Thioanhydride/isothiocyanate/epoxide ring-opening terpolymerisation: sequence selective enchainment of monomer mixtures and switchable catalysis†

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We report a new sequence selective terpolymerisation in which three monomers (butylene oxide (BO) A, PhNCS B and phtalic thioanhydride (PTA) C) are selectively enchainged into an (ABA/C)n sequence. PTA/PhNCS/BO ring-opening terpolymerisation (ROTERP) can be coupled with CS₂ ROTERP to generate tetrapolymers and with εDL ROP in switchable catalysis for blockpolymer synthesis.

Heteroatom-containing polymers have strong potential as degradable replacements for polymers based on saturated aliphatic backbones.1-3 Often such polymers are synthesised via the living ring opening polymerisation (ROP) of a heterocycle A to produce (A)n, such as poly(thio)esters, poly(thio)carbonate and poly(thio)ethers.2,4 In some cases, the ROP of three or four-membered heterocycles A can be coupled with the insertion of heteroallenes or cyclic anhydrides B to generate alternating copolymers (AB)n in so called ring-opening copolymerisation (ROCP).5,6 Prominent examples include CO3/epoxide ROCOP forming polycarbonates or cyclic anhydride/epoxide ROCOP forming polyesters.7-9 However, sulfur-containing variants also exist and relevant to this study are recent reports on isothiocyanate/epoxide ROCOP forming poly(Al,Si)thioimidocarbonates),8,9 isothiocyanate/thiirane ROCOP forming poly(dithioimidocarbonates), cyclic thiioanhydride/epoxide ROCOP forming poly(ester-alt-thioesters).10-18 Recently we realised alternating ring-opening terpolymerisation (ROTERP) of ternary monomer mixtures comprising thiophatatic anhydride (PTA), CS₂ and butylene oxide (BO).19 Here, a simple lithium catalyst (e.g. lithiumbenzyloxide LiOBn or lithiumhexamethyldisilazide LiHMDS) selectively forms poly(ester-alt-alt-dithioimidocarbonates) in up to 98% selectivity with respect to the erroneous thioester links from PTA/BO ROCOP. The polymer shows an unusual “head-to-head-alt-tail-to-tail” selectivity, meaning that ester groups sit next to tertiary carbon centres while trithiocarbonates sit next to secondary carbon centres. Our results indicated that PTA/CS₂/BO ROTERP formally derives from Li catalysed CS₂/BO ROCOP reported by Werner and co-workers.17 Both polymerisations are enabled by a O/S exchange reaction step in which an alkoxide chain end from a ring opened epoxide A isomerizes into a thiolate through incorporation of the oxygen into the adjacent link and this isomerised incorporated epoxide (forming a link akin insertion of a virtual thirane) has been termed Α.17,20 Thus, we hypothesised that other monomer combinations which undergo Li catalysed ROCOP could be suitable for sequence selective ROTERP. Xiong and coworkers very recently reported RNCS/epoxide ROCOP mediated by a simple LiO′Bu catalyst and accordingly we hypothesised that isothiocyanates could undergo ROTERP (Fig. 1).12

Therefore, we investigated the terpolymerisation of PTA/PhNCS/BO at different monomer ratios and catalyst loadings at 80 °C with lithiumbenzyloxide (LiOBn) as the catalyst (Table 1). Gratifyingly, we find that mixtures comprising 15 eq. PhNCS and 5 eq. BO with LiOBn loadings of 1-8 mol% per equivalent of PTA form poly(ester-alt-alt-alt-dithioimidocarbonates) in quantitative polymer selectivity. The 1H NMR spectrum (Fig. 2) of the polymer shows two main aryl resonances corresponding to a symmetrically substituted terephthalate unit (δ = 7.67 and 7.47 ppm) in an approximate 4 : 5 integrative ratio with respect to the NPh aromatic resonances indicating a 1 : 1 ratio of the two aromatic moieties.

Furthermore, there is one main resonance for the CH2R2 protons (δ = 5.23 ppm) in a 2 : 4 integrative ratio and one main signal for the CH3R2 protons (δ = 3.80-2.90 ppm) in a 4 : 4 integrative ratio to the aromatic terephthalate signals respectively. The 13C[1H] NMR spectrum (Fig. 2) reveals that the links formed are primarily dithioimidocarbonates R-S-C(≡NPh)-S-R (δ = 159.1 ppm, ≈31% of all links) and esteresters R-C(=O)-O-R (δ = 166.6 ppm, ≈64% of all links), alongside minor tho-
Due to overlapping $^1$H NMR resonances for the monothioimidocarbonate $R$-O resonance, in alkoxide chain ends or from insertion of thiolate chain from incomplete O/S exchange and insertion of PTA or PhNCS between PTA/BO and PhNCS/BO and are inferred to result or LiOCH$_2$Ph (LiOBn).

Table 1

<table>
<thead>
<tr>
<th>LiOBn : PTA : BO : PhNCS$^d$</th>
<th>Time [min]</th>
<th>PTA conversion$^b$</th>
<th>Coupling selectivity$^c$</th>
<th>Ester selectivity$^d$</th>
<th>$M_n$,$e$ [kDa] ($\Delta$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 : 6 : 32 : 93</td>
<td>2</td>
<td>99%</td>
<td>&gt;99%</td>
<td>97.5%</td>
<td>2.51 (1.15)</td>
</tr>
<tr>
<td>1 : 13 : 62 : 187</td>
<td>3</td>
<td>98%</td>
<td>&gt;99%</td>
<td>97.5%</td>
<td>5.5 (1.17)</td>
</tr>
<tr>
<td>1 : 25 : 125 : 375</td>
<td>6</td>
<td>98%</td>
<td>&gt;99%</td>
<td>97.5%</td>
<td>9.94 (1.20)</td>
</tr>
<tr>
<td>1 : 50 : 250 : 750</td>
<td>30</td>
<td>95%</td>
<td>&gt;99%</td>
<td>97.5%</td>
<td>13.61 (1.33)</td>
</tr>
<tr>
<td>1 : 100 : 500 : 1500</td>
<td>480</td>
<td>86%</td>
<td>&gt;99%</td>
<td>97.5%</td>
<td>14.0 (1.49)</td>
</tr>
<tr>
<td>1 : 50 : 250 : 125</td>
<td>30</td>
<td>&gt;99%</td>
<td>&gt;99%</td>
<td>91%</td>
<td>22.18 (1.32)</td>
</tr>
<tr>
<td>1 : 50 : 250 : 250</td>
<td>30</td>
<td>&gt;99%</td>
<td>&gt;99%</td>
<td>94%</td>
<td>21.6 (1.29)</td>
</tr>
<tr>
<td>1 : 50 : 250 : 500</td>
<td>30</td>
<td>&gt;99%</td>
<td>&gt;99%</td>
<td>95%</td>
<td>17.61 (1.28)</td>
</tr>
<tr>
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<td>480</td>
<td>86%</td>
<td>&gt;99%</td>
<td>95%</td>
<td>17.61 (1.28)</td>
</tr>
<tr>
<td>1 : 50 : 250 : 750$^f$</td>
<td>30</td>
<td>0%</td>
<td>—</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>1 : 50 : 250 : 375$^h$</td>
<td>30</td>
<td>&gt;99%</td>
<td>95%</td>
<td>96%</td>
<td>18.09 (1.38)</td>
</tr>
</tbody>
</table>

$a$ Copolymerisation at $T = 80 \, ^\circ\mathrm{C}$, LiOBn generated in situ from LiHMDS and BnOH (see ESI$^\dagger$).

$b$ Determined by comparison of the relative integrals, in the normalised $^1$H NMR spectrum (CDCl$_3$, 25 °C, 400 MHz), of terephthalate CH resonance due to (co)terminal polymer and PTA.

$c$ Determined by comparison of the relative integrals, in the normalised $^1$H NMR spectrum (CDCl$_3$, 25 °C, 400 MHz), of tertiary CH resonance due to (co)terminal polymer and cyclic thiimidocarbonate and polyether.

$d$ Determined by comparison of the relative integrals, in the normalised $^1$H NMR spectrum (CDCl$_3$, 25 °C, 400 MHz), of tertiary CH resonance due to (co)terminal polymer and cyclic thiimidocarbonate and polyether.

$e$ Determined by comparison of the relative integrals, in the normalised $^1$H NMR spectrum (CDCl$_3$, 25 °C, 400 MHz), of tertiary CH resonance due to (co)terminal polymer and cyclic thiimidocarbonate and polyether.

$f$ Determined by SEC (size-exclusion chromatography) measurements conducted in THF, using narrow MW polystyrene standards to calibrate the instrument.

$g$ NaHMDS was employed in place of PhNCS.

$h$ KHMDMS was employed in place of LiHMDS.
rearrangement and propagation from alkoxide intermediates. In these cases, we also observe the onset of PhNCS/BO ROCOP to form poly(monothioimidocarbonates) once all PTA is consumed as confirmed by $^1$H (Fig. S6†) and $^{13}$C NMR (Fig. S7†) spectroscopy. This leads to poly(monothioimidocarbonate) blocks forming adjacent to the ROTERP blocks which also explains the higher molecular weights obtained at the same PTA loadings. The new monomer combination is furthermore compatible with PTA/CS$_2$/BO ROTERP. Tetrapolymerisation of PTA and BO with PhNCS and CS$_2$ results in poly(ester-alt-ester-alt-heterocarbonate) formation in which both RNCS (forming dithioimidocarbonates) and CS$_2$ (forming trithiocarbonates) are incorporated into the polymer. NMR (Fig. S11–S15†) and FTATR-IR (Fig. S17†) clearly show the presence of trithiocarbonate ($\delta = 222.8$ ppm, $\tilde{\nu} = 1063$ cm$^{-1}$) links in addition to the other links from PTA/PhNCS/BO ROTERP. Corresponding to Table 1 run #11, a polymer comprising 67% arylester, 23% trithiocarbonate, 6% dithioimidocarbonate, 3.5% thioester and 0.5% monothioimidocarbonate links is produced (96% ROTERP links and 4% ROCOP errors). Apparently, PTA/PhNCS/BO ROP/TERP follows a similar polymerisation mechanism (Fig. S31†) as the one proposed for PTA/CS$_2$/BO ROTERP. It involves a central O/S rearrangement step and preferential insertion of PTA into alkoxide chain ends and of the heteroallene into thiolate chain ends at which point a mixture of heteroallenes can be employed which are then both incorporated at this insertion step. Interestingly, although PhNCS and CS$_2$ are employed in an equimolar ratio, CS$_2$ is incorporated preferentially in 77% selectivity. We could confirm this observation by varying the initial PhNCS:CS$_2$ ratio in which CS$_2$ was always incorporated to a greater degree than employed in the initial monomer feed (see Fig. S18†). In ROCOP, switchable catalysis has been established as an elegant and valuable tool to synthesise blockpolymers with useful material properties. Here a suitable catalyst first mediates the ROP of for example cyclic esters (e.g. $\varepsilon$DL forming PDL) with epoxides present in the mixture until the second ROCOP monomer (e.g. CO$_2$) is added causing immediate termination of ROP and the onset of (e.g. CO$_2$/epoxide forming polycarbonate) ROCOP to form a ROCOP block connected to the ROP polymer. We recently showed that ROTERP is also suitable for switchable catalysis in combination with the lithium catalysed ROP of $\varepsilon$DL.$^{29}$ Having identified a new heteroallene that undergoes ROTERP, we were intrigued whether this monomer combination is also suitable for the construction of blockpolymers via switchable catalysis. Accordingly, we added PhNCS (750 eq. per LiOBn) and PTA (50 eq.) to polymerising $\varepsilon$DL (50 eq.) in BO (250 eq.) after 15 min at room temperature which completely and immediately stops the occurrence of $\varepsilon$DL ROP. Heating to 80 °C initiates ROTERP and a poly(ester-alt-ester-alt-dithioimidocarbonate) block grows from the PDL-chain-end until the reaction is stopped after 30 min. Under these conditions the ROTERP block consists of 64% arylester links, 31% dithioimidocarbonate links and 2.5% erroneous thioester and monothioimidocarbonate links, respectively (95% ROTERP links and 5% ROCOP errors). Switchable catalysis and block polymer formation were established by various methods: (i) no $\varepsilon$DL is consumed after ROTERP starts (Fig. S21†) and the $^{13}$C$^\text{1H}$ PDL ($\delta = 173.2$ ppm, Fig. S23†) remains unchanged showing that ROP stops and that no transesterification between blocks occurs; (ii) the number averaged molecular weight shifts from $M_n = 8.26$ ($D = 1.11$) to 14.57 kg mol$^{-1}$ ($D = 1.24$, Fig. 3), which shows the growth of existing chains rather than the initiation of new ones; (iii) $^{31}$P end group analysis shows the consumption of all PDL end groups (Fig. S30†); (iv) the composition of the resulting blockpolymer remains unchanged through multiple precipitations from DCM/MeOH and THF/pentane supporting that the blocks are joint; (vii) DSC analysis exhibits two $T_g$’s at $-46.5$ °C for the ROP block and 35.5 °C for the ROTERP block suggesting microphase separation in the solid-state which could be confirmed by AFM (Fig. 3); (viii) TGA analysis shows a stepwise thermal decomposition profile with two $T_d$$_{\text{onset}}$ at approximately 235 °C for the ROTERP block and 300 °C for the ROP block (Fig. 3). In conclusion, we have identified a new
monomer combination that undergoes lithium catalysed sequence selective terpolymerisation. Mixtures of PTA/PhNCS/BO forming poly(ester-alt-ester-alt-dithioimidocarbonate)s in up to 95% selectivity with respect to the erroneous links from PhNCS/BO and PTA/BO ROCOP. This has enabled the synthesis of complex tetrapolymers from quarternary monomer mixture or via switchable catalysis. Our results establish ROTERP as a valuable methodology for the synthesis of heteroatom containing blockpolymers.

**Conflicts of interest**

There are no conflicts of interests.

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**Notes and references**


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**Fig. 3** (Left) εDL ROP to ROTERP switchable catalysis sequence. (Right) overlayed SEC traces before and after switch as well as TGA, DSC and 3d image of AFM tapping topography (height range 4 nm) of the obtained blockpolymer. 3d display of topography is overlaid with a false colour map of the blockpolymer’s components.