Design and Synthesis of a Cyclic Double-Grafted Polymer Using Active Ester Chemistry and Click Chemistry via A “Grafting onto” Method

Meng Liu 1, Lu Yin 1, Shuangshuang Zhang 1,*, Zhengbiao Zhang 1, Wei Zhang 1,*,2 and Xiulin Zhu 1,2

1 Suzhou Key Laboratory of Macromolecular Design and Precision Synthesis, Jiangsu Key Laboratory of Advanced Functional Polymer Design and Application, State and Local Joint Engineering Laboratory for Novel Functional Polymeric Materials, College of Chemistry, Chemical Engineering and Materials Science, Soochow University, Suzhou 215123, China; 20164209036@stu.suda.edu.cn (M.L.); yinlu0701@163.com (L.Y.); zhengbiaozhang@suda.edu.cn (Z.Z.); xlzhu@suda.edu.cn (X.Z.)

2 Global Institute of Software Technology, No 5. Qingshan Road, Suzhou National Hi-Tech District, Suzhou 215163, China

* Correspondence: sgny19910327@163.com (S.Z.); weizhang@suda.edu.cn (W.Z.);
Tel.: +86-512-6588-4243 (W.Z.); Fax: +86-512-6588-2787 (W.Z.)

Received: 4 January 2019; Accepted: 24 January 2019; Published: 1 February 2019

Abstract: Combing active ester chemistry and click chemistry, a cyclic double-grafted polymer was successfully demonstrated via a “grafting onto” method. Using active ester chemistry as post-functionalized modification approach, cyclic backbone (c-P2) was synthesized by reacting propargyl amine with cyclic precursor (poly(pentafluorophenyl 4-vinylbenzoate), c-PPF4VB6.5k). Hydroxyl-containing polymer double-chain (l-PS-PhOH) was prepared by reacting azide-functionalized polystyrene (l-PSN3) with 3,5-bis(propynyloxy)phenyl methanol, and further modified by azide group to generate azide-containing polymer double-chain (l-PS-PhN3). The cyclic backbone (c-P2) was then coupled with azide-containing polymer double-chain (l-PS-PhN3) via CuAAC reaction to construct a novel cyclic double-grafted polymer (c-P2-g-Ph-PS). This research realized diversity and complexity of side chains on cyclic-grafted polymers, and this cyclic double-grafted polymer (c-P2-g-Ph-PS) still exhibited narrow molecular weight distribution ($M_w/M_n < 1.10$).

Keywords: cyclic double-grafted polymer; topological architecture; active ester chemistry; “grafting onto” approach

1. Introduction

Grafted polymers, in conjunction with conventional linear polymers, have fascinating topological macromolecular structure with various side chains along their main backbones. They exhibit unique and remarkable properties by controlling chemical components of main backbones or side chains, the length of side chains, and grafting density [1,2]. Grafted polymers have been thoroughly probed as the precursors of drug-delivery materials [3], biosensors [4], nanowires [5] and nanotubes [6]. According to the structure of polymeric main backbones, grafted polymers can be roughly divided into linear-, star-, cyclic-, dendritic-, hyperbranched-grafted polymers and so on. With continuous development of new synthesis technologies, novel grafted polymers have exhibited more and more different properties and exploration for new strategies to construct novel grafting polymers have never stopped [7–15].
Comparing linear counterparts, cyclic polymers without any terminals, as another kind of topological architecture, demonstrate remarkable and unique properties, such as a smaller hydrodynamic volume, lower intrinsic viscosity, a higher glass transition temperature, higher rate of crystallization and a higher density [16–22]. In general, the research of cyclic polymers can be mainly divided into three parts: (1) constructing complex cyclic topological structure; (2) exploring new synthetic methods and strategies of cyclic polymers; (3) comparing the performance difference between cyclic polymers and linear counterparts [23–27]. Recent studies related to cyclic polymers have displayed that partial bio-materials containing cyclic structures could reduce cytotoxicity [28], increase transfection efficiency [29] or improve drug loading and releasing capacity [30,31].

Cyclic-grafted polymers consisting of one cyclic backbone and various side chains, as one kind of grafted polymers, have potential applications in biomaterials [32–34]. There are three main categories for producing cyclic-grafted polymers: (1) “grafting through” approach, the polymerization of cyclic-macromonomers [35]; (2) “grafting from” approach, the growth of side chains from a cyclic-macronitiator backbone [36–42]; (3) “grafting onto” approach, the addition of ready-made polymeric chains to a cyclic backbone by high-effective chemical reactions or supramolecular assembly, such as esterification reaction [43], click chemistry [44–48], active ester chemistry [49,50], Suzuki coupling reaction [51] and metallo-supramolecular interactions [52]. In the “grafting onto” approach, cyclic backbone and grafting chains can be independently synthesized and characterized. Diversity and complexity of side chains probably contribute to constructing functionalized cyclic topologies and exploring their potential applications.

In this work, combing active ester chemistry and click chemistry, we constructed a cyclic double-grafted polymer successfully via the “grafting onto” approach as shown in Scheme 1. Using active ester chemistry, a cyclic backbone (c-P2) was synthesized by post-functionalized modification by reacting propargyl amine with cyclic precursor (poly(pentafluorophenyl 4-vinylbenzoate), c-PPF4VB6.5k) (Scheme S1). Additionally, hydroxyl-containing polymer double-chain (l-PS-PhOH) was prepared by reacting azide-functionalized polystyrene (l-PSN3) with 3,5-bis(propynyloxy)phenyl methanol, and further modified by azide group to generate azide-containing polymer double-chain (l-PS-PhN3). The cyclic backbone (c-P2) was then coupled with prepared polymer double-chain (l-PS-PhN3) using CuAAC reaction to successfully construct a novel cyclic double-grafted polymer (c-P2-g-Ph-PS). This research realized diversity and complexity of side chains on cyclic-grafted polymers, and this cyclic double-grafted polymer (c-P2-g-Ph-PS) still exhibited narrow molecular weight distribution.

Scheme 1. Synthetic routes of a cyclic double-grafted polymer.
2. Materials and Methods

2.1. Materials

3,5-Dihydroxybenzyl alcohol (HWRK CHEM, Beijing, China, 98%), 3-(trimethylsilyl)propargyl bromide (Energy Chemical, Shanghai, China, 97%), 18-crown-6 (Sinopharm Chemical Reagent, Suzhou, China, CP), diphenyl azidophosphate (DPPA) (Alfa Aesar, Shanghai, China, 97%), 1,8-diazabicyc[5.4.0]undec-7-ene (DBU) (Tokyo Chemical Industry Co., Ltd., 98%), pentamethyldiethylenetriamine (PMDETA) (Energy Chemical, Shanghai, China, 99%) and copper(0) powder (Sinopharm Chemical Reagent, Suzhou, China, 99.9%) were purchased and used as received. Copper(I) bromide (CuBr) (Sinopharm Chemical Reagent, Suzhou, China, 99%) was washed by the mixture solution (acetic acid/deionized water, v/v = 5/95) and anhydrous ethanol many times, and then dried in a vacuum. The solvents including acetone, dichloromethane (CH$_2$Cl$_2$), hydrochloric acid (HCl), toluene, ethyl acetate (EA), petroleum ether (PE), tetrahydrofuran (THF), dimethylformamide (DMF) and methanol (MeOH) were used as received without any purification process.

2.2. Characterizations

All the $^1$H NMR and $^{13}$C NMR spectra were measured on a Bruker (300 MHz) Nuclear Magnetic Resonance spectrometer (Bruker, USA). All the average molecular weights ($M_n$) and molecular weight distributions ($M_w/M_n$) were measured by TOSOH HLC-8320 size exclusion chromatography (SEC, Tosoh Corporation, Japan). The recycling preparative HPLC Mode LC-9260NEXXT (often called as Prep-SEC, Tosoh Corporation, Japan) was utilized to purify crude polymers. A Bruker TENSOR-27 FT-IR spectrometer was utilized to measure FT-IR spectra (Bruker, USA). Matrix assisted laser desorption ionization/time of flight mass spectra (MALDI TOF MS) (Bruker, USA) were gained by using an UltrafleXtreme MALDI TOF mass spectrometer. The UV-light resource was considered using one low-pressure lamp purchased from Beijing China Education Au-light Co. Ltd (CEL-LPH120-254, 120 W, Beijing, China). All the parameters and measure conditions of these spectrometers are shown in detail in the supporting information.

2.3. Synthesis of 3,5-bis(propargyloxy)benzyl Alcohol

3,5-Dihydroxybenzyl alcohol (1.4 g, 10 mmol), 3-(trimethylsilyl)propargyl bromide (4.2 g, 22 mmol), 18-crown-6 (264 mg, 1 mmol), K$_2$CO$_3$ (3.45 g, 25 mmol) and acetone (80 mL) were added into a round-bottom flask in nitrogen atmosphere. This mixture solution was placed in oil-bath at 80 °C for 48 h. The mixed solution was filtered to remove indissoluble solid, and the filtrate was then concentrated. The solution was extracted with CH$_2$Cl$_2$ for three times and washed by 1 mol/L HCl and brine. The collected organic phase was dried using MgSO$_4$. After evaporating the solution, the crude product was purified by silica gel chromatography (eluent: petroleum ether/ethyl acetate = 4/1) to get a white solid (1.2 g, yield: 57.3%). $^1$H NMR (CDCl$_3$, 300 MHz, ppm, Figure S1): 6.63 (a, 2H), 6.54 (b, 1H), 4.67 (c, 6H), 2.53 (d, 2H), 1.70 (e, 1H). $^{13}$C NMR: (CDCl$_3$, 75 MHz, ppm): 158.92, 143.70, 106.31, 101.58, 78.49, 75.81, 65.14, 56.01.

2.4. Synthesis of Hydroxyl-Containing Polymer Double-Chain (l-PS-PhOH)

3,5-Bis(propargyloxy)benzyl alcohol (10.81 g, 0.05 mmol), azide-functional polystyrene (l-PS-N$_3$, 257.5 mg, 0.103 mmol), PMDETA (17.84 mg, 0.103 mmol), toluene (5 mL) and a magnetic stirrer were added into a 10 mL ampoule in nitrogen atmosphere. CuBr (14.8 mg, 0.103 mmol) and Cu (3.18 mg, 0.05 mmol) were added into above ampoule. The mixture solution was stirred at ambient temperature for 1.5 h in nitrogen atmosphere. After that, the polymer was precipitated in anhydrous methanol and dried in a vacuum (249.4 mg, yield: 92.9%). The crude polymer double-chain was purified by Prep-SEC to get hydroxyl-containing polymer double-chain (l-PS-PhOH, $M_{n,SEC}$ = 5000 g/mol, $M_w/M_n$ = 1.04).
2.5. Synthesis of Azide-Containing Polymer Double-Chain (l-PS-PhN₃)

Hydroxyl-containing polymer double-chain (l-PS-PhOH, 100 mg, 0.02 mmol) was dissolved in DMF (1 mL) and put in an ampoule (5 mL) containing a magnetic stirrer in nitrogen atmosphere. The ampoule was wrapped in aluminum foil to avoid light. DPPA (110.08 mg, 0.4 mmol) and DBU (60.90 mg, 0.4 mmol) were added into the ampoule under nitrogen atmosphere. The ampoule was placed in an oil-bath at 80 °C for 24 h. The mixed solution was purified by passing through a short Al₂O₃ column, precipitated in anhydrous methanol and dried under vacuum. (l-PS-PhN₃, 93.4 mg, yield: 93.4%, \(M_{n,SEC} = 5000 \text{ g/mol}, M_w/M_n = 1.04\)).

2.6. Synthesis of Cyclic Double-Grafted Polymer (c-P2-g-Ph-PS)

Cyclic polymer (c-P₂, 2.3 mg, \(5 \times 10^{-4}\) mmol), linear polymer (l-PS-PhN₃, 81 mg, \(1.62 \times 10^{-2}\) mmol), PMDETA (5.62 mg, \(3.24 \times 10^{-2}\) mmol), mixture solvent (THF = 2 L, DMF = 1 mL) and a magnetic stirrer were added into a 10 mL ampoule in nitrogen atmosphere. CuBr (4.64 mg, \(3.24 \times 10^{-2}\) mmol) and Cu (1.72 mg, \(2.7 \times 10^{-2}\) mmol) were added into above ampoule. The solution was reacted at ambient temperature in nitrogen atmosphere. After 24 h, the polymer was precipitated in anhydrous methanol and dried in a vacuum. The crude double-grafted polymer was further purified by Prep-SEC to get final cyclic double-grafted polymer (c-P₂-g-Ph-PS, 33.6 mg, yield: 40.33%, \(M_{n,SEC} = 30,700 \text{ g/mol}, M_w/M_n = 1.04\)).

3. Results and Discussion

3.1. Synthesis of l-PS-PhOH and l-PS-PhN₃

Hydroxyl-containing polymeric double-chain (l-PS-PhOH) was synthesized by reacting 3,5-bis(propargyloxy)benzyl alcohol (Scheme 2) and azide-functionalized polystyrene (l-PS-N₃) by virtue of Copper-catalyzed azide/alkyne cycloaddition (CuAAC) reaction. The synthesis and characterization of l-PS-N₃ (\(M_{n,SEC} = 2500 \text{ g/mol}, M_w/M_n = 1.05\)) was shown in our previous publication [50]. The usage of slightly excessive l-PS-N₃ was necessary in the process of preparing l-PS-PhOH, the gained crude l-PS-PhOH needs to be easily purified by Prep-SEC.

![Scheme 2](image.png)

Scheme 2. The structure of 3,5-bis(propargyloxy)benzyl alcohol.

Hydroxyl-containing polymer double-chain (l-PS-PhOH) was verified by SEC, NMR, MALDI TOF MS and FT-IR spectroscopy. As shown in Figure 1, corresponding to l-PS-N₃, \(^1\)H NMR spectra of l-PS-PhOH showed that the characteristic signal of the methine hydrogen (–CH(Ph)–, \(f\)) shifted from 3.8–4.1 ppm to 4.9–5.2 ppm completely. A new peak was clearly observed at 4.6 ppm, which was assigned to the benzylic hydrogen (–CH₂–, \(i\)). In addition, the \((f+h)/i/b\) integration ratio is close to 6/2/4, which means the successful formation of l-PS-PhOH. The number average molecular weight of l-PS-PhOH (\(M_{n,SEC} = 5000 \text{ g/mol}, M_w/M_n = 1.04\)) was twice than that of l-PS-N₃ (\(M_{n,SEC} = 2500 \text{ g/mol}, M_w/M_n = 1.05\)), and the molecular weight distribution remained at 1.04, which also indicated the successful preparation of l-PS-PhOH. MALDI TOF MS (Figure 3) provided direct and powerful evidence for the formation of polymeric double-chain. The typical experimental peak \(m/z\) value (4715.97 Da) accords with the theoretical calculating value ([40M+Na]⁺, 4715.74 Da), in accordance with 40 repeat units of l-PS-PhOH with one sodium cation. The difference value of two adjacent experimental peaks is consistent with the \(m/z\) value of a styrene. Furthermore, in Figure 4, the complete disappearance of signals from azide groups (2094 cm\(^{-1}\)) also proved the formation of l-PS-PhOH. All the above results confirmed the successful preparation of hydroxyl-containing polymer double-chain (l-PS-PhOH) without residue of l-PS-N₃.
Figure 1. $^1$H NMR (300 MHz) spectra of l-PS-N$_3$ and l-PS-PhOH in CDCl$_3$.

![1H NMR spectra](image1)

Figure 2. Size exclusion chromatography (SEC) curves of l-PS-N$_3$ and l-PS-PhOH. Tetrahydrofuran (THF) as the eluent and PS as calibration standards.

![SEC curves](image2)

Figure 3. Expanded matrix assisted laser desorption ionization/time of flight mass spectra (MALDI-TOF MS) of l-PS-PhOH with the inserted full spectrum.

![MALDI-TOF MS](image3)
Azide-containing polymer double-chain (l-PS-PhN3) was synthesized by hydroxyl-containing polymer double-chain (l-PS-PhOH) under the system of DPPA/DBU mixtures. Azide-containing polymer double-chain (l-PS-PhN3) was verified by SEC, NMR, MALDI TOF MS and FT-IR spectroscopy. Comparing to the spectrum of l-PS-PhOH (Figure 1), 1H NMR spectrum of l-PS-PhN3 (Figure 5) demonstrated that the benzylic hydrogen (–CH2–, i) shifted from 4.6 to 4.2 ppm completely, which indicated the complete formation of azide-containing polymeric double-chain (l-PS-PhN3). After azidation, the (f+h)/i/b integration ratio still kept at 6/2/4, which also indicated the successful preparation of l-PS-PhN3. From SEC curves (Figures 2 and 6), there are no obvious changes before and after azidation. The average molecular weight of l-PS-PhN3 is 4900 g/mol and the molecular weight distribution remained at 1.03. In FT-IR spectrum (Figure 4), the vibrational absorption peak from azide group of l-PS-PhN3 appeared at 2094 cm⁻¹. MALDI TOF MS provided persuasive evidence for successful formation of l-PS-PhN3. Figure 7 exhibited two main peak distributions, accurately assigned to polymeric double-chain (l-PS-PhN3). The typical experimental peak m/z value (such as 4712.91 Da), is consistent with the theoretical calculating value ([40 M-N₂ + Na]⁺, 4712.74 Da). The difference value of two adjacent experimental peaks (103.94 Da) accords with the value of one styrene. These results demonstrated the successful preparation of l-PS-PhN3.

Figure 4. FT-IR spectra of l-PS-N3, l-PS-PhOH and l-PS-PhN3.

3.2. Synthesis of l-PS-PhN3 and c-P2-g-Ph-PS

Figure 5. 1H NMR spectra of l-PS-PhN3 and c-P2-g-Ph-PS in CDCl₃.
The active ester chemistry, such as the nucleophilic substitution of activated ester bearing pentafluorophenyl groups with diverse amines, is one kind of high-effective chemical reaction, which is often utilized as one post-modification technology for constructing functional polymers that cannot be obtained by conventional polymerization technologies [53]. Here, we chose propargylamine as the amine to react with cyclic poly(pentafluorophenyl 4-vinylbenzoate) (c-PPF4VB6.5k) for synthesizing functional cyclic polymer (c-P2). All the synthesis and characterizations of c-PPF4VB6.5k and c-P2 are shown in supporting information in detail (Figures S2–S8).

Furthermore, functional cyclic polymer (c-P2) was used as cyclic polymeric backbone to react with polymer double-chain (l-PS-PhN3) via CuAAC reaction for constructing cyclic double-grafted polymer (c-P2-g-Ph-PS). The crude cyclic double-grafted polymer (c-P2-g-Ph-PS) was purified by Prep-SEC and further characterized by NMR and SEC. As shown in Figure 5, 1H NMR spectrum of c-P2-g-Ph-PS indicated the successful preparation of cyclic double-grafted polymer by the integration ratio from 1H NMR spectrum, but the difference between l-PS-PhN3 and c-P2-g-Ph-PS suggested the successful preparation of cyclic double-grafted polymer. Additionally, according to the SEC curves of c-P2, l-PS-PhN3 and c-P2-g-Ph-PS (Figure 6), the shifts toward high molecular weight field can be observed clearly, demonstrating the successful formation of cyclic double-grafted polymer. The molecular weight of cyclic double-grafted polymer (c-P2-g-Ph-PS) was 30,700 g/mol and the molecular weight still stayed at 1.04.

![Figure 6. SEC curves of c-P2, l-PS-PhN3 and c-P2-g-Ph-PS. THF as the eluent and PS as calibration standards.](image-url)
4. Conclusions

A novel cyclic topological architecture, a cyclic double-grafted polymer, was successfully constructed using active ester chemistry and click chemistry via a “grafting onto” method. Cyclic backbone (c-P2) was synthesized by reacting propargyl amine with cyclic precursor (c-PPF4VB6.5k) using active ester chemistry as a post-modification approach. Hydroxyl-containing polymer double-chain ([l-PS-PhOH]) was prepared by reacting azide-functionalized polymer chain ([l-PSN3]) with 3,5-bis(propynloxy)phenyl methanol, and further azide-modified to generate azide-containing polymer double-chain ([l-PS-PhN3]) and well characterized by SEC, NMR and MALDI TOF MS. Finally, this cyclic backbone (c-P2) was coupled with azide-containing polymer double-chain ([l-PS-PhN3]) using CuAAC reaction to successfully construct a novel cyclic double-grafted polymer (c-P2-g-Ph-PS). Notably, this cyclic double-grafted polymer (c-P2-g-Ph-PS) still exhibited a narrow molecular weight distribution. On the basis of our previous work, this research realized diversity and complexity of side chains from cyclic-grafted polymers, which could eventually enrich the topological architecture and provide a new platform for constructing amphiphilic cyclic-brush polymers with amphiphilic polymeric double-chains along the cyclic backbone.

Supplementary Materials: The following are available online at http://www.mdpi.com/2073-4360/11/2/240/s1. Scheme S1. Illustration of synthetic routes of cyclic polymer (c-PPF4VB6.5k) and linear polystyrene (l-PS-N3), Figure S1 1H NMR (300 MHz) spectrum of 3,5-bis(propargyloxy)benzyl alcohol; Figure S2. 1H NMR (300 MHz) spectra of linear polymer (l-PPF4VB6.5k) and cyclic polymer (c-PPF4VB6.5k); Figure S3. 13C NMR (75 MHz) spectrum of linear polymer (l-PPF4VB4.0k); Figure S4. SEC curves of l-PPF4VB4.0k, c-PPF4VB4.0k and c-P2. THF was used as the eluent and PS standards for the calibration; Figure S5. Expanded MALDI-TOF mass spectrum of l-PPF4VB6.5k with inserted full spectrum; Figure S6. UV-vis spectrum of RAFT agent, l-PPF4VB6.5k and c-PPF4VB6.5k; Figure S7. Pictures of l-PPF4VB6.5k and c-PPF4VB6.5k; Figure S8. 1H NMR (300 MHz) spectrum of cyclic polymer (c-P2).

Author Contributions: M.L., L.Y. and S.Z. conducted all the experiments; S.Z. analyzed all the research data and wrote this manuscript alone; Z.Z., X.Z. and W.Z. designed the research content and provided suggestions for revising this manuscript.

Acknowledgments: The financial supports from the National Nature Science Foundation of China (21234005) and the Priority Academic Program Development of Jiangsu Higher Education Institutions (PAPD) are gratefully acknowledged.

Conflicts of Interest: All the authors declare that there are no conflict of interest for this research.
References

1. Feng, C.; Li, Y.; Yang, D.; Hu, J.; Zhang, X.; Huang, X. Well-defined graft copolymers: From controlled synthesis to multipurpose applications. Chem. Soc. Rev. 2011, 40, 1282–1295. [CrossRef] [PubMed]

2. Sheiko, S.S.; Sumerlin, B.S.; Matyjaszewski, K. Cylindrical molecular brushes: Synthesis, characterization, and properties. Prog. Polym. Sci. 2008, 33, 759–785. [CrossRef]

3. Johnson, J.A.; Lu, Y.Y.; Burts, A.O.; Xia, Y.; Durrell, A.C.; Tirrell, D.A.; Grubbs, R.H. Drug-loaded, bivalent-bottle-brush polymers by graft-through ROMP. Macromolecules 2010, 43, 10326–10335. [CrossRef] [PubMed]

4. Heinrich, C.D.; Thelakkat, M. Poly-(3-hexylthiophene) bottlebrush copolymers with tailored side-chain lengths and high charge carrier mobilities. J. Mater. Chem. C 2016, 4, 5370–5378. [CrossRef]

5. Yuan, J.; Xu, Y.; Walther, A.; Bolisetty, S.; Schumacher, M.; Schmalz, H.; Ballauff, M.; Müller, A.H. Water-soluble organo-silica hybrid nanowires. Nat. Mater. 2008, 7, 718–722. [CrossRef] [PubMed]

6. Huang, K.; Rzayev, J. Well-defined organic nanotubes from multicomponent bottlebrush copolymers. J. Am. Chem. Soc. 2009, 131, 6880–6885. [CrossRef] [PubMed]

7. Fischer, K.; Schmidt, M. Solvent-Induced Length Variation of Cylindrical Brushes. Macromol. Rapid Commun. 2001, 22, 787–791. [CrossRef]

8. Gu, L.; Shen, Z.; Feng, C.; Li, Y.; Lu, G.; Huang, X.; Wang, G.; Huang, J. Synthesis of PPEGMEA-γ-PMAA densely grafted double hydrophilic copolymer and its use as a template for the preparation of size-controlled superparamagnetic Fe₃O₄/polymer nano-composites. J. Mater. Chem. 2008, 18, 4332–4340. [CrossRef]

9. Xu, Y.; Bolisetty, S.; Ballauff, M.; Müller, A.H.E. Switching the morphologies of cylindrical polycation brushes by ionic and supramolecular inclusion complexes. J. Am. Chem. Soc. 2009, 131, 1640–1641. [CrossRef] [PubMed]

10. Lu, G.; Liu, H.; Gao, H.; Feng, C.; Li, Y.; Huang, X. Construction of semi-fluorinated amphiphilic graft copolymer bearing a poly(2-methyl-1,4-bistrifluorovinyloxybenzene) backbone and poly(ethylene glycol) side chains via the grafting-onto strategy. RSC Adv. 2015, 5, 39668–39676. [CrossRef]

11. Xu, B.; Gu, G.; Feng, C.; Jiang, X.; Hu, J.; Lu, G.; Zhang, S.; Huang, X. (PAA-γ-PS)-co-PPEGMEMA asymmetric polymer brushes: Synthesis, self-assembly, and encapsulating capacity for both hydrophobic and hydrophilic agents. Polym. Chem. 2016, 7, 613–624. [CrossRef]

12. Xu, B.; Liu, Y.; Sun, X.; Hu, J.; Shi, P.; Huang, X. Semifluorinated Synergistic Nonfouling/Fouling-Release Surface. ACS Appl. Mater. Interfaces 2017, 9, 16517–16523. [CrossRef] [PubMed]

13. Feng, C.; Huang, X. Polymer brushes: Efficient synthesis and applications. Acc. Chem. Res. 2018, 51, 2314–2323. [CrossRef] [PubMed]

14. Chan, S.Y.; Choo, W.S.; Young, D.J.; Lohn, X.J. Thixotropic supramolecular pectin-poly(ethylene glycol) methacrylate (PEGMA) hydrogels. Polymers 2016, 8, 404. [CrossRef]

15. Wang, Y.; Baily, T.S.; Hong, M.; Chen, E.Y.-X. Stereoregular brush polymers and graft copolymers by chiral zirconocene-mediated coordination polymerization of P3HT macromers. Polymers 2017, 9, 139. [CrossRef]

16. Shin, E.J.; Jeong, W.; Brown, H.A.; Koo, B.J.; Hedrick, J.L.; Waymouth, R.M. Crystallization of cyclic polymers: Synthesis and crystallization behavior of high molecular weight cyclic poly(ε-caprolactone)s. Macromolecules 2011, 44, 2773–2779. [CrossRef]

17. Lee, C.-U.; Li, A.; Ghale, K.; Zhang, D. Crystallization and melting behaviors of cyclic and linear polypeptoids with alkyl side chains. Macromolecules 2013, 46, 8213–8223. [CrossRef]

18. Clarson, S.J.; Semlyen, J.A. Cyclic polysiloxanes: 1. Preparation and characterization of poly(phenylmethylsiloxane). Polymer 1986, 27, 1633–1636. [CrossRef]

19. Guo, L.; Zhang, D. Cyclic poly(α-peptoid)s and their block copolymers from N-heterocyclic carbene-mediated ring-opening polymerizations of N-substituted N-carboxyloxazolidinones. J. Am. Chem. Soc. 2009, 131, 18072–18074. [CrossRef]

20. Culkin, D.A.; Jeong, W.; Cshony, S.; Gomez, E.D.; Balsara, N.P.; Hedrick, J.L.; Waymouth, R.M. Zwitterionic Polymerization of Lactide to Cyclic Poly(Lactide) by Using N-Heterocyclic Carbene Organocatalysts. Angew. Chem. Int. Ed. 2007, 46, 2627–2630. [CrossRef]

21. Bielawski, C.W.; Benitez, D.; Grubbs, R.H. An “endless” route to cyclic polymers. Science 2002, 297, 2041–2044. [CrossRef] [PubMed]
22. Qiu, X.-P.; Tanaka, F.; Winnik, F.M. Temperature-induced phase transition of well-defined cyclic poly(N-isopropylacrylamide) in aqueous solution. *Macromolecules* 2007, 40, 7069–7071. [CrossRef]
23. Laurent, B.A.; Grayson, S.M. Synthetic approaches for the preparation of cyclic polymers. *Chem. Soc. Rev.* 2009, 38, 2202–2213. [CrossRef] [PubMed]
24. Jia, Z.; Monteiro, M.J. Cyclic polymers: Methods and strategies. *J. Polym. Sci. Part A Polym. Chem.* 2012, 50, 2085–2097. [CrossRef]
25. Tezuka, Y. Topological polymer chemistry for designing multicyclic macromolecular architectures. *Polym. J.* 2012, 44, 1159–1169. [CrossRef]
26. Hoskins, J.N.; Grayson, S.M. Cyclic polyesters: Synthetic approaches and potential applications. *Polym. Chem.* 2011, 2, 289–299. [CrossRef]
27. Josse, T.; De Winter, J.; Gerbaut, P.; Coulembier, O. Cyclic Polymers by Ring-Closure Strategies. *Angew. Chem. Int. Ed.* 2016, 55, 13944–13958. [CrossRef]
28. Wei, H.; Chu, D.S.H.; Zhao, J.; Pahang, J.A.; Pun, S.H. Synthesis and evaluation of cyclic cationic polymers for nucleic acid delivery. *ACS Macro Lett.* 2013, 2, 1047–1050. [CrossRef]
29. Cortez, M.A.; Godbey, W.T.; Fang, Y.; Payne, M.E.; Cafferty, B.J.; Kosakowska, K.A.; Grayson, S.M. The synthesis of cyclic poly(ethylene imine) and exact linear analogues: An evaluation of gene delivery comparing polymer architectures. *J. Am. Chem. Soc.* 2015, 137, 6541–6549. [CrossRef]
30. Wan, X.; Liu, T.; Liu, S. Synthesis of amphiphilic tadpole-shaped linear-cyclic diblock copolymers via ring-opening polymerization directly initiating from cyclic precursors and their application as drug nanocarriers. *Biomacromolecules* 2011, 12, 1146–1154. [CrossRef]
31. Chen, B.; Jerger, K.; Fréchet, J.M.J.; Szoka, F.C., Jr. The influence of polymer topology on pharmacokinetics: Differences between cyclic and linear PEGylated poly(acrylic acid) comb polymers. *J. Contr. Release* 2009, 140, 203–209. [CrossRef] [PubMed]
32. Wei, H.; Wang, C.E.; Tan, N.; Boydston, A.J.; Pun, S.H. ATRP synthesis of sunflower polymers using cyclic multimacrorinitiators. *ACS Macro Lett.* 2015, 4, 938–941. [CrossRef] [PubMed]
33. Wang, C.E.; Wei, H.; Tan, N.; Boydston, A.J.; Pun, S.H. Sunflower polymers for folate-mediated drug delivery. *Biomacromolecules* 2015, 17, 69–75. [CrossRef] [PubMed]
34. Zhang, S.; Tezuka, Y.; Zhang, Z.; Li, N.; Zhang, W.; Zhu, X. Recent advances in the construction of cyclic grafted polymers and their potential applications. *Polym. Chem.* 2018, 9, 677–686. [CrossRef]
35. Xia, Y.; Boydston, A.J.; Grubbs, R.H. Synthesis and direct imaging of ultrahigh molecular weight cyclic brush polymers. *Angew. Chem. Int. Ed.* 2011, 50, 5882–5885. [CrossRef] [PubMed]
36. Jia, Z.; Fu, Q.; Huang, J. Synthesis of amphiphilic macromacrophase graft copolymers consisting of a poly(ethylene oxide) ring and multi-polystyrene lateral chains. *Macromolecules* 2006, 39, 5190–5193. [CrossRef]
37. Pang, X.; Wang, G.; Jia, Z.; Liu, C.; Huang, J. Preparation of the amphiphilic macro-rings of poly(ethylene oxide) with multi-polystyrene lateral chains and their extraction for dyes. *J. Polym. Sci. Part A Polym. Chem.* 2007, 45, 5824–5837. [CrossRef]
38. Fan, X.; Wang, G.; Huang, J. Synthesis of macrocyclic molecular brushes with amphiphilic block copolymers as side chains. *J. Polym. Sci. Part A Polym. Chem.* 2011, 49, 1361–1367. [CrossRef]
39. Zhang, K.; Tew, G.N. Cyclic brush polymers by combining ring-expansion metathesis polymerization and the ‘grafting from’ technique. *ACS Macro Lett.* 2012, 1, 574–579. [CrossRef]
40. Yao, W.; Li, Y.; Feng, C.; Lu, G.; Huang, X. Synthesis of a sun-shaped amphiphilic copolymer consisting of a cyclic perfluorocyclobutyl aryl ether-based backbone and lateral PMAA side chains. *RSC Adv.* 2014, 4, 52105–52116. [CrossRef]
41. Zhang, S.; Yin, L.; Zhang, W.; Zhang, Z.; Zhu, X. Synthesis of diverse cyclic-brush polymers with cyclic polystyrene as a universal template via a grafting-from approach. *Polym. Chem.* 2016, 7, 2112–2120. [CrossRef]
42. Tu, X.; Meng, C.; Liu, Z.; Sun, L.; Zhang, X.; Zhang, M.; Sun, M.; Ma, L.; Liu, M.; Wei, H. Synthesis and phase transition of poly(N-isopropylacrylamide)-based thermo-sensitive cyclic brush polymer. *Polymers* 2017, 9, 301. [CrossRef]
43. Li, H.; Jérôme, R.; Lecomte, P. Amphiphilic sun-shaped polymers by grafting macrocyclic copolyesters with PEO. *Macromolecules* 2008, 41, 650–654. [CrossRef]
44. Zhang, K.; Lackey, M.A.; Wu, Y.; Tew, G.N. Universal cyclic polymer templates. *J. Am. Chem. Soc.* 2011, 133, 6906–6909. [CrossRef] [PubMed]
45. Lahasky, S.H.; Serem, W.K.; Guo, L.; Garno, J.C.; Zhang, D. Synthesis and characterization of cyclic brush-like polymers by N-heterocyclic carbene-mediated zwitterionic polymerization of N-propargyl N-carboxyanhydride and the grafting-to approach. *Macromolecules* 2011, 44, 9063–9074. [CrossRef]

46. Li, A.; Lu, L.; Li, X.; He, L.; Do, C.; Garno, J.C.; Zhang, D. Amidine-mediated zwitterionic ring-opening polymerization of N-alkyl N-carboxyanhydride: Mechanism, kinetics, and architecture elucidation. *Macromolecules* 2016, 49, 1163–1171. [CrossRef]

47. Du, P.; Li, A.; Li, X.; Zhang, Y.; Do, C.; He, L.; Rick, S.W.; John, V.T.; Kumar, R.; Zhang, D. Aggregation of cyclic polypeptoids bearing zwitterionic end-groups with attractive dipole–dipole and solvophobic interactions: A study by small-angle neutron scattering and molecular dynamics simulation. *Phys. Chem. Chem. Phys.* 2017, 19, 14388–14400.

48. Cai, T.; Yang, W.J.; Neoh, K.-G.; Kang, E.-T. Preparation of jellyfish-shaped amphiphilic block-graft copolymers consisting of a poly(e-caprolactone)-block-poly(pentafluorostyrene) ring and poly (ethylene glycol) lateral brushes. *Polym. Chem.* 2012, 3, 1061–1068. [CrossRef]

49. Wang, D.; Xiao, L.; Zhang, X.; Zhang, K.; Wang, Y. Emulsion templating cyclic polymers as microscopic particles with tunable porous morphology. *Langmuir* 2016, 32, 1460–1467. [CrossRef]

50. Zhang, S.; Cheng, X.; Wang, J.; Zhang, Z.; Zhang, W.; Zhu, X. Synthesis of a cyclic-brush polymer with a high grafting density using active ester chemistry via the “grafting onto” approach. *Polym. Chem.* 2018, 9, 5155–5163. [CrossRef]

51. Wang, Y.; Zhang, S.; Wang, L.; Zhang, W.; Zhou, N.; Zhang, Z.; Zhu, X. The Suzuki coupling reaction as a post-polymerization modification: A promising protocol for construction of cyclic-brush and more complex polymers. *Polym. Chem.* 2015, 6, 4669–4677. [CrossRef]

52. Zhang, K.; Zha, Y.; Peng, B.; Chen, Y.; Tew, G.N. Metallo-supramolecular cyclic polymers. *J. Am. Chem. Soc.* 2013, 135, 15994–15997. [CrossRef] [PubMed]

53. Das, A.; Theato, P. Activated ester containing polymers: Opportunities and challenges for the design of functional macromolecules. *Chem. Rev.* 2016, 116, 1434–1495. [CrossRef] [PubMed]