



Cite this: *RSC Adv.*, 2023, 13, 14539

Received 13th April 2023

Accepted 9th May 2023

DOI: 10.1039/d3ra02447d

rsc.li/rsc-advances

Water soluble macrocyclic host for recognition of *N*-methylquinolinium salts in water†

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In this paper, we reported the synthesis of water soluble macrocyclic arenes **1** containing anionic carboxylate groups. It was found that host **1** could form a 1 : 1 complex with *N*-methylquinolinium salts in water. Moreover, the complexation and decomplexation of the complexes between host and the guests could be achieved by changing the pH of the solution, and the process could also be observed by naked eyes.

Macrocyclic arenes composed of hydroxyl-substituted aromatic rings bridged by methylene or methenyl groups have attracted considerable attention for their wide application in molecular recognition and assembly, biomedicine, materials science, and adsorption separation.^{1–9} Good examples are calixarenes,^{10–12} pillar[n]arenes,^{13–16} prism[5]arene,¹⁷ pagoda[n]arene,^{18,19} geminiarenes,^{20,21} tiara[5]arene,^{22,23} biphen[n]arenes,^{24–26} helic[6]arene^{27–29} and so on.^{30,31} Although a large number of macrocyclic arenes have been reported in the last decade. However, reports on the synthesis and applications of water soluble macrocyclic arenes are very limited.^{32–34} Water soluble macrocyclic arenes have the ability to recognize and assemble in water, which is of practical significance for understanding and mimicking biological processes.^{35–38} By introducing the water soluble groups into suitable positions on the macrocyclic arenes, water-soluble macrocyclics can be easily obtained. Water soluble carboxylatopillar[n]arenes^{39–42} and psulfonatocalix[n]arenes^{43–45} have gained maximum popularity for their easy preparation, π -electron rich hydrophobic cavity, ideal candidates for host–guest binding with cationic guests and application in fabricating responsive supramolecular systems and as nanovalves for trapping and releasing cargo molecules. Moreover, water-soluble carboxylatobiphen[n]arenes and cationic biphen[4,5]arenes^{46,47} were also been reported. Despite these seminal reports, exploring new water soluble macrocyclic arenes is still very urgent and of great value.

Recently, our group reported the easy synthesis of phenanthrene[2]arene and found that it has a larger cavity,⁵ which can encapsulate benzene molecules. Inspired by these results, we deduce that if the cavity can be further expanded and then water-soluble groups are introduced, water soluble hosts with

the abilities to form different kinds of complexes with guests in water could be obtained. 1,1':2',1''-Terphenyl can be regarded as the product of broken triphenylene at position 12a,12b. Thus, using 1,1':2',1''-terphenyl as the building block for the construction of macrocyclic arenes, the cavity should be larger than that of phenanthrene[2]arene. In addition, the introduction of anionic carboxylate groups into the synthesized macrocyclic arenes will certainly improve its water solubility. Herein, we report the synthesis of 1,1':2',1''-terphenyl-based water soluble macrocyclic arenes **1** (Fig. 1). ¹H NMR experiments demonstrated that host **1** could form 1 : 1 complexes with *N*-methylquinolinium salts in water. Furthermore, the complexation and decomplexation of the complexes between host **1** and the guests could be chemically controlled by the addition of acid/base, and the process could also be observed by naked eye.

The synthetic route for host **1** was outlined in Scheme 1 and contains four steps. Firstly, compound **4** was obtained by the Suzuki coupling reaction of the commercially available 4,4'-dibromo-1,1':2',1''-terphenyl with 2,5-dimethoxy phenyl borate in the presence of Pd(PPh₃)₄ as the catalyst in 74% yield. Then, macrocyclic arenes **3** was synthesized in moderate yield by treatment of **4** with paraformaldehyde and boron trifluoride diethyl etherate in dichloromethane at room temperature. *per*-Hydroxylated macrocyclic arenes can be quantitatively obtained by the cleavage of the methyl groups in macrocyclic arenes **3** through the reaction with excess BBr₃ in CH₂Cl₂. **2** was prepared in 43% yield by nucleophilic substitution reaction of *per*-

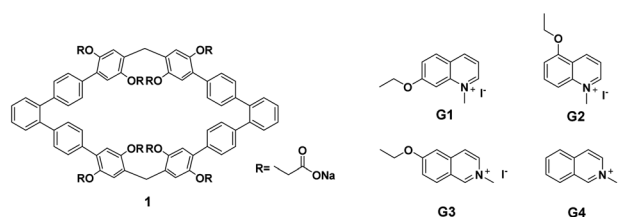
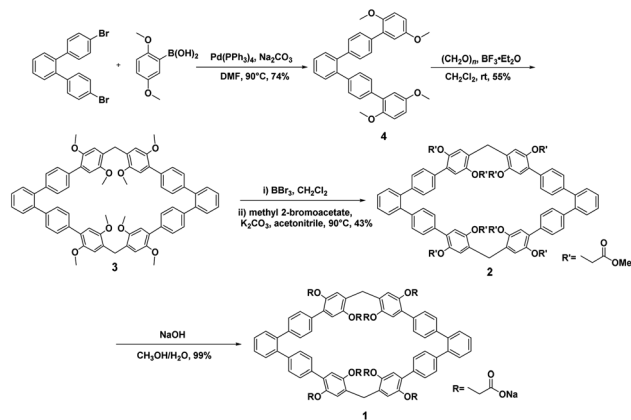


Fig. 1 Structures of host **1** and guests **G1**–**G4**.

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† Electronic supplementary information (ESI) available: Synthesis of **1**–**4**; the ¹H and ¹³C NMR spectra of **1**–**4**; CCDC: 2255801 for **3**. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d3ra02447d>





Scheme 1 Synthesis of host 1.

hydroxylated macrocyclic arenes and methyl bromoacetate in the presence of excess of K_2CO_3 . Finally, water soluble host **1** was obtained nearly quantitatively by hydrolysis of **2** in the presence of NaOH in CH_3OH/H_2O . The structure of host **1** was first confirmed by 1H NMR, ^{13}C NMR as well as HRMS spectra.

We tried to obtain the single crystals of host **1** that suitable for X-ray analysis, but failed. Luckily, single crystals of **3** that suitable for X-ray analysis was obtained by slow evaporation of the solution in $CHCl_3$, providing unambiguous evidence for the construction of new 1,1':2',1''-terphenyl-based macrocyclic arenes. As shown in Fig. 2, the formed macrocyclic arenes **3** has a large cavity with the size of $7.630 \text{ \AA} \times 7.249 \text{ \AA}$. The distance between the two dimethoxybenzene units that attached to the 1,1':2',1''-terphenyl is measured to be $10.532 \text{ \AA}$. Moreover, from the *c*-axis direction of **3** packing structure, a channel-like architecture is formed. Since macrocyclic arenes **3** has larger cavity, therefore, macrocyclic arenes **1** should also have a large cavity that can be used to complex guest molecules.

Consequently, the complex ability of host **1** toward *N*-methylquinolinium salt **G1** were first investigated by 1H NMR experiments. As shown in Fig. 3, when **1** and **G1** were mixed at a 1 : 1 molar ratio in D_2O , a new set of proton signals that different from **1** and **G1** were observed on 1H NMR spectrum, indicating that a new complex **1@G1** was formed. The protons H_1 and H_2 corresponding to **1** shift upfield by 0.05 and 0.16 ppm respectively, which could be attributed to the strong shielding

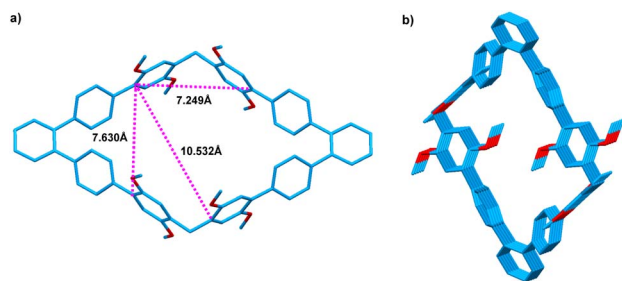


Fig. 2 Crystal structures of **3** (a) cavity size; (b) packing of **3** view along the *c* axis; Solvents and hydrogen atoms not involved in the non-covalent interactions are omitted for clarity.

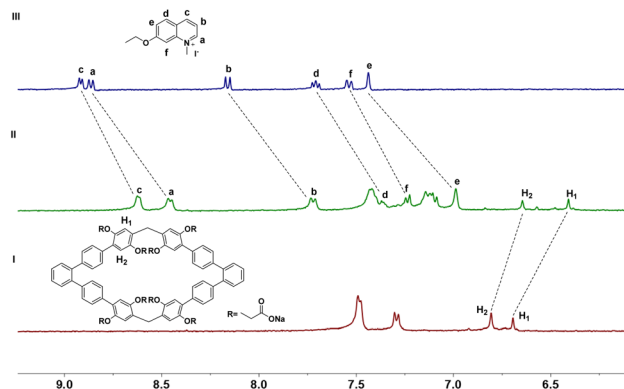


Fig. 3 Partial 1H NMR spectra (400 MHz, D_2O , 298 K) of (I) free **1**, (II) **1** and 1.0 equiv. of **G1**, and (III) free **G1**. $[1]_0 = 1.0 \text{ mM}$.

effect of **G1**. Meanwhile, the a–f proton signals corresponding to **G1** also showed a significantly upfield shift, which might be due to the strong shielding effect of the aromatic rings in host **1**. These results indicating that a new complex **1@G1** could be formed. In addition, the spectrum of complex **1@G1** showed only one set of resonances, which indicated that the complexation and decomplexation between host **1** and **G1** were a fast exchange on the NMR time scale at room temperature. Similar to the **G1**, **G2–G4** can also form 1 : 1 complex with host **1** in the similar complexation mode to that of complex **1@G1**. To further get insights into the complexation process between **1** and **G1**, 1H NMR spectroscopic titrations experiments were then carried out. By monitoring the change of proton H_1 corresponding to **1** upon the addition of **G1**, a 1 : 1 complex between **1** and **G1** was formed by the mole ratio plot. The binding constant K_a of the complex **1@G1** was determined to be $375.5 \pm 23.1 \text{ M}^{-1}$ using the BindFit software. The binding constant K_a of the complex **1@G2**, **1@G3** and **1@G4** were calculated to be $421.7 \pm 19.5 \text{ M}^{-1}$, $385.4 \pm 28.4 \text{ M}^{-1}$ and $193.5 \pm 19.7 \text{ M}^{-1}$, respectively.

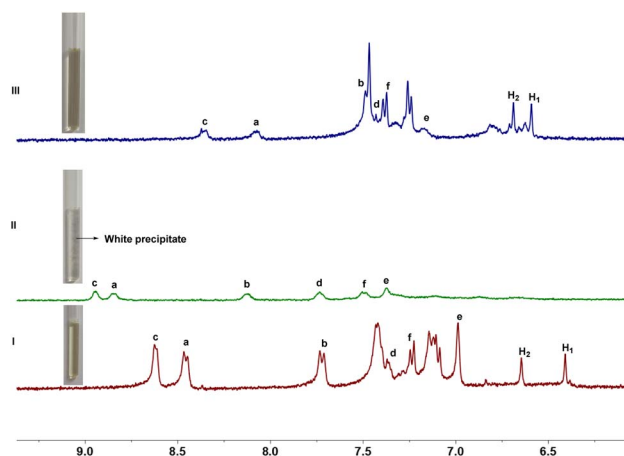


Fig. 4 Partial 1H NMR spectra (400 MHz, D_2O , 295 K) and photographs of (I) **1** and 1.0 equiv. of **G1**, (II) after addition of 2.0 μL of aqueous DCl solution (20 wt%) to I, and (III) after addition of 1.0 μL of aqueous NaOD solution (40 wt%) to II. $[1]_0 = 1.0 \text{ mM}$.



Since the conversion of anionic carboxylate groups and neutral carboxylic groups can be controlled by the addition of acid and base. Therefore, we wonder whether the complexation and decomplexation of complex **1@G1** can be regulated by acid and base. In order to verify our hypothesis, ¹H NMR experiments were carried out. After the addition of aqueous DCl to the solution of complex **1@G1**, the solution changed from neutral to acidic, the protons of H₁ and H₂ corresponding to **1** disappeared, while the a-f proton signals corresponding to **G1** shifted downfield to their original values, which suggested the decomplexation of complex **1@G1**. Moreover, white precipitates were observed, indicating the formation of water-insoluble protonated host **1**. When aqueous NaOD was added into the above mixture, the solution changed from acidic to neutral, the signals of protons a-f in **G1** shifted upfield to their complexed values, while the signal of protons H₁ and H₂ in host **1** appeared again and the white precipitates disappeared, which indicated the reformation of complex **1@G1**. After the addition of NaOD, the chemical shift of host and guest protons could not be completely restored, which may due to the formation of NaCl in solution. Since the anion in the guest molecule is iodine, after adding DCl and NaOD, there are iodine and chlorine anions in the solution, resulting in changes in the chemical shift of the formed complex. The appearance and disappearance of the precipitates might be mainly due to the protonation and deprotonation of carboxylate groups on host **1**. Addition of aqueous DCl resulted in the precipitation of water-insoluble protonated host **1**, and led to the decomplexation of complex **1@G1**. However, upon the addition of aqueous NaOD, the carboxylic acid groups of host **1** were converted to carboxylate groups, and complex **1@G1** was reformed. Thus, the complexation and decomplexation of complex **1@G1** could be reversibly controlled by acid/based and observed by naked eye (Fig. 4). Similar to the case of complex **1@G1**, the complexation and decomplexation process of complexes **1@G2**, **1@G3** and **1@G4** can also be reversibly controlled by acid/based and observed by naked eye.

Conclusions

In summary, we have successfully synthesized 1,1':2',1'-terphenyl-based water soluble macrocyclic arenes **1** that containing of anionic carboxylate groups, and studied its binding ability toward *N*-methylquinolinium salts in water. It was found that host **1** could form 1 : 1 complex with *N*-methylquinolinium salts in water. Since the conversion of anionic carboxylate groups and neutral carboxylic groups can be controlled by the addition of acid and base. Thus, the complexation and decomplexation of formed complexes can be achieved by the addition of acid and base and the process can be observed by naked eye. The high affinity ability in water and controllable host-guest system of this new water-soluble host could be utilized to fabricate functional supramolecular assemblies, which are ongoing in our laboratory.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We are grateful for the financial support from the National Natural Science Foundation of China (No. 21602055 and 51772091); Natural Science Foundation of Hunan Province (No. 2017JJ3094). Research Project on Teaching Reform of Ordinary Colleges and Universities in Hunan Province: [2018] (No. 436: 670).

Notes and references

- 1 C. F. Chen and Y. Han, *Acc. Chem. Res.*, 2018, **51**, 2093–2106.
- 2 T. Schrader and A. D. Hamilton, *Functional Synthetic Receptors*, Wiley-VCH, Weinheim, Germany, 2005.
- 3 F. Zeng, L. Cheng, G.-C. Ou, L.-L. Tang and M.-H. Ding, *J. Org. Chem.*, 2022, **87**, 3863–3867.
- 4 M.-H. Ding, J. Liao, L.-L. Tang, G.-C. Ou and F. Zeng, *Chin. Chem. Lett.*, 2021, **32**, 1665–1668.
- 5 F. Zeng, L. Cheng, W.-J. Zhang, L.-L. Tang and X.-F. Wang, *Org. Chem. Front.*, 2022, **9**, 3307–3311.
- 6 F. Zeng, X.-S. Xiao, S.-F. Gong, L. Yuan and L.-L. Tang, *Org. Chem. Front.*, 2022, **9**, 4829–4833.
- 7 Q. Shi, X. Wang, B. Liu, P. Qiao, J. Li and L. Wang, *Chem. Commun.*, 2021, **57**, 12379–12405.
- 8 H. Yao, S.-Y. Li, H. Zhang, X.-Y. Pang, J.-L. Lu, C. Chen, W. Jiang, L.-P. Yang and L.-L. Wang, *Chem. Commun.*, 2023, **59**, 5411–5414.
- 9 F. Zeng, L.-L. Tang, H. Yu, F.-P. Xu and L. Wang, *Chin. Chem. Lett.*, 2023, DOI: [10.1016/j.ccl.2023.108304](https://doi.org/10.1016/j.ccl.2023.108304).
- 10 P. Neri, J. L. Sessler and M.-X. Wang, *Calixarenes and Beyond*, Springer, Cham, Switzerland, 2016.
- 11 W. Sliwa and C. Kozłowski, *Calixarenes and Resorcinarenes: Synthesis, Properties and Applications*, Wiley-VCH, Weinheim, Germany, 2009.
- 12 Y. Ding, J. Jiao, B. Sun, Z. Yang, L. Chen and L. Wang, *Chin. Chem. Lett.*, 2021, **32**, 3539–3543.
- 13 T. Ogoshi, S. Kanai, S. Fujinami, T.-A. Yamagishi and Y. Nakamoto, *J. Am. Chem. Soc.*, 2008, **130**, 5022–5023.
- 14 T. Ogoshi, *Pillararenes*, Royal Society of Chemistry, Cambridge, U.K., 2016.
- 15 T. Ogoshi, T.-A. Yamagishi and Y. Nakamoto, *Chem. Rev.*, 2016, **116**, 7937–8002.
- 16 M. Xue, Y. Yang, X. Chi, Z. Zhang and F. Huang, *Acc. Chem. Res.*, 2012, **45**, 1294–1308.
- 17 P. D. Sala, R. D. Regno, C. Talotta, A. Capobianco, N. Hickey, S. Geremia, M. De Rosa, A. Spinella, A. Soriente, P. Neri and C. Gaeta, *J. Am. Chem. Soc.*, 2020, **142**, 1752–1756.
- 18 X.-N. Han, Y. Han and C.-F. Chen, *J. Am. Chem. Soc.*, 2020, **142**, 8262–8269.
- 19 X.-N. Han, Q.-S. Zong, Y. Han and C.-F. Chen, *CCS Chem.*, 2021, **3**, 738–750.
- 20 J. R. Wu and Y. W. Yang, *J. Am. Chem. Soc.*, 2019, **141**, 12280–12287.
- 21 J. R. Wu, G. Wu and Y. W. Yang, *Acc. Chem. Res.*, 2022, **55**, 3191–3204.
- 22 W. Yang, K. Samanta, X. Wan, T. U. Thikekar, Y. Chao, S. Li, K. Du, J. Xu, Y. Gao, H. Zuillhof and A. C.-H. Sue, *Angew. Chem., Int. Ed.*, 2020, **59**, 3994–3999.



- 23 M. Guo, X. Wang, C. Zhan, P. D. Drouhard, W. Li, K. Du, M. A. Olson, H. Zuillhof and A. C.-H. Sue, *J. Am. Chem. Soc.*, 2018, **140**, 74–77.
- 24 H. Chen, J. Fan, X. Hu, J. Ma, S. Wang, J. Li, Y. Yu, X. Jia and C. Li, *Chem. Sci.*, 2015, **6**, 197–202.
- 25 Z.-Y. Zhang and C. Li, *Acc. Chem. Res.*, 2022, **55**, 916–929.
- 26 K. Xu, Z. Y. Zhang, Z. Zhou and C. Li, *Chin. Chem. Lett.*, 2022, **33**, 2451–2454.
- 27 G.-W. Zhang, P.-F. Li, Z. Meng, H.-X. Wang, Y. Han and C.-F. Chen, *Angew. Chem., Int. Ed.*, 2016, **55**, 5304–5308.
- 28 Q. Shi and C.-F. Chen, *Org. Lett.*, 2017, **19**, 3175–3178.
- 29 Q. Shi, Y. Han and C.-F. Chen, *Chem.-Asian J.*, 2017, **12**, 2576–2582.
- 30 J. Q. Wang, Y. Han and C.-F. Chen, *Chem. Commun.*, 2021, **57**, 3987–3990.
- 31 J. Li, H.-Y. Zhou, Y. Han and C.-F. Chen, *Angew. Chem., Int. Ed.*, 2021, **60**, 21927–21933.
- 32 L. Escobar and P. Ballester, *Chem. Rev.*, 2021, **121**, 2445–2514.
- 33 F. Zeng and C.-F. Chen, *Org. Biomol. Chem.*, 2015, **13**, 1988–1991.
- 34 S. Kubik, *Chem. Soc. Rev.*, 2010, **39**, 3648–3663.
- 35 J. Murray, K. Kim, T. Ogoshi, W. Yao and B. C. Gibb, *Chem. Soc. Rev.*, 2017, **46**, 2479–2496.
- 36 W. Wang, Z. Li, C. Song, J. Yang and Y. W. Yang, *Molecules*, 2022, **23**, 8554.
- 37 J. Mendoza, L. Cruz, V. Freitas, F. Pina and N. Basilio, *Molecules*, 2021, **26**, 5389.
- 38 J.-R. Wu, G. Wu, Z. Cai, D. Li, M. H. Li, Y. Wang and Y.-W. Yang, *Molecules*, 2022, **27**, 6259.
- 39 S. Dasgupta and P. S. Mukherjee, *Org. Biomol. Chem.*, 2017, **15**, 762–772.
- 40 T. Ogoshi, M. Hashizume, T. Yamagishi and Y. Nakamoto, *Chem. Commun.*, 2010, **46**, 3708–3710.
- 41 Z. Li, J. Yang, G. Yu, J. He, Z. Abliz and F. Huang, *Chem. Commun.*, 2014, **50**, 2841–2843.
- 42 J. Yang, X. Chi, Z. Li, G. Yu, J. He, Z. Abliz, N. Li and F. Huang, *Org. Chem. Front.*, 2014, **1**, 630–633.
- 43 D. S. Guo and Y. Liu, *Acc. Chem. Soc.*, 2014, **47**, 1925–1934.
- 44 Z. Liu, X. Sun, X. Dai, J. Li, P. Li and Y. Liu, *J. Mater. Chem. C*, 2021, **9**, 1958–1965.
- 45 L. Memmi, A. Lazar, A. Brioude, V. Ball and A. W. Coleman, *Chem. Commun.*, 2001, 2474–2475.
- 46 X. Huang, X. Zhang, T. Qian, J. Ma, L. Cui and C. Li, *Beilstein J. Org. Chem.*, 2018, **14**, 2236–2241.
- 47 J. Ma, Q. Meng, X. Hu, B. Li, S. Ma, B. Hu, J. Li, X. Jia and C. Li, *Org. Lett.*, 2016, **18**, 5740–5743.

