high molar mass population was assigned to the formation of PE homopolymer taking place when all the PMMA_n chain transfer agents have been consumed.⁴⁸

However, under the conditions employed here, one might ask whether the consumption of the PMMA_{16} unsaturated chain-ends is indeed only related to the reaction of polyethylenyl radicals formed in water (this is made possible by the solubility of ethylene in water from 0.3 g L⁻¹ at 4.5 bars to 4.6 g L⁻¹ at 110 bar at 75 °C⁵¹) and entering the particles to react with PMMA_{16} . An additional experiment was thus carried out under the exact same conditions but without ethylene

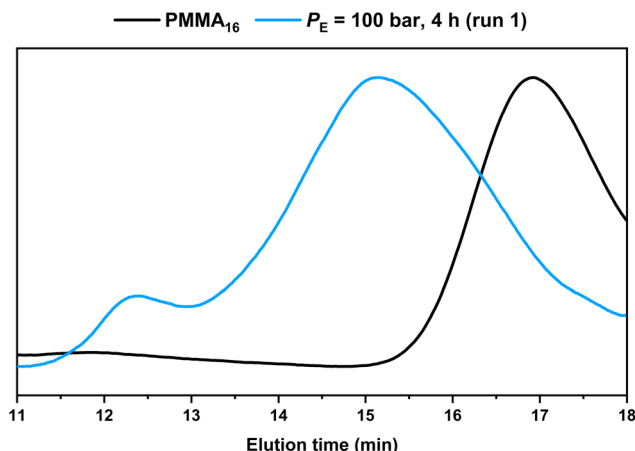


Fig. 3 HT-SEC chromatograms of the dry extract from the PMMA_{16} seed and from the latex obtained by ethylene emulsion polymerization in the presence of PMMA_{16} (run 1, Table 2).

pressure (run 2, Table 2) to further investigate this point. After 2 h of reaction at 80 °C, a plateau of 27% was reached for the consumption of the PMMA_{16} vinyl end groups. Consumption of vinyl end groups of PMMA_n was not observed when similar experiments were carried out in DMC in the presence of AIBN. The nature of the primary radicals generated (carbon- or oxygen-centered for DMC or emulsion polymerization, respectively) might be put forward to explain this different behavior. It is worth mentioning here that in emulsion polymerization of MMA initiated with APS, depending on the initial conditions (notably the initiator and surfactant concentration), PMMA oligomers will form in water with a degree of polymerization lower than 10 (the DP for surface activity, z , is close to 5, and the oligomers remain water-soluble up to a DP of 10 (j_{crit})⁵²). The formation of our PMMA seeds is based on CCTP for which the majority of the chains are not initiated by a charged group. The corresponding critical DPs should then be much lower and the amount of PMMA oligomers remaining in water small. The isolation of the aqueous phase of PMMA_{16} latex by centrifugation confirmed nevertheless the presence of some oligomers that quantitatively reacted after 2 h at 80 °C in the presence of APS (Fig. S1†). However, a reaction between the water-soluble radicals generated from APS with these water-soluble species could reasonably not account for the 27% of double bond consumption mentioned above. SEC analyses of the dry extract recovered from run 2 (Table 2) indeed revealed a significant conversion of the lowest molar mass species into higher molar mass ones shifted in the molar mass main distribution (Fig. 4). The fraction of double bonds consumed in run 2 is thus probably explained also by the reaction of part of the PMMA_{16} chains with entering radicals after initiation with APS as explained above. The shift of molar mass distribution observed in Fig. 3 when ethylene is polymerized is thus very likely related to the polyethylenyl radicals entering the particles and reacting with PMMA_{16} as expected. The following investigations aim at demonstrating that $\text{PMMA-}b\text{-PE}$ are effectively formed during this process.

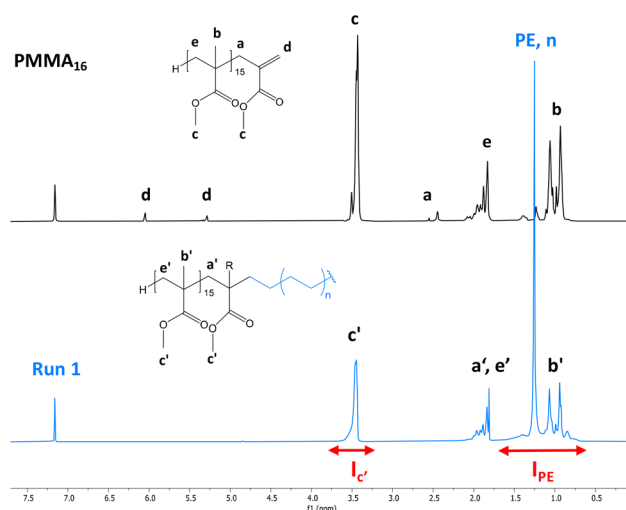


Fig. 2 ^1H NMR spectra (TCE/ C_6D_6 at 90 °C) of the dry extract (a) from the PMMA_{16} seed latex and (b) from the latex obtained by ethylene emulsion polymerization in the presence of PMMA_{16} (run 1, Table 2).



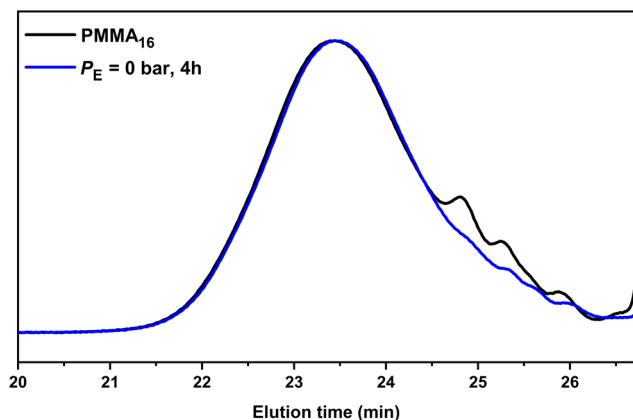


Fig. 4 SEC-THF chromatograms of the seed PMMA₁₆ and the polymer obtained after 4 h of reaction without ethylene pressure (run 2, Table 2).

Kinetic study

The influence of the molar mass of the PMMA_n latexes (*i.e.*, the value of *n*) on the polymerization kinetics was first investigated. A series of polymerizations were conducted at 80 °C and 100 bar of ethylene with both the PMMA₁₆ and PMMA₃₆ latexes, but with two different solids content (5 wt% for PMMA₁₆ and 10 wt% for PMMA₃₆) in order to keep the same amount of vinyl functions in both systems. Other parameters were kept identical to those described above with PMMA₁₆. As the reactions were conducted in a pressurized autoclave, the withdrawal of aliquots during the experiment was not possible. Therefore, each point of the kinetics (Table 2) represents an experiment. The trends observed through the data obtained (Fig. 5) are indeed showing the good consistency of the different experiments.

For both the PMMA₁₆ and PMMA₃₆ seeds, the double bonds on the methacrylic oligomers are quickly consumed at the beginning of the reaction (> 90% after 2 h) until almost being entirely consumed. Again, DP_{PE} was calculated with eqn (1) by assuming that the obtained product is a PMMA-*b*-PE block copolymer. The integral of the methoxy signal from PMMA at 3.5 ppm (*I*_c) was fixed at a value of 48 for PMMA₁₆ and 108 for PMMA₃₆ (Fig. 6). As shown in Fig. 5b, the solids content starts to strongly increase only when most of PMMA_n is consumed. This might be showing rapid termination of growing polyethylenyl radicals with PMMA_n and formation of PE homopolymer when all PMMA_n has been consumed as previously observed in DMC.⁴⁸ After 4 h, an increase of both solids content and DP_{PE} was observed for the two PMMA_n (after 4 h: $\tau_{s,f}$ = 7.4 wt% and DP_{PE} = 26 and 16.6 wt% and 123 for PMMA₁₆ and PMMA₃₆, respectively).

This is indeed consistent with HT-SEC analyses (Fig. 7) that shows the shift of the chromatograms towards higher molar masses with polymerization times in both cases and as long as PMMA_n remains. Once PMMA_n has been consumed, a second population of much higher molar mass appears. As observed in DMC, block copolymers seem to be formed early during the polymerization, and PMMA_n seems to be similarly acting as

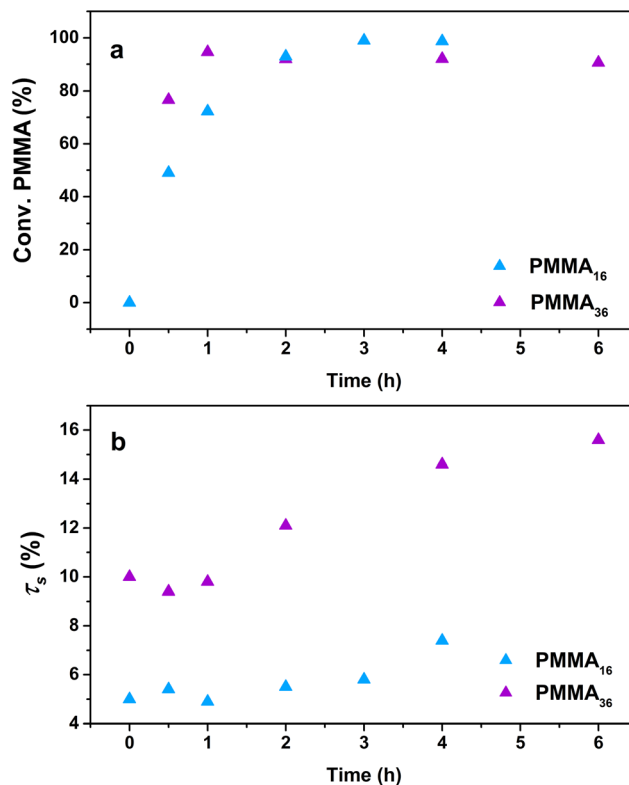


Fig. 5 Evolution of (a) PMMA_n conversion and (b) solids content with polymerization time at 80 °C and 100 bar of ethylene pressure (respectively runs 3 and 4, Table 2).

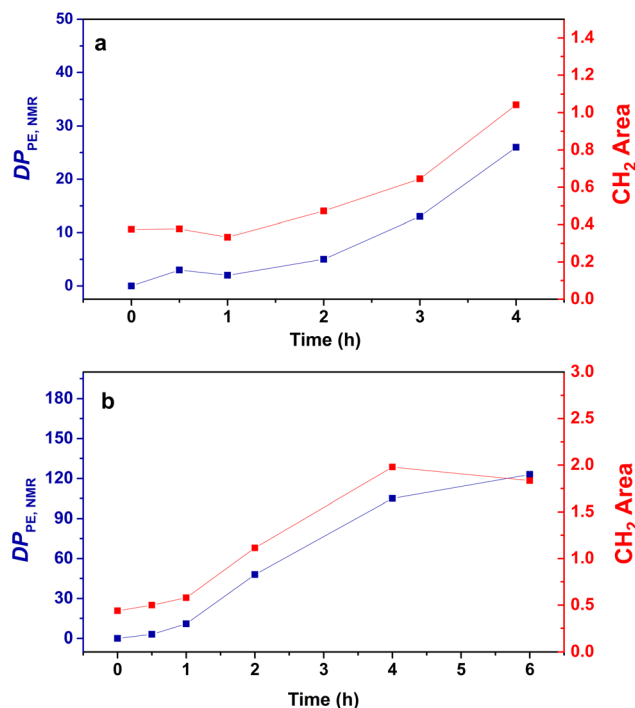


Fig. 6 Evolution of the DP_{PE} determined by NMR (blue) and the area of the absorption band of CH₂ determined by FT-IR (red) with the polymerization time for the copolymers obtained with (a) PMMA₁₆ and (b) PMMA₃₆ (respectively runs 3 and 4, Table 2).



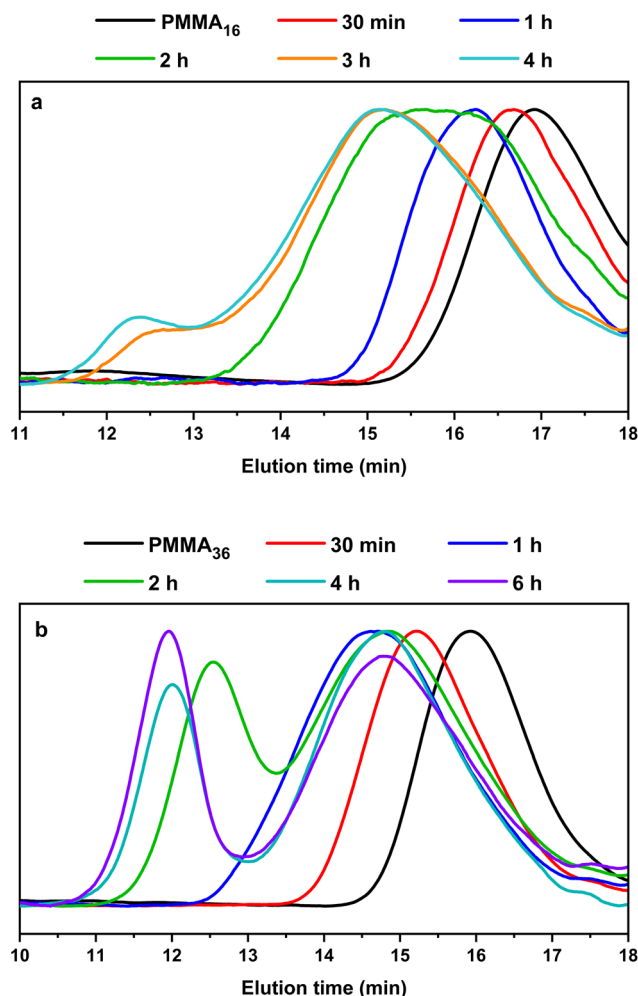


Fig. 7 HT-SEC chromatograms of the polymers obtained during seeded emulsion polymerization of ethylene with (a) PMMA₁₆ and (b) PMMA₃₆, at 80 °C and 100 bar of ethylene pressure after the indicated polymerization time (respectively runs 3 and 4, Table 2).

degradative chain transfer agent by terminating the growing PE chains entering the particles. When the reaction time increases, the amount of unsaturated oligomer decreases and the probability for a polyethylenyl radical to react on the double bond of the methacrylic oligomer is lower, so the PE block is longer. When all the vinyl functions are consumed, homopolymerization of ethylene might then take place.

No control experiment of ethylene polymerization that could be performed without PMMA_n in the exact same conditions can be compared to these experiments to show that this high molar mass population represents PE chains. Indeed, a typical radical emulsion polymerization of ethylene would not be of relevance here since it would not represent the seeded emulsion polymerization systems used. To confirm the nature of this second population, the final products were analyzed by FT-IR spectroscopy. The evolution of the absorption band area of the CH₂ stretching from ethylene units (3000–2800 cm⁻¹) was measured by normalizing the area of

the absorption band of C=O from MMA units (1735 cm⁻¹) (see Fig. S3†). If we compare the evolution of DP_{PE,NMR} and the CH₂ area with the polymerization time (Fig. 6), we can observe two different regimes of PE production. A first slow regime (up to one hour) where methacrylic oligomers are still available in the medium, and a second regime for which the production rate of PE is higher, when there are no methacrylic oligomers remaining. The transition to this second regime is also correlated with the appearance of a second high molar mass population in HT-SEC observed for elution times between 11 and 13 min (Fig. 7). This transition is observed after 2 h of polymerization for PMMA₁₆ and after 1 h for PMMA₃₆. These additional results show that a high molar mass PE homopolymer is formed when all the methacrylic oligomers have been consumed. The molar masses measured by high-temperature SEC in PS equivalent are particularly high, exceeding several hundred thousand grams per mol.⁵³

Thermal analysis was conducted on the copolymer dried extracts. The cooling curves plotted in Fig. 8 show that for short polymerization times (up to 1 h), an increase of *T_g* of the PMMA block is clearly observed (Table 2). In our case, the addition of ethylene units to a low molar mass PMMA extends the polymer chains leading to chain stiffening and hence an increase of the *T_g* value. This behavior can be interpreted by Fox and Flory⁵⁴ with the use of the following equation:

$$T_g = T_{g,\alpha} - \frac{K}{M_n} \quad (2)$$

where *K* is a constant which depends on the nature of the polymer and *T_{g,α}* the maximum glass transition temperature that can be reached at a theoretical infinite molar mass.

Fox and Flory's relation also explains the lower increase of *T_g* value of our copolymers in the case of PMMA₃₆ since its molar mass is higher and so its *T_g* value is closer to the *T_{g,α}*. For both PMMA₁₆ and PMMA₃₆, a first population of chains crystallizing at 42 °C is visible at 2 h for PMMA₁₆ and at 1 h for PMMA₃₆, followed by a second at higher temperature (55–56 °C) for longer reaction times (4 h for the two PMMA_n). The population of chains crystallizing at 55–56 °C may correspond to PE chains since they appear at the same time a second population of high molar mass chains is observed in HT-SEC (Fig. 7). The other population could mostly correspond to PMMA-*b*-PE chains, where DP_{PE} are smaller and thus crystallize at lower temperature. The thermal behavior of these copolymers was compared to the behavior an ethylene homopolymer obtained by radical emulsion polymerization with the same initiating and stabilization system (see ESI for details on its synthesis†) and to a physical blend of PMMA₁₆ and the same PE homopolymer. Firstly, the PE sample (dark) and the PMMA₁₆/PE physical blend (blue) behave similarly when subjected to the same temperature profile (Fig. 9). A single crystallization peak is observed at close values, at 61 °C for the blend and at 64 °C for the PE homopolymer. The thermal behaviour of the copolymers synthesized with PMMA₁₆ is different. The crystallization of the sample obtained after 3 h of polymerization (orange) occurs at a significantly lower temperature than



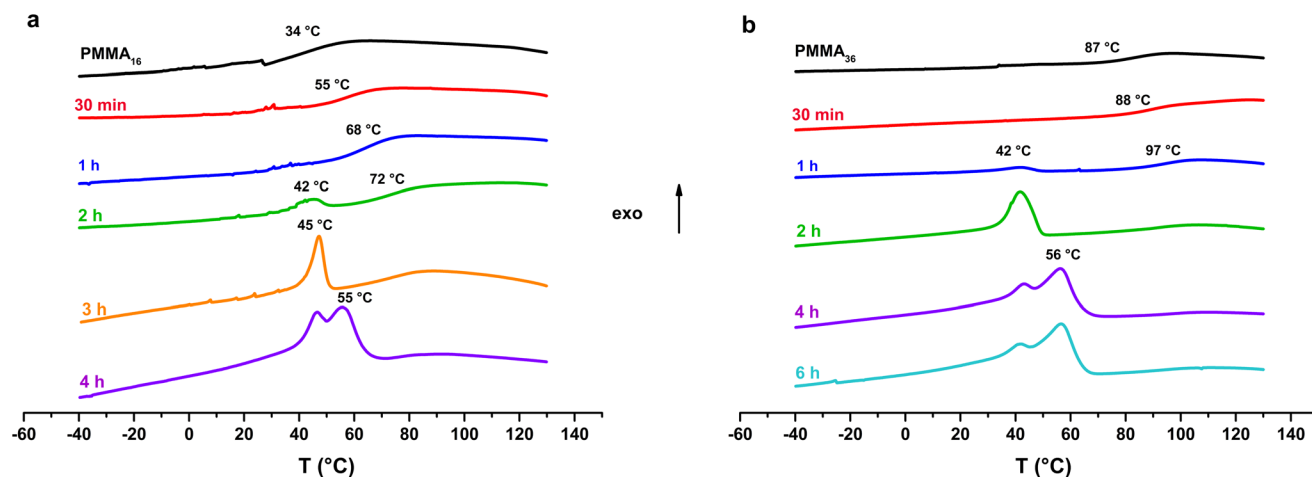


Fig. 8 DSC analysis (cooling) of the samples obtained with (a) PMMA₁₆ and (b) PMMA₃₆ (respectively runs 3 and 4, Table 2).

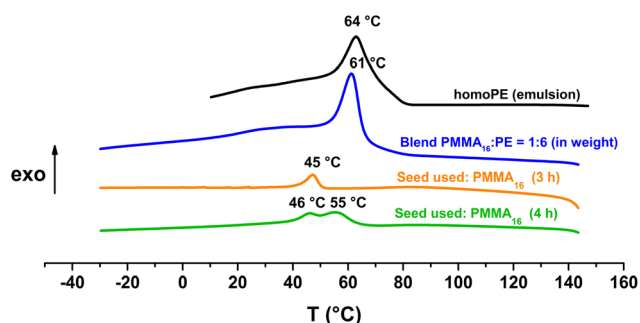


Fig. 9 DSC analyses (cooling) of the polymers obtained after polymerization with PMMA₁₆ as a seed (run 3-t₄ and run 1 in Table 2), a blend of PMMA₁₆ and PE (1 : 6 in weight), and homoPE obtained by radical emulsion polymerization in water.

that of the blend (45 °C vs. 61 °C for the blend), showing that PMMA interferes with crystallization of the PE segments and proving that it is indeed bound to PE. By comparing the thermograms of PE and of the copolymer obtained after 4 h of reaction (green), the second crystallization is most likely caused by PE chains formed when all the PMMA₁₆ has been consumed.

Finally, the colloidal features of the obtained dispersions, stable in each case, were investigated by DLS and TEM analyses.

The particle sizes determined by DLS (Fig. 10) do not significantly increase during the polymerization for the reaction with PMMA₁₆ even after a long polymerization time (142 nm at the beginning and 155 nm after 4 h), contrary to PMMA₃₆ where the particle size slightly increases (101 nm at the beginning and 117 nm after 6 h). Surprisingly, the polydispersity index decreases with the polymerization time. The reaction starts with a quite polydisperse latex, but during the reaction the size distribution seems to become narrower around higher particle sizes.

Cryo-TEM analysis of the different latexes revealed that both the PMMA₁₆ and PMMA₃₆ particles were spherical and

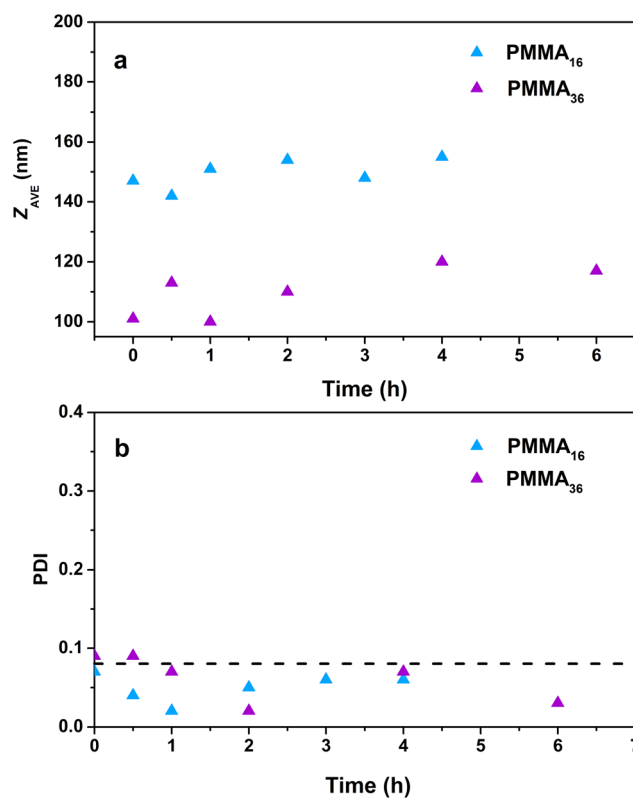


Fig. 10 Evolution of the (a) particle size and (b) polydispersity index with polymerization time for polymer latexes obtained during seeded emulsion polymerization of ethylene with (a) PMMA₁₆ and (b) PMMA₃₆, at 80 °C and 100 bar of ethylene pressure (respectively runs 3 and 4, Table 2).

mostly polydisperse (Fig. 11a and Fig. 12a, respectively), explaining the discrepancies between the particle diameters determined by DLS and TEM. As they are initially polydisperse, it is difficult to conclude on any secondary nucleation that could occur during ethylene polymerization, especially when



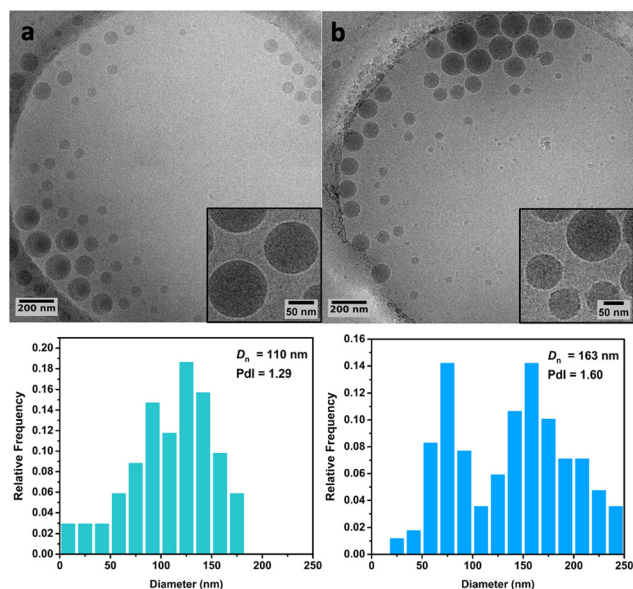


Fig. 11 Cryo-TEM images and particle size distributions of (a) the PMMA₁₆ seed particles and (b) of the particles obtained after 3 h of polymerization at 80 °C under 100 bar of ethylene pressure with PMMA₁₆ seed (run 3-*t*₄, Table 2).

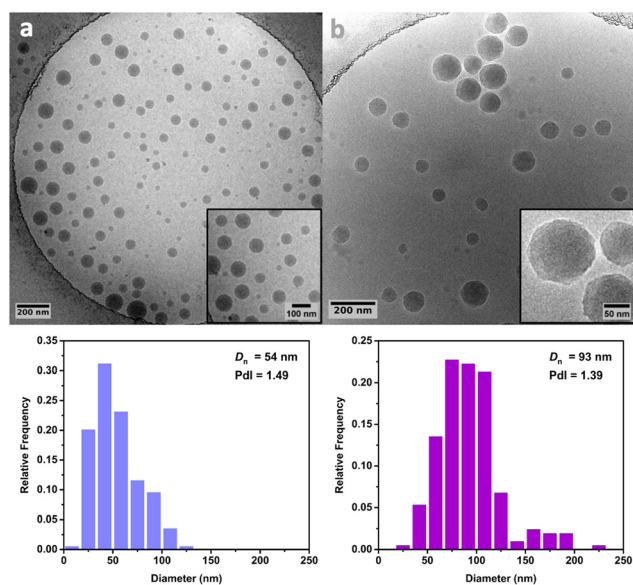


Fig. 12 Cryo-TEM images and particle size distributions of (a) the PMMA₃₆ seed particles and (b) the particles obtained after 4 h of polymerization at 80 °C under 100 bar of ethylene pressure with PMMA₃₆ seed (run 4-*t*₄, Table 2).

PE homopolymer starts to form. However, the particle surface seems to change when PE is formed, being more heterogeneous with some facets (Fig. 11b and 12b). This type of surface was already observed for PE particles by TEM in our group,^{50,55} when PE lamellas prohibit the formation of perfect spheres. However, we do not observe any phase separation

with PE excrescences as already observed when PE was polymerized from PS seeds.⁵⁶

Solids content effect

In our previous study polymerizations were performed in DMC,⁴⁸ and it was shown that the amount of methacrylic oligomers added greatly affected the yield and thus the molar mass of the PE block formed. Thus, this part aims to study the behavior of our system when we increase the initial solids content ($\tau_{s,i}$) of the PMMA₁₆ seeds from 5 to 10 wt% while keeping all the other parameters constant (run 5, Table 2).

Increasing $\tau_{s,i}$ to 10 wt% still led to the formation of a stable latex, and neither the particle size nor the polydispersity index are affected by this change. But the polymerization degree of PE and thus the molar mass are. Indeed, after 4 h of polymerization DP_{PE} decreases from 26 down to 3. In DMC, when polymerizations were carried out with a molar ratio of $[PMMA_{11}]:[AIBN]:[ethylene] = 3:1:1900$ instead of $1:1:1900$, a drastic drop in yield but also in DP_{PE} was observed (after 6 h, DP_{PE} of 221 down to 44 for $[PMMA_{11}]:[AIBN]$ ratios increasing from 1 to 3).⁴⁸ Thus, our results seem to be consistent with the results obtained in DMC and a degradative chain transfer role of PMMA_n in ethylene emulsion polymerization. Indeed, by increasing the solids content, although the overall amount of vinyl functions is higher, the concentration of double bonds per particle, where the polymerization takes place, remains the same. For the same concentration of initiator, the time needed to consume the double bonds increases. Thus, after 4 hours, homopolymerization of ethylene has consumed more ethylene monomer at 5 wt% solids and the chains will then be longer in that case. In brief, when increasing the solids content, shorter PE chains are produced before being terminated reflecting in a decrease of DP_{PE} .

Polymerization of ethylene in the presence of a PMMA seed synthesized by aqueous radical emulsion polymerization

With the aim of comparing the behavior of the PMMA seeds obtained by CCTP to a PMMA seed obtained by free radical emulsion polymerization in water, a PMMA latex was synthesized under conventional emulsion polymerization conditions, using SDS as surfactant and KPS as thermal initiator (PMMA-FRP, Table S1†). The conversion of MMA was almost complete (96%) and a stable latex was obtained.

This latex was then diluted to set the initial solids content at 5 wt% and used as a seed in ethylene polymerization conducted at 80 °C, under 100 bar of ethylene for 4 h (run 6, Table 2). All of the parameters were kept constant between this reaction and run 1 (Table 2).

The increase of solids content with PMMA-FRP seed is significantly higher than with the PMMA₁₆ CCTP seed (9.8 wt% with PMMA-FRP against 2.4 wt% with PMMA₁₆). The latex remained stable and the particle size increased from 42 nm for the seeds to 68 nm for the final particles. It is worth mentioning that, during a MMA free radical polymerization, disproportionation reactions occur leading to around half of the



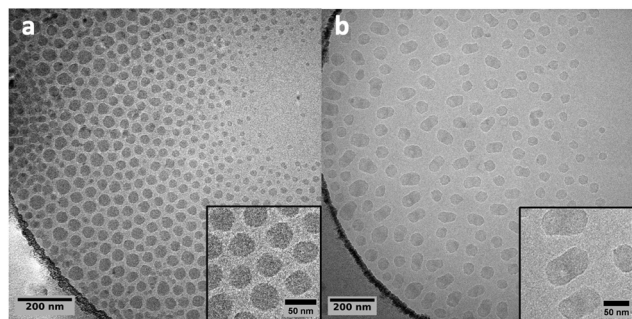


Fig. 13 Cryo-TEM images of (a) PMMA-FRP particles (Table S2†) and (b) the particles obtained after 4 h of polymerization at 80 °C under 100 bar with PMMA-FRP seed (run 6, Table 2).

chains carrying an unsaturated chain ends.⁵⁷ Thus, this seed cannot be strictly speaking considered as a real “blank” experiment without double bonds. Furthermore, the molar mass of the chains for PMMA-FRP seed is much higher than the one for the PMMA₁₆ seed ($M_n = 138\,000\text{ g mol}^{-1}$ and $M_n = 1600\text{ g mol}^{-1}$, respectively). So, for a fixed initial solids content, there are fewer double bonds in the PMMA-FRP seed. Accordingly, the probability/efficiency for an ethylene oligomer to terminate on a double bond is probably lower with these FRP seeds than with the PMMA₁₆ explaining the above obtained results.

This experiment was mainly realized to compare the morphology of the obtained particles to those obtained from PMMA seeds produced by CCTP. Fig. 13 shows the cryoTEM images of the PMMA-FRP seed and of the particles obtained after 4 h of ethylene polymerization. The morphology of the latter looks elongated as made of two incompatible phases that would separate. This morphology is in sharp contrast with that of the particles obtained when using PMMA-CCTP seeds (Fig. 11 and 12) and for which the chemical link between PMMA and PE blocks seems to prevent phase separation.

Conclusions

The seeded radical emulsion polymerization of ethylene was performed using ω -unsaturated PMMA oligomer latexes synthesized by CCTP. In each case, stable latexes were produced. Thus, this chemistry previously performed in organic DMC solution was successfully transposed to a polymerization in aqueous dispersed media and yielding stable latexes. As in DMC, the chain end double bonds carried by PMMA_n synthesized by CCTP are quickly consumed while ethylene is polymerized. For short reaction times, PMMA-*b*-PE block copolymers were produced directly in the particles, while for longer polymerization times PE homopolymer was additionally formed. Depending on the solids content of the seed latex, the molar mass of the PE block could be adjusted. Cryo-TEM analyses of the final latex confirmed that the chains in the particles do not phase separate. The synthesis of PMMA-*b*-PE copolymers in emulsion polymerization, a widely used indus-

trial process, is thus now easily accessible by simply post-polymerization treatment of ω -unsaturated PMMA oligomer latexes synthesized industrially by CCTP for many years. Such copolymers may be useful for compatibilization. The strategy should be easily transposable to other kind of ω -unsaturated oligomers synthesized by CCTP, and thus enable the formation of new PE-based block copolymers in emulsion. To the best of our knowledge, the synthesis of such aqueous copolymer latexes is unprecedented.

Conflicts of interest

There are no conflicts of interest to declare.

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References

- 1 Z. Chen and M. Brookhart, *Acc. Chem. Res.*, 2018, **51**, 1831–1839.
- 2 C. J. Kay, P. D. Goring, C. A. Burnett, B. Hornby, K. Lewtas, S. Morris, C. Morton, T. McNally, G. W. Theaker, C. Waterson, P. M. Wright and P. Scott, *J. Am. Chem. Soc.*, 2018, **140**, 13921–13934.
- 3 C. Dommange, F. D'Agosto and V. Monteil, *Angew. Chem., Int. Ed.*, 2014, **53**, 6683–6686.
- 4 A. Wolpers, C. Bergerbit, B. Ebeling, F. D'Agosto and V. Monteil, *Angew. Chem., Int. Ed.*, 2019, **58**, 14295–14302.
- 5 A. Wolpers, F. Baffie, V. Monteil and F. D'Agosto, *Macromol. Rapid. Commun.*, 2021, **42**, 2100270.
- 6 Y. Nakamura, B. Ebeling, A. Wolpers, V. Monteil, F. D'Agosto and S. Yamago, *Angew. Chem., Int. Ed.*, 2018, **57**, 305–309.
- 7 A. Wolpers, F. Baffie, L. Verrieux, L. Perrin, V. Monteil and F. D'Agosto, *Angew. Chem., Int. Ed.*, 2020, **59**, 19304–19310.
- 8 F. Baffie, M. Lansalot, V. Monteil and F. D'Agosto, *Polym. Chem.*, 2022, **13**, 2469–2476.
- 9 C. Bergerbit, F. Baffie, A. Wolpers, P.-Y. Dugas, O. Boyron, M. Taam, M. Lansalot, V. Monteil and F. D'Agosto, *Angew. Chem.*, 2020, **132**, 10471–10476.
- 10 J. Delorme, O. Boyron, P.-Y. Dugas, P.-E. Dufils, D. J. Wilson, V. Monteil, F. D'Agosto and M. Lansalot, *Polym. Chem.*, 2020, **11**, 7410–7420.
- 11 A. A. Gridnev and S. D. Ittel, *Chem. Rev.*, 2001, **101**, 3611–3660.
- 12 S. Slavin, K. McEwan and D. M. Haddleton, in *Polymer Science: A Comprehensive Reference*, Elsevier, 2012, pp. 249–275.



- 13 D. M. Haddleton, E. Depaquis, E. J. Kelly, D. Kukulj, S. R. Morsley, S. A. F. Bon, M. D. Eason and A. G. Steward, *J. Polym. Sci., Part A: Polym. Chem.*, 2001, **39**, 2378–2384.
- 14 D. M. Haddleton, D. R. Maloney, K. G. Suddaby Adam Clarke and S. N. Richards, *Polymer*, 1997, **38**, 6207–6217.
- 15 W. H. Moser, G. D. Joly, L. R. Krepski, B. N. Gaddam, A. S. Abuelyaman, B. D. Craig, T. D. Dunbar, C. Cao, J. D. Oxman, A. Falsafi and H. T. Bui, United States, US20150238389A1, 2015.
- 16 G. D. Joly, A. S. Abuelyaman, R. S. Davidson, T. D. Jones, B. N. Gaddam and S. J. Moench, World Intellectual Property Organization, WO2014116461A1, 2014.
- 17 W. H. Moser, A. R. Fornof, G. D. Joly, S. Yurt, L. R. Krepski, A. S. Abuelyaman, A. Falsafi, B. D. Craig, J. D. Oxman and B. N. Gaddam, World Intellectual Property Organization, WO2016133668A1, 2016.
- 18 W. H. Moser, A. R. Fornof, G. D. Joly, S. Yurt, L. R. Krepski, A. S. Abuelyaman, A. Falsafi, B. D. Craig, J. D. Oxman and B. N. Gaddam, United States, US20200031961A1, 2020.
- 19 G. D. Joly, L. R. Krepski, A. R. Fornof, S. Yurt, B. N. Gaddam and A. S. Abuelyaman, United States, US20150203658, 2015.
- 20 N. R. Bagnall, M. H. Jones, G. C. Jernigan, C. Routt, L. C. Dar and B. T. Worrell, *J. Am. Chem. Soc.*, 2023, **145**, 14202–14207.
- 21 D. Kukulj, T. P. Davis, K. G. Suddaby, D. M. Haddleton and R. G. Gilbert, *J. Polym. Sci., Part A: Polym. Chem.*, 1997, **35**, 859–878.
- 22 D. Kukulj, T. P. Davis and R. G. Gilbert, *Macromolecules*, 1997, **30**, 7661–7666.
- 23 K. G. Suddaby, D. M. Haddleton, J. J. Hastings, S. N. Richards and J. P. O'Donnell, *Macromolecules*, 1996, **29**, 8083–8091.
- 24 D. M. Haddleton, D. R. Morsley, J. P. O'Donnell and S. N. Richards, *J. Polym. Sci., Part A: Polym. Chem.*, 1999, **37**, 3549–3557.
- 25 S. A. F. Bon, D. R. Morsley, J. Waterson, D. M. Haddleton, M. R. Lees and T. Horne, *Macromol. Symp.*, 2001, **165**, 29–42.
- 26 S. C. J. Pierik, B. Smeets and A. M. van Herk, *Macromolecules*, 2003, **36**, 9271–9274.
- 27 P. B. Zetterlund and D. R. D'hooge, *Macromolecules*, 2019, **52**, 7963–7976.
- 28 N. M. B. Smeets, J. P. A. Heuts, J. Meuldijk, M. F. Cunningham and A. M. van Herk, *Macromolecules*, 2009, **42**, 7332–7341.
- 29 N. M. B. Smeets, J. P. A. Heuts, J. Meuldijk, M. F. Cunningham and A. M. van Herk, *Macromolecules*, 2009, **42**, 6422–6428.
- 30 N. M. B. Smeets, J. P. A. Heuts, J. Meuldijk and A. M. van Herk, *J. Polym. Sci., Part A: Polym. Chem.*, 2008, **46**, 5839–5849.
- 31 A. A. Gridnev, *Polym. J.*, 1992, **24**, 613–623.
- 32 G. Patias, A. M. Wemyss, S. Efstathiou, J. S. Town, C. J. Atkins, A. Shegiwal, R. Whitfield and D. M. Haddleton, *Polym. Chem.*, 2019, **10**, 6447–6455.
- 33 D. J. Krasznai, T. F. L. McKenna, M. F. Cunningham, P. Champagne and N. M. B. Smeets, *Polym. Chem.*, 2012, **3**, 992–1001.
- 34 N. G. Engelis, A. Anastasaki, G. Nurumbetov, N. P. Truong, V. Nikolaou, A. Shegiwal, M. R. Whittaker, T. P. Davis and D. M. Haddleton, *Nat. Chem.*, 2017, **9**, 171–178.
- 35 L. Nurmi, J. Lindqvist, R. Randev, J. Syrett and D. M. Haddleton, *ChemComm*, 2009, 2727–2729.
- 36 C. J. Atkins, G. Patias, J. S. Town, A. M. Wemyss, A. M. Eissa, A. Shegiwal and D. M. Haddleton, *Polym. Chem.*, 2019, **10**, 646–655.
- 37 P. Cacioli, D. G. Hawthorne, R. L. Laslett, E. Rizzardo and D. H. Solomon, *J. Macromol. Sci., Part A: Pure Appl. Chem.*, 1986, **23**, 839–852.
- 38 L. M. Muratore, K. Steinhoff and T. P. Davis, *J. Mater. Chem.*, 1999, **9**, 1687–1691.
- 39 B. Yamada, P. B. Zetterlund and E. Sato, *Prog. Polym. Sci.*, 2006, **31**, 835–877.
- 40 B. Yamada and S. Kobatake, *Prog. Polym. Sci.*, 1994, **19**, 1089–1131.
- 41 B. Yamada, F. Oku and T. Harada, *J. Polym. Sci., Part A: Polym. Chem.*, 2003, **41**, 645–654.
- 42 G. Nurumbetov, N. Engelis, J. Godfrey, R. Hand, A. Anastasaki, A. Simula, V. Nikolaou and D. M. Haddleton, *Polym. Chem.*, 2017, **8**, 1084–1094.
- 43 L. Hutson, J. Krstina, C. L. Moad, G. Moad, G. R. Morrow, A. Postma, E. Rizzardo and S. H. Thang, *Macromolecules*, 2004, **37**, 4441–4452.
- 44 J. Krstina, G. Moad, E. Rizzardo, C. L. Winzor, C. T. Berge and M. Fryd, *Macromolecules*, 1995, **28**, 5381–5385.
- 45 A. Shegiwal, A. M. Wemyss, E. Liarou, J. Town, G. Patias, C. J. Atkins, A. Marathianos, D. W. Lester, S. Efstathiou and D. M. Haddleton, *Eur. Polym. J.*, 2020, **125**, 109491.
- 46 I. Schreur-Piet and J. P. A. Heuts, *Polym. Chem.*, 2017, **8**, 6654–6664.
- 47 L. Chen, L. Yan, Q. Li, C. Wang and S. Chen, *Langmuir*, 2010, **26**, 1724–1733.
- 48 F. Baffie, G. Patias, A. Shegiwal, F. Brunel, V. Monteil, L. Verrieux, L. Perrin, D. M. Haddleton and F. D'Agosto, *Angew. Chem., Int. Ed.*, 2021, **60**, 25356–25364.
- 49 A. Shegiwal, A. M. Wemyss, M. A. J. Schellekens, J. de Bont, J. Town, E. Liarou, G. Patias, C. J. Atkins and D. M. Haddleton, *J. Polym. Sci., Part A: Polym. Chem.*, 2019, **57**, E1–E9.
- 50 G. Billuart, E. Bourgeat-Lami, M. Lansalot and V. Monteil, *Macromolecules*, 2014, **47**, 6591–6600.
- 51 E. J. Bradbury, D. McNulty, R. I. Savage and E. E. McSweeney, *Ind. Eng. Chem.*, 1952, **44**, 211–212.
- 52 I. A. Maxwell, B. R. Morrison, D. H. Napper and R. G. Gilbert, *Macromolecules*, 1991, **24**, 1629–1640.
- 53 T. Williams and I. M. Ward, *J. Polym. Sci., Part B: Polym. Lett.*, 1968, **6**, 621–624.
- 54 T. G. Fox and P. J. Flory, *J. Polym. Sci.*, 1954, **14**, 315–319.
- 55 E. Grau, P.-Y. Dugas, J.-P. Broyer, C. Boisson, R. Spitz and V. Monteil, *Angew. Chem., Int. Ed.*, 2010, **49**, 6810–6812.
- 56 G. Billuart, PhD thesis, Université de Lyon, 2015.
- 57 G. Ayrey and C. G. Moore, *J. Polym. Sci.*, 1959, **36**, 41–53.

