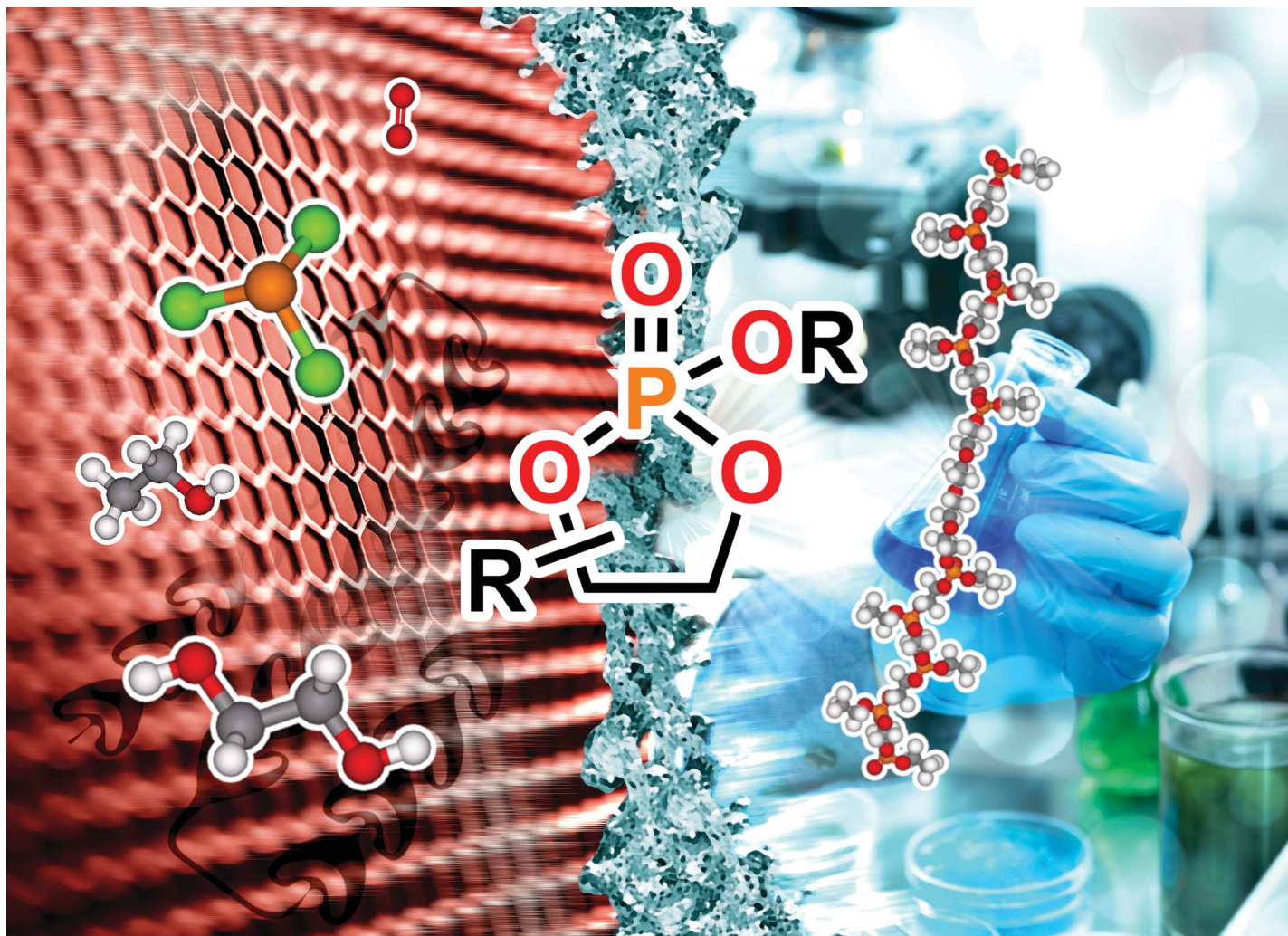




Showcasing research from Professor Monbaliu's laboratory,  
Center for Integrated Technology and Organic Synthesis,  
University of Liège, Belgium.

Accelerating the end-to-end production of cyclic phosphate  
monomers with modular flow chemistry

A compact and safe process for the efficient production  
of cyclic phosphate monomers. This multidisciplinary work  
was carried out within a joint European research program  
(Interreg EMR IN FLOW).

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# Accelerating the end-to-end production of cyclic phosphate monomers with modular flow chemistry†

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The biocompatibility, tunable degradability and broad functionalities of polyphosphoesters and their potential for biomedical applications have stimulated a renewed interest from Chemistry, Medicinal Chemistry and Polymer Sciences. Commercial applications of polyphosphoesters as biomaterials are still hampered because of the time and resource-intensive sourcing of their corresponding monomers, in addition to the corrosive and sensitive nature of their intermediates and by-products. Here, we present a groundbreaking challenge for sourcing the corresponding cyclic phosphate monomers by a different approach. This approach relies on the use of continuous flow technologies to intensify the end-to-end preparation of cyclic phosphate monomers with a semi-continuous modular flow platform. The applied flow technology mitigates both safety and instability issues related to the more classical production of cyclic phosphate monomers. The first flow module allows safe synthesis of a library of cyclic chlorophosphite building blocks and features in-line <sup>31</sup>P NMR real-time monitoring. After optimization on the microfluidic scale, this first module is successfully transposed toward mesofluidic scale with a daily throughput of 1.88 kg. Downstream of the first module, a second module is present, allowing the quantitative conversion of cyclic chlorophosphites with molecular oxygen toward chlorophosphate derivatives within seconds. The two modules are concatenable with a downstream semi-batch quench of intermediate chlorophosphate with alcohols, hence affording the corresponding cyclic phosphate monomers. Such a continuous flow setup provides considerable unprecedented advantages to safely and efficiently synthesize a library of versatile high value-added cyclic phosphate monomers at large scale. These freshly produced monomers can be successfully (co) polymerized, using either batch or flow protocols, into well-defined polyphosphoesters with assessed thermal properties and cytotoxicity.

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## Introduction

With major roles as building blocks of the genetic information through deoxy- and ribonucleic acids and as cell energy currency in metabolic processes through adenosine triphosphate, polyphosphoester (PPE) structures are ubiquitous in Biology.<sup>1</sup> Contrastingly to polyesters (*i.e.* based on carboxylic acid esters), synthetic PPEs have only found scarce industrial

applications such as flame retardants.<sup>2–4</sup> However, PPEs have recently gained a renewed interest from the academic community in light of their potential biomedical applications<sup>5</sup> such as drug delivery systems<sup>6–12</sup> and tissue engineering.<sup>13–17</sup>

Synthetic PPEs are especially appealing for their tunable (bio)degradability and biocompatibility properties.<sup>2,18</sup> A convenient synthetic method to obtain PPEs involves the ring opening polymerization (ROP) of cyclic phosphate monomers (CPMs), which enables the preparation of high molecular weight materials with narrow polydispersities.<sup>19</sup> Among the various CPMs, 5-membered ring monomers are currently more popular compared to their 6-membered ring counterparts. The ring strain of the 5-membered CPMs makes them more reactive during polymerization, yet also more sensitive to hydrolysis, and prone to efficient organocatalyzed ROP.<sup>2</sup> In sharp contrast to classical aliphatic polyesters, the physicochemical properties of PPEs, such as hydrophobicity,<sup>20,21</sup> can conveniently be adjusted through straightforward modifications of the pendant side-chain on the pentavalent phosphorous during the

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upstream monomer preparation. Moreover, introduction of a suitable functional pendant side-chain also enables post-polymerization modifications<sup>22,23</sup> and introduction of bioactive molecules,<sup>24,25</sup> which further expands the scope of applications of PPEs.

One of the most significant challenges met during the production of PPEs using ROP strategies actually relates to the sourcing of CPMs (Fig. 1). The preparation of such monomers is usually performed by a resource- and time-intensive 3-step procedure. The preparation of CPMs starts from  $\text{PCl}_3$  and a suitable diol (**1a**) to yield a cyclic chlorophosphite adduct **2a** with the concomitant release of 2 equiv. of gaseous  $\text{HCl}$ . Next, the chlorophosphite intermediate is further oxidized into its cyclic chlorophosphate counter-part **3a**. The CPM is obtained after a final functionalization using an adapted alcohol.<sup>18,26</sup> This process involves the use of hazardous  $\text{PCl}_3$  as starting material, corrosive and sensitive intermediates (chlorophosphite(*a*)tes **2a** and **3a**) and produces large amounts of  $\text{HCl}$  as a by-product. These circumstances lead to significant safety concerns upon scalability trials.

To address these issues that challenge upscaling a modular semi-continuous flow platform dedicated to the production of CPMs has been developed. The platform allows to produce CPMs with unprecedented versatility, efficiency and scalability starting from bulk chemicals. The modular flow platform features the concatenation (*i.e.*, the connection within a same uninterrupted system) of 3 chemical steps and mitigates the inherent safety hazards related to production of large amounts of by-product  $\text{HCl}$ , as well as to handling of corrosive intermediates and reactants. Moreover, this approach also significantly reduces the overall process time for production of chlorophosphite(*a*)te intermediates in a continuous fashion (batch: 5 h to 4 d; this flow protocol: <2 min). This study also demonstrates that model CPMs are successfully polymerized through either batch or continuous flow protocols. The absence of cytotoxicity of selected PPE samples is also assessed, as well as their thermal properties. This contribution unlocks the production of CPMs and consequent PPEs, and thus opens new avenues to access novel biomaterials and large-scale production of industrially

relevant PPEs by enabling their safer and efficient end-to-end production.

## Results and discussion

### Continuous flow preparation of cyclic chlorophosphites and analogs

The preparation of cyclic chlorophosphites in batch typically relies on the method of Lucas *et al.*,<sup>27</sup> which involves the careful addition of a diol to a solution containing  $\text{PCl}_3$  with the subsequent release of  $\text{HCl}$  (2 equiv.). Even with appropriate care during the addition process, reckless release of  $\text{HCl}$  gas raises significant safety concerns upon scale-up. Taking these hazards into account prompted us to adapt the synthesis of chlorophosphites under continuous flow conditions for enabling a smoother and more controllable generation of  $\text{HCl}$  gas, as well as a fast and efficient production of cyclic chlorophosphites and analogs.

The first exploratory optimizations involved the preparation of model compound **2a** through the addition of neat ethylene glycol (**1a**) to a concentrated solution of  $\text{PCl}_3$  in  $\text{CH}_2\text{Cl}_2$  at room temperature. The outlet of the reactor was directly connected to a closed surge with a flow of an inert gas to continuously flush gaseous  $\text{HCl}$  out of the reactor effluent toward an alkaline aqueous trap. Starting from a 1 M  $\text{PCl}_3$  solution, reducing the residence time to 1 min allowed complete conversion of  $\text{PCl}_3$  (Table 1, entries 1–4). Next, to maximize the output, the concentration of the  $\text{PCl}_3$  solution was increased (entries 5 and 6) with an appropriate adjustment of the counter-pressure to avoid  $\text{HCl}$  degassing inside the coil reactor. A starting 3 M solution (entry 5) maintained complete conversion and excellent selectivity, whilst affording a 76% isolated yield after fractional distillation under reduced pressure. A very high hourly productivity of 14.8 g was obtained using a 1 mL internal volume coil reactor, despite its minimal footprint. Further increasing the concentration of the starting solution (entry 6) led to a biphasic regime and to the appearance of major side-products; thus, the concentration was set to 3 M for the  $\text{PCl}_3$  solution as a reference for the following experiments. A variety of solvents were assessed for the production of **2a**, yet leading to similar (entries 7, 9 and 10) or lower (entry 8) yields. 2-Methyl-tetrahydrofuran (MeTHF) emerged as a more sustainable alternative to  $\text{CH}_2\text{Cl}_2$ , as it is biobased and more favorable than  $\text{CH}_2\text{Cl}_2$  according to the CHEM21 selection guide.<sup>28</sup> MeTHF also supersedes THF in terms of health and environmental scoring. In-line  $^{31}\text{P}$  NMR was conveniently implemented downstream the reactor for real-time reaction monitoring, hence accelerating the optimization process as well as improving process safety (see ESI, Section 2.5.1†). Implementation of in-line monitoring indeed contributes to ensure that hazardous  $\text{PCl}_3$  is fully converted and that no major impurities are present, while avoiding time-consuming high-field NMR analysis that could lead to a degradation of the crude sample containing a sensitive chlorophosphite. Chlorophosphite **2a** was detected as a sole peak ( $\sim 168$  ppm) during in-line monitoring while the absence of  $\text{PCl}_3$  ( $\sim 219$  ppm) was confirmed. No other peaks were observed by  $^{31}\text{P}$  NMR.

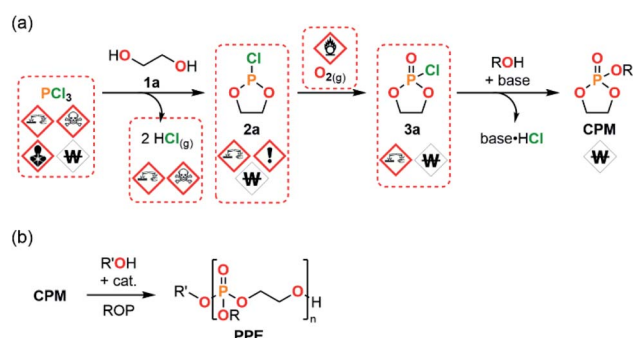
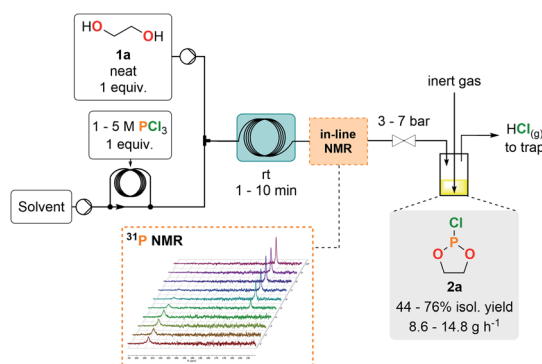


Fig. 1 (a) 3-step synthetic pathway commonly used for the preparation of CPMs involving hazardous and/or sensitive reactants, intermediates and by-products. (b) Ring-opening polymerization strategy used for the production of PPEs starting from CPMs.

Table 1 Optimization of the continuous flow preparation of cyclic chlorophosphite **2a**<sup>a</sup>

| Entry          | Res. time [min] | Solvent                         | PCl <sub>3</sub> conc. [mol L <sup>-1</sup> ] | P [bar] | Conv. PCl <sub>3</sub> <sup>b</sup> [%] | Isol. yield [%] | Prod. [g h <sup>-1</sup> ] |
|----------------|-----------------|---------------------------------|-----------------------------------------------|---------|-----------------------------------------|-----------------|----------------------------|
| 1              | 10              | CH <sub>2</sub> Cl <sub>2</sub> | 1                                             | 3       | >99                                     | n.d.            | n.d.                       |
| 2              | 5               | CH <sub>2</sub> Cl <sub>2</sub> | 1                                             | 3       | >99                                     | n.d.            | n.d.                       |
| 3              | 2.5             | CH <sub>2</sub> Cl <sub>2</sub> | 1                                             | 3       | >99                                     | n.d.            | n.d.                       |
| 4              | 1               | CH <sub>2</sub> Cl <sub>2</sub> | 1                                             | 3       | >99                                     | n.d.            | n.d.                       |
| 5              | 1               | CH <sub>2</sub> Cl <sub>2</sub> | 3                                             | 5       | >99                                     | 76              | 14.8                       |
| 6 <sup>c</sup> | 1               | CH <sub>2</sub> Cl <sub>2</sub> | 5                                             | 7       | >99                                     | n.d.            | n.d.                       |
| 7              | 1               | MTBE                            | 3                                             | 5       | >99                                     | 62              | 12.2                       |
| 8              | 1               | MeCN                            | 3                                             | 5       | >99                                     | 44              | 8.6                        |
| 9              | 1               | THF                             | 3                                             | 5       | >99                                     | 72              | 14.0                       |
| 10             | 1               | MeTHF                           | 3                                             | 5       | >99                                     | 74              | 14.4                       |

<sup>a</sup> General conditions: room temperature, continuous flow coil reactor of 1 mL of internal volume. <sup>b</sup> Determined by <sup>31</sup>P NMR. <sup>c</sup> Two phases present in the effluent of the reactor, impurities detected by <sup>31</sup>P NMR. MTBE = methyl *tert*-butyl ether, MeCN = acetonitrile, THF = tetrahydrofuran, MeTHF = 2-methyltetrahydrofuran.

With a highly productive and compact continuous flow system that mitigates the hazardous preparation of **2a** in hand, the scope of the procedure was next extended to a variety of cyclic chlorophosphites and analogs **2**. Starting from various 1,2-diol and derivatives **1** (including potentially biobased derivatives **1a–g, j, l, m**)<sup>29–33</sup> allowed to rapidly generate a library of chlorophosphites bearing a range of backbone modulations (Fig. 2). Good to high isolated yields were obtained for most examples, except for **2e** for which the final vacuum distillation led to significant decomposition (see ESI, Section 2.4.1†). Switching from a 1,2-diol to a 1,2-dithiol (**1h**) accessed dithiaphospholane derivative **2h** in 58% of isolated yield, which was reported for the functionalization of nucleosides<sup>34,35</sup> and the preparation of phospholipid analogs.<sup>36</sup>

With diverse applications as a phosphitylation agent for the preparation of glycosyl donors<sup>37</sup> or the derivatization of lignin samples for characterization by <sup>31</sup>P NMR,<sup>38–40</sup> the preparation of **2i** was next envisioned. Exploring the reaction with hindered 1,2-diol such as pinacol (**1i**) led to a major loss of selectivity and precluded the isolation of the product. According to the prior Art,<sup>41</sup> highly sterically congested pinacol-like diols are associated with a much slower formation rate toward the corresponding cyclic chlorophosphites, hence allowing for competitive formation of a biphosphite derivative. The latter, in presence of HCl, underwent dealkylation eventually leading to the formation of 2 equivalents of cyclic H-phosphonate. Attempts to avoid

formation of such side-products relied on addition of a base to the reaction mixture; however, in order to avoid extensive precipitation of hydrochlorides upon continuous flow operation, organic bases known to form ionic liquids under protonation, such as 1-methylimidazole, were privileged.<sup>42</sup> Minor adaptations from the original continuous flow system were implemented with the selection of acetonitrile as a solvent to allow solubilization of the ionic liquid (see ESI, Section 2.5.2†). Chlorophosphite **2i** was isolated in 66% yield using the adapted system accordingly, leading to a hourly productivity of 7.3 g. With such an adaptation, preparation of more demanding building blocks **2j–m** was foreseen (Fig. 2). The six-membered ring dioxaphosphinane **2j** was successfully produced in the presence of 1-methylimidazole, leading to a 53% isolated yield. This derivative is used in the preparation of various six-membered CPMs.<sup>2,18</sup> Oxathiaphospholane **2k** and oxazaphospholidine **2i** were produced in average to high yields, thus even further broadening the scope. Finally, the synthesis of the diazaphospholidine derivative **2m** was developed but required additional adaptations to avoid clogging of the continuous flow system. Switching to 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU) as a base and lowering the concentration of the starting feed solutions prevented precipitation of solid residues, therefore enabling the isolation of **2m** in a moderate yield (43%). This continuous flow protocol offers a compact solution for on-demand production of versatile cyclic P(III) adducts while enabling a gram-scale production in minutes



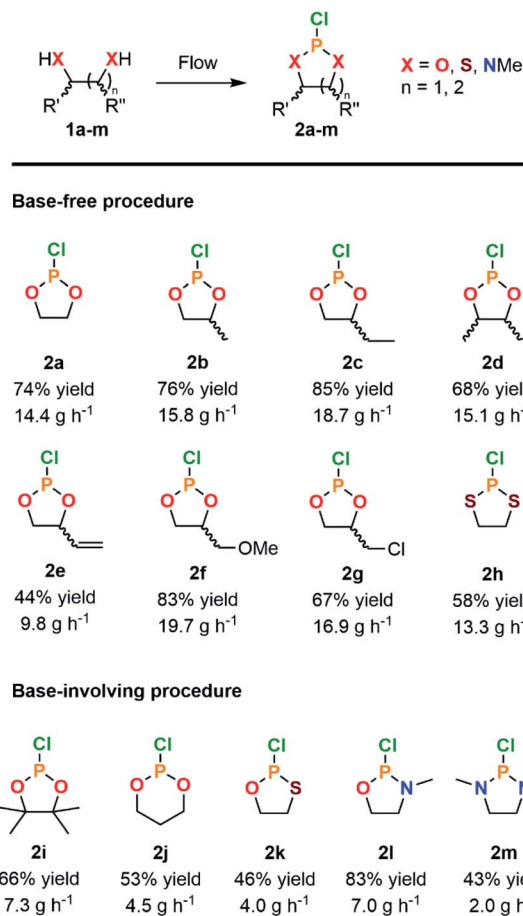


Fig. 2 Scope of cyclic adducts **2a–m** obtained from the continuous flow reaction of  $\text{PCl}_3$  with various diols and analogs through a base-free (see Table 1, entry 10) or a base-involving (see ESI, Section 2.5.2†) procedure.

and mitigating the safety issues related with synthesis of such derivatives.

### Scalability of the continuous flow preparation of cyclic chlorophosphite **2a**

The optimized procedure for the generation of cyclic chlorophosphites in a microfluidic environment bears significant potential for further scalability trials.<sup>43,44</sup> To this end, the preparation of **2a** was transposed in a commercial glass mesofluidic continuous flow reactor (see ESI, Section 2.4.3†). The reactor setup consisted in two glass fluidic modules (FMs), fluidically connected in series (5.4 mL total internal volume). The production of **2a** was conveniently handled with the controlled release of the  $\text{HCl}$  gas by-product through a dome-type back-pressure regulator (4 bar) downstream the reactor. The effluent was collected in a closed vessel under a continuous flow of argon to flush out  $\text{HCl}$  toward an alkaline aqueous trap. The mesofluidic scale was associated with an isolated yield of 74% and a daily production of 1.88 kg under safe conditions ( $\text{STY} = 348 \text{ kg per L per day}$ ).

### Cyclic phosphate monomers (CPMs) production in a semi-continuous flow platform

The concatenation of the following steps toward functionalized CPMs was envisioned. The oxidation of cyclic chlorophosphites is typically carried out with a stream of gaseous molecular oxygen which is bubbled in an organic solution containing the chlorophosphite. The reaction time needed for this batch procedure typically varies between 8 h to 4 days.<sup>8,26,45</sup> Continuous flow oxidation of the model compound **2a** was optimized using molecular oxygen. The lab-scale continuous flow setup included the upstream generation of **2a**, which was further reacted with oxygen in a second PFA coil reactor (see ESI, Section 2.5.3†). Quick analysis of the effluent by a benchtop  $^{31}\text{P}$  NMR allowed to qualitatively control the conversion and selectivity of the reaction. A significant improvement of the selectivity was observed upon using a coil reactor with a smaller internal diameter and an arrowhead-type micromixer system. The mixing efficiency appeared as a much powerful leverage than residence time for the oxidation step, indeed, the slug flow regime of the gas–liquid system combined with a medium counter-pressure (15 bar), a smaller internal diameter of the coil reactor and a specific arrowhead micromixer (see Table S1 in the ESI†) leads to an improved interfacial area, mixing and solubility of oxygen within the liquid phase and, thus, to high mass transfer and improved performances. Quantitative conversion with 59% of selectivity toward **3a** was obtained under 65 °C within 21 s of residence time in the presence of 4 equiv. of molecular oxygen. Despite these very promising results, chlorophosphate **3a** rapidly degraded after collection.

It was therefore decided to directly implement the last condensation reaction to access CPMs by directly collecting the reactor effluent in a surge containing an alcohol and a base. Such a fully concatenated protocol would not only enable the neutralization of the remaining  $\text{HCl}$  gas, but also the reaction of the intermediate chlorophosphate to form the desired CPM directly from  $\text{PCl}_3$  and a diol. It was indeed foreseen that avoiding the handling of intermediate chlorophosphi(a)tes **2** and **3** would significantly increase process safety and efficiency. Using the optimized conditions for the preparation of **2a** and **3a** (see Table 1, entry 10 and ESI, Table S3, entry 8†), a semi-

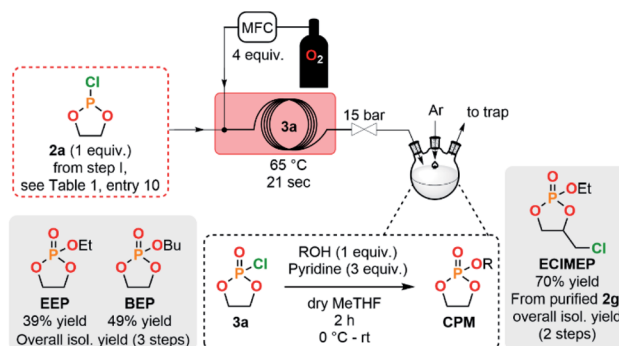


Fig. 3 Concatenated semi-continuous flow process for the production of cyclic phosphate monomers (see ESI, Fig. S2 and S3† for the detailed setups).

continuous flow system was developed (Fig. 3). The upstream generation of **2a** allowed its direct and quick conversion to **3a** and a final functionalization toward a CPM in the semi-batch reactor using an adapted alcohol in presence of pyridine. This flexible process allowed generation of various CPMs from bulk chemicals by conveniently switching the alcohol present in the collection surge. Two model monomers **EEP** and **BEP** were isolated in less than a day starting from  $\text{PCl}_3$  and **1a** using this procedure and required only one final purification step. Moreover, the overall isolated yields obtained (39% and 49% for **EEP** and **BEP**, respectively) over the 3 chemical steps outperform the typical output achieved in batch (17–21%).<sup>8,46</sup> The concatenation under continuous flow indeed contributes to reducing the number of purifications required and has thus a positive impact on the overall yield. Minor traces of a cyclic *H*-phosphonate were noticed in samples of **EEP** and **BEP** (see ESI, Section 2.5.4†) potentially highlighting a partial hydrolysis of **2a** in the system.<sup>47</sup> Starting directly from purified **2a** led to a high purity monomer free of *H*-phosphonate impurities through a 2-step semi-continuous flow procedure (see ESI, Section 2.5.5†).

A novel **ECIMEP** monomer bearing an ethoxy pendant group, additionally to a chloromethyl functionality linked to the cyclic backbone, was successfully isolated using an adapted procedure (see Fig. 3 and ESI, Section 2.4.5†). The 2-step process started from purified **2g**, which allowed to avoid side-products from the synthesis of the chlorophosphite and, subsequently, to obtain high purity **ECIMEP** in 70% of isolated yield over 2 chemical steps. A batch procedure for the preparation of the monomer **ECIMEP** was additionally developed (see ESI, Sections 2.5.6–2.5.8†).

### Batch and flow (co)polymerization of cyclic phosphate monomers (CPMs)

The development of a semi-continuous flow system for the convenient and safer preparation of CPMs paves the way toward a better accessibility to PPEs as source for potential flame-retardants or biomedical applications at lab- and industrial-scale. To demonstrate the potential of this methodology, **EEP** was first selected as a reference monomer to assess its polymerizability. **EEP** ROP was first carried out in a batch reactor using DBU and a thiourea derivative (TU) as a well-established organocatalytic system<sup>48</sup> in the presence of tetraethylene glycol (**TEG**) as an initiator (Table 2, entries 1–3). The **EEP** monomer produced through the semi-continuous flow platform was successfully polymerized accordingly (entry 1) with a good control of the macromolecular parameters. A well-defined polymer **TEG-PEEP** homopolymer with a narrow dispersity ( $D = 1.14$ ) and an average molar mass of  $13\,600\text{ g mol}^{-1}$  close to the targeted value ( $12\,000\text{ g mol}^{-1}$ ) was obtained. The small molar mass difference was attributed to unavoidable transesterification side-reactions at high monomer conversion.<sup>48</sup>

Polymerization reactions can also benefit from continuous flow systems, leading to lower dispersity and improved reproducibility by taking advantage of better thermal management and mass transfer.<sup>49</sup> The homopolymerization of **EEP** was further explored under microfluidic conditions (entries 2 and 3)

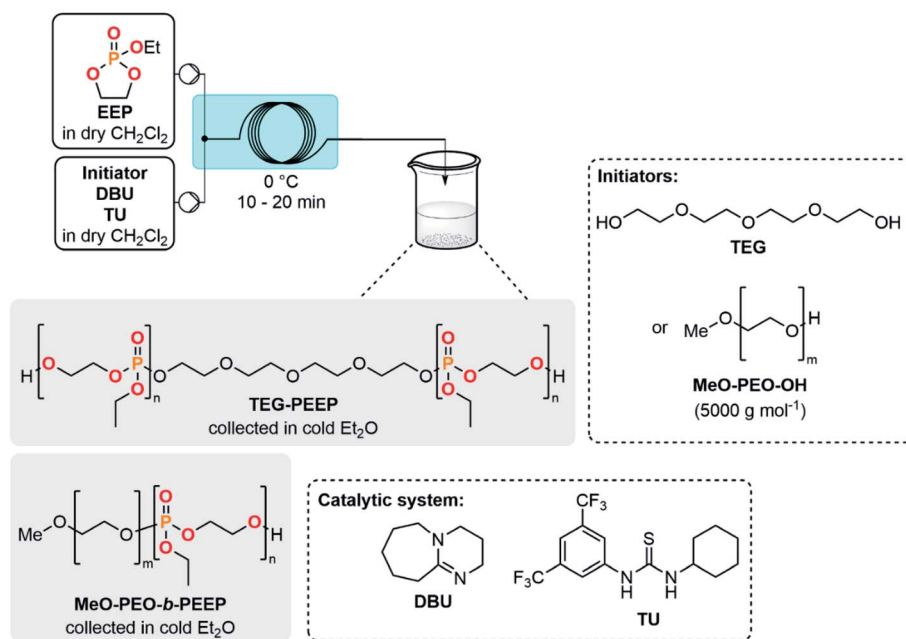
with inspiration from the study of Junkers *et al.*<sup>23</sup> The outlet of the reactor was directly collected in cold  $\text{Et}_2\text{O}$ , triggering precipitation of PPE. A residence time of 10 min at  $0\text{ }^\circ\text{C}$  (entry 2) led to a molar mass of  $5300\text{ g mol}^{-1}$  and a low dispersity ( $D = 1.06$ ). Interestingly, increasing the residence time to 20 min (entry 3) led to an increase of molar mass ( $M_n\text{ NMR} = 9400\text{ g mol}^{-1}$ ) while keeping a low polydispersity index ( $D = 1.11$ ). The molecular weight distributions were monomodal for **TEG-PEEP** homopolymers obtained under flow conditions, whereas a shoulder toward high molecular weight was observed for the **TEG-PEEP** sample obtained under batch conditions (see ESI, Sections 2.8.1–2.8.3†).

Design of novel block or graft copolymers based on the association of poly(ethylene oxide) (PEO) and PPEs have already demonstrated their potential for applications such as drug delivery systems<sup>9,24,25,50–52</sup> and hydrogels.<sup>53,54</sup> High reproducibility of continuous flow procedures is particularly appealing for preparation of such biomaterials to face demanding regulatory policies required for potential commercialization phases.

Using a PEO-based macroinitiator (**MeO-PEO-OH**,  $5000\text{ g mol}^{-1}$ ) in conjunction with a DBU/TU catalytic system and based on the batch study of Jérôme *et al.*,<sup>20</sup> the polymerization of **EEP** was carried out toward the formation of a **MeO-PEO-*b*-PEEP** diblock copolymer under continuous flow conditions (entry 4). Performing the reaction at  $0\text{ }^\circ\text{C}$  within a residence time of 20 min led to the targeted  $M_n$  of  $7000\text{ g mol}^{-1}$  toward a well-defined block copolymer ( $D = 1.05$ ). Cytotoxicity assays of **MeO-PEO-*b*-PEEP** on bovine fibroblast cells and human umbilical vein endothelial cells revealed that no cytotoxic effects were observed for a concentration of up to  $0.1\text{ mg mL}^{-1}$  of copolymer (see ESI, Section 2.5.12†). These results are in agreement with data collected in a previous study regarding PEO–PPE diblock copolymers,<sup>55</sup> which paves the way for a potential biomedical application.

Following the encouraging results collected for the polymerization of **EEP**, the polymerization of **ECIMEP** (obtained from **2g**, see Fig. 3) was investigated for the production of novel PPEs. Indeed, the presence of a chloromethyl functionality on the polymer backbone will first modulate the physicochemical properties of the corresponding poly(**ECIMEP**) and additionally enable further post-polymerization derivatization as similarly described in literature for poly( $\epsilon$ -caprolactone).<sup>56,57</sup> Moreover, the introduction of a chloromethyl moiety directly linked to a carbon of the polymer main-chain could render a more stable functionality after derivatization over elimination by hydrolysis compared to a more conventional functionalization through the pendant side-chain linked to the phosphorus atom which can be degraded under basic or acidic conditions.<sup>58</sup> For this purpose, homopolymerization of **ECIMEP** monomer was carried out in a batch process with similar conditions used for **EEP**. Unfortunately, these conditions are not adapted for the polymerization of **ECIMEP** and did not lead to the formation of the expected homopolymer. This might be explained by a higher steric hindrance compared to the **EEP** monomer but also by the potential formation of a secondary alcohol at the  $\omega$  chain-end after the initiation step, which could be insufficiently reactive to polymerize further with a second **ECIMEP**. Nevertheless,



Table 2 Batch and continuous flow polymerization trials of EEP<sup>a</sup>

| Entry | Reactor | Initiator  | EEP equiv. | DBU equiv. | TU equiv. | Reaction time <sup>b</sup> [min] | M <sub>n</sub> NMR <sup>c</sup> [g mol <sup>-1</sup> ] | M <sub>n</sub> SEC <sup>d</sup> [g mol <sup>-1</sup> ] | D (SEC) <sup>e</sup> | M <sub>n</sub> <sup>theo f</sup> [g mol <sup>-1</sup> ] |
|-------|---------|------------|------------|------------|-----------|----------------------------------|--------------------------------------------------------|--------------------------------------------------------|----------------------|---------------------------------------------------------|
| 1     | Batch   | TEG        | 80         | 15.8       | 4.6       | 30                               | 13 600                                                 | 3300                                                   | 1.14                 | 12 000                                                  |
| 2     | Flow    | TEG        | 80         | 15.8       | 4.6       | 10                               | 5300                                                   | 1700                                                   | 1.06                 | 12 000                                                  |
| 3     | Flow    | TEG        | 80         | 15.8       | 4.6       | 20                               | 9400                                                   | 2100                                                   | 1.11                 | 12 000                                                  |
| 4     | Flow    | MeO-PEO-OH | 13         | 5          | 4.6       | 20                               | 7000                                                   | 8300                                                   | 1.05                 | 7000                                                    |

<sup>a</sup> General conditions: temperature = 0 °C, atmospheric pressure, 1 equiv. of initiator, for the batch experiment 6.58 mmol of EEP used, for the flow experiments: conc.-EEP = 1.10 (entries 2 and 3) or 1.65 (entry 4) mol L<sup>-1</sup>. <sup>b</sup> Defined as residence time for continuous flow experiments. <sup>c</sup> Average molar mass of the (co)polymer determined by <sup>1</sup>H NMR as follow: M<sub>n</sub> = DP<sub>EEP</sub> × 152 + 194.23 for TEG-PEEP and M<sub>n</sub> = DP<sub>EEP</sub> × 152 + 5000 for MeO-PEO-b-PEEP. <sup>d</sup> Determined by SEC according to a polystyrene calibration. <sup>e</sup> Molecular weight distribution (PDI) determined by SEC. <sup>f</sup> Targeted molar mass of the (co)polymer after complete monomer conversion: M<sub>n</sub> = n<sub>EEP</sub>/n<sub>initiator</sub> × 152 + 194.23 for TEG-PEEP and M<sub>n</sub> = n<sub>EEP</sub>/n<sub>initiator</sub> × 152 + 5000 for MeO-PEO-b-PEEP.

random copolymerization of ECIMEP with EEP using benzyl alcohol (BzOH) as an initiator was successful when the molar ratio of ECIMEP in the co-monomer feed was lower or equal to 0.50 (Table 3, entries 1 and 2). BzO-PEEP-co-PECIMEP copolymers of two different compositions and molar mass were prepared. Monomodal molecular weight distribution with low dispersity were obtained for both copolymers.

Regarding the effects of the chloromethyl functionality on the physicochemical properties of both BzO-PEEP-co-PECIMEP copolymers as compared to a BzO-PEEP homopolymer reference (entry 3), a limited influence on the value of the glass transition temperature (*T<sub>g</sub>*) of the copolymers was observed. Both copolymers are amorphous with a *T<sub>g</sub>* varying between -70 and -60 °C. Concerning the thermal stability, measured by TGA

Table 3 Batch (co)polymerization trials toward BzO-PEEP-co-PECIMEP and BzO-PEEP<sup>a</sup>

| Entry | Target polymer      | Molar fraction of ECIMEP in the comonomer feed | Molar fraction of ECIMEP <sup>b</sup> | M <sub>n</sub> NMR <sup>c</sup> [g mol <sup>-1</sup> ] | M <sub>n</sub> SEC <sup>d</sup> [g mol <sup>-1</sup> ] | D <sup>e</sup> (SEC) | T <sub>g</sub> <sup>f</sup> [°C] |
|-------|---------------------|------------------------------------------------|---------------------------------------|--------------------------------------------------------|--------------------------------------------------------|----------------------|----------------------------------|
| 1     | BzO-PEEP-co-PECIMEP | 0.50                                           | 0.50                                  | 11 300                                                 | 4000                                                   | 1.15                 | -69.7                            |
| 2     | BzO-PEEP-co-PECIMEP | 0.25                                           | 0.31                                  | 4700                                                   | 2300                                                   | 1.08                 | -61.5                            |
| 3     | BzO-PEEP            | 0                                              | 0                                     | 3300                                                   | 2600                                                   | 1.05                 | -71.9                            |

<sup>a</sup> General conditions: initiator = benzyl alcohol (BzOH, 0.1 mmol), n<sub>EEP</sub> + n<sub>ECIMEP</sub> = 17.5 mmol, n<sub>DBU</sub> = 6.5 mmol, n<sub>TU</sub> = 2.3 mmol, temperature = 0 °C, reaction time = 30 min. <sup>b</sup> Molar fraction of ECIMEP subunits in the copolymer determined by <sup>1</sup>H NMR. <sup>c</sup> Average molar mass of the (co) polymer determined by <sup>1</sup>H NMR as follow: M<sub>n</sub> = DP<sub>EEP</sub> × 152 + DP<sub>ECIMEP</sub> × 201 + 108. <sup>d</sup> Determined by SEC according to a polystyrene calibration. <sup>e</sup> Molecular weight distribution (PDI) determined by SEC. <sup>f</sup> Glass transition temperature determined by DSC.



(see ESI, Sections 2.8.5–2.8.7†), the presence of the chloromethyl moiety modifies the degradation profile of **BzO-PEEP-co-PECIMEP** copolymers with an increase of the percentage of combustion residues at 600 °C compared to **BzO-PEEP**. These results pave the way for potential applications of these new copolymers as flame-retardant materials which are currently investigated in a follow-up study.

## Conclusions

A semi-continuous flow modular platform is described and allows convenient and safer preparation of CPMs directly from bulk chemicals. The first module was dedicated to the preparation of cyclic chlorophosphite building blocks with high productivities despite a small footprint system. This procedure allowed to efficiently mitigate the safety concerns inherently associated with the preparation of CPMs, while benefiting from real-time  $^{31}\text{P}$  NMR monitoring. The process was further expanded to a large scope of cyclic chlorophosphites and (O,S)-, (O,N)-, (N,N)- and (S,S)-analogs. The preparation of a model cyclic chlorophosphite was transposed to pilot-scale with a mesofluidic reactor upon minor parameters adjustments leading to a daily productivity of ~2 kg. Next, the concatenation of the first module was implemented with (a) a downstream continuous flow oxidation using molecular oxygen and (b) a subsequent final functionalization using an alcohol in a semi-batch reactor. The oxidation step toward a cyclic chlorophosphate was intensified by taking advantage of a high mixing efficiency of the continuous flow system leading to a residence time of 21 s to achieve a quantitative conversion of the chlorophosphite. Finally, both cyclization and oxidation modules were telescoped with a downstream semi-batch reactor to conveniently perform the final functionalization toward CPMs using an adapted alcohol. The fully integrated semi-continuous flow platform allowed the production of CPMs while mitigating significant safety hazards. Next, a model CPM, **EEP**, obtained from the semi-continuous flow platform was successfully (co)polymerized by batch and continuous flow procedures leading to well-defined and biocompatible PPEs. Finally, **ECIMEP**, a novel CPM bearing a chloromethyl functionality was successfully copolymerized leading to new PPEs with promising thermal properties. With the development of a semi-continuous flow platform for the efficient end-to-end production of CPMs, we forecast an accelerated and easier access to PPEs and their industrial applications by enabling a safer, more convenient and scalable preparation of the source monomers.

## Data availability

All experimental supporting data are provided in the ESI.†

## Author contributions

RM planned, executed, optimized the experiments in batch and flow for the preparation of CPMs and wrote the manuscript. RM and RR performed the polymerization experiments in batch and

flow. RR executed the assessment of thermal properties of PPEs. NMSVDA and DGMM are responsible for the biocompatibility studies and related cytotoxicity assays and provided feedback to the manuscript. CJ advised and proofread the manuscript. JCMM supervised the project, advised, and corrected the manuscript.

## Conflicts of interest

There are no conflicts to declare.

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## Notes and references

- 1 C. T. Walsh, in *The Chemical Biology of Phosphorus*, ed. K. J. Weissman, Royal Society of Chemistry, Cambridge, 2020, pp. 3–26.
- 2 K. N. Bauer, H. T. Tee, M. M. Velencoso and F. R. Wurm, *Prog. Polym. Sci.*, 2017, **73**, 61–122.
- 3 A. B. Morgan and J. W. Gilman, *Fire Mater.*, 2013, **37**, 259–279.
- 4 J. L. Olmedo-Martínez, L. Meabe, R. Riva, G. Guzmán-González, L. Porcarelli, M. Forsyth, A. Mugica, I. Calafel, A. J. Müller, P. Lecomte, C. Jérôme and D. Mecerreyes, *Polym. Chem.*, 2021, **12**, 3441–3450.
- 5 S. Monge, B. Cannicconi, A. Graillet and J.-J. Robin, *Biomacromolecules*, 2011, **12**, 1973–1982.
- 6 Y. P. Borguet, S. Khan, A. Noel, S. P. Gunsten, S. L. Brody, M. Elsabahy and K. L. Wooley, *Biomacromolecules*, 2018, **19**, 1212–1222.
- 7 C. J. McKinlay, R. M. Waymouth and P. A. Wender, *J. Am. Chem. Soc.*, 2016, **138**, 3510–3517.
- 8 S. Schöttler, G. Becker, S. Winzen, T. Steinbach, K. Mohr, K. Landfester, V. Mailänder and F. R. Wurm, *Nat. Nanotechnol.*, 2016, **11**, 372–377.
- 9 F. Zhang, S. Zhang, S. F. Pollack, R. Li, A. M. Gonzalez, J. Fan, J. Zou, S. E. Leininger, A. Pavia-Sanders, R. Johnson, L. D. Nelson, J. E. Raymond, M. Elsabahy, D. M. P. Hughes,



- M. W. Lenox, T. P. Gustafson and K. L. Wooley, *J. Am. Chem. Soc.*, 2015, **137**, 2056–2066.
- 10 T.-M. Sun, J.-Z. Du, Y.-D. Yao, C.-Q. Mao, S. Dou, S.-Y. Huang, P.-Z. Zhang, K. W. Leong, E.-W. Song and J. Wang, *ACS Nano*, 2011, **5**, 1483–1494.
- 11 J. Liu, W. Huang, Y. Pang, X. Zhu, Y. Zhou and D. Yan, *Biomacromolecules*, 2010, **11**, 1564–1570.
- 12 H. Elzeny, F. Zhang, E. Ali, H. Fathi, S. Zhang, R. Li, M. A. El-Mokhtar, M. A. Hamad, K. Wooley and M. Elsabahy, *Drug Des., Dev. Ther.*, 2017, 483–496.
- 13 B. M. Watson, F. K. Kasper and A. G. Mikos, *Biomed. Mater.*, 2014, **9**, 025014.
- 14 S. Kootala, M. Tokunaga, J. Hilborn and Y. Iwasaki, *Macromol. Biosci.*, 2015, **15**, 1634–1640.
- 15 S. Kunomura and Y. Iwasaki, *J. Biomater. Sci., Polym. Ed.*, 2019, **30**, 861–876.
- 16 A. E. Polloni, V. Chiaradia, R. J. F. C. do Amaral, C. Kearney, B. Gorey, D. de Oliveira, J. V de Oliveira, P. H. H. de Araújo, C. Sayer and A. Heise, *Polym. Chem.*, 2020, **11**, 2157–2165.
- 17 R. Riva, U. Shah, J.-M. Thomassin, Z. Yilmaz, A. Lecat, A. Colige and C. Jérôme, *Biomacromolecules*, 2020, **21**, 349–355.
- 18 G. Becker and F. R. Wurm, *Chem. Soc. Rev.*, 2018, **47**, 7739–7782.
- 19 K. D. Troev, *Polyphosphoesters: Chemistry and Applications*, Elsevier, London, 2012.
- 20 Z. E. Yilmaz, S. Vanslambrouck, S. Cajot, J. Thiry, A. Debuigne, P. Lecomte, C. Jérôme and R. Riva, *RSC Adv.*, 2016, **6**, 42081–42088.
- 21 H. Li, R. Zhou, J. He, M. Zhang, J. Liu, X. Sun and P. Ni, *Bioconjugate Chem.*, 2021, **32**, 2095–2107.
- 22 Y. Iwasaki and K. Akiyoshi, *Biomacromolecules*, 2006, **7**, 1433–1438.
- 23 E. Baeten, S. Vanslambrouck, C. Jérôme, P. Lecomte and T. Junkers, *Eur. Polym. J.*, 2016, **80**, 208–218.
- 24 J. Zou, F. Zhang, S. Zhang, S. F. Pollack, M. Elsabahy, J. Fan and K. L. Wooley, *Adv. Healthcare Mater.*, 2014, **3**, 441–448.
- 25 S. Zhang, J. Zou, M. Elsabahy, A. Karwa, A. Li, D. A. Moore, R. B. Dorshow and K. L. Wooley, *Chem. Sci.*, 2013, **4**, 2122–2126.
- 26 G. Becker and F. R. Wurm, *Tetrahedron*, 2017, **73**, 3536–3540.
- 27 H. J. Lucas, F. W. Mitchell and C. N. Scully, *J. Am. Chem. Soc.*, 1950, **72**, 5491–5497.
- 28 D. Prat, A. Wells, J. Hayler, H. Sneddon, C. R. McElroy, S. Abou-Shehada and P. J. Dunn, *Green Chem.*, 2016, **18**, 288–296.
- 29 M. Sutter, E. Da Silva, N. Duguet, Y. Raoul, E. Métay and M. Lemaire, *Chem. Rev.*, 2015, **115**, 8609–8651.
- 30 R. Gérardy, D. P. Debecker, J. Estager, P. Luis and J. C. M. Monbaliu, *Chem. Rev.*, 2020, **120**, 7219–7347.
- 31 W. H. Faveere, S. Van Praet, B. Vermeeren, K. N. R. Dumoleijn, K. Moonen, E. Taarning and B. F. Sels, *Angew. Chem., Int. Ed.*, 2021, **60**, 12204–12223.
- 32 N. N. Tshibalonza, R. Gérardy, Z. Alsafr, G. Eppe and J.-C. M. Monbaliu, *Green Chem.*, 2018, **20**, 5147–5157.
- 33 R. Morodo, R. Gérardy, G. Petit and J.-C. M. Monbaliu, *Green Chem.*, 2019, **21**, 4422–4433.
- 34 H. Saneyoshi, T. Mashimo, K. Hatano, Y. Ito and H. Abe, *Tetrahedron Lett.*, 2013, **54**, 1080–1083.
- 35 V. Zelikman, J. Pelletier, L. Simhaev, A. Sela, F.-P. Gendron, G. Arguin, H. Senderowitz, J. Sévigny and B. Fischer, *J. Med. Chem.*, 2018, **61**, 3939–3951.
- 36 C. L. Franklin, H. Li and S. F. Martin, *J. Org. Chem.*, 2003, **68**, 7298–7307.
- 37 R. Pongdee, B. Wu and G. A. Sulikowski, *Org. Lett.*, 2001, **3**, 3523–3525.
- 38 A. Granata and D. S. Argyropoulos, *J. Agric. Food Chem.*, 1995, **43**, 1538–1544.
- 39 M. Nagy, M. Kosa, H. Theliander and A. J. Ragauskas, *Green Chem.*, 2010, **12**, 31–34.
- 40 Q. Schmetz, C. Muzyka, T. Berchem and A. Richel, *Green Chem.*, 2020, **22**, 7031–7046.
- 41 A. Zwierzak, *Can. J. Chem.*, 1967, **45**, 2501–2512.
- 42 S. K. Kashani, R. J. Sullivan, M. Andersen and S. G. Newman, *Green Chem.*, 2018, **20**, 1748–1753.
- 43 R. Gérardy, N. Emmanuel, T. Toupay, V.-E. Kassian, N. N. Tshibalonza, M. Schmitz and J.-C. M. Monbaliu, *Eur. J. Org. Chem.*, 2018, **2018**, 2301–2351.
- 44 R. Morodo, P. Bianchi and J.-C. M. Monbaliu, *Eur. J. Org. Chem.*, 2020, **2020**, 5236–5277.
- 45 R. S. Edmundson, *Chem. Ind.*, 1962, 1828–1829.
- 46 Q. Wu, D. Zhou, R. Kang, X. Tang, Q. Yang, X. Song and G. Zhang, *Chem.-Asian J.*, 2014, **9**, 2850–2858.
- 47 T. Biela, S. Penczek, S. Slomkowski and O. Vogl, *Makromol. Chem., Rapid Commun.*, 1982, **3**, 667–671.
- 48 B. Clément, B. Grignard, L. Koole, C. Jérôme and P. Lecomte, *Macromolecules*, 2012, **45**, 4476–4486.
- 49 N. Zaquen, M. Rubens, N. Corrigan, J. Xu, P. B. Zetterlund, C. Boyer and T. Junkers, *Prog. Polym. Sci.*, 2020, **107**, 101256.
- 50 F. Zhang, J. A. Smolen, S. Zhang, R. Li, P. N. Shah, S. Cho, H. Wang, J. E. Raymond, C. L. Cannon and K. L. Wooley, *Nanoscale*, 2015, **7**, 2265–2270.
- 51 Y.-C. Wang, L.-Y. Tang, Y. Li and J. Wang, *Biomacromolecules*, 2009, **10**, 66–73.
- 52 J. Wang, D. Li, W. Tao, Y. Lu, X. Yang and J. Wang, *Biomacromolecules*, 2019, **20**, 1740–1747.
- 53 J.-Z. Du, T.-M. Sun, S.-Q. Weng, X.-S. Chen and J. Wang, *Biomacromolecules*, 2007, **8**, 3375–3381.
- 54 W. Wang, S. Liu, B. Chen, X. Yan, S. Li, X. Ma and X. Yu, *Biomacromolecules*, 2019, **20**, 3672–3683.
- 55 S. Vanslambrouck, R. Riva, B. Ucar, V. Préat, M. Gagliardi, D. G. M. Molin, P. Lecomte and C. Jérôme, *Molecules*, 2021, **26**, 1750.
- 56 R. Riva, S. Lenoir, R. Jérôme and P. Lecomte, *Polymer*, 2005, **46**, 8511–8518.
- 57 R. Riva, S. Schmeits, C. Jérôme, R. Jérôme and P. Lecomte, *Macromolecules*, 2007, **40**, 796–803.
- 58 J. Baran and S. Penczek, *Macromolecules*, 1995, **28**, 5167–5176.

