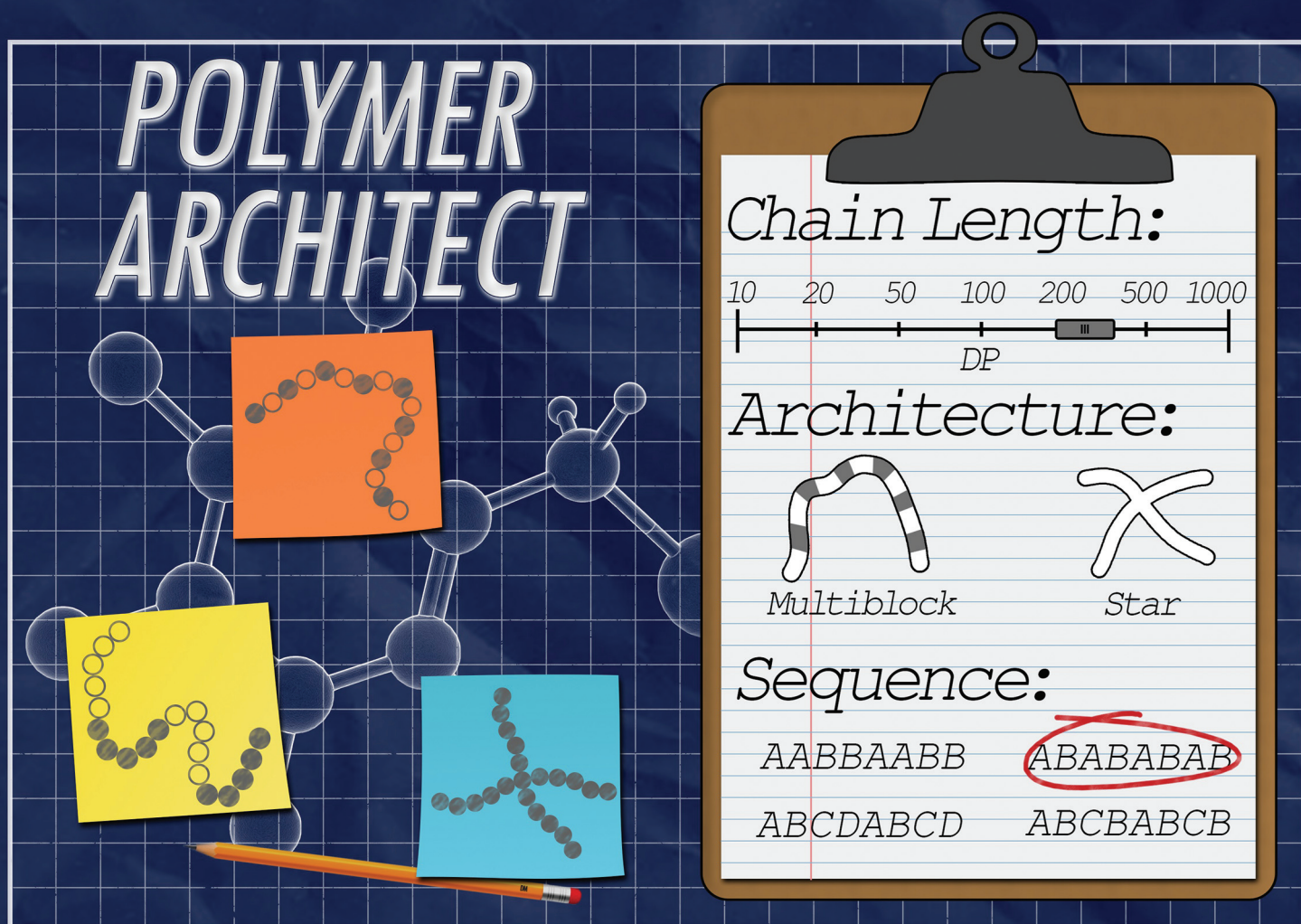


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One-pot synthesis of amphiphilic multiblock
poly(2-oxazoline)s *via para*-fluoro-thiol click reactions

PAPER

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[View Journal](#) | [View Issue](#)Cite this: *Polym. Chem.*, 2021, **12**, 6392One-pot synthesis of amphiphilic multiblock poly(2-oxazoline)s via *para*-fluoro-thiol click reactions†Tieshuai Zhao, Ben Drain,  Gokhan Yilmaz and C. Remzi Becer  *

A clickable initiator, pentafluoro benzyl bromide, has been investigated for the cationic ring opening polymerization of poly(2-oxazolines). Additionally, the clickable alpha end group was then utilized in a *para*-fluoro-thiol click reaction to synthesise linear diblock, tetrablock, multiblock copolymers as well as star shaped poly(2-oxazoline)s using dithiol compounds as terminating agents. Thus, a one-pot approach combining the *para*-fluoro-thiol click reaction and direct termination of the poly(2-oxazoline) living chain end with 4,4'-thiobisbenzenethiol has been performed to prepare multiblock copolymers of poly(2-ethyl-2-oxazoline) (PEtOx) and poly((2-ethyl-2-oxazoline)-*b*-(2-methyl-2-oxazoline)) (PEtOx-*mb*-PMeOx). All obtained polymers were characterized by Size Exclusion Chromatography (SEC), ¹H Nuclear Magnetic Resonance (NMR) and Matrix-Assisted Laser Desorption/Ionization-Time of Flight (MALDI-ToF) mass spectrometry. Last but not least, the self-assembly properties of prepared amphiphilic polymers were studied with DLS and TEM. Nanoparticles with a diameter ranging from 184 nm to 250 nm were observed in TEM for PEtOx-*mb*-PMeOx copolymers.

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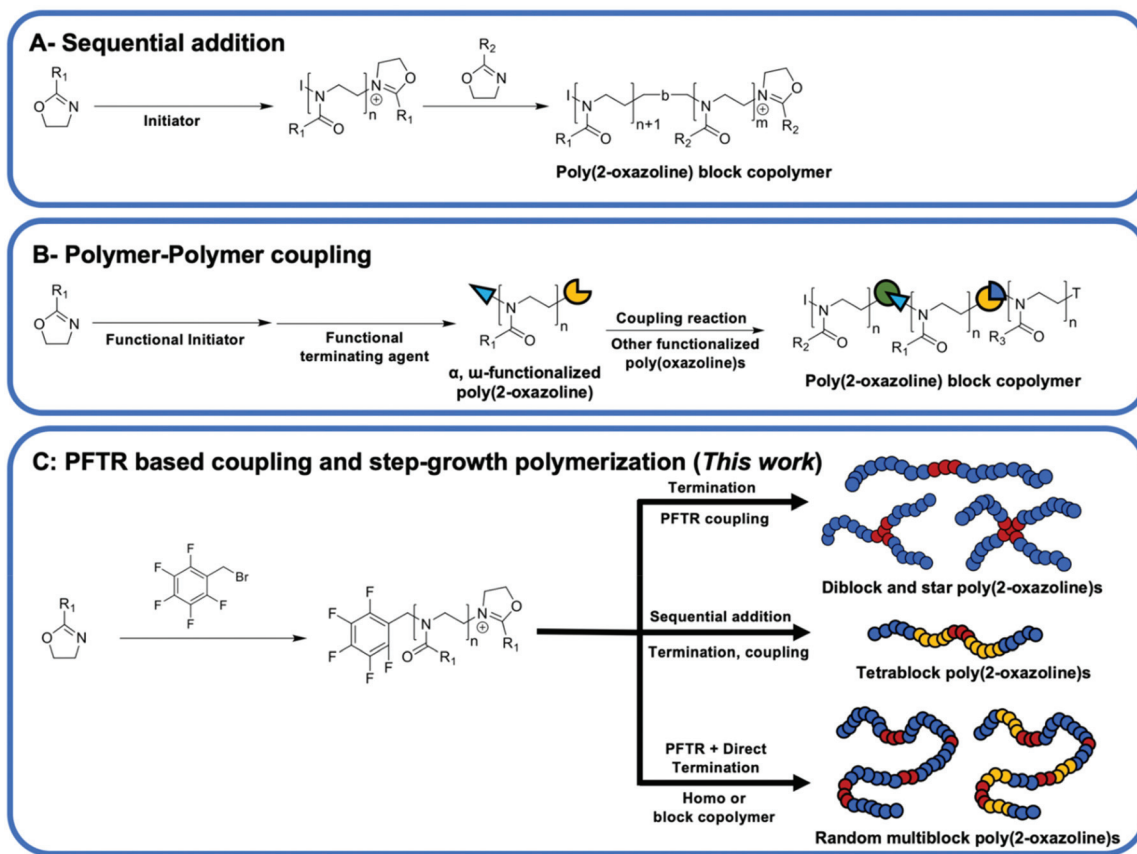
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Introduction

Poly(2-oxazoline)s(POx) have attracted great attention as they possess various properties including biocompatibility,^{1–8} thermal responsive ability,^{9–12} tunable hydrophilicity^{9,13} and adjustable cytotoxicity^{7,14,15} that are useful for biomedical and industrial applications such as adhesives¹⁰ and lubricants.¹⁶ Synthesis of poly(2-oxazoline)s with well-defined chain length, and end groups has been achieved through a living cationic ring-opening polymerization (CROP) process that can be initiated by various commercially available initiators such as tosylates, triflates, acid halides, and benzyl halides.¹⁷ Thus, the living chain ends have been further utilized in chain extension reactions to obtain block copolymers^{10,13,18,19} or alternatively have been functionalized to introduce functional groups for post-polymerization modification (PPM) reactions.^{20–22}

Owing to the biocompatibility of POx, synthesis of poly(2-oxazoline) based block copolymers with amphiphilic structures to form polymeric micelles or nanoparticles have been on the focus of many research groups.^{23–28} Preparation of diblock, triblock and tetrablock POx and subsequent manufacturing of micelles and nanoparticles have been already

reported.^{10,12,13,18,29–31} There are two main methods for the synthesis of 2-oxazoline based block copolymers. These methods are sequential addition of monomers to the polymerization mixture after complete consumption of the previous monomer, and the coupling of end-functionalized polymers¹¹ (Scheme 1A). The former method (aka sequential addition method) was the most widely employed one since one-pot process is more convenient to obtain multiblock polymers.³² However, the chain length of block polymers synthesized *via* this method is relatively



Scheme 1 Synthetic approaches to prepare poly(2-oxazoline) based block copolymers.

synthesis of block copolymers and dendrimers of POxs.^{11,40,41} The clickable azide or alkyne groups are introduced through the use of a functional initiator⁴² or by termination⁴³ of the living cationic chain end to prepare functionalized POx chains and subsequently reacted with other polymers to prepare linear diblock,^{13,18,44} linear triblock,¹⁰ and star shaped polymers.⁴² However, the undesired strong interaction between amide groups on POx backbone and Cu catalyst creates a real challenge in the purification of the final product.⁴⁵

Utilizing metal-free click reactions such as thiol-ene reaction⁴⁶ and Cu-free strain-promoted azide-alkyne cycloaddition (SPAAC) click reactions⁴⁷ to synthesize block copolymers have also been reported. Besides, studies on α - and ω -terminal functionalized POxs usually led to synthesis of diblock copolymer or triblock copolymer due to low coupling efficiency of POxs,^{17,22} while the selected polymer-polymer coupling reaction utilized in reports was mainly CuAAC click reaction^{13,18} with some studies employed thiol-based click reactions to prepare star, grafted and block copolymer of POxs.^{48–50} The use of *para*-fluoro-thiol click reaction to prepare POx block copolymers was not reported. Additionally, research on the synthesis of multi-block POxs using a difunctionalized POx precursor *via* a step growth approach remains absent (Scheme 1C).

Herein, we introduce the use of pentafluoro benzyl bromide (PFBB) as a functional initiator for the CROP of poly(2-ethyl-2-

oxazoline) (PEtOx). Clickable functionality allows the preparation of linear diblock, star shaped and multiblock poly(2-oxazolines). Utilizing commercially available dithiol and multi-thiol compounds, linear diblock and star shaped polymers of PEtOx were synthesized *via para*-fluoro-thiol click reaction (PFTR) approach. The block copolymer consists of 2-ethyl-2-oxazoline and 2-methyl-2-oxazoline (MeOx) was also prepared using PFBB as an initiator, and this resulted in a tetrablock copolymer of P(EtOx-*b*-MeOx) with a dithiol compound in between the blocks. Furthermore, a combination of the PFTR and direct termination of the cationic chain end of PEtOx or P(EtOx-*b*-MeOx) with dithiol as the nucleophile gives rise to a multiblock poly(2-oxazoline)s within 10 minutes in a one-pot reaction. The synthesized diblock and multiblock poly(2-oxazoline)s have presented amphiphilic structures and therefore the self-assembly properties of these polymers were investigated using transmission electron microscopy (TEM).

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understand the polymerization kinetics. Initial reaction conditions were used a [EtOx]:[PFBB] ratio of 60:1, giving a theoretical degree of polymerization (DP) of 60. The polymerization was performed in a microwave reactor at 140 °C, at a monomer concentration of 4.0 M, and using dry acetonitrile (MeCN) as the solvent. These conditions were widely employed in other kinetic investigations for the CROP of 2-oxazolines with other commercially available initiators such as methyl tosylate,⁵¹ benzyl bromide,⁵² and acyl halides.⁵³ This allowed a direct comparison of the results including rate constant of polymerization to previously reported values. A stock solution of EtOx and PFBB in dry MeCN at 4.0 M concentration was prepared and equal amounts of stock solution were distributed into a series of 2–5 mL microwave vials under inert and anhydrous conditions. These vials were then transferred to the microwave reactor, heated to 140 °C, and quenched with a sodium hydroxide/methanol solution at predetermined time points. ¹H NMR and SEC analysis were conducted to determine the monomer conversion, the molar mass of obtained polymers at different time points as well as the polymerization rate constant (Fig. 1). Monomer conversion was calculated based on integration of EtOx peaks in the ¹H NMR (ESI, Fig. S1†). ¹H NMR spectrum collected at each time point are shown in Fig. S1.† ¹H NMR results show the shift and split of the peak for the methylene group of pentafluoro benzyl bromide and the consumption of EtOx.

The plot of $\ln([M]_0/[M]_t)$ versus time depicted in Fig. 1A show a pseudo-linear relationship indicating that the concentration of active and propagating chains is constant through-

out the polymerization. However, closer inspection of data points suggesting a slightly sigmoidal shape, which indicates the slight existence of possible slow initiation and termination. Nevertheless, the semi-logarithmic kinetic plot together with the SEC results demonstrates the living character of the polymer chains under these reaction conditions. In addition to this, the reaction time vs. conversion plot in Fig. 1B showing the linear dependence at low conversion, and the $M_{n,SEC}$ vs. conversion plot depicted in Fig. 1C shows a linear dependence further confirming a living, controlled chain growth polymerization and absence of termination or chain transfer reactions. Dispersity values were below 1.20 throughout the kinetics investigation, indicating the polymerization has proceeded in a controlled fashion. The decreasing trend of dispersity values with increasing molecular weight also fits the theoretical trend for living polymerization.⁵⁴ The linear trends in both Fig. 1A and 1C are in good agreement with experimental data with R^2 values over 0.99 were obtained. The retention time change in Fig. 1D together with the $M_{n,SEC}$ values indicated the growing character of the polymer chain, while the experimental $M_{n,SEC}$ value has stayed lower than the theoretical value. For example, at $t = 12$ min where the monomer conversion was around 93%, the $M_{n,theo}$ value was 5700 Da, while the measured $M_{n,SEC}$ was only 5000 Da. This slight difference may be due to the PMMA calibration standards or due to effect of pentafluoro benzyl initiating group on overall hydrodynamic volume of the polymer.

The apparent propagation rate of polymerization (k_{app}) was determined based on the first order kinetics following (ESI, eqn (S2)†). While based on the assumption of concentration of initiator and living propagating chain ends are equal and given concentration of initiator at t_0 is $[I]_0$, k_{app} was calculated as $3.38 \times 10^{-3} \text{ L (mol s}^{-1}\text{)}$. This kinetic study has shown that PFBB is an excellent initiator for CROP of EtOx, although the reaction rate is rather slow compared to benzyl bromide, which was reported by Schubert *et al.* as $1.14 \times 10^{-1} \text{ L (mol s}^{-1}\text{)}$ at 160 °C.⁵¹ As the counter ion in each initiating system remains the same, possible reason for a moderate polymerization rate could be the pentafluoro benzyl group is more electron withdrawing and sterically large than a benzyl group, which makes the cationic species generated in initiation and propagation more energetically unfavorable.

Before exploitation of the clickable pentafluoro benz

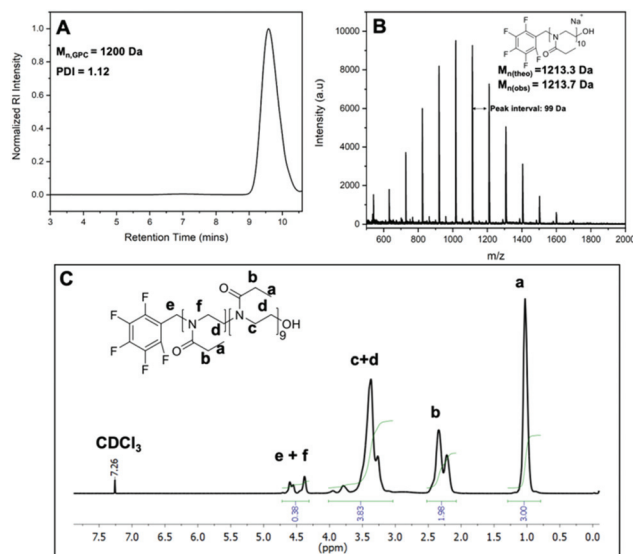


Fig. 2 (A) SEC measurement in THF as an eluent, (B) MALDI-ToF MS, and (C) ^1H NMR analysis of PFB-PETox₁₀. The SEC trace and the distribution shown in MALDI-ToF indicates a DP10 polymer. In the ^1H NMR, corresponding peaks were labelled, and part of the backbone hydrogens (f) shifted to lower field at around 4.5 ppm from c+d at 3.45 ppm due to initiating group effect.

major distribution that fits to the theoretical molar mass value of PETox with pentafluoro benzyl and hydroxyl end groups was observed. For instance, the observed peak at 1113.9 Da was close to the theoretical calculations of the sodium salt of PETox with 9 repeat units and possessing -OH and PFB end groups. Similarly, the peak at 1213.7 Da was close to the same structure with 10 repeat units and the distance between the peaks was 99 Da, which is equal to the molar mass of EtOx. It should be noted that in the ^1H NMR analysis the total integration ratio of peaks was 9.2. This corresponds to 90 protons of ethyl groups and 2 protons of methylene group from the initiator, which is in good agreement with a degree of polymerization of 10. However, the integration ratio of peaks from 4.2 to 4.7 ppm to the side chain methyl group (peak a, Fig. 2C) does not fit the theoretical ratio of 1:15. That is because of the two backbone hydrogens closest to the α -end of polymer has split from the main peak c in Fig. 2C and move towards larger chemical shift (peak f Fig. 2C). Peaks from 4.2 to 4.7 ppm include peak e from the initiator methylene group and peak f from the closest backbone methylene group.

Synthesis of extended linear and star shaped polymers via PFTR click reaction

As reported in our earlier work, PFTR click reaction can be utilized in preparation of step growth polymers with PFB and dithiol compounds.⁵⁵ Thus, it is natural to investigate the possibility of using this highly efficient coupling strategy to synthesize extended linear, tetra-block and star shaped polymers of EtOx and MeOx utilizing various dithiol and multi-thiol compounds.

Table 1 List of polymers, their architectures and molar mass data

No	Polymer name	Architecture	$M_{n,SEC}$ (Da)	$\bar{D}$
P1	P(EtOx) ₁₀	Linear	1200	1.12
P2	P(EtOx ₁₀)-A*-P(EtOx ₁₀)	Extended linear	2200	1.21
P3	P(EtOx ₁₀)-B*-P(EtOx ₁₀)	Extended linear	2300	1.23
P4	P(EtOx ₁₀)-C*-P(EtOx ₁₀)	Extended linear	2100	1.20
P5	D*-P(EtOx ₁₀) ₃			

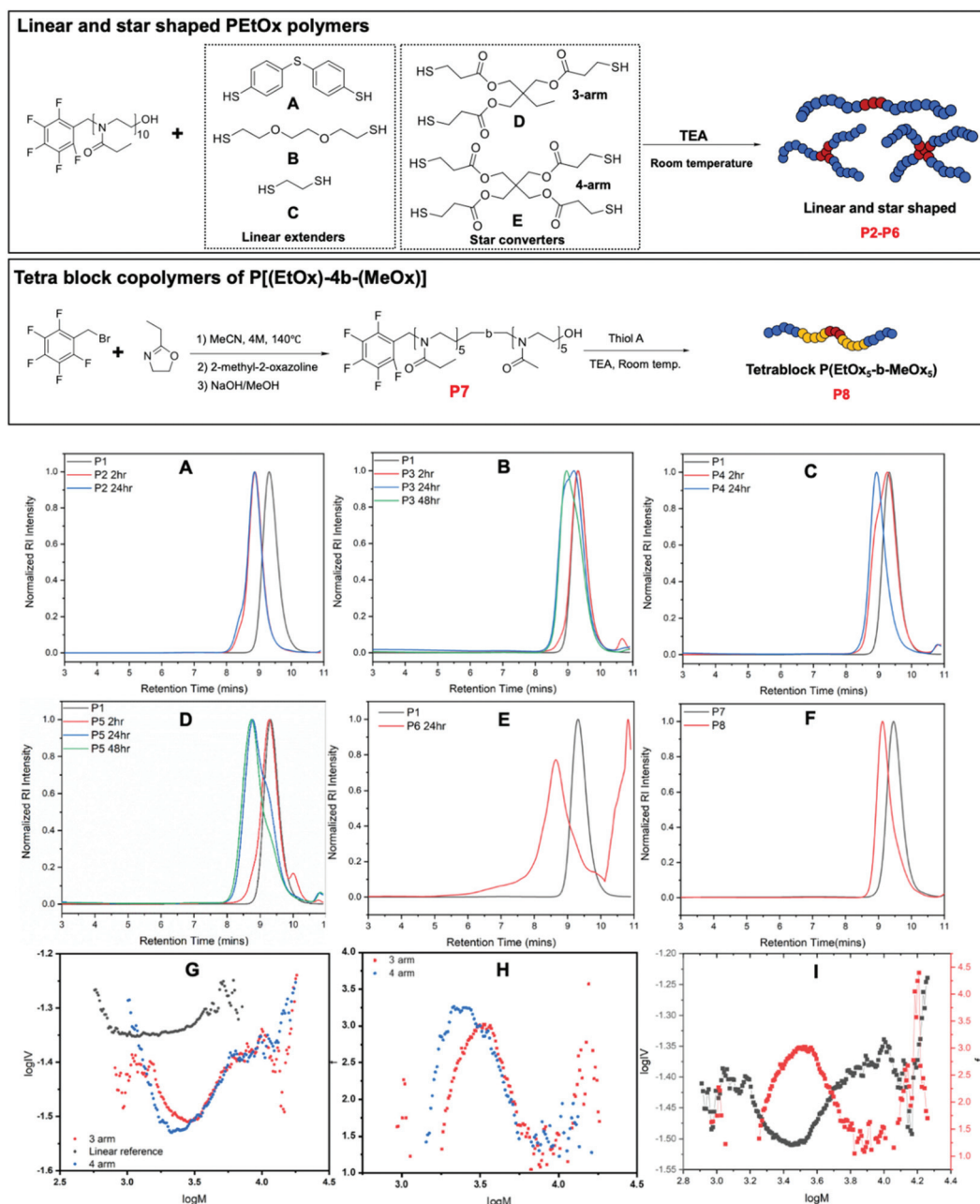
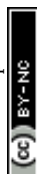


Fig. 3 Top: Reaction schemes for the synthesis of extended linear, star shaped, and tetrablock copolymers. Bottom SEC traces and advanced SEC analysis of extended linear, star shaped and tetrablock poly(2-oxazoline)s. (A–F) Corresponding SEC traces of PFTR coupling reaction of **P1**, **P7** with Thiols (A–E). (G–I) Viscosity analysis of 3-arm and 4-arm star polymers **P5** and **P6**, respectively. (G) Mark–Houwink plot for the 3 arm (**P5** in red) and 4 arm (**P6** in blue) star polymers formed via the PTFR click reaction. A reference PETox polymer **P1** is shown in black for comparison. (H) Functionality plot for both 3-arm and 4-arm star polymers showing how the number of arms varies with molar mass. Average number of arms was calculated for 3-arm and 4-arm star polymers, which are 2.2 and 2.3, respectively. (I) Overlapped Mark–Houwink plot and functionality plot of 3-arm star polymers showing the correspondence of decreased viscosity and increased number of arms.

This suggests that the reactions proceed albeit at a slower rate than dithiol A and the coupling efficiency might not be high. For dithiol C, a significant increase in hydrodynamic volume was observed after 2 hours but the overlay of SEC traces with **P1** suggests the coupling reaction was not complete (Fig. 3C). The complete shift of SEC trace was observed after 24 hours without any leftover homopolymers present.

3-Arm star polymer formation with thiol D (Fig. 3D) shows a small change in hydrodynamic volume after 2 hours, while a significant shift of the polymer peak was observed after 24 hours with a distorted curve shape that has a low molecular weight contribution. This indicates the presence polymer chains without a pentafluoro benzyl initiating group, or the tri-functional core did not fully react with pentafluoro benzyl



groups and thus generated 2-arm linear polymers. In the case of 4-arm star polymer formation with thiol E (Fig. 3E) significant increase of hydrodynamic volume was observed with after 24 hours with a tail present at high molecular weight. This might be the consequence of star-star coupling.

Preparation of block copolymers of EtOx and MeOx, P(EtOx-*b*-MeOx), was also performed using PFBB as the initiator. The synthesis of block copolymers was monitored by ^1H NMR and SEC. Collected results were shown in ESI, Fig. S2.† This serves as the proof of concept of block copolymer synthesis of poly(2-oxazoline)s using PFBB as the initiator and adds potential for easy tetra block copolymer synthesis, which was then investigated, and corresponding SEC results were shown in Fig. 3F. A significant shift of polymer peaks indicating the coupling reaction was happening and generated tetrablock polymers, although the low molecular weight tailing in the SEC trace of **P8** suggesting the existence of unreacted diblock polymers. ^1H NMR and MALDI-ToF characterization results of the coupling reaction of PFB-P(EtOx) with dithiol A is displayed in the ESI (Fig. S3†). Analysis of the results for the coupling reaction of PFB-P(EtOx) with dithiol A has shown that the coupling reaction has a good efficiency as the major distribution in the MALDI-ToF MS that was close to the calculated theoretical molecular weight of the extended linear polymer (**P2**). Additionally, aromatic protons of dithiol C have been observed in the ^1H NMR and this along with the significant change in hydrodynamic volume in the SEC chromatogram support the creation of an extended linear polymer. The existence of other desired extended linear polymers (**P3** and **P4**) was proved by MALDI-ToF characterization and is shown in Fig. S4 and S5,† respectively. Unfortunately, it was not possible to obtain a MALDI-ToF MS analysis of the tetrablock polymer **P8** P(EtOx₅-MeOx₅)₂-A; however, it was fully analyzed by ^1H NMR (Fig. S6†). It was noticeable that, although the SEC and ^1H NMR results indicate the success of coupling reactions, MALDI-ToF analysis of **P2** and **P4** show significantly deviated results with a difference between theoretical and experimental value of 3 to 7 Da. This indicates the current MALDI-ToF system, using dithranol or *trans*-2-[3-(4-*tert*-butylphenyl)-2-methyl-2-propenylidene] malononitrile (DCTB) as a matrix, sodium trifluoro acetate as a salt, and PMMA calibration standard, might not be the optimum system for analyzing these two polymers.

As the MALDI-ToF analysis of **P5** and **P6** did not provide a spectrum with acceptable signal-to-noise ratio, advanced SEC was employed to analyze obtained star polymers. Further analysis to study the degree of branching and number of arms of the obtained star polymers was performed using advanced SEC equipped with an RI and viscosity detectors. SEC viscosity analysis data for **P5** and **P6** are shown in Fig. 3G–I. Fig. 3G shows Mark–Houwink plot for both star polymers together with the linear reference polymer. Prior to any calculations for the number of arms is undertaken it is critical to qualitatively analyze the plots. This is because to obtain numerical values from intrinsic viscosity data an assumptive value for the structural factor, ϵ , is required. Firstly, if we consider the star polymer is prepared from a trifunctional core then the

maximum number of arms would be 3 and that only a 1-arm, 2-arm or 3-arm structures are possible. Therefore, it can be assumed that the minima of viscosity represents 3-arm polymers because a 1-arm and 2-arm appear equivalent to a linear reference in a Mark–Houwink plot. However, there are data points between the reference and the minima which is due to co-elution. Here, co-elution is likely because the arms only have a theoretical molecular weight of 1000 Da whereas a 3-arm star has a theoretical molecular weight of 3000 Da meaning their hydrodynamic sizes are not significantly different. Thus, leftover arms, 1-arm and 2-arm polymers will co-elute with the star polymer and will lead to intrinsic viscosity (η) values that are a weighted average of the species eluting. As will be shown later this leads to species that appear to have arms between 1–3 but this in reality is a mixture. In the same way we can qualitatively interpret the 4-arm star shaped polymer prepared from a tetrafunctional thiol core. The minima is slightly lower than that of the 3 arm star although

ions to the *para*-fluorine has been reported as an important step in this *para*-fluoro-thiol click reaction (PFTR). The termination step in the synthesis of polyoxazoline involves the use of a nucleophile to attack the oxazolinium ion to end-cap the propagating chain. This stops the chain growth and installs functional groups on omega chain end. Various nucleophiles including amines, alcohols and thiolate salts have been reported to be utilized in the end-functionalization of polyoxazolines.⁶⁴ Both termination reaction of living chain ends and PFTR are highly efficient reactions. They require a nucleophilic attack and by polymerizing a 2-oxazoline with PFBB as the initiator, the living propagating chain could be treated as a A-

B functionalized macromonomer. Thus, using a dithiol as a bi-nucleophile to simultaneously terminate living chain ends and take part in PFTR could result in the preparation of multiblock oxazoline polymers in one-pot. It should also be noted that by using sequential addition of monomers to synthesis poly(2-oxazoline) block polymers, it is also possible to easily synthesize multiblock poly(2-oxazoline)s (Fig. 4).

In order to investigate the combination of cationic chain end termination with a thiol and performing *para*-fluoro thiol click reaction simultaneously, dithiol A was selected owing to relatively faster reaction kinetics of aromatic thiols. PEtOx with [monomer] to [initiator] molar ratio of 10 to 1 was synthesized

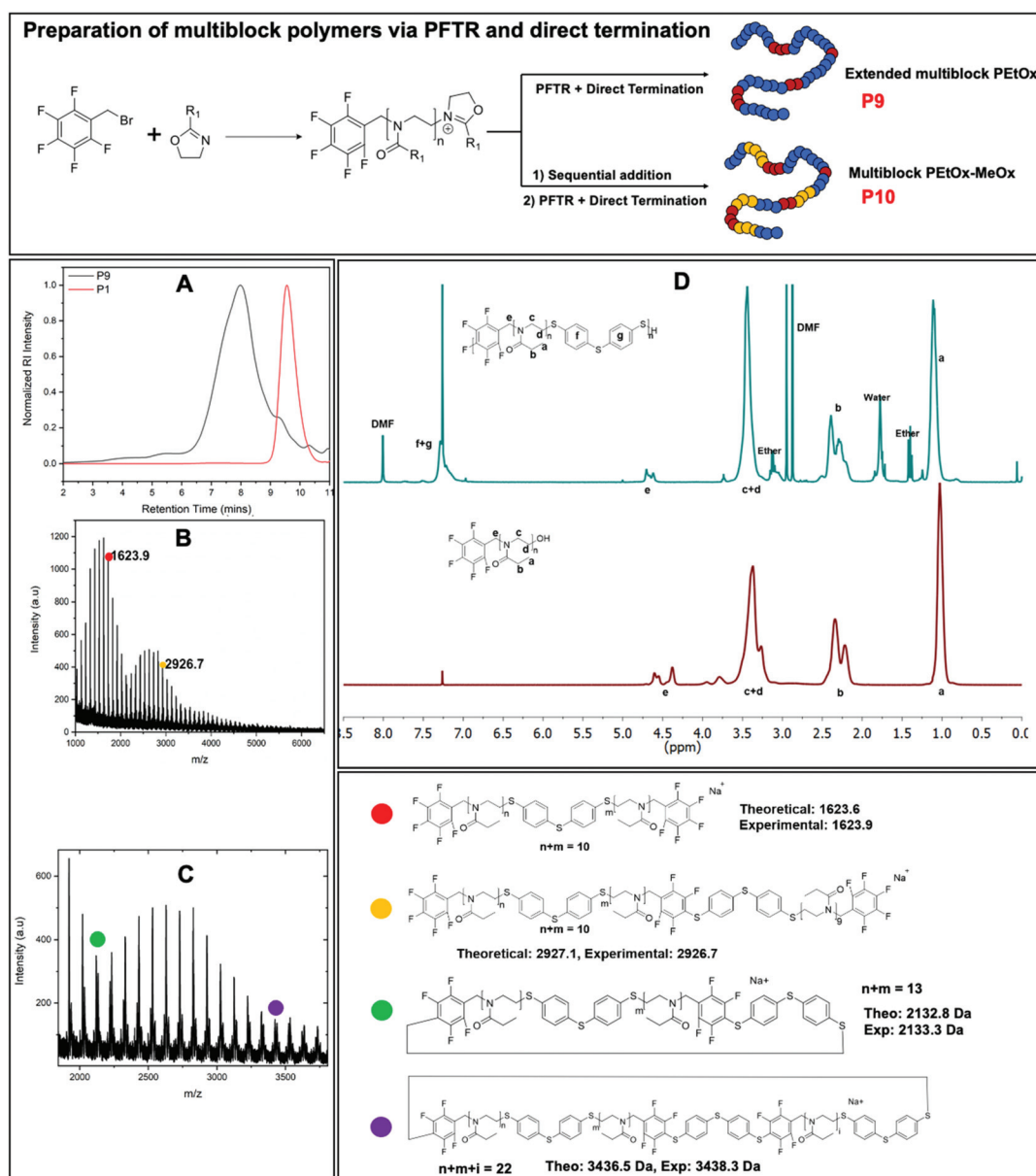


Fig. 4 Preparation of multiblock PET

under microwave heating using PFBB as the initiator. The same reaction conditions were used as discussed in the kinetic study section. After 6 minutes of polymerization the reactor was cooled down to 40 °C with compress air. Then a vacuum dried and nitrogen degassed solution of dithiol A together with dry TEA and DMPP in dry DMF was added into the polymerization reaction. The molar ratio of reactants and base were kept at [Dithiol A]:[Initiator]:[TEA]:[DMPP] = 100:100:205:1.5. The reaction mixture was then heated for another 3 minutes at 70 °C. The obtained polymer (**P9**) was analyzed with ¹H NMR, SEC and MALDI-ToF MS. A significant increase of hydrodynamic volume and average molecular weight distribution were observed (Fig. 4A), indicating the formation of a multiblock PEOx. Longer microwave heating did not lead to an increase in hydrodynamic volume, indicating that this was the full extent of reaction. The polymer was purified by precipitation in cold diethyl ether, filtration and drying in the vacuum oven. ¹H NMR analysis was carried out, which showed the presence of aromatic peaks indicative of the dithiol in the polymer backbone (Fig. 4D). MALDI-TOF MS spectra for the multiblock polymer (Fig. 4B) mainly shows peaks of low molecular weight, this is likely due to multiple aromatic and fluorinated aromatic rings making ionization of the polymer more difficult, and polymers with higher molar mass are known to be more difficult to ionize.^{65–68} Nevertheless, observed values prove that the direct termination and the PFTR click reaction are both happening and resulting in polymer growth (Fig. 4B and C). For example, as shown in Fig. 4B the mass of peak observed at 1623.9 Da fits to the sodium salt of a diblock polymer with two pentafluoro benzyl end groups that are connected *via* the termination with dithiol A and have 10 EtOx repeating units in total. Additionally, the peak observed at 2926.7 Da fits to the value of the sodium salt of the polymer at 1623.9 Da having one aromatic dithiol reacted with one pentafluoro benzyl group, and the aromatic dithiol end-caps another PEOx chain with 9 repeating units and a pentafluoro benzyl bromide group, indicates the mixed ways of chain growth presenting in the multiblock PEOx formation. The zoom in analysis of secondary distributions (Fig. 4C) revealed potential cyclic structures formation as shown with two example peaks at 2133.3 Da and 3438.3 Da, which is an inevitable side reaction in step-growth polymerization. These results are suggesting the one-pot direct termination approach for preparing multiblock PEOx was successful.

Based on the approach of synthesizing multiblock PEOx with dithiol A, preparation of multiblock P(EtOx-*b*-MeOx) (**P10**) with dithiol A was performed using the same procedure. Similar results were obtained based on the ¹H NMR and SEC analysis. (Fig. S7†) The multiblock copolymer **P10** does not have a very large hydrodynamic volume change compared to the starting block polymer of PFB-P(EtOx-MeOx). A possible reason for this might be due to the chain end livingness of the P(EtOx-*b*-MeOx) that might be hampered while adding the second block, that changed the stoichiometry of step-growth polymerization and restricts the size of final polymer.

Self-assembly properties of amphiphilic polymer **P2**, **P8**, **P9** and **P10**

The results discussed so far indicate that PFBB is a useful clickable initiator for the synthesis of desired diblock, star and multiblock polymers of PEOx utilizing α -end and ω -end functionalities. Using a microwave assisted polymerization strategy, it is possible to obtain a multiblock P(EtOx-*b*-MeOx) in one pot in 10 minutes, which is very fast and convenient. Among obtained polymers, diblock polymer **P2**, tetrablock polymer **P8** and multiblock polymer **P9** and **P10** have an amphiphilic backbone structure with aromatic and fluorinated aromatic rings. Thus, the thermal responsive and self-assembly properties of these polymers were investigated. Surprisingly, no evidence for thermal responsiveness was found in THF. Moreover, these polymers were not soluble in water which is likely due to the presence of fluorinated and non-fluorinated aromatic rings within the backbone structure. The self-assembly behavior of these polymers is discussed in following section.

The self-assembly properties of **P2**, **P8**, **P9** and **P10**

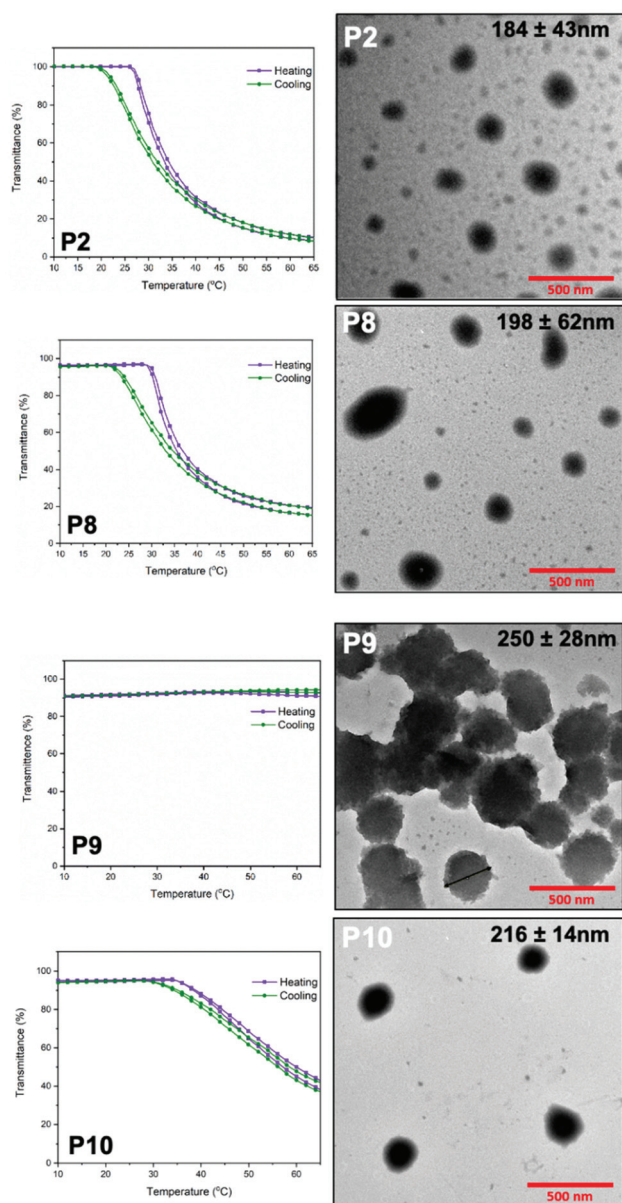


Fig. 5 Transmittance measurements at 550 nm (left) and TEM measurements (right) of P2, P8, P9 and P10.

measurements obtained at 550 nm from 10 °C to 90 °C have shown that the reversible thermal aggregation of nanoparticle solutions of **P2**, **P8** and **P10** occurs in the temperature range of 25–35 °C, indicated by the decrease in transmittance value. The nanoparticle solution of **P9** did not show a thermal aggregation behavior, most likely due to longer chain length of **P9** that provides more hydrophilic POx chains and require larger thermodynamic driving force to aggregate. Obtained transmittance results also indicate the nanoparticle aggregation of **P9** observed in TEM possibly happened during the TEM sample preparation process. The self-assembly and thermal aggregation behaviors of these nanoparticle solutions show the importance of controlled chain length and dispersity to adjust the

hydrophilicity of polymers to obtain nanoparticles that do not aggregate at a physiological temperature range.

Conclusions

In summary, 2,3,4,5,6-pentafluoro benzyl bromide was investigated as a clickable initiator for the cationic ring opening polymerization of 2-oxazoline copolymers and shown to be feasible not only for the controlled synthesis of PETox but also its further modification using *para*-fluoro-thiol click reaction. The utilization of a clickable pentafluoro benzyl group to synthesize extended linear, tetrablock and star polymers has been examined. Furthermore, we investigated the possibility of using a poly(2-oxazoline) with a pentafluoro benzyl group and utilizing the living chain end for the synthesis of multiblock polymers using a dithiol. Thus, poly(2-oxazoline) multiblock polymers were obtained and the self-assembly properties of obtained amphiphilic polymers were studied. Finally, poly(2-oxazoline) based nanoparticles sized from 184 nm to 250 nm were observed *via* TEM analysis. This work enriched the arsenal of clickable poly(2-oxazoline)s and provided access to preparation of different architecture of poly(2-oxazoline)s with a metal free, base catalyzed click reaction. As obtained multiblock polymers are not soluble in water, the partial hydrolysis of these polymers to adjust the hydrophilicity of multiblock polymers would be interesting future work as it might change the solubility as well as the self-assembly architecture of these polymers, and possibly produce useful polymers for drug delivery applications.

Author contributions

TZ has designed the project and performed the experiments, BD has contributed in the SEC analysis and GY has performed the TEM measurements. CRB has contributed in the design and supervision of the project.

Conflicts of interest

There are no conflicts to declare.

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