

Cite this: *Chem. Sci.*, 2019, 10, 9057

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

Macrocycles of higher *ortho*-phenylenes: assembly and folding†

Zacharias J. Kinney,  Viraj C. Kirinda  and C. Scott Hartley *

Higher-order structure in abiotic foldamer systems represents an important but largely unrealized goal. As one approach to this challenge, covalent assembly can be used to assemble macrocycles with foldamer subunits in well-defined spatial relationships. Such systems have previously been shown to exhibit self-sorting, new folding motifs, and dynamic stereoisomerism, yet there remain important questions about the interplay between folding and macrocyclization and the effect of structural confinement on folding behavior. Here, we explore the dynamic covalent assembly of extended *ortho*-phenylenes (hexamer and decamer) with rod-shaped linkers. Characteristic ^1H chemical shift differences between cyclic and acyclic systems can be compared with computational conformer libraries to determine the folding states of the macrocycles. We show that the bite angle provides a measure of the fit of an *o*-phenylene conformer within a shape-persistent macrocycle, affecting both assembly and ultimate folding behavior. For the *o*-phenylene hexamer, the bite angle and conformer stability work synergistically to direct assembly toward triangular [3 + 3] macrocycles of well-folded oligomers. For the decamer, the energetic accessibility of conformers with small bite angles allows [2 + 2] macrocycles to be formed as the predominant species. In these systems, the *o*-phenylenes are forced into unusual folding states, preferentially adopting a backbone geometry with distinct helical blocks of opposite handedness. The results show that simple geometric restrictions can be used to direct foldamers toward increasingly complex folds.

Received 17th June 2019
Accepted 9th August 2019

DOI: 10.1039/c9sc02975c

rsc.li/chemical-science

Introduction

The structures of biomacromolecules demonstrate that molecular folding is an effective strategy for the generation of large-scale structural complexity. Decades of study have thus provided many examples of abiotic foldamers¹ that adopt well-defined secondary structures, ranging from peptides that hew closely to their biochemical inspirations to aromatic oligomers that offer distinctly different structures and properties.^{2–6} However, despite the importance of hierarchical structure in biochemistry, tertiary structure remains rare in abiotic foldamers, especially in the context of non-peptidic systems.^{7–11} That is, folding into helices, and to some extent other secondary structures,¹² is fairly common, but these locally folded segments are rarely studied within larger architectures.

Combining foldamer subunits within a macrocycle represents a simple strategy toward higher-order structure, placing multiple secondary structures into well-defined spatial relationships (note that this approach is distinct from macrocycles that themselves fold^{13–18} or with a single folded unit^{19,20}).

Department of Chemistry & Biochemistry, Miami University, Oxford, OH 45056, USA.
E-mail: scott.hartley@miamioh.edu

† Electronic supplementary information (ESI) available: Supplemental figures and discussion referred to in the text, experimental procedures, NMR spectra, and computational data. See DOI: 10.1039/c9sc02975c

Foldamer-based macrocycles raise questions about how the foldamer moieties interact when confined within a larger structure, and about how folding and macrocyclization affect each other. For example, Huc has demonstrated homochiral self-sorting in macrocycles of aromatic oligoamides,²¹ and Chmielewski and Otto have very recently demonstrated a dynamic library that yields an emergent folding pattern.²² In related work, Sawada and Fujita have used foldamers as the components of complex assemblies based on metal coordination.^{23,24}

Our work has focused on the *o*-phenylenes, simple polyphenylenes²⁵ that adopt helical conformations in solution driven by (offset) arene–arene stacking interactions parallel to the helical axis.^{26–34} In previous work on *o*-phenylene-based macrocycles, we showed that amino-functionalized *o*-phenylene tetramers could be co-assembled with a series of rod-shaped dialdehyde linkers, giving, for example, the [3 + 3] macrocycles **oP⁴(Phen)₃₊₃** and **oP⁴(DPB)₃₊₃** shown in Chart 1 (*i.e.*, 3 *o*-phenylenes + 3 linkers).³⁵ The resulting twisted macrocycles^{36–40} are shape-persistent but have well-defined degrees of conformational freedom *via* the foldamer moieties. They exhibit rich stereochemical behavior, with both homochiral *D*₃-symmetric and heterochiral *C*₂-symmetric diastereomers distinguishable by NMR spectroscopy. Later work showed that



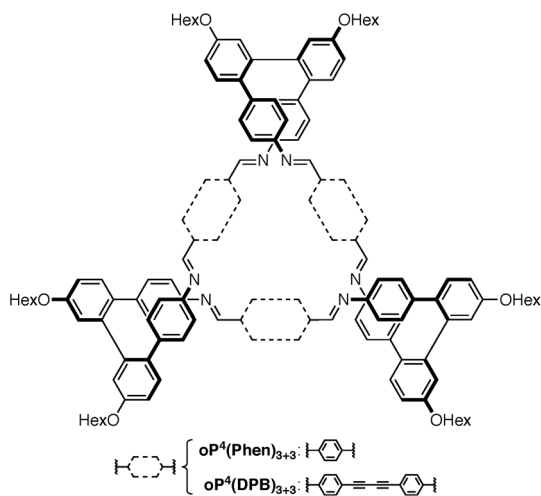
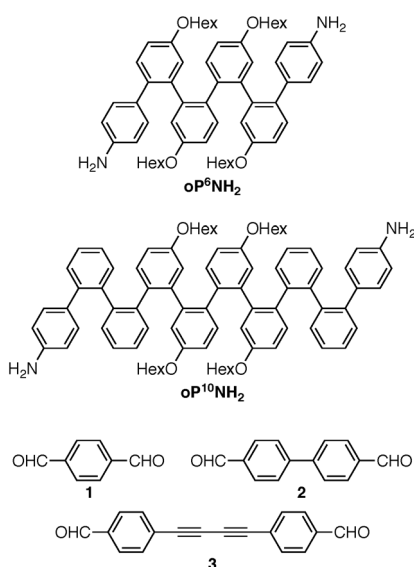


Chart 1 Selected previously reported twisted macrocycles.

variation in linker geometry can be used to direct the assembly toward $[2 + 2]$ and $[1 + 1]$ macrocycles.⁴¹

These results provided a proof of concept for the use of dynamic covalent chemistry to assemble *o*-phenylene foldamers into closed architectures (*i.e.*, macrocycles and cages). However, *o*-phenylene tetramers are the absolute simplest that can fold, with only a single relevant degree of conformational freedom (the central biaryl bond). In a sense, they cannot misfold: an *o*-phenylene tetramer can adopt only two possible backbone geometries, not counting enantiomers, both of which are helical.

Understanding the relationships between folding, self-assembly, and macrocyclic structure requires components with more-complex conformational energy landscapes. Here, we extend our results to include longer *o*-phenylene components **oP⁶(NH₂)** and **oP¹⁰(NH₂)**, shown in Chart 2, which have

Chart 2 *o*-Phenylenes and linker components.

been co-assembled with rod-shaped linkers **1**, **2**, and **3**. Increasing the length of the *o*-phenylenes to six or ten repeat units greatly increases the complexity of these systems: the hexamer gives two full turns of the helix when perfectly folded and can potentially adopt ten unique backbone geometries (five folding states and their enantiomers). The decamer gives 3.3 turns of the helix when perfectly folded and can potentially adopt 74 unique geometries (37 folding states and their enantiomers). This added complexity has a substantial effect on both self-assembly and the folding behavior of the macrocycles. Importantly, it affords new insights into how the subcomponents' folding propensities relate to the efficiency of assembly and how they respond to structural constraints imposed by the macrocycle. The behavior can be explained using simple models derived from conformational energy landscapes of the parent oligomers.

Results and discussion

Synthesis of components

The diamino *o*-phenylenes **oP⁶(NH₂)** and **oP¹⁰(NH₂)** were synthesized as shown in Scheme 1. Known²⁹ hydroxy-functionalized *o*-phenylene tetramer **4** was first alkylated to give **5**. This oligomer could then be terminated by Suzuki coupling to give **oP⁶(NH₂)**, or extended further through Suzuki coupling (**6**) and then triflation to give octamer **7**. We had originally intended to uniformly functionalize the longer oligomer with hexyloxy groups. However, despite multiple attempts, we were unable to couple **5** with the appropriate 10-hydroxy-10,9-boroxarophenanthrene⁴² derivative. It is likely that steric hindrance between the larger hexyloxy groups with the (predominantly folded) conformations of **5** prevented coupling. This limitation turned out to be useful, however, as the

Scheme 1 Synthesis of *o*-phenylenes **oP⁶(NH₂)** and **oP¹⁰(NH₂)**.

sequence of distinct repeat units in $\text{oP}^{10}(\text{NH}_2)$ allowed them to be more easily distinguished by NMR spectroscopy (see below).

The dialdehyde linkers were either commercially available (1) or were prepared according to literature procedures (2 and 3).^{43,44}

Hexamer assembly

We first examined the assembly of the *o*-phenylene hexamer $\text{oP}^6(\text{NH}_2)$, using the same conditions as previously used for the assembly of tetra(*o*-phenylene) macrocycles.³⁵ As shown in Fig. 1a, $\text{oP}^6(\text{NH}_2)$ (1.1 eq.) was combined with linkers 1, 2, or 3 (1.0 eq.) in chloroform (1.5 mM aldehyde) in the presence of TFA (0.10 eq.) and 3 Å molecular sieves. Monitoring of the reaction by ¹H NMR spectroscopy indicated that, in all three cases, the reaction mixtures reached a steady state after 5 d. At this point, the reactions were quenched with triethylamine to stop assembly and facilitate analysis.^{45–47} Gel permeation chromatography (GPC, Fig. 1b) of the resulting crude products showed a single predominant peak, indicating that assembly occurs efficiently in all three cases to give a single species (or set of species of very similar hydrodynamic radii). MALDI mass

spectra (Fig. 1c) indicated that the major products are the [3 + 3] macrocycles $\text{oP}^6(\text{Phen})_{3+3}$, $\text{oP}^6(\text{BIP})_{3+3}$, and $\text{oP}^6(\text{DPB})_{3+3}$. These macrocycles could be isolated by semi-preparative GPC in moderate yields,⁴⁸ and were then fully characterized by NMR spectroscopy and mass spectrometry.

From the GPC and mass spectrometry results, it is clear that *o*-phenylene hexamer $\text{oP}^6(\text{NH}_2)$ assembles effectively with rod-shaped linkers into quasi-triangular [3 + 3] macrocycles, much like the previously reported *o*-phenylene tetramers in Chart 1. What role does their folding play in macrocyclization?

The conformational analysis of *o*-phenylenes has been reviewed elsewhere.²⁶ Briefly, for an *o*-phenylene with *n* repeat units, the backbone folding state will be defined by the *n* – 3 internal biaryl dihedral angles. To a first approximation, these dihedrals can assume four values, approximately –55°, +55°, +135°, and –135°, which we refer to as the A, A', B, and B' states, respectively. Every second ring along the backbone must be nearly coplanar; it follows that within a single *o*-phenylene molecule only the A/B or A'/B' states can coexist. Thus, the total population of oligomers can be divided into two enantiomeric pools of oligomers (A/B and A'/B'). The limiting conformations, AA...A and BB...B, are both helical; the AA...A conformer is a compact left-handed helix and BB...B is an extended right-handed helix. The overall conformational distribution is then governed by two simple principles: (1) each "A" state results in a single arene–arene stacking interaction, worth on the order of 0.5 kcal mol^{–1} for unsubstituted *o*-phenylenes (in CDCl₃ around 0 °C). (2) The "ABA" sequence of dihedrals is strongly disfavored because it introduces unfavorable steric clashes. Thus, the "perfectly folded" AA...A state is favored for simple *o*-phenylenes.^{27,49}

This model is admittedly simplified: it fails, for example, to account for steric effects of substituents, and the twisting of the backbone in longer oligomers can certainly lead to additional conformational stabilization *via* other interactions beyond arene–arene stacking (*e.g.*, edge-to-face arene–arene interactions).³³ However, it provides a very useful framework within which to understand *o*-phenylene folding, and can be refined by using DFT calculations to more-accurately predict conformer energies.⁴⁹

An *o*-phenylene hexamer can therefore adopt five unique conformations in the A/B pool (and their enantiomers in the A'/B' pool), shown in Fig. 2a for (unsubstituted) hexa(*o*-phenylene). While the AAA conformer typically predominates, both the AAB and BAB conformers are usually observable.^{26,49,50}

In the context of macrocyclization, we also must consider the fit to the product geometry.⁵¹ For example, for a shape-persistent triangular [3 + 3] macrocycle, we expect the corners to make angles of roughly 60°. ⁵² The potential for each folding state to fit into this architecture can be approximated *via* its "bite" (or "bend") angle β ; that is, the angle made by the points of attachment to the *o*-phenylene moiety. This is a straightforward application of the concept of the minimization of strain in the assembly of macrocycles and cages, well-known in both metal-coordination and dynamic covalent systems.^{51–55}

Both the relative stabilities of different *o*-phenylene folding states E_{rel} and bite angles β are straightforward to predict using

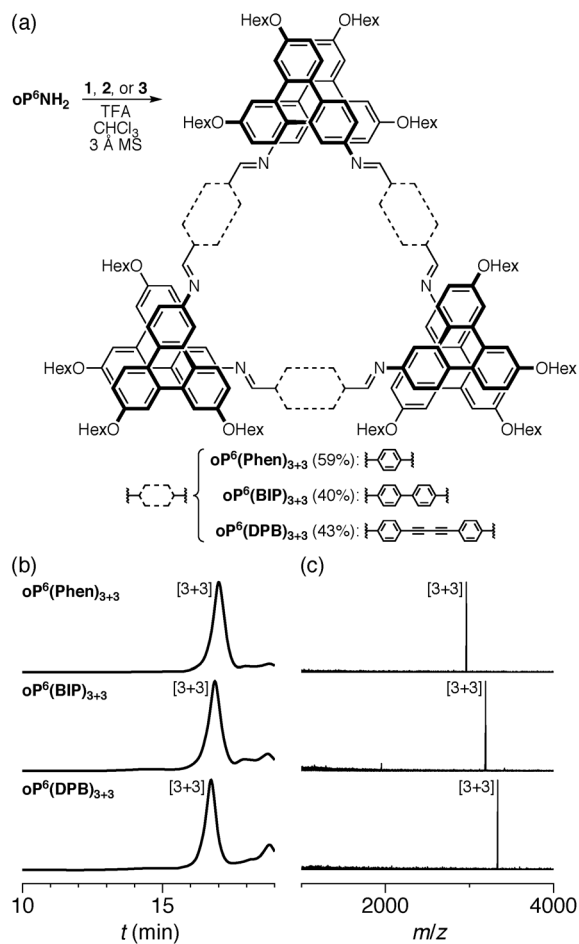


Fig. 1 (a) Assembly of $\text{oP}^6(\text{NH}_2)$ with linkers 1, 2, or 3. (b) GPC chromatograms of the (quenched) crude reaction mixtures after 5 d (RI detection). (c) MALDI mass spectra of the crude reaction mixtures.





Fig. 2 (a) Possible backbone geometries of hexa(*o*-phenylene) (PCM(CHCl₃)/B97-D/cc-pVDZ). (b) Calculated bite angles β and relative stabilities E_{rel} of the conformers. The region with $\beta \approx 60^\circ$, optimum for a [3 + 3] macrocycle, is highlighted. Inset: vectors used to calculate β .

DFT optimizations. For simplicity and efficiency, and anticipating the more-complex deca(*o*-phenylene) systems discussed below, we decided to focus on the unsubstituted hexa(*o*-phenylene); while the substituents will certainly affect the stabilities of *o*-phenylene conformers, the trend in stabilities and the bite angles should be consistent. The optimized geometries, shown in Fig. 2a, were obtained at the PCM(CHCl₃)/B97-D/cc-pVDZ level.⁵⁶ This method was found to reproduce the results of the previously used B97-D/TZV(2d,2p) method^{49,57} but in a fraction of the computational time (about $\frac{1}{5}$ for hexa(*o*-phenylene)). Note, however, that the calculated E_{rel} values from both methods overestimate the differences in stability that are actually observed in solution. They should nevertheless accurately predict the relative stabilities, which is sufficient for the analysis presented here. Bite angles β were calculated from the geometries in Fig. 2a using the vectors shown in the inset (see ESI†).

A plot of E_{rel} against β , shown in Fig. 2b, makes it immediately clear why **oP⁶(NH₂)** assembles efficiently into [3 + 3] macrocycles with linkers **1**, **2**, or **3**. A reversible system at equilibrium will favor macrocycles that balance their strain against size (*i.e.*, smaller macrocycles are favored entropically but can only form if they do not introduce significant strain).^{51,58} For this system, [1 + 1] macrocycles (the smallest conceivable) should be impossible given the geometries of **oP⁶(NH₂)** and any of the linkers. In their simplest form, [2 + 2] macrocycles would require $\beta \approx 0^\circ$,⁵² which is not found for any possible conformers of hexa(*o*-phenylene). The next-largest [3 + 3] macrocycle requires $\beta \approx 60^\circ$, and here multiple conformers have well-

matched intrinsic bite angles, including the most-stable AAA geometry ($\beta = 68^\circ$). Thus, this size is favored both entropically and enthalpically and should be assembled efficiently, as was observed.

NMR spectroscopy provides insight into how the *o*-phenylenes are folded within the macrocycles. One of the most useful features for the conformational analysis of *o*-phenylenes is that their backbones are typically in slow conformational exchange on the NMR time scale around room temperature (but fast exchange on the lab time scale).^{26,59} Thus, their NMR spectra are a superposition of contributions from different folding states. Importantly, the ¹H chemical shifts are very sensitive to the geometry, in some cases varying by >1 ppm. In general, this dependence can be quantitatively predicted using *ab initio* methods. NMR spectroscopy therefore allows the folding states of *o*-phenylenes to be determined and quantified. While crystallography would have provided useful structural information, it would be very challenging to crystallize the array of macrocyclic geometries expected in solution (see below), and even then the prevalence of each species in the overall population would not be determined. Since our focus was the behavior of mixtures of stereoisomers and folding states, the oligomers were decorated with hexyloxy groups to promote solubility and crystals suitable for X-ray diffraction were not obtained.

We synthesized **oP⁶(M)**, an acyclic analogue of the *o*-phenylene hexamer moieties within the macrocycles, as a reference point (see ESI†). Its ¹H NMR spectrum (CDCl₃, 0 °C) is shown in Fig. 3 (top).⁶⁰ As expected, the spectrum is relatively complicated, with contributions from misfolded states in the alkoxy (*i.e.*, -CH₂O-, 3–4 ppm), aromatic, and imine regions. EXSY experiments show that the protons giving rise to these minor signals are in exchange with those for major signals, confirming that they are different conformational states and not impurities (see ESI†). It was straightforward to assign the signals corresponding to the major (twofold-symmetric) conformer using standard 2D NMR experiments (COSY, HSQC, HMBC). Following a strategy now used many times for *o*-phenylenes,^{26,59} a simplified computational model of **oP⁶(M)** was optimized in all five possible backbone geometries at the PCM(CHCl₃)/B97-D/cc-pVDZ level and NMR isotropic shieldings were calculated for each at the PCM(CHCl₃)/WP04/6-31G(d) level.^{61,62} Comparison of the experimental and predicted NMR data confirmed that the major conformer does indeed correspond to the AAA state (see ESI†). By analogy with previous results,^{26,49,50,59,63} the minor conformers were assigned to the AAB and BAB geometries; this is probably most easily seen in the imine region around 8.5 ppm, which shows two equal-intensity imine signals for an unsymmetrical conformer (*i.e.*, AAB) and a single smaller signal for a symmetrical conformer (BAB), labeled in Fig. 3.

The ¹H NMR spectra of macrocycles **oP⁶(Phen)₃₊₃**, **oP⁶(BIP)₃₊₃**, and **oP⁶(DPB)₃₊₃** are also shown in Fig. 3. A cursory inspection indicates that the spectra are all similar (disregarding the signals associated with the linkers themselves), and similar to that of **oP⁶(M)**. This similarity suggests that the folding behavior has not been dramatically perturbed; however, there are some clear differences. First, the spectrum of **oP⁶(Phen)₃₊₃**, and to a lesser extent **oP⁶(BIP)₃₊₃**, is more complex





Fig. 3 ^1H NMR spectra (500 MHz, 0°C) of $\text{oP}^6(\text{M})$, $\text{oP}^6(\text{Phen})_{3+3}$, $\text{oP}^6(\text{BIP})_{3+3}$, and $\text{oP}^6(\text{DPB})_{3+3}$. ^1H assignments for the major conformers of $\text{oP}^6(\text{M})$ and $\text{oP}^6(\text{DPB})_{3+3}$ are shown (blue), as are imine ^1H assignments to backbone geometries (orange). Assignments of the imine signals of $\text{oP}^6(\text{Phen})_{3+3}$ to the homochiral D_3 and heterochiral C_2 stereoisomers are also labeled.

than the others. Close inspection of the imine signals (inset in Fig. 3) shows that there are far more signals than in the spectrum for $\text{oP}^6(\text{M})$, suggesting many distinct species in solution. In our previous work on $\text{oP}^4(\text{Phen})_{3+3}$, we found that the *p*-phenylene linker was sufficiently rigid that the different diastereomeric forms of the macrocycle could be identified;³⁵ that is, the single imine signal for the D_3 -symmetric homochiral (*PPP*, *MMM*) stereoisomer was readily distinguished from the three imine signals for the C_2 -symmetric heterochiral (*PMM*, *MPP*) stereoisomers. The behavior of $\text{oP}^6(\text{Phen})_{3+3}$ appears to be similar, except that there is also the added complication of oligomer misfolding. Indeed, the four most-prominent imine signals are consistent with the homochiral- D_3 (1 signal) and heterochiral- C_2 (3 signals) configurations of the perfectly folded (AAA)₃ macrocycle (inset in Fig. 3), with the smaller peaks corresponding to macrocycles with at least one of the *o*-phenylenes misfolded. The D_3 - and C_2 -symmetric stereoisomers of the (AAA)₃ configuration appear to be present in approximately the 1 : 3 ratio attributable to symmetry alone.

Clearly, there is little stereochemical control in these systems, as was expected from the previous results for $\text{oP}^4(\text{Phen})_{3+3}$. However, the specific folding state of the hexa(*o*-phenylene) moieties within macrocycles $\text{oP}^6(\text{Phen})_{3+3}$, $\text{oP}^6(\text{BIP})_{3+3}$, and $\text{oP}^6(\text{DPB})_{3+3}$ is a separate issue. *o*-Phenylene

conformations with close to the optimum β should be preferred by the [3 + 3] macrocycle (Fig. 2b, shaded region). This analysis suggests that, of the three conformers that should be observed in solution, AAA, AAB, and BAB, only the AAA and AAB geometries should fit.

The coexistence of multiple stereoisomers of each macrocycle complicates assignment of the ^1H NMR spectra of $\text{oP}^6(\text{Phen})_{3+3}$ and $\text{oP}^6(\text{BIP})_{3+3}$. Fortunately, the diphenylbutadiyne linker in $\text{oP}^6(\text{DPB})_{3+3}$ is sufficiently flexible⁶⁴ that the different stereoisomers no longer give distinct ^1H NMR signals. That is, multiple stereoisomers of $\text{oP}^6(\text{DPB})_{3+3}$ presumably coexist, but they give indistinguishable spectra because the linker does not communicate the configuration between *o*-phenylenes (as had been previously observed for $\text{oP}^4(\text{DPB})_{3+3}$).³⁵ Thus, analysis is no more difficult than it was for $\text{oP}^6(\text{M})$: as before, each of the aromatic protons on the backbone of the major conformer could be assigned using standard 2D NMR experiments (see ESI†). The assignments, shown in Fig. 3, are a close match to those for $\text{oP}^6(\text{M})$, suggesting that the *o*-phenylene subunits of $\text{oP}^6(\text{DPB})_{3+3}$ are also in the AAA configuration.

It is conceivable that other backbone geometries could give similar NMR spectra to the AAA state of an *o*-phenylene hexamer. We therefore compared the ^1H chemical shifts in $\text{oP}^6(\text{DPB})_{3+3}$ and $\text{oP}^6(\text{M})$ more quantitatively (this analysis also anticipates the much more complex case of the deca(*o*-phenylenes) discussed below). We focused on the chemical shift differences $\Delta\delta$ for analogous protons between the *o*-phenylene within the macrocycle and in the model $\text{oP}^6(\text{M})$. For example, for proton H_{1a} (see Fig. 3 for labeling), the experimentally observed $\Delta\delta_{1a}^{\text{exp}}$ is given by

$$\Delta\delta_{1a}^{\text{exp}} = \delta_{1a}(\text{oP}^6(\text{DPB})_3) - \delta_{1a}(\text{oP}^6(\text{M})) \quad (1)$$

where $\delta_{1a}(\text{oP}^6(\text{DPB})_3)$ and $\delta_{1a}(\text{oP}^6(\text{M}))$ are the chemical shifts of proton H_{1a} in $\text{oP}^6(\text{DPB})_{3+3}$ and the major conformer of $\text{oP}^6(\text{M})$, respectively. The experimental $\Delta\delta^{\text{exp}}$ values should reflect the chemical shift differences between the (possibly misfolded) *o*-phenylenes in the macrocycle and the AAA state that was already established for $\text{oP}^6(\text{M})$.

We focus on $\Delta\delta$ (rather than δ) because substituent effects on chemical shift should cancel out: while the hexyloxy groups and imines should significantly affect the chemical shifts of nearby protons, their contributions to $\Delta\delta$, which are primarily dependent on the orientations of the protons with respect to the shielding zones of nearby aromatic rings, should be negligible. This assumption allows us to carry out computational work on the library of unsubstituted conformers and apply the results to experimental systems regardless of *o*-phenylene substitution pattern or structural context, ignoring, for example, the relative orientations of attached functionality.

Computational $\Delta\delta^{\text{calc}}$ values were then predicted for each of the five optimized geometries of hexa(*o*-phenylene) (Fig. 2) at the GIAO/PCM(CHCl_3)/WP04/6-31G(d) level. They were calculated relative to the AAA conformer; for example, for proton H_{1a} in conformer AAB, the predicted difference in chemical shift $\Delta\delta_{1a}^{\text{calc}}(\text{AAB})$ is given by



$$\Delta\delta_{1a}^{\text{calc}}(\text{AAB}) = \sigma_{1a}(\text{AAA}) - \sigma_{1a}(\text{AAB}) \quad (2)$$

where $\sigma_{1a}(\text{AAA})$ and $\sigma_{1a}(\text{AAB})$ are the isotropic shieldings of H_{1a} in the AAA and AAB conformers, respectively.

The predicted $\Delta\delta^{\text{calc}}$ vary from the trivial case of the AAA conformer, for which, by definition, $\Delta\delta^{\text{calc}} = 0$ for all protons, to more than 2 ppm for some key protons in other conformers. Each conformer gives a unique pattern of $\Delta\delta_{1a}^{\text{calc}}$, shown in Fig. 4a, that should serve as a fingerprint for the folding state of an *o*-phenylene moiety. The experimental differences $\Delta\delta^{\text{exp}}$ for the major *o*-phenylene conformer in $\text{OP}^6(\text{DPB})_{3+3}$, shown in Fig. 4b, are all very close to 0 (≤ 0.06 ppm). They are very clearly the best match for the AAA configuration, with a root-mean-squared deviation (RMSD) of only 0.02 ppm between the experimental and predicted $\Delta\delta$. As shown in Fig. 4b, this match is substantially better than the next-best comparison (to AAB, RMSD = 0.43 ppm), significant at the 99.9% confidence level (see ESI for details†).

The regions of the spectra corresponding to the alkoxy groups (3–4 ppm), also shown in Fig. 3, provide corroborating evidence that the *o*-phenylenes remain well-folded within the macrocycles. These signals should be largely unaffected by changes in linker structure or relative *o*-phenylene stereochemistry but are very sensitive to *o*-phenylene folding state.⁶⁵ These regions of the spectra are very similar for $\text{OP}^6(\text{M})$ and all three macrocycles, indicating similar overall folding behavior in all cases.

Although the *o*-phenylenes within the macrocycles remain predominantly folded into the AAA state, there are differences in the relative populations of the different conformers. This is most easily seen in the imine regions of the spectra in Fig. 3. For $\text{OP}^6(\text{M})$, integration of the imine singlets gives a ratio of 54 : 39 : 7 for AAA : AAB : BAB. In contrast, for $\text{OP}^6(\text{DPB})_{3+3}$ we obtain 69 : 31 : 0, which is significantly different even taking into account the 5–10% error typically assumed for NMR integration. This result is consistent with expectations based on Fig. 2: the AAA state ($\beta = 68^\circ$) was expected to be the best match for the [3 + 3] macrocyclic geometry, and indeed it is amplified in the macrocycle. The AAB state is still observed ($\beta = 84^\circ$), although it is less populated, but the BAB state ($\beta = 132^\circ$) is eliminated entirely.

This result implies that the bite angle is a useful parameter for assessing the fit of an *o*-phenylene conformer within a shape-persistent architecture. In our previous work on *o*-phenylene tetramer macrocycles (e.g., $\text{OP}^4(\text{DPB})_{3+3}$), we had noted that macrocyclization led to improved folding.³⁵ This conclusion was drawn on the basis of characteristic changes in chemical shifts for key protons in rapid exchange.⁶⁶ Macrocycle $\text{OP}^6(\text{DPB})_{3+3}$ confirms this result by allowing direct observation of a change in the conformational population.

To visualize the structure of the *o*-phenylene hexamer macrocycles, we performed geometry optimizations on $\text{OP}^6(\text{DPB})_{3+3}$, a simplified version of $\text{OP}^6(\text{DPB})_{3+3}$ with the alkoxy groups removed, at the PCM(CHCl_3)/B97-D/cc-pVDZ level. Of course, even with structural simplifications and the relative simplicity of the linkers, there remain many conformational degrees of freedom that must be accounted for. Thus, the larger structures were built up by systematically optimizing key dihedral angles for smaller fragments, as described in the ESI.† We stress that the strategy does not guarantee the identification of the global conformational energy minimum; however, it should provide a good indication of the fit of the *o*-phenylene within the macrocyclic framework.

The geometries of $\text{OP}^6(\text{DPB})_{3+3}$ in two of its configurations, homochiral-(AAA)₃ and homochiral-(AAA)₂(AAB), are shown in Fig. 5. The heterochiral-(AAA)₃ conformer (not shown) is energetically very similar to the homochiral isomer ($\Delta E =$

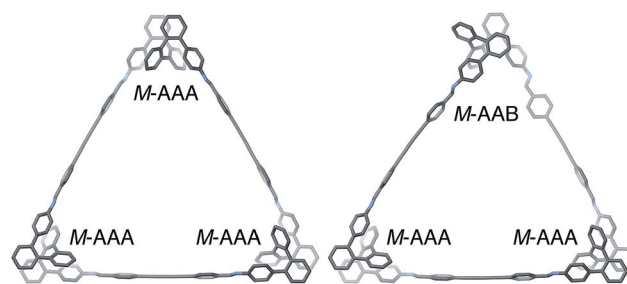


Fig. 5 Optimized geometries (PCM(CHCl_3)/B97-D/cc-pVDZ) of $\text{OP}^6(\text{DPB})_{3+3}$ in its homochiral-(AAA)₃ (left) and homochiral-(AAA)₂(AAB) (right) configurations.

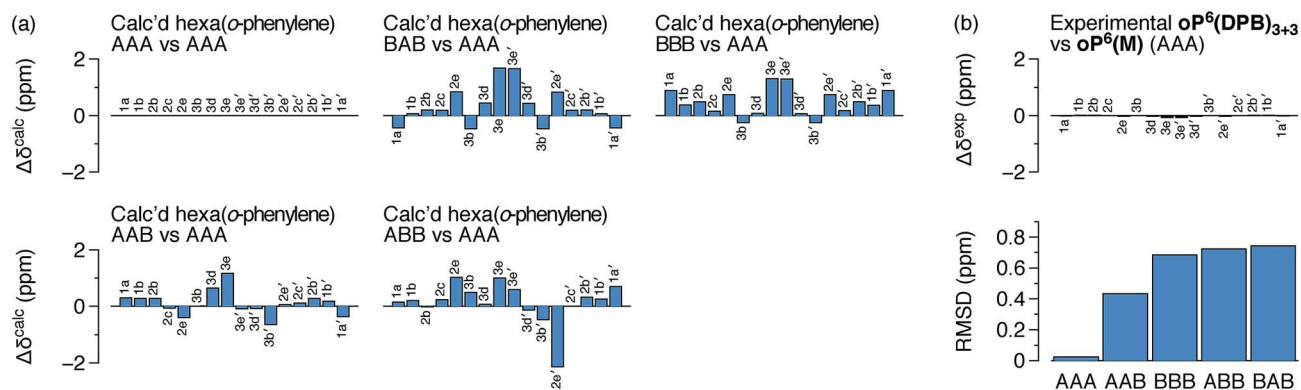


Fig. 4 (a) Predicted $\Delta\delta^{\text{calc}}$ for each possible backbone configuration of hexa(*o*-phenylene) relative to the parent AAA. (b) $\Delta\delta^{\text{exp}}$ for $\text{OP}^6(\text{DPB})_{3+3}$ vs. $\text{OP}^6(\text{M})$ (AAA) and RMSD of predicted vs. experimental differences for each possible backbone configuration.



0.01 kcal mol⁻¹). Conversely, the mixed homochiral-(AAA)₂-(AAB) geometry is predicted to be less stable than the fully folded (AAA)₃-homochiral macrocycle by 3.6 kcal mol⁻¹. This is significantly larger than the inherent preference for the AAA conformer over AAB (2.5 kcal mol⁻¹ at the same level of theory), suggesting that the poorer fit of the AAB state, because of its less-well-matched β , introduces additional strain (*i.e.*, by forcing the *o*-phenylene moiety away from its optimum AAB geometry).

Decamer assembly

We then turned to the assembly of the **oP**¹⁰(NH₂) system, as shown in Fig. 6a. Reaction with linkers **1**, **2**, and **3** was carried out under the same conditions as for **oP**⁶(NH₂). In all three cases, assembly was found to occur more slowly, with the best results obtained after 11 d. At this point, GPC analysis of the quenched reaction mixtures, shown in Fig. 6b, gave qualitatively similar results for all three linkers. Two distinct peaks corresponding to two major products (overlapping at approximately 17 min) were observed along with a broad peak corresponding to larger species. Beyond 11 d, additional molecular sieves had to be added to maintain the same distribution of products but no net improvement could be obtained. Thus, we cannot conclusively say that the GPC traces in Fig. 6b represent equilibrium distributions for these systems. It is, however, clear that assembly is less efficient for **oP**¹⁰(NH₂) compared to **oP**⁶(NH₂). MALDI mass spectra of the crude reaction mixtures show dominant peaks corresponding to the [2 + 2] macrocycles along

with small signals for the [3 + 3] macrocycles in two cases (Fig. 6c). Careful purification by multiple semi-preparative GPC injections allowed the two major products to be isolated in reasonable yields and conclusively identified as the [2 + 2] and [3 + 3] macrocycles by NMR and mass spectrometry. Isolated yields of the [2 + 2] macrocycles increase in the order **oP**¹⁰(Phen)₂₊₂ < **oP**¹⁰(BIP)₂₊₂ < **oP**¹⁰(DPB)₂₊₂. As the GPC chromatograms of the crude mixtures are very similar in all three cases, this trend likely reflects greater ease of purification of the macrocycles with longer linkers rather than greater efficiency of self-assembly.

Of course, the conformational landscape available to an unconstrained *o*-phenylene decamer is substantially larger than of a hexamer, with 37 possible folding states (not including enantiomers). A library of all of these possibilities for the unsubstituted deca(*o*-phenylene) was optimized as before (PCM(CHCl₃)/B97-D/cc-pVDZ). The corresponding relative stabilities E_{rel} and bite angles β are plotted in Fig. 7. Like hexa(*o*-phenylene) (Fig. 2), the most stable conformer of deca(*o*-phenylene) is the perfectly folded state (AAAAAAA). This conformer has a $\beta = 69^\circ$ and would be expected to fit well into a roughly triangular [3 + 3] macrocycle. Unlike hexa(*o*-phenylene), deca(*o*-phenylene) has many folding geometries with β near 0°, several of which should be energetically accessible (bearing in mind that the E_{rel} overestimate the absolute energetic differences between conformers). These smaller bite angles should be a good match to the smaller [2 + 2] architecture.

The complexity of the deca(*o*-phenylene) conformational energy landscape explains why mixtures are obtained from assembly of **oP**¹⁰(NH₂) with linkers **1**, **2**, or **3** as well as the slight preference for assembly into [2 + 2] macrocycles. As briefly discussed for the hexamer-based system, the equilibria between macrocycles of various *o*-phenylene geometries and sizes will be governed, at least to a first approximation, by (1) the entropic favorability of smaller macrocycles, as commonly exploited in self-assembly (including dynamic covalent assembly),⁵¹ and (2) the strain associated with misfolding.⁶⁷ For the *o*-phenylene decamer system, Fig. 7 predicts that these two effects work against each other: while the AAAAAAA conformer is the most inherently stable, it is not sufficiently favored to overcome the

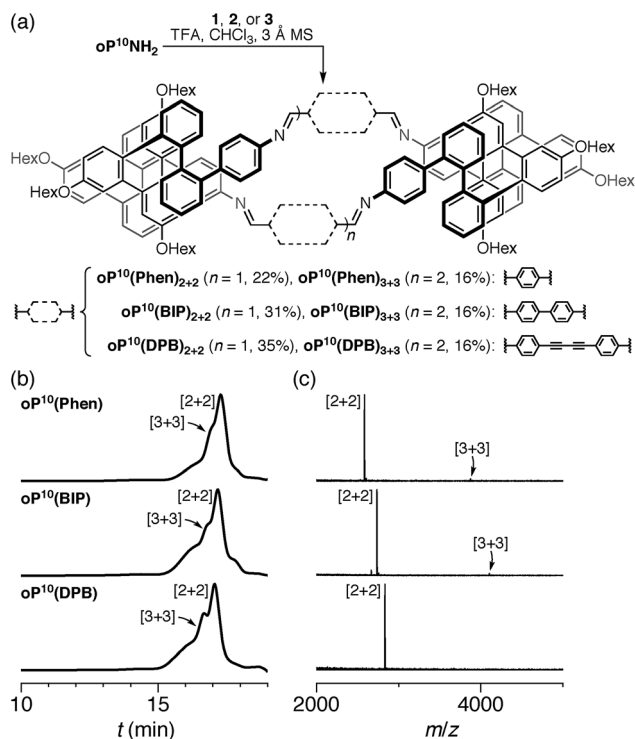


Fig. 6 (a) Assembly of **oP**¹⁰(NH₂) with linkers **1**, **2**, and **3**. (b) GPC chromatograms of the (quenched) crude reaction mixtures after 11 d (RI detection). (c) MALDI mass spectra of the crude reaction mixtures.



Fig. 7 Calculated E_{rel} vs. β for all conformers of deca(*o*-phenylene) (PCM(CHCl₃)/B97-D/cc-pVDZ). Selected conformations are labeled.



entropic favorability of (misfolded) smaller macrocycles. This behavior contrasts with that of the shorter *o*-phenylene tetramers and hexamers, which offer no conformers that can reasonably fit within a macrocycle smaller than [3 + 3].

This inherent frustration implies that confinement within smaller architectures forces the foldamer subunits toward structurally complex misfolded states. We therefore explored the folding of the *o*-phenylene moieties as before. Model compound $\text{oP}^{10}(\text{M})$ was prepared for comparison (see ESI†), with its ^1H NMR spectrum shown in Fig. 8 (top). While complex, the signals corresponding to its major conformer are straightforward to assign and the geometry can be established as AAAAAAA by comparison with DFT predictions of chemical shifts (see ESI†). Nevertheless, there are several other observed conformers, most easily seen in the imine region of the spectrum. Integration shows that approximately 25% of the population is perfectly folded.

The ^1H NMR spectra of the macrocycles resulting from the three linkers are shown in the ESI.† Close examination of the alkoxy group region ($-\text{CH}_2\text{O}-$) suggests that the folding of the *o*-phenylenes within a particular macrocycle size ([2 + 2] or [3 + 3]) is similar, but that there are differences between the two sizes themselves. The aromatic regions of the ^1H NMR spectra of macrocycles resulting from linkers 1 and 2 ($\text{oP}^{10}(\text{Phen})_{2+2}$, $\text{oP}^{10}(\text{Phen})_{3+3}$, $\text{oP}^{10}(\text{BIP})_{2+2}$, and $\text{oP}^{10}(\text{BIP})_{3+3}$) are complex, again reflecting the coexistence of many stereoisomers. We therefore focused on the macrocycles from more-flexible linker 3 ($\text{oP}^{10}(\text{DPB})_{2+2}$ and $\text{oP}^{10}(\text{DPB})_{3+3}$), which again give more easily

interpreted spectra, shown in Fig. 8, since the stereoisomers are indistinguishable.

The behavior of the [3 + 3] macrocycle $\text{oP}^{10}(\text{DPB})_{3+3}$ is straightforward and closely parallels that of $\text{oP}^6(\text{DPB})_{3+3}$. While its ^1H NMR signals are slightly broadened, there is very little difference between its spectrum and that of $\text{oP}^{10}(\text{M})$ in both the alkoxy and aromatic regions. Explicit assignments of the aromatic protons in the major conformer were made through analysis of 2D spectra. Differences in chemical shifts $\Delta\delta^{\text{exp}}$ were calculated relative to the AAAAAAA conformer of $\text{oP}^{10}(\text{M})$ and were all ≤ 0.02 ppm. These were then compared to predicted changes in chemical shifts $\Delta\delta^{\text{calc}}$ for the full 37-conformer library relative to the AAAAAAA conformer of deca(*o*-phenylene). The best match was, unsurprisingly, to the AAAAAAA conformer of the library, with a RMSD of 0.01 ppm. Thus, the *o*-phenylene decamers in the [3 + 3] macrocycle remain well-folded, with little difference in behavior relative to freely mobile acyclic oligomers. Integration of the imine region indicates a slight increase in preference for the well-folded conformer within the macrocycle, with 40% of the oligomers in the AAAAAAA state (*vs.* 25%).

Folding in the [2 + 2] macrocycle $\text{oP}^{10}(\text{DPB})_{2+2}$ is very different. A quick inspection of the ^1H NMR spectra (Fig. 8, bottom) shows that there is a marked difference in overall appearance, including in the alkoxy region.⁶⁸ While challenging, it was possible to assign all of the significant signals in the aromatic region of the spectrum to two principal conformers that are present in an approximately 2 : 1 ratio.⁶⁹ The major conformer, whose assignments are shown in Fig. 8, is unsymmetrical, with unique ^1H signals along the entirety of its backbone. The minor conformer is twofold symmetric (see ESI† for assignments).

The differences in chemical shift $\Delta\delta^{\text{exp}}$ between both the major and minor conformers of $\text{oP}^{10}(\text{DPB})_{2+2}$ and the AAAAAAA state of $\text{oP}^{10}(\text{M})$ were then determined and are shown in Fig. 9a (top). They are substantially larger than was observed in $\text{oP}^6(\text{DPB})_{3+3}$ and $\text{oP}^{10}(\text{DPB})_{3+3}$. For example, proton H_{3e} in the major conformer is shifted upfield by 1.1 ppm (to 4.74 ppm), indicating that it must be deep within the shielding zone of a nearby aromatic ring. It was therefore clear that neither conformer corresponds to the AAAAAAA state. The experimental $\Delta\delta^{\text{exp}}$ were then compared with the $\Delta\delta^{\text{calc}}$ values predicted for the library of deca(*o*-phenylene) conformers. Clear matches were identified: the major conformer corresponds to the AAAABBB folding state, whereas the minor conformer corresponds to the BAAAAAB folding state. In both cases the similarity between the experimental data and the NMR predictions for the parent deca(*o*-phenylene) conformer library is striking (Fig. 9a). Quantitative comparisons to the full library were made by comparing the RMSD values for each possible conformer, as shown in Fig. 9b. For both of the experimentally observed folds, the best matches have RMSD values on the order of 0.1 ppm, typical for NMR comparisons made for *o*-phenylenes.²⁶ The second-best matches are significantly worse (>0.3 ppm), with the differences statistically significant at a $>99\%$ confidence level (see ESI†).

The AAAABBB and BAAAAAB folding states are highlighted in the plot of E_{rel} *vs.* β in Fig. 7. They are indeed among the

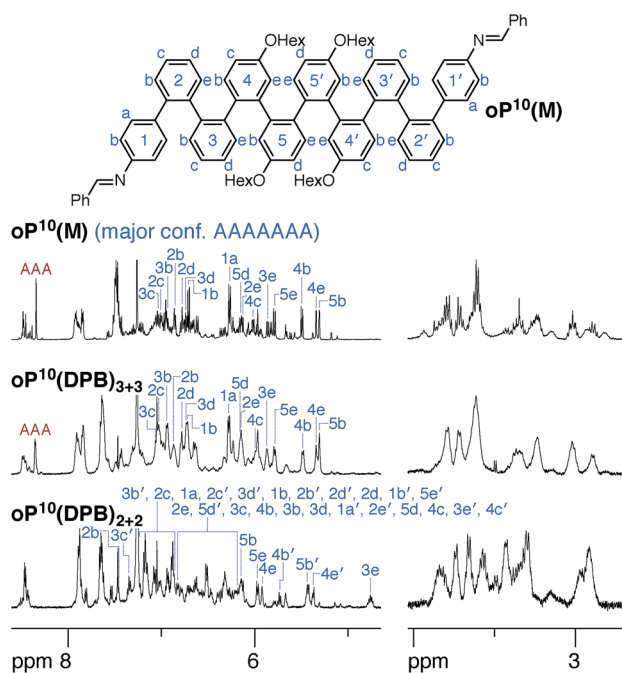


Fig. 8 ^1H NMR spectra (500 MHz, 0°C) of $\text{oP}^{10}(\text{M})$, $\text{oP}^{10}(\text{DPB})_{2+2}$, and $\text{oP}^{10}(\text{DPB})_{3+3}$. ^1H assignments for the major conformers are shown (blue). For $\text{oP}^{10}(\text{DPB})_{2+2}$, the assignments that are grouped together are listed in order from least- to most-shielded; chemical shifts for a minor conformation were also assigned (not shown).



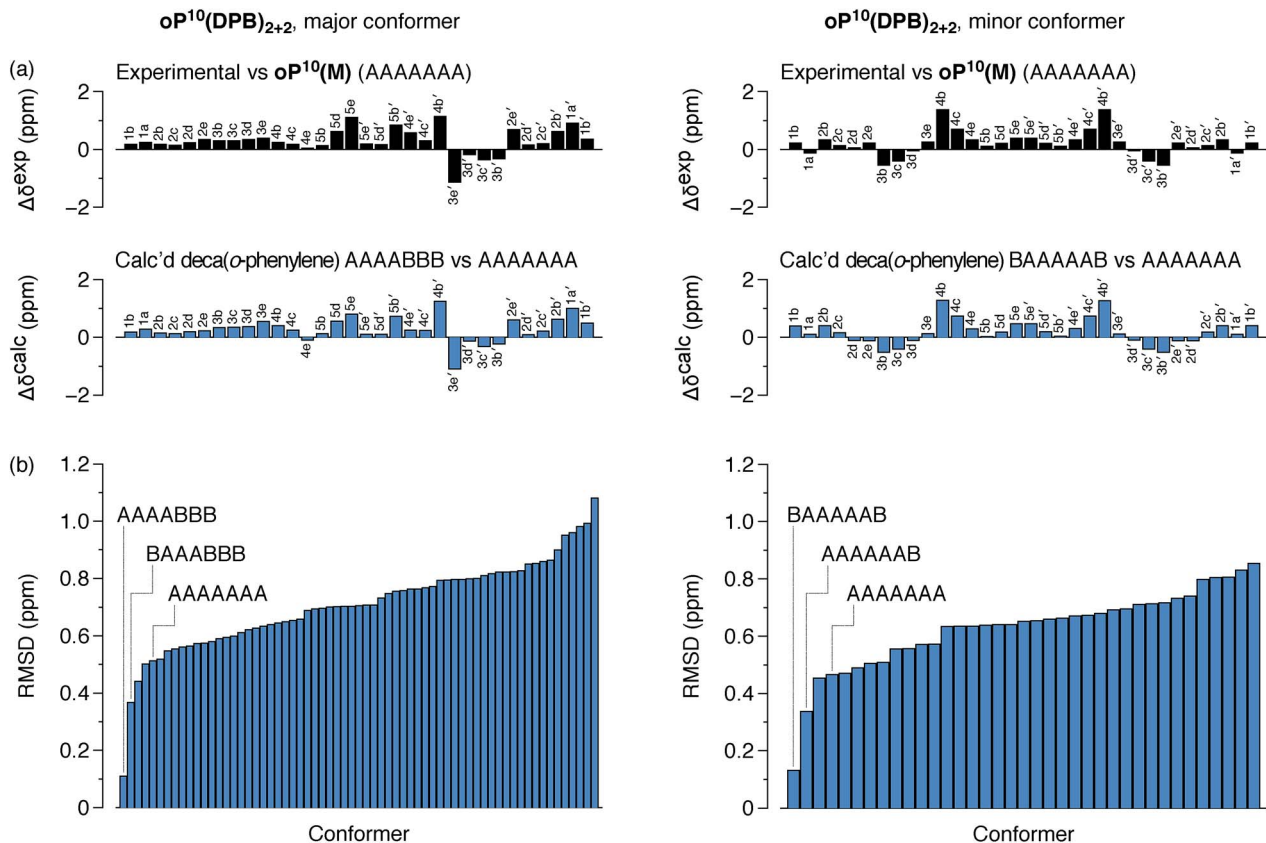


Fig. 9 (a) Experimental $\Delta\delta^{\text{exp}}$ for $oP^{10}(\text{DPB})_{2+2}$, major and minor *o*-phenylene conformers, vs. $oP^{10}(\text{M})$ in its AAAAAAA conformer, and predicted $\Delta\delta^{\text{calc}}$ for the parent deca(*o*-phenylene), AAAABBB and BAAAAAB conformers, vs. AAAAAA. (b) RMSD of $\Delta\delta^{\text{exp}}$ values for each deca(*o*-phenylene) backbone configuration vs. the experimental $\Delta\delta^{\text{calc}}$. The AAAABBB configuration is a better fit for the major conformer than BAAAABB at the >99.9% confidence level, and the BAAAAAB configuration is a better fit than AAAAAAB for the minor conformer at the 99.7% confidence level.

configurations that would be predicted to be good matches to the $[2 + 2]$ geometry, with low bite angles ($\beta < 30^\circ$) and relatively good stabilities. To confirm that they should be well-accommodated within a $[2 + 2]$ macrocycle, we optimized the geometries of the (AAAABBB)₂, (BAAAAAB)₂, and (AAAABBB)(-BAAAAAB) configurations of a simplified version of $oP^{10}(\text{DPB})_{2+2}$ with the alkoxy groups removed, $oP^{10}(\text{DPB})_{2+2}$, at the PCM(CHCl₃)/B97-D/cc-pVDZ level. All possible relative configurations of the *o*-phenylenes and, in the case of (AAAABBB)₂, their relative direction (*i.e.*, parallel or antiparallel) were considered. The most stable geometries for each species are shown in Fig. 10. Clearly, the assumption that the $[2 + 2]$ architecture requires $\beta \approx 0^\circ$ is an oversimplification, with the linkers crossing in all three geometries. The results do confirm, however, that both observed *o*-phenylene folding states are good matches to this size. The product presumably comprises a mixture of these geometries, but unfortunately we cannot be more specific at this time.

Fig. 7 suggests that there are other configurations that would also have been expected to be competitive, with smaller bite angles, higher stabilities, or both compared to the observed conformers (*e.g.*, AAABBBA or ABBAAB). Why are these not observed? There are two likely explanations. First, the true optimum β , as mentioned above, is not predictable by the

macrocycle stoichiometry alone. Because the $[2 + 2]$ macrocycles can adopt geometries with crisscrossed linkers, somewhat larger bite angles are tolerated (*e.g.*, 43° in the homochiral-(AAAABBB)₂ macrocycle). Second, the stabilities in Fig. 7 are predicted for the unsubstituted deca(*o*-phenylene). Substituted oligomers will experience additional interactions involving the substituents that are not accounted for in these models. For example, steric clashes between the relatively large hexyloxy groups will disfavor some conformers with congested structures. Indeed, optimization of several of the alternate backbone geometries with explicit methoxy groups indicates that they are less stable than would have been predicted using the library of parent conformers (see ESI†). This is an important caveat; however, the optimization of the geometries of unsubstituted oligomers is so much simpler, and the NMR predictions so useful, that explicit consideration of the substituents does not seem worthwhile.

Unlike the *o*-phenylene tetramer and hexamer systems, the assembly and structure of the $oP^{10}(\text{NH}_2)$ -derived macrocycles demonstrates that new structural complexity can emerge in these systems once the baseline conformational behavior is sufficiently complex. The distribution of products highlights an important feature of the *o*-phenylenes: the overall driving force for folding is relatively weak; thus, it is competitive with the



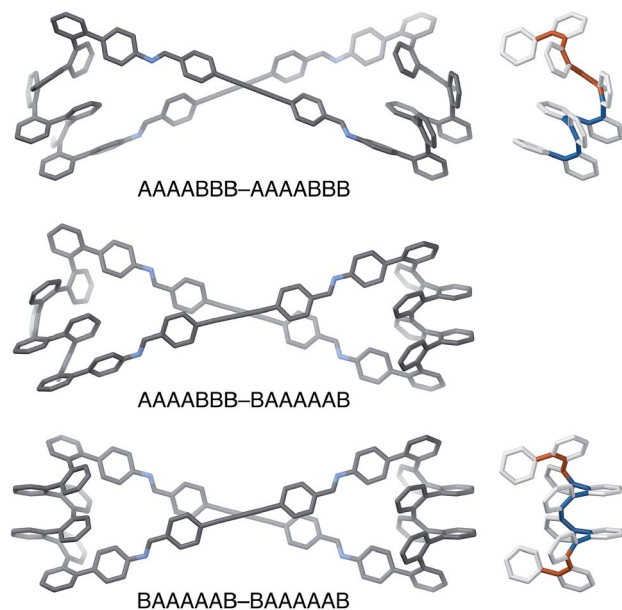


Fig. 10 Left: Optimized geometries (PCM(CHCl₃)/B97-D/cc-pVDZ) of $\text{oP}^{10}(\text{DPB})_{2+2}$ in its homochiral-antiparallel-(AAAABBB)₂ (top), homochiral-(AAAABBB)(BAAAAAB) (middle), and homochiral-(BAAAAAB)₂ (bottom) configurations. Right: *o*-phenylene geometries with the A...A (blue) and B...B (orange) folding states highlighted on the backbone.

entropic driving force for smaller macrocycles. It follows that thermodynamically controlled approaches can be used to prepare closed architectures with the foldamers in misfolded states. While quantitative yields may not be feasible, once isolated these species offer added structural complexity, with the foldamer moieties restricted to a predictable region of their conformational energy landscapes. In principle, it should be possible to use perturbations (*e.g.*, host-guest binding) to select for macrocycles of various sizes,⁷⁰ and it could also be possible to perturb assembly by tuning the *o*-phenylene folding propensity.⁶³

The minor *o*-phenylene folding observed in $\text{oP}^{10}(\text{DPB})_{2+2}$, BAAAAAB, is not unexpected given that fraying at the ends of *o*-phenylenes is commonly observed.⁵⁹ The major state, AAAABBB, is much more unusual, however, and shows that simple geometric constraints can be used to access folding patterns that are otherwise unavailable. This geometry has a fascinating structure: the “AAAA” end is folded by analogy with all other *o*-phenylenes, into a compact helix with arene-arene stacking. In contrast, the “BBB” end is pulled into the extended helix that has not, to our knowledge, been observed for simple *o*-phenylenes (but is commonly found for the analogous poly(quinoxaline-2,3-diyls)^{71,72}). *o*-Phenylene moieties in this conformation thus have two distinct helical blocks twisting in opposite directions, left then right handed for AAAABBB, as is shown in Fig. 10, or right then left handed for A'A'A'B'B'B'.

Unlike the hierarchical structure in proteins, the AAAABBB folding in $\text{oP}^{10}(\text{DPB})_{2+2}$ does not result from structural complexity programmed into the *o*-phenylenes themselves. The reactant $\text{oP}^{10}(\text{NH}_2)$ is symmetrical and its sequence does not code for anything more complex than the perfectly folded

AAAAAAA pattern, as is clear from model $\text{oP}^{10}(\text{M})$. Instead, the *o*-phenylene moiety in the macrocycle adopts a more complex geometry as a consequence of the constraints on folding imposed by the larger architecture *via* β . Of course, the fact that this geometry is structurally interesting, with opposing helical twists, is presumably just a coincidence; this just happens to be one of the conformers with the appropriate β . *o*-Phenylenes of other lengths would likely adopt very different folding motifs in strained macrocycles, but this should be predictable, to some extent, by determining the stabilities and bite angles of conformer libraries for the parent acyclic oligomers.

Conclusions

In summary, *o*-phenylene hexamer $\text{oP}^6(\text{NH}_2)$ and decamer $\text{oP}^{10}(\text{NH}_2)$ have been co-assembled with a series of rod-shaped linkers 1, 2, and 3. Assembly occurs effectively in the case of $\text{oP}^6(\text{NH}_2)$ to give [3 + 3] macrocycles. Analysis of NMR spectra shows that within the macrocycles, the *o*-phenylene moieties remain predominantly folded into the AAA state, with a greater proportion of perfectly folded oligomers than is observed for an analogous acyclic model compound ($\text{oP}^6(\text{M})$) but little stereocontrol. In contrast, $\text{oP}^{10}(\text{NH}_2)$ assembles in all three cases into a mixture of [2 + 2] and [3 + 3] macrocycles. In the [3 + 3] macrocycles, the behavior of the *o*-phenylene decamers directly parallels that of the hexamers and (previously reported) tetramers: an increased preference for the perfectly folded AAAAAA state, and again no evidence for stereocontrol. In the [2 + 2] macrocycles, the oligomers are forced to misfold, and instead adopt the AAAABBB state as their major conformation. Not only has this state not been previously observed (nor expected) for acyclic oligomers, but it is an unusual fusion of two helical blocks with distinct geometries and opposite twist senses.

The trade-off between inherent conformer stability (E_{rel}) and the bite angle (β) made by the linkages to the *o*-phenylene appears to be the key to understanding the relationship between folding and macrocyclization in these systems. Both of these parameters are predictable using *ab initio* calculations of conformer libraries of unsubstituted *o*-phenylenes. These libraries also provide a powerful framework for relating the NMR spectra to folding state *via* substituent-independent changes in chemical shifts ($\Delta\delta$), which provide a spectral fingerprint that can be used to identify new folding patterns.

The work described here does not yet represent a viable strategy toward high yields of structurally complex and well-defined species. Most importantly, we still lack control over the absolute and relative stereochemistry of the *o*-phenylene moieties. Nevertheless, the results show that foldamers within shape-persistent macrocycles can exhibit folding behavior that is distinctly different from the analogous acyclic species in solution. This strategy offers the potential of generating structural complexity efficiently and predictably through geometric restrictions on folding rather than, for example, monomer sequence. Studies of the effect of tuning the overall folding propensity on assembly and efforts to improve stereocontrol in these systems are ongoing.



Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We thank the National Science Foundation (CHE-1608213) for support of this work, the Volwiler Distinguished Research Professorship, and Dr Dominik Konkolewicz for the use of his analytical GPC.

Notes and references

- 1 S. H. Gellman, *Acc. Chem. Res.*, 1998, **31**, 173–180.
- 2 D. J. Hill, M. J. Mio, R. B. Prince, T. S. Hughes and J. S. Moore, *Chem. Rev.*, 2001, **101**, 3893–4011.
- 3 G. Guichard and I. Huc, *Chem. Commun.*, 2011, **47**, 5933–5941.
- 4 D.-W. Zhang, X. Zhao, J.-L. Hou and Z.-T. Li, *Chem. Rev.*, 2012, **112**, 5271–5316.
- 5 T. A. Martinek and F. Fülöp, *Chem. Soc. Rev.*, 2012, **41**, 687–702.
- 6 B. A. F. Le Bailly and J. Clayden, *Chem. Commun.*, 2016, **52**, 4852–4863.
- 7 S. De, B. Chi, T. Granier, T. Qi, V. Maurizot and I. Huc, *Nat. Chem.*, 2018, **10**, 51–57.
- 8 N. Delsuc, M. Hutin, V. E. Campbell, B. Kauffmann, J. R. Nitschke and I. Huc, *Chem.–Eur. J.*, 2008, **14**, 7140–7143.
- 9 K. L. George and W. S. Horne, *Acc. Chem. Res.*, 2018, **51**, 1220–1228.
- 10 N. Delsuc, S. Massip, J.-M. Léger, B. Kauffmann and I. Huc, *J. Am. Chem. Soc.*, 2011, **133**, 3165–3172.
- 11 H.-Y. Hu, J.-F. Xiang, Y. Yang and C.-F. Chen, *Org. Lett.*, 2008, **10**, 69–72.
- 12 L. Sebaoun, V. Maurizot, T. Granier, B. Kauffmann and I. Huc, *J. Am. Chem. Soc.*, 2014, **136**, 2168–2174.
- 13 P. Reiné, J. Justicia, S. P. Morcillo, S. Abbate, B. Vaz, M. Ribagorda, Á. Orte, L. Álvarez de Cienfuegos, G. Longhi, A. G. Campaña, D. Miguel and J. M. Cuerva, *J. Org. Chem.*, 2018, **83**, 4455–4463.
- 14 K. Urushibara, Y. Ferrand, Z. Liu, H. Masu, V. Pophristic, A. Tanatani and I. Huc, *Angew. Chem., Int. Ed.*, 2018, **57**, 7888–7892.
- 15 R. Katoono, Y. Tanaka, K. Fujiwara and T. Suzuki, *J. Org. Chem.*, 2014, **79**, 10218–10225.
- 16 R. Katoono, K. Kusaka, K. Fujiwara and T. Suzuki, *Chem.–Asian J.*, 2014, **9**, 3182–3187.
- 17 F. Chen, T. Tanaka, Y. Hong, W. Kim, D. Kim and A. Osuka, *Chem.–Eur. J.*, 2016, **22**, 10597–10606.
- 18 T. Hjelmgaard, L. Nauton, F. De Riccardis, L. Jouffret and S. Faure, *Org. Lett.*, 2018, **20**, 268–271.
- 19 N. Fuentes, A. Martin-Lasanta, L. Alvarez de Cienfuegos, R. Robles, D. Choquesillo-Lazarte, J. M. García-Ruiz, L. Martínez-Fernández, I. Corral, M. Ribagorda, A. J. Mota, D. J. Cárdenas, M. C. Carreño and J. M. Cuerva, *Angew. Chem., Int. Ed.*, 2012, **51**, 13036–13040.
- 20 H. Wu, A. Acharyya, Y. Wu, L. Liu, H. Jo, F. Gai and W. F. DeGrado, *ChemBioChem*, 2018, **19**, 902–906.
- 21 C. Tsiamantas, X. de Hatten, C. Douat, B. Kauffmann, V. Maurizot, H. Ihara, M. Takafuji, N. Metzler-Nolte and I. Huc, *Angew. Chem., Int. Ed.*, 2016, **55**, 6848–6852.
- 22 B. Liu, C. G. Pappas, E. Zangrando, N. Demitri, P. J. Chmielewski and S. Otto, *J. Am. Chem. Soc.*, 2019, **141**, 1685–1689.
- 23 T. Sawada, A. Matsumoto and M. Fujita, *Angew. Chem., Int. Ed.*, 2014, **53**, 7228–7232.
- 24 T. Sawada, M. Yamagami, K. Ohara, K. Yamaguchi and M. Fujita, *Angew. Chem., Int. Ed.*, 2016, **55**, 4519–4522.
- 25 B. A. G. Hammer and K. Müllen, *Chem. Rev.*, 2016, **116**, 2103–2140.
- 26 C. S. Hartley, *Acc. Chem. Res.*, 2016, **49**, 646–654.
- 27 S. Ando, E. Ohta, A. Kosaka, D. Hashizume, H. Koshino, T. Fukushima and T. Aida, *J. Am. Chem. Soc.*, 2012, **134**, 11084–11087.
- 28 E. Ohta, H. Sato, S. Ando, A. Kosaka, T. Fukushima, D. Hashizume, M. Yamasaki, K. Hasegawa, A. Muraoka, H. Ushiyama, K. Yamashita and T. Aida, *Nat. Chem.*, 2011, **3**, 68–73.
- 29 G. N. Vemuri, R. R. Pandian, B. J. Spinello, E. B. Stopler, Z. J. Kinney and C. S. Hartley, *Chem. Sci.*, 2018, **9**, 8260–8270.
- 30 Y. Tokoro, N. Ohtsuka, A. Kusakabe and S.-i. Fukuzawa, *Eur. J. Org. Chem.*, 2017, 2353–2358.
- 31 D. Lotter, A. Castrogiovanni, M. Neuburger and C. Sparr, *ACS Cent. Sci.*, 2018, **4**, 656–660.
- 32 A. Muraoka and H. Kiwaki, *Chem. Lett.*, 2018, **47**, 487–489.
- 33 D. Lehnher, C. Chen, Z. Pedramrazi, C. R. DeBlase, J. M. Alzola, I. Keresztes, E. B. Lobkovsky, M. F. Crommie and W. R. Dichtel, *Chem. Sci.*, 2016, **7**, 6357–6364.
- 34 Y. Mizukoshi, K. Mikami and M. Uchiyama, *J. Am. Chem. Soc.*, 2015, **137**, 74–77.
- 35 Z. J. Kinney and C. S. Hartley, *J. Am. Chem. Soc.*, 2017, **139**, 4821–4827.
- 36 R. Katoono, S. Kawai, K. Fujiwara and T. Suzuki, *Chem. Sci.*, 2015, **6**, 6592–6600.
- 37 H. Sato, J. A. Bender, S. T. Roberts and M. J. Krische, *J. Am. Chem. Soc.*, 2018, **140**, 2455–2459.
- 38 A. M. Castilla, W. J. Ramsay and J. R. Nitschke, *Acc. Chem. Res.*, 2014, **47**, 2063–2073.
- 39 A. U. Malik, F. Gan, C. Shen, N. Yu, R. Wang, J. Crassous, M. Shu and H. Qiu, *J. Am. Chem. Soc.*, 2018, **140**, 2769–2772.
- 40 G. Naulet, L. Sturm, A. Robert, P. Dechambenoit, F. Röhricht, R. Herges, H. Bock and F. Durola, *Chem. Sci.*, 2018, **9**, 8930–8936.
- 41 Z. J. Kinney and C. S. Hartley, *Org. Lett.*, 2018, **20**, 3327–3331.
- 42 Q. J. Zhou, K. Worm and R. E. Dolle, *J. Org. Chem.*, 2004, **69**, 5147–5149.
- 43 M. Kozáková, M. Buděšínský and J. Hodačová, *Synth. Commun.*, 2005, **35**, 161–167.
- 44 Y. B. Borzdina, E. Mostovich, V. Enkelmann, B. Wolf, P. T. Cong, U. Tutsch, M. Lang and M. Baumgarten, *J. Mater. Chem. C*, 2014, **2**, 6618–6629.
- 45 Although catalytic amounts of primary amines, which would be expected in the present systems even after quenching



- with NEt_3 , can catalyze transimination by themselves (ref. 46 and 47), previous work has shown that this is not significant under the conditions used here (ref. 35).
- 46 M. Ciaccia, S. Pilati, R. Cacciapaglia, L. Mandolini and S. Di Stefano, *Org. Biomol. Chem.*, 2014, **12**, 3282–3287.
- 47 M. Ciaccia, R. Cacciapaglia, P. Mencarelli, L. Mandolini and S. Di Stefano, *Chem. Sci.*, 2013, **4**, 2253–2261.
- 48 The macrocycles were carefully purified through multiple GPC injections; thus, the isolated yields are likely low compared to the actual efficiency of assembly.
- 49 S. M. Mathew, J. T. Engle, C. J. Ziegler and C. S. Hartley, *J. Am. Chem. Soc.*, 2013, **135**, 6714–6722.
- 50 S. M. Mathew and C. S. Hartley, *Macromolecules*, 2011, **44**, 8425–8432.
- 51 C. Yu, Y. Jin and W. Zhang, *Dynamic Covalent Chemistry*, John Wiley & Sons, Ltd, Chichester, UK, 2017, pp. 121–163.
- 52 R. Chakrabarty, P. S. Mukherjee and P. J. Stang, *Chem. Rev.*, 2011, **111**, 6810–6918.
- 53 W. Zhang and J. S. Moore, *Angew. Chem., Int. Ed.*, 2006, **45**, 4416–4439.
- 54 J. Bunzen, J. Iwasa, P. Bonakdarzadeh, E. Numata, K. Rissanen, S. Sato and M. Fujita, *Angew. Chem., Int. Ed.*, 2012, **51**, 3161–3163.
- 55 T. P. Money Penny II, A. Yang, N. P. Walter, T. J. Woods, D. L. Gray, Y. Zhang and J. S. Moore, *J. Am. Chem. Soc.*, 2018, **140**, 5825–5833.
- 56 S. Grimme, *J. Comput. Chem.*, 2006, **27**, 1787–1799.
- 57 S. E. Wheeler, *Acc. Chem. Res.*, 2013, **46**, 1029–1038.
- 58 G. M. Whitesides, J. P. Mathias and C. T. Seto, *Science*, 1991, **254**, 1312–1319.
- 59 C. S. Hartley and J. He, *J. Org. Chem.*, 2010, **75**, 8627–8636.
- 60 NMR spectra throughout this work have been run at 0 °C, which typically gives slightly sharper peaks.
- 61 K. W. Wiitala, T. R. Hoye and C. J. Cramer, *J. Chem. Theory Comput.*, 2006, **2**, 1085–1092.
- 62 R. Jain, T. Bally and P. R. Rablen, *J. Org. Chem.*, 2009, **74**, 4017–4023.
- 63 S. Mathew, L. A. Crandall, C. J. Ziegler and C. S. Hartley, *J. Am. Chem. Soc.*, 2014, **136**, 16666–16675.
- 64 S. Toyota, *Chem. Rev.*, 2010, **110**, 5398–5424.
- 65 In the spectrum of $\text{oP}^6(\mathbf{M})$, individual signals vary by nearly 1 ppm over different conformers.
- 66 *o*-Phenylene tetramers are in fast backbone conformational exchange on the NMR time scale.
- 67 We do not expect there to be significant differences in entropy between specific foldamer conformers (beyond those associated with their symmetries) since they are composed of rigid subunits and conformationally well-defined. Likewise, the behavior of the linkers is not expected to be strongly dependent on macrocycle size; this is consistent with the very similar outcomes of assembly for all systems examined thus far (*e.g.*, as assessed by GPC).
- 68 The ^1H NMR spectrum of $\text{oP}^{10}(\text{DPB})_{2+2}$ was also acquired in acetone- d_6 . The appearance was very similar to that in CDCl_3 , suggesting at least qualitatively similar folding behavior in both solvents.
- 69 Here, “significant” signals are defined as those with cross-peaks observable in an HSQC spectrum and related signals from an HMBC spectrum.
- 70 P. T. Corbett, J. Leclaire, L. Vial, K. R. West, J.-L. Wietor, J. K. M. Sanders and S. Otto, *Chem. Rev.*, 2006, **106**, 3652–3711.
- 71 Y. Ito, E. Ihara, M. Murakami and M. Sisido, *Macromolecules*, 1992, **25**, 6810–6813.
- 72 Y. Nagata, H. Hasegawa, K. Terao and M. Sugimoto, *Macromolecules*, 2015, **48**, 7983–7989.

