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In situ monitoring of PISA morphologies†

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Polymerization-induced self-assembly (PISA) is a facile method to obtain block copolymer aggregates with defined morphologies. However, the transitions between these morphologies have been difficult to monitor directly in real-time during the polymerization. Herein, we describe a straightforward and readily accessible *in situ* method to monitor the evolution of nanostructure *via* changes in internal hydrophobicity during the PISA process using a polymer-tethered pyrene fluorescent probe. We were able to correlate morphological transitions with changes of the pyrene emission and gain unprecedented insight into the evolution of core hydrophobicity during PISA.

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Introduction

Amphiphilic block copolymers have been widely used to produce self-assembled nanostructures.^{1–4} In solution, these assemblies adopt not only spherical micellar morphologies but also higher-order structures, such as cylindrical micelles (worms) and vesicles to name a few.^{5–8} In particular, polymerization-induced self-assembly (PISA) has been used as a versatile method to obtain block copolymer particles in a highly tunable fashion at high concentrations.^{9,10} During PISA, a solvophilic precursor polymer is chain extended with a monomer that forms a solvophobic block. At a certain critical degree of polymerization (DP) of the solvophobic block, aggregation becomes energetically favorable and causes assembly of the polymer chains into multimolecular aggregates.^{11,12} Controlled radical polymerization techniques, such as reversible addition–fragmentation chain transfer (RAFT)^{13–15} or atom transfer radical polymerization (ATRP),^{5,16} have made the synthesis of well-defined block copolymers relatively straightforward, and for this reason, are widely adopted to conduct PISA.^{9,17–19}

Many approaches have focused on broadening the scope of PISA systems by introducing a wide range of polymer functionalities^{20–24} and exploring increasingly complex morphologies,^{25,26} such as framboidal shapes.²⁷ Such nanoparticles have been utilized for a range of applications from coatings²⁸ to nanomedicine.^{20,23,29–31}

With these promising applications, a better understanding of the morphological progression from molecularly dissolved unimers to complex assembled structures could enable us to better predict, design, and direct these polymerizations. However, the sensitivity of the self-assemblies to fluctuations of temperature and concentration imposes major restrictions on the feasibility and reproducibility of *ex situ* morphology characterization. Both dry state transmission electron microscopy (TEM) and dynamic light scattering (DLS), which are commonly used to characterize PISA morphologies, are often conducted at room temperature, and require dilution or drying of the samples, which may alter assembly size and morphology. To circumvent the problems associated with *ex situ* methods, liquid-cell transmission electron microscopy^{32,33} and small-angle X-ray scattering^{34,35} have been adapted, using specialized setups and systems, for monitoring PISA *in situ*. However, the practicality and availability of these methods is a limitation in some cases.

Aggregation-dependent fluorescent probes, such as pyrene, have long been used to observe the onset of self-assembly in block copolymer micelles, *i.e.* the critical micelle concentration (CMC).^{36–40} When a ground state pyrene monomer (M) is excited (M*) at $\lambda_{\text{max}} \sim 345$ nm, three distinct monomeric emission bands are observed at $I_1 \sim 375$ nm, $I_3 \sim 385$ nm, and $I_5 \sim 395$ nm upon radiative decay. The I_1/I_3 ratio can be used to determine the changes in hydrophobicity in the local environment surrounding the pyrene – as the I_1 emission band is sen-

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sitive to solvent polarity, and the I_3 emission band is relatively insensitive. Upon micellization, the presence of pyrene embedded in the hydrophobic core will lead to a change in the I_1/I_3 ratio. When plotted over a range of concentrations, the CMC can be determined by the inflection point in the graph.⁴¹ If the excited pyrene (M^*) is spatially proximal to another pyrene in its ground state (M), the pair can stack together to form a dimer ($M-M^*$). This complex ($M-M^*$) is referred to as an excimer and has an emission band around 475 nm (I_{exi}).^{42–44} The ratio of excimer to monomer has been used to monitor intra- and intermolecular interactions in proteins and membranes.^{45–47}

There is considerable utility in employing aggregation-dependent fluorescent probes to interrogate supramolecular assembly and morphology. For example, Brédas and co-workers have elucidated how the intermolecular arrangements of organic π -conjugated molecules dictate electronic and optical properties.⁴⁸ Importantly, π -conjugated systems often stack in columns and form face-to-face interactions (H -type aggregates) that have a low fluorescence quantum yield. Efforts have been made to enhance the emission of π -conjugated materials by modifying the π - π stacking arrangement *via* installation of bulky side groups.⁴⁹ Utilizing supramolecular interactions to alter the arrangement of these systems, in particular, the adoption of a slipped stacking arrangement (J -type aggregates) can result in dramatic improvement in fluorescence quantum yield. Zhang, Wang, and co-workers reported that by introducing sterically demanding groups, bis(pyrene) derivatives could adopt a J -type aggregation which led to an almost 30-fold improvement in excimer emission.⁵⁰ Importantly, the sterically bulky groups led these aggregates to adopt a dot-shaped (spherical) morphology with high excimer fluorescence, while the bis-pyrenes in the absence of sterically demanding groups formed sheet-like morphologies and

adopted an H -type aggregation with much lower excimer emission. These examples highlight how changes in fluorescence excimer emission can be followed to provide insight into nano-scale transformations in supramolecular morphology.

In this article, we describe a straightforward and practical method to monitor changes in morphology and core hydrophobicity of the amphiphilic block copolymer polymer assemblies that form and evolve during the PISA process. Leveraging the aggregation-induced change in the emission spectrum of the fluorescent probe pyrene, we show that the ratio between the monomer (I_{mon} at 375 nm) and excimer emission (I_{exi} at 475 nm) can be used to study morphological changes during PISA. At the same time, the I_1/I_3 ratio could also be used to monitor the hydrophobicity within the cores of nanoparticles prepared during PISA.^{51,52} Importantly, all this information can be obtained *in situ*, using a standard benchtop fluorometer.

Results and discussion

We synthesized a poly(N,N -dimethylacrylamide) (PDMA) macro-chain-transfer agent (macroCTA) with a carboxylic acid functionality on the Z group for the selective installation of the pyrene probe in proximity to the core-forming hydrophobic block of the polymer chain.²⁵ Size-exclusion chromatography (SEC) indicated the resulting macroCTA had a number-average molar mass (M_n) of 7900 g mol⁻¹ and dispersity (D) = 1.09 (Fig. S1†). To install the fluorescent pyrene probe at the growing chain end of the PDMA macroCTA, the amine-functionalized pyrene was coupled to the carboxylic acid on the Z-group of the macromolecular trithiocarbonate using standard peptide coupling conditions (see Fig. 1 and S1–16† for details).



Fig. 1 Synthetic outline of the synthesis of the macro-chain-transfer agent (macroCTA) and the polymerization-induced self-assembly (PISA) process. (a) Synthesis of PDMA macroCTA (1), pyrene-labeled PDMA macroCTA (2) and the chain extension polymerisation using diacetone acrylamide (DAAm) to yield the amphiphilic block copolymer poly(N,N -dimethylacrylamide)-*block*-poly(diacetone acrylamide) (PDMA-*b*-PDAAm) (3). (b) Schematic representation of the morphologies observed during PISA upon chain extension of the PDMA macroCTA with DAAm.



Previously, Figg *et al.* showed that to access higher-order morphologies in aqueous DMA/diacetone acrylamide (DAAm)-based PISA systems, the incorporation of hydrophilic DMA monomer into the hydrophobic DAAm block was crucial to increase core hydration and chain mobility.⁵³ As an alternative approach, we decided to alter the solvent polarity by using a 1:1 (v/v) mixture of methanol and water to facilitate morphology transitions without the need for an additional comonomer. Additionally, we added non-functionalized PDMA macroCTA (**1**) to the pyrene-functionalized PDMA macroCTA (**2**) to prevent fluorescent quenching and to minimize the contribution of the hydrophobic pyrene chain end in the assembly process. The ratio of **2** to **1** was maintained at 0.2 throughout the experiments. All the polymerizations were conducted at 70 °C under constant stirring using 4,4'-azobis(4-cyanovaleric acid) (ACVA) as a thermal radical initiator in a 1:1 (v/v) mixture of methanol and water.

Initially, a range of DPs of the hydrophobic block was targeted to evaluate the scope of this PISA system (Table S1†). Each PISA reaction achieved full monomer conversion within 5 h, as confirmed by ¹H NMR spectroscopy. Size-exclusion chromatography (SEC) of the chain extension products showed some residual uninitiated macroCTA. The deconvoluted SEC chromatogram revealed that the chain extension (PDMA-Pyr-*b*-PDMA) resulted in a blocking efficiency of 91%, see Fig. S16.† Despite this, we still observed a range of morphologies from spheres (DP = 100), worms/worm network (DP = 200) to vesicles (DP = 300), as confirmed by dry state TEM (Fig. 2b). Importantly, control experiments with 100% non-functionalized macroCTA showed similar morphologies at the same DPs (Table S1 and Fig. S18†), suggesting the pyrene residues play only a minimal role in governing self-assembly. It is worth noting that keeping the percentage of pyrene-functionalized macroCTA low (20%) was critical in ensuring that the pyrene aggregation did not affect the overall morphologies obtained during the PISA process. Increasing ratios of functionalized macroCTA led to different morphologies when compared to the control studies without pyrene-functionalized PDMA (**2**). Most notably, higher ratios of pyrene-functionalized macroCTA

led to kinetically trapped spheres throughout the same polymerization.

After showing that we could obtain a range of morphologies with the chain extension of the PDMA macroCTA using only DAAm, we sought to study the emission behavior of the pyrene probes in morphological transitions during PISA. Using the same polymerization conditions, we targeted a final DAAm block with DP of 300, which generated vesicles in the previous experiments (Fig. 2b). Specifically, two polymerization sets under identical conditions, but one in sealed glass vials and one in the fluorometer, were conducted to correlate monomer conversion, nanoparticle morphology, and pyrene fluorescence during the PISA process. The polymerization set in glass vials was used to determine monomer conversion, molar mass, and morphology (Fig. 3a and Table S2†) by ¹H NMR spectroscopy, SEC, and TEM, respectively. The polymerization in the fluorometer allowed monitoring of the pyrene emission throughout the reaction. Notably, we found that the emission ratio of pyrene excimer to monomer ($I_{\text{exi}}/I_{\text{mon}}$), which is related to the stacking of the pyrene molecules, changed throughout the PISA process (Fig. 3b).

Previously, Wang *et al.* reported that the arrangement of stacked fluorophores affected the fluorescence quantum yield of the pyrene excimer.⁵⁰ We reasoned that during PISA, the pyrenes at the core of the self-assembly can undergo changes in their stacking orientation *i.e.*, packing mode and packing efficiency. Therefore, the sensitivity of pyrene excimer emission to minor changes in fluorophore stacking orientation was used to monitor the structural rearrangements that accompany morphological transitions during PISA.^{43,50,54}

After 15 min of the polymerization, we observed the aggregation of polymers into spherical micelles (Fig. 3a). As the self-assemblies transformed from spheres to worms after 30 min (DP = 108), we observed a decrease in the $I_{\text{exi}}/I_{\text{mon}}$ ratio. Throughout the PISA process, the transitions between different morphologies led to considerable structural rearrangements; we believe the fluctuations in the excimer ratio are due to changes in the packing efficiency and packing modes between the fluorophores during the reaction. The tran-



Fig. 2 Size-exclusion chromatography (SEC) and transmission electron microscopy (TEM) characterization of polymers and assemblies formed via PISA. (a) SEC traces of PDMA macroCTA chain extended with DAAm to different targeted degrees of polymerization (DP) (50, 100, 200, and 300). (b) TEM images of spheres at a PDAAm DP of 100 (red, left), worms at PDAAm DP of 200 (orange, middle), and vesicles at a PDAAm DP of 300 (dark yellow, right). Scale bar is set to 500 nm. All PISA experiments for TEM were carried out to full conversion and crosslinked at 70 °C using a 10 wt% O-alkyl bishydroxylamine crosslinker relative to DAAm monomer to preserve the morphology for TEM analysis.





Fig. 3 Correlation of polymerization kinetics (obtained by NMR spectroscopy), PISA morphology (obtained by *ex situ* TEM), and pyrene fluorescence (obtained by fluorescence spectroscopy). (a) TEM images of assemblies from polymerization aliquots at different conversions: spheres (21% red, top left), worms (36% orange, middle left), and vesicles (51% yellow, bottom left, 60% green, top right, 84% blue, middle right, and 93% purple, bottom right). Scale bars represent 1 μm. The ratio of (b) pyrene excimer to monomer emission ($I_{\text{excimer}}/I_{\text{monomer}}$) and (c) pyrene monomer emission bands I_1 to I_3 (I_1/I_3) throughout the PISA process.

sitions during the classical PISA process (sphere-to-worm and worm-to-vesicles) are hypothesized to go through different mechanisms. The sphere to worm transition is often proposed to take place by collisions between self-assembled aggregates.⁵⁵ In contrast, the worm to vesicle phase is hypothesized to proceed through a more continuous process. Blanz *et al.* showed that the worm to vesicle transition proceeds *via* several intermediates: the branched network of worms leads to the partial fusion of worms, which then form nascent bilayers (as evidenced by the presence of 'octopi'- and 'jellyfish'-like intermediate structures) that eventually form pure vesicles.⁵⁶ Nevertheless, both transitions require structural rearrangements in the polymer self-assembly.

Upon further polymerization of the DAAM block to DP = 153 at 45 min, vesicles were formed, which again led to a major change in the pyrene emission and, in this case, an increase in the $I_{\text{excimer}}/I_{\text{monomer}}$ ratio. These results suggest that transitions involving cylindrical structures are associated with changes in chain arrangements, which affect the fluorophore stacking and can be readily detected by changes in the $I_{\text{excimer}}/I_{\text{monomer}}$ ratio. Also, further increases of the PDAAm block size from this point onwards (DP = 181 at 60 min to DP = 280 at

180 min) showed only a small change in the $I_{\text{excimer}}/I_{\text{monomer}}$ ratio, consistent with no further significant change in morphology. It is worth noting that this gradual increase in the excimer ratio corresponds well with the fact that the hydrophobic block is continuing to grow, and in turn, the packing parameter, which affects the pyrene stacking arrangement, is still increasing throughout the polymerization.¹¹

The pyrene monomer emission can also be used to gain information about the hydrophobicity in the local environment of the fluorescent probe. A decrease in the intensity ratio of the emission bands I_1 to I_3 is indicative of an increasingly hydrophobic local environment. From these studies, we can monitor the change in hydrophobicity near the growing chain end in the core of the assemblies throughout the polymerization. Upon chain extension of the PDMA macroCTA with DAAM, the pyrene on the active chain end experiences a progressively more hydrophobic environment, as shown in the decreasing I_1/I_3 ratio within the first 40 min of the polymerization (Fig. 3c). Interestingly, at 45 min a slight increase of the I_1/I_3 ratio was observed during the worm-to-vesicle transition, which suggests a slight increase in hydrophilicity that coincides with the formation of a bilayer vesicular structure



where water is on the inside and outside of the bilayer. Notably, after the polymer chains have rearranged into vesicles the hydrophobicity (as indicated by the I_1/I_3 ratio) remains relatively constant as the pyrene moieties are buried in the center of the bilayer, presumably with very little solvation. These data suggest this non-invasive technique can monitor, in tandem, both changes in the self-assembly and hydration in the cores.

Conclusions

In summary, we demonstrated the use of pyrene as a fluorescent tag to monitor morphology and hydrophobicity during the PISA process. We were able to relate morphological transitions during the PISA process to changes in pyrene excimer-to-monomer ratio, which is dependent on the spatial proximity of pyrene molecules. Furthermore, the ratio of the I_1 and I_3 emission bands of the pyrene monomer provided unprecedented insight into the hydrophobicity at the growing chain end at the assembled core. In the future, we believe this straightforward and non-invasive *in situ* technique could be used to better understand and predict morphological progressions during the PISA process.

Conflicts of interest

There are no conflicts to declare.

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