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The morphology transition from micelles to vesicles of a solution-state self-assembled block copolymer, containing a fluorescent dye at the core–shell interface, has been induced by an addition–elimination reaction using a thiol, and has been shown to be coupled to a simultaneous ON-to-OFF switch in particle fluorescence.

Precise control over solution-state self-assembled polymer morphologies is currently of great interest to the research community. Various morphologies such as spherical micelles, cylinders, rods and vesicles can be formed by the self-assembly of amphiphilic block copolymers in selective solvents.¹ Block copolymer composition and properties control the morphology that is adopted in solution by the amphiphile. Conventional self-assembled morphologies are based on hydrophilic–hydrophobic repulsive interactions,² and as a result, self-assembled nanostructures formed from stimuli-responsive polymers are able to undergo morphology transitions induced by external stimuli such as pH, temperature and light.^{3–9} Importantly, the responsive behavior of self-assemblies enables these new materials to find applications in nanotechnology and/or drug delivery.^{10–15} Moreover, self-assembled nanostructures having fluorescent properties are also of interest given the desire to track such species in applications such as nanomedicine.^{16–18}

Previously, Baker *et al.* have shown that the conversion of dibromomaleimide (DBM) to dithiomaleimide (DTM) is highly efficient, and that by an excess of thiol further conversions can occur due to retention of the double bond in the DTM motif.¹⁹ Furthermore, we previously reported that DTM's functionalized with alkyl thiols are highly fluorescent whilst those with aromatic substituents show a significant decrease in fluorescence emission.²⁰ Recently, we demonstrated that the DTM motif can be introduced into block copolymers as a highly emissive fluorescent self-reporting

probe *via* a dual ring-opening polymerization and reversible addition-fragmentation chain transfer polymerization (ROP/RAFT) initiator.²⁰ Aqueous solution-state self-assembly of these amphiphilic block copolymers results in DTM incorporation at the core–shell interface of spherical micelles.²¹ Herein, we utilize the reactivity of the DTM group to induce a morphology transition from spherical micelles to vesicles which occurs simultaneously with an ON-to-OFF switch of fluorescence emission (Fig. 1). The change in fluorescence and morphology is induced by an addition–elimination reaction which removes the hydrophobic segment that is connected to the DTM functional group at the block copolymer interface, and replaces it with an aromatic substituent. This subtle change in chemistry of the DTM group has a significant effect on the properties of the assembly. We propose the use of this triggered change in fluorescence and overall self-assembled structure at the DTM group as an accessible read-out for the change in chemistries within a block copolymer in a polymeric nanostructure. As such we propose this approach has interesting potential scope for use in sensing and also tracking applications.

We first synthesized an amphiphilic block-dye-block copolymer *via* a combination of ROP²² and RAFT polymerization²³ utilizing a dual ROP/RAFT initiator, **1** (Scheme 1). The design of this initiator species ensures that the DTM group is located between

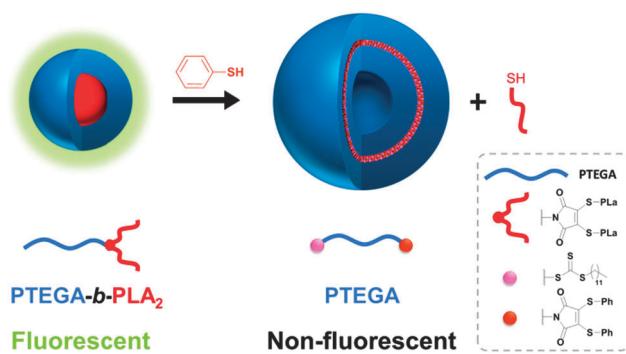


Fig. 1 Schematic of the morphology change induced by an addition–elimination reaction of the micelle with thiophenol.

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the hydrophobic and the hydrophilic blocks, allowing the addition–elimination reaction to be coupled with both a morphology transition and a fluorescence ON-to-OFF switch. The structure and properties of the amphiphilic copolymers were carefully chosen to enable a significant modification of the hydrophilic/hydrophobic balance alongside a fluorescence emission decrease by changing the nature of the hydrophobic segment. As the hydrophobic segments were connected to the DTM motif as the thiol ligands, subsequent addition–elimination with a thiol following self-assembly would allow elimination of the hydrophobic blocks.

The DTM-containing ROP/RAFT dual initiator, **1**, was synthesized from a 2,3-DBM-functionalized RAFT agent.²⁴ Reaction with mercaptoethanol and triethylamine gives a fluorescent DTM-functionalized ROP/RAFT dual initiator (**1**) with two hydroxyl groups allowing ROP²² of *rac*-lactide to be performed to afford polymer **1'**. This polymer was then chain-extended to afford the diblock copolymer [poly(ethylene glycol monomethyl ether methacrylate)]-*b*-[poly(D,L-lactide)], **2**, see Scheme 1. The well-defined fluorescent block-dye-block copolymer, PTEGA-*b*-PLA₂, **2**, was fully characterized using NMR spectroscopy and SEC analysis (M_n (NMR) = 33.1 kDa, M_n (SEC, DMF) = 19.5 kDa, D_M = 1.42).

The self-assembly of copolymer, **2**, into micelles, **3**, was achieved *via* direct dissolution of the copolymer in purified 18.2 MΩ cm water at a concentration of 1 mg mL⁻¹. The fluorescence spectra of the spherical micelles **3** in 18.2 MΩ cm water shows an excitation maxima at 405 nm, and an emission maxima at 510 nm, which are similar to previously reported DTM polymer systems.²¹ The size and morphology of the micelles was confirmed by light scattering and microscopy analysis. Multi-angle laser light scattering (MA-LLS) indicated R_h = 26 nm (see ESI†) and dry-state transmission electron microscopy (TEM) analysis on graphene oxide, a very thin support that does not require staining, suggests a spherical morphology, see Fig. 2a.²⁵

The micelle solution, **3**, was treated with 20 equiv. of thiophenol, and then purified by exhaustive dialysis (MWCO = 1 kDa). Further experiments with a range of aromatic thiols indicated that the

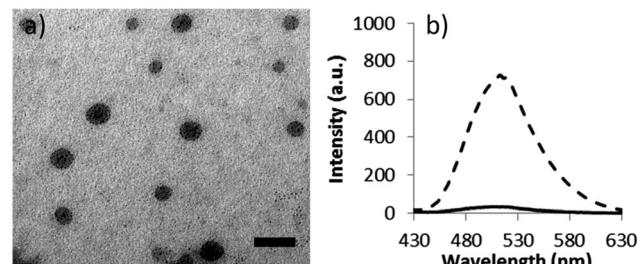
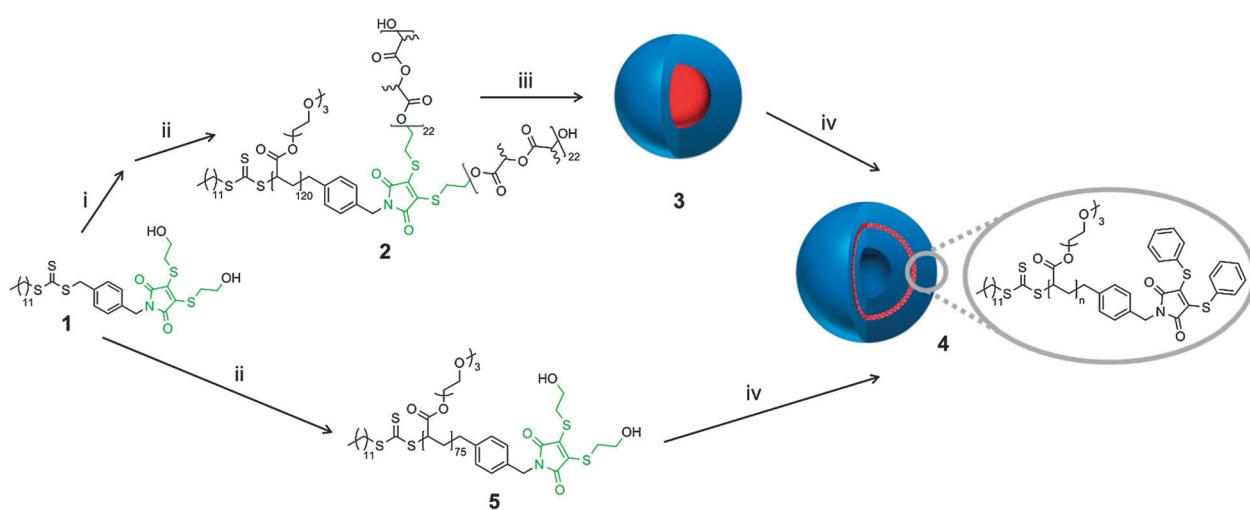


Fig. 2 (a) Representative unstained TEM image of micelles (scale bar = 50 nm) **3** and (b) fluorescence emission spectra before (dash line) and after the reaction (solid line), with excitation at 405 nm.

reaction also works efficiently with a small excess of thiol. The thiol underwent addition to the DTM group, with corresponding elimination to afford thio-terminated poly(*rac*-lactide), **7**. Given that the thio-PLA residue is insoluble in water, the solution was then centrifuged to remove the thio-PLA precipitate (see ESI† for ¹H NMR spectrum).

To characterize the particle morphology of assembly **4a**, Multi-Angle Laser Light Scattering (MA-LLS) was performed to determine the radius of gyration R_g and hydrodynamic radius R_h of the assemblies, **4a**. The ratio of R_g/R_h gives an indication of the nanostructure morphology, with 0.775 indicating a solid micelle and 1 indicating a hollow vesicular structure.²⁶ By interpreting the data collected in static light scattering (SLS) mode, using CONTIN analysis, the radius of gyration R_g was determined to be 51 nm. From the dynamic light scattering (DLS) data, R_h was found to be 56 nm. For nanostructure **4a** the R_g/R_h was calculated to be 0.91, which suggests that nanostructures formed are hollow vesicular particles (see ESI†). We propose that the vesicles' hydrophobic layer is composed of both the substituted maleimide group (containing the -SPh ligands) and dodecyl end group (the RAFT agent Z-group). This is consistent with previous reports which have shown that hydrophilic polymers with hydrophobic aromatic and aliphatic end-groups can self-assemble into nanoparticles, including vesicles.^{27–30}



Scheme 1 Preparation of polymers **2** and **5** from CTA **1**, preparation of micelles **3**, and the addition–elimination reaction of **3** and **5** which results in a morphology transition to afford vesicle **4**. Conditions: (i) *rac*-lactide, thiourea, *(−)*-sparteine, CH_2Cl_2 ; (ii) AIBN, CHCl_3 , TEGA at 60 °C; (iii) direct dissolution in water; (iv) thiophenol.



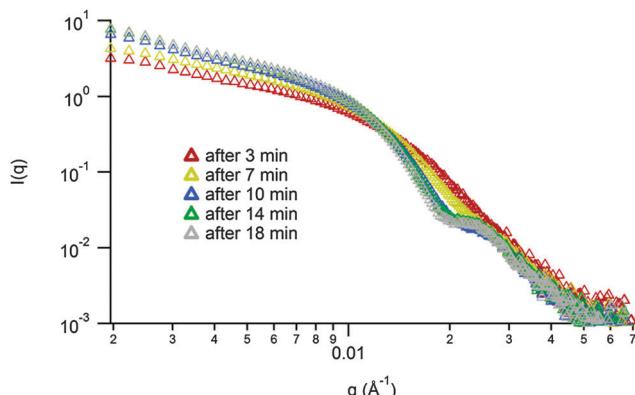


Fig. 3 SAXS profiles for the *in situ* micelle-to-vesicle transition.

Moreover, examination of the emission spectrum of the resultant solution of **4a** indicated a drastic decrease of the fluorescence as a consequence of the modification at the DTM reactive center. At the same excitation wavelength, the comparison of the emission spectra of the solution before and after the reaction (measured at the same concentration) showed a decrease in the intensity of the maxima (510 nm) from 730 to 30 a.u., see Fig. 2b.

Unfortunately, attempts to image the vesicles by dry-state TEM were not possible as the nanostructures were not stable to dehydration. However, to further probe the proposed micelle-to-vesicle transition upon thiol addition synchrotron SAXS experiments were performed. The transition described in Scheme 1 part (iv) was performed in the beamline and the *in situ* kinetics of the morphology transition were examined (Fig. 3). This has the advantage that it allows for the monitoring of the transition without the need for removal of insoluble PLA, drying and suspension of the nanostructures (as was necessary for MA-LLS analysis). Analysis of the SAXS curves over a 20 minute time period indicates a change of morphology. At the beginning, a spherical morphology was observed. An increase in the size of the morphologies in solution happened promptly (less than 10 min) and a form factor fit indicated the formation of vesicles as well as the presence of random chains in solution (thio-PLA in solution) (see ESI[†] for the different fittings of the SAXS curves over time).

To further probe the vesicle formation which was observed in the transition from **3** to **4a**, the addition–elimination reaction was performed on a range of model homopolymers (data shown for PTEGA₇₅, M_n (NMR) = 16.2 kDa, M_n (SEC) = 13.5 kDa, D_M = 1.19), **5**, synthesized *via* RAFT polymerization of TEGA utilizing **1** as a chain transfer agent (CTA). DLS analysis of aqueous solutions of the initial homopolymer **5** (which possesses α -diol and ω -dodecyl end-groups), indicated the presence of unimers in solution. However, after the addition–elimination reaction between **5** and an excess of thiophenol, well-defined nanostructures were observed, **4b**. MA-LLS was performed, and values of R_g and R_h were extracted from the results. By interpreting the data collected in SLS mode, the R_g equals 63 nm and from the DLS data, the R_h equals 68 nm and hence the R_g/R_h obtained was 0.93, which once again suggests that the nanostructures formed in this reaction are vesicles. Similar to particles **4a** which result from reaction of the micelles (**3**) with thiophenol, the reaction of the homopolymer **5** with thiophenol

forms particles **4b** of a similar size and morphology. This is understandable as the addition–elimination reaction of **3** and **5** would be expected to give the same resultant homopolymer, namely P(TEGA) homopolymer with α -Ph and ω -dodecyl end-groups, which would be expected to assemble into a similar morphology. As observed for the addition–elimination reaction with the micelles, the fluorescence emission of the homopolymer solution again underwent a fast (15 min by analysis of the 535 nm emission) ON-to-OFF switch during the reaction with thiophenol, see ESI.[†]

In conclusion, we have shown that an amphiphilic block-dye-block copolymer containing a DTM group can undergo a fast morphology transition from spherical micelles to vesicles. This is triggered by the addition of thiophenol to the DTM group, with corresponding elimination of the hydrophobic blocks. A unimer-to-vesicle transition also occurs for a DTM group-containing homopolymer. In both cases addition of thiophenol leads to a simultaneous fluorescence ON-to-OFF switch. This approach is extremely versatile and could be tuned for utilization with a range of aromatic thiols and self-assembled systems which contain the DTM functional group. We suggest that such a simultaneous ON–OFF switch and morphological reorganization could be readily applied as a tracking mechanism and also as a mechanism for monitoring release in biological and/or synthetic self-assembled systems.

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