



Cite this: *Polym. Chem.*, 2023, **14**, 4275

Received 10th August 2023,

Accepted 31st August 2023

DOI: 10.1039/d3py00929g

rsc.li/polymers

Amine-bearing cyclic ketene acetals for pH-responsive and degradable polyesters through radical ring-opening polymerisation†

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A series of previously unreported amine-bearing cyclic ketene acetals (CKAs) furnished with different alkyl-substituents were prepared. These CKAs pave the way for a new range of pH-responsive and degradable polyesters from radical ring-opening polymerisation (RROP). CKA 2-methylene-1,3,6-trioxocane (MTC) was copolymerised with the amine-bearing CKAs to obtain a range of new degradable smart copolymers. Varying the type and amount of amine-bearing CKA allowed to fine-tune the pK_a value of the polymers, showcasing the versatility of this approach.

Following the increasing issues with plastic waste, polyesters like poly(caprolactone) (PCL) or poly(lactic acid) (PLA) and similar materials have become increasingly important materials owing to the biodegradable ester units in the polymer backbone. They found numerous applications in tissue engineering, drug delivery or commodity materials.¹ Traditional synthetic routes towards polyesters are either by polycondensation or ring-opening polymerisation.² However, the former tends to generate polyesters with low molecular weight and the latter method requires rigorous synthetic procedures (especially from ionic polymerization) to afford pure polymers.³ Proposed by Bailey *et al.*, radical ring-opening polymerisation (RROP) of cyclic ketene acetals (CKAs) has received considerable attention and is regarded as an alternative tool to generate polyesters.⁴ RROP possesses the robustness of radical polymerisations, yields degradable polyesters^{2,5} and allows for functional groups (e.g. amines) within the polymer backbone.⁶

Homopolymerisations of CKAs are known to generate polyesters (bio)degrading under basic or enzymatic conditions.^{7,8} Until now, 2-methylene-1,3-dioxepane (MDO), 2-methylene-1,3,6-trioxocane (MTC) and 2-methylene-4-phenyl-1,3-dioxolane (MPDL) are the CKAs with the highest publication

record.^{9–13} These CKAs are generally characterised by an almost complete degree of ring-opening and a known structure in terms of the main repeating unit and branching positions from inter- or intramolecular H-transfer.^{13–16} Some of these polymers already have an additional functionality like the semi-crystallinity of poly(MDO) and the higher hydrophilicity of poly(MTC) as compared to other polyesters from RROP.^{12–14} However, the lack of responsive groups limits their range of potential applications like drug delivery.¹⁷

One strategy to circumvent this issue is the copolymerisation of CKAs with various responsive vinylic monomers.^{5,18–20} This strategy introduced ester linkages into the vinyl polymer backbone to confer (bio)degradability² and responsive vinylic monomers endowed responsive properties.^{21,22} CKAs including MDO and BMDO have been successfully combined with oligoethylene glycol methacrylate and *N,N*-dimethylacrylamide for water solubility,^{23–25} *N*-isopropylacrylamide for temperature responsiveness^{26,27} and 2-(diethylamino)ethyl methacrylate for pH-responsiveness.^{23,28} CKA-bearing copolymers have been developed in the field of functional and degradable nanoparticles,^{29–31} hydrogels,³² 3D printing materials,³³ and coatings.^{34,35} However, the unfavorable CKA/vinylic copolymerisation parameters caused a composition drift towards gradient copolymer, which prohibited the randomly distributed ester units in the polymer backbone. This will therefore lead to an incomplete degradation into a mixture of oligomers.^{36,37} Hence, a reproducible method towards responsive and fully degradable polyesters from RROP is still a challenge.

Our group has focused on fully degradable polyesters purely based on CKAs,^{6,8,15,38} including amine-bearing CKAs for pH-responsive polyesters. This includes the amine-bearing *N*-iso-propyl-2-methylene-1,3,6-dioxazocane (*i*Pr-MAC, 7c), allowing for pH-responsive and biodegradable nanoparticles.⁶ With a cloud point of 6.0, the pK_a of the polymerised *i*Pr-MAC was similar to the well-known pH-responsive poly(diisopropyl aminoethyl methacrylate) (PDPA).³⁹ Copolymerisation of 7c *i*Pr-MAC with 4,7-dimethyl-2-methylene-1,3-dioxepane (DMMDO) led to copolymers with an increased molecular weight, but the pH-

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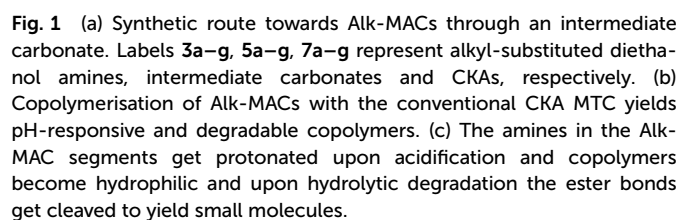
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† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d3py00929g>

These alkyl-modified diols **3a–g** and ethyl chloroformate **4** then reacted in the presence of triethylamine (TEA) to yield the intermediate cyclic carbonates **5a–g**. Reaction conditions, including reaction temperature, feeding speed of TEA, feed

Methylenation of all cyclic carbonates **5a–g** (1.0 equiv.) with the Petasyl reagent **6** (2.3 equiv.) were performed in THF at 65 °C for 22 h. All the new CKAs were obtained in a low yield (from 6% for **7d** over 11–15% for **7a–b**, **7e–g** to 25% for **7c** (Table S1†)), representing the delicacy of the carbonate route. ^1H , ^{13}C NMR and HR-ESI-MS measurements confirmed the successful preparation of these new molecules (Fig. 2 and Fig. S12–18†). As depicted in Fig. 2, signals **a–d** could be assigned to the CKA ring. From the core CKA functionality, the tertiary carbon atom **c** between the oxygen atoms was positioned at 164 ppm for the ^{13}C NMR spectrum (Fig. S12–S18†). The intense singlet **d** in the range of 3.53–3.61 ppm in the ^1H NMR spectra and new signal at 69 ppm in the ^{13}C NMR spectra represented *exo*-methylene group and hence proved the formation of the MACs. Signals **1–4** were attributed to the protons on the alkyl-substituents and remained mainly unchanged from the carbonates for all substituents. As compared to other chemical shifts of the starting diols and intermediate substances, protons **a**, **a'** and **2**, **2'** of the chiral ^sBu -MAC **7e** and the corresponding carbonate **5e** showed an additional splitting (see in Fig. 2 and S8†), which is typical for chiral molecules.

Additionally, theoretical calculations of the molecular structures of all intermediate carbonate and corresponding CKAs were performed using software package GAMESS with basis set STO-6G,⁴⁰ simulating conditions of 0 K and vacuum. These helped us in understanding the energetically optimal structure of the carbonates **5a–g** and CKAs **7a–g** (see in the insets in Fig. 2 and Fig. S20†). They revealed that, except for the chiral ^sBu derivatives **5e** and **7e**, the side chain on the amine stretched away from the carbonate or CKA ring. Both chiral derivatives, however, presented an additional structural distinctiveness. Here, the ^sBu-side group tended to bend inwards to the eight-membered ring, which might lead to significant steric hindrance around the tertiary amine. However, as the



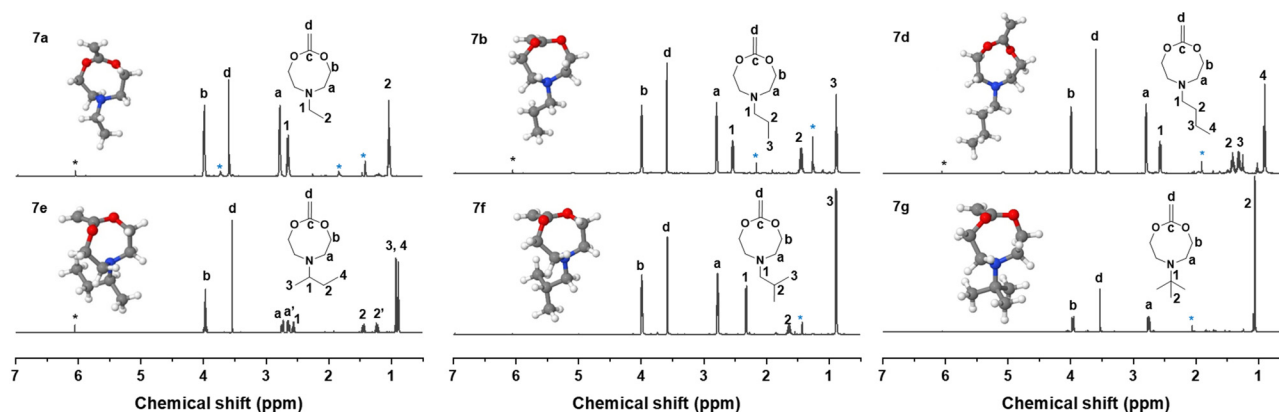


Fig. 2 ^1H NMR spectra of all new Alk-MACs **7a–b**, **7d–g** with their respective alkyl side groups. The inserts are energetic optimal structures of CKAs calculated at 0 K, under vacuum; red balls represent oxygen atoms, grey balls represent carbon atoms, white balls represent hydrogen atoms, blue balls represent nitrogen atoms. The asterisked signals in the ^1H NMR spectra represent the trace of Petasis reagent (black, 6.05 ppm), solvent and impurities (blue, *tert*-butanol: 1.28 ppm, acetone: 2.17 ppm, THF: 1.85 and 3.76 ppm).

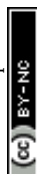
calculations were performed at simulated 0 K, the effect on the molecule at room temperature remained unknown.

Copolymerisation of the previously reported ^iPr -MAC **7c** and conventional CKA MTC was the starting point to fabricate the pH-responsive polyesters through free RROP. By varying feed ratio of **7c** to MTC from 5/95, 10/90, 25/75 to 50/50 (mol%), copolymer composition was modulated. After polymerising either thermally or under UV light for 24 h, purified copolymers were obtained through dialysis. All copolymers were named according to the feed ratio of Alk-MACs and the polymerisation method. For instance, ^iPr -MAC-5T represents a **7c**/MTC mixture with a feed ratio of 5/95 that was subjected to thermal polymerisation.

To determine the monomer conversion and the final molar fractions, crude and purified copolymers were analysed by ^1H NMR spectroscopy (Fig. S21–24†). Signals attributed to protons on the P(MTC) parts were consistent with reported signal assignments.¹⁴ For the P(^iPr -MAC) segment, the signal centered at 2.93 ppm appeared in all the samples, which is attributed to the α -proton on the ^iPr side chain. This signal was significant, as the chemical shift remained virtually unchanged for the monomer and the polymer and confirmed the successful incorporation of the ^iPr group through all attempts. After workup, the ester signal of P(MTC) (4.20 ppm) and amine signal of P(^iPr -MAC) (2.93 ppm) were integrated to calculate the molar fractions in the copolymers (Fig. S23 and section 3.3 of the ESI†). Intriguingly, the final molar fraction of ^iPr -MAC in the copolymers was always higher than the feed ratio, indicating that ^iPr -MAC was more reactive than MTC. Additionally, UV-irradiated polymerisation consistently led to a higher ^iPr -MAC content in the copolymers as compared to thermal polymerization, implying an even greater imbalance of reactivities (Table 1). Copolymers of ^iPr -MAC and MTC could hence be gradient copolymers, which means copolymers first enriched with ^iPr -MAC. Considering the omnipresent dispersity within polymers from free radical polymerisation, the

partial presence of homopolymers could not be excluded. DOSY measurements of the bimodal copolymer ^iPr -MAC-5T was carried out (Fig. S26†), and showed that the self-diffusion coefficients related to ^iPr -MAC and MTC motifs within the main diffusion peak and thus substantiated the successful formation of a copolymer. Additionally, MTC branching units occurred by intramolecular (MTC back-biting) or intermolecular H-abstraction reactions can also be observed. The characteristic signals around 1.20 ppm are related to the methyl end groups from each MTC branching unit (Fig. S24†).^{13,41} The SEC chromatograms of the resultant copolymers showed that with increasing ^iPr -MAC content, copolymers exhibited a broader distribution ranging from 1.3 for 5 mol% to over 11 for 50 mol% (Fig. S29† and Table 1). While thermal initiation showed a consistent increase in dispersity with rising ^iPr -MAC content, tendency was less clear for polymerizations under UV irradiation, again implying a more reliable reaction under purely thermal conditions. The corresponding results should be treated with caution owing to the complex polymeric structures including ring-opened and ring-retaining structures as well as branched and linear units in all possible combinations from MTC and ^iPr -MAC. 2D NMR measurements of sample copolymers (Fig. S27 and 28†) still allowed for allocating some major signals. As the ester bonds of MTC and the amines of ^iPr -MAC could be confirmed, both structural arguments for the targeted degradability and pH-responsiveness were present.

Following the more reliable reaction, thermal polymerisation was applied to other Alk-MAC/MTC pairs. A feed ratio of 25/75 (mol%, Alk-MAC/MTC) was applied to yield an Alk-MAC content between 25 and 50 mol% as this would ensure pH-responsiveness and also the MTC content necessary to keep structural diversity within reason. The purified copolymers were analysed by ^1H - and ^{13}C NMR spectroscopy and are comprised in Fig. S32–S33.† Similar to the ^iPr -MAC/MTC copolymers, the high amount of possible substructures prohibited



Name	$f_{\text{MAC},0}/f_{\text{MTC},0}$ ^a	% Conv. MAC/MTC ^b	$F_{\text{MAC}}/F_{\text{MTC}}$ ^c	M_n ^d (kg mol ⁻¹)	M_w ^d (kg mol ⁻¹)	D^d	pK _a ^e
ⁱ Pr-MAC-5T	5/95	67/35	7/93	25.8	32.9	1.3	—
ⁱ Pr-MAC-10T	10/90	40/28	14/86	26.6	43.5	1.6	—
ⁱ Pr-MAC-25T	25/75	48/25	34/66	32.2	155	4.8	7.0 ± 0.1
ⁱ Pr-MAC-50T	50/50	35/37	69/31	18.2	193	11	6.5 ± 0.1
ⁱ Pr-MAC-5U	5/95	80/67	10/90	16.3	47.4	2.9	—
ⁱ Pr-MAC-10U	10/90	100 ^f /38	19/81	29.3	68.2	2.3	—
ⁱ Pr-MAC-25U	25/75	64/15	55/45	9.2	31.5	3.4	6.9 ± 0.2
ⁱ Pr-MAC-50U	50/50	100 ^f /44	80/20	16.5	248	15	6.7 ± 0.2
Et-MAC-25T	25/75	22/17	37/63	17.8	44.8	2.5	5.7 ± 0.3
ⁿ Pr-MAC-25T	25/75	27/26	26/74	15.3	42.4	2.8	6.1 ± 0.2
ⁿ Bu-MAC-25T	25/75	22/17	36/64	11.8	126	11	6.6 ± 0.3
^s Bu-MAC-25T	25/75	29/13	45/55	49.1	420	8.6	6.6 ± 0.3
ⁱ Bu-MAC-25T	25/75	30/17	30/70	21.5	183	8.5	6.4 ± 0.2
^t Bu-MAC-25T	25/75	60/19	50/50	10.8	65.4	6.1	7.2 ± 0.3

All copolymers contained the aimed-for amount of Alk-MAC and pH-responsiveness was then characterised by dissolving each copolymer in a HCl solution, titrating it from pH 2 towards pH 10. This process deprotonated the ammonium units and gave tertiary amines, changing the solubility of the copolymers from hydrophilic (transparent solution) to hydro-

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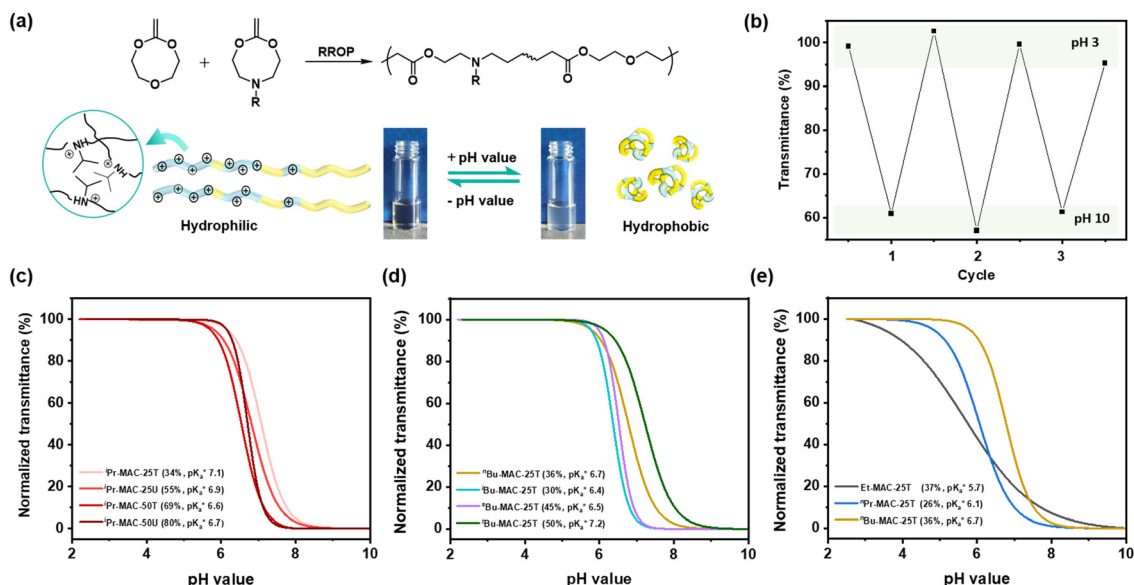


Fig. 3 (a) Transparent copolymer solutions became turbid when the pH value increased. (b) Reversible deprotonation–protonation process of ⁱPr-MAC-25T copolymer solution detected by UV-Vis measurements over three cycles. (c)–(e) Evolution of % transmittance of copolymers solution over pH, showing dependencies of (c) amount of incorporated ⁱPr-MAC, (d) bulkiness of the butyl substitute (ⁿBu, ⁱBu, ^sBu, ^tBu), (e) length of the *n*-alkyl substituent (Et, ⁱPr, ⁿBu). The pK_a values shown in the figure caption were obtained from the single Boltzmann fit of all three measurements of respective copolymer, and therefore were slightly different from the average pK_a value listed in the Table 1 (for a comparison of both values, see Table S13 in the ESI†).

following the different measurement principles. Owing to the higher hydrophilicity of MTC, all pK_a values were considerably higher than the one recorded for the ⁱPr-MAC/DMMDO copolymer (found at pH 3.5),^{6,13,14} meeting a major goal of this study.

For all Bu-MAC copolymers (Fig. 3d), the MAC content ranged from 30% in ⁱBu-MAC-25T to 50% in ^tBu-MAC-25T, being in a comparable range. The bulkiest residue of ^tBu-MAC-25T exhibited the highest pK_a of 7.2 despite having the highest amount of Bu-MAC units. All three other Bu-MACs were relatively close to each other with pK_a values of 6.4, 6.5 and 6.7 for ⁱBu-MAC-25T, ^sBu-MAC-25T and ⁿBu-MAC-25T, respectively. This implicated that the steric hindrance of the ^tBu residue caused this higher pK_a value.

Copolymers with linear substituents showed that Et-MAC-25T exhibited a significantly broader reduction region of

% transmittance than ⁱPr-MAC-25T and ⁿBu-MAC-25T. As molecular weights remained within reason, this change could be attributed to the length of the side chain. Corresponding pK_a values increased from 5.7 for Et-MAC-25T over 6.1 for ⁱPr-MAC-25T to 6.7 for ⁿBu-MAC-25T. It can hence be concluded that longer alkyl substituents promoted a higher pK_a value. This could again be attributed to the increasing steric hindrance of a larger substituent, which goes in-line with the shift of the pK_a values observed within the series of butyl-substituents.

Selected copolymers were then subjected to hydrolytic degradation under accelerated condition. The copolymers were dissolved in THF/MeOH (67/33 vol%) with 5 wt% KOH and kept stirring at room temperature for 24 h to ensure complete degradation. Degradation was then proven in multiple ways: (i) ¹H NMR measured in CDCl₃ and D₂O: signals around 4.2 ppm

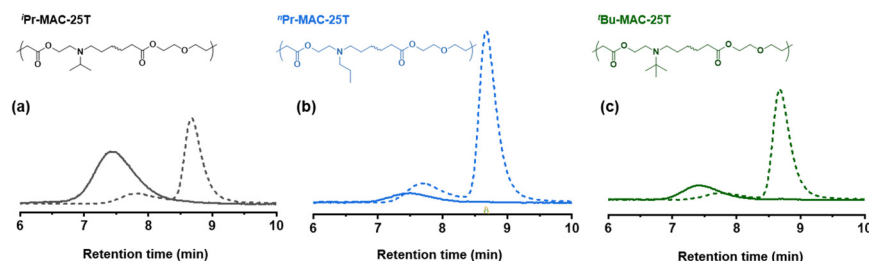
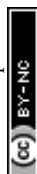


Fig. 4 SEC analysis of copolymers before (solid lines) and after hydrolytic degradation (dashed lines) under accelerated conditions. Light scattering signals of (a) ⁱPr-MAC-25T, (b) ⁱPr-MAC-25T and (c) ^tBu-MAC-25T are depicted.



Conclusions

Author contributions

Conflicts of interest

Acknowledgements

Deutsche Forschungsgemeinschaft (GA2051/7-1) for financial support. We thank the Howdle group at the University of Nottingham for granting access to their NMR facilities.

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