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Poly(ethylene glycol)-*b*-poly(vinyl acetate) block copolymer particles of various morphologies via RAFT/MADIX aqueous emulsion PISA

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The polymerization-induced self-assembly (PISA) of amphiphilic diblock copolymers of poly(ethylene glycol)-*b*-poly(vinyl acetate) (PEG-*b*-PVAc) in water was achieved through macromolecular design via interchange of xanthate (MADIX) polymerization in emulsion. Poly(ethylene glycol) with a dithiocarbonate (xanthate) end-group (PEG-X) was synthesized by post-modification of commercial poly(ethylene glycol). The chain extension of PEG-X was then conducted in water using vinyl acetate (VAc) to produce PEG-*b*-PVAc amphiphilic diblock copolymers. Their *in situ* self-assembly led to stable latexes particles. The molar mass of the PVAc block was varied to trigger the formation of various morphologies including spheres, vesicles and large compound vesicles.

Introduction

Polymerization-induced self-assembly (PISA)^{1,2} is a concept that takes advantage of a living polymerization technique to chain extend preformed solvophilic polymer chains with either solvophobic (emulsion) or solvophilic (dispersion) monomer units. In both cases, the formed second block is however solvophobic, and simultaneously to its growth the resulting block copolymers can self-assemble into nanoparticles. PISA has been largely employed in the recent years to produce block copolymer nanoparticles in water, either in dispersion or emulsion conditions, using mainly reversible addition-fragmentation chain transfer (RAFT).² In addition, by varying the nature and molar mass of the two blocks and/or the solids content, different morphologies such as spheres, worms or vesicles, already depicted when self-assembling preformed block copolymers in a selective solvent of one of the two blocks,³ can also be produced by PISA in water up to high solids content.⁴

While these morphologies seem to be easily obtained in aqueous dispersion polymerization by simply varying the molar mass of the hydrophobic block or the targeted solids content,⁵ emulsion polymerization reveals to be more challenging to reach higher-order morphologies different from spheres.⁶⁻¹⁹ As now generally accepted, these morphologies result from the fusion of the original block copolymer spherical micelles and further chain rearrangements.²⁰ In dispersion polymerization, the core monomer is soluble in water and swells its polymer,²¹

which favors fusion and rearrangement events. In emulsion polymerization,²² although the core monomer is also able to swell its polymer, its hydrophobic character induces the presence of monomer droplets. A small fraction is also dissolved in water. Therefore, in emulsion PISA, once the first block copolymer micelles are formed the monomer has to diffuse from these droplets through the water phase to feed the particles with monomer. This diffusion is thus allowed but also limited by the solubility of the monomer in water, which directly impacts the plasticization of the particles. This simple observation naturally led researchers to consider that hydrophobic monomers exhibiting high water solubility should favor the fusions and rearrangements of the chains mentioned above, and thus the formation of various morphologies by PISA in aqueous emulsion. This was effectively shown when using methyl methacrylate (MMA) instead of styrene (respective water solubilities at 50 °C: 15 g L⁻¹²³ vs. 0.45 g L⁻¹²⁴) in a RAFT-mediated PISA system using poly(methacrylic acid-co-poly(ethylene glycol) methyl ether methacrylate) as hydrophilic macromolecular chain transfer agent (macroCTA). Under exactly the same conditions (pH 3.5), emulsion polymerization of styrene systematically led to spheres whatever the targeted degree of polymerization (DP, from 150 to 570) of the PS block,²⁵ while the use of MMA allowed the formation of spheres, fibres and vesicles for DP of 200, 300 and 500, respectively.⁹ This strategy was recently confirmed by Armes and coworkers who also obtained non-spherical morphologies by polymerizing in emulsion 4-hydroxybutyl methacrylate (HBMA, 20 g L⁻¹ at 70 °C),¹³ glycidyl methacrylate (GlyMA, 24–25 g L⁻¹ at 80 °C),¹⁴ and 2-methoxyethyl methacrylate (MOEMA, 19.6 g L⁻¹ at 70 °C).¹⁶

With a solubility in water of 26 g L⁻¹ at 50 °C,²⁶ vinyl acetate (VAc) is one of the most hydrophilic monomers among the monomers classically employed in aqueous emulsion polymerization, and

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poly(vinyl acetate) (PVAc) latexes find industrial applications in various domains such as adhesives, paints, and coatings.²⁷ The particles obtained in this process are systematically spherical. Taking advantage of the best attributes of PISA for VAc emulsion polymerization is thus very appealing in order to generate new types of non-spherical PVAc particles. Vinyl acetate is a less activated monomer and its RAFT polymerization requires the use of specific macroCTA carrying dithiocarbamate or dithiocarbonate (xanthate, X) chain ends.²⁸ In this last case, the polymerization process is termed macromolecular design via interchange of xanthate (MADIX). To the best of our knowledge, only five papers depict the use of MADIX-mediated PISA for less activated monomers.^{17, 29–32} Two of them focus on VAc emulsion polymerization, conducted with dithiocarbonate end functionalized macroCTAs. In the first case, a dithiocarbonate end functionalized dextran (Dext-X) was used.²⁹ The functionalization rate was rather low (ca. 30%), and when employed to mediate emulsion polymerization of VAc, spherical particles were obtained. Their formation and stabilization were possible by a fraction of in situ formed dextran-*b*-PVAc block copolymers. In the second study, performed in our group, emulsion polymerization of VAc was carried out in the presence of a PEG quantitatively functionalized with a dithiocarbonate end group (PEG-X).³⁰ Using a redox initiation at 25 °C, kinetics was slow, but the control of the polymerization was shown by the increase of the molar mass upon VAc conversion and low dispersities that remained around 2, as expected. Here again, however, spherical PVAc particles composed of PEG-*b*-PVAc were systematically observed. The choice of the low temperature initiating conditions was conditioned by the comparative use in the same paper of poly(*N*-acryloylmorpholine) and poly(*N*-vinylpyrrolidone) macroCTAs, the thermal instability of the latter having been demonstrated earlier.³³

Building on these results, we anticipated that conducting PEG-X mediated VAc polymerization at higher temperature may not only lead to faster kinetics but also favor the softening of the particles considering both the glass transition temperature (T_g) of PVAc (ca. 30 °C)³⁴ and the improved plasticization with VAc. In fine, this could favor fusions and rearrangements of the polymer chains to achieve the yet unattained higher-order morphologies.

Emulsion polymerizations of VAc were thus conducted in the presence of PEG-X at 70 °C using 2,2'-azobis(2-amidinopropane) dihydrochloride (AIBA) as initiator at a solids content of 10 wt%, targeting different DP for the PVAc block (DP_{target} from 50 to 200). A systematic characterization of the macromolecular and colloidal features of the resulting latexes was performed. Particular attention was paid to the development of robust systems leading to higher-order morphologies.

Materials and Methods

Materials.

Vinyl acetate (VAc, 99%, Aldrich), 2,2'-azobis(2-amidinopropane) dihydrochloride (AIBA, 97%, Aldrich),

poly(ethylene glycol) methyl ether (PEG-OH, M_n = 2000 g mol⁻¹, Aldrich), *O*-ethyl xanthic acid potassium salt (98%, Alfa Aesar), triethylamine (99%, Aldrich), 2-bromopropionyl bromide (97%, Aldrich), dichloromethane (ACS reagent, Aldrich), ammonium chloride (99.5%, ACS reagent, Aldrich), hydrogen carbonate (ACS reagent, Aldrich), sodium chloride (99%, ACS reagent, Aldrich), anhydrous magnesium sulfate (99.5%, Alfa Aesar) were used as received. Water was deionized before use (Purelab Classic UV, Elga LabWater).

Experimental procedures

Preparation of PEG-X

Synthesis of 2-bromopropionic acid [poly(ethylene glycol) methyl ether] (PEG-Br). Poly(ethylene glycol) methyl ether (40.1 g, 2.0×10^{-2} mol) was placed in a three-neck round bottom flask and stirred with triethylamine (8 mL, 5.9×10^{-2} mol) in dichloromethane (100 mL). The mixture was immersed in an ice bath and the 2-bromopropionyl bromide (5 mL, 4.8×10^{-2} mol) was added dropwise. The mixture was stirred at room temperature overnight. The white precipitate was filtered off. The resulting solution was washed with saturated ammonium chloride (2 × 50 mL), saturated hydrogen carbonate (2 × 50 mL), water (2 × 50 mL), saturated sodium chloride (2 × 50 mL) and eventually dried over anhydrous magnesium sulfate. The solvent was evaporated under vacuum. The polymer was recovered by precipitation in diethyl ether, yielding a white powder.

Synthesis of PEG-X. PEG-Br (30 g, 1.4×10^{-2} mol) was dissolved in dichloromethane (100 mL) in a 250 mL round bottom flask placed in an oil bath. *O*-ethyl xanthic acid potassium salt (4.85 g, 3.0×10^{-2} mol) was added portion-wise. The mixture was stirred at room temperature overnight. The precipitate was filtered off. The resulting solution was washed with saturated ammonium chloride (2 × 50 mL), saturated hydrogen carbonate (2 × 50 mL), water (2 × 50 mL), saturated sodium chloride (2 × 50 mL) and eventually dried over anhydrous magnesium sulfate. The solvent was evaporated under vacuum. The polymer was recovered by precipitation in diethyl ether, yielding a white powder that was characterized by ¹H and ¹³C nuclear magnetic resonance (See Figures S1–S3 in ESI).

General procedure for the emulsion polymerization of VAc in the presence of PEG-X.

All emulsion polymerizations were performed using the following protocol. VAc, PEG-X, AIBA and deionized water were introduced in a 50 mL one-neck round bottom flask, which was immersed in an ice bath. After degassing of the medium for 15 min under argon, the flask was placed in an oil bath previously heated at 70 °C. After 90 min under stirring at about 400 rpm, the reaction was quenched by immersion of the flask in an ice bath and exposure to air. VAc consumption was followed by gravimetric analysis of samples withdrawn from the medium at different times.

Analytical techniques

Nuclear magnetic resonance (NMR). ¹H and ¹³C spectra were recorded on a Bruker Avance III 400 MHz spectrometer (¹H: 400

MHz; ^{13}C : 100 MHz) at 25 °C. The samples were diluted in CDCl_3 at concentrations of about 30 mg mL^{-1} for ^1H and 90 mg mL^{-1} for ^{13}C . NMR spectra were recorded with a 5 mm BBFO+ probe with a z-gradient coil.

Size exclusion chromatography (SEC). SEC 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 °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 Polymer Standard Service columns (SDVB, 5 μm , 300 x 7.5 mm) and a guard column. The average molar masses (number-average molar mass, M_n and weight-average molar mass, M_w) and the molar mass dispersity, $\bar{D} = M_w/M_n$, were calculated from the RI signal with a calibration curve based on polystyrene standards (Polymer Standards Service).

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

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 in the Philips CM120 microscope operating at an accelerating voltage of 120 kV (Centre Technologique des Microstructures (CTμ), platform of the Université Claude Bernard Lyon 1, Villeurbanne, France). The number-average diameter (D_n), the weight-average diameter (D_w) and the diameter dispersity (D_w/D_n) were determined from the TEM images using the ImageJ software and the following equations:

$$D_n = \frac{\sum_i n_i \times D_i}{\sum_i n_i} \text{ and } D_w = \frac{\sum_i n_i \times D_i^4}{\sum_i n_i \times D_i^3}$$

Results and discussion

PEG-X mediated emulsion polymerization of VAc for a DP_{target} of 51

As mentioned in the introduction, PEG-*b*-PVAc spherical particles were already successfully obtained through PEG-X mediated VAc emulsion polymerization in water at 25 °C.³⁰ Building on those preliminary results, a new set of experiments was implemented using an azo initiator (AIBA), setting the temperature at 70 °C, and targeting different DP for the PVAc segment. The solids content was set to 10 wt%.

Table 1. Experimental conditions and results for the aqueous emulsion polymerisation of VAc in presence of PEG-X macroCTA.^a

Exp	DP_{target}^b	t (min)	τ^{th} (wt%) ^c	τ (wt%) ^c	Conv. (%) ^d	DP_{exp}^e	M_n^{th} (g mol^{-1}) ^f	M_n^{SEC} (g mol^{-1}) ^g	\bar{D}^h	M_n^{NMR} (g mol^{-1}) ⁱ	Z_{ave} (nm) Pdl ^j	Morphology ^k (D_n (nm)/ D_w/D_n)
A1	51	80	10.4	9.8	93	47	6400	5460	1.5	6300	22 0.05	S (15.9/1.04)
A2	62	120	10.4	9.8	95	59	7400	8300	1.7	7230	65 0.50	S+V
A3	75	125	10.2	9.6	94	71	8400	7930	1.8	8130	113 0.21	S+V
A4	100	130	10.0	9.4	92	92	10200	9450	2.1	11800	203 0.16	V
A5	152	80	10.2	9.7	95	144	14700	14360	2.1	14450	197 0.07	V ^l
A6	199	90	10.1	9.1	94	187	18400	17920	2.3	18270	-	LCV
B1	51	80	15.4	14.8	95	48	6400	6930	1.5	6300	-	S (17.3/1.04)
B2	76	90	15.1	14.9	97	73	8600	8800	1.8	7780	-	S+V(+W)
B3	100	80	15.0	14.6	97	97	10600	10730	1.9	11200	-	V

^a All experiments were performed at 70 °C with $[\text{PEG-X}]/[\text{AIBA}] = 3$. The final pH was around 3 for each latex (for an initial pH ranging between 4.5 and 5. The slight decrease of the pH value can likely be attributed to formation of some acetic acid resulting from hydrolysis of VAc, which is however very limited at pH ca. 4.5³⁴). ^b DP_{target} is defined by the experimental ratio: $[\text{PEG-X}]/[\text{VAc}]$. ^c Theoretical solids content (τ^{th}) and experimental final solids content measured by gravimetric analysis (τ). ^d Conversion calculated from the solids content. ^e Experimental DP calculated from the conversion. ^f Theoretical number-average molar mass of the block copolymer calculated using the conversion and the targeted DP. ^g Experimental number-average molar mass of the block copolymer determined by SEC-THF using a calibration based on polystyrene standards. ^h Dispersity of the block copolymers, M_w/M_n , determined by SEC-THF. ⁱ Experimental number-average molar mass of the block copolymers determined by ^1H NMR. ^j Intensity-average diameter and polydispersity of the final latex from DLS. ^k Main Morphologies based on cryo-TEM images on the final latex: S = spheres, W = worms, V = vesicles, LCV = Large compound vesicles. For the spheres, the number-average diameter and dispersity index determined by statistical analysis of the TEM images are also indicated (Figure S17). ^l For this experiment, multilamellar vesicles were also observed (see A5 cryo-TEM image, Figure 6).

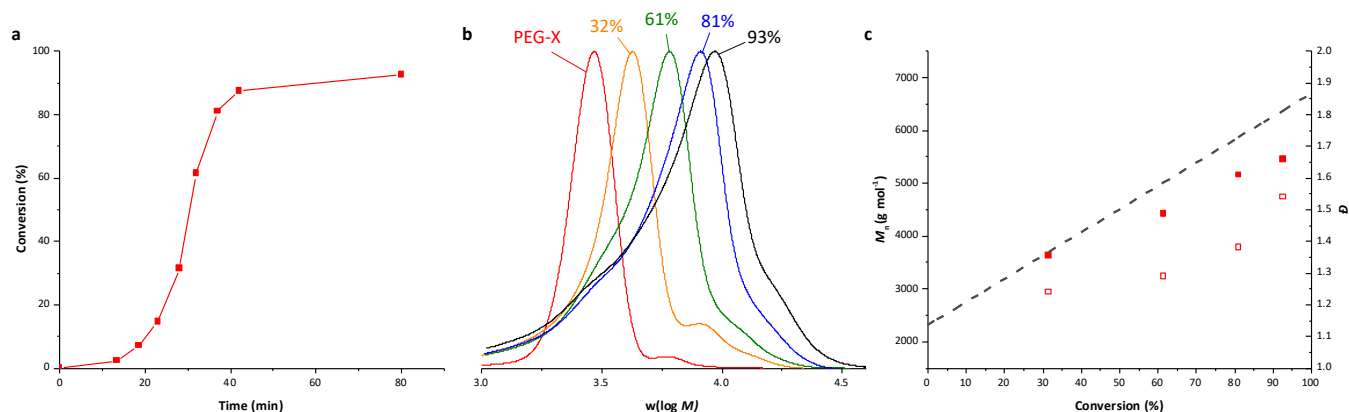


Figure 1. PEG-X mediated VAc emulsion polymerization for $DP_{\text{target}} = 51$ (entry A1, Table 1). a) Evolution of conversion versus time, b) SEC traces (RI detection) for samples withdrawn during the polymerization. c) Corresponding evolution of the number-average molar masses M_n (plain squares) and dispersity ($D = M_w/M_n$, hollow squares) with VAc conversion (the dashed line corresponds to the theoretical M_n values).

A first experiment was conducted targeting a DP of 51 (entry A1, Table 1). As expected, and in strong contrast with our previous works for which 80% conversion was reached in 50 h, the polymerization went to conversions higher than 90% in less than 60 min (Figure 1a). SEC traces of samples withdrawn during the polymerization show the presence of more than one population (Figure 1b). Accordingly, the dispersities are somewhat high and increase with the conversion (Figure 1c).

Two phenomena could explain these results. (i) A shoulder on the high molar mass side of the molar mass distribution (MMD) is visible. It is shifted, together with the whole distribution, towards high molar masses upon VAc conversion. It is worth mentioning here that PEG-X MMD features such a shoulder too, yet of very small intensity (Figure S4). Indeed, this shoulder probably corresponds to dihydroxy telechelic PEG contamination formed as a side product during anionic polymerization of ethylene oxide (traces of water initiation). The presence of these species probably led to the formation of a small amount of dixanthate telechelic PEG after functionalization of PEG-OH (X-PEG-X), visible on the SEC analyses using both RI and UV detections at 270 nm (Figure S4). During the VAc emulsion polymerization, these chains could thus be extended into PVAc-*b*-PEG-*b*-PVAc triblock copolymers and would correspond to the shoulder observed on the high molar mass of the distribution (Figure S5). (ii) The MMD is also featuring a tailing on the low molar mass side. VAc's inclination to produce head-to-head (HH) units during the polymerization is documented in the literature and has been particularly discussed in the case of RAFT/MADIX.^{35–38} The corresponding chains terminated by $-\text{CH}(\text{OAc})-\text{CH}_2-\text{X}$ are much less reactivatable than the expected $-\text{CH}_2-\text{CH}(\text{OAc})-\text{X}$. Indeed, the tailing observed on the low molar mass is thus not ascribed to unreacted PEG-X (see below), although it is worth mentioning that degradation of PEG-X during the reaction or slow reinitiation could contribute to this tailing. Rather, $-\text{CH}(\text{OAc})-\text{CH}_2-\text{X}$ terminated block copolymers probably accumulate. This is indeed consistent with the SEC traces recorded using UV at 270 nm (Figure S5) that show that the chains forming this tailing are also carrying a xanthate moiety. All in all, either (i) or (ii) assumption gives rise to side populations that are nevertheless

amphiphilic block copolymers containing PEG and PVAc segments that will be involved in the self-assembly process.

The ^1H NMR spectrum (Figure 2) performed on the powder obtained after drying the latex confirms the presence of signals at about 3.15 ppm and 4.05 ppm, which are, respectively, characteristics of HH units linked to a xanthate and of dead chains.³⁶

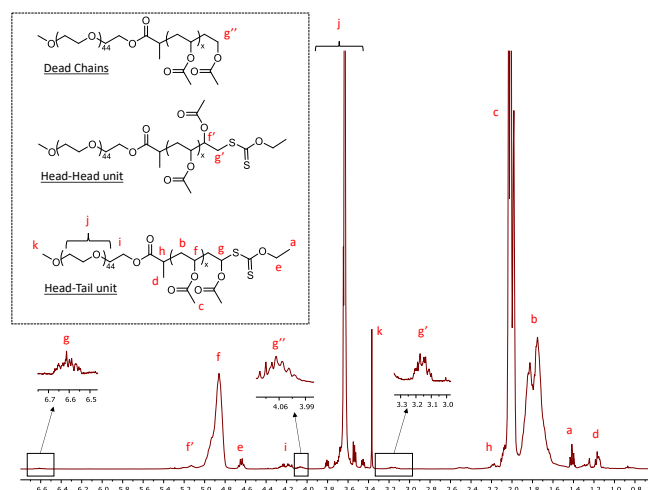


Figure 2. ^1H NMR spectrum (CDCl_3 , 256 scans) of the dried latex obtained from entry A1, Table 1, after 93% conversion.

The full assignment of this ^1H NMR spectrum shows that no trace of remaining PEG-X could be detected indicating that the majority of these chains have been chain extended. The ^1H NMR assignments further allow to calculate the final molar mass of the formed chains by the relative integration of the signal attributed to the protons f at 4.90 ppm compared to the signal attributed to the protons of the methoxy group at 3.37 ppm. It shows an almost perfect fit with the expected theoretical values (Table 1, A1). Those data are additionally consistent with the molar mass data collected from SEC that show a linear increase of the number-average molar masses upon VAc conversion (Figure 1c). A good match between expected and experimental values is obtained although the values obtained using polystyrene calibration might be slightly flawed. Nevertheless,

the consistency between the molar mass values obtained from ^1H NMR, SEC and theory is a good indication that the degradation of PEG-X suspected above is low (if any), and that the majority of the chains have been chain extended during VAc polymerization. Those results allow confirming that the expected amphiphilic block copolymers were formed.

The polymerization A1 in Table 1 visually led to a clear solution (Figure 3, $\text{DP}_{\text{exp}} = 47$). However, as shown by the cryo-TEM image (Figure 4) and DLS results (Table 1), the obtained medium is a latex made of isometric spherical particles with a diameter of ca. 20 nm. Remarkably, it can be noticed that the particle diameter is close to the length of two back-to-back PVAc chains on a regular stretched zigzag conformation (i.e. 23.7 nm, see Figure S17 in ESI for a detailed calculation). It can thus be acknowledged that those spherical particles are made of PEG-*b*-PVAc amphiphilic block copolymers resulting of their direct self-assembly, which is consistent with the expected PISA mechanism.

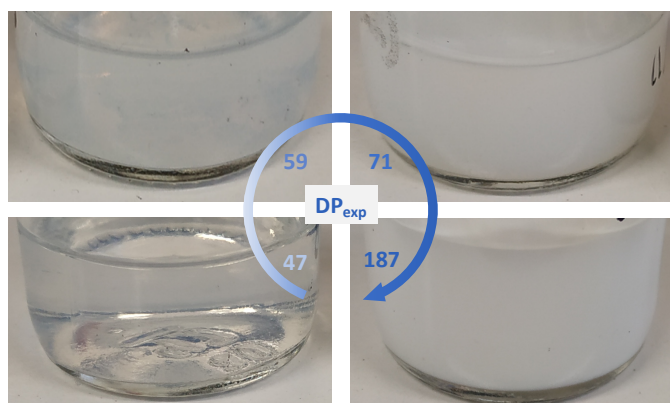


Figure 3. Digital pictures of the crude latexes A1, A2, A3 and A6 (Table 1).

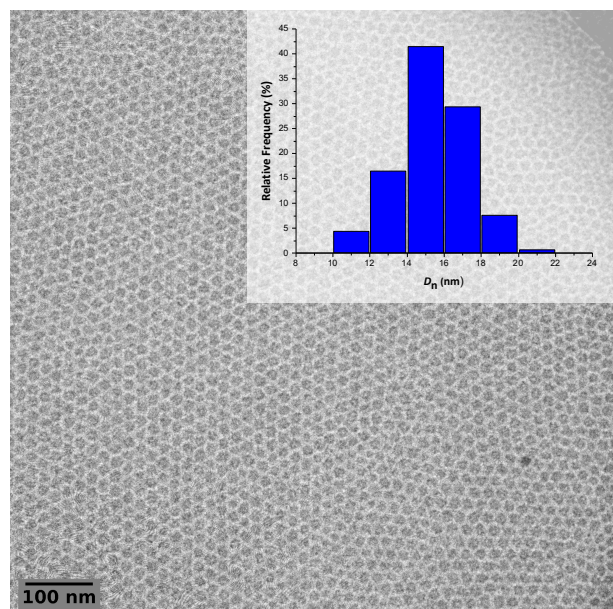


Figure 4. Representative cryo-TEM image of the latex obtained from entry A1 in Table 1 and corresponding particle size distribution of the crude latex.

Influence of the targeted DP of PVAc block

$\text{DP}_{\text{target}}$ was then increased to investigate the impact on the PEG-*b*-PVAc self-assembly. A set of experiments was thus performed varying $\text{DP}_{\text{target}}$ of the PVAc block between 50 to 200 (entries A1 to A6, Table 1). The total solids content was kept at 10 wt%.

For all the aforementioned syntheses, and as exemplary shown in Figure 5a, the polymerization went to almost completion (> 92%) in less than 60 min. The kinetics of the different experiments remain however difficult to compare one with each other in detail since the initiator, monomer and PEG-X concentrations were modified simultaneously to maintain the solids content constant. Nevertheless, the very similar profiles obtained are consistent with a rapid compartmentalization of the reaction within particles (nucleation).

Overall, all the experiments showed a good blocking efficiency as shown by the linear increases of number-average molar masses with the conversion (Figure 5b). Once again, the molar masses obtained using SEC are not fully matching the expected ones probably due to the use of polystyrene standards, which are not well suited for the particular nature of the PEG-*b*-PVAc block copolymers. However, the molar masses calculated according to the ^1H NMR spectra of the final latexes are in good agreement with the theoretical ones (Table 1).

The SEC traces obtained for all these experiments are similar to those of latex A1 (Figures S6-S10). However, the dispersity tends to increase with both the conversion and the targeted degree of polymerization of the PVAc block. This might be related to the above-mentioned side formation of PVAc-*b*-PEG-*b*-PVAc triblock copolymers, the impact of which on the dispersity is probably exacerbated when the DP of the PVAc segment increases.

The obtained latexes showed different aspects depending on the DP_{exp} of the PVAc segment. As mentioned above, the synthesis with the lowest DP (A1, $\text{DP}_{\text{exp}} = 47$) led to a completely transparent dispersion. However, as the DP increases, the latex becomes slightly translucent (A2, $\text{DP}_{\text{exp}} = 59$) before showing the aspect of a classical milky dispersion (A3, $\text{DP}_{\text{exp}} = 71$ to A6, $\text{DP}_{\text{exp}} = 187$) (Figure 3). DLS analyses of these dispersions are quite consistent with these observations, with a particle diameter that increases with the final DP (Table 1). However, as shown by the polydispersity index, the DLS results tend to show that the objects present in the latexes are either very dispersed in size or that not only spherical particles are present.

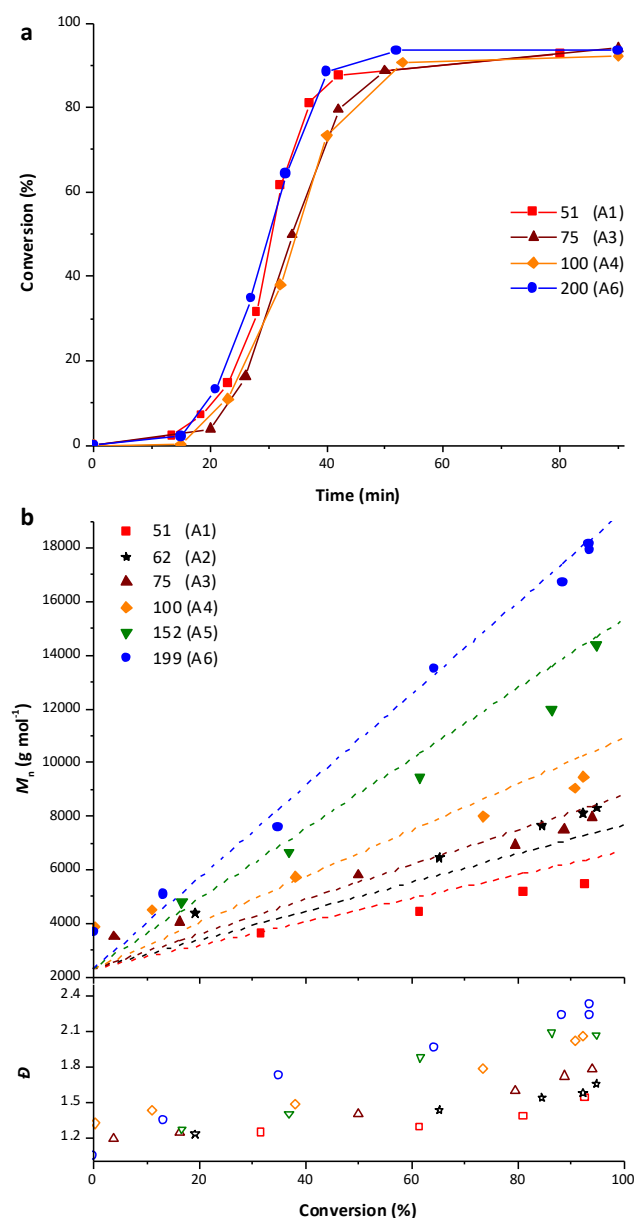


Figure 5. PEG-X mediated VAc emulsion polymerizations when targeting different DPs for the PVAc block. Evolution of a) conversion versus time and b) number-average molar masses M_n (plain squares) and dispersity ($D = M_w/M_n$, hollow squares) with VAc conversion (the dashed lines correspond to the theoretical M_n values). See Table 1 for details.

Further analyses of the latexes by cryo-TEM however show that the in situ formed amphiphilic block copolymers self-assemble into nano-objects of various sizes and morphologies (Figure 6). As the DP of the hydrophobic PVAc block increases, the morphology of the particles shifts from isometric spherical particles (A1, $\text{DP}_{\text{exp}} = 47$) to vesicles (A4, $\text{DP}_{\text{exp}} = 92$) and then to higher-order morphologies (A6, $\text{DP}_{\text{exp}} = 187$). For a DP_{exp} comprised between 59 and 92 (A2, A3 and A4), the latexes are

composed of vesicles and spherical particles, the size and proportion of vesicles increasing with DP_{exp} . Further increase of the hydrophobic block molar mass gives access to higher-order morphologies such as distorted or multilamellar vesicles (A5, $\text{DP}_{\text{exp}} = 144$) and eventually to “large compound vesicles” (LCVs, A6, $\text{DP}_{\text{exp}} = 187$) (see additional cryo-TEM images in Figure S11).

Morphology prediction in PISA is difficult since the mechanisms involved are complex and depend on many parameters. Nonetheless, a reproducible pattern for the morphological transitions has now been settled through various studies.^{2, 20, 39-41} It is agreed that initial spherical particles merge to form worm-like particles which themselves give rise to branched “octopi-like” structures as a transition state to vesicles. The morphologies observed match with this established pattern, and morphologies of even higher-order than vesicles are observed. However, according to this pattern, the worm/fibre morphology that is not seen here should be formed for DP between 47 (A1, spheres) and 59 (A2, spheres + vesicles). The narrowness of this DP window is consistent with the PISA literature concerning the formation of worm/fibre particles.⁴²

Nevertheless, to the best of our knowledge, these experiments are the first depicting the obtention of non-spherical morphologies by VAc PISA in emulsion. Indeed, as mentioned above, the morphologies obtained in emulsion are most of the time restrained to kinetically trapped spheres, the number of reports of higher-order morphologies remaining limited.² The formation of these morphologies in PISA is the result of particle collision/fusion phenomena (vide supra).²⁰ As such, their formation is directly linked with the mobility of the polymer chains. In this specific system, several factors may be involved. Especially, the solubility of VAc in water is most certainly crucial. As mentioned in the introduction, the importance of the monomer solubility in water for the formation of non-spherical objects in PISA emulsion-based systems was recently highlighted. Cockram *et al.* obtained unusual “monkey nut” particles through RAFT-mediated emulsion polymerization in water using a core forming monomer, namely HBMA, with a water solubility higher than those usually employed (20 g L^{-1} at 70°C).¹³ The authors postulated that a higher monomer aqueous solubility would lead to a more efficient plasticization of the hydrophobic core, which would facilitate the rearrangement of the polymer chains. The same group then obtained well-defined worms by emulsion polymerization of GlyMA, which has a similar aqueous solubility ($24\text{--}25 \text{ g L}^{-1}$ at 80°C),¹⁴ and both worms and vesicles with MOEMA (19.6 g L^{-1} at 70°C).¹⁶ With a water solubility of 26 g L^{-1} (at 50°C), VAc provides a system with comparable features. Thus, the results obtained through this work tend to confirm the determining role of this parameter to trigger the formation of particles with higher-order morphologies.

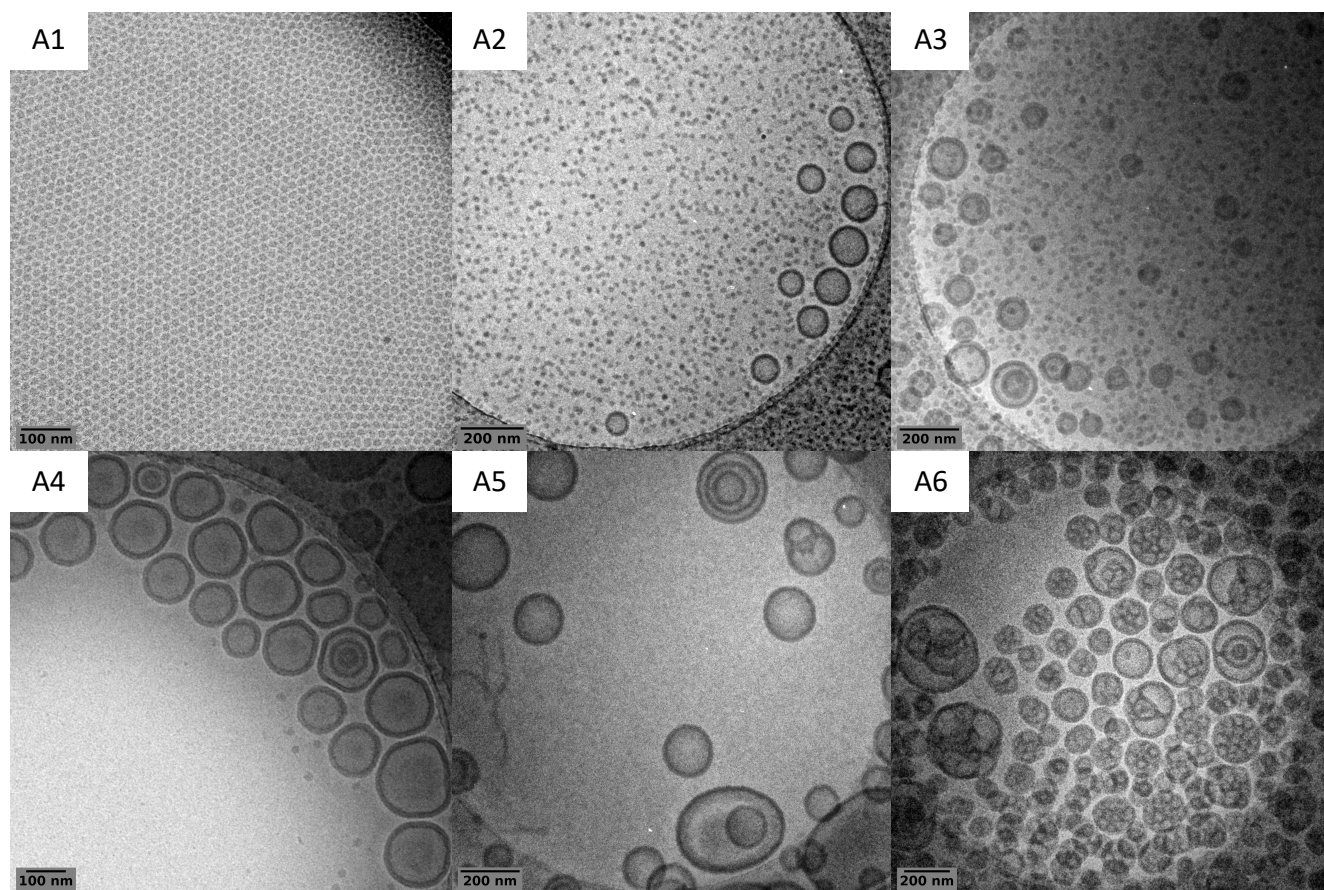


Figure 6. Representatives cryo-TEM images of the crude latexes A1-A6 (Table 1).

Influence of the solids content

To investigate further this system and identify conditions for which worms/fibres could be obtained, three additional syntheses were performed at a higher solids contents, that is 15 wt%, targeting a DP of 51, 76 and 100 for the PVAc block (Entries B1, B2 and B3 in Table 1, respectively). The same good control (Figures S12-S16) over the chain extension of PEG-X with VAc is observed as attested by the nice match between expected and experimental molar mass values obtained upon VAc conversions, and the rather low dispersities (Table 1). In addition, the morphology evolution as the DP increases is similar to the one reported in the previous series of experiments (10 wt% solids content), with a transition from spherical particles (B1, $DP_{exp} = 48$) to vesicles (B3, $DP_{exp} = 97$) as shown by the cryo-TEM images (Figure 7). However, the increase in the solids content allowed triggering the formation of short worms along with the spheres and vesicles in B2 ($DP_{exp} = 73$).

The coexistence of the three morphologies indicate that finding the experimental window (i.e. the DP) for which the worm/fibre morphology can be exclusively attained is very narrow. The

difficulty in obtaining pure worm/fibre morphology in this system probably lies on the fact that the underpinning formation mechanisms are intricate. Indeed, in addition to VAc hydrophilicity, PVAc's low T_g (ca. 30 °C) should also play a significant role. The synthesis temperature is fixed way above the T_g of the PVAc, which implies that the hydrophobic block keeps a certain mobility ensuring a labile structure for the self-assemblies. This provides more opportunities for morphological transitions that we did not meet in our seminal works on PEG-X mediated VAc emulsion polymerization, where the polymerizations were performed at 25 °C. Eventually, the nature of the hydrophilic block plays an important role in the self-assembly and rearrangement mechanism. More specifically, the relatively low steric repulsion between the PEG corona compared to other hydrophilic macroCTA, including charged ones, would enhance the effective collision of the assemblies and thus promote their rearrangement.⁴³ The fine tuning of these parameters should allow to find conditions leading to pure worm/fibre morphologies. These investigations are underway in our laboratories.

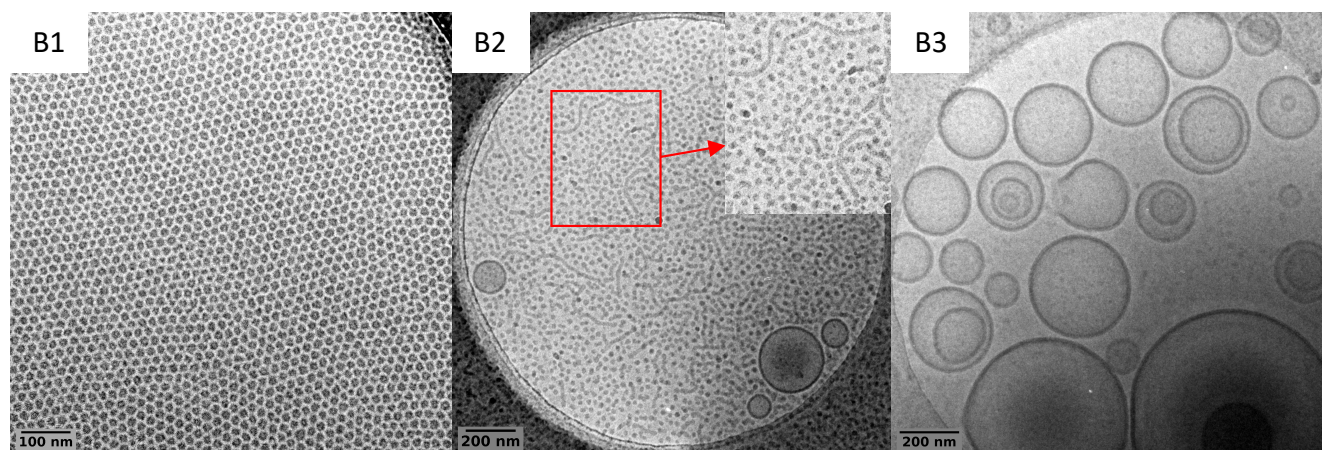


Figure 7. Representatives cryo-TEM images of the crude latexes B1-B3 (Table 1).

Conclusions

The aqueous MADIX-mediated emulsion polymerization of vinyl acetate was conducted using a poly(ethylene glycol) bearing a xanthate chain-end as hydrophilic macroCTA. The study focused mainly on synthesis performed at 70 °C with a targeted solids content of 10 wt%, which led to relatively well controlled polymerizations, with a good blocking efficiency, and to the formation of stable latexes. Increasing the DP of the PVAc block allowed to trigger morphological transitions of the self-assembled PEG-*b*-PVAc nano-objects, from spherical particles for a DP of 47 to a mix of spheres and vesicles for DP comprised between 62 and 100, and eventually to higher-order morphologies, notably large compound vesicles for DP of 187. Increasing the solids content to 15 wt%, worm morphology was obtained for a DP_{exp} of 73, however coexisting with spheres and vesicles.

The access to non-spherical morphologies depicted here for the first time for VAc emulsion polymerization was interpreted as the result of the high water solubility of VAc compared to other hydrophobic monomers classically employed in emulsion polymerization. Although the overall picture of the phenomena impacting the formation of a specific morphology is still to provide, adjusting the polymerization temperature and solids content may allow to isolate pure PVAc worm/fibre particles and eventually establish a phase diagram for this system.

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Conflicts of interest

There are no conflicts to declare.

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