

## RESEARCH ARTICLE

View Article Online  
View Journal | View IssueCite this: *Mater. Chem. Front.*,  
2018, 2, 597

## Solid-state self-inclusion complexation behaviour of a pillar[5]arene-based host–guest conjugate†

T. Ogoshi,<sup>ib</sup>\*<sup>ab</sup> T. Furuta,<sup>a</sup> Y. Hamada,<sup>a</sup> T. Kakuta<sup>a</sup> and T. Yamagishi<sup>a</sup>

A host–guest conjugate consisting of a pillar[5]arene and an ethylene moiety containing a triazole group at one end and a perfluorooctyl group at the other end was synthesized. The host–guest conjugate displayed unusual real-time scale solid-state self-inclusion complexation behaviour. The host–guest conjugate formed a de-threaded supramolecular structure only in solution, even at high concentrations. However, heating the solid-state host–guest conjugate led to the formation of a self-inclusion complex structure at a real-time scale; at 100 °C, it took ca. 20 h to form the self-inclusion complex at the equilibrium state at a conversion of 85%. The self-inclusion complex converted to the de-threaded form at a real-time scale when it was stored in solvents with small molecules, which worked as competitive guests.

Received 3rd December 2017,  
Accepted 24th January 2018

DOI: 10.1039/c7qm00553a

rsc.li/frontiers-materials

## Introduction

A self-inclusion complex, pseudo[1]rotaxane, which consists of a macrocyclic host–guest conjugate, is one of the simpler supramolecular structures among various supramolecular structures such as rotaxanes, catenanes, polyrotaxanes, and polycatenanes.<sup>1–12</sup> The host–guest conjugates exhibit a fast or slow exchange between the self-included and de-threaded supramolecular structure when exposed to different external stimuli. As the concentration of the host–guest conjugate increases, self-assembly normally leads to a mixture of a self-inclusion complexes and linear and cyclic supramolecular oligomers and polymers. Crown ethers,<sup>13–15</sup> cyclodextrins,<sup>16–18</sup> cucurbit[*n*]urils,<sup>19,20</sup> calix[*n*]arenes,<sup>21–23</sup> and pillar[*n*]arenes<sup>24–34</sup> have been utilized as hosts to make host–guest conjugates. The kinetics of the self-inclusion complexation process and stability of the self-inclusion complex can be controlled by tuning the relationship between the cavity size of the macrocyclic hosts and length, size, and functional groups on the guest. In most cases, the change in the supramolecular structure in solution can be monitored using <sup>1</sup>H NMR and optical measurements such as UV, fluorescence and CD measurements. In this study, we discovered an unusual solid-state self-inclusion complexation behaviour when we synthesized a new pillar[5]arene-based host–guest conjugate.

<sup>a</sup> Graduate School of Natural Science and Technology, Kanazawa University, Kakuma-machi, Kanazawa, 920-1192, Japan. E-mail: ogoshi@se.kanazawa-u.ac.jp

<sup>b</sup> WPI Nano Life Science Institute, Kanazawa University, Kakuma-machi, Kanazawa, 920-1192, Japan

† Electronic supplementary information (ESI) available: <sup>1</sup>H and <sup>13</sup>C NMR spectra of **1**, concentration-dependent <sup>1</sup>H NMR spectra of **1** in the de-threaded form and self-inclusion complex, de-threading behaviour of **1s** in various deuterated solvents and calculations. See DOI: 10.1039/c7qm00553a

Pillar[*n*]arenes,<sup>35–37</sup> which were first reported by our group,<sup>38</sup> form stable host–guest complexes with linear alkanes containing electron-withdrawing groups at both ends.<sup>39–43</sup> Thus, we used an ethylene moiety with two different electron-withdrawing groups, a triazole and a perfluorooctyl group, at each end as the guest to make the host–guest conjugate **1** with 9 cyclohexyl groups (Cy, Fig. 1). The host–guest conjugate **1** only formed the de-threaded structure **1d** in solution. Interestingly, heating **1d** in the solid state led to formation of a self-inclusion complex structure (**1s**). The change in the supramolecular structure in the solid state occurred at a real-time scale: it took ca. 20 h to reach the equilibrium state by heating at 100 °C. By controlling the temperature, 85% conversion of **1d** to **1s** was achieved at the equilibrium state. Reports of changes to a supramolecular structure in the solid-state are very rare. Horie and co-workers reported pseudorotaxanes that act as a thermal- and photo-driven molecular switch in the single-crystal state.<sup>44,45</sup> Monitoring of macrocyclic ring motion of [2]rotaxane using <sup>2</sup>H solid state

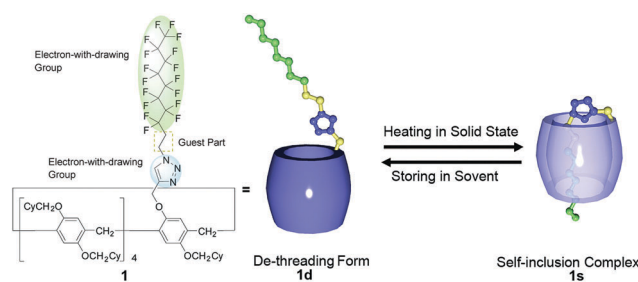


Fig. 1 Chemical structure of pillar[5]arene-based host–guest conjugate and supramolecular structural changes between de-threading form **1d** and self-inclusion complex **1s**. Cy: cyclohexyl group.

NMR in metal–organic frameworks was reported by Loeb and co-workers.<sup>46</sup> Takata and co-workers reported thermal-induced solid-state ring-shuttling in side-chain-type polyrotaxanes.<sup>47</sup> However, to the best of our knowledge, there are no examples of solid-state supramolecular structural change of host–guest conjugates from a de-threaded to a self-inclusion complex structure by heating. Furthermore, **1s** was very stable; even at high concentrations, **1s** did not change to other supramolecular structures such as linear and cyclic supramolecular oligomers and polymers. This is an unusual phenomenon because supramolecular structures change from self-inclusion complexes to linear and cyclic supramolecular oligomers and polymers with increasing concentration.<sup>22–26</sup> The selective synthesis of self-inclusion complexes has been reported using cyclodextrins and pillar[5]arenes as wheels,<sup>17,18,30–34</sup> but the complexation to form the self-inclusion complexes was mainly in solution. To the best of our knowledge, this is the first example of self-inclusion complexation by external stimuli in the solid state, while there are reports of self-inclusion complexes in solid state.<sup>30–34</sup> Furthermore, **1s** was converted back to **1d** by storing **1s** in solution. The structural change also occurred at a real-time scale; it took *ca.* 20 h to reach the equilibrium state at 25 °C. In this study, we investigated the mechanism of the unusual self-inclusion complexation in the solid state and de-threading behaviour in solution.

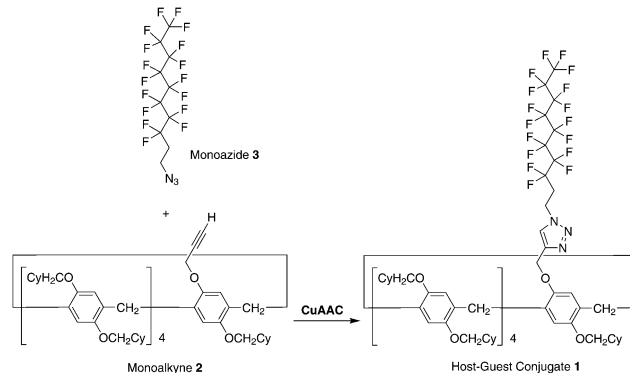
## Experimental section

### Materials and methods

Pillar[5]arene containing a monoalkyne group (**2**) and a monoazide containing ethylene and perfluorooctyl moieties (**3**) were synthesized according to previous reports.<sup>24,48</sup> Solution <sup>1</sup>H NMR spectra were recorded at 500 MHz with a JEOL-ECA500 spectrometer. Powder X-ray diffraction (PXRD) measurements were performed on a Rigaku Smart Lab high-resolution diffractometer. Differential scanning calorimetry (DSC) was performed using a PerkinElmer Jade DSC calorimeter equipped with an Intra-cooler II under a flow of dry nitrogen and with a heating rate of 10 °C min<sup>-1</sup>.

### Synthetic procedures

The synthesis of the pillar[5]arene-based host–guest conjugate **1** is shown in Scheme 1. Tris[(1-benzyl-1*H*-1,2,3-triazol-4-yl) methyl]-amine (173 mg, 0.321 mmol) and Cu(CH<sub>3</sub>CN)<sub>4</sub>PF<sub>6</sub> (121 mg, 0.321 mmol) were added to a solution of the monoalkyne **2**<sup>24</sup> (485 mg, 0.321 mmol) and monoazide **3**<sup>48</sup> (200 mg, 4.10 mmol) in dichloromethane (20 mL). The mixture was stirred at 25 °C for 48 h. The resulting solution was concentrated under vacuum. Column chromatography (silica gel; hexane/dichloromethane = 3 : 1) afforded a white solid in the de-threaded form (**1d**, 155 mg, 0.0774 mmol, yield: 24%). <sup>1</sup>H NMR (CDCl<sub>3</sub>, 500 MHz, ppm): δ = 7.63 (s, 1H, triazole unit), 6.74–7.00 (m, 10H, phenyl), 5.25, 5.23, 5.08, 5.06 (dd, 2H, methylene), 4.65, 4.67, 4.67 (t, 2H, methylene), 3.36–3.88 (m, 28H, methylene), 2.81–2.91 (m, 2H, methylene), 1.05–2.05 (m, 99H, cyclohexyl group).



Scheme 1 Synthesis of the host–guest conjugate **1** by copper(I)-catalysed alkyne–azide cycloaddition (CuAAC) “click” reaction between the monoalkyne **2** and monoazide **3**.

<sup>13</sup>C NMR (CDCl<sub>3</sub>, 125 MHz, ppm): δ = 150.6, 149.9, 148.7, 145.8, 128.8, 128.6, 128.5, 128.4, 128.3, 128.2, 128.1, 127.9, 127.8, 122.6, 115.4, 115.1, 115.0, 114.9, 114.7, 74.2, 74.1, 73.9, 62.8, 42.3, 38.6, 38.5, 38.4, 32.0, 31.9, 30.5, 30.4, 30.2, 30.0, 29.5, 29.2, 29.0, 26.7, 26.1, 26.0. HRESIMS: *m/z* calcd for C<sub>111</sub>H<sub>200</sub>F<sub>17</sub>N<sub>3</sub>Na<sub>1</sub>O<sub>10</sub> [M + Na]<sup>+</sup>: 2025.0478, found 2025.0474.

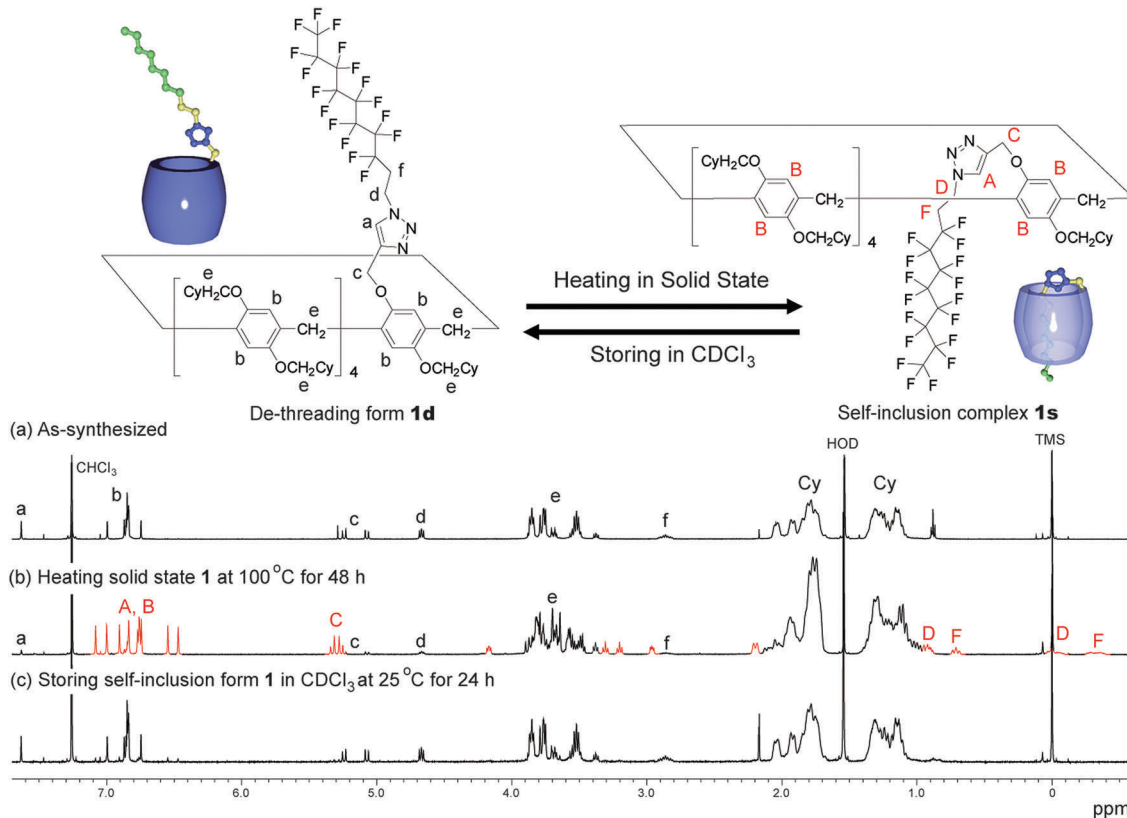
### Monitoring the self-complexation and de-threading behaviour

An as-synthesised sample of **1d** in the solid state was heated at various temperatures. After heating the sample in the solid state, the sample was cooled to 25 °C, dissolved in CDCl<sub>3</sub>, then the <sup>1</sup>H NMR spectrum of the sample was obtained after 3 min. De-threading occurred in CDCl<sub>3</sub>, but it took several hours (*ca.* 20 h at 25 °C) to reach equilibrium; therefore, <sup>1</sup>H NMR spectra obtained quickly after dissolving the sample in CDCl<sub>3</sub> can be used to determine the supramolecular structure formed in the solid state. The de-threading process was monitored by <sup>1</sup>H NMR measurements by storing the mixture (**1s/1d** = 85/15), which was prepared by heating the solid state **1d** at 100 °C for 48 h, at various temperatures in CDCl<sub>3</sub>. The ratio between **1s** and **1d** was determined by calculating the integration ratio between the <sup>1</sup>H NMR signal c from **1s** and the signal c from **1d** (Fig. 2).

## Results and discussion

### Unusual solid-state self-inclusion complexation behaviour of pillar[5]arene-based host–guest conjugate

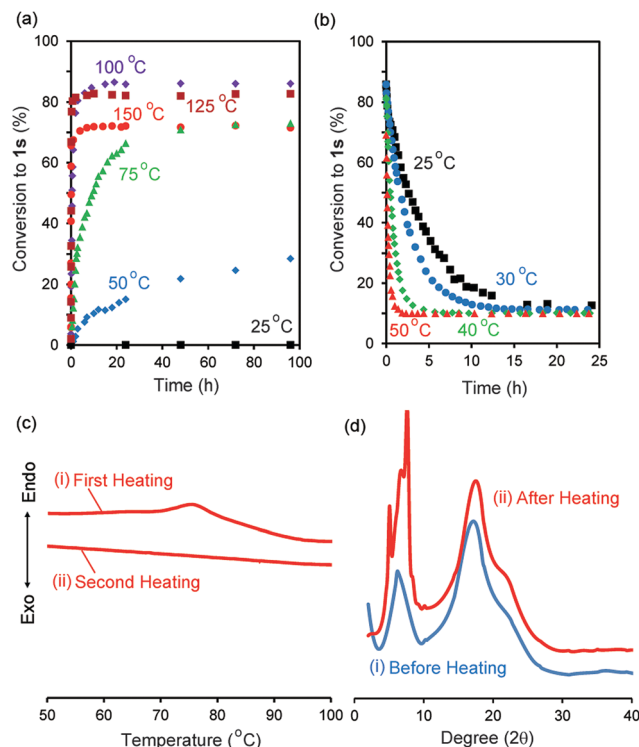
We synthesized the host–guest conjugate **1** by the copper(I)-catalysed alkyne–azide cycloaddition (CuAAC) click reaction between a pillar[5]arene containing a monoalkyne group (**2**) and a monoazide containing ethylene and perfluorooctyl moieties (**3**) (Scheme 1). To investigate the assembled structure of **1**, we obtained the <sup>1</sup>H NMR spectra of **1** at various concentrations (Fig. S3, ESI<sup>†</sup>). However, the proton peaks of **1** did not change even at a high concentration, which indicated that supramolecular polymerization and self-inclusion complexation did not take place. The as-synthesised **1** formed the de-threaded structure **1d**. This was owing to the bulkiness of the perfluorooctyl



**Fig. 2**  $^1\text{H}$  NMR spectra of the host–guest conjugate **1**. As-synthesized (a), after heating of **1** in the solid state at  $100\text{ }^\circ\text{C}$  for 48 h, dissolving the solid sample in  $\text{CDCl}_3$ , and obtaining the spectrum after 3 min (b), and 24 h (c).

moieties in **1**. The cavity size of pillar[5]arenes is *ca.*  $4.7\text{ \AA}$ , which accommodates a linear hydrocarbon.<sup>49,50</sup> The size of the perfluorooctyl moiety is slightly larger than the cavity size of a pillar[5]arene (see ESI† in details). The other possible reason for the de-threaded structure of **1d** is the electron repulsion force between the electron-rich cavity of the pillar[5]arene<sup>43</sup> and the electronegativity of the perfluorooctyl moiety. The electron repulsion force would prevent threading of the perfluorooctyl moiety into the pillar[5]arene cavity to form a self-inclusion complex and cyclic and linear supramolecular oligomers and polymers. However, surprisingly, new proton signals (red) appeared along with the original proton signals of **1d** after heating of **1d** at  $100\text{ }^\circ\text{C}$  for 48 h in the solid state, which indicated that complexation occurred. No solvent molecules were included in the cavity of the pillar[5]arene host–guest conjugate **1**, which confirmed by  $^1\text{H}$  NMR measurement of **1d** (Fig. 1S). To investigate the supramolecular structure after the heating, we investigated the concentration-dependent  $^1\text{H}$  NMR spectra after heating **1d** in the solid state. However, even at high concentrations, the new set of proton signals did not change (Fig. S6, ESI†), which indicated that the supramolecular structure of **1** after the heating was the self-inclusion complex **1s**, which was very stable; **1s** did not form cyclic and linear supramolecular oligomers and polymers even at high concentrations. We investigated the effect of the temperature on the conversion from **1d** to **1s** in the solid state (Fig. 3a). At  $25\text{ }^\circ\text{C}$ ,

**1s** was not formed even after a significant period (150 h). **1s** was slowly formed, but the reaction did not reach a thermodynamic equilibrium upon heating of **1d** at  $50\text{ }^\circ\text{C}$  for 95 h. At  $100\text{ }^\circ\text{C}$ , it took 20 h to form **1s** in a thermodynamic equilibrium; the conversion to **1s** at the equilibrium state was 85%. The conversion to **1s** decreased with increasing reaction temperature above  $100\text{ }^\circ\text{C}$ . The time required to reach a thermodynamic equilibrium became shorter with increasing reaction temperature. These results were used to determine the thermodynamic ( $\Delta H^\ddagger$  and  $\Delta S^\ddagger$ ) parameters based on Eyring plots (Fig. S4, ESI†).  $\Delta H^\ddagger$  and  $\Delta S^\ddagger$  were  $70.8\text{ kJ mol}^{-1}$  and  $-124.2\text{ J mol}^{-1}\text{ K}^{-1}$ , respectively. The calculated  $\Delta G^\ddagger$  and half-life time  $t_{1/2}$  at  $50\text{ }^\circ\text{C}$  were  $110.9\text{ kJ mol}^{-1}$  and 23.4 h, respectively. We could monitor the self-inclusion complexation behaviour of **1** in the solid state at a real time-scale. To understand the reason why **1** formed the self-inclusion complex **1s** by heating, we calculated the electrostatic potential profile of the guest (ESI† in details). The ethylene moieties in the guest are positive because triazole and perfluorooctyl groups at both ends of the guest part are negative. In the case of pillar[5]arenes, the inner surfaces of the cavities are negative, and the outer surfaces of the cavities and rims are almost neutral.<sup>43</sup> Thus, pillar[5]arenes capture the positive moieties of the ethylene moieties in the guest part. This was consistent with the experimental data: the proton signals from the ethylene moieties in the guest part (signals d and f) showed a large up-field shift after the formation of the self-inclusion complex.



**Fig. 3** Percentage conversions to **1s** versus time (a) in solid state **1d** at 25 °C (black squares), 50 °C (blue diamonds), 75 °C (green triangles), 100 °C (purple diamonds), 125 °C (brown squares), 150 °C (red circles) and (b) in  $\text{CDCl}_3$  at 25 °C (black squares), 30 °C (blue circles), 40 °C (green diamonds), 50 °C (red triangles). (c) DSC heating curves in (i) first and (ii) second heating processes. (d) PXRD patterns (i) before and (ii) after heating at 100 °C for 48 h.

However, the fluoro atoms have high negative charge (Fig. S12, ESI<sup>†</sup>); therefore, there would be electron repulsion between the perfluorooctyl group and the negatively charged pillar[5]arene cavity. In addition, the size of the perfluorooctyl group was slightly larger than the cavity size of pillar[5]arenes, because the cavity size of the pillar[5]arenes was just able to accommodate the linear hydrocarbons.<sup>49</sup> Thus, heating was necessary to form the self-inclusion complex to induce threading of the perfluorooctyl groups into the pillar[5]arene cavity.

#### Assembled structural change by self-inclusion complex in solid state by heating

To investigate the solid-state structural change of the supramolecular assembly from **1d** to **1s**, DSC measurements were carried out (Fig. 3c). In the first heating of **1d** (Fig. 3c(i)), an endothermic peak was observed from 70 °C to 90 °C. However, an endothermic peak was not observed in the second heating (Fig. 3c(ii)). No endothermic peak was observed in the first and second heating of a mixture of **1s** and **1d** (**1s/1d** = 85/15, Fig. S5, ESI<sup>†</sup>). Thus, the endothermic peak observed in the first heating of **1d** resulted from formation of **1s**. Fig. 3d shows the powder X-ray diffraction patterns before and after heating of **1d** in the solid state at 100 °C for 48 h. New diffraction peaks appeared in the small angle region around 7–10 degrees upon heating,

which indicated that the structure of the assemblies change from **1d** to **1s** even in the solid state.

#### De-threading in solution

Assembly **1s** was stable in the solid state, but de-threading slowly occurred when **1s** was stored in  $\text{CDCl}_3$ . Fig. 2b and c show the  $^1\text{H}$  NMR spectra after dissolving **1s** (**1s/1d** = 85/15) in  $\text{CDCl}_3$  after 3 min and 24 h. The new set of the proton signals from **1s** was observed after 3 min, but almost disappeared after 24 h, indicating that **1s** changed to **1d** by dissolving of **1s** over several hours in  $\text{CDCl}_3$ . **1s** was slowly converted to **1d** at 25 °C (Fig. 3b). The time required to reach thermodynamic equilibrium became shorter with increasing reaction temperature. The de-threading was accelerated by heating in solution. These results were used to determine the thermodynamic ( $\Delta H^\ddagger$  and  $\Delta S^\ddagger$ ) parameters based on Eyring plots (Fig. S8, ESI<sup>†</sup>).  $\Delta H^\ddagger$  and  $\Delta S^\ddagger$  were 72.3  $\text{kJ mol}^{-1}$  and  $-82.5 \text{ J mol}^{-1} \text{ K}^{-1}$ , respectively. The calculated  $\Delta G^\ddagger$  and half-life time  $t_{1/2}$  at 25 °C were 90.7  $\text{kJ mol}^{-1}$  and 2.89 h, respectively. We also monitored the de-threading behaviour of the complex in  $\text{CDCl}_3$  in a real-time scale.

The effect of different solvents on the supramolecular structure change was also investigated. When  $\text{CD}_2\text{Cl}_2$  was used as a solvent (Fig. S9, ESI<sup>†</sup>), the set of signals attributed to **1s** also disappeared after 24 h, as observed in  $\text{CDCl}_3$  (Fig. 2c), which indicated that de-threading occurred in  $\text{CD}_2\text{Cl}_2$ . In contrast, the  $^1\text{H}$  NMR signals of **1s** did not change even after 24 h when **1s** was dissolved in deuterated 1,1,2,2-tetrachloroethane and cyclohexane (Fig. S10 and S11, ESI<sup>†</sup>). The affinity of the solvent molecules for the pillar[5]arene cavity depends not only on the polarity of the solvent but also on the molecular size of the solvent molecules. The cavity size of pillar[5]arenes is 4.7 Å; thus, pillar[5]arenes can form complexes with solvents with a small molecule size but cannot form complexes with solvents with a large molecule size.<sup>51</sup> The inclusion of the small  $\text{CH}_2\text{Cl}_2$  molecules into the cavity of pillar[5]arenes has been completely characterized by single X-ray crystal analysis.<sup>52</sup> The size of the  $\text{CD}_2\text{Cl}_2$  and  $\text{CDCl}_3$  molecules is small enough to form complexes with pillar[5]arenes; thus, de-threading occurred in these solvents. In contrast, the 1,1,2,2-tetrachloroethane and cyclohexane molecules were too bulky to fit into the pillar[5]arene cavity. Therefore, no de-threading occurred in these solvents.

## Conclusions

Unusual real-time scale solid-state self-inclusion complexation behaviour was observed when a pillar[5]arene-based host-guest conjugate was heated in the solid state. The inclusion of a superhydrophobic perfluorooctyl group in the guest in the solid state would be used in surface chemistry to control contact angles. The self-inclusion complex converted to the de-threaded form at a real-time scale when it was stored in solution in solvents with small molecules, which work as competitive guests. The real-time scale structural changes would be useful for controlling supramolecular assembly and systems at a real-time scale. This is now under investigation.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This work was partially supported by Grant-in-Aid for Scientific Research on Innovative Areas (2601):  $\pi$ -System Figuration (15H00990 and 17H05148), Kiban B (16H04130) from MEXT Japan and Kanazawa University Chozen Project.

## Notes and references

- P. L. Anelli, N. Spencer and J. F. Stoddart, *J. Am. Chem. Soc.*, 1991, **113**, 5131–5133.
- W. R. Dichtel, O. Š. Miljanić, W. Zhang, J. M. Spruell, K. Patel, I. Aprahamian, J. R. Heath and J. F. Stoddart, *Acc. Chem. Res.*, 2008, **41**, 1750–1761.
- A. Harada, A. Hashidzume, H. Yamaguchi and Y. Takashima, *Chem. Rev.*, 2009, **109**, 5974–6023.
- F. M. Raymo and J. F. Stoddart, *Chem. Rev.*, 1999, **99**, 1643–1664.
- S. A. Nepogodiev and J. F. Stoddart, *Chem. Rev.*, 1998, **98**, 1959–1976.
- Z. Niu and H. W. Gibson, *Chem. Rev.*, 2009, **109**, 6024–6046.
- G. Wenz, B.-H. Han and A. Müller, *Chem. Rev.*, 2006, **106**, 782–817.
- M. Xue, Y. Yang, X. Chi, X. Yan and F. Huang, *Chem. Rev.*, 2015, **115**, 7398–7501.
- S. Dong, B. Zheng, F. Wang and F. Huang, *Acc. Chem. Res.*, 2014, **47**, 1982–1994.
- M. Fujita, *Acc. Chem. Res.*, 1999, **32**, 53–61.
- M. J. Langton and P. D. Beer, *Acc. Chem. Res.*, 2014, **47**, 1935–1949.
- J.-P. Sauvage, *Acc. Chem. Res.*, 1998, **31**, 611–619.
- H. V. Schroder, J. M. Wollschlager and C. A. Schalley, *Chem. Commun.*, 2017, **53**, 9218–9221.
- B. Zheng, F. Wang, S. Dong and F. Huang, *Chem. Soc. Rev.*, 2012, **41**, 1621–1636.
- Y. Kohsaka, Y. Koyama and T. Takata, *Angew. Chem., Int. Ed.*, 2011, **50**, 10417–10420.
- Y. Inoue, P. Kuad, Y. Okumura, Y. Takashima, H. Yamaguchi and A. Harada, *J. Am. Chem. Soc.*, 2007, **129**, 6396–6397.
- A. Harada, A. Miyawaki, P. Kuad, Y. Takashima and H. Yamaguchi, *J. Am. Chem. Soc.*, 2008, **130**, 17062–17069.
- Y. Inoue, M. Miyauchi, H. Nakajima, Y. Takashima, H. Yamaguchi and A. Harada, *J. Am. Chem. Soc.*, 2006, **128**, 8994–8995.
- N. Zhao, G. O. Lloyd and O. A. Scherman, *Chem. Commun.*, 2012, **48**, 3070–3072.
- B. Vinciguerra, L. Cao, J. R. Cannon, P. Y. Zavalij, C. Fenselau and L. Isaacs, *J. Am. Chem. Soc.*, 2012, **134**, 13133–13140.
- C. Gargiulli, G. Gattuso, A. Notti, S. Pappalardo and M. F. Parisi, *Tetrahedron Lett.*, 2011, **52**, 6460–6464.
- C. Capici, Y. Cohen, A. D'Urso, G. Gattuso, A. Notti, A. Pappalardo, S. Pappalardo, M. F. Parisi, R. Purrello, S. Slovak and V. Villari, *Angew. Chem., Int. Ed.*, 2011, **50**, 11956–11961.
- R. M. Yebeutchou, F. Tancini, N. Demitri, S. Geremia, R. Mendichi and E. Dalcanale, *Angew. Chem., Int. Ed.*, 2008, **47**, 4504–4508.
- T. Ogoshi, T. Furuta and T. Yamagishi, *Chem. Commun.*, 2016, **52**, 10775–10778.
- N. L. Strutt, H. C. Zhang, M. A. Giesener, J. Y. Lei and J. F. Stoddart, *Chem. Commun.*, 2012, **48**, 1647–1649.
- Z. Zhang, Y. Luo, J. Chen, S. Dong, Y. Yu, Z. Ma and F. Huang, *Angew. Chem., Int. Ed.*, 2011, **50**, 1397–1401.
- Y. Wang, J. F. Xu, Y. Z. Chen, L. Y. Niu, L. Z. Wu, C. H. Tung and Q. Z. Yang, *Chem. Commun.*, 2014, **50**, 7001–7003.
- L. Liu, L. Wang, C. Liu, Z. Fu, H. Meier and D. Cao, *J. Org. Chem.*, 2012, **77**, 9413–9417.
- Z. Zhang, G. Yu, C. Han, J. Liu, X. Ding, Y. Yu and F. Huang, *Org. Lett.*, 2011, **13**, 4818–4821.
- X. Wu, M. Ni, W. Xia, X.-Y. Hu and L. Wang, *Org. Chem. Front.*, 2015, **2**, 1013–1017.
- Y. F. Guan, P. Y. Liu, C. Deng, M. F. Ni, S. H. Xiong, C. Lin, X. Y. Hu, J. Ma and L. Y. Wang, *Org. Biomol. Chem.*, 2014, **12**, 1079–1089.
- Y. Chen, D. R. Cao, L. Y. Wang, M. Q. He, L. X. Zhou, D. Schollmeyer and H. Meier, *Chem. – Eur. J.*, 2013, **19**, 7064–7070.
- L. Chen, Z. Li, Z. Chen and J. L. Hou, *Org. Biomol. Chem.*, 2013, **11**, 248–251.
- Y. Han, G.-F. Huo, J. Sun, J. Xie, C.-G. Yan, Y. Zhao, X. Wu, C. Lin and L. Wang, *Sci. Rep.*, 2016, **6**, 28748.
- T. Ogoshi, T. Yamagishi and Y. Nakamoto, *Chem. Rev.*, 2016, **116**, 7937–8002.
- M. Xue, Y. Yang, X. D. Chi, Z. B. Zhang and F. H. Huang, *Acc. Chem. Res.*, 2012, **45**, 1294–1308.
- N. L. Strutt, H. C. Zhang, S. T. Schneebeli and J. F. Stoddart, *Acc. Chem. Res.*, 2014, **47**, 2631–2642.
- T. Ogoshi, S. Kanai, S. Fujinami, T. Yamagishi and Y. Nakamoto, *J. Am. Chem. Soc.*, 2008, **130**, 5022–5023.
- C. Li, *Chem. Commun.*, 2014, **50**, 12420–12433.
- X. Shu, S. Chen, J. Li, Z. Chen, L. Weng, X. Jia and C. Li, *Chem. Commun.*, 2012, **48**, 2967–2969.
- K. Han, Y. Zhang, J. Li, Y. Yu, X. Jia and C. Li, *Eur. J. Org. Chem.*, 2013, 2057–2060.
- X. Y. Shu, J. Z. Fan, J. Li, X. Y. Wang, W. Chen, X. S. Jia and C. J. Li, *Org. Biomol. Chem.*, 2012, **10**, 3393–3397.
- T. Ogoshi and T. Yamagishi, *Chem. Commun.*, 2014, **50**, 4776–4787.
- K.-J. Chen, Y.-C. Tsai, Y. Suzaki, K. Osakada, A. Miura and M. Horie, *Nat. Commun.*, 2016, **7**, 13321.
- M. Horie, Y. Suzaki, D. Hashizume, T. Abe, T. Wu, T. Sassa, T. Hosokai and K. Osakada, *J. Am. Chem. Soc.*, 2012, **134**, 17932–17944.
- K. Zhu, V. N. Vukotic, C. A. O'Keefe, R. W. Schurko and S. J. Loeb, *J. Am. Chem. Soc.*, 2014, **136**, 7403–7409.

- 47 N. Zhu, K. Nakazono and T. Takata, *Chem. Commun.*, 2016, **52**, 3647–3649.
- 48 S. Périno, C. Contino-Pépin, S. Jasseron, M. Rapp, J.-C. Maurizis and B. Pucci, *Bioorg. Med. Chem. Lett.*, 2006, **16**, 1111–1114.
- 49 T. Ogoshi, R. Sueto, K. Yoshikoshi, Y. Sakata, S. Akine and T. Yamagishi, *Angew. Chem., Int. Ed.*, 2015, **54**, 9849–9852.
- 50 Z. B. Zhang, B. Y. Xia, C. Y. Han, Y. H. Yu and F. H. Huang, *Org. Lett.*, 2010, **12**, 3285–3287.
- 51 T. Ogoshi, T. Akutsu, D. Yamafuji, T. Aoki and T. Yamagishi, *Angew. Chem., Int. Ed.*, 2013, **52**, 8111–8115.
- 52 T. Boinski and A. Szumna, *Tetrahedron*, 2012, **68**, 9419–9422.