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Anionic polymerization and polyhomologation: an ideal combination to synthesize polyethylene-based block copolymers[†]

Hefeng Zhang,^a Nazeeha Alkayal,^a Yves Gnanou^b and Nikos Hadjichristidis*^a

A novel one-pot methodology combining anionic polymerization and polyhomologation, through a "bridge" molecule (BF₃OEt₂), was developed for the synthesis of polyethylene (PE)-based block copolymers. The anionically synthesized macroanion reacts with the "bridge" molecule to afford a 3-arm star (trimacromolecular borane) which serves as an initiator for the polyhomologation.

Polyethylene (PE)-based materials are very important to modern life covering a wide spectrum of applications from commodity plastics (e.g. packaging, bottles) to precision-processed biomaterials (e.g. total joint replacement and medical devices).1 In order to better understand the behaviour and improve the performance of these materials, welldefined (controllable molecular weight, narrow molecular weight and structural distribution) PE-based polymers are needed.² Unfortunately, anionic polymerization which gives access to well-defined polymers when chain growth occurs in a "living" fashion is not compatible with ethylene, the monomer of PE.3 The only way to prepare well-defined PEs by anionic polymerization is by hydrogenation of the corresponding anionically prepared PBd-1,4. However, the produced PE possesses butylene units originating from the hydrogenation of the unavoidable 1,2-units (minimum 5%) formed during the chain growth.⁴ Recently, Shea developed a novel polymerization methodology leading to perfectly linear PEs.⁵ The general reaction scheme involves the formation of an organoboron zwitterionic complex between a methylide (monomer) and a trialkylborane Lewis acid (initiator) which breaks down by the intramolecular 1,2-migration. As a consequence, the methylene group of methylide is randomly inserted one by one into the three branches of the trialkylborane leading to a 3-arm PE star. The resulting star is subsequently oxidized/hydrolysed to give perfectly OH-end-capped linear PEs. Molecular weights from 500 to 5.0×10^5 and the polydispersity index (PDI) typically less than 1.1 have

been reported. For this type of polymerization Shea coined the name polyhomologation or C1 polymerization. $^{5\text{--}7}$

By using functionalized ylides and/or organoboranes, functionalized linear and nonlinear homo/copolymers, such as α -hydroxyl- ω -allylic PE, 3-armed α -hydroxyl PE star, PE-*co*-PP(polypropylene) and cyclic PE have been synthesized.⁷ Until now, polyhomologation has been combined with controlled/living radical and ring opening polymerization reactions to afford PE-based block copolymers such as PS-*b*-PE, PEO-*b*-PE diblock and PE-*b*-PDMS-*b*-PE triblock copolymers, *etc.* (PS: polystyrene; PEO: polyethylene oxide; PDMS: polydimethylsiloxane).⁸

Here, we report a novel one-pot methodology combining anionic polymerization and polyhomologation, through a "bridge" molecule (BF₃OEt₂), for the synthesis of polyethylene (PE)-based block copolymers. The synthetic approach involves: (a) the synthesis of a 3-arm star polymer (trimacromolecular borane, macroinitiator) by reacting living macroanions with BF₃OEt₂, (b) the *in situ* polyhomologation of dimethylsulfoxonium methylide, with the macroinitiator to produce a 3-arm star block copolymer of PE, and (c) the oxidation/hydrolysis by trimethylamine N-oxide dihydrate (TAO) to afford the PE-based block copolymers. As examples the syntheses of PBd-*b*-PE (PBd: polybutadiene) and PS-*b*-PE are shown in Scheme 1. The details are given in the ESL[†]

A living macroanion, PBdLi or PSLi, was first synthesized in cyclohexane at room temperature and then reacted (30% excess) with BF_3OEt_2 (0.7 M solution in diethyl ether) to afford a 3-arm boron-linked star. Due to the high sensitivity



Scheme 1 Combination of anionic polymerization and polyhomologation leading to PBd-b-PE and PS-b-PE block copolymers (BF₃OEt₂: "bridge" molecule).

^a King Abdullah University of Science and Technology, Division of Physical Sciences & Engineering, KAUST Catalysis Center, Polymer Synthesis Laboratory, Thuwal, Kingdom of Saudi Arabia

Kingaom of Sauai Arabia

^b King Abdullah University of Science and Technology, Division of Physical Sciences & Engineering, Thuwal, Kingdom of Saudi Arabia.

E-mail: nikolaos.hadjichristidis@kaust.edu.sa

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Fig. 1 The products of coupling of PBd and PS anions with BF_3OEt_2 decompose during SEC measurements (THF at 35 °C).

of boranes to air (oxygen and moisture) it was not possible to analyse the stars formed by size exclusion chromatography (SEC), since the borane species decompose during analysis (Fig. 1). The peaks corresponding to the products of coupling of PBd and PS anions with BF_3OEt_2 almost disappeared after 24 and 18 hours, respectively, and thus the decomposition of these macromolecular boranes (2- and 3-arm stars) was confirmed.

In order to avoid the decomposition, $(PBd)_3B$ and $(PS)_3B$ 3-arm stars were directly used after synthesis *in situ* for the polymerization of dimethylsulfoxonium methylide in toluene at 70 °C and 80 °C, respectively.

The dimethylsulfoxonium methylide (monomer) was prepared in refluxed tetrahydrofuran (THF) at 70 $^{\circ}$ C from trimethylsulfoxonium chloride in the presence of sodium hydride according to Corey's method (yield: 60–80%).⁹ After synthesis, THF was switched to toluene, the polymerization solvent.

The polyhomologation of dimethylsulfoxonium methylide with the star-like macroinitiators, (PBd)₃B and (PS)₃B, resulted in the corresponding 3-arm star block copolymers (PBd-b-PE)3B and (PS-b-PE)3B, which could not be analyzed by SEC due to reasons explained above. Oxidation/hydrolysis of the resulting starblock copolymers with trimethylamine N-oxide dihydrate (TAO) led to the linear w-hydroxyl PBd-b-PE and PS-b-PE. Parallel experiments led to the conclusion that the methylide monomer cannot be polymerized either by the macroanion (PBdLi and PSLi) or by BF3OEt2. This indicates that only the boron-linked 3-arm star polymer has the potential for polyhomologation. The initiating efficiency of the different boron-based 3-arm stars depends on the chemical nature/steric hindrance of the arms. Macroinitiators based on polybutadiene initiate the polyhomologation much faster than PS₃B. For example, polyhomologation leading to DP = 1700 could be completed in less than 10 min at 70 °C with (PBd)₃B as an initiator, whereas one hour was needed when PS3B was used at 80 °C. After polyhomologation and cooling to room temperature, the toluene solution turned cloudy thus indicating successful formation of a PE block (Fig. 2).

The syntheses of PBd-*b*-PE and PS-*b*-PE were monitored by HT(high temperature)-SEC in trichlorobenzene (TCB) at 150 $^{\circ}$ C (Fig. 3).



Fig. 2 PS-b-PE solution in toluene after polyhomologation at 80 °C and 25 °C.

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Fig. 3 Syntheses of PBd-*b*-PE and PS-*b*-PE monitored by HT-SEC using TCB as an eluent at 150 °C. ^a The negative peak was shown with positive style for better comparison (dn/dc of PE and PBd are negative in TCB).

 Table 1
 PE-based block copolymers synthesized by combining anionic polymerization and polyhomologation

	PBd or PS block		PE block		(PBd or PS)-b-PE			
Sample	${M_{ m n} \over (imes 10^3)}$	PDI _{HT-SEC}	$\overline{M_{ m n,theor.}}_{ m (imes 10^3)^c}$	$M_{ m n,NMR} \ (imes 10^3)$	PDI _{HT-SEC}	$T_{\rm m}$ (°C)	T_{g} (°C)	UCST (°C) ^g
PBd-b-PE PS-b-PE	5.9^{a} 6.5^{b}	1.05 1.09	10.5 3.3	24.0^{d} 11.9 ^e	1.14 1.11	101 102		40–55 40–55

^{*a*} The molecular weight was determined by MALDI-ToF (Fig. S1, ESI). ^{*b*} The molecular weight was determined by HT-SEC using TCB as an eluent at 150 °C with a flow rate of 0.8 mL min⁻¹ calibrated by PS standards. ^{*c*} The molecular weight was calculated from the ratio of the ylide to the initiator. ^{*d*} $M_{n,PF,NMR} = 14 \times DP_{PBd} \times (A_{0.8-2.4ppm} - 0.6 \times A_{6.9-7.4ppm}) - 2 \times A_{4.8-5.6ppm}).$ ^{*d*} $M_{n,PF,NMR} = 14 \times DP_{PS} \times (A_{0.8-2.4ppm} - 0.6 \times A_{6.4-7.4ppm})/(2.5 \times A_{6.4-6.3ppm}).$ ^{*f*} The glass transition of the PS block was hidden by the melting process of the PE block. ^{*g*} Upper critical solution temperature (UCST) was measured in toluren (3 mg mL⁻¹) on a UV-Vis instrument with a manual temperature controller.

It is clear that the peak appearing after polyhomologation moves to the higher molecular weight region (lower elution volumes), the lower molecular weight peak corresponding to the excess precursor eliminated after fractionation. For this reason the polyhomologation products were dissolved in toluene at high temperature to give a clear and transparent solution, followed by cooling to room temperature. The solution was then centrifuged in the presence of a small amount of selective solvent for PBd (petroleum ether) and PS (acetone) to facilitate the separation. The absence of the PBd and PS precursors (Fig. 3) proves the success of the fractionation (Table 1).

Successful syntheses of diblock copolymers have been confirmed by Fourier-transformed infrared spectroscopy (FTIR) (Fig. 4). In the case of PBd-*b*-PE, the peaks at 964 and 910 cm⁻¹ characteristic of and attributable to ==C-H bending vibration in PBd blocks were found.

In the DSC curve (Fig. 5) of the PBd-*b*-PE diblock copolymer, a melt temperature of 101 $^{\circ}$ C attributable to the PE block was



Fig. 4 FTIR spectra of PBd-*b*-PE and PS-*b*-PE copolymers, ^aPE standard with a molecular weight of 2000.



Fig. 5 DSC curves of PBd-b-PE and PS-b-PE diblock copolymers.



Fig. 6 Thermal responses of PBd-*b*-PE and PS-*b*-PE copolymers revealed by ¹H NMR in toluene- d_8 (left) at different temperatures and UCST measurements in toluene (3 mg mL⁻¹) (right).

found, as well as the glass transition to the PBd block at *ca.* -85 °C. In the case of PS-*b*-PE, the glass transition of the PS block was covered by the melting process of the PE block.

The PE fingerprint is clearly seen from the thermal behavior of the PE copolymers in toluene monitored by UV-Vis measurements at 450 nm. The graphs (transmittance as a function of temperature) of PBd-*b*-PE and PS-*b*-PE are shown in Fig. 6. A phase transition occurs at 40–55 °C upon heating the PBd-*b*-PE solution (3 mg mL⁻¹): the transmittance then increases from ~40% to 100%. For PS-*b*-PE, the phase transition is much more significant as the transmittance increases from 2% to 100% upon increasing the temperature to 40–55 °C.

The PE fingerprint was also evident in the thermal responses (shielding effect) of diblock copolymers in ¹H NMR spectra (Fig. 6). The single and sharp peak at 1.4 ppm, attributable to the protons in PE backbone, compared to vinyl protons in the 1,4-type double bond in the PBd block at 5.5 ppm, increases with temperature, which means that more and more PE blocks dissolved in the solvent. The area ratio of the peaks at 1.4 ppm to the one at 4.8–5.6 ppm was used to calculate the ratio of the PE block in the copolymers using: $M_{n,PE,NMR} = 14 \times DP_{PEd} \times (A_{0.8-2.4ppm} - 0.6 \times A_{6.9-7.4ppm} - 2 \times A_{4.8-5.6ppm})/A_{4.8-5.6ppm}$. Correspondingly, the peaks of aryl protons due

to the PS blocks at 6.4–6.9 ppm were employed to calculate the respective contents of PS and PE in PS-*b*-PE copolymers. Surprisingly, the molecular weights calculated in this way were higher than expected. For PBd-*b*-PE, the molecular weight of 24.0×10^3 drawn from NMR for the PE block is 2.3 times higher than the expected value of 10.5×10^3 . The deviation is more obvious for PS-*b*-PE (Table 1). The higher molecular weight observed indicates that the macroinitiator concentration was lower than the theoretical one, meaning that the efficiency of the linking reaction (living polymers with BF₃OEt₂) was about 30%. Other reasons for this disagreement in molecular weights should be excluded since the PDIs of the final block copolymers were very low (Table 1).

In conclusion, a novel strategy, based on anionic polymerization combined with polyhomologation through a "bridge" molecule (BF₃OEt₂), has been proposed for the synthesis of PE-based block copolymers. The examples described in this work (PBd-*b*-PE and PS-*b*-PE) are only two out of the many that can be prepared by this general methodology. A few are given below: ABE, (AB)_{2or3}CE, (ABC)_{2or3}DE, ABCDE, (ABCD)_{2or3}FE, *etc.* (A, B, C, D, E, F: different blocks, E: polyethylene). The potential of this method is limited only by our imagination.

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