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Direct routes to functional RAFT agents from substituted *N*-alkyl maleimides†

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***N*-substituted maleimides have become an indispensable tool for the synthesis of bioconjugates and functional materials. Herein, we present three strategies for the incorporation of *N*-alkyl substituted maleimides into RAFT agents and show that these maleimide-derived CTAs can be used to easily introduce a range of chemical functionality at the β -position of polymer chains, resulting in α,β,ω -functional RAFT polymers. With both functional maleimides and RAFT agents that are increasingly available on the market, the approach presented in this study could facilitate the synthesis of end-functional macromolecules and will complement well the range of existing synthetic routes, including those utilising *N*-substituted maleimides, to functional polymeric materials.**

Introduction

Maleimides are versatile molecular tools utilised by chemists and biologists to engineer functional materials and (bio)conjugates. Due to their unique reactivity, maleimides are often part of ligation strategies to incorporate molecular fragments with specific functionalities – e.g. fluorophores, spin probes, polymer chains and drugs – into (bio)molecules with complementarily reactive residues.^{1,2} This is typically done through (i) thio-Michael and (ii) Diels–Alder reactions, and (iii) use of *N*-substituted maleimides as comonomers in radical polymerisations. Due to its high efficiency and, under specific conditions, selectivity, thio-Michael reaction between maleimide

and sulfhydryl-containing (bio)molecules is often utilised as the ligation strategy for the synthesis of bioconjugates. For example, 8 of the 11 currently approved antibody drug conjugates (ADCs) used as anticancer therapeutics,^{3,4} as well as clinically PEGylated Fab' fragment Certolizumab pegol, targeting tumour necrosis factor alpha (TNF- α) and clinically utilised to treat a range of inflammatory autoimmune diseases,⁵ are generated through maleimide-cysteine coupling. Moreover, the ability of maleimides to act as electron-deficient dienophiles in Diels–Alder (DA) reactions resulted in their use to produce a variety of functional low molar mass intermediates,⁶ bioconjugates,⁷ and polymer-based materials.⁸

In radical polymerisations, *N*-substituted maleimides possess a rather peculiar reactivity, in that they have a little or no tendency for homopolymerisation, but very rapidly cross-propagate in the presence of electron-rich monomers such as styrenes.⁹ This unique behaviour has been exploited in reversible-deactivation radical polymerisations (RDRP) to insert monomers at predetermined positions of polymer chains, where *N*-substituted maleimides are added at definite time points during the polymerisation reaction and, due to their preferential crossover propagation, they are very rapidly incorporated into specific short segments of the growing polymer chains.¹⁰ The Lutz group have pioneered this approach and very elegantly utilised *N*-substituted maleimides to control monomer sequence distribution in polymers synthesised by RDRP, inserting functionalities at precisely defined points of polymer chains.^{9,11–21} For example, 1-alkyne and activated ester functionalities have been used to generate polymer chains with defined intramolecularly crosslinked regions,¹³ including well-defined structures such as α -shaped²¹ and more complex¹⁹ macromolecular origamis. Bulky substituents such as functional dendrons¹⁵ and, following post-polymerisation modification, poly(ethylene oxide) side chains¹¹ can also be introduced within specific segments of polymer chains. Harrison and co-workers have then investigated the level of monomer insertion precision that can be obtained using chain polymerization, and estimated that improved control

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can be attained for longer chains, controlling the relative position of a specific monomer with respect to the total chain length.²²

Taking advantage from their very low tendency to homopolymerize, Stayton and co-workers successfully inserted single *N*-substituted maleimide units at the ω (*Z*)-terminal chain end of RAFT polymers with very good efficiencies.²³ Xu *et al.* utilised photoinduced electron/energy transfer (PET) to develop a single unit monomer insertion (SUMI) strategy based on RAFT polymerisation, where three different monomers, including *N*-substituted maleimides, are sequentially added, to generate well-defined trimers with high control over monomer sequence.²⁴ Longer discrete and stereospecific oligomers were subsequently synthesised by Xu and Boyer following a similar approach, under red light irradiation and ZnTPP as the photocatalyst, using indenes and *N*-substituted maleimides as electron donor and electron acceptor monomers, respectively.²⁵

RAFT polymerisation is ideally suited for the synthesis of heterotelechelic polymers, that is, polymer chains with different functionalities at their α and ω chain-ends, which are derived from the R and Z groups of the chain-transfer agents (CTAs) chosen for their synthesis.^{26–28} However, for specific applications an additional functionality may be required at one of the two polymer chain-ends, to give α,β,ω -functional polymer chains. In this work, we addressed this need and describe three synthetic routes where functional *N*-alkyl maleimides are utilised to insert specific functionalities between the R and Z groups of trithiocarbonate CTAs. Following characterisation of these CTAs, we then show that they can be successfully employed to generate α,β,ω -functional polymers using acrylates and acrylamide model monomers. We believe that our approach complements well the existing routes to functional heterotelechelic (co)polymers and, with the increasing number of *N*-substituted maleimides that are increasingly becoming available it could become a valuable strategy among the wide toolbox of synthetic strategy to produce functional polymeric materials.

Results and discussion

Design of *N*-alkyl maleimide-derived chain-transfer agents

To produce the required maleimide-derived chain-transfer agents (**Mal-CTAs**) the three synthetic strategies illustrated in Fig. 1 were designed. These routes rely on a single unit monomer insertion approach, which involves the addition of a single monomer molecule to a preformed RAFT agent,²⁹ or a bis(thioacyl) disulfide. In Route 1a, oxidative coupling of trithiocarbonate salts (**D**)³⁰ using $K_3Fe(CN)_6$ or I_2 afforded the bis(thioacyl) disulfides (**A**). Reaction of the latter with 1.8 equivalent of AIBN, used here as a representative model azo-initiator, and *N*-alkyl maleimides (**B**) at 70 °C in toluene, afforded the desired (**Mal-CTAs**) (chemical structures shown in Chart 1).

Route 1b starts from a preformed trithiocarbonate CTA (**C**), thus it is particularly suited for use with commercially available RAFT agents, *e.g.* 2-cyano-2-propyl dodecyl trithiocarbo-

nate CTA (**7**) used in this work. These can be converted into the final *N*-alkyl maleimide-derived chain-transfer agents **Mal-CTAs** in one step, by reaction with 0.05 eq. of a radical azo initiator. With an increasing number of trithiocarbonate CTAs, as well as bis(thioacyl) disulfides, that are now becoming available on the market, strategies 1a and 1b provide a direct access to the desired **Mal-CTAs** from readily accessible precursors. If required, trithiocarbonate CTAs starting materials (**C**) can also be obtained from bis(thioacyl) disulfides (**A**) by treatment with a radical azo initiator.²⁹ However, as already discussed, in this case Route 1a allows to bypass this extra step, affording the desired *N*-alkyl maleimide-derived **Mal-CTAs** directly from bis(thioacyl) disulfides precursors (**A**).

Route 2 starts from trithiocarbonate salts (**D**), prepared from thiols and CS_2 , again following well-established protocols²⁹ (ESI, Scheme S1†). Reaction of these with an α -halogen propionic ester, in this case methyl 2-bromopropionate, afforded the intermediates with general structure (**E**), which were then treated with 0.05–0.010 eq. of AIBN to give the desired *N*-alkyl maleimide-derived **Mal-CTAs**.

Fig. 1 (bottom) also shows the chemical structure of all intermediates utilised in this work. Hydrophobic (**10a**) and hydrophilic (**10b**) and (**10c**) *N*-alkyl substituted maleimides were chosen for these proof-of-principle experiments as model maleimide monomers (Fig. 1). (**10a**) possesses a diglyceride-like structure, and its incorporation at the β position results in polymers with two fatty acid ester moieties at their chain-end. These materials possess a molecular structure largely analogous to that of α -diglyceride polymers described by Hawker and co-workers,³¹ and to that of phospholipid-terminated polymers utilised as membrane-inserting macromolecules for the modification of artificial lipid bilayers,³² and plasma membranes of selected cells both *in vitro*^{33–36} and *in vivo*.³⁷ Maleimides (**10b**) and (**10c**) were chosen as representative example of a hydrophilic maleimide monomer, with free hydroxyl groups available for further chemical modification either prior or after incorporation into the desired functional polymers. Both maleimide derivatives were synthesised by reacting a furan-maleic anhydride Diels–Alder adduct with an appropriate amine in methanol in the presence of Et_3N , followed by removal of furan protecting group by retro Diels–Alder reaction in refluxing toluene, to give the maleimide monomers (**10a**), (**10b**), (**10c**) used in this study (ESI, Scheme S2†).

Hydrophilic and hydrophobic bis(thioacyl) disulfides (**3**) and (**6**), respectively, were used for Route 1a. It should be noted that although in this study bis(dodecylsulfanylthiocarbonyl) disulfide (**6**) was synthesised as described in the ESI,† this intermediate is also commercially available, which further simplifies access to the desired **Mal-CTAs**. The same consideration also applies to 2-cyano-2-propyl dodecyl trithiocarbonate CTA (**7**), which was chosen as the starting point for Route 1b.

Route 2 started from sodium 2-hydroxyethyl carbonotrithioate (**1**) and sodium dodecyl carbonotrithioate (**4**), which following reaction with 2-bromo methylpropionate in acetone at ambient temperature gave the trithiocarbonate CTAs (**2**) and



Maleimide-derived chain-transfer agents

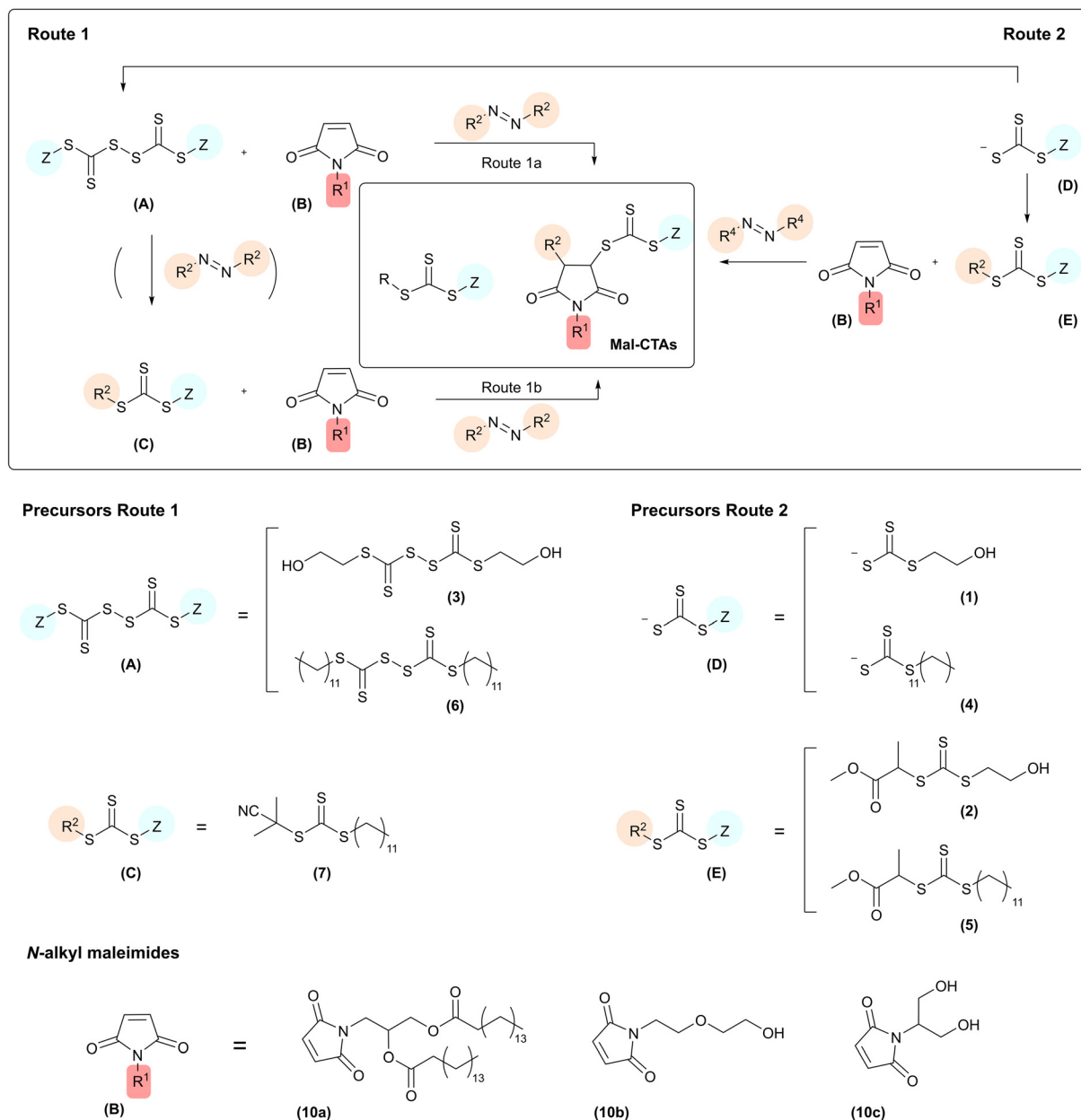


Fig. 1 Top: Synthetic routes to *N*-alkyl-substituted maleimide-derived chain-transfer agents (**Mal-CTAs**) synthesised in this study. Bottom: precursors utilised to produce the **Mal-CTAs**.

(5) respectively. Finally, treatment of the latter with 0.05–0.010 eq. of AIBN at 70 °C in the presence of *N*-alkyl substituted maleimides (**10a**) and (**10b**) afforded the required maleimide-derived chain-transfer agents (**Mal-CTAs**). The chemical structure of all **Mal-CTAs** synthesised in this work is shown in Chart 1. Overall yields for isolated **Mal-CTAs** are in the 22–73% range (Tables S1 and S2[†]), which is in line with previous work by Coote, Boyer, Xu and co-workers, who reported an overall yield of 56% for a single insertion of *N*-phenylmaleimide (PMI) into *n*-butyl benzyl trithiocarbonate (BBTC) CTA, using zinc tetraphenylporphyrin (ZnTPP) as the photocatalyst, after 20 h of

red-light irradiation ($\lambda_{\text{max}} = 635 \text{ nm}$) and subsequent purification by silica chromatography. These authors also reported that further increasing the [maleimide]:[CTA] ratio not only did not improve the reaction yield, but rather led to more by-products due to multiple maleimide monomer insertions.²⁵ For the synthesis of (**11a–c**) and (**12a–b**), the complex pattern of signals in the ¹H NMR spectra of these RAFT agents do not allow to easily detect potential traces of azo initiator-derived by-products in the crude reaction mixtures, although their presence before the purification steps in principle cannot be ruled out.



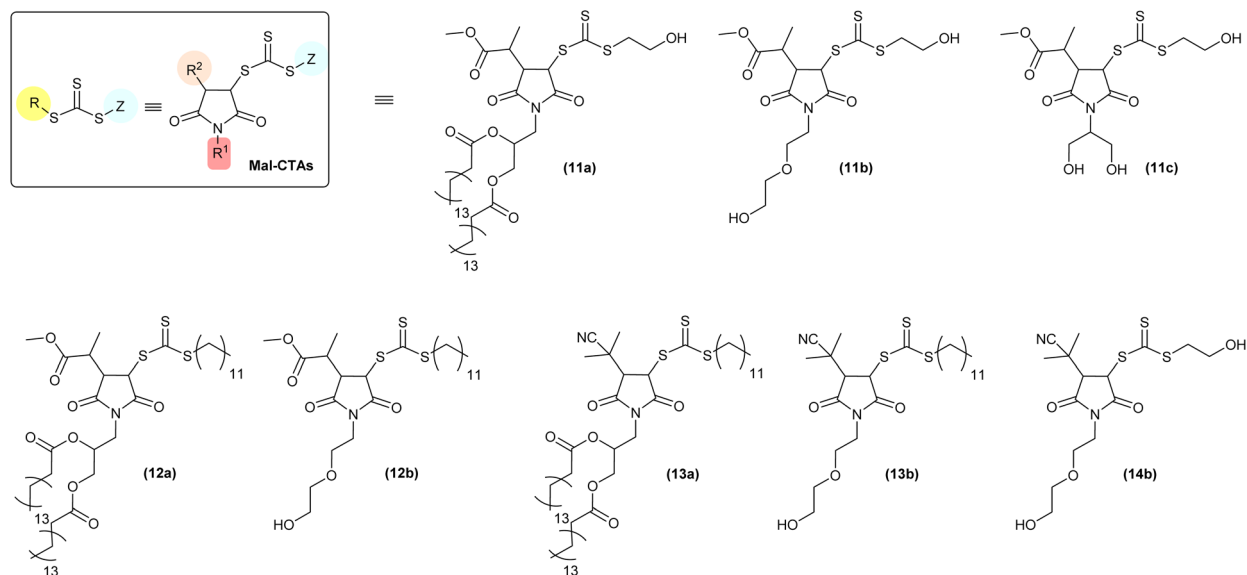


Chart 1 Maleimide-derived chain-transfer agents (Mal-CTAs) synthesised in this work.

In terms of reaction mechanism and characterisation of the final **Mal-CTA**, the two sp^2 -hybridised CH vinyl carbon atoms of maleimide precursors are prochiral, that is, each can be converted into a chiral centre by addition of a further substituent. ^1H NMR analysis of **(13b)** is discussed here as a representative example, as this **Mal-CTAs** does not possess any chiral centres in R group of the precursor, or in the *N*-alkyl substituent R^1 of the maleimide monomer, which would result in more complex spectra due to the presence of additional stereoisomers.

In principle, addition of two substituent to maleimide vinyl group could occur in *cis* (*syn*)- or *trans* (*anti*)-fashions. The presence of a single pattern of signals for $-\text{CHSC}(\text{S})\text{SZ}-$ ($\delta = 4.72$ ppm, d, $J_{\text{Hf-Hg}} = 6.2$ Hz, H_f) and $-\text{CHC}(\text{CH}_3)_2\text{CN}$ (CHR , $\delta = 3.22$ ppm, d, $J_{\text{Hg-Hf}} = 6.2$ Hz, H_g) groups in the ^1H NMR spectrum of **(13b)** indicated that addition of R and Z groups of its CTA precursor (or R group from the azo initiator and Z from bis(thioacyl) disulfides) to the maleimide vinyl group was stereoselective, leading exclusively to a single pair of enantiomers (Fig. 2, top). Although experiments such as single-crystal X-ray diffraction to assign the absolute configuration of the stereocentres of the adducts were not carried out in this study, both reaction mechanism and ^1H NMR spectral features of our **Mal-CTAs** closely resemble those of the *trans*-(BBTC-PMI) adduct described by Coote, Boyer, Xu and co-workers²⁵ strongly suggest that **(13b)** exists as a mixture of *trans*-stereoisomers, namely enantiomers (*R,S*) and (*S,R*), at their newly generated saturated 5-membered rings. Moreover, the J coupling constant $J_{\text{Hf-Hg}} = 6.2$ Hz observed for the two CH at the newly generated stereogenic centres is in line with those reported in previous studies ($J = 6.4$ – 6.5 Hz) for *trans*-adducts formed by reaction between substituted maleimides and xanthates,^{38–40} further suggesting a *trans*-configuration for **(13b)** and the other **Mal-CTAs** synthesised in this work.

This is consistent with a mechanism by which a 2-(2-methylpropanenitrile) radical is first added to the maleimide ring, providing significant steric hindrance on the face at which this first addition occurred. As a result, addition of the second substituent occurs at the less sterically encumbered opposite face of the 5-membered rings, yielding a *trans* (*anti*)-adduct. ^1H and ^{13}C NMR spectra of all **Mal-CTAs**, along with 2D COSY and HSQC spectra are shown in the ESI, Fig. S42–S73.† Conversely when a relatively less hindered 2-methylpropanoate group was added to the maleimide vinyl group the presence of a small amounts of the minor *cis* (*syn*) adduct was also observed (Fig. 2, bottom), with signals for the $-\text{CHSC}(\text{S})\text{SZ}$ proton in their ^1H NMR spectrum shifted at lower fields, and with larger J coupling values of around 9 Hz, consistent with a smaller torsion angle (5 – 10°) for the *cis* (*syn*) isomer. NOESY analysis further supported this assignment, by showing that, for the *trans* (*anti*), but not *cis* (*syn*) isomer, the $-\text{CHSC}(\text{S})\text{SZ}$ proton is spatially close to the methyl group of the 2-methylpropanoate substituent (Fig. S50 and S58†).

Next, to confirm their suitability to mediate RAFT polymerisations, selected **Mal-CTAs** were utilised for the polymerisation of (i) *n*-butyl acrylate (BA) and (ii) *N*-hydroxyethylacrylamide (HEA), 4-acryloylmorpholine (NAM), and *N,N*-dimethylacrylamide (DMA) as model acrylate and acrylamide monomers, respectively, initially at a $[\text{monomer}]_0 : [\text{Mal-CTA}]_0 = 50 : 1$ (Fig. 3). Polymerisations of BA were carried out using the relatively hydrophobic **Mal-CTAs** **(11a)** and **(12a)** in DMF, using AIBN as the radical initiator, whereas for HEA, NAM and DMA hydrophilic acrylamide monomers and more hydrophilic CTA **(11b)** Perrier's ultrafast RAFT conditions,⁴¹ with VA-044 azo initiator in water/1,4-dioxane, were chosen (Table S3†). All polymerisations were found to be controlled, with molar mass distributions $\mathcal{D} < 1.20$ (Table 1, runs 15–19). Using *N,N*-dimethylacrylamide (DMA) as a representative acrylamide monomer,



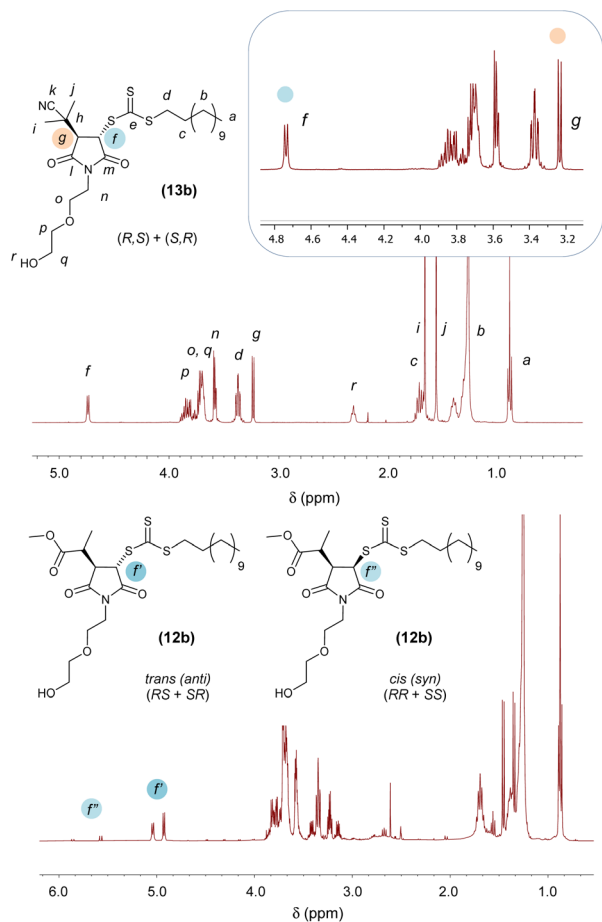


Fig. 2 Top: ^1H NMR spectrum of racemic *trans* (*anti*)-adduct Mal-CTA (**13b**) in CDCl_3 . Inset: magnification of the 3.2–4.8 ppm region, highlighting the signals of $-\text{HCSC}(\text{S})\text{SZ}$ and $-\text{CHC}(\text{CH}_3)_2\text{CN}$ protons at the stereogenic centres. Bottom: ^1H NMR spectrum of adduct Mal-CTA (**12b**) in CDCl_3 , showing the presence of minor *cis* (*syn*) adduct – *anti* : *sin* 92 : 8. Signal splitting is due to the additional chiral centre in the 2-methylpropionate substituent.

polymers with increasing DP – $[\text{monomer}]_0 : [\text{Mal-CTA}]_0 = 50, 100, 200,$ and $500 : 1$ – were then synthesised.

A slight shoulder at low molecular masses was observed in the SEC traces of the two shorter polymers (**11b**)-DMA₃₉ and (**11b**)-DMA₉₇, potentially suggestive of either a small proportion of early irreversible termination, or slow consumption of the RAFT agent in the pre-equilibrium step. (Mal-CTA)-DMA_{*n*} materials with higher molar mass (**11b**)-DMA₁₈₄ and (**11b**)-DMA₄₇₅ had comparably more symmetrical traces, and again good control over molar mass dispersity was achieved (Table 1, runs 19–22).

To further examine the controlled nature of these polymerisations, we periodically sampled a DMA polymerisation, mediated by (**13b**) at 15, 30, 60, 120 and 360 min. Consistent with conventional RAFT agents, the polymerisation with Mal-CTA (**13b**) exhibited a linear evolution of molar mass with conversion. SEC chromatograms displayed narrow molar mass distributions for each time point ($D < 1.20$, Fig. S84[†]). Furthermore, the first order kinetic plot also exhibited the typical linear increase then plateauing due to termination, with a small acceleration between 120 and 360 min likely due to the gel effect at high conversion.

To extend the range of monomer types and Mal-CTAs evaluated, we also conducted a polymerisation using (**13b**) with methyl methacrylate (MMA) targeting a DP of 125. Interestingly the resulting polymer displayed a broad molar mass distribution ($D = 2.05$) and $M_{n,\text{SEC}}$ (51.0 kg mol^{-1}) significantly higher than the theoretical molar mass (10.5 kg mol^{-1}), suggesting methacrylate polymerisations may not be controlled with this Mal-CTA (Fig. S85 and Table S4[†]). We theorised this was due to the insertion of the maleimide between the typically stabilising nitrile based reinitiating group and the trithiocarbonate. This would lead to a more stable radical on the MMA initiator derived chains than the maleimide reinitiating group on the Mal-CTA resulting in poor CTA participation in the polymerisation and thus high molar masses with high dispersity. To investigate this, we conducted an identical MMA polymerisation with CTA (**7**), the precursor to Mal-CTA (**13b**). In this case, the polymerisation was well controlled, displaying low dispersity ($D = 1.19$) while the

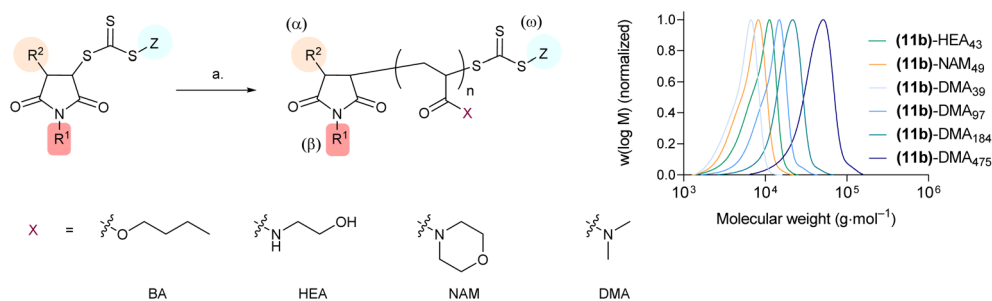


Fig. 3 Synthesis and SEC characterisation of α,β,ω -functional polymers produced from maleimide-derived CTAs. Reagents and conditions. a. for (Mal-CTA)-(BA)_{*n*} – $[\text{BA}]_0 : [\text{Mal-CTA}]_0 = 50 : 1$, DMF, AIBN, 80 °C, 90 min; for (Mal-CTA)-(HEA)_{*n*} and (Mal-CTA)-(NAM)_{*n*} – $[\text{monomer}]_0 : [\text{Mal-CTA}]_0 = 50 : 1$, water : 1,4-dioxane, VA-044, 100 °C, 15 min; for (Mal-CTA)-(DMA)_{*n*} – $[\text{DMA}]_0 : [\text{Mal-CTA}]_0 = 50, 100, 200, 500 : 1$, DMF water : 1,4-dioxane, VA-044, 75 °C, 15 min. HEA = *N*-hydroxyethylacrylamide, NAM = 4-acryloylmorpholine, DMA = *N,N*-dimethylacrylamide. Right: SEC traces for the analysis of (Mal-CTA)-(HEA)_{*n*}, (Mal-CTA)-(NAM)_{*n*}, and (Mal-CTA)-(DMA)_{*n*} polymers, in DMF + 0.1% LiBr as the mobile phase. (Mal-CTA)-(BA)_{*n*} were analysed using CHCl_3 as the mobile phase and are shown in the ESI.[†]



Table 1 Composition and molar mass distribution of the maleimide-based polymers synthesized in this study

Code	Polymer	Mal-CTA	Monomer	$[M]_0 : [Mal-CTA]_0$	Conversion ^a (%)	DP _{theo} ^b	$M_{n, SEC}$ (kg mol ⁻¹)	D
15	(11a)-BA ₄₈	(11a)	BA	50	97	48	7.2 ^c	1.17 ^c
16	(12a)-BA ₃₄	(12a)	BA	50	69	34	7.6 ^c	1.15 ^c
17	(11b)-HEA ₄₃	(11b)	HEA	50	87	43	7.8 ^d	1.1 ^d
18	(11b)-NAM ₄₉	(11b)	NAM	50	98	49	6.1 ^d	1.17 ^d
19	(11b)-DMA ₃₉	(11b)	DMA	50	79	39	5.1 ^d	1.15 ^d
20	(11b)-DMA ₉₇	(11b)	DMA	100	97	97	10.7 ^d	1.20 ^d
21	(11b)-DMA ₁₈₄	(11b)	DMA	200	92	184	16.9 ^d	1.17 ^d
22	(11b)-DMA ₄₇₅	(11b)	DMA	500	95	475	35.8 ^d	1.24 ^d

^a Determined by ¹H NMR, by comparing the area of the vinyl signals of residual monomer with that of polymer pendant chains. ^b Theoretical degree of polymerisation was calculated from monomer conversion and initial $[M]_0 : [Mal-CTA]_0$. ^c Estimated by SEC analysis, using chloroform as the mobile phase and PMMA narrow standards. ^d Estimated by SEC analysis, using DMF + 0.1 w/w% LiBr as the mobile phase and PMMA narrow standards.

theoretical ($M_{n, th} = 10.5$ kg mol⁻¹) and experimental ($M_{n, SEC} = 13.2$ kg mol⁻¹) were in closer agreement (Fig. S84 and Table S4†). Overall, this highlights a potential limitation for the use of these **Mal-CTAs** to mediate the polymerisation of methacrylate monomers, while being able to efficiently control acrylate and acrylamide polymerisations.

Conclusions

In summary, in this study we present three general strategies for the synthesis of functional chain-transfer agents starting from *N*-alkyl substituted maleimides. Typically heterotelechelic polymers can be produced by RAFT polymerisation by using chain-transfer agents with functional R groups and subsequently modifying the thiocarbonylthio group at the ω -chain-end.^{27,42} By using appropriate *N*-substituted maleimides and common RAFT agents and precursors, the simple approach we present allows for a facile introduction of an additional residue at the β -position of a polymer chain, resulting in α, β, ω -functional polymers, potentially further expanding the range of applications of RAFT polymers. Moreover, an increasing number of maleimide-containing derivatives are becoming available, largely driven by their application in biology and conjugation chemistry. At the same time, many RAFT agents are also now available on the market. Our strategy can therefore allow for the production of a range of functional chain-transfer agents – incorporating for example fluorophores, ligands, peptide/proteins or drug molecules – in one simple step starting from commercially available maleimide derivatives and RAFT agents. We believe that the strategies presented in this study complements well the existing techniques which utilise *N*-substituted maleimides for the synthesis of functional polymers and will further expand the palette of tools available to polymer chemists for the design of functional CTAs and resulting polymeric materials.

Author contributions

G. M., S. S. and R. C. designed the experiments. R. C. carried out the experiments with the assistance of E. R., F. M., and

L. S. for the synthesis of **Mal-CTAs**, R. F. for the synthesis and characterisation of Mal-CTA-derived polymers, P. G. for the polymerisation reactions, K. B. and H. W. for the NOESY experiments. G. M., S. S. and R. C. wrote, reviewed, and edited the manuscript.

Conflicts of interest

There are no conflicts to declare.

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