

Cite this: *Chem. Sci.*, 2023, 14, 3311

All publication charges for this article have been paid for by the Royal Society of Chemistry

# Degradable polyisoprene by radical ring-opening polymerization and application to polymer prodrug nanoparticles†

Maëlle Lages,<sup>a</sup> Théo Pesenti,<sup>a</sup> Chen Zhu,<sup>a</sup> Dao Le,<sup>a</sup> Julie Mougin,<sup>a</sup> Yohann Guillaneuf<sup>b</sup> and Julien Nicolas<sup>a\*</sup>

Radical ring-opening polymerization (rROP) has received renewed attention to incorporate cleavable linkages into the backbones of vinyl polymers, especially from cyclic ketene acetals (CKAs). Among the monomers that hardly copolymerize with CKAs are (1,3)-dienes such as isoprene (I). This is unfortunate since synthetic polyisoprene (PI) and its derivatives are the materials of choice for many applications, in particular as elastomers in the automotive, sport, footwear, and medical industries, but also in nanomedicine. Thionolactones have been recently proposed as a new class of rROP-compatible monomers for insertion of thioester units in the main chain. Herein, we report the synthesis of degradable PI by rROP via the copolymerization of I and dibenzo[c,e]oxepane-5-thione (DOT). Free-radical polymerization as well as two reversible deactivation radical polymerization techniques were successfully used for the synthesis of (well-defined) P(I-co-DOT) copolymers with adjustable molecular weights and DOT contents (2.7–9.7 mol%). Reactivity ratios of  $r_{\text{DOT}} = 4.29$  and  $r_1 = 0.14$  were determined, suggesting preferential incorporation of DOT in comparison to I. The resulting P(I-co-DOT) copolymers were successfully degraded (from –47% to –84% decrease in  $M_n$ ) under basic conditions. As a proof of concept, the P(I-co-DOT) copolymers were formulated into stable and narrowly dispersed nanoparticles, showing similar cytocompatibility on J774.A1 and HUVEC cells compared to their PI counterparts. Furthermore, Gem-P(I-co-DOT) prodrug nanoparticles were synthesized by the “drug-initiated” method and exhibited significant cytotoxicity on A549 cancer cells. P(I-co-DOT) and Gem-P(I-co-DOT) nanoparticles were degraded under basic/oxidative conditions by bleach and under physiological conditions in the presence of cysteine or glutathione.

Received 23rd September 2022  
Accepted 20th February 2023

DOI: 10.1039/d2sc05316k

rsc.li/chemical-science

## Introduction

Degradable vinyl polymers are receiving considerable attention due to their application in the biomedical field (e.g., safety of injected materials and facilitated excretion),<sup>1</sup> but also to address environmental issues (e.g., environmental preservation and the circular plastic economy).<sup>2–4</sup> In this context, there has been a renewed interest in radical ring-opening copolymerization (rROP)<sup>5</sup> as a means of imparting degradability to vinyl polymers through the “cleavable comonomer” approach, particularly with cyclic ketene acetals (CKAs) which are precursors to ester groups in the polymer backbone.<sup>6</sup> Yet, CKAs exhibit important limitations such as: (i) a low reactivity

towards important classes of vinyl monomers (e.g., acrylates, styrenics, methacrylates, etc.), thus requiring at best a large excess of CKAs in the comonomer feed; (ii) an often significant proportion of ring-retained CKAs during polymerization, resulting in non-degradable acetal units in the copolymer backbone and (iii) a very poor stability in protic solvents or in the presence of traces of water, thus severely limiting their use for instance in polymerization in aqueous dispersed media.<sup>5,6</sup>

Among the monomers that hardly copolymerize with CKAs are (1,3)-dienes such as isoprene (I). Synthetic polyisoprene (PI) and its derivatives are the materials of choice for many applications, in particular as elastomers in the automotive, sport, footwear and medical industries,<sup>7,8</sup> with a global PI market size estimated at USD 2.15 billion in 2020.<sup>9</sup> PI-based materials have also been extensively studied for application in colloids,<sup>10–12</sup> materials science/nanotechnology<sup>13–16</sup> and drug delivery owing to their biocompatibility and structural analogy with natural terpenoids.<sup>17–22</sup> Moreover, the inclusion of double bonds in the polymer backbone provides an opportunity for further functionalization via silylation or thiol-ene chemistry.<sup>23,24</sup> Even if PI can be degraded by thermal,<sup>25</sup> chemical (e.g., ozone,<sup>16,26</sup>

<sup>a</sup>Université Paris-Saclay, CNRS, Institut Galien Paris-Saclay, 17 Avenue des Sciences, 91400 Orsay, France. E-mail: julien.nicolas@universite-paris-saclay.fr; Web: <https://twitter.com/julnicolas>; Tel: +33 1 80 00 60 81

<sup>b</sup>Aix-Marseille-Univ., CNRS, Institut de Chimie Radicale, UMR 7273, F-13397 Marseille, France

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d2sc05316k>

chloranil,<sup>27</sup> periodic acid,<sup>28</sup> potassium persulfate,<sup>29</sup> Grubbs catalysts,<sup>30,31</sup> and photo-oxidation<sup>32,33</sup> and biological (enzymatic<sup>34–36</sup> and bacterial<sup>37</sup> pathways (although they cannot occur under healthy physiological conditions), insertion of labile functionalities into PI backbones, allowing their degradation and potential recyclability, has never been a success. DFT calculations indeed recently predicted very unfavourable reactivity ratios for the free-radical polymerization (FRP) of I with main CKAs (e.g.,  $r_{\text{MDO}} = 0.02$  and  $r_{\text{I}} = 9.5$ ;  $r_{\text{BMDO}} = 0.006$  and  $r_{\text{I}} = 157$  at 70 °C),<sup>38</sup> suggesting very little incorporation of CKA units into the PI backbone. Experimental results confirmed the very low insertion of 2-methylene-1,3-dioxepane (MDO) while almost no open 5,6-benzo-2-methylene-1,3-dioxepane (BMDO) units were present. Therefore, an efficient pathway for PI degradation *via* rROP would certainly have major implications in many different areas such as sustainable materials and biomedical applications.

Thionolactones have been recently reported as a new class of cyclic monomers capable of polymerization by rROP and more specifically *via* a thiocarbonyl-addition-ring opening (TARO) mechanism, enabling insertion of labile thioester groups into the polymer backbone.<sup>5</sup> In particular, it has been shown that dibenzo[*c,e*]oxepane-5-thione (DOT) exhibited superior reactivity towards several vinyl monomers (e.g., acrylates,<sup>39,40</sup> acrylamides,<sup>41</sup> acrylonitrile,<sup>39</sup> styrene (S)<sup>42</sup> and maleimide<sup>43</sup>), extensive stability in protic solvents and in aqueous media, and quantitative ring opening.<sup>39–41,43,44</sup> Furthermore, it allows to diversify degradation pathways; from hydrolysis<sup>41,42,44,45</sup> to aminolysis<sup>39–41</sup> and thiolysis.<sup>43</sup> Very recently, DOT was copolymerized with *n*BA or S by either conventional emulsion polymerization<sup>44</sup> and polymerization-induced self-assembly,<sup>46</sup> while DOT was also used to impart chemical recyclability to polystyrene.<sup>47</sup> Moreover, it has been shown that DOT-containing copolymers can be degraded in the presence of physiologically relevant concentrations of cysteine or glutathione,<sup>48</sup> which allows DOT-containing copolymers to be considered in nanomedicine.

Herein, we report the successful rROP of I and DOT by both FRP and two reversible deactivation radical polymerization techniques: nitroxide-mediated polymerization<sup>49</sup> (NMP) and

reversible addition–fragmentation chain-transfer<sup>50</sup> (RAFT) polymerization (Fig. 1). Owing to favourable reactivity ratios, we demonstrated the formation of thioester-containing copolymers that can be readily degraded under basic conditions. To show the versatility and broad applicability of this copolymerization system, we also reported the formulation of the resulting copolymers into nanoparticles as well as the synthesis of degradable polymer prodrug nanoparticles for anticancer therapy.

## Experimental part

### Materials

Isoprene (I, 99%), dicumyl peroxide (DCP, 98%), potassium hydroxide (KOH, 90%), dioxane (anhydrous, 99.8%) and isopropylamine ( $\geq 99\%$ ) were purchased from Sigma-Aldrich and used as received. (*N-tert*-Butyl-*N*-(1-diethoxyphosphoryl-2,2-dimethylpropyl)aminoxy)-propionic acid alkoxyamine (Bloc-Builder MA, BB) was kindly supplied by Arkema. 1,5,7-triazabicyclo[4.4.0]dec-5-ene (TBD,  $>98\%$ ) was purchased from TCI. Deuterated chloroform ( $\text{CDCl}_3$ ) and tetrahydrofuran ( $d_8$ -THF) were purchased from Eurisotop. Tetrahydrofuran (THF, HPLC grade), methanol (MeOH, HPLC) and chloroform ( $\text{CHCl}_3$ , HPLC grade) were obtained from VWR Chemicals. Hydrochloric acid (HCl, 37%) was supplied by Carlo-Erba. 2.5% Active chlorine bleach solution was purchased from Leroy Merlin (France). Dibenzoc[*c,e*]oxepane-5-thione (DOT),<sup>39</sup> gemcitabine-AMA-SG1 (Gem-AMA-SG1)<sup>19</sup> and 2-ethylsulfanylthiocarbonylsulfanylpropionic acid ethyl ester (ETSPE)<sup>51</sup> were prepared as reported elsewhere. Pressure tubes (Ace Glass 8648\_164, 15 mL-capacity, fitted with a plunger valve and thermowell) were purchased from Sigma-Aldrich.

### Analytical methods

**Nuclear magnetic resonance (NMR) spectroscopy.** NMR spectroscopy was performed in 5 mm diameter tubes in  $\text{CDCl}_3$  or  $d_8$ -THF at 25 °C.  $^1\text{H}$  NMR spectroscopy was performed on a Bruker Avance 300 spectrometer at 300 MHz with 350 scans. The chemical shift scale was calibrated based on the internal solvent signals ( $\delta = 7.26$  ppm) for  $\text{CDCl}_3$  and the TMS internal



Fig. 1 Synthesis of degradable vinyl polymer nanoparticles *via* radical ring opening copolymerization (rROP) of dibenzoc[*c,e*]oxepane-5-thione (DOT) and isoprene (I) in solution followed by nanoprecipitation.



standard ( $\delta = 0$  ppm) for  $d_8$ -THF.  $^{31}\text{P}$  NMR spectroscopy was conducted in  $\text{CDCl}_3$  on a Bruker Avance 400 spectrometer operating at a frequency of 161.9 MHz with a 5 mm gradient BBFO probe (31P-109Ag/1H). The NMR spectra were recorded with proton decoupling (1D sequence with inverse gated  $^1\text{H}$ -decoupling) with a spectral width of 32 467 Hz, an acquisition time of 1 s and a relaxation delay of 34 s. The chemical shift scale was calibrated based on added diethyl phosphite at  $\delta = 7.1$  ppm.

**Size exclusion chromatography (SEC).** For kinetic monitoring, SEC on dry extracts was performed on a Tosoh EcoSEC HLC-8320 GPC with two columns from Agilent (PL-gel MIXED-D  $300 \times 7.5$  mm, bead diameter 5  $\mu\text{m}$ ; linear part 400 to 400 000  $\text{g mol}^{-1}$ ). Analyses were performed at 35  $^\circ\text{C}$  in chloroform (HPLC grade) at a flowrate of 1  $\text{mL min}^{-1}$  using toluene as a flowrate marker. A conventional calibration curve was constructed based on polystyrene (PS) standards (peak molar masses:  $M_p = 575$ –126 500  $\text{g mol}^{-1}$ ) from Polymer Laboratories. A PI calibration curve was constructed by converting the PS standard peak molecular weights,  $M_{\text{PS}}$ , to PI molecular weights,  $M_{\text{PI}}$ , using MarkHouwink–Sakurada (MHS) constants determined for both polymers in  $\text{CCl}_4$  at 25  $^\circ\text{C}$ . For PI, the MHS constants used were  $K_{\text{PI}} = 2.44 \times 10^{-4}$  and  $\alpha_{\text{PI}} = 0.712$ . For PS,  $K_{\text{PS}} = 7.1 \times 10^{-4}$  and  $\alpha_{\text{PS}} = 0.54$  ( $M_w < 16\,700$   $\text{g mol}^{-1}$ ) or  $K_{\text{PS}} = 1.44 \times 10^{-4}$  and  $\alpha_{\text{PS}} = 0.713$  ( $M_w \geq 16\,700$   $\text{g mol}^{-1}$ ).<sup>52,53</sup> This technique allowed  $M_n$  (the number-average molar mass),  $M_w$  (the weight-average molar mass), and  $M_w/M_n$  (the dispersity,  $D$ ) to be determined.

For purified and degraded samples, SEC was performed at 30  $^\circ\text{C}$  with two columns from Agilent (PL-gel MIXED-D;  $300 \times 7.5$  mm; bead diameter, 5  $\mu\text{m}$ ; linear part, 400–400 000  $\text{g mol}^{-1}$ ) and a differential refractive index detector (Spectrasystem RI-150 from Thermo Electron Corp.), using  $\text{CHCl}_3$  as an eluent at a flowrate of 1  $\text{mL min}^{-1}$  and toluene as a flowrate marker. A conventional calibration curve was constructed based on PS standards (peak molar masses:  $M_p = 575$ –126 500  $\text{g mol}^{-1}$ ) from

Polymer Laboratories. A PI calibration curve was constructed in the same manner as described above. SEC of degraded copolymers was performed in the presence of 0.1% (v/v) of trifluoroacetic acid (TFA, 99%) in  $\text{CHCl}_3$  (in both the mobile phase and the sample) to avoid the formation of aggregates and/or interaction with the columns and sulfhydryl or hydroxyl chain ends.

**Dynamic light scattering (DLS) and zeta potential ( $\zeta$ ).** Intensity-averaged nanoparticle diameter ( $D_z$ ), polydispersity index (PDI) and zeta potential were measured by DLS with a nano ZS from Malvern (173 $^\circ$  scattering angle) at a temperature of 25  $^\circ\text{C}$ . Nanoparticle dispersions were diluted in MilliQ water (dilution 1/10 (v/v)) prior to analysis. The surface charge of the nanoparticles was investigated by  $\zeta$ -potential (mV) measurement at 25  $^\circ\text{C}$  after dilution (1/10 v/v) with 1 mM NaCl, using the Smoluchowski equation.

**Cryogenic transmission electron microscopy (cryo-TEM).** The morphologies of P(I-co-DOT) and PI nanoparticles were observed by cryo-TEM. 5  $\mu\text{L}$  of the nanoparticle suspension was diluted at 1.2  $\text{mg mL}^{-1}$  just prior analysis and deposited onto a Lacey Formvar/carbon 300 mesh copper grid (Ted Pella). The excess was manually blotted with a filter paper and the residual thin film was immediately frozen by plunging into liquid ethane and cooled down at liquid nitrogen temperature using a Leica EM-CPC cryopluger. Observation was performed using a JEOL 2100HC microscope (JEOL Europe) or a JEOL 2200FS field emission microscope (JEOL USA) operating under an acceleration voltage of 200 kV in zero-loss mode (slit was 20 eV). High-magnification images ( $2\text{k} \times 2\text{k}$  pixels) were recorded using a CCD camera (Gatan, Inc.) with Digital Micrograph software.

## Synthetic procedures

**Synthesis of poly(isoprene-co-dibenzo[*c,e*]oxepane-5-thione) (P(I-co-DOT), F0–F2) by free-radical polymerization in solution.** In a typical procedure (F1, Table 1), I (4 mL, 2.73 g, 40 mmol), DOT (0.091 g, 0.404 mmol,  $f_{\text{DOT},0} = 0.01$ ), DCP (0.025 g,

Table 1 Experimental conditions for the synthesis of P(I-co-DOT) copolymers by rROP and their macromolecular characteristics

|      | Entry           | $f_{\text{DOT},0}$ (mol%) | $t$ (h) | $\text{DP}_{n,\text{th}}$ | Conv. I <sup>a</sup> (wt%) | $M_{n,\text{th}}^b$ ( $\text{g mol}^{-1}$ ) | $M_n^c$ ( $\text{g mol}^{-1}$ ) | $M_w^c$ ( $\text{g mol}^{-1}$ ) | $D^c$ | $F_{\text{DOT}}^d$ (mol%) |
|------|-----------------|---------------------------|---------|---------------------------|----------------------------|---|---------------------------------|---------------------------------|-------|---------------------------|
| FRP  | F0              | 0                         | 30      | —                         | 37                         | —   | 7200                            | 11 000                          | 1.53  | 0                         |
|      | F1              | 1                         | 30      | —                         | 38                         | —   | 10 000                          | 16 400                          | 1.64  | 2.7                       |
|      | F2              | 2                         | 30      | —                         | 40                         | —   | 10 400                          | 17 000                          | 1.63  | 4.6                       |
| NMP  | N0              | 0                         | 48      | 100                       | 33                         | 2700  | 2400                            | 2900                            | 1.19  | 0                         |
|      | N1              | 1                         | 48      | 100                       | 37                         | 2900  | 3100                            | 3900                            | 1.25  | 2.7                       |
|      | N2              | 2                         | 48      | 100                       | 40                         | 3100  | 3300                            | 4400                            | 1.32  | 4.4                       |
|      | N3              | 2                         | 48      | 215                       | 36                         | 5700  | 5300                            | 8200                            | 1.54  | 5.3                       |
|      | N4              | 2                         | 48      | 435                       | 21                         | 6600  | 4800                            | 7100                            | 1.48  | 9.7                       |
|      | N5 <sup>e</sup> | 2                         | 16      | 100                       | 17                         | 1800  | 1800                            | 2200                            | 1.29  | 6.6                       |
|      | N6              | 3                         | 20      | 500                       | 18                         | 6500  | 7900                            | 8400                            | 1.07  | 6.1                       |
|      | N7 <sup>e</sup> | 3                         | 20      | 500                       | 22                         | 8100  | 9300                            | 10 700                          | 1.15  | 6.9                       |
| RAFT | R0              | 0                         | 24      | 500                       | 11                         | 3900  | 3400                            | 3900                            | 1.16  | 0                         |
|      | R1              | 2                         | 24      | 500                       | 10                         | 3800  | 4500                            | 6100                            | 1.36  | 5.5                       |
|      | R2              | 2                         | 72      | 600                       | 55                         | 22 900                                      | 11 000                          | 18 200                          | 1.67  | 3.9                       |

<sup>a</sup> Determined by gravimetry. <sup>b</sup> Determined according to:  $M_{n,\text{th}} = M_{n,\text{BB}} + \text{DP}_{n,\text{th}} \times \text{conv. MW}_I$ . <sup>c</sup> Determined by SEC on precipitated samples.

<sup>d</sup> Determined by  $^1\text{H}$  NMR in  $d_8$ -THF of precipitated samples by integrating the 8H (Ar) of open DOT at 6.7–8 ppm and protons from PI (1H for the (1,4) conformation at 5–5.5 ppm, 1H for the (1,2) conformation at 5.5–5.9 ppm and 4.4–5 ppm combining 2H of (1,2) and 2H of (3,4) conformations). For F1–F2 and R1–R2, the aromatic signal of DCP at 7.0–7.5 ppm was subtracted from the aromatic signal of DOT according to the DCP/PI ratio calculated on the F0 and R0 spectrum respectively. <sup>e</sup> The Gem-AMA-SG1 alkoxyamine was used instead of the BB alkoxyamine.



0.093 mmol) and dioxane (5 mL) were introduced in a pressure tube fitted with a plunger valve and thermowell. The solution was subjected to three cycles of freeze-thaw degassing, and then backfilled with argon. The tube was placed in an oil bath at 115 °C for 30 h at 300 rpm and then cooled down at room temperature. The I conversion was determined by gravimetry. The copolymer was purified by three successive precipitations in MeOH to remove unreacted DOT and dried under vacuum at room temperature. The resulting copolymer was analysed by  $^1\text{H}$  NMR in  $d_8$ -THF and by SEC in  $\text{CHCl}_3$ .

The same procedure was adapted by varying the initial feed ratio of DOT ( $f_{\text{DOT},0}$ ) as follows: **F0** [I (4 mL, 2.73 g, 40 mmol) and DCP (0.025 g, 0.093 mmol) in 5 mL of dioxane] and **F2** [I (4 mL, 2.73 g, 40 mmol), DOT (0.185 g, 0.816 mmol,  $f_{\text{DOT},0} = 0.02$ ) and DCP (0.025 g, 0.093 mmol) in 5 mL of dioxane].

The determination of the reactivity ratios was performed according to the following protocol. In a typical procedure, I (0.08 mL, 54 mg, 0.79 mmol), DOT (179 mg, 0.79 mmol,  $f_{\text{DOT},0} = 0.5$ ), DCP (0.9 mg, 0.0036 mmol), anhydrous DMSO (61 mg, 0.79 mmol) used as the internal reference, and dioxane (7 mL) were introduced in a pressure tube fitted with a plunger valve and thermowell. The solution was subjected to three cycles of freeze-thaw degassing, and then backfilled with argon. The tube was placed in an oil bath at 115 °C for 1 h 30 and then cooled down at room temperature. The monomer conversions (kept at a low level; 34% maximum) were determined by  $^1\text{H}$  NMR in  $\text{CDCl}_3$  by integrating the signal at 6.4 ppm for I (dd, 1H), the signal at 8.2 ppm for DOT (d, 1H), and the signal of the internal reference at 2.6 ppm ( $d_6$ -DMSO, s, 6H). The copolymer was then purified by two successive precipitations in MeOH to remove unreacted DOT and dried under vacuum at room temperature. The resulting copolymer was analysed by  $^1\text{H}$  NMR in  $d_8$ -THF to determine  $F_{\text{DOT}}$ . Different  $f_{\text{DOT},0}$  values were considered (0.1, 0.2, 0.3, 0.4, 0.5, 0.6, 0.7 and 0.9) by keeping a total monomer concentration of 1.13 M (Table S1†). Evolution of  $F_{\text{DOT}}$  with  $f_{\text{DOT},0}$  was then plotted, and the reactivity ratios were determined using nonlinear least-squares (NLLS) analysis performed with the CONTOUR software.<sup>54</sup>

**Synthesis of P(I-co-DOT) (N0–N4 and N6) by NMP in solution.** The experimental protocol was adapted from a previous report.<sup>55</sup> In a typical procedure (**N2**, Table 1), a mixture of I (6 mL, 4.086 g, 60 mmol, 100 eq.), DOT (0.277 g, 1.22 mmol, 2 eq.,  $f_{\text{DOT},0} = 0.02$ ), BB (0.229 g, 0.60 mmol, 1 eq.) and dioxane (6 mL) was divided into 6 pressure tubes fitted with a plunger valve and thermowell. Each solution was subjected to three cycles of freeze-thaw degassing, and then backfilled with argon. The tubes were placed in an oil bath at 115 °C for 1, 3, 7, 16, 30 and 48 h, and then cooled down at room temperature. The I conversion was determined by gravimetry and the dry samples were analysed by SEC. The copolymer obtained at 48 h was purified by three successive precipitations in MeOH to remove unreacted DOT, dried under vacuum at room temperature and analysed by  $^1\text{H}$  NMR in  $d_8$ -THF and by SEC in  $\text{CHCl}_3$ .

The same procedure was then adapted by varying  $f_{\text{DOT},0}$  and the targeted number-average degree of polymerization of I ( $\text{DP}_{n,\text{th}}$ ) as follows: **N0** [I (6 mL, 4.086 g, 60 mmol, 100 eq.), BB (0.229 g, 0.60 mmol, 1 eq.) and dioxane (6 mL)]; **N1** [I (6 mL,

4.086 g, 60 mmol, 100 eq.), DOT (0.137 g, 0.61 mmol, 1.01 eq.,  $f_{\text{DOT},0} = 0.01$ ), BB (0.229 g, 0.60 mmol, 1 eq.) and dioxane (6 mL)]; **N3** [I (6 mL, 4.086 g, 60 mmol, 215 eq.), DOT (0.277 g, 1.22 mmol, 4.4 eq.,  $f_{\text{DOT},0} = 0.02$ ), BB (0.106 g, 0.279 mmol, 1 eq.) and dioxane (6 mL)]; **N4** [I (6 mL, 4.086 g, 60 mmol, 435 eq.), DOT (0.277 g, 1.22 mmol, 8.88 eq.,  $f_{\text{DOT},0} = 0.02$ ), BB (0.053 g, 0.138 mmol, 1 eq.) and dioxane (6 mL)] and **N6** [I (1.50 mL, 1.02 g, 15 mmol, 500 eq.), DOT (0.105 g, 0.46 mmol, 15.5 eq.,  $f_{\text{DOT},0} = 0.03$ ), BB (0.0114 g, 0.03 mmol, 1 eq.) and dioxane (1.50 mL)]. For **N3'** [I (6 mL, 4.086 g, 60 mmol, 215 eq.), DOT (0.277 g, 1.22 mmol, 4.4 eq.,  $f_{\text{DOT},0} = 0.02$ ), BB (0.106 g, 0.279 mmol, 1 eq.), dioxane (6 mL) and anhydrous DMSO (0.056 mL, 0.123 g, 1.5 mmol)], anhydrous DMSO was added as an internal reference to determine the DOT conversion by  $^1\text{H}$  NMR in  $\text{CDCl}_3$ .

**Synthesis of P(I-co-DOT) (R0–R2) by RAFT polymerization in solution.** The experimental protocol was adapted from a previous report.<sup>56</sup> In a typical procedure (**R1**, Table 1), I (2.86 g, 42 mmol, 500 eq.), DOT (0.193 g, 0.85 mmol, 10.2 eq.,  $f_{\text{DOT},0} = 0.02$ ), ETSPE (0.02 g, 0.084 mmol, 1 eq.) and DCP (0.0045 g, 0.016 mmol, 0.2 eq.) were introduced in a pressure tube fitted with a plunger valve and thermowell. The solution was subjected to three cycles of freeze-thaw degassing, and then backfilled with argon. The tube was placed in an oil bath at 115 °C for 24 h and then cooled down at room temperature. The I conversion was determined by gravimetry. The copolymer was purified by three successive precipitations in MeOH to remove unreacted DOT and dried under vacuum at room temperature. The resulting copolymer was analysed by  $^1\text{H}$  NMR in  $d_8$ -THF and by SEC in  $\text{CHCl}_3$ .

The same procedure was then adapted by varying  $f_{\text{DOT},0}$ ,  $\text{DP}_{n,\text{th}}$  and reaction time as follows: **R0** [I (2.86 g, 42 mmol, 500 eq.), ETSPE (0.02 g, 0.084 mmol, 1 eq.) and DCP (0.0045 g, 0.016 mmol, 0.2 eq.)] and **R2** [I (1.47 g, 21.6 mmol, 600 eq.), DOT (0.099 g, 0.441 mmol, 12.25 eq.,  $f_{\text{DOT},0} = 0.02$ ), ETSPE (0.008 g, 0.036 mmol, 1 eq.) and DCP (0.0019 g, 0.007 mmol, 0.2 eq.) for 72 h].

**Synthesis of gemcitabine-poly(isoprene-co-dibenzo[c,e]oxepane-5-thione) (Gem-P(I-co-DOT), N5 and N7) by NMP in solution.** The experimental protocol was adapted from a previous report.<sup>19</sup> In a typical procedure (**N5**, Table 1), a solution of I (0.85 mL, 0.578 g, 85 mmol, 100 eq.), DOT (0.039 g, 0.17 mmol, 2.04 eq.,  $f_{\text{DOT},0} = 0.02$ ), and Gem-AMA-SG1 (0.052 g, 0.085 mmol, 1 eq.) was added into 0.85 mL of dioxane and introduced in a pressure tube fitted with a plunger valve and thermowell. The solution was subjected to three cycles of freeze-thaw degassing, and then backfilled with argon. The tube was placed in an oil bath at 115 °C for 16 h and then cooled down at room temperature. The I conversion was determined by gravimetry. The copolymer was purified by three successive precipitations in MeOH to remove unreacted DOT and dried under vacuum at room temperature. The resulting copolymer was analysed by  $^1\text{H}$  NMR in  $\text{CDCl}_3$  and by SEC in  $\text{CHCl}_3$ . The drug content in gemcitabine (Gem) was determined according to:  $\% \text{Gem} = \text{MW}_{\text{Gem}} / M_{n,\text{Gem-PI}}$  with  $\text{MW}_{\text{Gem}} = 263.2 \text{ g mol}^{-1}$ .

The same procedure was then adapted by targeting  $\text{DP}_{n,\text{th}} = 500$  as follows: **N7** [I (1.30 mL, 0.885 g, 13 mmol, 500 eq.)], DOT





(0.091 g, 0.40 mmol, 15 eq.,  $f_{\text{DOT},0} = 0.03$ ), Gem-AMA-SG1 (0.016 g, 0.026 mmol, 1 eq.) and 1.30 mL of dioxane.

### Nanoparticle preparation

Nanoparticles were prepared by the nanoprecipitation technique. Briefly, 10 mg of polymer **N0–N5** (Table 1) were dissolved in 2 mL of THF and added dropwise to 3 mL MilliQ water under stirring. THF was removed under vacuum to reach a final concentration of 3.3 mg mL<sup>−1</sup>. For **N6**, 20 mg of polymer (Table 1) were dissolved in 8 mL of dioxane and added dropwise to 12 mL MilliQ water under stirring. Dioxane was removed under vacuum to reach a final concentration of 1.7 mg mL<sup>−1</sup>. For **N7**, volumes of dioxane and water were divided by two to reach a final concentration of 3.3 mg mL<sup>−1</sup>. Intensity-average diameter ( $D_z$ ) and zeta potential measurements were carried out in triplicate.

### Degradation procedures

**Degradation of the copolymers by KOH.** 10 mg of copolymer were solubilized in 0.5 mL of DCM prior to adding 0.5 mL of a 5 wt% aqueous solution of KOH in MeOH. The solution was stirred for 16 h at room temperature and dried under vacuum. HCl solution (1 M) was then added to the organic solution followed by extraction of the aqueous layer (three times). Solvent was removed under reduced pressure. The degradation products were analysed by SEC.

**Degradation of the copolymers by isopropylamine.** 10 mg of copolymer were solubilized in 1 mL of THF prior to adding 0.4 mL of isopropylamine. The solution was stirred for 16 h at room temperature. THF and isopropylamine were removed under vacuum. The degradation products were analysed by SEC.

**Degradation of the copolymers by TBD.** 10 mg of copolymer were solubilized in 0.5 mL of THF prior to adding 0.5 mL of a 5 wt% solution of TBD in THF. The solution was stirred for 16 h at room temperature and dried under vacuum. The organic solution was washed three times with HCl solution (1 M) to remove TBD and then dried under vacuum until constant weight. The degradation products were analysed by SEC.

**Degradation of the nanoparticles by bleach solution.** In a typical procedure, 1 mL of nanoparticle suspensions **N6** (Table 4, 1.7 mg mL<sup>−1</sup>) and **N7** (Table 4, 3.3 mg mL<sup>−1</sup>) was added to 2 mL of 2.5% active chlorine bleach solution. The solution was stirred for 12 days at room temperature. The product was extracted with 2 mL CHCl<sub>3</sub>. The organic solvent was dried with MgSO<sub>4</sub>, filtered, and evaporated under reduced pressure. The degradation products were analysed by SEC.

**Degradation of the nanoparticles by cysteine and glutathione (GSH).** Degradant solutions were prepared by dissolving cysteine or glutathione and tris(carboxyethyl phosphine) TCEP with a 10/1 thiol/TCEP molar ratio in PBS pH 7.4. The pH of solutions was adjusted by adding 0.5 M NaOH solution to 7.4 for cysteine solution and 7.2 for glutathione solution. PBS was added to adjust the concentration of thiol to 20 mM. 1 mL of **N6** (Table 4, 1.7 mg mL<sup>−1</sup>) or 0.5 mL of **N7** (Table 4, 3.3 mg mL<sup>−1</sup>) and 0.5 mL water were added to 1 mL thiol 20 mM solution resulting in 10 mM thiol solutions. The solutions were stirred at

37 °C for 15 days. The mixture was extracted with 2 mL CHCl<sub>3</sub>. The organic solvent was dried with MgSO<sub>4</sub>, filtered, and evaporated under reduced pressure. The degradation products were analysed by SEC.

### In vitro cytotoxicity

**Cell lines and cell culture.** Human umbilical vein endothelial cells (HUVEC), mouse monocyte macrophage cells (J774.A1) and lung cancer cells (A549) were obtained from the American Type Culture Collection (Molsheim, France) and maintained as recommended. HUVEC cells were grown in Dulbecco's modified Eagle medium (DMEM) with high glucose supplemented with 10% fetal bovine serum (FBS), penicillin (100 U mL<sup>−1</sup>) and streptomycin (100 U mL<sup>−1</sup>). J774.A1 cells were grown in Roswell Park Memorial Institute medium (RPMI) 1640 supplemented with 10% FBS, penicillin (100 U mL<sup>−1</sup>) and streptomycin (100 U mL<sup>−1</sup>). A549 cells were cultured in Roswell Park Memorial Institute medium (RPMI) and supplemented with 10% FBS and penicillin (100 U mL<sup>−1</sup>). Cells were maintained in a humid atmosphere at 37 °C with 5% CO<sub>2</sub>.

**Cytotoxicity assay (MTT).** MTT [3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl tetrazolium bromide] was used to test the cytotoxicity of the different nanoparticles *via* cell viability measurement. Briefly, cells (5 × 10<sup>3</sup> per well) were seeded in 96-well plates. After a 24 h incubation, cells were exposed to a series of increasing concentrations of nanoparticles. After 72 h, 20 µL of MTT solution (5 mg mL<sup>−1</sup> in PBS) was added to the medium in each well. The plates were incubated for 1 h at 37 °C and the medium was removed after centrifugation. 200 µL of DMSO were then added to each well to dissolve the precipitates. Absorbance was measured at 570 nm using a plate reader (Metertech Σ 960, Fisher Bioblock, Illkirch, France). The percentage of surviving cells was calculated as the absorbance ratio of treated to untreated cells. All experiments were set up in sextuplicate to determine means and SDs.

**Cytotoxicity of the degradation products.** The degraded copolymers by TBD were dissolved in CHCl<sub>3</sub> and filtered 3 times to remove the residual salts. They were dried under vacuum until constant weight. Due to the insolubility of the degradation products in water, they were dispersed in aqueous media as follows: For J774.A1 cells, the degradation products were dissolved in DMSO and added drop by drop to MilliQ water under stirring to reach a final concentration of 5 v% of DMSO. A control experiment with a 5 v% solution of DMSO in MilliQ water was performed on cells and no cytotoxicity was shown in the concentration range considered. For HUVEC cells, the degradation products were dissolved in THF and added dropwise to MilliQ water under stirring. THF was removed under vacuum.

## Results and discussion

### Synthesis of DOT-containing polyisoprene

Copolymerization of DOT with I was first investigated by free-radical copolymerization in dioxane at 115 °C (Fig. 2). Initial molar fractions in DOT ( $f_{\text{DOT},0}$ ) up to 2 mol% were considered





Fig. 2 Synthesis of poly(isoprene-co-dibenzo[c,e]oxepane-5-thione) (P(I-co-DOT)) copolymers by radical ring-opening copolymerization (rROP) of DOT and I in dioxane at 115 °C by: (a) free-radical polymerization (FRP); (b) nitroxide-mediated polymerization (NMP) and (c) reversible addition-fragmentation chain-transfer (RAFT) polymerization.

due its low solubility in dioxane. After 30 h of copolymerization, experiments **F1** ( $f_{\text{DOT},0} = 0.01$ ) and **F2** ( $f_{\text{DOT},0} = 0.02$ ) resulted in  $\sim 40\%$  conversion of I and  $M_n$ s of  $\sim 10\,000\text{ g mol}^{-1}$  (Table 1).

Homopolymerization of I under identical polymerization conditions gave similar results (**F0**, Table 1), suggesting no retardation effect of DOT with I, in contrast to other vinyl monomers.<sup>39,44</sup>  $^1\text{H}$  NMR spectroscopy was then performed on the purified (co)polymers, showing successful insertion of DOT units to an extent of  $F_{\text{DOT}} = 2.7$  and 4.6 mol%, for experiments **F1** and **F2**, respectively (Fig. 3a and Table 1). Importantly, CKAs required a much higher initial feed ratio of 75 mol% to only incorporate 7 mol% of MDO in the PI backbone.<sup>38</sup> This was supported by theoretical DFT calculations that reported  $r_{\text{MDO}} = 0.02$  and  $r_{\text{I}} = 9.5$  (or  $r_{\text{BMDO}} = 0.006$  and  $r_{\text{I}} = 157$ ) at 70 °C for copolymerization between MDO (or BMDO) and I.<sup>38</sup> This therefore suggested a superior reactivity of DOT towards I compared to CKAs. Reactivity ratios of this copolymerization system were estimated using a NLLS method performed with the CONTOUR software,<sup>54</sup> giving  $r_{\text{DOT}} = 4.29$  and  $r_{\text{I}} = 0.14$  (the 95% joint confidence interval is given in Fig. S1†). This suggested a significant preferential incorporation of DOT over I and a high heterogeneity of the DOT fraction between chains if high conversion is reached (the latter expected from a FRP process). Similar trends, which can be explained by the structural similarity between DOT and RAFT agents, have been previously shown for the copolymerization of DOT with other vinyl monomers, such as *N,N*-dimethylacrylamide ( $r_{\text{DOT}} = 1.89$  and  $r_{\text{DMAM}} = 0.34$ ), *N*-2,3,4,5,6-pentafluorophenylmaleimide ( $r_{\text{DOT}} = 0.198$  and  $r_{\text{PFPMI}} = 0.0078$ ) and *N*-phenylmaleimide ( $r_{\text{DOT}} = 0.348$  and  $r_{\text{PhMI}} = 0.0136$ ).<sup>41</sup>

Copolymerizations *via* NMP and RAFT were then investigated to yield well-defined P(I-co-DOT) copolymers. NMP of I using BB as a SG1-based alkoxyamine was performed with  $f_{\text{DOT},0} = 1$  and 2 mol% in dioxane and with  $\text{DP}_{n,\text{th}} = 100$  (**N0–N2**, Table 1). 30–40% conversions of I were obtained after 48 h, with  $M_n$  values in the 2400–3300  $\text{g mol}^{-1}$  range and low dispersities ( $D = 1.19$ – $1.32$ ), indicating a good control even in the presence of DOT. Interestingly, no noticeable retardation effect from DOT was obtained, despite the increasing initial molar fraction of DOT in the comonomer feed<sup>39,44</sup> (Fig. 3c). At  $f_{\text{DOT},0} = 2$  mol%, different  $M_{n,\text{th}}$  were targeted by varying the  $\text{DP}_{n,\text{th}}$  from 100 to 435 (**N2–N4**, Fig. 3c–e, Table 1). Interestingly, the

copolymerizations exhibited all the characteristics of a controlled system, as confirmed by: (i) the linear evolutions of the  $M_n$  values with I conversion, in pretty good agreement with the theoretical ones (Fig. 3d) and (ii) the low dispersities ( $D < 1.2$ ) throughout the reactions (Fig. 3d and S2†). The final copolymers were then purified and analysed by  $^1\text{H}$  NMR (Table 1, Fig. S3†), which successfully demonstrated insertion of open DOT units into the PI backbone at 4.4 and 5.3 mol% for **N2** and **N3**, respectively. The final amount of DOT in the copolymer was correlated to: (i) the initial stoichiometry, as the higher  $f_{\text{DOT},0}$ , the higher the  $F_{\text{DOT}}$  (see **N1** and **N2**, Table 1) and (ii) the I conversion, as the lower the I conversion, the higher the  $F_{\text{DOT}}$ , in agreement with the reactivity ratios and thus the expected composition drift of P(I-co-DOT) copolymers (see **N2–N4**, Table 1). In order to gain further insight into this copolymerization system, the evolutions of the individual conversions of DOT and I, as well as  $F_{\text{DOT}}$  during the copolymerization were studied for **N3**. Note that **N3** was repeated in the presence of DMSO as the internal reference and noted as **N3'** (Fig. 3e and S4†). It appeared that DOT was indeed consumed faster than I, leading to 85% DOT conversion and 37% I conversion after 48 h. As expected, the evolution of  $F_{\text{DOT}}$  progressively decreased from 15.5 to 4.3 mol% after 48 h in agreement with a gradient composition.

**N3'** and **N0** were also analysed by  $^{31}\text{P}$  NMR to perform in-depth chain-end characterization and give crucial insights into the nature of the copolymer terminal sequences and the living chain fractions.<sup>57</sup> Interestingly, the phosphorus signals (24.4–24.6 ppm) from **N3'** and **N0** had the same shape and were characteristic of I-SG1 terminal sequences (Fig. S5†),<sup>53</sup> thus ruling out accumulation of DOT as the last monomer unit and supporting its faster incorporation in the copolymer compared to I. By integrating these phosphorus signals and that of diethyl phosphite (7.1 ppm) used as an internal standard, the livingness of both (co)polymers was estimated to be  $\sim 50\%$ .

P(I-co-DOT) copolymers were also synthesized by RAFT polymerization using ETSPE as a RAFT agent<sup>56</sup> and DCP as a source of radicals. After 24 h with  $f_{\text{DOT},0} = 0.02$ , 10% of I conversion was achieved giving  $M_n = 4500\text{ g mol}^{-1}$  and  $D = 1.36$  (**R1**, Table 1). The absence of DOT under the same conditions (**R0**, Table 1) led to very similar macromolecular characteristics, despite obtaining a lower dispersity ( $D = 1.16$ ). To obtain higher





**Fig. 3** (a)  $^1\text{H}$  NMR spectrum (300 MHz,  $d_8$ -THF) in the 0–8 ppm region of P(I-co-DOT) copolymers F0, F1 and F2 (Table 1). (b) Evolution of molar DOT content in copolymers with molar DOT in the monomer feed with nonlinear least-squares fitted curves with  $r_{\text{DOT}} = 4.29$  and  $r_1 = 0.14$ . (c) Evolution of I conversion vs. time as a function of  $f_{\text{DOT},0}$  (N0, N1 and N2, Table 1). (d) Evolution of  $M_n$  and  $\bar{D}$  vs. I conversion as a function of  $DP_{n,\text{th}}$  (N2, N3 and N4, Table 1). (e) Evolution of I conversion, DOT conversion and average molar fraction of DOT ( $F_{\text{DOT}}$ ) in the copolymer vs. time (N3').





Table 2 Macromolecular characteristics of P(l-co-DOT) copolymers obtained by rROP before and after degradation

| Entry           | Degradation by KOH <sup>d</sup>   |                                   |                              |  | Degradation by isopropylamine <sup>e</sup>              |   |   |   | Degradation by TBD <sup>f</sup>                         |   |
|-----------------|-----------------------------------|-----------------------------------|------------------------------|--|---|---|---|---|---|---|
|                 | $M_n^a$<br>(g mol <sup>-1</sup> ) | $M_w^a$<br>(g mol <sup>-1</sup> ) | $F_{\text{DOT}}^b$<br>(mol%) | $M_{n,\text{deg},\text{th}}^c$<br>(g mol <sup>-1</sup> ) | $M_{n,\text{deg},\text{KOH}}$<br>(g mol <sup>-1</sup> ) | $M_{w,\text{deg},\text{KOH}}$<br>(g mol <sup>-1</sup> ) | $M_{n,\text{deg},\text{isoprop}}$<br>(g mol <sup>-1</sup> ) | $M_{w,\text{deg},\text{isoprop}}$<br>(g mol <sup>-1</sup> ) | $M_{n,\text{deg},\text{TBD}}$<br>(g mol <sup>-1</sup> ) | $M_{w,\text{deg},\text{TBD}}$<br>(g mol <sup>-1</sup> ) |
|                 |                                   |                                   |                              |  | (% $M_n$ loss)  | (% $M_w$ loss)  | (% $M_n$ loss)  | (% $M_w$ loss)  | (% $M_n$ loss)  | (% $M_w$ loss)  |
| FRP             |                                   |                                   |                              |  |   |   |   |   |   |   |
| F0              | 7200                              | 11 000                            | 1.53                         | 0  | 7200 (0)  | 11 000 (0)  | 7200 (0)  | 10 900 (-1)   | 7200 (0)  | 10 900 (-1)   |
| F1              | 10 000                            | 16 400                            | 1.64                         | 2.7  | 5300 (-47)  | 10 800 (-34)  | 9600 (-4)   | 14 400 (-12)  | 2700 (-73)  | 4800 (-70)  |
| F2              | 10 400                            | 17 000                            | 1.63                         | 4.6  | 4100 (-61)  | 8100 (-52)  | 9600 (-8)   | 15 600 (-8)   | 1700 (-84)  | 3200 (-81)  |
| NMP             |                                   |                                   |                              |  |   |   |   |   |   |   |
| N0              | 2400                              | 2900                              | 1.19                         | 0  | 2400 (0)  | 2900 (0)  | 2400 (0)  | 2900 (0)  | 2400 (0)  | 2900 (0)  |
| N1              | 3100                              | 3900                              | 1.25                         | 2.7  | 1600 (-48)  | 2700 (-30)  | 3100 (0)  | 3900 (0)  | 1400 (-55)  | 2400 (-38)  |
| N2              | 3300                              | 4400                              | 1.32                         | 4.4  | 1600 (-52)  | 2800 (-36)  | 3300 (0)  | 4400 (0)  | 1300 (-61)  | 2200 (-50)  |
| N3              | 5300                              | 8200                              | 1.54                         | 5.3  | 1700 (-68)  | 2900 (-64)  | 5200 (-2)   | 7600 (-7)   | 1100 (-80)  | 2300 (-72)  |
| N4              | 4800                              | 7100                              | 1.48                         | 9.7  | 1000 (-79)  | 1600 (-77)  | 4700 (-3)   | 6700 (-6)   | 800 (-83)   | 1700 (-76)  |
| N5 <sup>g</sup> | 1800                              | 2200                              | 1.29                         | 6.6  | —   | —   | —   | —   | 700 (-61)   | 1000 (-55)  |
| N6              | 7900                              | 8400                              | 1.07                         | 6.1  | 1700 (-78)  | 2640 (-68)  | —   | —   | —   | —   |
| N7 <sup>g</sup> | 9300                              | 10 700                            | 1.15                         | 6.9  | 2100 (-71)  | 4650 (-56)  | —   | —   | —   | —   |
| RAFT            |                                   |                                   |                              |  |   |   |   |   |   |   |
| R0              | 3400                              | 3900                              | 1.16                         | 0  | 3400 (0)  | 4300 (0)  | —   | —   | 3400 (0)  | 4400 (0)  |
| R1              | 4500                              | 6100                              | 1.36                         | 5.5  | 1300 (-71)  | 3500 (-43)  | —   | —   | 900 (-80)   | 1800 (-70)  |
| R2              | 11 000                            | 18 200                            | 1.67                         | 3.9  | 3300 (-70)  | 7600 (-58)  | —   | —   | 2300 (-79)  | 4300 (-76)  |

<sup>a</sup> Determined by SEC on precipitated samples. <sup>b</sup> Determined by <sup>1</sup>H NMR in *d*<sub>8</sub>-THF of precipitated samples by integrating the 8H (Ar) of open DOT at 6.7–8 ppm and protons from P1 (1H for the (1,4) conformation at 5–5.5 ppm, 1H for the (1,2) conformation at 5.5–5.9 ppm and 4.4–5 ppm combining 2H of (1,2) and 2H of (3,4) conformations). For F1–F2 and R1–R2, the aromatic signal of DCP at 7.0–7.5 ppm was subtracted from the aromatic signal of DOT according to the DCP/P1 ratio calculated on the F0 and R0 spectrum respectively. <sup>c</sup> Determined according to:  $M_{n,\text{deg},\text{th}} = [(1 - F_{\text{DOT}}) / F_{\text{DOT}}] \times \text{MW}_{\text{DOT}} + \text{MW}_{\text{P1}}$ . <sup>d</sup> Determined by SEC after degradation by KOH for 16 h. <sup>e</sup> Determined by SEC after degradation by isopropylamine for 16 h. <sup>f</sup> Determined by SEC after degradation by TBD for 16 h. <sup>g</sup>  $M_n$  loss is calculated according to: %  $M_n$  loss =  $[(M_n - M_{n,\text{deg}}) / M_n] \times 100$ . <sup>h</sup> The Gem-AMA-SG1 alkoxyamine was used instead of the BB alkoxyamine.



molecular weights, the copolymerization reaction was then carried out for 72 h with  $DP_{n,th} = 600$  (**R2**, Table 1) instead of 500. This time, 55% conversion was obtained as well as  $M_n = 11\,000\text{ g mol}^{-1}$  with however a higher dispersity ( $D = 1.67$ ). The DOT contents in the P(I-co-DOT) copolymers were determined to be 5.5 and 3.9 mol% for **R1** and **R2**, respectively (Table 1 and Fig. S6†), which is also in line with the faster incorporation of DOT compared to I. Therefore, these results confirmed the ability of DOT to be inserted into the PI backbone by FRP, NMP and RAFT polymerization mechanisms.

Interestingly, the values of  $F_{DOT}$  vs. conversion for **F2**, **N2–N5**, **N3'** and **R1–R2** are in fairly good agreement with the theoretical evolution of  $F_{DOT}$  vs. conversion plotted using a PREDICI numerical simulation<sup>58</sup> that predicts the compositional drift occurring during the copolymerization between vinyl and cyclic monomers, with  $r_{DOT}$ ,  $r_I$  and  $f_{DOT,0}$  as the input parameters (Fig. S7†). This observation therefore validates the values of the reactivity ratios that we determined. It is also important to note that the copolymerization behaviour is dramatically different when the theoretical evolution of  $F_{MDO}$  vs. conversion is plotted using the corresponding reactivity ratios<sup>38</sup> and the same  $f_{MDO,0}$  value as for DOT. Indeed,  $F_{MDO}$  values stay below 0.003 regardless of conversion, in contrast to  $F_{DOT}$  values that range from 0.12 to 0.02 with conversion. This result highlights the strong added value of DOT over CKA in incorporating weak bonds into polyisoprene chains.

### Degradation of P(I-co-DOT) copolymers

Degradation of the copolymers was then performed to confirm the successful insertion of thioester groups. Different degradation conditions were used (e.g., KOH, TBD and isopropylamine)<sup>39–44</sup> to investigate hydrolysis and aminolysis as

two distinct degradation modalities (in addition to physiological<sup>48</sup> and oxidative<sup>59</sup> degradations of thioester-containing copolymers). In general, whatever the polymerization methods used to generate the copolymers, significant degradation was observed after their treatment with either KOH or TBD for 16 h (Table 2, Fig. 4, S8 and S9†). The decrease in  $M_n$  and  $M_w$  ranged from –47 to –79% with KOH and from –55 to –84% with TBD (Table 2). Conversely, the  $M_n$  of PI homopolymers (**F0**, **N0** and **R0**) after degradation under the same conditions stayed constant (Fig. S8–S10† and Table 2). The degradation was also in agreement with the DOT content as increasing  $F_{DOT,0}$  from 0.027 (**F1**, Table 2) up to 0.046 (**F2**, Table 2) led to a higher decrease in  $M_n$  and  $M_w$  in the presence of TBD or KOH compared to **F1**. A similar trend was observed for copolymers obtained by NMP (**N1–N4**, Table 2). In addition,  $M_{n,deg}$  values were in fairly good agreement with theoretical  $M_{n,s}$  values of the copolymers after degradation ( $M_{n,deg,th}$ , Table 2). However, no degradation was shown in the presence of isopropylamine, which could be correlated to the strong hydrophobicity of PI, similar to that observed with styrene.<sup>42,44</sup>

### Application to nanoparticles and polymer prodrug nanoparticles

Polymer nanoparticles for drug delivery purposes are usually obtained by formulation of preformed polymers such as aliphatic polyesters, poly(alkyl cyanoacrylate), *etc.*<sup>60–64</sup> These polymers, which are few in number, are chosen because of their biocompatibility and biodegradability compared to non-degradable materials, which can accumulate in the body and induce prohibitive side effects. In order to diversify the range of polymers suitable for biomedical applications, the design of (bio)degradable vinyl polymers is currently of great interest.<sup>1</sup>

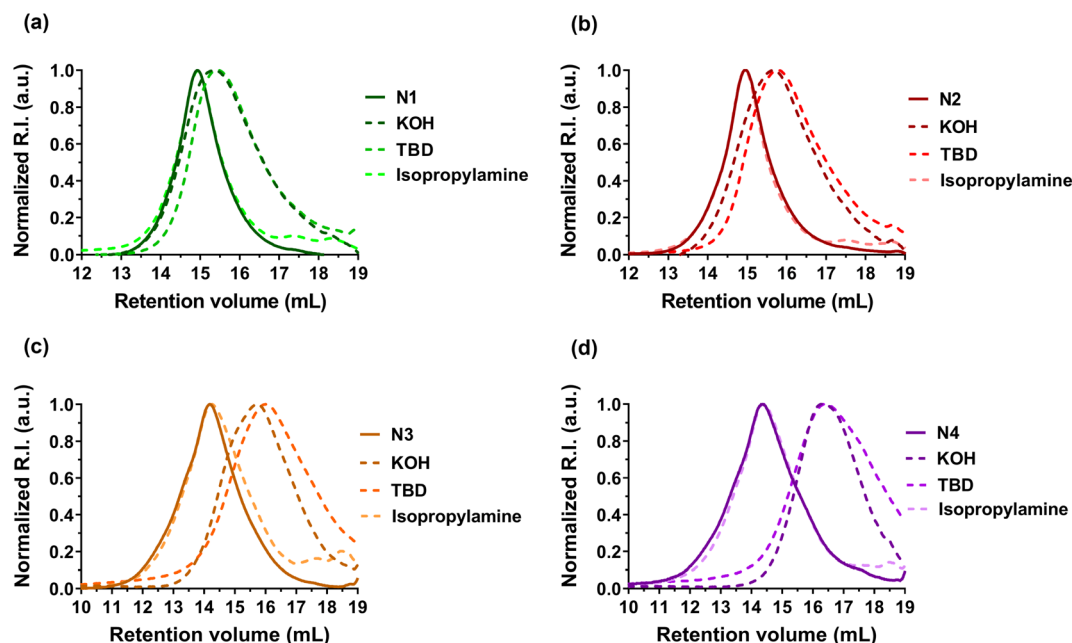


Fig. 4 Evolution of the SEC chromatograms of P(I-co-DOT) copolymers: (a) **N1**, (b) **N2**, (c) **N3**, and (d) **N4**, after degradation under basic (KOH or TBD) or aminolytic (isopropylamine) conditions.



**Table 3** Colloidal and macromolecular characteristics of P(I-co-DOT) nanoparticles obtained by nanoprecipitation before and after degradation by TBD

| Entry | $D_z^a$ (nm) | PDI <sup>a</sup> | $\zeta$ (mV) | % $M_n$ loss, deg.TBD <sup>b</sup> | % $M_w$ loss, deg.TBD <sup>b</sup> | $\bar{D}^b$ |
|-------|--------------|------------------|--------------|------------------------------------|------------------------------------|-------------|
| N0    | 150          | 0.09             | −56          | 0                                  | −3                                 | 1.17        |
| N1    | 140          | 0.15             | −27          | −45                                | −30                                | 1.58        |
| N2    | 145          | 0.10             | −50          | −50                                | −26                                | 2.30        |
| N3    | 130          | 0.07             | −46          | −56                                | −48                                | 2.05        |
| N4    | 105          | 0.09             | −27          | −59                                | −39                                | 2.24        |

<sup>a</sup> Determined by DLS. <sup>b</sup> Determined by SEC after degradation of the nanoparticles in water with TBD for 48 h.

We illustrated the potential of the I/DOT copolymerization system for drug delivery purposes by first investigating the formulation of P(I-co-DOT) copolymers into nanoparticles (N0–N4, Table 3). Narrowly dispersed nanoparticles were successfully formed by nanoprecipitation, exhibiting an average diameter between 105 and 150 nm (Table 3 and Fig. S11†). The nanoparticle diameters also tended to decrease with the increase of the copolymer chain length, in agreement with previous data on PI nanoparticles.<sup>19</sup> This appeared to be attributed to an increase in hydrophobic interactions between PI chains as they lengthen, making the core of the nanoparticles more compact. All nanoparticles also exhibited strongly negative zeta potentials (Table 3), suggesting efficient electrostatic stabilisation, which was confirmed by their remarkable long-term colloidal stability up to one month (Table 3 and Fig. 5a, S12†). Cryo-TEM analysis also revealed the formation of nanoparticles with spherical morphologies (Fig. 5b). Overall,

insertion of thioester units in the PI backbone did not impact the colloidal characteristics (size, stability and morphology) of the nanoparticles compared to their non-degradable PI counterparts.<sup>19</sup>

Direct degradation of the nanoparticles was then attempted in the presence of TBD for 48 h at room temperature. Despite the strong hydrophobicity of the copolymer that may prevent efficient access of TBD to the hydrophobic core of nanoparticles, the decrease in  $M_n$  and  $M_w$  reached 40–59% and 26–48%, respectively (N1–N4, Table 3). The higher the  $F_{DOT}$ , the greater the degradation, even if this trend was less pronounced than that for the copolymers, probably for reasons of accessibility to the thioester group in the nanoparticulate state. However, longer degradation times and/or higher concentrations of TBD may have certainly enhanced the degradation of the nanoparticles.



**Fig. 5** (a) Evolution of the intensity-average diameter ( $D_z$ ) and polydispersity index (PDI) with time measured by DLS of P(I-co-DOT) nanoparticles (N2, N3 and N4, Table 3). (b) Representative cryo-TEM images of nanoparticles N0 and N2.



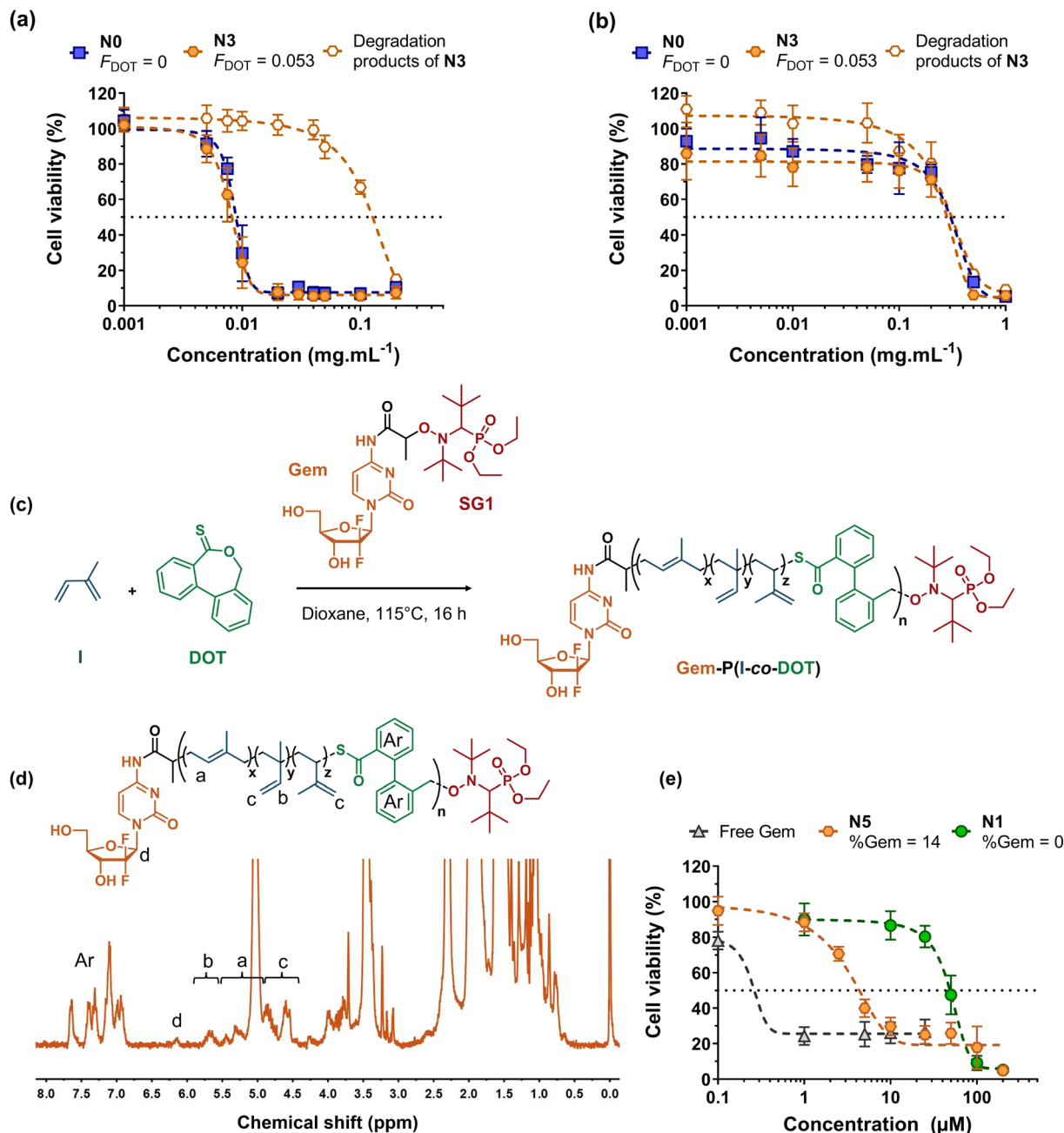


Fig. 6 Cell viability of: (a) murine macrophage (J774.A1) and (b) human endothelial cell (HUVEC) lines as a function of the concentration of nanoparticles N3 and N0 (Table 3) and of N3 degradation products. (c) Synthesis of the Gem-P(I-co-DOT) polymer prodrug by radical ring-opening polymerization (rROP) of dibenzo[c,e]oxepane-5-thione (DOT) and isoprene (I) via nitroxide-mediated polymerization using a gemcitabine-functionalized macroalcoxyamine based on the nitroxide SG1 (Gem-AMA-SG1). (d)  $^1\text{H}$  NMR spectrum (300 MHz,  $d_8$ -THF) in the 0–8 ppm region of the Gem-P(I-co-DOT) polymer prodrug N5 (Table 1). (e) Cell viability of the lung cancer cell line (A549) as a function of the concentration of Gem-P(I-co-DOT) nanoparticles (N5, Table 1), P(I-co-DOT) nanoparticles (N1, Table 1) and free gemcitabine.

Cell viability experiments were then performed on two different healthy cell lines (J774.A1 and HUVEC), during which P(I-co-DOT) nanoparticles or the degradation products of the corresponding copolymers were evaluated (Fig. 6a and b). Half-maximal inhibitory concentrations ( $\text{IC}_{50}$ ) of cell proliferation of  $7.9 \mu\text{g mL}^{-1}$  and  $0.31 \text{ mg mL}^{-1}$  were obtained for J774.A1 and HUVEC cells, respectively. These values are identical to the  $\text{IC}_{50}$  values obtained for PI nanoparticles (Fig. 6a and b),

demonstrating the absence of the cytotoxicity effect from DOT units. Moreover, the degradation products exhibited similar  $\text{IC}_{50}$  values compared to the nanoparticles for HUVEC cells and much higher  $\text{IC}_{50}$  values on J774.A1 cells ( $0.11 \text{ mg mL}^{-1}$ ). These results are very encouraging and let envision the use of P(I-co-DOT) as a carrier material for the design of degradable polymer nanoparticles for drug delivery.

Of the many polymer-based drug delivery systems developed to date, polymer-drug nanocarriers, in which the drug is covalently bound to the polymer, are the most promising because they circumvent the limitations associated with physical encapsulation of drugs (*e.g.*, burst release, poor drug loading, *etc.*).<sup>65–68</sup> While grafting drugs to a preformed polymer is the most popular strategy to produce polymer prodrugs, the “drug-initiated” method,<sup>69</sup> which consists in growing a polymer from a drug to obtain one drug molecule at the extremity of a well-defined polymer chain, has recently received increasing attention as an easy and scalable route to yield high drug loading, surfactant-free, vinyl polymer prodrug nanoparticles.<sup>17,19,21,22,70–72</sup> For instance, this has been applied to the synthesis of the Gem-polyisoprene prodrug by NMP, whose nanoparticles exhibited *in vitro* and *in vivo* anticancer activities on different cancer cell lines. However, they were not degradable, which may severely limit their development because of potential accumulation in the body and harmful side effects. Herein, we propose to apply the I/DOT copolymerization system to the synthesis of degradable Gem-P(I-*co*-DOT) prodrug nanoparticles (Fig. 6c). To install a Gem moiety at the  $\alpha$ -position of P(I-*co*-DOT) copolymer chains, the copolymerization of I and DOT ( $f_{\text{DOT},0} = 0.02$ ) was initiated by the Gem-AMA-SG1 alkoxyamine. Gem-AMA-SG1 was obtained by coupling unprotected Gem to the AMA-SG1 alkoxyamine, resulting in an amide bond that can be selectively cleaved by cathepsin B, whose over-expression is correlated with invasive and metastatic cancers.<sup>73,74</sup> After 16 h (11% conv. in I), this yielded Gem-P(I-*co*-DOT) copolymers with  $M_n = 1800 \text{ g mol}^{-1}$  and  $D = 1.29$  (N5, Table 1).  $^1\text{H}$  NMR analysis of N5 confirmed the synthesis of the expected structure (Fig. 6d), particularly *via* the presence of Gem (peak d) and the characteristic proton signals of the PI

backbone, and gave  $M_{n,\text{NMR}} = 2400 \text{ g mol}^{-1}$  (by integrating peak *a* from PI and peak *d* from Gem, Fig. 6d) in rather good agreement with the  $M_{n,\text{SEC}}$  value (Table 1). Targeting such a low  $M_n$  allowed the prodrug to exhibit a Gem loading as high as 14 wt% with a  $F_{\text{DOT}}$  value of 6.6 mol% (Fig. 6d and Table 1). TBD-assisted degradation of Gem-P(I-*co*-DOT) prodrug N5 led to a decrease in  $M_n$  and  $M_w$  of 61% and 55%, respectively (very close to the theoretical degradation), thus confirming the presence of thioester groups in the PI backbone and their successful cleavage (Table 2 and Fig. S13†). After nanoprecipitation, narrowly dispersed nanoparticles were obtained with  $D_z = 140 \text{ nm}$  and a low PDI (0.11). They also showed a greater negative zeta potential value ( $-68 \text{ mV}$ ) than their drug-free counterparts (Table 3), suggesting the (partial) presence of Gem moieties at the surface nanoparticle surface, in agreement with molecular modelling experiments<sup>75</sup> for polymer prodrugs obtained by the “drug-initiated” method.

Gem-P(I-*co*-DOT) prodrug nanoparticles were then tested for their *in vitro* cytotoxicity on human lung carcinoma (A549) cells. Whereas drug-free P(I-*co*-DOT) nanoparticles gave an  $\text{IC}_{50}$  of  $49 \mu\text{M}$ , the prodrug nanoparticles led to a 35-fold-decrease in  $\text{IC}_{50}$  ( $1.4 \mu\text{M}$ ) (Fig. 6e). This demonstrates the cytotoxic activity of Gem-P(I-*co*-DOT) prodrug nanoparticles. As expected, free Gem exhibited a much greater cytotoxicity ( $\text{IC}_{50} = 0.25 \mu\text{M}$ ), since the free drug is immediately active whereas a prodrug must release the drug for it to be active. However, Gem is known to be quickly deaminated by deoxycytidine deaminase,<sup>76</sup> thus preventing its direct administration *in vivo* and requiring the use of protecting nanocarriers/prodrugs.

After establishing the degradation of P(I-*co*-DOT) nanoparticles by TBD (Table 3), the final step of this work was to study their degradation under physiological conditions, which



Fig. 7 SEC chromatograms of: (a) P(I-*co*-DOT) polymer (N6, see Table 1) and (b) Gem-P(I-*co*-DOT) prodrug (N7, see Table 1) after degradation in the presence of KOH (5 wt%, 20 °C) on the polymer and chlorine bleach solution (2.5%, 20 °C) on the nanoparticles. SEC chromatograms of: (c) P(I-*co*-DOT) polymer (N6, see Table 1) and (d) Gem-P(I-*co*-DOT) prodrug (N7, see Table 1) after degradation in the presence of cysteine (pH 7.4, PBS, 37 °C) and glutathione (pH 7.2, PBS, 37 °C) on the nanoparticles.



**Table 4** Colloidal and macromolecular characteristics of P(I-co-DOT) **N6** and Gem-P(I-co-DOT) **N7** nanoparticles obtained by nanoprecipitation before and after degradation by bleach solution, cysteine or glutathione (GSH)

| Entry                 | $D_z^a$<br>(nm) | PDI <sup>a</sup> | $\zeta$<br>(mV) | Degradation by bleach solution <sup>b</sup>                    |  |           | Degradation by cysteine <sup>c</sup>                             |  |           | Degradation by GSH <sup>d</sup>                             |   |           |
|-----------------------|-----------------|------------------|-----------------|--|--|-----------|--|--|-----------|---|---|-----------|
|                       |                 |                  |                 | $M_{n,deg,bleach}$<br>(g mol <sup>-1</sup> )<br>(% $M_n$ loss) | $M_{w,deg,bleach}$<br>(g mol <sup>-1</sup> )<br>(% $M_w$ loss) | $\bar{D}$ | $M_{n,deg,cysteine}$<br>(g mol <sup>-1</sup> )<br>(% $M_n$ loss) | $M_{w,deg,cysteine}$<br>(g mol <sup>-1</sup> )<br>(% $M_w$ loss) | $\bar{D}$ | $M_{n,deg,GSH}$<br>(g mol <sup>-1</sup> )<br>(% $M_n$ loss) | $M_{w,deg,GSH}$<br>(g mol <sup>-1</sup> )<br>(% $M_w$ loss) | $\bar{D}$ |
| <b>N6</b>             | 185             | 0.08             | -37             | 1800 (-77)   | 2900 (-65)   | 1.60      | 6300 (-20)   | 6950 (-17)   | 1.10      | 6500 (-18)  | 7200 (-14)  | 1.11      |
| <b>N7<sup>e</sup></b> | 130             | 0.08             | -41             | 2600 (-65)   | 3900 (-63)   | 1.47      | 6600 (-19)   | 8300 (-11)   | 1.25      | 7100 (-12)  | 8600 (-8)   | 1.21      |

<sup>a</sup> Determined by DLS. <sup>b</sup> Determined by SEC after degradation of the nanoparticles by 2.5% active chlorine bleach solution during 12 days at 20 °C.

<sup>c</sup> Determined by SEC after degradation of the nanoparticles in the presence of cysteine in PBS pH 7.4 during 15 days at 37 °C. <sup>d</sup> Determined by SEC after degradation of the nanoparticles in the presence of GSH in PBS pH 7.2 during 15 days at 37 °C. <sup>e</sup> The Gem-AMA-SG1 alkoxyamine was used instead of BB alkoxyamine.

is a key issue when developing biodegradable drug delivery systems. To this end, two model copolymers were synthesized: P(I-co-DOT) **N6** ( $M_n = 7900$  g mol<sup>-1</sup> and  $F_{DOT} = 6.1$  mol%) and Gem-P(I-co-DOT) **N7** ( $M_n = 9300$  g mol<sup>-1</sup> and  $F_{DOT} = 6.9$  mol%). They were then formulated by nanoprecipitation into nanoparticles (Table 4) exhibiting high colloidal stability for up to at least 14 days (**N6**:  $D_z = 185$  nm, PDI = 0.08 and **N7**:  $D_z = 130$  nm, PDI = 0.08).

Spurred by the physiological thiolytic degradability of DOT-containing copolymers,<sup>48</sup> the nanoparticles were subjected to degradation in the presence of: (i) chlorine bleach solution (2.5% sodium hypochlorite, 20 °C), which mimics both oxidative and base-catalysed hydrolysis due to the oxidative and alkaline nature of bleach, respectively<sup>59</sup> and (ii) cysteine (10 mM, pH 7.4, PBS, 37 °C) or glutathione (10 mM, pH 7.2, PBS, 37 °C), which mimic reductive degradations under physiological conditions. Note that using degradation under alkaline conditions is a routine procedure in rROP, which is used to anticipate and predict the degradation of rROP-derived copolymers in the long run. For instance, this is usually performed with ester-containing copolymers obtained by rROP of CKA and vinyl monomers.<sup>77–80</sup>

Degradation of nanoparticles **N6** and **N7** under oxidative/alkaline conditions (Fig. 7a,b and Table 4) resulted in a significant decrease in  $M_n$  and  $M_w$  (~60–80% after 12 days), which was similar to the degradation of the corresponding copolymers under alkaline conditions (KOH 5 wt%, see Table 2). More importantly, degradation of nanoparticles **N6** and **N7** under physiological reductive conditions successfully occurred (Table 4). As expected, the degradation kinetics were slower (~10–20% decrease in  $M_n$  or  $M_w$  after 15 days) than under oxidative/alkaline conditions, but significantly faster than the hydrolytic degradation under physiological conditions of most CKA-containing copolymers, which typically degrade over several months and even a year.<sup>81–83</sup> Overall, these results confirmed that P(I-co-DOT) and Gem-P(I-co-DOT) nanoparticles could be degraded under physiological conditions.

## Conclusion

We successfully synthesized a small library of thionolactone-containing polyisoprene by radical ring-opening

copolymerization of isoprene and DOT, under either FRP, NMP and RAFT polymerization conditions. The resulting P(I-co-DOT) copolymers exhibited predictable  $M_n$ , low dispersity and tunable DOT contents (2.7–9.7 mol%) simply by adjusting the copolymerization conditions. The copolymers were significantly degraded under basic conditions, which confirmed the successful insertion of open DOT units in the PI chains. Cyto-compatible, narrowly dispersed, surfactant-free nanoparticles were also prepared by nanoprecipitation of P(I-co-DOT) copolymers, with average diameters suitable for drug delivery purposes. As a proof of concept, well-defined Gem-P(I-co-DOT) polymer prodrug nanoparticles were produced, which exhibited significant cytotoxicity on A549 cancer cells, as well as degradation under oxidative/alkaline conditions and reductive physiological conditions. In conclusion, DOT represents a promising monomer that can easily introduce labile thioester groups into PI chains *via* rROP and advantageously broaden the range of degradable vinyl materials, not only in nanomedicine but also for other applications involving PI.

## Data availability

The authors declare that all data supporting the findings of this study are available within the article and ESI,† and raw data files are available from the corresponding author upon request.

## Author contributions

M. L.: investigation, writing – original draft, data curation, visualization. C. Z.: investigation, data curation. T. P.: investigation, data curation. D. L.: investigation, data curation. J. M.: investigation, data curation. Y. G.: formal analysis, software, writing – review & editing. J. N.: conceptualization, visualization, writing – review & editing, funding acquisition, supervision, resources.

## Conflicts of interest

There are no conflicts to declare.



## Acknowledgements

The authors acknowledge the financial support from the Agence Nationale de la Recherche (grant number ANR-18-CE06-0014 CKAPART). We thank Stéphanie Denis (IGPS, Université Paris-Saclay) for technical assistance in cell culture experiments and Dr Sylvain Trepout (Institut Curie, Orsay, France) and PICT-Ibisa for the experiments using a JEOL 2200FS TEM. We thank Pr. Alex Van Herk from Eindhoven University of Technology for sharing the CONTOUR software. This work has benefited from the facilities and expertise of the Platform for Transmission Electronic Microscopy of I2BC (Université Paris-Saclay).

## References

- 1 V. Delplace and J. Nicolas, *Nat. Chem.*, 2015, **7**, 771–784.
- 2 R. Wei, T. Tiso, J. Bertling, K. O'Connor, L. M. Blank and U. T. Bornscheuer, *Nat. Catal.*, 2020, **3**, 867–871.
- 3 A. Rahimi and J. M. García, *Nat. Rev. Chem.*, 2017, **1**, 0046.
- 4 The future of plastic, *Nat. Commun.*, 2018, **9**, 2157, <https://doi.org/10.1038/s41467-018-04565-2>.
- 5 T. Pesenti and J. Nicolas, *ACS Macro Lett.*, 2020, **9**, 1812–1835.
- 6 A. Tardy, J. Nicolas, D. Gïmes, C. Lefay and Y. Guillaueuf, *Chem. Rev.*, 2017, **117**, 1319–1406.
- 7 S. Nazhat, S. Parker, M. Patel and M. Braden, *Biomaterials*, 2001, **22**, 2411–2416.
- 8 W. H. Kaminsky and B., *Plast. Eng.*, 2005, **70**, 333–380.
- 9 A. Region Europe, *And Segment Forecasts, 2021–2028*.
- 10 S. Förster and E. Krämer, *Macromolecules*, 1999, **32**, 2783–2785.
- 11 J. K. Wegrzyn, T. Stephan, R. Lau and R. B. Grubbs, *J. Polym. Sci., Part A: Polym. Chem.*, 2005, **43**, 2977–2984.
- 12 J. W. Bartels, S. I. Cauët, P. L. Billings, L. Y. Lin, J. Zhu, C. Fidge, D. J. Pochan and K. L. Wooley, *Macromolecules*, 2010, **43**, 7128–7138.
- 13 S. C. Schmidt and M. A. Hillmyer, *Macromolecules*, 1999, **32**, 4794–4801.
- 14 J. Wootthikanokkhan and B. Tongrubai, *J. Appl. Polym. Sci.*, 2003, **88**, 921–927.
- 15 K. S. Murthy, Q. Ma, E. E. Remsen, T. Kowalewski and K. L. Wooley, *J. Mater. Chem.*, 2003, **13**, 2785–2795.
- 16 C. Cheng, K. Qi, E. Khoshdel and K. L. Wooley, *J. Am. Chem. Soc.*, 2006, **128**, 6808–6809.
- 17 D. Vinciguerra, A. Degrassi, L. Mancini, S. Mura, J. Mougin, P. Couvreur and J. Nicolas, *Biomacromolecules*, 2019, **20**, 2464–2476.
- 18 D. Vinciguerra, M. Jacobs, S. Denis, J. Mougin, Y. Guillaueuf, G. Lazzari, C. Zhu, S. Mura, P. Couvreur and J. Nicolas, *J. Controlled Release*, 2019, **295**, 223–236.
- 19 S. Harisson, J. Nicolas, A. Maksimenko, D. T. Bui, J. Mougin and P. Couvreur, *Angew. Chem., Int. Ed.*, 2013, **52**, 1678–1682.
- 20 Y. Bao, E. Guégain, V. Nicolas and J. Nicolas, *Chem. Commun.*, 2017, **53**, 4489–4492.
- 21 Y. Bao, E. Guégain, J. Mougin and J. Nicolas, *Polym. Chem.*, 2018, **9**, 687–698.
- 22 Y. Bao, T. Boissenot, E. Guégain, D. Desmaële, S. Mura, P. Couvreur and J. Nicolas, *Chem. Mater.*, 2016, **28**, 6266–6275.
- 23 S. Schlögl, M.-L. Trutschel, W. Chassé, I. Letofsky-Papst, R. Schaller, A. Holzner, G. Riess, W. Kern and K. Saalwächter, *Polymer*, 2014, **55**, 5584–5595.
- 24 W. M. Gramlich and M. A. Hillmyer, *Polym. Chem.*, 2011, **2**, 2062–2067.
- 25 F. Chen and J. Qian, *Fuel*, 2002, **81**, 2071–2077.
- 26 S.-Y. Chen, Y. Huang and R. C.-C. Tsiang, *J. Polym. Sci., Part A: Polym. Chem.*, 2008, **46**, 1964–1973.
- 27 E. I. Baptista, G. M. Campese, N. L. Zalloum, A. F. Rubira and E. C. Muniz, *Polym. Degrad. Stab.*, 1998, **60**, 309–315.
- 28 P. Berto, S. Grelier and F. Peruch, *Polym. Degrad. Stab.*, 2018, **154**, 295–303.
- 29 C. W. Phetphaisit and P. Phinyocheep, *J. Appl. Polym. Sci.*, 2003, **90**, 3546–3555.
- 30 S. S. Solanky, I. Campistron, A. Laguerre and J.-F. Pilard, *Macromol. Chem. Phys.*, 2005, **206**, 1057–1063.
- 31 S. Ouardad and F. Peruch, *Polym. Degrad. Stab.*, 2014, **99**, 249–253.
- 32 J. A. Blach, G. S. Watson, W. K. Busfield and S. Myhra, *Polym. Int.*, 2002, **51**, 12–20.
- 33 C. Adam, J. Lacoste and J. Lemaire, *Polym. Degrad. Stab.*, 1991, **32**, 51–69.
- 34 S. Sato, Y. Honda, M. Kuwahara and T. Watanabe, *Biomacromolecules*, 2003, **4**, 321–329.
- 35 M. Enoki, Y. Doi and T. Iwata, *Macromol. Biosci.*, 2003, **3**, 668–674.
- 36 A.-L. Altenhoff, S. Thierbach and A. Steinbüchel, *Biodegradation*, 2021, **32**, 113–125.
- 37 M. D. Chengalroyen and E. R. Dabbs, *J. Polym. Environ.*, 2013, **21**, 874–880.
- 38 A. Tardy, N. Gil, C. M. Plummer, C. Zhu, S. Harisson, D. Siri, J. Nicolas, D. Gïmes, Y. Guillaueuf and C. Lefay, *Polym. Chem.*, 2020, **11**, 7159–7169.
- 39 N. M. Bingham and P. J. Roth, *Chem. Commun.*, 2019, **55**, 55–58.
- 40 R. A. Smith, G. Fu, O. McAteer, M. Xu and W. R. Gutekunst, *J. Am. Chem. Soc.*, 2019, **141**, 1446–1451.
- 41 N. M. Bingham, Q. U. Nisa, S. H. L. Chua, L. Fontugne, M. P. Spick and P. J. Roth, *ACS Appl. Polym. Mater.*, 2020, **2**, 3440–3449.
- 42 N. Gil, B. Caron, D. Siri, J. Roche, S. Hadiouch, D. Khedaoui, S. Ranque, C. Cassagne, D. Montarnal and D. Gïmes, *Macromolecules*, 2022, **55**, 6680–6694.
- 43 M. P. Spick, N. M. Bingham, Y. Li, J. de Jesus, C. Costa, M. J. Bailey and P. J. Roth, *Macromolecules*, 2020, **53**, 539–547.
- 44 P. Galanopoulou, N. Gil, D. Gïmes, C. Lefay, Y. Guillaueuf, M. Lages, J. Nicolas, M. Lansalot and F. D'Agosto, *Angew. Chem., Int. Ed.*, 2022, **61**, e202117498.
- 45 N. Gil, C. Thomas, R. Mhanna, J. Mauriello, R. Maury, B. Leuschel, J.-P. Malval, J.-L. Clément, D. Gïmes and C. Lefay, *Angew. Chem., Int. Ed.*, 2021, **134**, e202117700.
- 46 M. Lages, N. Gil, P. Galanopoulou, J. Mougin, C. Lefay, Y. Guillaueuf, M. Lansalot, F. D'Agosto and J. Nicolas, *Macromolecules*, 2022, **55**, 9790–9801.
- 47 G. R. Kiel, D. J. Lundberg, E. Prince, K. E. L. Husted, A. M. Johnson, V. Lensch, S. Li, P. Shieh and J. A. Johnson, *J. Am. Chem. Soc.*, 2022, **144**, 12979–12988.



- 48 N. Bingham, Q. Nisa, P. Gupta, N. Young, E. Vellio and P. Roth, *Biomacromolecules*, 2022, **23**, 2031–2039.
- 49 J. Nicolas, Y. Guillaneuf, C. Lefay, D. Bertin, D. Gigmes and B. Charleux, *Prog. Polym. Sci.*, 2013, **38**, 63–235.
- 50 S. Perrier, *Macromolecules*, 2017, **50**, 7433–7447.
- 51 M. R. Wood, D. J. Duncalf, S. P. Rannard and S. Perrier, *Org. Lett.*, 2006, **8**, 553–556.
- 52 S. Allorio, S. Pispas, E. Siakali-Kioulafa and N. Hadjichristidis, *J. Polym. Sci., Part B: Polym. Phys.*, 1995, **33**, 2229–2234.
- 53 S. Harriison, P. Couvreur and J. Nicolas, *Macromolecules*, 2011, **44**, 9230–9238.
- 54 M. Van Den Brink, A. M. Van Herk and A. L. German, *J. Polym. Sci., Part A: Polym. Chem.*, 1999, **37**, 3793–3803.
- 55 S. Harriison, P. Couvreur and J. Nicolas, *Macromol. Rapid Commun.*, 2012, **33**, 805–810.
- 56 V. Jitchum and S. Perrier, *Macromolecules*, 2007, **40**, 1408–1412.
- 57 V. Delplace, S. Harriison, A. Tardy, D. Gigmes, Y. Guillaneuf and J. Nicolas, *Macromol. Rapid Commun.*, 2014, **35**, 484–491.
- 58 D. Gigmes, P. H. M. Van Steenberge, D. Siri, D. R. D'hooge, Y. Guillaneuf and C. Lefay, *Macromol. Rapid Commun.*, 2018, **39**, 1800193.
- 59 O. Ivanchenko, S. Mazières, S. Harriison and M. Destarac, *Polym. Chem.*, 2022, **13**, 5525–5529.
- 60 C. Vauthier, C. Dubernet, E. Fattal, H. Pinto-Alphandary and P. Couvreur, *Adv. Drug Delivery Rev.*, 2003, **55**, 519–548.
- 61 A. Kumari, S. K. Yadav and S. C. Yadav, *Colloids Surf., B*, 2010, **75**, 1–18.
- 62 E. M. Pridgen, R. Langer and O. C. Farokhzad, *Nanomedicine*, 2007, **2**, 669–680.
- 63 K. S. Soppimath, T. M. Aminabhavi, A. R. Kulkarni and W. E. Rudzinski, *J. Controlled Release*, 2001, **70**, 1–20.
- 64 M. Sumana, A. Thirumurugan, P. Muthukumaran and K. Anand, in *Integrative Nanomedicine for New Therapies*, Springer, 2020, pp. 231–246.
- 65 V. Delplace, P. Couvreur and J. Nicolas, *Polym. Chem.*, 2014, **5**, 1529–1544.
- 66 I. Ekladios, Y. L. Colson and M. W. Grinstaff, *Nat. Rev. Drug Discovery*, 2019, **18**, 273–294.
- 67 M. L. Girase, P. G. Patil and P. P. Ige, *Int. J. Polym. Mater. Polym. Biomater.*, 2020, **69**, 990–1014.
- 68 P. Thakor, V. Bhavana, R. Sharma, S. Srivastava, S. B. Singh and N. K. Mehra, *Drug Discovery Today*, 2020, **25**, 1718–1726.
- 69 J. Nicolas, *Chem. Mater.*, 2016, **28**, 1591–1606.
- 70 Y. Bao and J. Nicolas, *Polym. Chem.*, 2017, **8**, 5174–5184.
- 71 E. Guégain, J. Tran, Q. Deguettes and J. Nicolas, *Chem. Sci.*, 2018, **9**, 8291–8306.
- 72 D. Vinciguerra, S. Denis, J. Mougin, M. Jacobs, Y. Guillaneuf, S. Mura, P. Couvreur and J. Nicolas, *J. Controlled Release*, 2018, **286**, 425–438.
- 73 C. S. Gondi and J. S. Rao, *Expert Opin. Ther. Targets*, 2013, **17**, 281–291.
- 74 D. Dheer, J. Nicolas and R. Shankar, *Adv. Drug Delivery Rev.*, 2019, **151–152**, 130–151.
- 75 P. Gao, J. Nicolas and T. Ha-Duong, *J. Am. Chem. Soc.*, 2021, **143**, 17412–17423.
- 76 V. Heinemann, Y.-Z. Xu, S. Chubb, A. Sen, L. W. Hertel, G. B. Grindey and W. Plunkett, *Cancer Res.*, 1992, **52**, 533–539.
- 77 G. G. Hedir, C. A. Bell, N. S. Jeong, E. Chapman, I. R. Collins, R. K. O'Reilly and A. P. Dove, *Macromolecules*, 2014, **47**, 2847–2852.
- 78 G. G. Hedir, C. A. Bell, R. K. O'Reilly and A. P. Dove, *Biomacromolecules*, 2015, **16**, 2049–2058.
- 79 C. Zhu, S. Denis and J. Nicolas, *Chem. Mater.*, 2022, **34**, 1875–1888.
- 80 V. Delplace, E. Guégain, S. Harriison, D. Gigmes, Y. Guillaneuf and J. Nicolas, *Chem. Commun.*, 2015, **51**, 12847–12850.
- 81 J. Undin, A. Finne-Wistrand and A.-C. Albertsson, *Biomacromolecules*, 2014, **15**, 2800–2807.
- 82 E. Guégain, J.-P. Michel, T. Boissenot and J. Nicolas, *Macromolecules*, 2018, **51**, 724–736.
- 83 J. Tran, T. Pesenti, J. Cressonnier, C. Lefay, D. Gigmes, Y. Guillaneuf and J. Nicolas, *Biomacromolecules*, 2019, **20**, 305–317.

