



Cite this: *Polym. Chem.*, 2015, **6**, 1944

Templated polymerizations on solid supports mediated by complementary nucleoside interactions†

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Template-directed radical polymerizations on solid supports are presented, which take advantage of complementary nucleoside interactions in non-polar solvents. The templates are prepared through copper-mediated living radical polymerization allowing control over the length and dispersity of the polymer bound to the support. We report a degree of control over subsequent polymerization of a complementary base pair monomer in the presence of the template not observed in its absence. Templates achieved from uridine, adenosine, cytidine and guanosine substituted methacrylate monomers were successfully prepared. Uridine derived templates were found to provide good fidelity of replication by incorporating mainly adenosine monomer in the final polymer, from a mixture of monomers. The latter was separated from its template and characterized demonstrating the preparation of polymers of approximately half of the template length. A model for the templating process is proposed.

Received 22nd December 2014,
Accepted 26th December 2014

DOI: 10.1039/c4py01783h

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Introduction

The controlled, sequence-specific replication of DNA using a template-directed mechanism is central to life.¹ The apparent simplicity of this templating process has inspired synthetic^{2–6} and semi-synthetic polymerization systems^{7–14} that use hydrogen bonding as a key mediating interaction between generational strands.^{15,16} Indeed, studies using RNA¹⁷ and DNA^{18,19} have demonstrated that enzyme-free extension of primers is possible with remarkably high fidelity. In addition, small interfering- and micro-RNAs that are transcribed from DNA continue to be uncovered, together with novel mechanisms to understand and control templating and translation processes.²⁰ Hence, oligomeric and polymeric molecules that include functionality capable of interacting in a specific way with nucleic acids²¹ and other cellular targets²² are of great significance and continue to present synthetic, biophysical and biological challenges.

To date, various post-polymerization modification methods and polymerization techniques have been applied for the preparation of synthetic nucleic acid analogues as summarized by Inaki¹⁵ and Smith.²³ More recently McHale *et al.* high-

lighted achievements in this field with a focus on poly(nucleobase)s and poly(nucleoside)s macromolecules obtained by controlled (radical) polymerization techniques.²⁴ Amongst them, atom transfer radical polymerization (ATRP) was predominantly employed for different nucleobase-^{25,26} and nucleoside-based monomers^{27,28} and initiators.²⁹ In addition nitroxide-mediated radical polymerization (NMP)^{6,30} and reversible addition-fragmentation chain transfer (RAFT) polymerization³¹ have been reported to yield well-defined poly(nucleobase) systems.

Furthermore, a range of monomer units and template strands ranging from DNA,³² RNA or DNA partnered with thioester peptide nucleic acids (tPNAs),³³ to synthetic constructs such as ribonucleoside and nucleobase bearing acrylates²⁸ and methacrylates^{25,27,34} have been used to prepare templates for polymerization studies. Lynn and co-workers demonstrated the templated polymerization of amine-containing thymidine monomeric units in solution³² and on solid supports using reductive amination to form the polymer backbone.³⁵ For this purpose, complementary nucleobase-pairing interactions between a natural DNA template, namely (dAp)_n, and the synthetic monomer were exploited to enable sequence- and chain-length-specific polymerization. In other studies, synthetic templates prepared by, *e.g.* free radical polymerization,³⁶ NMP⁶ and ring-opening metathesis polymerization (ROMP) were used.³⁷ Well-defined daughter poly(nucleobase)s were obtained with chain lengths and dispersities similar to the template polymer³⁷ and high molar masses *via* segregation of propagating chains in discrete micelle cores, respectively.⁶

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† Electronic supplementary information (ESI) available: Experimental data of monomer synthesis, preparation of adenosine, cytidine and guanosine templates and templated polymerization. See DOI: 10.1039/c4py01783h



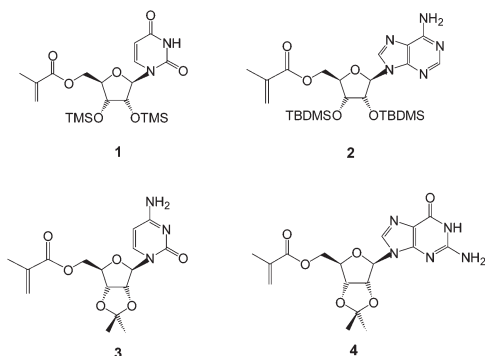


Fig. 1 Structures of the methacryloylribonucleosides prepared, including uridine (**1**),²⁷ adenosine (**2**),²⁷ cytidine (**3**) and guanosine (**4**) derivatives.

Following our initial report on templated ATRP of complementary nucleoside monomers²⁸ it became clear that a template, which allowed the resultant polymer–polymer complex to be separated more readily would be extremely beneficial, as would a template of controlled length and dispersity. In order to address these possibilities we opted to use ATRP, which we had previously shown to be compatible with the growth of methacryloyl nucleoside polymers both in solution²⁷ and from a solid support.³⁸

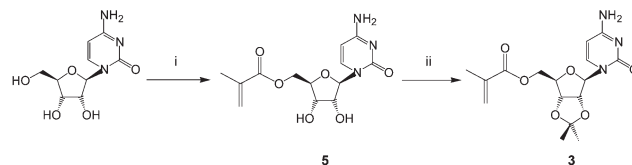
Herein, we describe the synthesis and characterization of the more challenging cytidine and guanosine ribonucleoside-based methacrylate monomers, to extend the uridine and adenosine materials prepared in our earlier work²⁷ (Fig. 1). Copper-mediated living radical polymerization was employed for the preparation of poly(methacryloyl nucleoside)s bound to polystyrene bead supports (Wang-type resin). The choice of this solid support allowed the cleavage of the polymer and thus its detailed characterization. Furthermore, the potential of the individual templates for templated polymerizations was investigated.†

Results and discussion

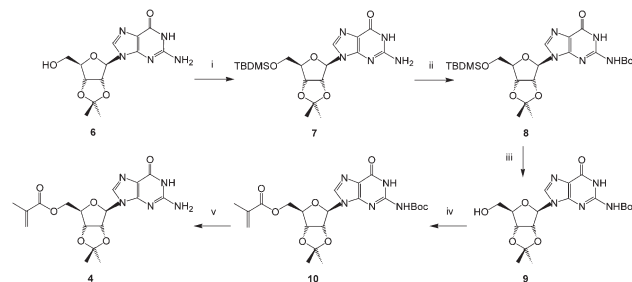
Synthesis of monomers

The monomers chosen for this study were the 5'-methacryloylribonucleosides (Fig. 1). The preparation of uridine **1** and adenosine **2** derivatives has been described previously²⁷ using an immobilized *Candida antarctica* lipase-promoted acylation reaction.³⁹ The cytidine **3** and guanosine **4** derivatives were prepared according to Schemes 1 and 2.

Cytidine was reacted with methacryloylacetone oxime in the presence of *C. antarctica* lipase⁴⁰ (Novozyme 435®) in dioxane and the product **5** was then protected as the 2',3'-acetonide⁴¹ **3** in reasonable yield (ESI†). Unlike Moris and Gotor's original report⁴⁰ we found this enzyme-promoted reaction⁴² to allow direct 5'-O-acylation of cytidine. The acetonide protecting group was synthetically accessible and found to provide accep-



Scheme 1 Synthesis of the cytidine derivative **3**: (i) methacryloylacetone oxime, Novozyme 435®, dioxane, radical inhibitor, 60 °C, 22 h (47%); (ii) *p*-toluenesulfonic acid, triethylorthoformate, acetone, r.t., 43 h (55%).



Scheme 2 Synthesis of the guanosine derivative **4**: (i) TBDMSCl, Et₃N, CH₂Cl₂, r.t., 72 h (88%); (ii) (Boc)₂O, DMF, DMAP, 60 °C, 48 h (46%); (iii) TBAF 1 M in THF, r.t., 2 h (99%); (iv) methacryloylacetone oxime, Novozyme 435®, dioxane, radical inhibitor, 60 °C, 24 h (78%); (v) TFA : CH₂Cl₂ (1 : 1), r.t., 2 h (58%).

table solubility in non-polar solvents in order to maximize templating hydrogen bonding and π -interactions. We found that the corresponding acrylate was not accessible by this route and required a lengthier protection sequence,⁴³ to be reported elsewhere. Attempts to acylate guanosine in a regioselective fashion using this enzyme were also unsuccessful under a variety of conditions and again necessitated protection of the amino function as follows (Scheme 2 and ESI†).

Guanosine was protected as the 2',3'-acetonide⁴³ **6** and then the 5'-hydroxyl protected as its TBDMS-ether **7**. The heterocyclic amine was subsequently protected with a Boc group to give **8** and the silyl ether removed using tetrabutylammonium fluoride furnishing **9**. Acylation of this newly exposed 5'-hydroxyl function using the methacryloylacetone oxime proceeded smoothly in the presence of Novozyme 435® giving the fully protected monomer **10**.

In contrast, attempts to use non-enzymatic acylation methods including the anhydride⁴⁴ or chloride led to intractable byproducts. The need to protect the exocyclic amine on the purine base is readily rationalized on the grounds that this group is more nucleophilic than the 5'-hydroxyl and may be better positioned to interact with the proposed serine-bound acylating reagent in the active site of the enzyme.⁴⁵ Finally, deprotection of the amine gave the methacryloylguanosine **4** in reasonable overall yield and high purity.

Preparation of solid-support polymer templates

The method chosen for the growth of the polymer on the support was copper-mediated living radical polymerization as

† Preliminary communication: M. Garcia, Ph.D. Thesis, University of Warwick, 2003.



used previously for the preparation of silica-supported poly(methacryloyl nucleoside)s.³⁸ In order to explore the scope of the template-directed polymerization on solid supports and exploit also the greater association constant of the G-C base pair as compared to the A-U pair, a full set of poly(methacryloyl nucleosides) bound to the resin were prepared. The supports used in the current work were polystyrene Wang-type resins that allowed acidic cleavage of the template polymer for analysis. The Wang initiator resin **11** (1.00 mmol g⁻¹ initial loading) was prepared by analogy with a method used for derivatisation of a PEG support⁴⁵ and it was possible to monitor the reaction by the appearance of the ester carbonyl stretch at 1734 cm⁻¹ in the infrared (IR) spectrum. Polymerizations of the four different monomers (1–4) with the Wang resin solid support initiator **11** were carried out using copper(i) bromide as a catalyst and *N*-(*n*-propyl)-2-pyridylmethanimine as ligand in toluene in which the Wang resin has good swelling properties (Scheme 3).⁴¹

In all cases, a mass increase after polymerization was observed, 180, 220, 240 and 120% for the polymerization of **1**, **2**, **3** and **4**, respectively. The polymerizations were followed by IR spectroscopy (Fig. 2A), observing the appearance of a broad NH signal (3600–3100 cm⁻¹) and a shift and an increase in the intensity of the carbonyl signal (1689 cm⁻¹). Furthermore, gel phase ¹H NMR provided valuable information about the successful polymerization, as exemplarily shown for poly(methacryloyl uridine) bonded to Wang resin (**12**) (Fig. 2).

In order to characterize the poly(methacryloyl uridine) further, it was cleaved from the resin by treatment with trifluoroacetic acid. It was found that the silyl protecting groups of the uridine and adenosine units were also removed under these conditions, making the product polymers insoluble in tetrahydrofuran (THF) and also not very soluble in dimethylformamide (DMF). Full re-protection of this polymer with TMS groups was not achieved. In contrast, the isopropylidene-protecting groups of the cytidine and guanosine moieties were

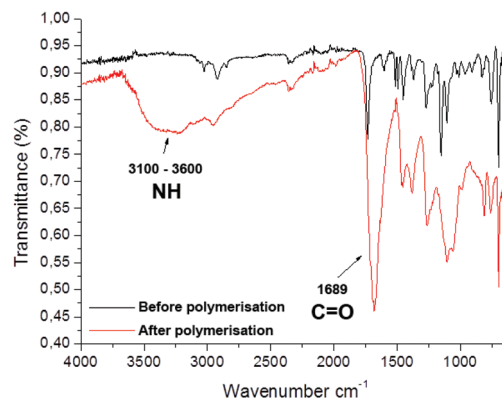
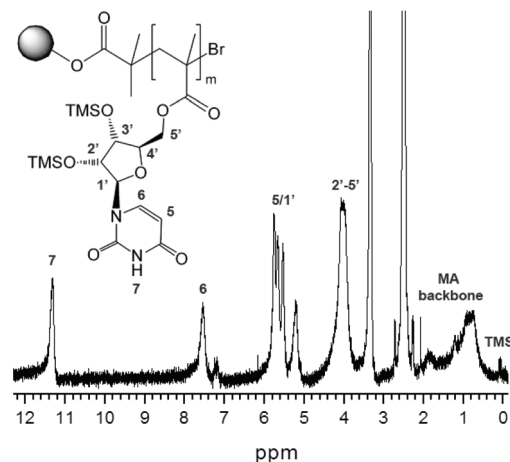
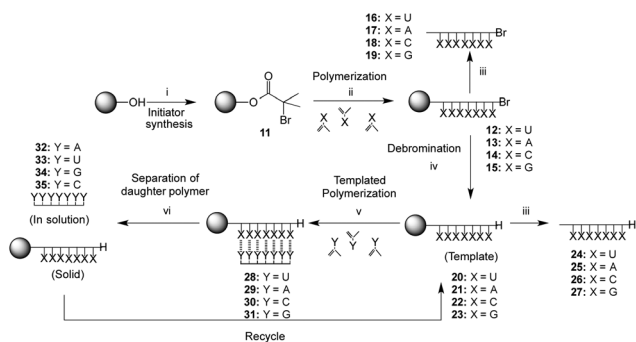


Fig. 2 Gel phase ¹H NMR (top) and IR (bottom) spectrum of poly(methacryloyl uridine) **12** bound to Wang resin.

not affected. In order to perform templated polymerizations, debromination of the resin bound polymers with tributyltin hydride and AIBN as initiator was deemed necessary, as the terminal bromide could inhibit radical propagation. The debromination of the polymer bound to the resin was followed by IR, which indicated the loss of the C–Br bond by the disappearance of the band around 700 cm⁻¹. Evidence for successful debromination of poly(methacryloyl cytidine) and poly(methacryloyl guanosine) was less assured since a C–Br peak was still discernible after treatment with tributyltin hydride in these cases. Prior to carrying out the template polymerization, the debrominated polymers were cleaved from the resin for full characterization. Reasonable molar masses and relatively low dispersities were found for uridine, cytidine and guanosine based polymers (Table 1), which is in accordance with a living mechanism for polymer growth.

Adenosine polymers **17** and **25** were found to be less well defined in agreement with observations made previously²⁸ using adenosine monomers. This could be due to (i) coordination of the copper(i) catalyst by amine moieties in monomer and polymer (ii) adenine's tendency to aggregate in solution and hydrogen bond to itself, including through Hoogsteen interactions, generating a less well solvated environment in which the polymerization takes place. For comparison, it has



Scheme 3 Solid support functionalization, template synthesis and templated polymerization of complementary monomer. 'X' and 'Y' represent the methacryloylribonucleosides (Fig. 1), 'Y' being the complementary ribonucleoside to 'X'; Reagents and conditions: (i) bromoisobutyryl-bromide, Et₃N, CH₂Cl₂, r.t., 22 h; (ii) Y, CuBr, *N*-(*n*-propyl)-2-pyridylmethanimine, toluene, 110 °C, 64 h; (iii) TFA/CH₂Cl₂ (1/1), r.t., 18 h; (iv) Bu₃SnH, AIBN, toluene, 100 °C, 20 h; (v) Y, AcOEt–toluene (2 : 1), AIBN, 60 °C, 42 h; (vi) CF₃CH₂OH, r.t., 1 h.



Table 1 SEC data for acid cleaved poly(methacryloyl uridine) **24**, poly(methacryloyl adenosine) **25**, poly(methacryloyl cytidine) **26** and poly(methacryloylguanosine) **19**

Polymer	M_n (g mol ⁻¹)	D	DP ^c
24 ^a	10 500	1.21	17
25 ^a	28 700	2.01	51
26 ^a	15 700	1.29	45
19 ^b	12 400	1.26	32

^a Measured by SEC (solvent: DMF). ^b Bromopolymer; it was not possible to prepare the debrominated polymer in sufficient quantity to allow for GPC assay. ^c DP calculated from M_n .

been shown by Rotello and co-workers that triazine polymers undergo folding due to intramolecular hydrogen bonding to give highly compact micelle-like structures.⁴⁶

Polymerizations in the presence of the solid-support polymer templates

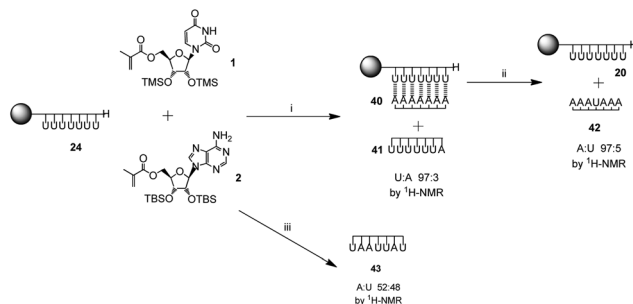
The prepared poly(nucleoside) modified resins, **20** (uridine), **21** (adenosine), **22** (cytidine) and **23** (guanosine), represent excellent candidates for the templated polymerization of the complementary nucleoside monomers, allowing for a straightforward separation of the template polymer from the resin. The complementary monomers were polymerized in the presence of the respective template resin under free radical conditions using AIBN as initiator. First, the silyl protected methacryloyl adenosine **2** was polymerized with the poly(methacryloyl uridine) template **20** to generate complex **28** with a ratio of adenosine to uridine of 1 : 1, as determined by gel phase ¹H NMR. It was possible to confirm the presence of the hydrogen bonds between these polymers by IR in the solid state, observing significant shifts in the amino N-H region [from 3223 (polymer) and 3260 (monomer) cm⁻¹ to 3396, 3207 and 3063 cm⁻¹] and carbonyl [from 1668 (polymer) and 1685 (monomer) to 1693 and 1634 cm⁻¹].

The daughter polymer **32**, poly(methacryloyl adenosine), was separated from the template **20** by simply washing the complex **28** with trifluoroethanol and precipitation in diethyl ether. ¹H NMR and IR of both the regenerated template resin **20** and the daughter polymer **32** showed that, firstly, the template **20** can be re-used for further template polymerizations and, secondly, poly(methacryloyl adenosine) **32** was obtained. As a control experiment, adenosine homopolymer **36** was prepared in solution from methacryloyl adenosine monomer **2** under the same conditions employed for the template polymerization showing the same ¹H NMR and IR spectrum as the daughter polymer **32**. SEC measurements revealed molar masses and dispersities lower than those found in the corresponding parent polymer, poly(methacryloyl uridine) **24** (Table 2).⁴⁷ The average DP of the poly(methacryloyl adenosine) **32** was approximately half that of the poly(methacryloyl uridine) **24**, possibly caused by chain transfer during the polymerization, although the fact that this is not observed with a 1 : 1 mixture of methacryloyl uridine and methacryloyl adenosine monomers (Scheme 4) suggests one of several possible effects.

Table 2 SEC data for poly(methacryloyl uridine) parent polymer **24**, the daughter polymer **32** and poly(methacryloyl adenosine) homopolymer **37**

Polymer	M_n (g mol ⁻¹)	D	DP ^c
24 ^a	10 500	1.21	17.1
32 ^a	3400	1.15	7.3
32 ^b	4300	1.60	7.7
37 ^a	8000	1.29	14.1
37 ^b	4500	4.38	7.9

^a Measured by SEC (solvent: DMF). ^b Measured by SEC (solvent: THF). ^c DP calculated from M_n .



Scheme 4 Control experiment: templated polymerization of methacryloyl adenosine **2** using template **24** in the presence methacryloyl uridine **1**: (i) AIBN, toluene–ethyl acetate (1 : 2), 60 °C, 24 h; (ii) CF₃CH₂OH, r.t., 30 min. Solution polymerization of monomer **1** and **2** (iii): AIBN, toluene–ethyl acetate (1 : 2), 60 °C, 24 h.

Firstly, a model whereby the propagating adenosine daughter polymer initially complexes to one site on the immobilized poly(methacryloyl uridine) template can be proposed. The interactions between template and growing daughter polymer are relatively weak at this stage with the most stable polymer complex formed by maximizing hydrogen bonding (Fig. 3). This use of Hoogsteen-type triplex hydrogen bonding, well-known for adenine⁴⁸ strengthens the interaction whilst effectively reducing the available length of the template by a factor of two as it folds onto the propagating polymer.

Secondly, another possible explanation might be that only the shorter uridine polymers unfold or are solvated sufficiently on the resin to act as a template, leading to the observed

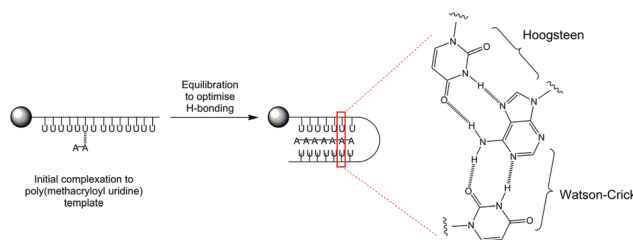


Fig. 3 Proposed templating mechanism with hydrogen-bonding interactions between poly(methacryloyl uridine) template and growing poly(methacryloyl adenosine) using Watson-Crick and Hoogsteen interactions.



shorter polyadenosine product in this case, although uridine does not significantly self-associate.

The templating process provided more control over the polymerization of methacryloyl adenosine **2** than the homopolymerization in absence of the template. This is evident when comparing the SEC analyses obtained for the daughter polymer **32** with the homopolymer **37** (Table 2). However, the inconsistency of these results with those obtained by gel phase ^1H NMR from complex **28** (a ratio of adenosine to uridine of 1 : 1) suggests, thirdly, that some polymer products may be retained on the support, despite the effective solvation provided by trifluoroethanol.

To further explore the limits of the templating effect of the immobilized poly(methacryloyl uridine) **20**, the polymerization was carried out both in the presence of the complementary monomer methacryloyl adenosine **2** and the monomer present in the template, *i.e.* the non-complementary methacryloyl uridine **1** (Scheme 4). Both monomers **1** and **2** were present in the same ratio and in equal amount to the accessible sites of the template **20**. An insoluble complex **40** was obtained which showed a ratio of uridine to adenosine of 1 : 1 by gel phase ^1H NMR. This was similar to the complex **28** obtained for the template polymerization of pure methacryloyl adenosine **2** and resembles the results obtained in solution for acryloyl monomers.²⁸ A soluble polymer **41**, which mostly contained uridine repeating units, was also obtained (uridine to adenosine 97 : 3 by ^1H NMR). Moreover, separation of the strands in complex **40** with trifluoroethanol showed a polymer **42** mostly composed of adenosine repeating units (adenosine to uridine 95 : 5 by ^1H NMR). The template **20** was also recovered. These results indicate that, as expected, poly(methacryloyl uridine) is a good template for the polymerization of methacryloyl adenosine with high selectivity.

When a statistical copolymerization was carried out in solution under the same conditions as those used for the template polymerization (monomers in equimolar amount in the presence of 3% AIBN with toluene as solvent), a copolymer **43** with ratio of uridine and adenosine of 52 : 48 by ^1H NMR was obtained (data not shown). Hence the observation that a greater proportion of adenosine is incorporated in the daughter polymer under templated conditions is good evidence that solid-supported polymerizations exert control through specific complementary ribonucleoside interactions over the polymerization process. This experiment is consistent with templating through a combination of both Watson-Crick and Hoogsteen interactions, depicted in Fig. 3 since incorporation of uridine would lack these stabilizing interactions.

Polymerization using immobilized poly(methacryloyl adenosine) **21** as template was performed with the complementary monomer, methacryloyl uridine **1**. Based on the increase in weight, similar yields were obtained as for the polymerization of methacryloyl adenosine **2** using immobilized poly(methacryloyl uridine) **20**. Furthermore, hydrogen bonds were observed between the polymers by IR. ^1H NMR and IR spectra of the poly(methacryloyl uridine) daughter polymer **33** were similar to those obtained of the cleaved immobilized

poly(methacryloyl uridine) **24**. SEC measurements of polymer **33** showed lower molar masses ($M_n = 2100 \text{ g mol}^{-1}$; $D = 1.7$) than those of its parent polymer **25**. This is in agreement with what was found for the acryloyl monomers²⁸ and it has been shown previously that the poly(acryloyl adenosine) shows a less good performance as template than poly(acryloyl uridine). Moreover, it appears that the complementary uridine by itself did not provoke sufficient unfolding of the adenosine polymer **21**. Inaki and co-workers have shown that polymers containing the adenine base are prone to intramolecular hydrogen bonding interactions.⁴⁹ When the poly(methacryloyl uridine) **36** was prepared by a non-templated polymerization process, SEC analysis of the corresponding homopolymer **36** showed a dispersity similar to that of polymer **33**. However, the molecular weight was higher ($M_n = 14\,400$; $D = 1.51$). This confirms that the presence of the poly(methacryloyl adenosine) as template inhibits the synthesis of poly(methacryloyl uridine).

In a similar manner poly(methacryloyl cytidine) **22** and poly(methacryloyl guanosine) **23** modified resins were employed for the template polymerization of their complementary monomers, methacryloyl guanosine **4** and methacryloyl cytidine **3**, respectively. IR measurements of the complexes demonstrated the presence of hydrogen bonds. Separation of the putative daughter polymers from the complexes afforded only unchanged monomers with no evidence of polymer. In both cases, template polymerization apparently did not take place under the conditions chosen and it seems the resins have inhibited the radical polymerization observed in the absence of the supported template. This effect seems not to be due to the presence of a bromine end-group on the resin bound polymer, nor specific to Wang resin since the radical polymerizations were carried out on resins treated with tributyltin hydride. It is entirely possible that self-association in both the cases of cytidine, where $(\text{C}\cdot\text{C})_n$ i-motif tetraplexes⁵⁰ are well recognized, especially in longer pyrimidine tracts, and guanosine, which is undoubtedly forming a variety of robust ribbon⁵¹ and quartet structures⁵² potentially inhibiting template effects. Other explanations include the isopropylidene protecting group adversely altering sugar or polymer backbone conformation.

Conclusions

The potential of immobilized poly(methacryloyl nucleosides) as templates for the polymerization of their complementary nucleoside monomers is presented for the first time. Methacryloyl cytidine and guanosine ribonucleoside monomers were prepared to add to uridine and adenosine, monomers available from our previous work and used in copper-mediated living radical polymerizations initiated by functional Wang resins. The poly(methacryloyl nucleoside) modified resins were characterized by IR and ^1H NMR spectroscopy. The choice of the resin enabled the acidic cleavage of the polymer and thus more complete characterization using solution NMR and SEC. All four immobilized polymers were applied in templated polymerizations of their respective complementary nucleoside



monomer exploiting complementary base pair interactions of uridine and adenosine as well as cytidine and guanosine. Poly-(methacryloyl uridine) templates were found to be suitable candidates for the templated polymerization of the complementary methacryloyl adenosine monomer. This result reflected what was observed in solution.²⁸ The great advantage of this approach, however, is the covalent linkage of the parent polymer to the solid support, which allows for a straightforward separation of the templated daughter polymer. The latter appeared to differ from the parent polymer in length and dispersity as measured by SEC, which was attributed to a dual hydrogen bonding complexation of the growing daughter polymer to the template through Watson-Crick, and possibly Hoogsteen-type, hydrogen bonding interactions. Competition experiments with non-complementary monomer and in the absence of the immobilized template, respectively, demonstrate the superiority of the template process. This particular templated polymerization shows that solid-supported polymerizations can provide good control over the polymerization process through specific hydrogen bonding interactions.

Our results contrast both with cellular replication in the presence of polymerases, transcription factors and other proteins which help maintain replicating oligonucleotide strands in their functional state⁵³ and with enzyme-free primer extensions.^{14,18,19} In summary, we demonstrate the utility of a polymer-supported template, and our experiments also highlight the potential for tuning solvent,⁵⁴ and chaperone molecules to create abiotic replicating systems.^{55,56}

Experimental

Materials and instrumentation

All reactions were carried out under an atmosphere of nitrogen using standard Schlenk techniques. The synthesis of 5'-methacryloyluridine **1**,²⁷ 5'-methacryloyl-adenosine **2**,²⁷ 5'-methacryloyl-cytidine **5**,²⁷ methacryloylacetone oxime,²⁷ Wang resin initiator **11**⁵⁷ and the ligand *N*-(*n*-pentyl)-2-pyridylmethanimine⁵⁸ were synthesized as previously reported and stored under anhydrous conditions prior to use. Copper(I) bromide (Aldrich, 98%) was purified according to the method of Keller and Wrcoff.⁵⁹ All other reagents were purchased from Aldrich or Lancaster Synthesis at the highest purity available and used without further purification unless otherwise stated. Solvents were of the highest grade available and were used as supplied. DSC was carried out on a Perkin Elmer Pyris 1 differential scanning calorimeter. TGA were recorded on a Perkin Elmer TGA 7 apparatus. Temperatures reported are T_{onset} . IR spectra were recorded on an Avatar 320 FT-IR fitted with a "Golden gate" attenuated total reflection (ATR) cell attachment. NMR spectra were recorded with a Bruker DPX 300. Chemical shifts (δ) are quoted in ppm using residual non-deuterated solvents as internal standard [d_6 -DMSO (^1H , δ : 2.50 ppm; ^{13}C , δ : 39.5 ppm)]. Coupling constants (J) are quoted in Hz. Molar mass distributions with PMMA standards in the range (200 to $6.85 \times 10^5 \text{ g mol}^{-1}$) used for specific calibration were measured

using size exclusion chromatography (SEC) at ambient temperature, on a system equipped with a guard column and on 2× mixed-C columns (Agilent) with differential refractive index detection using DMF as eluent, at a flow rate of 0.5 ml min^{-1} . Sample solutions were prepared by adding 2.0 ml of solvent to 4.0 mg of sample; leaving 72 h at 60 °C to dissolve. Molar mass distributions with PEO/PEG standards were carried out at RAPRA Polymer Laboratories. They were measured using size exclusion chromatography (SEC) at 80 °C, on a system equipped with a PLgel 2× mixed bed-B 30 cm and 10 μm columns with differential refractive index detection using DMF with 1% of lithium bromide as eluent, at a flow rate of 1.0 ml min^{-1} . Sample solutions were prepared by adding 10 ml of solvent to 20 mg of sample, leaving overnight and warming, if necessary, at 80 °C for 20 minutes to dissolve. After thorough mixing, the solutions were filtered through a 0.2 μm PTFE membrane prior to chromatography. Nomenclature of the compounds refers to a mixture of systematic and trivial nomenclature, which is in accordance with that used by other authors. The letters U, A, C and G during the text corresponds to methacryloyl uridine, methacryloyl adenosine, methacryloyl cytidine and methacryloyl guanosine, respectively. The letters U and A in the NMR refers to protons of uridine and adenosine, correspondingly. Yields of the polymerizations were determined gravimetrically.

Resin initiated polymerization of **1** (general procedure)

A mixture of the Wang initiator-resin **11** (300.0 mg, 0.30 mmol), 5'-*O*-methacryloyl-2',3'-*O*-trimethylsilyluridine **1** (1.37 g, 3.00 mmol) and copper(I) bromide (43.0 mg, 0.30 mmol) was placed in a Schlenk and de-aerated. The ligand *N*-(*n*-pentyl)-2-pyridylmethanimine (105.8 mg, 0.60 mmol) was added in de-aerated toluene (5.00 ml) and the reaction mixture was de-aerated four times using a freeze-pump-thaw cycle and then heated at 110 °C for 64 h. The solid was filtered, washed with water, methanol and acetone and dried *in vacuo* to afford the poly(methacryloyl uridine) substituted Wang resin **12** (849.3 mg, 120% weight increased) as a light brown solid; TGA 310.3, 355.8, 354.2, 461.0 and 540.0 °C; T_{onset} 244.5 °C; $\nu_{\text{max}}/\text{cm}^{-1}$ (solid): 3430 (N-H *st*), 2970 (C-H *st*), 1737 (C=O *st*), 1668 (C=O *st*), 1149 and 1103 (C-O *as st*), 1059 (C-O *sym st*), 698 (C-Br *st*); δ_{H} (d_6 -DMSO, gel phase, 300 MHz): 11.27 (1H, bs, NH), 7.58 [1H, bs, H(6)], 5.78–5.38 [2H, m, H(1') and H(5)], 4.33–3.78 [5H, m, H(2'–5')], 1.00–0.60 (5H, m, CH₃ and CH₂), 0.05 [9H, bs, Si(CH₃)₃], –0.05 [9H, bs, Si(CH₃)₃] ppm.

Cleavage of poly(methacryloyl uridine) from resin (general procedure)

Poly(methacryloyl uridine) substituted Wang resin **12** (95.1 mg) was swollen in a mixture of trifluoroacetic acid–dichloromethane (1 : 1, 4.0 ml). The solution was stirred at ambient temperature for 18 h, and then the resin was filtered and washed with more trifluoroacetic acid–dichloromethane (1 : 1, 3.0 ml). The filtrates were combined and cold diethyl ether was added (30 ml). The mixture was kept in the fridge



overnight to precipitate the product **16** which was filtered and washed with cold diethyl ether to afford the unprotected poly-5'-*O*-methacryloyluridine **16** (76. mg, 80%)⁶⁰ as a pale brown solid; TGA 98.0, 312.1 and 542.5 °C; T_{onset} 270.6 °C; $M_n = 7700$; $D = 1.19$ (SEC-DMF, PMMA calibrated); $\nu_{\text{max}}/\text{cm}^{-1}$ (solid): 3350 (N-H *st*), 3072 (C-H *st*), 1773 (C=O *st*), 1690 (C=O *st*), 1620 (C=C *st*), 1423 (N-H δ *ip*), 1338 (C-N *st*), 1139 and 1077 (C-O *as st*), 995 (C-O-C *st*); δ_{H} (d_6 -DMSO, 300 MHz): 11.39 (1H, bs, NH), 7.54 [1H, bs, H(6)], 5.76–5.65 [2H, m, H(1') and H(5)], 4.00–3.55 [5H, m, H(2'–5')], 1.13–0.76 (5H, m, CH₃ and CH₂) ppm.

Debromination of the poly(methacryloyl uridine) substituted Wang resin (general procedure)

A mixture of the poly-5'-*O*-methacryloyl-2',3'-*O*-trimethylsilyluridine substituted Wang resin **12** (600.0 mg, 388.1 mg polymer, 0.84 mmol polymer), tri-*n*-butyltin hydride (307.3 mg, 1.27 mmol), AIBN (18.0 mg) and de-aerated toluene (6.0 ml) was de-aerated in a Schlenk by four freeze–pump–thaw cycles. It was heated at 100 °C for 20 h. The resin was filtered, washed with water, methanol and acetone and dried *in vacuo* to give the debrominated poly-5'-*O*-methacryloyl-2',3'-*O*-trimethylsilyluridine substituted Wang resin **20** (546 mg) as a dark brown solid; TGA 313.7 and 408.1 °C; T_{onset} 230.1 °C; $\nu_{\text{max}}/\text{cm}^{-1}$ (solid): 1750 (C=O *st*), 1693 (C=O *st*), 1454 (N-H δ *ip*), 1252 (C-N *st*), 1152 and 1079 (C-O *as st*), 839 (C-O-C *st*); δ_{H} (d_6 -DMSO, gel phase, 300 MHz): 11.79 (1H, bs, NH), 7.85 [1H, bs, H(6)], 5.91–5.68 [2H, m, H(1') and H(5)], 4.03–3.58 [5H, m, H(2'–5')], 0.99–(–0.98) (23 H, m, 2Si(CH₃)₃, CH₃ and CH₂) ppm.

Cleavage of debrominated poly(methacryloyl uridine) from Wang resin

Following the general procedure with the debrominated poly(methacryloyl uridine) substituted Wang resin **20** (342.6 mg), and stirring in a mixture of trifluoroacetic acid–dichloromethane (1 : 1, 10.0 ml) at ambient temperature for 22 h to give the *title product* **24** (180 mg, 53%)⁶⁰ as a pale brown solid; TGA 189.9, 276.8, 304.7 and 411.5 °C; T_{onset} 247.5 °C; $M_n = 7800$, $D = 1.23$ (SEC-DMF, PMMA calibrated); $\nu_{\text{max}}/\text{cm}^{-1}$ (solid): 3223 (N-H and O-H *st*), 2970 (C-H *st*), 1668 (C=O *st*), 1464 (N-H δ *ip*), 1387 (C-N *st*), 1268, 1106 and 1077 (C-O *as st*); δ_{H} (d_6 -DMSO, 300 MHz): 11.39 (1H, bs, NH), 7.61 [1H, bs, H(6)], 5.93–5.66 [3H, m, H(1'), H(5) and OH(2')], 5.25 [1H, bs, OH(3')], 4.14–3.88 [5H, m, H(2'–5')], 1.78–0.74 (5H, m, CH₃ and CH₂) ppm.

Polymerization of **2** using an immobilized template of poly(methacryloyl uridine) (general procedure)

A solution of AIBN (0.3 ml of a 30.0 mg in 3.0 ml methanol solution) was placed in a Schlenk tube and the solvent removed *in vacuo*. 2',3'-*O*-*tert*-Butyldimethylsilyl-5'-*O*-methacryloyl adenosine **2** (159.7 mg, 0.28 mmol), the immobilized template of poly(methacryloyl uridine) **20** (200.0 mg, 129.7 mg polymer, 0.28 mmol polymer), ethyl acetate (4.0 ml) and de-aerated toluene (2.0 ml) were added. The reaction mixture was de-aerated four times using the freeze–pump–thaw cycle and heated at 60 °C for 42.5 h. It was then filtered and washed

with toluene and dried *in vacuo* to give the complex **28** (212 mg, 59%)⁶¹ as a cream colored solid. ¹H NMR (d_6 -DMSO, gel phase) of this complex **28** showed a 1 : 1 ratio of uridine to adenosine; TGA 336.5, 406.9 and 656.0 °C; T_{onset} 276.9 °C; $\nu_{\text{max}}/\text{cm}^{-1}$ (solid): 3396, 3207 and 3063 (N-H *hydrogen bonded*), 2928 and 2857 (C-H *st*), 1693 (C=O *st*), 1634 (C=O *st*), 1453 (N-H δ *ip*), 1373 (N-H *st*), 1249 (C-O-C *as st*), 1152 and 1077 (C-O *as st*), 836 (Si-CH₃), 751 (Si-C *st*); δ_{H} (d_6 -DMSO, gel phase, 300 MHz): 11.53 (1H, bs, NH), 8.34 [1H, bs, H(2A)], 8.08 [1H, bs, H(8A)], 7.44–7.16 [3H, m, NH₂ and H(6U)], 5.98–5.87 [1H, m, H(5U)], 5.59–5.32 [2H, m, 2H(1')], 4.50–3.70 [10H, m, 2H(2'–5')], 1.70–(–0.80) (58, m, 2Si(CH₃)₃, 2Si(CH₃)₂C(CH₃)₃, 2CH₃ and 2CH₂) ppm.

Separation of the daughter polymer poly(methacryloyl adenosine) **32** from its parent polymer the immobilised poly(methacryloyl uridine) **24** (general procedure)

The previous complex **28** (172.6 mg) was stirred in trifluoroethanol (5.0 ml) at ambient temperature for 1 h. It was then filtered and washed with more trifluoroethanol. The trifluoroethanol washings were removed under reduced pressure to give the poly-2',3'-*O*-*tert*-butyldimethylsilyl-5'-*O*-methacryloyl-adenosine **32** (48 mg, 30%)⁶² as a white solid. Analytical data for the solid supported parent polymer was identical to the parent polymer **24** prepared above. Analytical data of the daughter polymer **32** is as follows: TGA 349 °C; T_{onset} 275.7 °C; $M_n = 4300$, $D = 1.60$ (SEC-THF, PMMA calibrated); $M_n = 3400$, $D = 1.15$ (SEC-DMF, PMMA calibrated); $\nu_{\text{max}}/\text{cm}^{-1}$ (solid): 3330 (NH₂ *st*), 2931 (C-H *st*), 1694 (C=O *st*), 1636 (C=N *st*), 1472 (N-C=O *sym st* and NH *amide II*), 1251 (C-C-N *bending*), 1155 and 1078 (C-O *as st*), 834 (Si-CH₃), 776 (C-Si *st*); δ_{H} (d_6 -DMSO, 300 MHz): 8.38 [1H, bs, H(2)], 8.08 [1H, bs, H(8)], 7.29 (2H, bs, NH), 5.92–5.89 [1H, m, H(1')], 4.65–3.75 [5H, m, H(2'–5')], 1.30–(–0.6) (35H, m, 2Si(CH₃)₂C(CH₃)₃, CH₃ and CH₂) ppm.

Acknowledgements

We thank Dr Andrew Steward (University of Warwick) for help with polymer analyses. We thank EPSRC for a Fast Stream studentship (GR/L71933) held by MG and the University of Warwick for postdoctoral fellowships (AK, KK). MG thanks the Warwick-Wisconsin Exchange Fellowship for an award and we thank Professor S. H. Gellman (University of Wisconsin-Madison) in whose laboratory some of this work was carried out for additional support and guidance. DMH is a Royal Society Wolfson fellow.

Notes and references

- 1 J. D. Watson and F. H. C. Crick, *Nature*, 1953, **171**, 737–738.
- 2 G. Challa and Y. Y. Tan, *Pure Appl. Chem.*, 1981, **53**, 627–641.
- 3 K. Takemoto and Y. Inaki, *Adv. Polym. Sci.*, 1981, **41**, 1–51.
- 4 A. W. Schwartz and L. E. Orgel, *Science*, 1985, **228**, 585–587.
- 5 Y. Y. Tan, *Prog. Polym. Sci.*, 1994, **19**, 561–588.



- 6 R. McHale, J. P. Patterson, P. B. Zetterlund and R. K. O'Reilly, *Nat. Chem.*, 2012, **4**, 491–497.
- 7 L. E. Orgel, *Nature*, 1992, **358**, 203–209.
- 8 I. A. Kozlov, S. Pitsch and L. E. Orgel, *Proc. Natl. Acad. Sci. U. S. A.*, 1998, **95**, 13448–13452.
- 9 X. Y. Li and D. R. Liu, *Angew. Chem., Int. Ed.*, 2004, **43**, 4848–4870.
- 10 C. T. Calderone and D. R. Liu, *Curr. Opin. Chem. Biol.*, 2004, **8**, 645–653.
- 11 R. E. Kleiner, Y. Brudno, M. E. Birnbaum and D. R. Liu, *J. Am. Chem. Soc.*, 2008, **130**, 4646–4659.
- 12 Y. Brudno and D. R. Liu, *Chem. Biol.*, 2009, **16**, 265–276.
- 13 Y. Brudno, M. E. Birnbaum, R. E. Kleiner and D. R. Liu, *Nat. Chem. Biol.*, 2010, **6**, 148–155.
- 14 J. C. Blain, A. Ricardo and J. W. Szostak, *J. Am. Chem. Soc.*, 2014, **136**, 2033–2039.
- 15 Y. Inaki, *Prog. Polym. Sci.*, 1992, **17**, 515–570.
- 16 J. G. Schmidt, L. Christensen, P. E. Nielsen and L. E. Orgel, *Nucleic Acids Res.*, 1997, **25**, 4792–4796.
- 17 T. Inoue and L. E. Orgel, *Science*, 1983, **219**, 859–862.
- 18 E. Kervio, A. Hochgesand, U. E. Steiner and C. Richert, *Proc. Natl. Acad. Sci. U. S. A.*, 2010, **107**, 12074–12079.
- 19 E. Kervio, B. Claasen, U. E. Steiner and C. Richert, *Nucleic Acids Res.*, 2014, **42**, 7409–7420.
- 20 J. Kurreck, *Angew. Chem., Int. Ed.*, 2009, **48**, 1378–1398.
- 21 J. Niu, R. Hili and D. R. Liu, *Nat. Chem.*, 2013, **5**, 282–292.
- 22 L. Zhang, W. Liu, L. Lin, D. Chen and M. H. Stenzel, *Bio-macromolecules*, 2008, **9**, 3321–3331.
- 23 W. T. Smith, *Prog. Polym. Sci.*, 1996, **21**, 209–253.
- 24 R. McHale and R. K. O'Reilly, *Macromolecules*, 2012, **45**, 7665–7675.
- 25 H. J. Spijker, F. L. van Delft and J. C. M. van Hest, *Macromolecules*, 2007, **40**, 12–18.
- 26 H. J. Spijker, A. J. Dirks and J. C. M. Van Hest, *J. Polym. Sci., Part A: Polym. Chem.*, 2006, **44**, 4242–4250.
- 27 A. Marsh, A. Khan, D. M. Haddleton and M. J. Hannon, *Macromolecules*, 1999, **32**, 8725–8731.
- 28 A. Khan, D. M. Haddleton, M. J. Hannon, D. Kukulj and A. Marsh, *Macromolecules*, 1999, **32**, 6560–6564.
- 29 A. Likhitsup, S. Yu, Y.-H. Ng, C. L. L. Chai and E. K. W. Tam, *Chem. Commun.*, 2009, 4070–4072.
- 30 B. D. Mather, M. B. Baker, F. L. Beyer, M. A. G. Berg, M. D. Green and T. E. Long, *Macromolecules*, 2007, **40**, 6834–6845.
- 31 Y. Kang, A. Lu, A. Ellington, M. C. Jewett and R. K. O'Reilly, *ACS Macro Lett.*, 2013, **2**, 581–586.
- 32 Z. Y. Li, Z. Y. J. Zhang, R. Knipe and D. G. Lynn, *J. Am. Chem. Soc.*, 2002, **124**, 746–747.
- 33 Y. Ura, J. M. Beierle, L. J. Leman, L. E. Orgel and M. R. Ghadiri, *Science*, 2009, **325**, 73–77.
- 34 H. J. Spijker, A. J. Dirks and J. C. M. van Hest, *Polymer*, 2005, **46**, 8528–8535.
- 35 X. Li and D. G. Lynn, *Angew. Chem., Int. Ed.*, 2002, **41**, 4567–4569.
- 36 M. Hattori, H. Nakagawa and M. Kinoshita, *Makromol. Chem.*, 1980, **181**, 2325–2333.
- 37 P. K. Lo and H. F. Sleiman, *J. Am. Chem. Soc.*, 2009, **131**, 4182–4183.
- 38 A. Marsh, A. Khan, M. Garcia and D. M. Haddleton, *Chem. Commun.*, 2000, 2083–2084.
- 39 M. Ferrero and V. Gotor, *Chem. Rev.*, 2000, **100**, 4319–4348.
- 40 F. Moris and V. Gotor, *J. Org. Chem.*, 1993, **58**, 653–660.
- 41 R. W. Chambers, P. Shapiro and V. Kurkov, *J. Am. Chem. Soc.*, 1960, **82**, 970–975.
- 42 I. Lavandera, S. Fernandez, J. Magdalena, M. Ferrero, R. J. Kazlauskas and V. Gotor, *ChemBioChem*, 2005, **6**, 1381–1390.
- 43 R. W. Chambers, J. G. Moffatt and H. G. Khorana, *J. Am. Chem. Soc.*, 1957, **79**, 3747–3752.
- 44 M. J. Cooper, R. S. Goody, A. S. Jones, J. R. Tittensor and R. T. Walker, *J. Chem. Soc. C*, 1971, 3183–3187.
- 45 K. Jankova, X. Y. Chen, J. Kops and W. Batsberg, *Macromolecules*, 1998, **31**, 538–541.
- 46 R. Deans, F. Ilhan and V. M. Rotello, *Macromolecules*, 1999, **32**, 4956–4960.
- 47 The poly(methacryloyl adenosine) **32** obtained from the template polymerization was found to be very soluble in THF, so it was possible to perform full SEC analysis in both THF and DMF (PMMA as the standard).
- 48 R. K. Castellano, V. Gramlich and F. Diederich, *Chem. – Eur. J.*, 2002, **8**, 118–129.
- 49 Y. Inaki, K. Ebisutani and K. Takemoto, *J. Polym. Sci., Part A: Polym. Chem.*, 1986, **24**, 3249–3262.
- 50 C. H. Kang, I. Berger, C. Lockshin, R. Ratliff, R. Moyzis and A. Rich, *Proc. Natl. Acad. Sci. U. S. A.*, 1994, **91**, 11636–11640.
- 51 A. L. Webber, S. Masiero, S. Pieraccini, J. C. Burley, A. S. Tatton, D. Iuga, T. N. Pham, G. P. Spada and S. P. Brown, *J. Am. Chem. Soc.*, 2011, **133**, 19777–19795.
- 52 J. T. Davis and G. P. Spada, *Chem. Soc. Rev.*, 2007, **36**, 296–313.
- 53 T. H. Dickey, S. E. Altschuler and D. S. Wuttke, *Structure*, 2013, **21**, 1074–1084.
- 54 P. A. Korevaar, C. Schaefer, T. F. de Greef and E. W. Meijer, *J. Am. Chem. Soc.*, 2012, **134**, 13482–13491.
- 55 A. E. Engelhart and N. V. Hud, *Cold Spring Harbor Perspect. Biol.*, 2010, **2**, a002196.
- 56 S. I. Walker, M. A. Grover and N. V. Hud, *PLoS One*, 2012, **7**, e34166.
- 57 S. Angot, N. Ayres, S. A. F. Bon and D. M. Haddleton, *Macromolecules*, 2001, **34**, 768–774.
- 58 D. M. Haddleton, M. C. Crossman, B. H. Dana, D. J. Duncalf, A. M. Heming, D. Kukulj and A. J. Shooter, *Macromolecules*, 1999, **32**, 2110–2119.
- 59 R. N. Keller, H. D. Wrcoff and L. E. Marchi, in *Inorganic Syntheses*, John Wiley & Sons, Inc., 2007, pp. 1–4.
- 60 Based on the amount recovered from the resin.
- 61 Based on the amount of complex obtained and the amount of template and complementary monomer used.
- 62 Based on the amount of daughter polymer recovered and the amount of daughter monomer used for the templating process.

