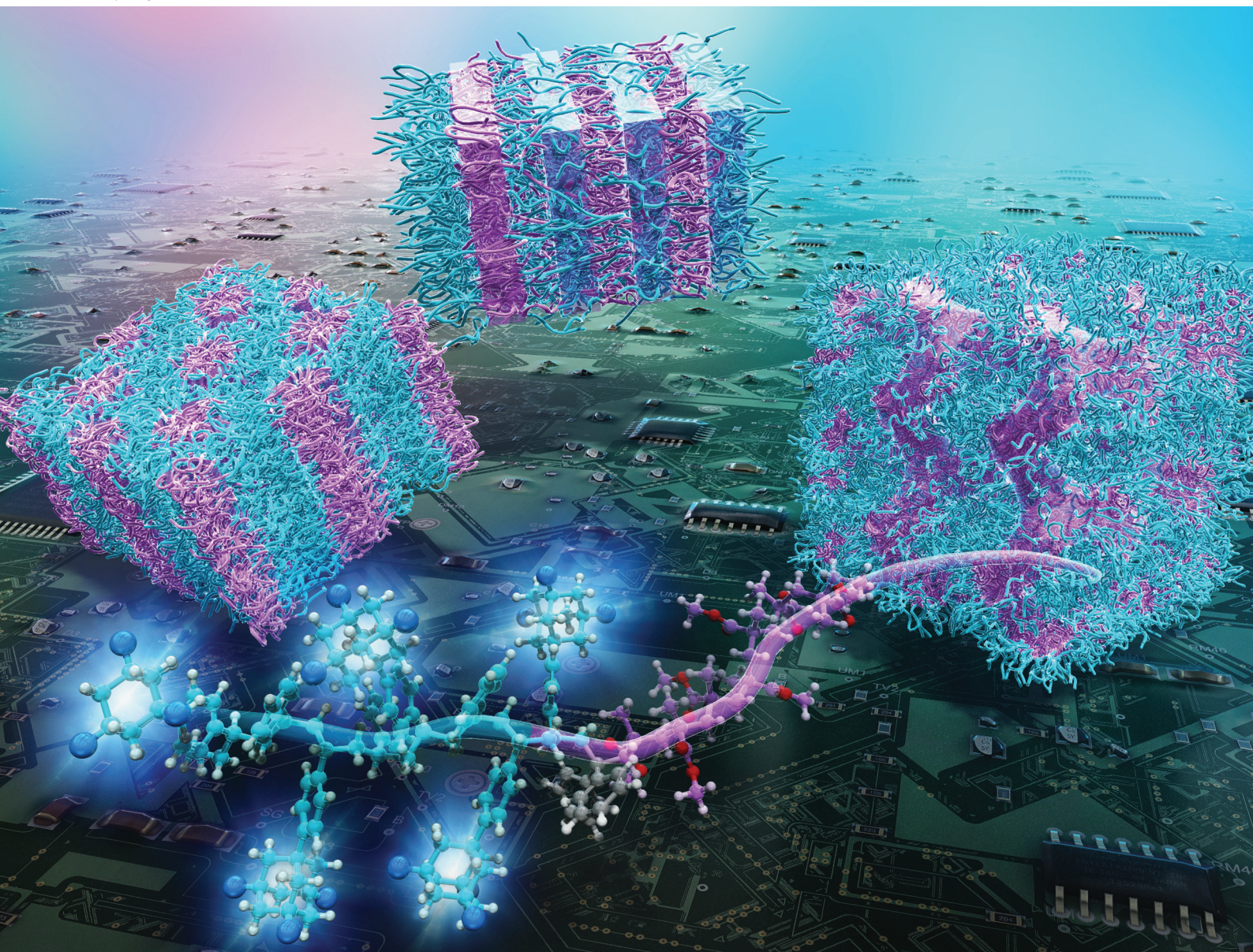


# Polymer Chemistry

Volume 14  
Number 22  
14 June 2023  
Pages 2655-2764

rsc.li/polymers



ISSN 1759-9962

## PAPER










Takuya Isono, Toshifumi Satoh *et al.*

Installation of the adamantyl group in polystyrene-*block*-poly(methyl methacrylate) *via* Friedel-Crafts alkylation to modulate the microphase-separated morphology and dimensions



Cite this: *Polym. Chem.*, 2023, **14**,  
2675

# Installation of the adamantyl group in polystyrene-*block*-poly(methyl methacrylate) *via* Friedel–Crafts alkylation to modulate the microphase-separated morphology and dimensions†

Takuya Isono, \*<sup>a</sup> Ema Baba, <sup>‡</sup> Shunma Tanaka, <sup>‡</sup> Ken Miyagi, <sup>c</sup> Takahiro Dazai, <sup>c</sup> Feng Li, <sup>a</sup> Takuya Yamamoto, <sup>a</sup> Kenji Tajima <sup>a</sup> and Toshifumi Satoh \*<sup>a</sup>

Although polystyrene-*block*-poly(methyl methacrylate) (PS-*b*-PMMA) is the most widely employed template and scaffold for constructing nanoscale patterns, its relatively small Flory–Huggins interaction parameter makes it challenging to realize microphase-separated structures with a periodicity (or domain spacing) of less than 20 nm. The direct post-polymerization modification of PS-*b*-PMMA can resolve this issue. In this study, we present the PS block post-polymerization modification of commercially available PS-*b*-PMMA through a metal-free Friedel–Crafts alkylation reaction with adamantanols to modulate its microphase-separated morphology and dimensions. The adamantyl group can be easily installed on the PS benzene ring, which increases the incompatibility with the PMMA block, enabling ordered microphase-separated structures, including lamellar, gyroid and hexagonally close-packed cylindrical structures, from low-molecular-weight PS-*b*-PMMA. Notably, a periodicity as low as approximately 16 nm was realized with this strategy, which is much lower than the lowest accessible periodicity of pristine PS-*b*-PMMA. Furthermore, we observed additional benefits from the installation of the adamantyl group, such as increased thermal stability, glass transition temperature and etching contrast.

Received 1st February 2023,  
Accepted 24th March 2023

DOI: 10.1039/d3py00113j

rsc.li/polymers

## Introduction

Block copolymers (BCPs) consisting of two chemically incompatible polymer blocks form periodic nanostructures known as microphase-separated structures. Depending on the total degree of polymerization ( $N$ ) and the volume fraction, BCPs show microphase-separated structures with different periodicities ranging from a few to hundreds of nanometers and different morphologies, such as lamellar, gyroid, hexagonally close-packed cylinders and body-centered cubic spheres.<sup>1–3</sup> Such tunable nanostructures have a wide range of applications, with the most important one being their use as structural templates for nanofabrication.<sup>3–6</sup> For example, applying perpendicularly orientated lamellar structures to a semi-

conductor substrate can allow fabrication of line-and-space patterns on the underlying substrate by the selective removal of one of the blocks and subsequent etching.<sup>7</sup> Perpendicularly orientated cylinders can also be used for fabricating holes and pillars on an underlying substrate.<sup>8,9</sup> Meanwhile, gyroid structures have been used as nanotemplates to create nanofiltration membranes,<sup>10</sup> efficient photovoltaics,<sup>11</sup> photonic crystals,<sup>12</sup> etc.<sup>13,14</sup>

Among the wide variety of BCPs available,<sup>6</sup> PS-*b*-PMMA has been the focus of intense research on nanotemplate applications. As the precise synthesis of PS-*b*-PMMA is achievable by living anionic polymerization, narrowly dispersed products with desirable molecular weights and compositions are made available on a commercial scale. This is extremely important to realize nanofabrications with high precision and reproducibility.<sup>15</sup> Moreover, the reasonable etching contrast between the PS and PMMA domains makes PS-*b*-PMMA more attractive and useful for constructing nanostructures as templates and scaffolds.<sup>16</sup> However, the major limitation of PS-*b*-PMMA is the difficulty in accessing microphase-separated structures with a periodicity of less than 20 nm, which limits its application in nanofabrications at the sub-10 nm resolution.<sup>17</sup> Such limitation stems from the relatively low Flory–Huggins interaction ( $\chi$ ) parameter of PS-*b*-PMMA.<sup>18</sup> Since the product of  $\chi$  and  $N$

<sup>a</sup>Faculty of Engineering, Hokkaido University, Sapporo 060-8628, Japan.  
E-mail: isono.t@eng.hokudai.ac.jp, satoh@eng.hokudai.ac.jp

<sup>b</sup>Graduate School of Chemical Sciences and Engineering, Hokkaido University, Sapporo 060-8628, Japan

<sup>c</sup>Lithography Process Materials Research Group, R&D Department, Tokyo Ohka Kogyo Co., Ltd, Kanagawa 253-0114, Japan

† Electronic supplementary information (ESI) available: Experimental details and additional data and discussions. See DOI: <https://doi.org/10.1039/d3py00113j>

‡ These two authors contributed equally to this work.



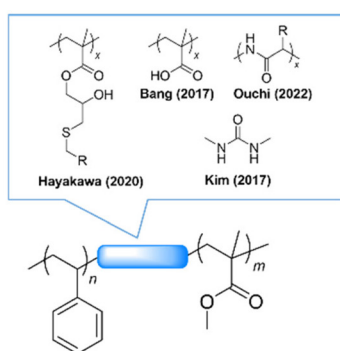
must be greater than 10.5 for microphase separation to occur in symmetrical BCPs, low- $\chi$  BCPs are required to have relatively high molecular weights to satisfy the " $\chi N > 10.5$ " requirement. Meanwhile, the periodicity of the microphase-separated structure, that is, the domain spacing ( $d$ ), is proportional to  $N^{2/3}$ , so the lower limit of  $d$  is determined by the  $\chi$  value of the BCP. In the case of PS-*b*-PMMA, the lowest theoretical molecular weight required for microphase separation is 28 000 g mol<sup>-1</sup>, resulting in the smallest accessible  $d$  of approximately 20 nm.<sup>18</sup>

To expand the accessible  $d$  and control morphology while maintaining the advantages of PS-*b*-PMMA, minor chemical modification of PS-*b*-PMMA should be an effective strategy. For example, the incorporation of short polar blocks, such as polymethacrylate derivatives and oligopeptides, between the PS and PMMA blocks was found to be effective for producing ordered microphase-separated structures with a  $d$  smaller than the smallest accessible one of unmodified PS-*b*-PMMA (Fig. 1a).<sup>19–21</sup> Moreover, Kim reported a reduction in the line edge roughness of the PS-*b*-PMMA nanopattern by incorporating a hydrogen bond-forming urea functionality into the junction point (Fig. 1a).<sup>22</sup> However, such modified PS-*b*-PMMA require multistep synthesis or a custom-made PS-*b*-PMMA possessing a reactive functional group. Meanwhile, our group reported the direct post-polymerization modification of commercially available (anionically synthesized) PS-*b*-PMMA to install polar functional groups at either the PMMA side chain or at the chain end (Fig. 1b).<sup>23–25</sup> We employed the ester-amide exchange reaction for the side-chain functionalization<sup>23</sup> and the terminal-selective transesterification reaction for the PMMA chain end functionalization.<sup>24,25</sup> In both cases, the

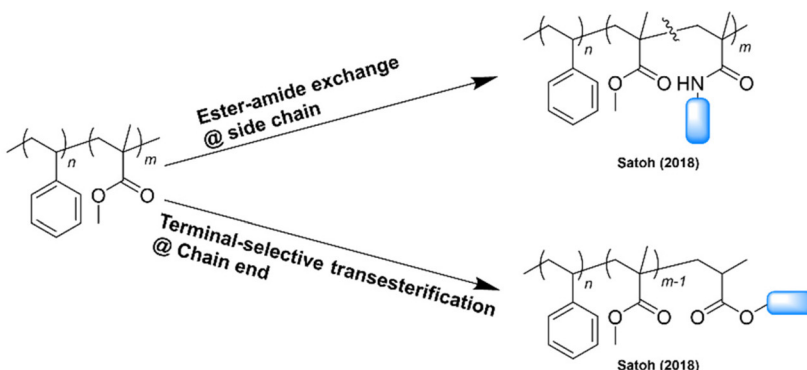
installation of a small proportion of polar functional groups (e.g., amide and oligosaccharide segments) enhanced the polarity of the PMMA block, which drastically promoted segregation of the PS block, resulting in ordered microphase-separated structure formation from low-molecular-weight PS-*b*-PMMA. The biggest advantage of the post-polymerization modification strategy is the easy access to a wide variety of PS-*b*-PMMA derivatives, involving different functional groups and a desirable degree of functionalization, in addition to a fixed backbone length and dispersity from a single starting material. This can lead to rapid discovery of suitable materials for nanofabrication applications using emerging material informatics. However, the post-polymerization modifications of pristine PS-*b*-PMMA reported thus far are based on the reactivity of the PMMA ester group, which limits the accessible chemical space of the PS-*b*-PMMA derivatives.

Herein, we focus on the PS block post-polymerization modification of commercially available PS-*b*-PMMA to modulate its microphase-separated morphology and dimensions (Fig. 1c). Although the benzene ring is chemically stable and much less reactive than other functional groups, such as ester groups, PS benzene ring modifications have been reported and widely implemented on a commercial scale. For example, sulfonation and chloromethylation of crosslinked polystyrene are well-known important steps for producing ion-exchange resins.<sup>26</sup> In addition, many other aromatic ring transformation reactions, such as hydrogenation (dearomatization),<sup>27</sup> halogenation,<sup>28</sup> nitration,<sup>29</sup> borylation<sup>30</sup> and acylation reactions,<sup>31,32</sup> have been applied to modify the PS benzene ring. Among them, the Friedel-Crafts alkylation reaction is a facile tool to

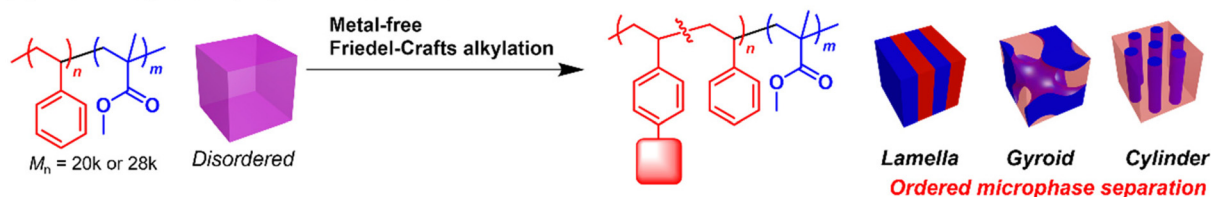
### (a) Junction point functionalization



### (b) Post-polymerization modification of PMMA block



### (c) This study: Post-polymerization modification of PS block



**Fig. 1** Chemical modification of PS-*b*-PMMA. (a) Previous works on the junction point functionalization. (b) Previous works on the PMMA side-chain and chain-end functionalization via post-polymerization modification of commercially available PS-*b*-PMMA. (c) This work, which focuses on post-polymerization modification of the PS block via the metal-free Friedel-Crafts alkylation reaction.



install hydrophobic functional groups on the PS block, which endows the modified PS-*b*-PMMA with a higher polarity difference between the blocks, thus enhancing the  $\chi$  value. Indeed, poly(4-*tert*-butylstyrene)-*b*-PMMA, which has an extra *tert*-butyl group on each PS benzene ring compared to conventional PS-*b*-PMMA, shows a higher  $\chi$  value, thus allowing access to a smaller  $d$  value.<sup>33</sup> Notably, metal-free Friedel–Crafts alkylation is readily realized by combining a 3° alcohol substrate with a strong Brønsted acid catalyst.<sup>34,35</sup> Such metal-free reactions are highly advantageous for reducing metal contamination, which is a prerequisite for semiconductor lithographic applications.

In this study, we investigated the installation of an adamantyl group on the PS block of low-molecular-weight PS-*b*-PMMA *via* metal-free Friedel–Crafts alkylation to control its morphology and dimensions. Interestingly, we found that the installation of an adamantyl group on the PS block resulted in higher thermal stability and higher etching resistance of the modified PS block. The adamantyl group also served as an additional hydrophobic functional group, which increased the  $\chi$  value of the modified PS-*b*-PMMA, thus allowing the formation of ordered microphase-separated structures with  $d$  values as low as 16 nm. Moreover, the volume fraction change through the adamantyl group installation led to a shift in the microphase-separated morphology from lamellar to gyroid and finally to a hexagonally close-packed cylinder structure. These findings contribute to the discovery of nanotemplate materials suitable for various nanofabrication applications.

## Experimental

The general procedure for the Friedel–Crafts alkylation reaction is as follows: in an Ar-filled glovebox, PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub> (300 mg, 1.47 mmol of styrene units, 1.00 eq.) and an adamantanol (0.735 mmol, 0.50 eq.) were placed in a Schlenk flask and dissolved in CH<sub>2</sub>Cl<sub>2</sub> (10 mL). After sealing with a rubber septum, the Schlenk flask was taken out from the glovebox. Triflic acid solution (1.10 mmol, 0.75 eq. in 10 mL CH<sub>2</sub>Cl<sub>2</sub>) was added into the polymer solution to start the reaction. The reaction mixture was stirred at room temperature for 1 h, after which the solution was poured into methanol to precipitate the polymer. The resulting precipitate was filtered and dried

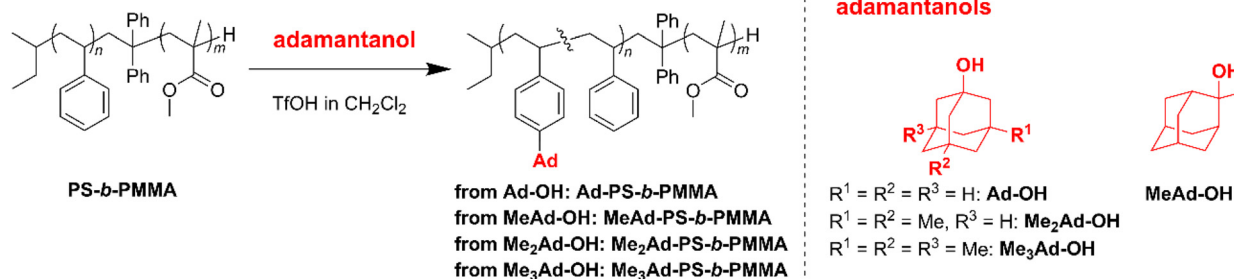
under vacuum to give the adamantly-modified PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub> as a white powder. See the ESI† for more details.

## Results and discussion

### Post-polymerization modification of the polystyrene block *via* the Friedel–Crafts alkylation reaction with adamantanols

The preferred adamantane source was 1-adamantanol (Ad-OH) (Scheme 1). Thus, the metal-free Friedel–Crafts alkylation reaction was carried out by treating PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub> with Ad-OH in the presence of triflic acid (TfOH) in CH<sub>2</sub>Cl<sub>2</sub> with a [styrene unit]<sub>0</sub>/[Ad-OH]<sub>0</sub>/[TfOH] ratio of 1.00/0.50/0.75 (run 1 in Table 1). The resulting product (Ad-PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub>) exhibited new signals at 1.77, 1.87 and 2.08 ppm in its <sup>1</sup>H NMR spectrum due to the adamantyl group, supporting the progress of the reaction (Fig. S1b†). Based on <sup>1</sup>H NMR analysis, the degree of side-chain functionalization ( $f$ : the average number of adamantyl groups attached per styrene unit) was determined to be 0.44. However, size exclusion chromatography (SEC) analysis revealed the presence of many high-molecular-weight by-products (line (b) in Fig. 2). This can be explained by crosslink formation through the adamantane moiety. Since multiple substitution reactions have been reported during the Lewis acid-catalyzed arylation of 1-bromoadamantane with benzene,<sup>36–38</sup> it is reasonably expected that crosslinking could occur in the same manner. Although the mechanism of the multiple substitution reaction is not mentioned in the literature, the reaction most likely proceeds *via* pathways involving carbocation formation from the 3° carbons of the adamantyl group.

Using the preliminary result, we employed commercially available 2-methyl-2-adamantanol (MeAd-OH) and 3,5-dimethyl-1-adamantanol (Me<sub>2</sub>Ad-OH) as the adamantane sources to reduce the possibility of the carbocation migration. Friedel–Crafts alkylation reactions were performed using these two adamantanol derivatives under the same conditions established above (runs 2 and 3 in Table 1). The SEC trace of the product obtained with MeAd-OH (MeAd-PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub>) exhibited high molecular weight elution peaks (line (c) in Fig. 2), indicating intermolecular crosslinking. In contrast, the product obtained with Me<sub>2</sub>Ad-OH indicated a suppressed crosslinking reaction, as evidenced by the clean SEC trend, even though a small high-molecular-weight elution peak was



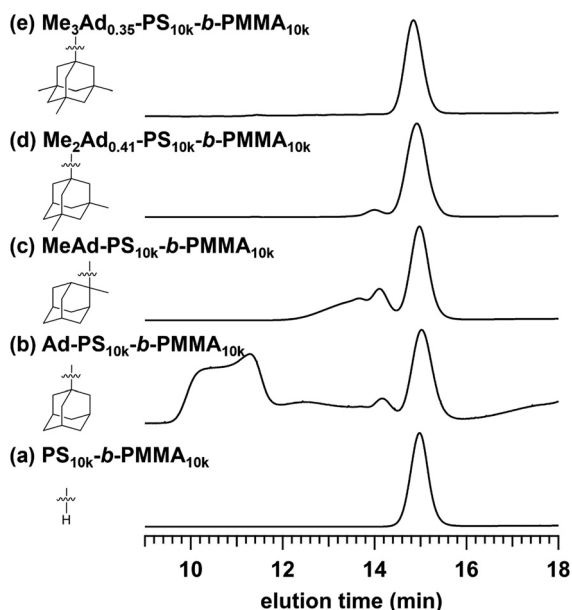
**Scheme 1** Metal-free Friedel–Crafts alkylation of PS-*b*-PMMA using adamantanols.



**Table 1** Friedel–Crafts alkylation reactions of PS-*b*-PMMA with various adamantanols<sup>a</sup>

Run	Sample name	[Styrene] <sup>b</sup> <sub>0</sub> /[adamantanol] <sub>0</sub> /[TfOH]	<i>f</i> <sup>c</sup>	<i>M</i> <sub>n</sub> <sup>d</sup>	<i>D</i> <sup>d</sup>	Yield (%)	$\phi_{\text{PS}}$ <sup>e</sup>
1	Ad-PS <sub>10k</sub> - <i>b</i> -PMMA <sub>10k</sub>	1.00/0.50/0.75	—	—	—	92.5	—
2	MeAd-PS <sub>10k</sub> - <i>b</i> -PMMA <sub>10k</sub>	1.00/0.50/0.75	—	26 400	1.44	69.6	—
3	Me <sub>2</sub> Ad <sub>0.41</sub> -PS <sub>10k</sub> - <i>b</i> -PMMA <sub>10k</sub>	1.00/0.50/0.75	0.41	21 400	1.06	90.1	0.65
4	Me <sub>3</sub> Ad <sub>0.35</sub> -PS <sub>10k</sub> - <i>b</i> -PMMA <sub>10k</sub>	1.00/0.50/0.75	0.35	22 200	1.02	97.7	0.64
5	Me <sub>2</sub> Ad <sub>0.07</sub> -PS <sub>10k</sub> - <i>b</i> -PMMA <sub>10k</sub>	1.00/0.10/0.15	0.07	19 300	1.02	92.1	0.55
6	Me <sub>2</sub> Ad <sub>0.20</sub> -PS <sub>10k</sub> - <i>b</i> -PMMA <sub>10k</sub>	1.00/0.20/0.30	0.20	20 300	1.03	84.9	0.60
7	Me <sub>2</sub> Ad <sub>0.29</sub> -PS <sub>10k</sub> - <i>b</i> -PMMA <sub>10k</sub>	1.00/0.30/0.45	0.29	21 600	1.07	87.1	0.62
8	Me <sub>2</sub> Ad <sub>0.38</sub> -PS <sub>10k</sub> - <i>b</i> -PMMA <sub>10k</sub>	1.00/0.40/0.60	0.38	24 200	1.35	83.1	0.64
9	Me <sub>3</sub> Ad <sub>0.63</sub> -PS <sub>10k</sub> - <i>b</i> -PMMA <sub>10k</sub>	1.00/0.60/0.90	0.63	24 000	1.06	96.5	0.70
10	Me <sub>3</sub> Ad <sub>0.86</sub> -PS <sub>10k</sub> - <i>b</i> -PMMA <sub>10k</sub>	1.00/0.80/1.20	0.86	25 400	1.06	84.6	0.73
11	Me <sub>2</sub> Ad <sub>0.08</sub> -PS <sub>14k</sub> - <i>b</i> -PMMA <sub>14k</sub>	1.00/0.10/0.15	0.08	26 600	1.02	86.6	0.56
12	Me <sub>2</sub> Ad <sub>0.14</sub> -PS <sub>14k</sub> - <i>b</i> -PMMA <sub>14k</sub>	1.00/0.20/0.30	0.14	28 000	1.03	89.5	0.58
13	Me <sub>2</sub> Ad <sub>0.28</sub> -PS <sub>14k</sub> - <i>b</i> -PMMA <sub>14k</sub>	1.00/0.30/0.45	0.28	30 300	1.09	92.4	0.62
14	Me <sub>2</sub> Ad <sub>0.09</sub> -PS <sub>4.2k</sub> - <i>b</i> -PMMA <sub>4.3k</sub>	1.00/0.10/0.15	0.09	9300	1.03	85.1	0.56
15	Me <sub>2</sub> Ad <sub>0.41</sub> -PS <sub>4.2k</sub> - <i>b</i> -PMMA <sub>4.3k</sub>	1.00/0.30/0.45	0.41	10 200	1.05	88.1	0.64
16	Me <sub>2</sub> Ad <sub>0.59</sub> -PS <sub>4.2k</sub> - <i>b</i> -PMMA <sub>4.3k</sub>	1.00/0.60/0.90	0.59	13 400	1.36	86.5	0.68

<sup>a</sup> Reaction conditions: Ar atmosphere; solvent, dichloromethane; temperature, r.t.; and reaction time, 1 h. <sup>b</sup> Styrene units in PS-*b*-PMMA. <sup>c</sup> Degree of substitution (*f*) was determined by quantitative <sup>13</sup>C NMR analysis of Me<sub>2</sub>Ad-PS-*b*-PMMA and by <sup>1</sup>H NMR analysis of Me<sub>3</sub>Ad-PS-*b*-PMMA. <sup>d</sup> Determined by SEC in THF, using PS standards. <sup>e</sup> Volume fraction of the modified PS block.



**Fig. 2** SEC traces of (a) PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub> and PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub> modified with (b) Ad-OH, (c) MeAd-OH, (d) Me<sub>2</sub>Ad-OH, and (e) Me<sub>3</sub>Ad-OH ([styrene unit]<sub>0</sub>/[adamantanol]<sub>0</sub>/[TfOH] = 1.00/0.50/0.75).

still visible (line (d) in Fig. 2). Nevertheless, the <sup>1</sup>H NMR spectrum showed new signals corresponding to the dimethyl-adamantyl group (Fig. S1c†), supporting the success of the Friedel–Crafts alkylation reaction. A logical outcome from the above results is that the best source of adamantane is 3,5,7-trimethyl-1-adamantanol (Me<sub>3</sub>Ad-OH), in which all 3° carbons are blocked by methyl groups. Indeed, Friedel–Crafts alkylation reactions using Me<sub>3</sub>Ad-OH with a [styrene unit]<sub>0</sub>/[adamantanol]<sub>0</sub>/[TfOH] ratio of 1.00/0.50/0.75 successfully produced side-chain-functionalized PS-*b*-PMMA (Me<sub>3</sub>Ad-PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub>) virtually without the high-molecular-weight elution peak due to intermolecular crosslinking (run 4 in Table 1, line

(e) in Fig. 2 for SEC, and Fig. S3† for <sup>1</sup>H NMR spectra). However, the limited commercial availability and the high price of Me<sub>3</sub>Ad-OH led us to use Me<sub>2</sub>Ad-OH for subsequent experiments.

Because the <sup>1</sup>H NMR signals of the Me<sub>2</sub>Ad groups overlapped with those of the polymer main chain, we employed an inverse-gated decoupling <sup>13</sup>C NMR spectrum to calculate the *f* value of Me<sub>2</sub>Ad-PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub> (Fig. 3). By comparing the signal intensities of the aromatic carbons (128–124 ppm) and adamantyl carbons (31.6–30.0 ppm), the *f* value was calculated to be 0.41. Based on the adamantane source, *f* value and the starting PS-*b*-PMMA, this sample was named Me<sub>2</sub>Ad<sub>0.41</sub>-PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub>.

Using the established protocol, we performed the Friedel–Crafts alkylation reactions with PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub> using Me<sub>2</sub>Ad-OH at various [Me<sub>2</sub>Ad-OH]<sub>0</sub>/[styrene unit]<sub>0</sub> ratios, from 0.10 to 0.50, to control the *f* value (runs 5–8 in Table 1). Note that a [TfOH]/[Me<sub>2</sub>Ad-OH]<sub>0</sub> ratio of 1.5 was applied for all of the reactions. Importantly, *f* increased linearly from 0.07 to 0.41 with an increasing [Me<sub>2</sub>Ad-OH]<sub>0</sub>/[styrene unit]<sub>0</sub> ratio, suggesting a high reaction yield and easy control of *f* (Fig. 4). The Fourier transform infrared (FT-IR) spectra of the products revealed a new characteristic absorption band at approximately 2900 cm<sup>-1</sup> due to the C–H stretching of the adamantyl side chain, and the peak intensity increased with increasing *f* (Fig. S2†). The SEC traces of the resulting products showed a sharp elution peak with a dispersity (*D*) typically well below 1.1 (Fig. 5). However, the increase in the [Ad-OH]<sub>0</sub>/[styrene unit]<sub>0</sub> ratio (typically >0.4) resulted in high-molecular-weight by-product formation. Friedel–Crafts alkylation reactions were also successfully applied to PS-*b*-PMMA with different molecular weights, that is, PS<sub>14k</sub>-*b*-PMMA<sub>14k</sub> and PS<sub>4.2k</sub>-*b*-PMMA<sub>4.3k</sub> (runs 11–16 in Table 1).

In addition, we attempted to incorporate an adamantyl side chain with *f* = 0.5, or higher, using Me<sub>3</sub>Ad-OH. Owing to the





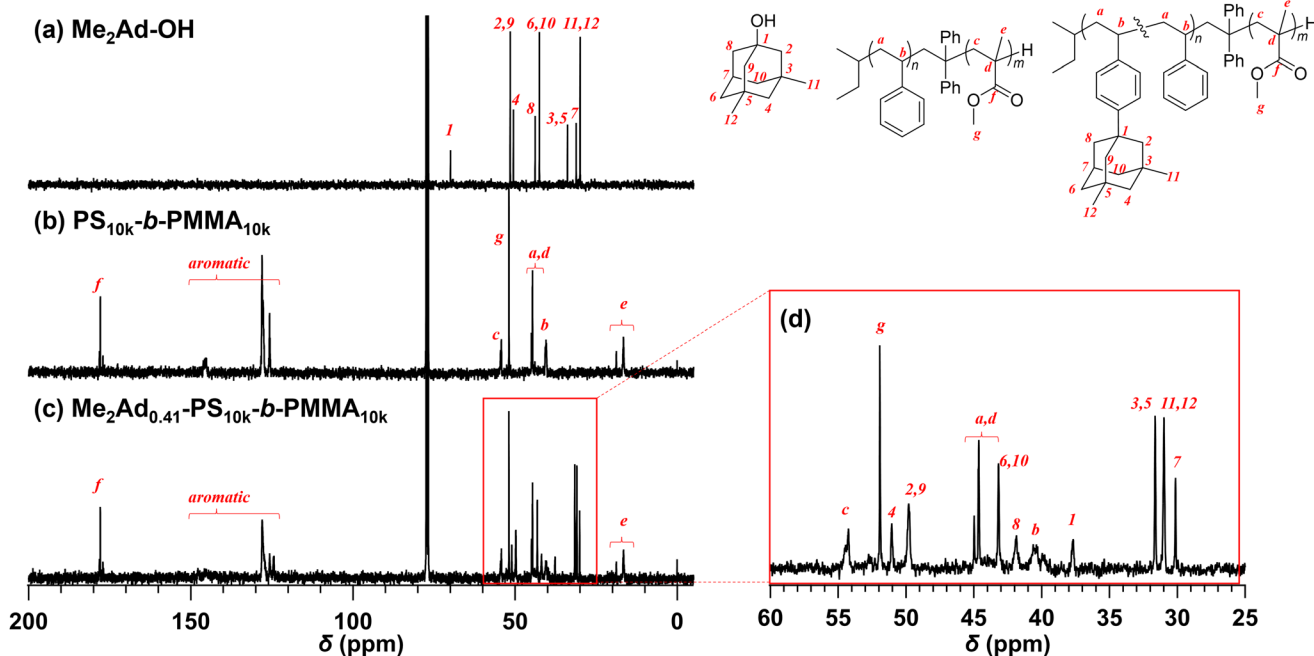


Fig. 3  $^{13}\text{C}$  NMR spectra of (a)  $\text{Me}_2\text{Ad-OH}$  and (b)  $\text{PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$  in  $\text{CDCl}_3$  (100 MHz), and (c) inverse-gated decoupling  $^{13}\text{C}$  NMR spectrum of  $\text{Me}_2\text{Ad}_{0.41}\text{-PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$  (lower) in  $\text{CDCl}_3$  (100 MHz). (d) The expansion from 60 to 25 ppm of the spectrum (c).

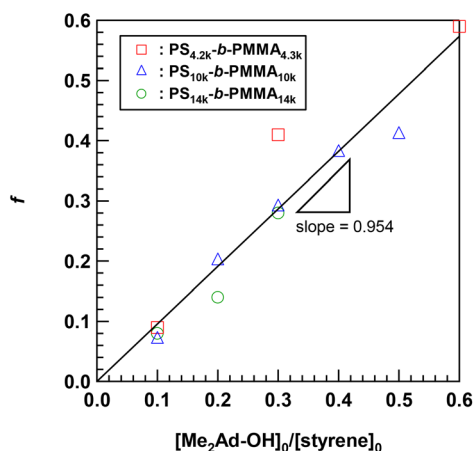


Fig. 4 Linear dependence of the  $f$  value on the initial ratio of  $\text{Me}_2\text{Ad-OH}$  to the styrene unit.

absence of free  $3^\circ$  carbons, Friedel-Crafts alkylation reactions on  $\text{PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$  with  $[\text{Me}_3\text{Ad-OH}]_0/[\text{styrene unit}]_0$  ratios of 0.60 and 0.80 successfully produced the desired products with  $f$  values of 0.63 and 0.86, respectively (runs 9 and 10 in Table 1). Notably, gelation was not observed in these reactions. This was in contrast to the reaction using  $\text{Me}_2\text{Ad-OH}$  at the same  $[\text{Me}_2\text{Ad-OH}]_0/[\text{styrene unit}]_0$  ratio, which produced a partially insoluble product (data not shown). However, even with  $\text{Me}_3\text{Ad-OH}$ , a small measure of the high-molecular-weight elution peak became pronounced with an increasing  $[\text{Me}_3\text{Ad-OH}]_0/[\text{styrene unit}]_0$  ratio (Fig. S3 $^\dagger$ ).

Overall, we successfully demonstrated an easy post-polymerization modification of the polystyrene block in  $\text{PS}\text{-}b\text{-PMMA}$ s

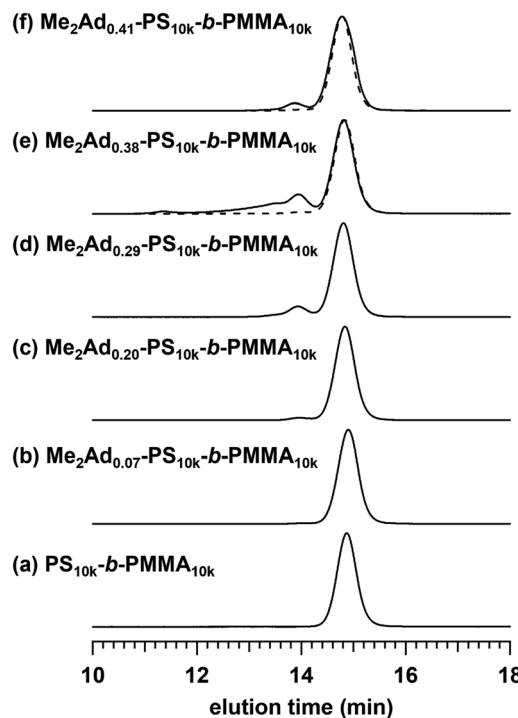


Fig. 5 SEC traces of  $\text{Me}_2\text{Ad-PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$  samples (b)-(f) and the corresponding  $\text{PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$  (a) (eluent, THF; flow rate,  $1.0\text{ mL min}^{-1}$ ). The dashed curves represent the SEC traces after preparative SEC purification.

via Friedel-Crafts alkylation. The synthesis results are summarized in Table 1. The volume fraction of the modified PS block ( $\phi_{\text{PS}}$ ) was calculated based on the density values of



PMMA ( $1.18 \text{ g cm}^{-3}$ ) and PS ( $1.05 \text{ g cm}^{-3}$ ). Here, the density of the modified PS block was assumed to be the same as that of pristine PS because a previous study confirmed that there is no significant difference in density between PS and poly[4-(1-adamantyl)styrene] homopolymers.<sup>39</sup>

### Thermal properties

To investigate the impact of the adamantyl side-chain modification, we assessed the thermal properties of a series of  $\text{Me}_2\text{Ad-PS}_{10k}\text{-}b\text{-PMMA}_{10k}$  copolymers with varied  $f$  values by thermogravimetric analysis (TGA) and differential scanning calorimetry (DSC). TGA revealed a slight increase in 5% and 10% weight loss temperatures ( $T_{d5}$  and  $T_{d10}$ , respectively) with an increasing  $f$  value ( $T_{d5}$ :  $348 \text{ }^\circ\text{C}$  for  $f = 0$  and  $355 \text{ }^\circ\text{C}$  for  $f = 0.41$ ;  $T_{d10}$ :  $356 \text{ }^\circ\text{C}$  for  $f = 0$  and  $364 \text{ }^\circ\text{C}$  for  $f = 0.41$ ) (Fig. S4†). This confirmed that the adamantyl side chain rendered PS- $b$ -PMMA more thermally stable. Ishizone reported a slightly increased  $T_{d10}$  in poly[4-(1-adamantyl)styrene] compared to that in pristine polystyrene,<sup>40</sup> which is consistent with our results. Through DSC analysis, we observed an increase in the glass transition temperature ( $T_g$ ) with increasing  $f$  (Fig. 6). A baseline shift was observed at  $108 \text{ }^\circ\text{C}$  in the second heating DSC curve of the parent  $\text{PS}_{10k}\text{-}b\text{-PMMA}_{10k}$ , which was assignable to the composite of  $T_g$  values of the PMMA and PS blocks. When the  $f$  value increased from 0.07 to 0.29, the composite  $T_g$

increased from  $116$  to  $133 \text{ }^\circ\text{C}$ . A further increase in the  $f$  value resulted in two separate  $T_g$  values, in which the lower  $T_g$  corresponding to the PMMA block was observed at  $132\text{--}133 \text{ }^\circ\text{C}$ . Meanwhile, the higher  $T_g$  corresponding to the modified PS block increased from  $145$  to  $167 \text{ }^\circ\text{C}$  with increasing  $f$  value. These results also demonstrate the increased thermal stability of the adamantyl-modified PS blocks. Such increased thermal stability is preferable for various nanotemplate applications.

### Etching resistance of adamantyl-modified PS

Since the adamantyl group is widely known as an etching-resistant functional group,<sup>40,41</sup> we predicted that the installation of the adamantyl group on the PS block could improve the etching resistance. To test this hypothesis, we prepared a series of  $\text{PS}_{10k}$  samples modified with  $\text{Me}_2\text{Ad-OH}$ , that is,  $\text{Me}_2\text{Ad}_{0.18}\text{-PS}_{10k}$  and  $\text{Me}_2\text{Ad}_{0.62}\text{-PS}_{10k}$ , in the same manner as the PS- $b$ -PMMA modification (Fig. S5 and S6†). Thin films of the modified PS, PS and PMMA homopolymers were prepared on a silicon substrate by spin coating, which was then subjected to  $\text{O}_2$  reactive ion etching for 15, 30, 60 and 120 s. The etching depth for all materials linearly increased with the etching time (Fig. 7). Importantly, the etching rate decreased with increasing adamantyl content. Notably, the etching resistance of  $\text{Me}_2\text{Ad}_{0.62}\text{-PS}$  was 7.7 and 2.0 times compared to PMMA and PS, respectively. These results confirm that the adamantyl group effectively enhanced the etching resistance of the PS block. This is another merit of installing an adamantyl group onto PS- $b$ -PMMA for nanofabrication.

### Microphase-separated structures

Using a series of  $\text{Me}_2\text{Ad}$ -functionalized PS- $b$ -PMMA, we investigated their microphase-separated structures in the bulk material by small-angle X-ray scattering (SAXS) and transmission electron microscopy (TEM). To study the effects of  $f$  on the microphase separation behavior, the  $\text{Me}_2\text{Ad-PS}_{10k}\text{-}b\text{-}$

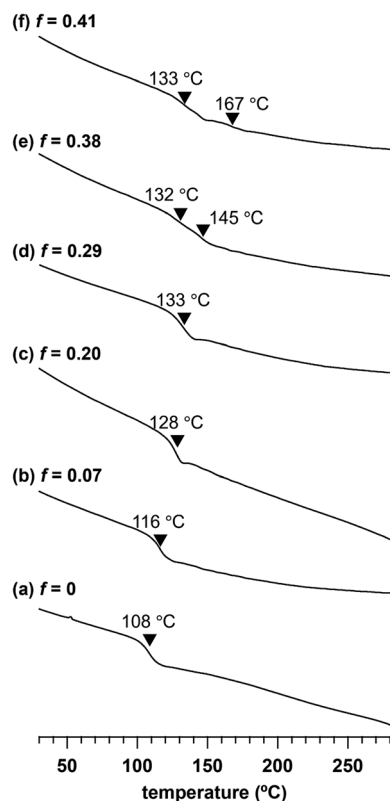


Fig. 6 Second heating DSC traces of (a)  $\text{PS}_{10k}\text{-}b\text{-PMMA}_{10k}$ , (b)  $\text{Me}_2\text{Ad}_{0.07}\text{-PS}_{10k}\text{-}b\text{-PMMA}_{10k}$ , (c)  $\text{Me}_2\text{Ad}_{0.20}\text{-PS}_{10k}\text{-}b\text{-PMMA}_{10k}$ , (d)  $\text{Me}_2\text{Ad}_{0.29}\text{-PS}_{10k}\text{-}b\text{-PMMA}_{10k}$ , (e)  $\text{Me}_2\text{Ad}_{0.38}\text{-PS}_{10k}\text{-}b\text{-PMMA}_{10k}$  and (f)  $\text{Me}_2\text{Ad}_{0.41}\text{-PS}_{10k}\text{-}b\text{-PMMA}_{10k}$ .

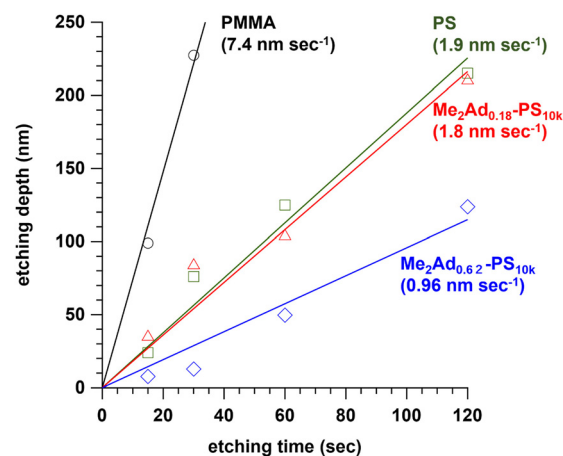
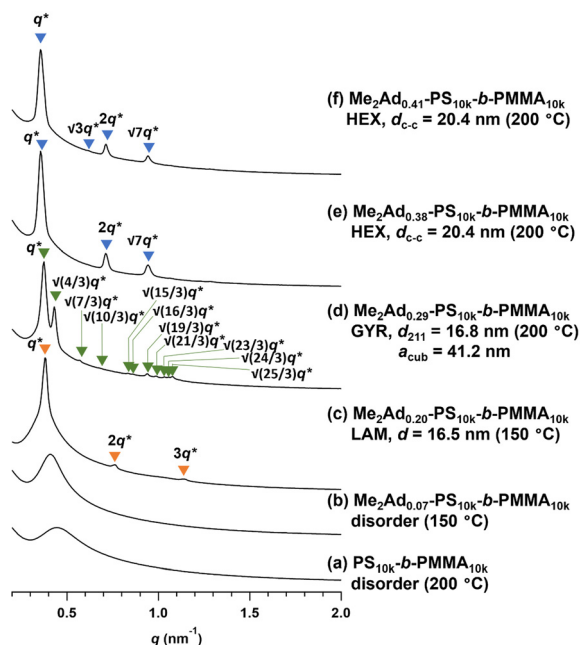


Fig. 7 Plots of etching depth of PMMA, PS,  $\text{Me}_2\text{Ad}_{0.18}\text{-PS}$  and  $\text{Me}_2\text{Ad}_{0.62}\text{-PS}$  thin films with increasing etching time. The values in parentheses represent the etching rate determined by the slope.





**Fig. 8** SAXS profiles of thermally annealed  $\text{Me}_2\text{Ad-PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$ . (a)  $\text{PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$ , (b)  $\text{Me}_2\text{Ad}_{0.07}\text{-PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$ , (c)  $\text{Me}_2\text{Ad}_{0.20}\text{-PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$ , (d)  $\text{Me}_2\text{Ad}_{0.29}\text{-PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$ , (e)  $\text{Me}_2\text{Ad}_{0.38}\text{-PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$  and (f)  $\text{Me}_2\text{Ad}_{0.41}\text{-PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$ . The value in the parentheses represents the annealing temperature.

$\text{PMMA}_{10\text{k}}$  samples were thermally annealed at 200 °C for 24 h and then subjected to SAXS measurements at room temperature. As we previously reported,<sup>23</sup> unmodified  $\text{PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$  did not show any ordered microphase-separated structures, as evidenced by the correlation of hole scattering in its SAXS profile, because  $\chi N$  was lower than the critical value owing to its low  $N$  content (line (a) in Fig. 8). Similarly, the 200 °C-annealed  $\text{Me}_2\text{Ad}_{0.07}\text{-PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$  and  $\text{Me}_2\text{Ad}_{0.20}\text{-PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$  samples did not show any higher-order scattering in their SAXS profiles, indicating the absence of ordered nanostructure formation. We also examined the SAXS profiles of the BCP annealed at 150 °C for 24 h (lines (b) and (c) in Fig. 8).  $\text{Me}_2\text{Ad}_{0.07}\text{-PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$  exhibited only a broad peak, confirming the absence of an ordered structure. In contrast, the  $\text{Me}_2\text{Ad}_{0.20}\text{-PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$  sample exhibited a sharp primary scattering peak ( $q^*$ ) at  $0.380\text{ nm}^{-1}$  together with higher ordered scattering peaks at the  $2q^*$  and  $3q^*$  positions. This scattering pattern is assignable to the lamellar morphology, and the domain spacing ( $d$ ) was calculated to be 16.5 nm using Bragg's equation ( $d = 2\pi/q^*$ ). This result successfully demonstrated that  $\chi N$  crossed the critical value ( $\chi N > 10.5$ ) upon the installation of the adamantyl group with  $f > 0.20$ . More importantly, a  $d$  of 16.5 nm was well below the smallest  $d$  achievable in pristine  $\text{PS-}b\text{-PMMA}$  (approximately 20 nm).<sup>18</sup> Thus, we can conclude that the introduction of the adamantyl group on the PS block is an easy and efficient strategy to realize smaller nanopatterns from low-molecular-weight  $\text{PS-}b\text{-PMMA}$ .

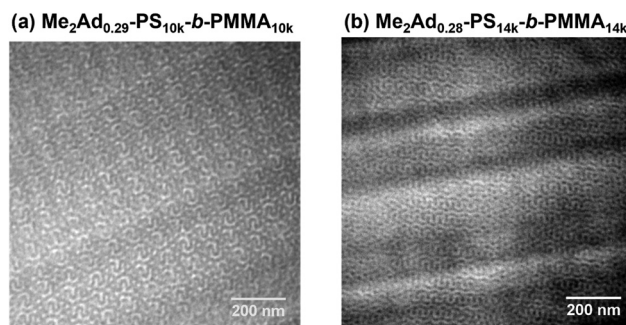
A further increase in the  $f$  value resulted in a shift of the microphase-separated morphology from lamellar (LAM) to

gyroid (GYR) and finally to hexagonally close-packed cylinder (HEX) structures. The 200 °C-annealed  $\text{Me}_2\text{Ad}_{0.29}\text{-PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$  samples exhibited a rather complicated scattering pattern, that is,  $q^*$ ,  $(4/3)^{1/2}q^*$ ,  $(7/3)^{1/2}q^*$ ,  $(8/3)^{1/2}q^*$ ,  $(10/3)^{1/2}q^*$ ..., which can be assigned to the GYR structure (Fig. 8d). The linear relationship in the plot for  $q_{hkl}$  vs.  $(h^2 + k^2 + l^2)^{1/2}$ , where  $h$ ,  $k$  and  $l$  are the Miller indices, validates the successful peak assignment (Fig. S7†). The cubic lattice parameter,  $a_{\text{cub}}$ , was calculated to be 41.2 nm using  $a_{\text{cub}} = 2\pi(h^2 + k^2 + l^2)^{1/2}/q_{hkl}$ . To further confirm the GYR structure, the nanostructures were examined using TEM (Fig. 9a). In the TEM images, double-wave morphologies, which are typical for GYR, were observed over a large area, which again confirmed the long-range order of GYR formation. Interestingly, the GYR structure was maintained for more than one year, as confirmed by SAXS measurements on the same sample left for one year under ambient conditions (Fig. S8†). This clearly demonstrates the highly stable nature of the GYR morphology. To date, ordered GYR structures have rarely been reported for  $\text{PS-}b\text{-PMMA}$  and solvent annealing or homopolymer blending are generally required to induce the GYR structure.<sup>42,43</sup>

For  $\text{Me}_2\text{Ad}_{0.38}\text{-PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$  and  $\text{Me}_2\text{Ad}_{0.41}\text{-PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$ , a sharp primary scattering peak ( $q^*$ ) together with higher-order scattering peaks at  $2q^*$  and  $\sqrt{7}q^*$  were observed in the SAXS profiles (lines (e) and (f) in Fig. 8). In addition, the  $\sqrt{3}q^*$  peak was barely observed. Considering the PS volume fraction, the scattering pattern could be reasonably assigned to the HEX morphology. The inter-cylinder distance ( $d_{\text{c-c}} = (2\pi/q^*)(4/3)^{1/2}$ ) was calculated to be 20.4 nm for both BCPs.

Thus, it was demonstrated that the installation of the adamantyl side chain is an efficient strategy to induce microphase separation even from the low-molecular-weight  $\text{PS-}b\text{-PMMA}$ , which originally did not form an ordered structure. Furthermore, the control of the  $f$  value enabled access to the desired morphologies involving LAM, HEX and even GYR. The shift in the morphology appeared to be the result of both the increase in  $N$  and  $\chi$  upon PS side-chain functionalization.

To gain an insight into the effect of a higher  $f$  on the microphase-separated structures, we performed SAXS measurements



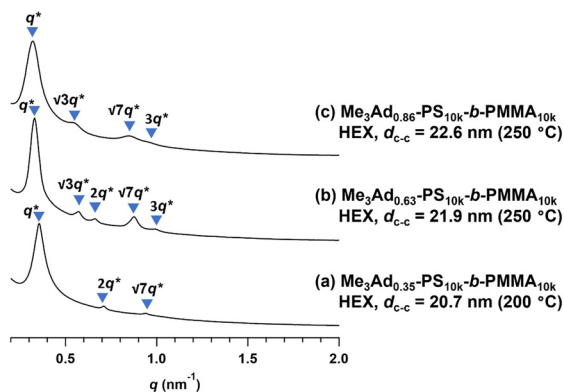
**Fig. 9** TEM micrographs of (a)  $\text{Me}_2\text{Ad}_{0.29}\text{-PS}_{10\text{k}}\text{-}b\text{-PMMA}_{10\text{k}}$  (annealed at 200 °C for 24 h) and (b)  $\text{Me}_2\text{Ad}_{0.28}\text{-PS}_{14\text{k}}\text{-}b\text{-PMMA}_{14\text{k}}$  (annealed at 250 °C for 24 h).



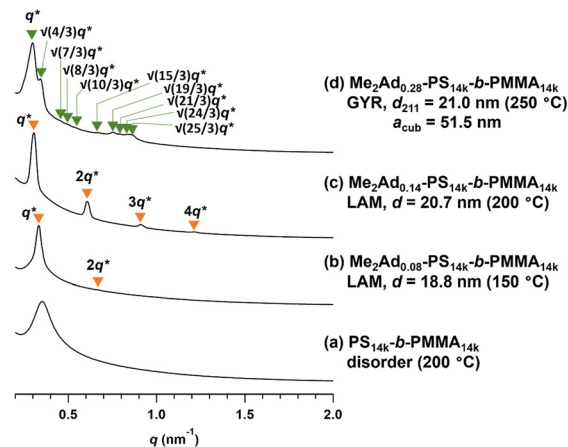


on Me<sub>3</sub>Ad-PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub>. For the 200 °C-annealed Me<sub>3</sub>Ad<sub>0.35</sub>-PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub>, the SAXS profile was assigned to the HEX structure with a  $d_{c-c}$  of 20.7 nm. Notably, the morphology and dimensions of this sample closely resemble those of Me<sub>2</sub>Ad<sub>0.38</sub>-PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub>. This confirmed that  $f$  is the most important parameter for tuning the microphase-separated structure, whereas the difference between the Me<sub>2</sub>Ad and Me<sub>3</sub>Ad groups has no significant impact. Considering the elevated  $T_g$  for the high- $f$  BCPs ( $T_g$  of the modified PS block: 176 and 197 °C for  $f = 0.63$  and  $0.86$ , respectively; see Fig. S9†), the thermal annealing of Me<sub>3</sub>Ad<sub>0.63</sub>-PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub> and Me<sub>3</sub>Ad<sub>0.86</sub>-PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub> was conducted at 250 °C for 24 h. The SAXS profiles of the samples also showed a typical scattering pattern assignable to the HEX morphology (Fig. 10). This result reveals that a further increase in  $f$  did not alter the resulting morphology when starting from the symmetric PS-*b*-PMMA.

SAXS measurements were also performed on the Me<sub>2</sub>Ad-PS<sub>14k</sub>-*b*-PMMA<sub>14k</sub> and Me<sub>2</sub>Ad-PS<sub>4.2k</sub>-*b*-PMMA<sub>4.3k</sub> series to verify the versatility of the proposed strategy. For Me<sub>2</sub>Ad-PS<sub>4.2k</sub>-*b*-PMMA<sub>4.3k</sub>, no nanostructure formation was observed, even after annealing at 150 °C, regardless of the  $f$  value, as indicated by the absence of a higher-order scattering peak (Fig. S10†). This suggests that the increments in  $N$  and  $\chi$  upon side-chain modification are not sufficient to cross the critical value for microphase separation ( $\chi N > 10.5$ ) for this low-molecular-weight PS-*b*-PMMA. A higher  $f$  value is necessary to induce the microphase separation. However, it is more challenging to synthesize such high- $f$  value samples because the intermolecular crosslinking reaction occurs more easily when the [Me<sub>2</sub>Ad-OH]<sub>0</sub>/[styrene]<sub>0</sub> ratio is increased, as mentioned before. In contrast, Me<sub>2</sub>Ad-PS<sub>14k</sub>-*b*-PMMA<sub>14k</sub> exhibited evidence of microphase separation even in the samples with  $f = 0.08$ . This indicates that the slight increase in  $\chi$  by adamantyl side-chain installation is sufficient to surpass the critical  $\chi N$  for this relatively high-molecular-weight PS-*b*-PMMA. The SAXS profiles of 150 °C-annealed Me<sub>2</sub>Ad<sub>0.08</sub>-PS<sub>14k</sub>-*b*-PMMA<sub>14k</sub> and 200 °C-



**Fig. 10** SAXS profiles of the thermally annealed Me<sub>3</sub>Ad-PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub>. (a) Me<sub>3</sub>Ad<sub>0.35</sub>-PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub>, (b) Me<sub>3</sub>Ad<sub>0.63</sub>-PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub> and (c) Me<sub>3</sub>Ad<sub>0.86</sub>-PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub>. The value in the parentheses represents the annealing temperature.



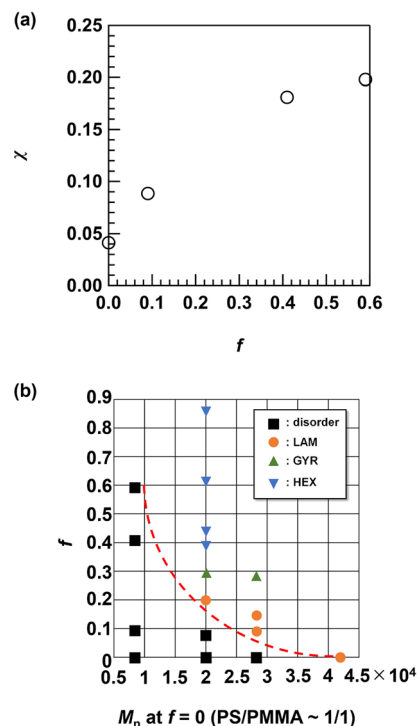
**Fig. 11** SAXS profiles of the thermally annealed Me<sub>2</sub>Ad-PS<sub>14k</sub>-*b*-PMMA<sub>14k</sub>. (a) PS<sub>14k</sub>-*b*-PMMA<sub>14k</sub>, (b) Me<sub>2</sub>Ad<sub>0.08</sub>-PS<sub>14k</sub>-*b*-PMMA<sub>14k</sub>, (c) Me<sub>2</sub>Ad<sub>0.14</sub>-PS<sub>14k</sub>-*b*-PMMA<sub>14k</sub> and (d) Me<sub>2</sub>Ad<sub>0.28</sub>-PS<sub>14k</sub>-*b*-PMMA<sub>14k</sub>. The value in the parentheses represents the annealing temperature.

annealed Me<sub>2</sub>Ad<sub>0.14</sub>-PS<sub>14k</sub>-*b*-PMMA<sub>14k</sub> exhibited scattering patterns associated with LAM structures with a  $d$  value of 18.8 and 20.7 nm, respectively (lines (b) and (c) in Fig. 11). A further increase in the  $f$  value resulted in the formation of the GYR structure, and the SAXS profile of the 250 °C-annealed Me<sub>2</sub>Ad<sub>0.28</sub>-PS<sub>14k</sub>-*b*-PMMA<sub>14k</sub> exhibited the characteristic scattering pattern assignable to the GYR structure (line (d) in Fig. 11). The characteristic morphologies observed in the TEM images also confirm the formation of a highly ordered GYR structure (Fig. 9b). SAXS analysis revealed  $a_{\text{cub}} = 51.5$  nm for Me<sub>2</sub>Ad<sub>0.28</sub>-PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub>. This GYR structure was also stable for more than one year (Fig. S8†).

### Phase behavior

The results described above clearly demonstrate the enhancement of the  $\chi$  value by side-chain modification. To investigate the change in the  $\chi$  value by changing the  $f$  value, we used the SAXS data of the disordered Me<sub>2</sub>Ad-PS<sub>4.2k</sub>-*b*-PMMA<sub>4.3k</sub> with different  $f$  values ( $f = 0.09, 0.41$  and  $0.59$ ) and PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub> ( $f = 0$ ) acquired at 200 °C for fitting with Leibler's mean-field theory.<sup>44</sup> These details are summarized in the ESI.† The result of the fitting for PS<sub>10k</sub>-*b*-PMMA<sub>10k</sub> yielded a  $\chi$  value of 0.032 at 200 °C, which is close to the reported value for PS-*b*-PMMA ( $\chi = 0.0282 + 4.46/T$ , where  $T$  is the absolute temperature;  $\chi = 0.038$  at 200 °C).<sup>18</sup> As shown in Fig. 12a, the  $\chi$  value increased with increasing  $f$  and reached an almost five-fold value ( $\chi \sim 0.20$ ) of that for the original PS-*b*-PMMA when  $f = 0.59$ . This confirmed the versatile control of the  $\chi$  value through optimization of the [adamantanol]<sub>0</sub>/[styrene unit]<sub>0</sub> ratio. The increased hydrophobicity of the polystyrene block stemming from the additional adamantyl group was responsible for the increase in the  $\chi$  value. As a consequence, poly(4-*tert*-butylstyrene)-*b*-PMMA also showed an increased  $\chi$  value (0.043 at 200 °C) over that of PS-*b*-PMMA.<sup>33</sup> Compared to the *tert*-butyl group, the adamantyl group appeared to contribute





**Fig. 12** (a) Dependence of the  $\chi$  value on  $f$  determined from the SAXS curve analysis of disordered Me<sub>2</sub>Ad-PS<sub>4.2k</sub>-*b*-PMMA<sub>4.3k</sub> at 200 °C. (b) Summary of the microphase-separated morphologies of the adamantyl-modified PS-*b*-PMMA.

more efficiently to the increased  $\chi$  value due to its higher carbon number.

The shift in the morphology from LAM to GYR and finally to HEX is reminiscent of the conventional BCP phase diagram.<sup>2</sup> As shown in Table 1, an increase in  $f$  results in an increase in  $\phi_{\text{PS}}$ . Thus, it is reasonable to deduce that the morphology shift is a direct result of increased  $\phi_{\text{PS}}$ . In this study, we did not observe a body-centered cubic (BCC) sphere morphology. To obtain the BCC morphology,  $\phi_{\text{PS}}$  must be higher than 0.8. Given that the original PS-*b*-PMMA has a symmetric volume fraction,  $\phi_{\text{PS}} > 0.8$ , this cannot be achieved even with  $f = 1.0$ . Nevertheless, we have demonstrated easy access to nanotechnologically interesting GYR and lithographically relevant LAM and HEX morphologies from a single PS-*b*-PMMA starting material. These results support the facile access to various nanotemplate structures by a judicious choice of the  $f$  value.

## Conclusions

We have successfully demonstrated PS block post-polymerization modification of low-molecular-weight PS-*b*-PMMA *via* a Friedel–Crafts alkylation reaction to induce microphase separation. In this study, an adamantane moiety was installed on the benzene ring of PS, and the degree of functionalization was easily controlled. One of the important features of this post-polymerization modification is that the reaction proceeds

without a metal catalyst, which is advantageous for microelectronic applications. The incorporation of the adamantyl group enhanced the incompatibility with the PMMA block, resulting in microphase separation into lamellar, gyroid and hexagonally close-packed cylindrical morphologies, even in the low-molecular-weight region. Notably, using this strategy, we achieved a domain spacing of approximately 16 nm, which was considerably lower than the lower limit of pristine PS-*b*-PMMA (~20 nm). Interestingly, the adamantyl group increased the thermal stability, glass transition temperature and etching resistance of the modified PS block, which are all preferable for nanofabrication applications. Because the presented post-polymerization modification reaction proceeds under mild reaction conditions, this strategy should be applied to a wide variety of PS-containing BCPs as well as tertiary alcohol substrates. Moreover, by combining this approach with the previously established direct PMMA post-polymerization modifications, the accessible chemical space of PS-*b*-PMMA derivatives can be vastly expanded, creating a large BCP library with a fixed total degree of polymerization and dispersity from a single PS-*b*-PMMA starting material. With the aid of an emerging materials informatics approach, a novel BCP material that is best suited for future nanofabrication applications can be obtained.

## Author contributions

Takuya Isono: conceptualization, project administration, investigation, supervision, writing – review and editing, funding acquisition. Ema Baba: investigation, methodology. Shunma Tanaka: investigation, methodology. Ken Miyagi: investigation. Takahiro Dazai: investigation. Feng Li: investigation, supervision. Takuya Yamamoto: investigation, supervision. Kenji Tajima: investigation, supervision. Toshifumi Satoh: writing – review and editing, supervision, project administration, funding acquisition.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This work was financially supported by a JSPS Grant-in-Aid for Scientific Research (B) (No. 20H02792 and 19H02769), the Photoexcitonix Project (Hokkaido University), the project of junior scientist promotion in Hokkaido University, and the Creative Research Institute (Hokkaido University). This work was performed with the approval of the Photon Factory Program Advisory Committee (Proposal No.: 2022G555 and 2021G0531). The authors thank Mr Toshiaki Ito (Hokkaido University, Japan) for his assistance with the TEM experiment.



## References

- 1 I. W. Hamely, *The Physics of Block Copolymers*, Oxford University Press, Oxford, 1988.
- 2 F. S. Bates and G. H. Fredrickson, *Phys. Today*, 1999, **52**, 32–38.
- 3 M. Lazzari, G. Liu and S. Lecommandoux, *Block Copolymers in Nanoscience*, Wiley-VCH Press, Weinheim, 2006.
- 4 H. Hu, M. Gopinadhan and C. O. Osuji, *Soft Matter*, 2014, **10**, 3867–3889.
- 5 H. Jung, W. H. Shin, T. W. Park, Y. J. Choi, Y. J. Yoon, S. H. Park, J. H. Lim, J. D. Kwon, J. W. Lee, S. H. Kwon, G. H. Seong, K. H. Kim and W. I. Park, *Nanoscale*, 2019, **11**, 8433–8441.
- 6 G. G. Yang, H. J. Choi, K. H. Han, J. H. Kim, C. W. Lee, E. I. Jung, H. M. Jin and S. O. Kim, *ACS Appl. Mater. Interfaces*, 2022, **14**, 12011–12037.
- 7 H. Tsai, J. W. Pitera, H. Miyazoe, S. Bangsaruntip, S. U. Engelmann, C.-C. Liu, J. Y. Cheng, J. J. Bucchingnano, D. P. Klaus, E. A. Joseph, D. P. Sanders, M. E. Colburn and M. A. Guillorn, *ACS Nano*, 2014, **5**, 5227–5232.
- 8 R. Ruiz, H. Kang, F. A. Detcheverry, E. Dobisz, D. S. Kercher, T. R. Albrecht, J. J. de Pablo and P. F. Nealey, *Science*, 2008, **321**, 936–939.
- 9 C. Tang, E. M. Lennon, G. H. Fredrickson, E. J. Kramer and C. Hawker, *J. Sci.*, 2008, **322**, 429–432.
- 10 S. Park, Y. Kim, H. Ahn, J. H. Kim, P. J. Yoo and D. Y. Ryu, *Sci. Rep.*, 2016, **6**, 36326.
- 11 E. J. W. Crossland, M. Kamperman, M. Nedelcu, C. Ducati, U. Wiesner, D.-M. Smilgies, G. E. S. Toombes, M. A. Hillmyer, S. Ludwigs, U. Steiner and H. A. Snaith, *Nano Lett.*, 2009, **8**, 2807–2812.
- 12 E. L. Lin, W. L. Hsu and Y. W. Chiang, *ACS Nano*, 2018, **12**, 485–493.
- 13 M. R. J. Scherer and U. Steiner, *Nano Lett.*, 2013, **13**, 3005–3010.
- 14 W.-C. Ma, W.-S. Huang, C.-S. Ku and R.-M. Ho, *J. Mater. Chem. C*, 2016, **4**, 840–849.
- 15 H. Lai, G. Huang, X. Tian, Y. Liu and S. Ji, *Polymer*, 2022, **249**, 124853.
- 16 M. G. Gusmão Cacho, P. Pimenta-Barros, M. Argoud, C. Navarro, K. Sakavuyi, R. Tiron and N. Possémé, *Microelectron. Eng.*, 2020, **230**, 111369.
- 17 L. Wan, R. Ruiz, H. Gao, K. C. Patel, T. R. Albrecht, J. Yin, J. Kim, Y. Cao and G. Lin, *ACS Nano*, 2015, **9**, 7506–7514.
- 18 Y. Zhao, E. Sivaniah and T. Hashimoto, *Macromolecules*, 2008, **41**, 9948–9951.
- 19 S. Woo, S. Jo, D. Y. Ryu, S. H. Choi, Y. Choe, A. Khan, J. Huh and J. Bang, *ACS Macro Lett.*, 2017, **6**, 1386–1391.
- 20 K. Wylie, L. Dong, A. Chandra, Y. Nabee and T. Hayakawa, *Macromolecules*, 2020, **53**, 1293–1301.
- 21 T. Yoshimura, T. Morishita, Y. Agata, K. Nagashima, K. Wylie, Y. Nabee, T. Hayakawa and M. Ouchi, *Macromolecules*, 2022, **55**, 2148–2159.
- 22 K. S. Lee, J. Lee, J. Kwak, H. C. Moon and J. K. Kim, *ACS Appl. Mater. Interfaces*, 2017, **9**, 31245–31251.
- 23 K. Yoshida, L. Tian, K. Miyagi, A. Yamazaki, H. Mamiya, T. Yamamoto, K. Tajima, T. Isono and T. Satoh, *Macromolecules*, 2018, **51**, 8064–8072.
- 24 K. Yoshida, S. Tanaka, T. Yamamoto, K. Tajima, R. Borsali, T. Isono and T. Satoh, *Macromolecules*, 2018, **51**, 8870–8877.
- 25 K. Yoshida, T. Yamamoto, K. Tajima, T. Isono and T. Satoh, *Polym. Chem.*, 2019, **10**, 3390–3398.
- 26 A. Ravve, *Principles of Polymer Chemistry*, Springer, New York, 3rd edn, 2012.
- 27 Z. Siddiqi and D. Sarlah, *J. Am. Chem. Soc.*, 2021, **143**, 21264–21269.
- 28 G. A. Crosby, N. M. Weinshenker and H. S. Uh, *J. Am. Chem. Soc.*, 1975, **97**, 2232–2235.
- 29 A. Philippides, P. M. Budd, C. Price and A. V. Cuncliffe, *Polymer*, 1993, **34**, 3509–3513.
- 30 J. Shin, S. M. Jensen, J. Ju, S. Lee, Z. Xue, S. K. Noh and C. Bae, *Macromolecules*, 2007, **40**, 8600–8608.
- 31 R. L. Letsinger, M. J. Kornet, V. Mahadevan and D. M. Jerina, *J. Am. Chem. Soc.*, 1964, **86**, 5163–5165.
- 32 B. Xue, P. P. Huang, M. Z. Zhu, S. Q. Fu, J. H. Ge, X. Li and P. N. Liu, *ACS Macro Lett.*, 2022, **11**, 1252–1257.
- 33 J. G. Kennemur, M. A. Hillmyer and F. S. Bates, *Macromolecules*, 2012, **45**, 7228–7236.
- 34 J. S. Olsson, T. H. Pham and P. Jannasch, *Macromolecules*, 2020, **53**, 4722–4732.
- 35 J. Y. Jeon, Z. Umstead, G. N. Kangovi, S. Lee and C. Bae, *Top. Catal.*, 2018, **61**, 610–615.
- 36 H. Newman, *Synthesis*, 1972, 692–693.
- 37 V. R. Reichert and L. J. Mathias, *Macromolecules*, 1994, **27**, 7015–7023.
- 38 H. Lim, M. C. Cha and J. Y. Chang, *Polym. Chem.*, 2012, **3**, 686–870.
- 39 B. J. Ree, S. Kobayashi, K. Heo, T. J. Lee, T. Satoh, T. Ishizone and M. Ree, *Polymer*, 2019, **169**, 225–233.
- 40 S. Kobayashi, T. Matsuzawa, S.-i. Matsuoka, H. Tajima and T. Ishizone, *Macromolecules*, 2006, **39**, 5979–5986.
- 41 J. M. Klopp, D. Pasini, J. D. Byers, C. G. Willson and J. M. J. Fréchet, *Chem. Mater.*, 2001, **13**, 4147–4153.
- 42 S. Park, Y. Kim, W. Lee, S.-M. Hur and D. Y. Kim, *Macromolecules*, 2017, **50**, 5033–5041.
- 43 J.-W. Hong, J.-H. Chang, I. C.-Y. Chang and Y.-S. Sun, *Soft Mater.*, 2021, **17**, 9189–9197.
- 44 L. Leibler, *Macromolecules*, 1980, **13**, 1602–1617.

