

Showcasing research from Professor Ok-Sang Jung's laboratory, Department of Chemistry, Pusan National University, Busan, Republic of Korea

Molecular balloon,  $\mathrm{Pd_6L_8}$  cages: recognition of alkyl sulfate surfactants

The inner cavity of Pd(II) molecular balloons was controlled by anion exchange of nitrate with alkyl sulfate. Contact angles for their crystalline solid surface were measured.





## ChemComm



### COMMUNICATION

View Article Online
View Journal | View Issue



Cite this: Chem. Commun., 2020, 56, 2841

Received 16th December 2019, Accepted 6th February 2020

DOI: 10.1039/c9cc09742b

rsc.li/chemcomm

# Molecular balloon, Pd<sub>6</sub>L<sub>8</sub> cages: recognition of alkyl sulfate surfactants†

Haeri Lee,‡ Dongwon Kim,‡ Hyejin Oh and Ok-Sang Jung \*\*D\*\*

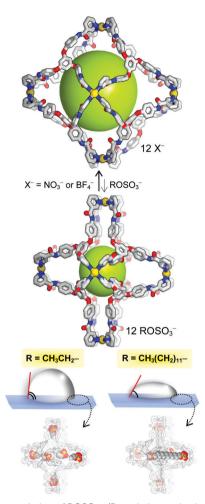
The unique molecular balloon system of [Pd<sub>6</sub>L<sub>8</sub>](NO<sub>3</sub>)<sub>12</sub> (an inner cavity of 19  $\times$  21  $\times$  25 Å<sup>3</sup>  $\rightleftharpoons$  13  $\times$  13  $\times$  13 Å<sup>3</sup>) was carried out *via* the anion exchange of nitrate with alkyl sulfates.

Reversible molecular encapsulation via the control of a trigger plays a major role in important chemical recognition processes including biological signal transduction and enzyme catalysis.<sup>1</sup> Recently, chemical scientists have attempted to exploit intermolecular interactions as an alternative means of significant control over the threshold of encapsulation.<sup>2</sup> Such interactions tend to mimic the supramolecular system that brings changes to the encapsulated substrates via the desired hydrophobic/ hydrophilic pores.<sup>3</sup> The encapsulated substrate can be activated by the influence of weak interactions or external stimuli. <sup>4</sup> Thus, research on the construction of desirable tailor-made large cages has been a hot issue, owing to diverse task-specific functions such as guest indicators, catalysts, solvent reservoirs, regionselectivity controllers, and drug-delivery systems.<sup>5</sup> Among various cage compounds, elegant cationic palladium(II) coordination cages have exhibited molecular recognition depending on the counteranionic nature and confined space.<sup>6</sup> Furthermore, postmodification of coordination cages for control of their physicochemical properties has attracted particular attention for its provision of facile variation of function.<sup>7</sup> However, to date, it remains difficult to include various guest molecules in coordination cages, due to the fixed size of the cavity. Both the internal space and the functional sites of such coordination cages have been known to be important factors for encapsulation and confinement effects.8 A series of alkyl sulfate salts of C<sub>n</sub>H<sub>2n+1</sub>OSO<sub>3</sub> have been used in ubiquitous detergents for dish-washing liquids, shower gels, shampoos, hair conditioners, fabric softeners,

Department of Chemistry, Pusan National University, Busan 46241, Republic of Korea. E-mail: oksjung@pusan.ac.kr; Fax: +82 51-5163522; Tel: + 82 51-5103240 † Electronic supplementary information (ESI) available: Experimental details and crystal structure determination. CCDC 1888048–1888051. HR-ESI-TOF-Mass spectra, TGA/DSC curves, NMR spectra, IR spectra and SEM images. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c9cc09742b ‡ Haeri Lee and Dongwon Kim contributed equally to this work.

cosmetics, medicines, toothpaste emulsifiers, and fire-extinguishing agents. Thus, recognition of and analytical techniques for alkyl sulfate surfactants are significant issues, owing to their widespread use.

Herein we present an unprecedented novel approach to the control of the uptake of alkyl sulfate surfactants based on flexible host-guest interaction between alkyl sulfates and large coordination cages. For the aggregates encapsulated via hydrogen bonds between sulfate groups and cage skeletons, the contact angles were measured according to the length of the alkyl sulfates. Electrostatic interaction between Pd<sup>2+</sup>···-OSO<sub>3</sub>R (3.92(1)-4.23(2) Å) forced to face to each other inside cage (Table S2, ESI†). A new  $C_3$ -symmetric N-donor ligand, 1,3,5tris(4-nicotinamide-phenoxy)benzene (L), was synthesized in three steps in a reasonable yield starting from phloroglucinol. The L was employed in the self-assembly of palladium(II) ions for the coordination cages  $[Pd_6L_8]^{12+}(X^-)_{12} (X^- = NO_3^-, 1.12NO_3;$  $X^- = BF_4^-$ , 1·12BF<sub>4</sub>) (Scheme 1). The reaction was monitored by <sup>1</sup>H NMR spectra that showed a significant downfield shift compared with that of the free L (Fig. S1-S5 and S8-S11, ESI†). High-resolution ESI-mass data at 1711.2797 ([1·12NO<sub>3</sub>-4NO<sub>3</sub>]<sup>4+</sup>, calcd 1711.276), 1356.6252 ([1·12NO<sub>3</sub>-5NO<sub>3</sub>]<sup>5+</sup>, calcd 1356.623) for  $1.12NO_3$  and at 1761.2153 ([ $1.12BF_4-4BF_4$ ]<sup>4+</sup>, calcd 1761.306), 1391.6473 ( $[1.12BF_4-5BF_4]^{5+}$ , calcd 1391.661), 1145.2067  $([1.12BF_4-6BF_4]^{6+}$ , calcd 1145.203) for  $1.12BF_4$  were consistent with the formation of the coordination cages. Their crystal structures were investigated via single-crystal X-ray structure determination. The crystalline solids of the cages are insoluble in common organic solvents such as chloroform, acetonitrile, and tetrahydrofuran, but are soluble in Me<sub>2</sub>SO. Subsequent reactions with the addition of tetrabutylammonium alkyl sulfates  $(ROSO_3^-NBu_4^+, R = ethyl-, octyl-, decyl-, dodecyl-, and tetradecyl-),$ [Pd<sub>6</sub>(ROSO<sub>3</sub>)<sub>6</sub>L<sub>8</sub>](ROSO<sub>3</sub>)<sub>6</sub> (2·12ROSO<sub>3</sub>) were attempted. It is noteworthy that the crystalline solid of 2.12ROSO3 has a much lower solubility in common organic solvents including N,N-dimethylformamide and Me<sub>2</sub>SO. In a trial entailing X-ray diffraction data collection, we could calculate unit cell parameters for 2·12C<sub>2</sub>H<sub>5</sub>OSO<sub>3</sub> and 2·12C<sub>14</sub>H<sub>29</sub>OSO<sub>3</sub> that were Communication ChemComm



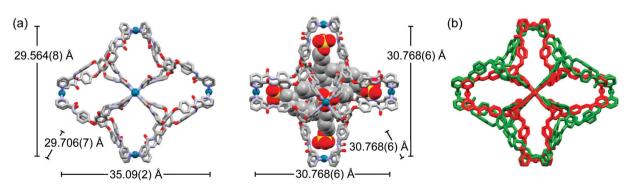
 $\begin{array}{ll} \textbf{Scheme 1} & \textbf{Encapsulation of ROSO_3}^- \ (\textbf{R} = \textbf{ethyl-, octyl-, decyl-, dodecyl-,} \\ \textbf{and tetradecyl-)} \ within present spherical coordination cages and different surface tensions depending on included alkyl chains of surfactant. \\ \end{array}$ 

similar to those of  $2\cdot12C_8H_{17}OSO_3$  and  $2\cdot12C_{12}H_{25}OSO_3$ , due to weak diffraction. The IR spectra for  $2\cdot12ROSO_3$  showed the S–O vibrational frequencies of the sulfate group (1361 and 1064 cm<sup>-1</sup>) and C–H from the aliphatic chains (1925–2852 cm<sup>-1</sup>) (Fig. S15, ESI†). Thermogravimetric analysis (TGA) and differential scanning calorimetry (DSC) results showed that the coordinate cages of  $1\cdot12NO_3$  were stable up to 367 °C and that the solvate

molecules were evaporated at 350 °C (two-step weight loss for solvate molecules: found, 4.7%; calcd, 9.3%, found, 24%; calcd, 27%).  $1\cdot12BF_4$  decomposed at 297 °C, up to the point, the solvate molecules were evaporated (two-step weight loss for solvate molecules: found, 8.1%; calcd, 7.9%, found, 20%; calcd, 25%).  $2\cdot12C_{12}H_{25}OSO_3$  started to be decomposed at temperatures above 420 °C, and the solvate molecules were fully evaporated by 300 °C (found, 23%; calcd, 22%).

In the crystal structures of the Pd<sub>6</sub>L<sub>8</sub> cages, the geometry of the palladium(II) ion was a typical square-planar arrangement with four N-donors from four L (Pd-N = 1.97(2)-2.048(6) Å). The diameter of all of the present cages was ~4 nm. However, the dimensions of the inner cavity for each cage were significantly different depending on the anions (19  $\times$  21  $\times$  25 Å<sup>3</sup> and  $13 \times 13 \times 13 \text{ Å}^3$  for  $1.12\text{NO}_3$  and  $2.12\text{C}_{12}\text{H}_{25}\text{OSO}_3$ , respectively) (Scheme 1 and Fig. S19, ESI†). The encapsulated alkyl sulfates were bound with skeletal cages via hydrogen bonds (RO<sub>3</sub>S-O···H-NCO = 2.11(1)-2.66(6) Å) for  $2.12C_8H_{17}OSO_3$  and 1.98(2)-2.38(1) Å for 2·12C<sub>12</sub>H<sub>25</sub>OSO<sub>3</sub> following the conformational change in the amide bonds, which was a driving force of the cage contraction (Fig. S19, ESI†). The solvent-accessible void volume was calculated as 63.0% (11695.6/18556.0 Å<sup>3</sup>) for 1·12NO<sub>3</sub>, 50.2% (26491.5/  $52684.0 \text{ Å}^3$ ) for  $2 \cdot 12C_8H_{17}OSO_3$ , and 45.3% (23932.4/52813.0 Å<sup>3</sup>) for 2·12C<sub>12</sub>H<sub>25</sub>OSO<sub>3</sub> based on PLATON. 11 For the 2·12C<sub>8</sub>H<sub>17</sub>OSO<sub>3</sub> and 2·12C<sub>12</sub>H<sub>25</sub>OSO<sub>3</sub> structures, a half amount of alkyl sulfates were found on the electron density map, which showed them to exist at the apical position of the palladium(II) inside the cages (Fig. 1b), whereas the opposite position was occupied by the Me<sub>2</sub>SO solvate molecules (Me<sub>2</sub>S =  $O \cdot \cdot \cdot Pd(II) = 2.96(2)$  Å for 2·12C<sub>8</sub>H<sub>17</sub>OSO<sub>3</sub> and 2.88(3) Å for 2·12C<sub>12</sub>H<sub>25</sub>OSO<sub>3</sub>).

In <sup>1</sup>H NMR spectra, the addition of sodium dodecyl sulfate to the Me<sub>2</sub>SO-d<sub>6</sub> solution of 1·12NO<sub>3</sub> produced significant chemical shifts in the protons of the amide and pyridyl moieties, indicating the contracted cages' strong hydrogen bonds between the sulfate groups and the cage-skeletal structures (Fig. 2c). That is, the intermolecular interactions between the coordination cages and alkyl sulfates resulted in downfield shifts. The gradual addition up to 12 equiv. of alkyl sulfate to the 1·12NO<sub>3</sub> solution attained the equilibrium state with a 2:3 integral ratio of contracted and expanded cages in the solution state along with a small amount of free L owing to partial dissociation of the cages in the Me<sub>2</sub>SO solution. However, either addition of an excessive amount of



 $\textbf{Fig. 1} \quad \text{Crystal structures and Pd} \cdot \cdot \cdot \text{Pd distances for } \textbf{1} \cdot 12\text{NO}_3 \text{ (a, left) and } \textbf{2} \cdot \text{C}_8 \text{H}_{15} \text{OSO}_3 \text{ (a, right)}. Overlay (b) of } \textbf{1} \cdot 12\text{NO}_3 \text{ (green) and } \textbf{2} \cdot \text{C}_8 \text{H}_{15} \text{OSO}_3 \text{ (red)}. \\ \textbf{2} \cdot \text{C}_8 \text{H}_{15} \text{OSO}_3 \text{ (a, right)}. \\ \textbf{3} \cdot \text{C}_8 \text{H}_{15} \text{OSO}_3 \text{ (red)}. \\ \textbf{3} \cdot \text{C}_8 \text{H}_{15} \text{OSO}_3 \text{ (a, right)}. \\ \textbf{4} \cdot \text{C}_8 \text{H}_{15} \text{OSO}_3 \text{ (red)}. \\ \textbf{5} \cdot \text{C}_8 \text{H}_{15} \text{OSO}_3 \text{ (red)}. \\ \textbf{5} \cdot \text{C}_8 \text{H}_{15} \text{OSO}_3 \text{ (red)}. \\ \textbf{6} \cdot \text{C}_8 \text{$ 

ChemComm Communication

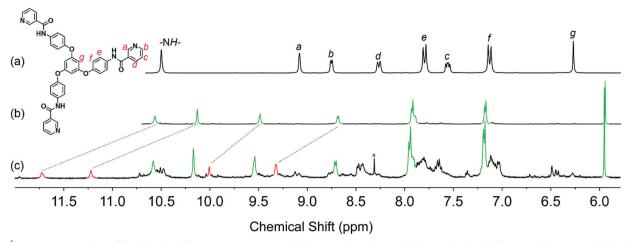


Fig. 2  $^{1}$ H NMR spectra for L (a), 1.12NO<sub>3</sub> (b), and after addition of 12 equiv. of NaC<sub>12</sub>H<sub>25</sub>OSO<sub>3</sub> into 1.12NO<sub>3</sub> (c) in Me<sub>2</sub>SO- $d_6$ . The 1.12NO<sub>3</sub> and 2.12C<sub>12</sub>H<sub>25</sub>OSO<sub>3</sub> resonances are indicated by the green and red signals, respectively

alkyl sulfate or rapid addition results in precipitation from the solution, owing to the low solubility. Of course, crystalline solids of 2.12ROSO3 are rarely soluble in common organic solvents. The significant solubility difference seems to come from low solvation energy between cage and solvent molecules, presumably owing to suitable interactions between cages and alkyl sulfates. The present IR spectra support our hypothesis that the anion exchange is reversible between NO<sub>3</sub><sup>-</sup> and alkyl sulfate (Fig. S21, ESI†).

In order to investigate the recognition and adsorption properties of the flexible cages, the contact angles of a droplet on the surface of fine-ground microcrystals (20-40 µm) were measured (Fig. S22, ESI†). The crystalline surface was prepared by spreading the ground microcrystals on a glass tape and subsequently flattening them onto slide glass. As shown in Fig. 3 and Table S3 (ESI†), the contact angles for a drop of water to 2·12C<sub>14</sub>H<sub>29</sub>OSO<sub>3</sub> (114.87°) were larger than those of cages containing 2·12C2H5OSO3 with ethyl analog (102.20°). Furthermore, contact angles increase according to the carbon number (n) of alkyl sulfates, thus indicating that the surface of the long chain crystals is more hydrophobic. In the present experimentation, the contact angles slightly decreased to 80.54°  $(n=2) \sim 100.80^{\circ}$  (n=14) after 10 min. By contrast, for a drop of 20% ethanol aqueous solution, the contact angles drastically decreased (from 99.94° to 35.84° on 2·12C<sub>14</sub>H<sub>29</sub>OSO<sub>3</sub>) after 10 min, as depicted in Fig. 3. The alcohol aqueous drop induced water adsorption into the void of the microcrystalline structure, as caused by hydrophilic structure transformation.<sup>12</sup> Furthermore, the reason it takes several minutes to wet the surface is that adsorption occurs in the heterogeneous phase. This process is an advanced method for surface modification of crystals via anion exchange by alkyl sulfate surfactants. Furthermore, the wettability of 2 crystals can be utilized as a molecular ruler for measurement of the length of alkyl sulfate chains.

In conclusion, we demonstrated an unusual system for contraction and expansion of flexible Pd<sub>6</sub>L<sub>8</sub> cages via inclusion and release of alkyl sulfate surfactants. Hydrogen bonds between surfactants and cage structures are the main driving force of cage transformation. Furthermore, the microcrystal surface of

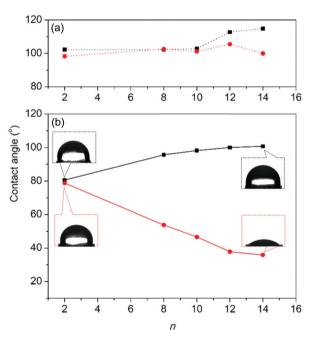


Fig. 3 Surfactant responsive contact angles on fine-ground crystal surface of  $2.12C_nH_{2n+1}OSO_3$  at 0 min (a) and after 10 min (b) (black line: drop of H<sub>2</sub>O; red line: mixture solution of water and ethanol (4:1)).

the Pd<sub>6</sub>L<sub>8</sub> cages presents the opportunity for tremendous modification of physical properties according to the chain length of sulfate anions. In particular, the relationship between surfactant recognition and hydrophobic properties has been demonstrated by dynamic contact angles showing surfactantenhanced spread. That is, these cages can be an unprecedented recognizer for C<sub>n</sub>H<sub>2n+1</sub>OSO<sub>3</sub> surfactants as well as a unique molecular ruler of alkyl chains via contact angles. This strategy, which involves an uncommon synergic cage effect, might find important applications for spatial and chain-length control entailing the release and retention of molecules.

This work was supported by National Research Foundation of Korea (NRF) grants funded by the Korean Government [MEST]

Communication ChemComm

(2016R1A2B3009532 and 2016R1A5A1009405). X-ray crystallography at the PLS-II 2D-SMC beamline was supported in part by MSIP and POSTECH.

### Conflicts of interest

There are no conflicts to declare.

### Notes and references

- 1 (a) C. Edlinger, T. Einfalt, M. Spulber, A. Car, W. Meier and C. G. Palivan, Nano Lett., 2017, 17, 5790; (b) G. Grossi, M. D. E. Jepsen, J. Kjems and E. S. Andersen, Nat. Commun., 2017, 8, 1.
- 2 (a) V. H. Pérez-Luna and O. González-Reynoso, Gels, 2018, 4, 61;
  (b) E. M. M. D. Valle, Process Biochem., 2004, 39, 1033;
  (c) J. Rebek, Jr., Angew. Chem., Int. Ed., 2005, 44, 2068.
- 3 (a) D. K. Singh, B. V. V. S. P. Kumar and M. Eswaramoorthy, Nanoscale, 2015, 7, 13358; (b) Y. Jang, R. Natarajan, Y. H. Ko and K. Kim, Angew. Chem., Int. Ed., 2014, 53, 1003; (c) D. Prochowicz, A. Kornowicz and J. Lewiński, Chem. Rev., 2017, 117, 13461.
- (a) A. J. Harnoy, I. Rosenbaum, E. Tirosh, Y. Ebenstein, R. Shaharabani,
   R. Beck and R. J. Amir, J. Am. Chem. Soc., 2014, 136, 7531; (b) I. K. Jeong,
   G. H. Gao, Y. Li, S. W. Kang and D. S. Lee, Macromol. Biosci., 2013,
   13, 946; (c) Y. Y. Xu, G. Li, W. Zhuang, H. C. Yu, Y. Hu and Y. Wang,
   J. Mater. Chem. B, 2018, 6, 7495.
- (a) D. Shetty, et al., ACS Appl. Mater. Interfaces, 2018, 10, 17359;
   (b) M. Hatano, T. Sakamoto, T. Mizuno, Y. Goto and K. Ishihara, J. Am. Chem. Soc., 2018, 140, 16253;
   (c) S. Kawano, T. Fukushima and K. Tanaka, Angew. Chem., Int. Ed., 2018, 130, 15043;
   (d) J. Guo, Y.-W. Xu, K. Li, L.-M. Xiao, S. Chen, K. Wu, X.-D. Chen, Y.-Z. Fan, J.-M. Liu and

- C.-Y. Su, Angew. Chem., Int. Ed., 2017, 56, 3852; (e) I. Ortiz-Rivera, M. Mathesh and D. A. Wilson, Acc. Chem. Res., 2018, 51, 1891.
- 6 (a) T. Murase, Y. Nishijima and M. Fujita, J. Am. Chem. Soc., 2012, 134, 162; (b) W. Cullen, M. C. Misuraca, C. A. Hunter, N. H. Williams and M. D. Ward, Nat. Chem., 2016, 8, 231; (c) S. Löffler, J. Lübben, L. Krause, D. Stalke, B. Dittrich and G. H. Clever, J. Am. Chem. Soc., 2015, 137, 1060; (d) Y. Nishioka, T. Yamaguchi, M. Yoshizawa and M. Fujita, J. Am. Chem. Soc., 2007, 129, 7000; (e) T. Y. Kim, R. A. S. Vasdev, D. Preston and J. D. Crowley, Chem. Eur. J., 2018, 24, 14878; (f) A. M. Castilla, T. K. Ronson and J. R. Nitschke, J. Am. Chem. Soc., 2016, 138, 2342; (g) J. E. M. Lewis, A. B. S. Elliott, C. J. McAdam, K. C. Gordon and J. D. Crowley, Chem. Sci., 2014, 5, 1833.
- 7 (a) M. Han, R. Michel, B. He, Y.-S. Chen, D. Stalke, M. John and G. H. Clever, Angew. Chem., Int. Ed., 2013, 52, 1319; (b) R. Custelcean, Chem. Soc. Rev., 2014, 43, 1813; (c) J. Mosquera, S. Zarra and J. R. Nitschke, Angew. Chem., Int. Ed., 2014, 53, 1556; (d) L. M. Mesquita, J. Anhäuser, D. Bellaire, S. Becker, A. Lützen and S. Kubik, Org. Lett., 2019, 21(16), 6442; (e) L. Escobar, D. Villarón, E. C. Escudero-Adán and P. Ballester, Chem. Commun., 2019, 55, 604.
- 8 C. M. Hong, D. M. Kaphan, R. G. Bergman, K. N. Raymond and F. D. Toste, *J. Am. Chem. Soc.*, 2017, **139**, 8013.
- 9 (a) M. Tsumadori, CD Proceedings 6th World Surfactant Congress CESIO, 2004, 196; (b) D. A. Korolchenko, E. N. Degaev and A. F. Sharovarnikov, ICMEA 2015, 2015, 23; (c) R. Vanyúr, L. Biczók and Z. Miskolczy, Colloids Surf., A, 2007, 299, 256.
- 10 (a) K. M. Ko, B. H. Chon, S. B. Jang and H. Y. Jang, J. Ind. Eng. Chem., 2014, 20, 228; (b) J. Huang. and H. R. Matthews, Anal. Biochem., 1990, 188, 114.
- 11 A. L. Spek, Acta Crystallogr., 2009, D65, 148.
- 12 (a) J. F. Wall and C. F. Zukoski, *Langmuir*, 1999, 15, 7432; (b) Y. Yuan and T. R. Lee, *Surface Sciences*, Springer, New York, 2013, vol. 51, p. 3; (c) X. Wang, J. Venzmer and E. Bonaccurso, *Langmuir*, 2016, 32, 8322.