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# Pd- $\eta^3$ -C<sub>6</sub>H<sub>9</sub> complexes of the Trost modular ligand: high nuclearity columnar aggregation controlled by concentration, solvent and counterion†

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Under optimised conditions, the Trost modular ligand (TML) series induces high levels of asymmetric induction in an extraordinarily wide range of reactions involving palladium  $\pi$ -allyl intermediates. Prior mechanistic investigations into reactions involving Pd- $\eta^3$ -C<sub>6</sub>H<sub>9</sub> intermediates have focussed on the monomeric 13-membered ring formed via P,P-chelation of the ligand to Pd. However, it is also recognised that ring-opening oligomerisation provides a pool of high nuclearity Pd- $\eta^3$ -C<sub>6</sub>H<sub>9</sub> species that, by affording a low level, or even the opposite sense, of asymmetric induction relative to the mononuclear species, are responsible for a reduction in selectivity under non-optimised conditions. Herein we describe an investigation by NMR spectroscopy, molecular mechanics, molecular dynamics, and small-angle neutron scattering (SANS), of a Pd- $\eta^3$ -C<sub>6</sub>H<sub>9</sub> cation bearing the 1,2-diaminocyclohexane TML ligand (2). Using both nondeuterated and perdeuterated (D<sub>47</sub>) isotopologues of the resulting complexes ([1]<sup>+</sup>), we show that a two-stage oligomerisation-aggregation process forms self assembled cylindrical aggregates of very high nuclearity (up to 56 Pd centres). We also investigate how concentration, solvent and counter-anion all modulate the extent of oligomerisation.

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## Introduction

The Trost modular ligand (TML) series<sup>1</sup> has been applied to an extraordinarily wide range of allylic alkylation (Tsuji-Trost) reactions.<sup>2</sup> Under carefully optimised conditions, these ligands frequently provide very high enantioselectivity in reactions that have proven challenging with other chiral ligands, particularly those involving cyclic allylic substrates.<sup>3–6</sup> These features have led to broad use of the TML in the synthesis of natural products,<sup>7</sup> as well as industrial application for the construction of

high-enantiopurity chiral building blocks. However, reactions involving the TML can exhibit memory effects,<sup>8,9</sup> and a high sensitivity of the enantioselectivity to reaction temperature, catalyst concentration, solvent and nucleophile counter-ion.



Our previous mechanistic studies of this system focussed on the monomeric cationic complex [1]<sup>+</sup>, in which a Pd( $\eta^3$ -C<sub>6</sub>H<sub>9</sub>) unit is chelated by the 1,2-diaminocyclohexane-derived TML ligand (2).<sup>10</sup> The monomeric cation [1]<sup>+</sup> was identified as an intermediate capable of leading to high asymmetric induction on attack of, for example, a malonate anion nucleophile, Scheme 1. Detailed NMR studies facilitated by isotopic labelling – in conjunction with MM-DFT simulations – led to a model<sup>10</sup> in which the amide units in the catalyst facilitate enantioselective ligand-accelerated catalysis.<sup>11</sup>

For the ligand-accelerated catalysis to function efficiently, cation [1]<sup>+</sup> requires a degree of flexibility. This flexibility is provided by the 13-membered chelate ring, but at a cost: complex [1]<sup>+</sup> can readily undergo ring-opening oligomerisation

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**Scheme 1** Asymmetric allylic alkylation of racemic 2-cyclohexenyl acetate; **2** = 1,2-diaminocyclohexane TML ligand, THAB = tetrahexylammonium bromide.



**Scheme 2** Oligomerisation of  $[1]^+$  erodes enantioselectivity during asymmetric Pd-catalysed allylic alkylation mediated by **2** (as Scheme 1). Nu = nucleophile.

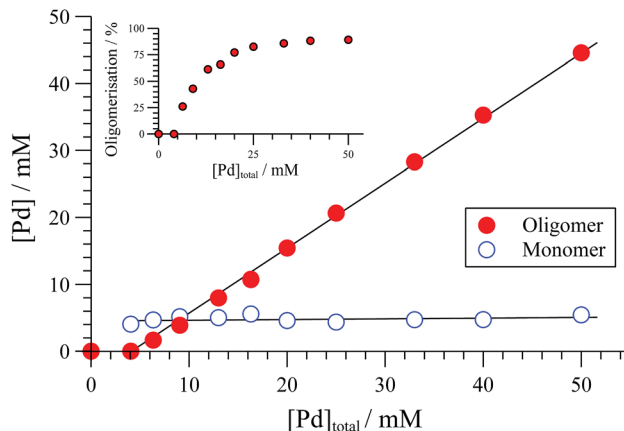
to generate polynuclear species  $([1]^+)_n$ , Scheme 2.<sup>12</sup> Competing nucleophilic attack on the oligomer, rather than the monomer  $[1]^+$  is, in part, responsible for a reduction in overall enantioselectivity under non-optimised conditions.<sup>13,14</sup>

To date, the structure and origin of the formation of these oligomeric species has not been studied in detail. Herein, we describe an investigation of the oligomerisation of  $D_0$  and  $D_{47}$  isotopologues of  $[1]^+$ , employing NMR spectroscopy, molecular mechanics (MM), molecular dynamics (MD), and contrast variation small-angle neutron scattering (SANS). The data obtained indicate that the impact of a first-stage of depletion of the monomeric species  $[1]^+$  from the catalyst pool, *via* cyclic oligomerisation, is amplified by a second-stage process involving columnar aggregation of the oligomers, leading to species with very high nuclearity (up to 56 Pd centres). The effects of solvent, ligand enantiopurity and counter-ion on the degree of aggregation are explored in detail, and it is concluded that a relatively small and restricted set of conditions facilitate dissolution of the complexes in a low aggregation state, consistent with the extensive optimisation frequently required for these catalyst systems.

## Results and discussion

### Preliminary NMR studies and synthesis of $[D_{47}]\text{-}[1][B(C_6F_5)_4]$

Despite extensive efforts,<sup>15</sup> we have been unable to crystallise any  $Pd(\eta^3\text{-C}_6\text{H}_5)$  complexes of **2**, in either oligomeric or monomeric forms. Indeed, to date, the only the X-ray crystal structures of Pd-



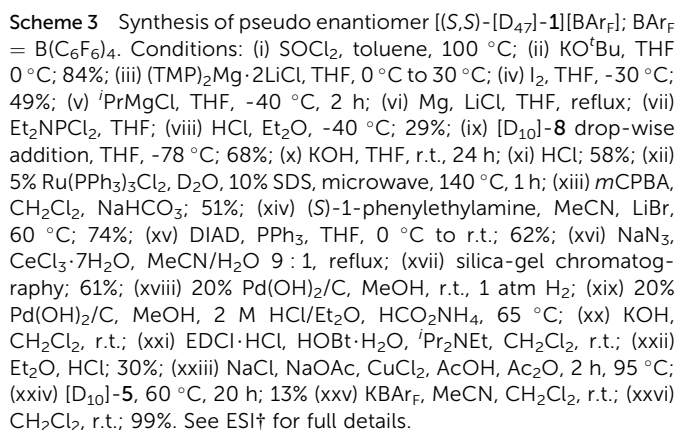
**Fig. 1** Speciation of  $[(R,R)\text{-}1][\text{BAR}_F]$  ( $\text{BAR}_F = \text{B}(\text{C}_6\text{F}_5)_4$ ) determined by  $^{31}\text{P}\{^1\text{H}\}$  NMR ( $\text{CD}_2\text{Cl}_2$ ,  $25^\circ\text{C}$ ). Solid lines through data are solely a guide to the eye. Inset shows proportion (%) of  $[\text{Pd}]_{\text{tot}}$  that is in oligomeric form.

allyl complexes of ligand **2**<sup>16</sup> are  $\eta^3\text{-C}_3\text{H}_5$  complexes with triflate counter anions: one a racemic tetranuclear cyclo-oligomer,<sup>12</sup> the other an acyclic dinuclear *bis-P,O*-chelate.<sup>17</sup>

The extent of solution-phase oligomerisation of cationic complexes of type  $[1]^+$  can be conveniently estimated by  $^{31}\text{P}\{^1\text{H}\}$  NMR spectroscopy.<sup>10,12</sup> Analysis of  $[(R,R)\text{-}1][\text{BAR}_4]$  complexes in  $\text{CH}_2\text{Cl}_2$ , where Ar =  $\text{C}_6\text{Cl}_5$ , 3,5- $(\text{CF}_3)_2\text{C}_6\text{H}_3$ , or  $\text{C}_6\text{F}_5$  (“ $\text{BAR}_F$ ”), indicates a maximum monomer concentration ( $[1]^+$ ) of about 4 mM, Fig. 1. In THF, the monomer maximum is lower (approx. 1.6 mM) and decreases as  $[\text{Pd}]_{\text{tot}}$  is raised above 10 mM. With smaller, less charge-diffuse, counter-anions such as chloride or triflate, the maximum monomer concentrations are lower still. We were unable to fit simple analytical solutions<sup>18</sup> for monomer-oligomer distributions to any of the  $^{31}\text{P}$  NMR data, indicative that physicochemical effects dominate over simple solution-phase equilibria, even at low  $[\text{Pd}]_{\text{tot}}$ .

We thus elected to study the oligomeric species by SANS – a technique that can be used for characterising the shape and dimensions of self-assembly structure and colloids.<sup>19</sup> We began with  $[(R,R)\text{-}1][\text{BAR}_F]$ ,<sup>20</sup> and, to aid the studies, also synthesised the perdeuterated enantiomeric complex  $[(S,S)\text{-}[D_{47}]\text{-}1][\text{BAR}_F]$ . Not only does this facilitate SANS in a non-deuterated solvent, thus providing greater neutron scattering contrast,<sup>19</sup> it also allows pseudo racemic and pseudo scalemic mixtures to be prepared by mixing  $[(S,S)\text{-}[D_{47}]\text{-}1][\text{BAR}_F]$  with  $[(R,R)\text{-}1][\text{BAR}_F]$ . The perdeuterated complex was synthesised from benzoic acid ( $[D_5]\text{-}3$ ), chlorobenzene ( $[D_5]\text{-}4$ ) and cyclohexene ( $[D_{10}]\text{-}5$ ), Scheme 3. A major hurdle was the *ortho*-metallation of ester  $[D_5]\text{-}6$  with  $(\text{TMP})_2\text{Mg}\cdot\text{LiCl}$ ,<sup>21</sup> which proceeded with an unexpectedly large net kinetic isotope effect ( $k_H/k_D \approx 30$ ).<sup>22</sup> This required an excess of base to be employed, and interfered with a planned direct phosphination of the metallated intermediate. Instead, the intermediate was trapped with  $\text{I}_2$ . The iodide  $[D_4]\text{-}7$  was then converted to a more conventional Grignard reagent,<sup>23</sup> before reaction with chlorophosphine  $[D_{10}]\text{-}8$ <sup>10,24</sup> to give phosphine  $[D_{14}]\text{-}9$ , and thus acid  $[D_{14}]\text{-}10$ .<sup>25</sup>

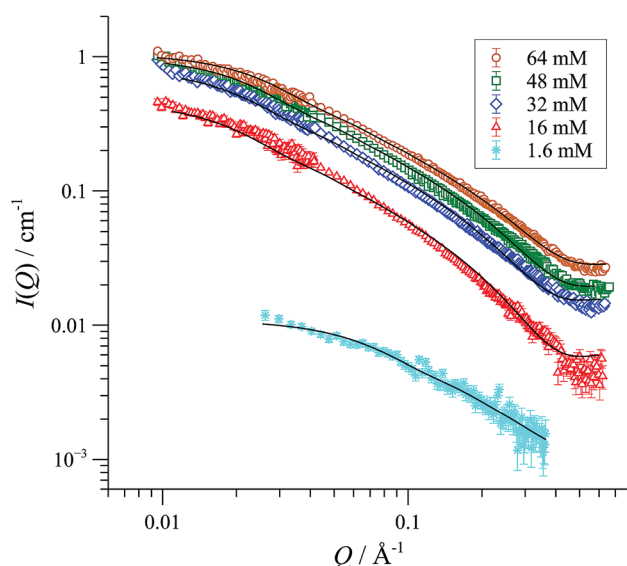




### SANS analysis of aggregation of $[1][\text{BAR}_\text{F}]$ in THF

We began with SANS analysis<sup>32</sup> of enantiopure  $[(R,R)\text{-1}][\text{Bar}_\text{F}]$  in THF- $\text{D}_8$  at 25 °C, with  $[\text{Pd}]_\text{tot}$  concentrations [6 to 64 mM, well above the oligomerisation threshold. The combined data sets,<sup>33</sup> presented here as plots of scattered intensity ( $I(Q)$ , y-axis) *versus* momentum transfer ( $Q$ , x-axis) were analysed with a range of different standard models for possible aggregate form factors, corresponding to various simple shapes,<sup>19</sup> Fig. 2.<sup>34</sup> This clearly identified the particles as cylindrical or rod-like, and Guinier analyses (see ESI, Fig. S69†) established the radii (8-9 Å) and lengths (150-200 Å) as essentially invariant across the range of  $[\text{Pd}]_\text{tot}$  explored. In other words, the number density of aggregates changes in response to  $[\text{Pd}]_\text{tot}$ , but not their average dimensions,<sup>35</sup> clearly indicative of a set of factors that tightly control the particle scale.

We have previously used  $^{31}\text{P}$  NMR spectroscopy to analyse the constitution of the solution-phase (*i.e.*, lower-order) oligomers generated from various complexes of type  $[\mathbf{1}][\text{BAR}_4]$  in  $\text{CD}_2\text{Cl}_2$ .<sup>10,12</sup> Using PPCOSY in combination with pairs of isotopically-differentiated ligands ( $[\text{D}_n\text{-}2]$ ), we were able to determine



**Fig. 2** SANS profiles of 1.6 to 64 mM (0.25–10.0 wt/vol%)  $\{[(R,R)\text{-}1][\text{BARf}]\}_n$  in THF- $\text{D}_8$ , 25 °C;  $I(Q)$  = scattered intensity,  $Q/\text{\AA}^{-1}$  = momentum transfer =  $(4\pi/\lambda)\sin(\theta/2)$ ; where scattering angle =  $\theta$  and neutron wavelength =  $\lambda$ . The fitted curves correspond to a cylindrical particle form factor. The sample with 1.6 mM  $[\text{Pd}]_{\text{tot}}$  has form factor fitted to radius  $\leq 11$  Å and length  $\leq 50$  Å. All other samples are fitted to radius 8–9 Å and length 150–200 Å.

that the oligomers are: i) non-chelated species (*i.e.* each of the ligands (**2**) in the oligomer are coordinated to two different Pd centres); ii) present in predominantly homochiral form (*i.e.*  $[(R,R)\text{-}1]^+$  and  $[(S,S)\text{-}1]^+$  oligomerise independently), and iii) contain no 'free' (*i.e.* not Pd-coordinated) P-centres in the ligand (**2**). Although a cyclic oligomer structure (Scheme 2) is fully consistent with these features, we were unable to determine the number ( $n$ ) of ring-opened monomer units incorporated within the cyclo-oligomer  $[(1)[\text{BAR}_4]]_n$ .

As  $[\text{Pd}]_{\text{tot}}$  in THF or  $\text{CH}_2\text{Cl}_2$  solutions of complexes of type  $[1][\text{BAR}_4]$  is increased, the  $^{31}\text{P}$  NMR bandshape of the signals arising from the cyclo-oligomer do not change in appearance, but the samples do become increasingly turbid. This behaviour suggests that in response to an increase in  $[\text{Pd}]_{\text{tot}}$ , cyclo-oligomers  $[(1)[\text{BAR}_4]]_n$  do not incorporate more monomer ( $n$ ), but instead aggregate to form larger particles,  $\{[(1)[\text{BAR}_4]]_n\}_m$ , containing ' $m$ ' cyclo-oligomers. It is these high nuclearity particles that are detected by SANS.<sup>12</sup>

In racemic or scalemic samples of  $[1][\text{BAR}_F]$ , the homochiral cyclo-oligomers,  $[(1)[\text{BAR}_F]]_n$ , could aggregate in three general forms: discrete homochiral, ordered heterochiral (*e.g.*, alternating or co-block), or statistically distributed. SANS data of mixtures of  $[(R,R)\text{-}1][\text{BAR}_F]$  and  $[(S,S)\text{-}1][\text{BAR}_F]$  representing enantiopure, scalemic, and racemic samples, was uniform across the series, within experimental error, Fig. 3. The absence of a change in particle number density<sup>36</sup> is consistent with a statistical cyclo-oligomer distribution in which there is no significant impact of homo- or hetero-chirality on the particle shape or size.

SANS data of the pseudo racemate  $[(S,S)\text{-}[\text{D}_{47}\text{-}1][\text{BAR}_F] + [(R,R)\text{-}[\text{D}_0\text{-}1][\text{BAR}_F]]$  in  $\text{H}_8\text{-THF}$ , and in  $\text{D}_8\text{-THF}$ , Fig. 4, and enantiomerically pure  $[(S,S)\text{-}[\text{D}_{47}\text{-}1][\text{BAR}_F]$  in  $\text{D}_0\text{-THF}$ , show that the fully and partially deuterated systems retain the concentration-independent, cylindrical aggregate shape. A major difference is, however, observed in the dimensions:  $[\text{D}_{47}\text{-}1][\text{BAR}_F]$  forms shorter (130 Å), slightly wider (10 Å radius) cylinders than  $[\text{D}_0\text{-}1][\text{BAR}_F]$ . The pseudo racemic mixture measures as an average of its precursors (9–10 Å radius, 150 Å length),



Fig. 3 Fitted SANS profiles and cylinder form factor fits for 32 mM enantio-pure, racemic and scalemic samples of  $[1][\text{BAR}_F]$  in  $\text{THF-D}_8$ , 25 °C.

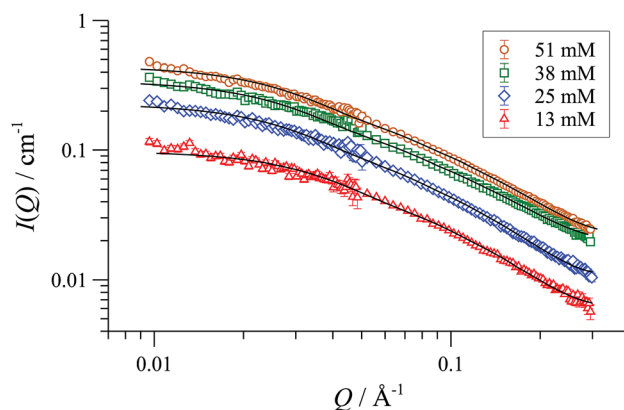


Fig. 4 SANS data and cylinder form factor fits for pseudo racemic complex  $[(S,S)\text{-}[\text{D}_{47}\text{-}1][\text{BAR}_F] + [(R,R)\text{-}[\text{D}_0\text{-}1][\text{BAR}_F]]$  in  $\text{THF-D}_8$  at 25 °C, at  $[\text{Pd}]_{\text{tot}}$  ranging from 13 to 51 mM.

again consistent with a statistical distribution of cyclo-oligomers in the aggregate.<sup>37</sup>

### Molecular mechanics (MM) and dynamics (MD)

Computational modelling was employed to probe the factors that control the aggregation phenomena, and to estimate the size ( $n$ ) and number of cyclo-oligomers in the particle ( $m$ ).<sup>38</sup> The number of atoms in the aggregates ( $>2,000$ , *vide infra*) means that density functional theory (DFT) calculations of their structures would demand currently unattainable computational resources and inordinate simulation times. On the other hand, molecular mechanics (MM) can provide a good approximation in just a small fraction of the computational time required by DFT. Although MM in general calculates only a steric energy, not the full free energy, and ignores bond dissociation energies, accurate results can be obtained from MM3 calculations provided that comparisons are isodesmic and, more importantly, isoparametric.<sup>39</sup> This requirement is fulfilled for all comparisons of ring oligomers and their non-covalent aggregates. We began by confirming that the structure of the 84-atom monomeric cationic P,P-chelate  $[1]^+$ , optimised at MM3 level of theory with dielectric constant  $\epsilon = 9.0$ , was almost identical to that obtained using DFT (B3LYP-D3) in a polarisable continuum model for dichloromethane.<sup>38</sup> Further calculations involving ion-pairs, oligomers and aggregates were then performed using MM3 to provide analysis of the energies involved, albeit at a coarse-grained level of detail.

Comparison of the MM3 optimised energies for monomeric P,P-chelate  $[1][\text{BAR}_F]$  with a series of homochiral cyclo-oligomers,  $[(1)[\text{BAR}_F]]_n$ , normalised by the number of Pd atoms ( $n$ , the  $x$ -axis in Fig. 5) confirmed cyclo-oligomerisation to be exothermic, and probably also exergonic up to tetramers ( $n = 4$ ). The cyclic dimer ( $n = 2$ ) still suffers from ring strain, and a more substantial stability is afforded by trimerisation ( $n = 3$ ) then tetramerisation<sup>40</sup> ( $n = 4$ ).

Further increase in oligomer ring size ( $n = 5, 6, 8$ ) yields a modest reduction in the system energy but generates species with significantly higher radii than the 8–9 Å cylinder radius detected by SANS. In summary, the tetranuclear species ( $n = 4$ ),





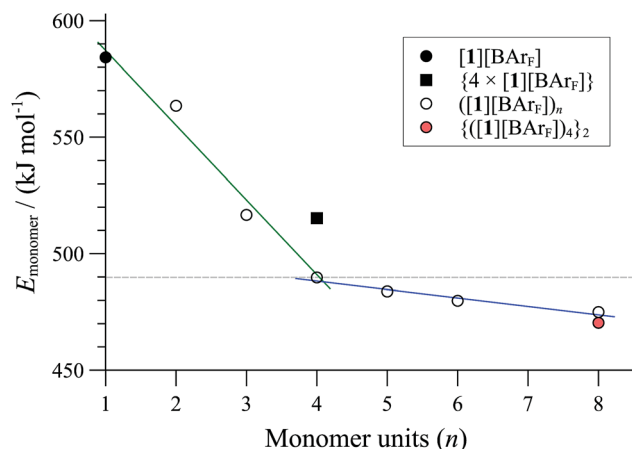


Fig. 5 MM3 energies ( $\epsilon = 9.0$ ) of cyclo-oligomeric complexes  $([1][\text{BARF}])_n$  as a function of monomers incorporated ( $n$ , x-axis).  $\{4 \times ([1][\text{BARF}])\}$  is a pre-orientated assembly of four monomeric P,P-chelates;  $\{([1][\text{BARF}])_4\}_2$  is an aggregate formed from two tetrameric cyclo-oligomers (see  $m = 2$  in Fig. 7).

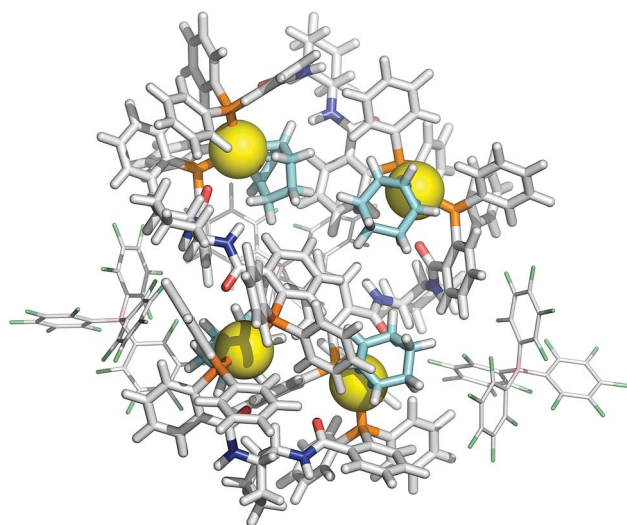


Fig. 6 MM3 structure of tetranuclear cyclo-oligomer  $([1][\text{BARF}])_4$ . For clarity, only two of the four  $\text{BARF}$  anions are shown. Colour coding: Pd, yellow; P, orange; O, red; N, dark blue; C, light grey; B, pink; F, green;  $\eta^3\text{-C}_6\text{H}_9$ , light blue.

Fig. 6, appears to be the dominant, if not exclusive, cyclo-oligomer, and aggregation of these cyclic tetramers  $([1][\text{BARF}])_4$  was thus probed by MM3 as a process to generate the cylindrical particles, Fig. 7.

Positively charged rod-like structures with dissociated or removed anions were estimated by MM3 to be very much higher in energy than those where the anions were closely associated with the cationic cyclo-oligomeric building blocks. Anion interactions were thus explored more deeply, and although  $\text{BARF}$  is considered a weakly coordinating anion,<sup>41</sup> the charge delocalisation over its surface reduces repulsive interactions with other  $\text{BARF}$  anions, and the presence of fluorine makes it significantly lipophilic. Indeed, the calculations indicated a favourable interleaving of the  $\text{BARF}$  anions in sandwich layers<sup>42</sup>



Fig. 7 MM3 energies ( $\epsilon = 9.0$ ) of  $\{([1][\text{BARF}])_4\}_m$  (normalised per monomer), as a function of number ( $m$ ) of cyclo-oligomeric tetramers aggregated.

between cationic cyclo-oligomers. The estimated formation energies, ( $\Delta E$ , Fig. 7) of such species  $\{([1][\text{BARF}])_4\}_m$  as a function of ' $m$ ' indicated that columnar aggregates are readily attainable, with the growing entropic cost ( $T\Delta S$ ) placing limits on the aggregate length.<sup>34,40</sup>

These conclusions were further probed by molecular-dynamics (MD) simulations in which the MM3-minimised structures  $\{([1][\text{BARF}])_4\}_m$  were computationally excited (300-700 K) over short periods (300 ps) to test the *relative* structural integrity of the aggregate as a function of ' $m$ '. In the low dielectric constant medium used for the model, most systems ( $m = 4$  to 16) did not undergo any significant changes in their tertiary structure at 300 K. As the energy input was increased the aggregate models exhibited varying degrees of structural deformation, undergoing rapid fragmentation at the highest energies. The most significant observations were made at intermediate energies (500-550 K): aggregates with  $m = 10$ -14 (*e.g.*, Fig. 8,  $m = 12$ ) retained a cylinder shape, albeit mildly distorted, over the full 300 ps simulation time, whereas higher or lower order aggregates significantly



Fig. 8 MM3 structure of  $\{([1][\text{BARF}])_4\}_{12}$ . The monomer  $[1][\text{BARF}]$  (top left), with palladium coloured orange, is shown for scale.



deformed, in some cases losing one or more  $\text{BAR}_\text{F}$  anions. The average dimensions of the MM3 aggregates with  $m = 10\text{--}14$  (radius  $8\text{--}9\text{ Å}$  and length  $150\text{--}200\text{ Å}$ ) are consistent with the particle dimensions determined by SANS, Fig. 2 and 3.

### The effect of counter-ion and solvent on shape and extent of aggregation

The effect of solvent type on aggregation was probed by MM,  $^{31}\text{P}$   $\{^1\text{H}\}$  NMR (Fig. 9) and SANS, also comparing  $[\mathbf{1}][\text{BAR}_\text{F}]$  with  $[\mathbf{1}][\text{OTf}]$  to explore the impact of counter-ion. The aggregation mode for  $[\mathbf{1}][\text{BAR}_\text{F}]$  determined by MM3, e.g., Fig. 8, involves multiple close-range electrostatic interactions that reduce the overall energy of the system. It is therefore not surprising that the solvent dielectric constant ( $\epsilon_r$ ) was found to modulate aggregation,<sup>43</sup> and thus also solubility, with precipitation being the ultimate consequence of strong aggregation.

As indicated in Fig. 9, both  $[\mathbf{1}][\text{OTf}]$  and  $[\mathbf{1}][\text{BAR}_\text{F}]$  readily oligomerise and aggregate in all of the solvents that were explored, becoming essentially insoluble at the extremes of  $\epsilon_r$ , (e.g., in alkanes, most ethers, chloroform, aromatic hydrocarbons, and at the opposite end of the scale, in water). The lipophilicity and charge-density of the anion also affects the solubility:  $[\mathbf{1}][\text{BAR}_\text{F}]$  (but not  $[\mathbf{1}][\text{OTf}]$ ) readily dissolves in THF, and at the opposite end of the  $\epsilon_r$  scale,  $[\mathbf{1}][\text{OTf}]$  (but not  $[\mathbf{1}][\text{BAR}_\text{F}]$ ) is soluble in aqueous-organic mixtures.

SANS was employed to explore how the macromolecular composition of aggregates  $\{([\mathbf{1}][\text{X}])_n\}_m$  is affected by solvation. Although  $[\mathbf{1}][\text{BAR}_\text{F}]$  is not soluble in organic-aqueous mixtures, SANS data were attainable in polar aprotic solvents (e.g., MeCN,  $\epsilon_r = 37.5$ ; and DMSO,  $\epsilon_r = 47$ ). This confirmed that cylindrical aggregates were still formed, but were significantly shorter than those in THF, Fig. 10. Medium length cylinders were detected in a 50 : 50 mixture of THF and acetonitrile, consistent with the intermediate solvent polarity ( $\epsilon_r \approx 23$ ). In all cases, the cylinders were of radius  $8\text{--}10\text{ Å}$ , strongly suggesting the prevalence of the tetranuclear cyclo-oligomer building blocks, with the solvent modulating only the aggregation number ' $m$ ':  $\{([\mathbf{1}][\text{BAR}_\text{F}])_4\}_m$ .



Fig. 10 SANS data and cylinder form factor fits for  $[\mathbf{1}][\text{BAR}_\text{F}]$  in various solvents at  $25\text{ °C}$ .

The  $[\mathbf{1}][\text{OTf}]$  aggregates behaved differently. Although, cylinders of radius  $8\text{--}10\text{ Å}$  were again detected in all cases, indicative of  $\{([\mathbf{1}][\text{OTf}])_4\}_m$  aggregates, the flexibility, lengths and charge distribution in the particles were very different to those formed from  $[\mathbf{1}][\text{BAR}_\text{F}]$ .

In  $\text{CD}_2\text{Cl}_2$ ,  $[\mathbf{1}][\text{OTf}]$  forms cylindrical aggregates (up to  $160\text{ Å}$ ; Fig. 11) apparently with a degree of flexibility, a phenomenon that can be attributed to the small and interactive triflate anion being less able to rigidify the structures than the larger and more lipophilic  $\text{BAR}_\text{F}$  anion. The anion effect became even more pronounced in media of higher dielectric constant. In acetonitrile-based solvent mixtures ( $\epsilon_r = 47\text{--}58$ ) the SANS data indicated an additional minor structure factor contribution ( $S(Q)$ ), consistent with weakly charged particles, Fig. 12. Weak repulsive interactions might arise from solvation-induced ion-pair separation of triflate from the cationic  $\text{Pd}(\text{II})$  oligomeric cores. The increased cationic repulsion between the cyclo-oligomers appears to result in much shorter cylinders, just  $30\text{ Å}$  in length, with misleadingly simple  $^{31}\text{P}$  NMR spectra.<sup>44</sup> Similar conclusions were drawn from MD simulations with the medium set at

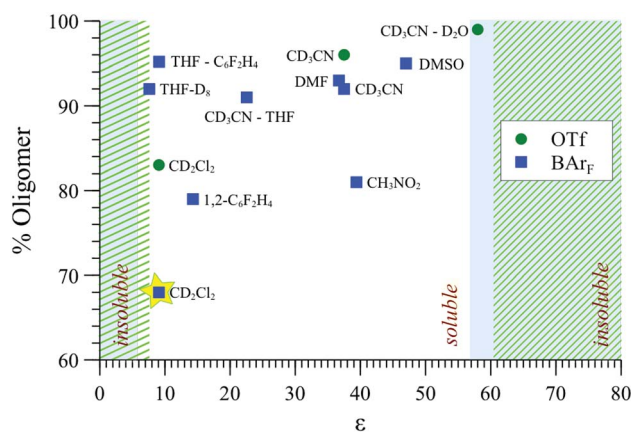


Fig. 9 Oligomerisation ( $^{31}\text{P}\{^1\text{H}\}$  NMR, %) of  $[\mathbf{1}][\text{BAR}_\text{F}]$  and  $[\mathbf{1}][\text{OTf}]$  at  $[\text{Pd}]_{\text{tot}} = 15\text{ mM}$  in solvents of various dielectric constant ( $\epsilon_r$ ); overlaid areas in green (OTf) and blue ( $\text{BAR}_\text{F}$ ) indicate regions where the complexes are insoluble.



Fig. 11 SANS data and flexible cylinder form factor fits for  $[\mathbf{1}][\text{OTf}]$  in  $\text{CD}_2\text{Cl}_2$  at  $25\text{ °C}$ ; fits are for radius  $8\text{--}9\text{ Å}$  and lengths:  $78\text{ Å}$  ( $13\text{ mM}$ );  $156\text{ Å}$  ( $25\text{ mM}$ ) and  $332\text{ Å}$  ( $50\text{ mM}$ ).



Fig. 12 SANS data and cylinder form factor fits (including an effective Hayter-Penfold structure factor  $S(Q)$  to account for charged particles) for  $[1][OTf]$  in  $CD_3CN$  at  $25^\circ C$ .

$\epsilon_r = 35$ : only the shortest aggregates  $\{([1][OTf])_4\}_{2-4}$  were structurally stable at elevated energies (500 K; 300 ps). All higher aggregates ( $m > 4$ ) underwent rapid fragmentation.

Finally, to probe the relevance of the higher aggregates to asymmetric alkylation (Scheme 1) SANS data were acquired on reaction mixtures in which  $[1][BARf]$  was employed as a pre-catalyst (10 mol%) for addition of tetrabutylammonium dimethylmalonate to cyclohexenylacetate in THF. While the effects of substrate background scattering, varying acquisition times and shorter Q-range slightly affected the data quality, it remained clear that the dominant structures in solution, for the whole duration of the catalytic process, were large cylinders. This result is consistent with previous conclusions that, in THF, the catalytic turnover proceeds *via* a small pool of highly-active monomeric catalyst species, in competition with cyclo-oligomers and aggregates.<sup>10,12</sup>

## Conclusions

Since the initial report that  $[1]^+$  readily oligomerises,<sup>13</sup> and that this is a largely undesirable property of an otherwise highly efficient catalyst, *e.g.*, Scheme 1, there has been limited understanding of the oligomer structures.<sup>10,12</sup> We have now identified, through NMR spectroscopy, SANS, and MM/MD simulations of  $[D_n]-1[BARf]$  ( $n = 0, 47$ ),<sup>45</sup> that monomeric cations  $[1]^+$  undergo chelate-opening to form a tetranuclear cyclo-oligomer  $([1][BARf])_4$ ; this being thermodynamically favoured over higher and lower nuclearity species. A second-stage process involving columnar-like interactions between the cyclo-oligomers then forms aggregates  $\{([1][BARf])_4\}_m$ . In THF these aggregates comprise alternating layers of cyclo-oligomer and interleaved  $BARf$  anions in the form of cylindrical particles ( $m = 10-14$ ; radius 8–9 Å; length 150–200 Å) containing up to 56  $Pd(\eta^3-C_6H_9)$  centres, ligands (2) and  $BARf$  anions.

The identity of the counter-anion has a pronounced effect on the proportion of oligomer generated from the monomer. Bulky, weakly-coordinating anions,<sup>46</sup> reduce the extent of oligomerisation, particularly in low polarity solvents that cannot

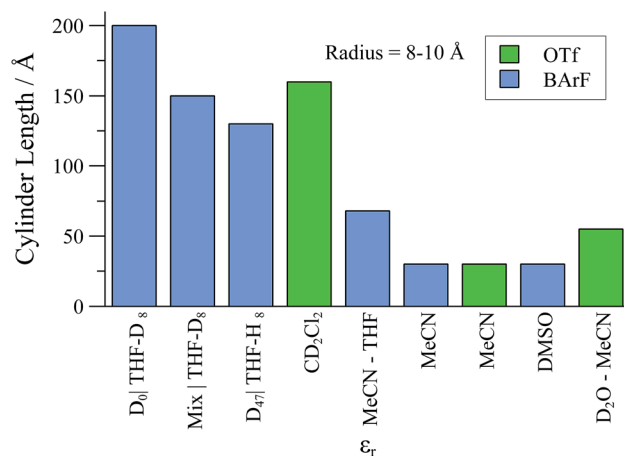


Fig. 13 Summary of upper-range aggregate cylinder lengths of  $[1][BARf]$  and  $[1][OTf]$  samples as determined by SANS experiments, in selected solvents., with  $[Pd]_{tot} = 25$  mM.

effectively stabilise charged particles. Here the role of the bulky and relatively lipophilic anions is to solvate the monomer  $[1]^+$ . Smaller harder, less lipophilic anions are less able to solvate the monomer, and have the indirect effect of shifting the equilibrium towards the oligomer; an undesirable feature for catalysis. The diminutive size of the anion also results in greater flexibility of the resulting columnar aggregates, which are more ionic in nature, reducing their solubility in less polar solvents.

Overall, although the solvent polarity, counter-anion (X), and net concentration ( $[Pd]_{tot}$ ) all affect the degree of oligomerisation and aggregation (Fig. 13) of monomer  $[1][X]$ , the solvent perhaps offers the greatest degree of scope for optimisation under the conditions of catalysis. In this regard,  $CH_2Cl_2$  is favourable: solutions can be virtually free of oligomer at ambient temperature, provided  $[Pd]_{tot} \leq 4$  mM. Intriguingly, SANS studies of ionic and non-ionic surfactants<sup>47</sup> have revealed specific solvent combinations that can lead to “dead zones” where aggregation is suppressed, even for concentrated solutions. If such “dead zone” solvent combinations can be found for complexes of type  $[1][X]$ , this may be highly advantageous for improving catalytic productivity whilst maintaining selectivity.

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## Notes and references

- 1 B. M. Trost, M. R. Machacek and A. Aponick, *Acc. Chem. Res.*, 2006, **39**, 747–760.
- 2 Z. Lu and S. Ma, *Angew. Chem., Int. Ed.*, 2007, **47**, 258–297.
- 3 B. M. Trost and R. C. Bunt, *J. Am. Chem. Soc.*, 1994, **116**, 4089–4090.





- 4 (a) S. Fuchs, V. Berl and J.-P. Lepoittevin, *Eur. J. Org. Chem.*, 2007, 7, 1145–1152; (b) M. J. Moschitto, D. N. Vaccarello and C. A. Lewis, *Angew. Chem., Int. Ed.*, 2015, **54**, 2142–2145.
- 5 B. M. Trost and F. D. Toste, *J. Am. Chem. Soc.*, 2003, **125**, 3090–3100.
- 6 B. M. Trost and G. Dong, *J. Am. Chem. Soc.*, 2006, **128**, 6054–6055.
- 7 B. M. Trost and N. Asakawa, *Synthesis*, 1999, 1491–1494.
- 8 B. M. Trost and R. C. Bunt, *J. Am. Chem. Soc.*, 1996, **118**, 235.
- 9 G. C. Lloyd-Jones and S. C. Stephen, *Chem.-Eur. J.*, 1998, **4**, 2539–2549.
- 10 C. P. Butts, E. Filali, G. C. Lloyd-Jones, P.-O. Norrby, D. A. Sale and Y. Schramm, *J. Am. Chem. Soc.*, 2009, **131**, 9945–9957.
- 11 D. J. Berrisford, C. Bolm and K. B. Sharpless, *Angew. Chem., Int. Ed.*, 1995, **34**, 1059–1070.
- 12 J. Eastoe, I. J. S. Fairlamb, J. M. Fernández-Hernández, E. Filali, J. C. Jeffery, G. C. Lloyd-Jones, A. Martorell, A. Meadowcroft, P.-O. Norrby, T. Riis-Johannessen, D. A. Sale and P. M. Tomlin, *Faraday Discuss.*, 2010, **145**, 27–47.
- 13 I. J. S. Fairlamb and G. C. Lloyd-Jones, *Chem. Commun.*, 2000, 2447–2448.
- 14 Stoichiometric reaction of  $[(R,R)\text{-}[1][\text{BAr}_4]]$  ( $\text{Ar} = 3,5\text{-(CF}_3)_2\text{C}_6\text{H}_3$ ) with tetrabutylammonium dimethylmalonate in THF at 21 °C:  $[\text{Pd}]_{\text{tot}} = 1 \text{ mM}$  affords alkylation product in 98% ee, whereas product of 6 % ee is obtained when  $[\text{Pd}]_{\text{tot}} = 27 \text{ mM}$ .<sup>10</sup>
- 15 Around 100 attempts were made to crystallise the following complexes ( $[(R,R)\text{-}]$  and *rac*  $[1][\text{X}]$ ,  $\text{X} = \text{BAr}_F$ ,  $\text{B}((3,5\text{-(CF}_3)_2\text{C}_6\text{H}_3)_4)$ ,  $[\text{Al}(\text{OC}(\text{CF}_3)_3)_4]$  and OTf) using diffusion, evaporation, cooling, seeding, co-crystallisation, and other methods, from a variety of binary and ternary solvent mixtures, including but not limited to chloroform,  $\text{CH}_2\text{Cl}_2$ /pentane, 1,2-dichloroethane/hexane, THF/heptane, 1,2-difluorobenzene/1,3-difluorobenzene, tetrachloroethane/hexafluorobenzene, MeCN/mesitylene, aqueous MeCN/MTBE, water, diethyl ether, etc. without success.
- 16 For examples of X-ray structures of Pd complexes of **2** that are inactive as catalysts for allylic alkylation, see: (a) K. R. Campos, M. Journet, S. Lee, E. J. J. Grabowski and R. D. Tillyer, *J. Org. Chem.*, 2005, **70**, 268–274; (b) C. Amatore, A. Jutand, L. Mensah and L. Ricard, *J. Organomet. Chem.*, 2007, **692**, 1457–1464.
- 17 C. P. Butts, J. Crosby, G. C. Lloyd-Jones and S. C. Stephen, *Chem. Commun.*, 2000, 1707–1708.
- 18 D. C. Smith and G. M. Gray, *J. Chem. Soc., Dalton Trans.*, 2000, 677.
- 19 J. Eastoe, *Surfactant Chemistry*, Wuhan University Press, Wuhan, China, 2005.
- 20 Perfluoro tetraphenylborate  $[\text{B}(\text{C}_6\text{F}_5)_4]^-$  (“ $\text{BAr}_F$ ”) was chosen to avoid SANS data being complicated by neutron scattering from protons or deuterons in the counter-ion, as this could then be used without perturbation in the perdeuterated complex.
- 21 C. J. Rohbogner, A. J. Wagner, G. C. Clososki and P. Knochel, *Org. Synth.*, 2009, **86**, 374–384.
- 22 The large KIE ( $k_{\text{H}}/k_{\text{D}} \approx 30$ , by comparison of approximate rates of metallation of the deuterated and non-deuterated substrates), may arise from multiple primary effects, involving the TMP-amide, and *in situ* generated  $[\text{D}]\text{-TMP}$ , or from tunnelling.
- 23 It was found that slow addition of  $[\text{D}10]\text{-8}$  to the Grignard drop-wise at  $-78^\circ\text{C}$ , reduced extensive by-product formation.
- 24 F. M. Piller, P. Appukkuttan, A. Gavryushin, M. Helm and P. Knochel, *Angew. Chem., Int. Ed.*, 2008, **47**, 6802–6806.
- 25 E. Filali, G. C. Lloyd-Jones and D. A. Sale, *Synlett*, 2009, 205–208.
- 26 K. Ishibashi and S. Matsubara, *Chem. Lett.*, 2007, **36**, 724–725.
- 27 M. Chandrasekhar, G. Sekar and V. K. Singh, *Tetrahedron Lett.*, 2000, **41**, 10079–10083.
- 28 A. K. Chakraborti, S. Rudrawar and A. Kondaskar, *Eur. J. Org. Chem.*, 2004, 3597–3600.
- 29 V. M. Mastranzo, E. Santacruz, G. Huelgas, E. Paz, M. V. Sosa-Rivadeneira, S. Bernès, E. Juaristi, L. Quintero and C. A. de Parrodi, *Tetrahedron: Asymmetry*, 2006, **17**, 1663–1670.
- 30 In contrast to the report of Singh,<sup>27</sup> hydrogenolysis with  $\text{Pd}(\text{OH})_2/\text{C}$  under acidic conditions in methanol failed to cleave the  $\alpha$ -methylbenzyl auxiliary. Instead, after reduction of the azide, the hydrogen source was changed to ammonium formate: (a) A. D. Brown, M. E. Bunnage, P. A. Glossop, K. James, C. A. L. Lane, R. A. Lewthwaite and D. A. Price, WO 2005/092840 PCT/IB2005/000619, 2005; (b) T. Bieg and W. Szeja, *Synthesis*, 1985, 76–77.
- 31 The dimer was prepared using the literature procedure [see: (a) B. M. Trost, P. E. Strege, L. Weber, T. J. Fullerton and T. J. Dietsche, *J. Am. Chem. Soc.*, 1978, **100**, 3407–3415; (b) S. Imaizumi, T. Matsuhisa and Y. Senda, *J. Organomet. Chem.*, 1985, **280**, 441–448]. However we identified degradation of the crude organometallic complex during the early work-up stages. Constant exposure of the crude material to air, avoiding vacuum operations until the complex has been purified, significantly increases the yield of **17**.
- 32 SANS experiments were conducted at the Institut Laue-Langevin (ILL) in Grenoble, France. Data was acquired in two detector movements on separate instrument: high-Q data on D16, and low-Q data on D22.
- 33 Very similar results were obtained on SANS2D at ISIS, UK using a different batch of complex, indicative of the reproducibility of the data.
- 34 Measurements were also conducted at higher and lower temperatures. Consistent with analysis by  $^{31}\text{P}\{^1\text{H}\}$  NMR spectroscopy, aggregation is reversible, and favoured by lower temperatures. No significant change in particle dimensions was detected by SANS within the relatively narrow range of temperatures explored (10–40 °C).
- 35 At higher concentrations the radius shrank slightly, possibly due to the accompanying increase in the ionic strength of the medium.<sup>12</sup>
- 36 It should be noted that most of the concentrations employed for the SANS experiments were, by necessity, well above the





- oligomerisation and aggregation threshold. Changes in particle number due to differential selectivity between homochiral and heterochiral aggregation of cyclo-oligomers may be small, and thus hard to distinguish.
- 37 The resolution of the SANS instrument was insufficient to completely rule out the possibility that the scattering profile is the composite of two curves arising from segregated (homochiral) cylinders.
- 38 (a) MM3 studies were conducted using Schrödinger MacroModel, version 9.9, Schrödinger, LLC, New York, NY, 2012, employing a modified form of the force field reported in: (b) H. Hagelin, M. Svensson, B. Åkermark and P.-O. Norrby, *Organometallics*, 1999, **18**, 4574–4583; (c) DFT studies were conducted using Jaguar, version 7.9, Software, Schrödinger, LLC, New York, NY, 2012.
- 39 See "Molecular Mechanics and Comparison of Force Fields", T. Liljefors, K. Gundertofte, P.-O. Norrby and I. Pettersson, in *Computational Medicinal Chemistry for Drug Discovery*, ed. P. Bultinck, J. P. Tollenaere, H. De Winter and W. Langenaeker, Marcel Dekker, New York, 2004, pp. 1–28.
- 40 Note that incorporation of each monomeric unit carries an entropic penalty of *ca.* 20–40 kJ/mol, not readily calculated by our current methods. To circumvent the entropy problem associated with molecularity, we compare assemblies with oligomers of the same size, and find that tetramers are favored over monomers  $[[[1][\text{BAR}_F]]_4 \text{ versus } \{4[[1][\text{BAR}_F]]\}$ , Figure 5] whereas an assembly of two tetramers is slightly favored over the octamer  $[[[1][\text{BAR}_F]]_4]_2 \text{ versus } [[1][\text{BAR}_F]]_8$ , Figure 5]. The dimensions of the MM3-minimised tetramer are similar to that of the Pd/ligand core of a cyclic tetranuclear complex  $[(\eta^3\text{-C}_3\text{H}_5\text{-Pd})_4(2)_4][\text{OTf}_4]$ , determined by single crystal X-ray diffraction.<sup>12</sup>
- 41 I. Krossing and I. Raabe, *Angew. Chem., Int. Ed.*, 2004, **43**, 2066–2090.
- 42 A variety of other anion arrangements explored by MM3 modelling did not reveal any plausible alternative modes of aggregation, that would be compatible with the SANS data.
- 43 The impact of solvent dielectric constant ( $\epsilon_r$ ) on ion-pair separation can be predicted by the Bjerrum length equation:  $\lambda_B = e^2 / (4\pi\epsilon_0\epsilon_r k_B T)$  which indicates that the ion separation distance,  $\lambda_B$ , is inversely proportional to the dielectric constant ( $\epsilon_r$ ) of the media in which the solute is dissolved. The equation presumes point charges, although the general principles may still apply to more complex systems such as  $[1][X]$ .
- 44 These systems present an apparently simple pair of doublets ( $J = 38 \text{ Hz}$ ), almost identical in appearance, but not chemical shift, to the pair of doublets ( $^{2\text{PD}}J_{\text{PP}}$ ) in the monomeric cationic chelate  $[1]^+$ .<sup>12</sup> However, it is very evident from the SANS data that the solutions are dominated by oligomers  $\{([1][\text{OTf}])_4\}_m$ , and from careful inspection of the  $^{31}\text{P}$  NMR spectrum that the monomer is indeed also present, but only in very low proportions ( $\leq 4\%$ ).
- 45 The enantiomerically pure species  $[[\text{D}_{47}]-1][\text{BAR}_F]$  is among the most synthetically complex perdeuterated species to have been prepared to date. For other examples see: (a) F. Aussenac, M. Laguerre, J.-M. Schmitter and E. J. Dufourc, *Langmuir*, 2003, **19**, 10468–10479; (b) R. G. Alken, *US pat.*, US2005069276A1, 2005; (c) M. Schulz, J. Hirschmann, A. Draksharapu, G. Bindra, S. Soman, A. Paul, R. Groarke, M. T. Pryce, M. S. Rau, W. R. Browne and J. G. Vos, *Dalton Trans.*, 2011, **40**, 10545–10552; (d) T. Abe, A. Miyazawa, H. Konno and Y. Kawanishi, *Chem. Phys. Lett.*, 2010, **491**, 199–202; (e) C. Lenges, P. White and M. Brookhart, *J. Am. Chem. Soc.*, 1999, **121**, 4385–4396; (f) M. Ogasawara, K. Takizawa and T. Hayashi, *Organometallics*, 2002, **21**, 4853–4861; (g) R. Brainard, T. Miller and G. Whitesides, *Organometallics*, 1986, **5**, 1481–1490.
- 46 All of the  $\text{BAR}_4$  anions tested, as well as  $\text{Al}(\text{OR}_F)_4$  ("Krossing anions": I. Krossing, *Chem.-Eur. J.*, 2001, **7**, 490–502), were effective in this regard.
- 47 M. J. Hollamby, R. Tabor, K. J. Mutch, K. Trickett, K., J. Eastoe, R. K. Heenan and I. Grillo, *Langmuir*, 2008, **24**, 12235–12240.

