

Cite this: *RSC Adv.*, 2017, 7, 37474

Received 18th July 2017

Accepted 21st July 2017

DOI: 10.1039/c7ra07915j

rsc.li/rsc-advances

## Facile synthesis of novel hybrid POSS biomolecules via "Click" reactions†

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A novel alkyne-terminated cubic-octameric POSS was synthesised in high yield (82–90%). The X-ray crystal structure revealed intra- and intermolecular hydrogen bonding between the amide groups of the arms. Hybrid biomaterials were synthesised in nearly quantitative yields via a click reaction with (i) azido-*N*-Fmoc-norleucine and (ii) 3'-azido-3'-deoxythymidine.

Among the most commonly studied scaffolds for developing hybrid biomaterials<sup>1–4</sup> is polyhedral oligomeric silsesquioxane (POSS). POSS units are symmetrical, three-dimensional cubic molecules, which are unique nanometer-sized hybrid inorganic-organic materials<sup>5,6</sup> with the formula (RSiO<sub>3/2</sub>)<sub>8</sub>, known as T<sub>8</sub>. POSS contains an inorganic inner siloxane nanocore, with the possibility of chemical functionalisation at each of the eight corners of the cubic unit. POSS units have been used extensively as scaffolds for the development of liquid crystals,<sup>7,8</sup> biocompatible materials,<sup>9</sup> catalysts<sup>10,11</sup> and dendrimers<sup>12</sup> and can also be used in cross-linking polymers.<sup>13</sup> Functionalisation of T<sub>8</sub> with different substituents has usually been achieved by hydro-silylation,<sup>14,15</sup> Heck,<sup>16–18</sup> and cross-metathesis<sup>16,19</sup> reactions.

Copper-catalyzed Azide-Alkyne Cycloaddition (CuAAC),<sup>20,21</sup> 'click' chemistry, is a simple method for coupling organic molecules containing azide and alkyne functional groups in high yields and its use in the fields of peptide and protein biomedical and material sciences is accelerating.<sup>22,23</sup> The click reaction has been used to synthesise POSS biomaterials such as hybrid POSS-PEG hydrogels<sup>24</sup> that support chondrocyte attachment and proliferation. Only one synthetic approach towards peptidyl silsesquioxanes using click chemistry has been reported to date.<sup>25–30</sup> focussing on the synthesis of octa(3-azidopropyl)polyhedral oligomeric silsesquioxane POSS-(N<sub>3</sub>)<sub>8</sub> (Fig. 1) and its reaction with a variety of alkynes.<sup>25–36</sup> The

preparation of octa(3-azidopropyl)polyhedral oligomeric silsesquioxane POSS-(N<sub>3</sub>)<sub>8</sub> requires a multi-step synthesis and suffers from a poor overall yield of 30–32%.<sup>32</sup>

The synthesis and click reaction of mono alkyne-heptaisobutyl POSS were first reported by Müller *et al.*<sup>37</sup> and Wu *et al.*<sup>38</sup> Müller *et al.*<sup>37</sup> have used aminopropylheptaisobutyl polyhedral oligomeric silsesquioxane (POSS) to produce mono-alkyne-POSS in three steps and the click coupling occurred between the azido-terminated polystyrenes and alkyne-POSS to afford mono-, di-, and pentafunctional POSS-containing hybrid polystyrenes as star-shaped telechelic POSS-containing hybrid polymers. Wu *et al.*<sup>38</sup> have prepared alkyne-functionalized-POSS in three steps from allyl-heptaisobutyl substituted-POSS and the product was linked to an azido-functionalized elastomer of poly(styrene-*b*-(ethylene-*co*-butylene)-*b*-styrene) (SEB-CH<sub>2</sub>N<sub>3</sub>) via a click coupling reaction to form a novel hybrid copolymer. These approaches focused mostly on the preparation of a hybrid copolymer based on mono-alkyne-heptaisobutyl and azido polystyrene. However, the preparation of octa-alkyne-terminated POSS and their click coupling reactions with azido-biomolecules to produce hybrid biomaterials has not been investigated yet. We believe that this approach will open a wide range of biomedical applications that were not accessible in the past.

Herein we report a novel synthetic approach to hybrid biomaterials based on octa-alkyne-terminated POSS and their reaction with (i) azido-*N*-Fmoc-L-norleucine and (ii) 3'-azido-3'-deoxythymidine, using the CuAAC reaction. Our approach

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† Electronic supplementary information (ESI) available: Detailed experimental procedures, <sup>1</sup>H, <sup>13</sup>C, <sup>19</sup>F, <sup>29</sup>Si NMR, IR, MALDI-TOF MS, elemental analysis, and analytical data for all new compounds. For ESI and crystallographic data in CIF or other electronic format. CCDC 1514595. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c7ra07915j

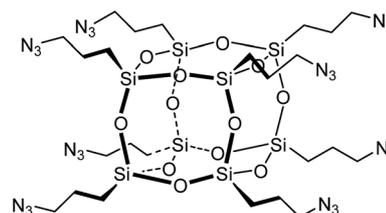
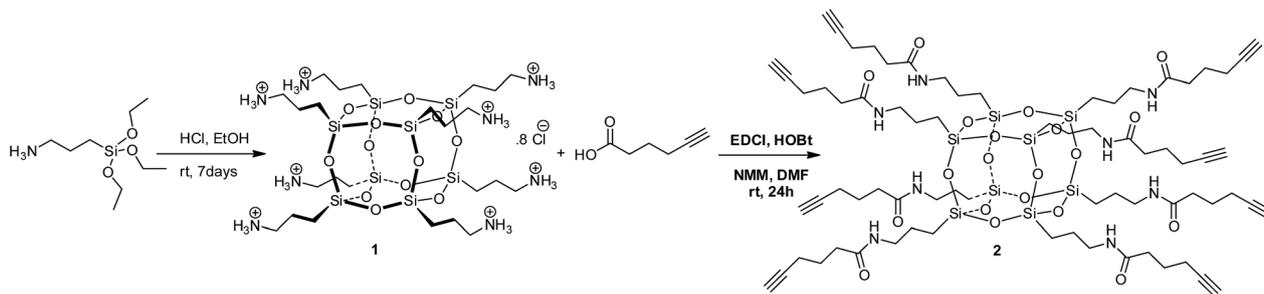


Fig. 1 Octa(3-azidopropyl)POSS.





Scheme 1 Synthesis of octa-alkyne-terminated POSS (2).

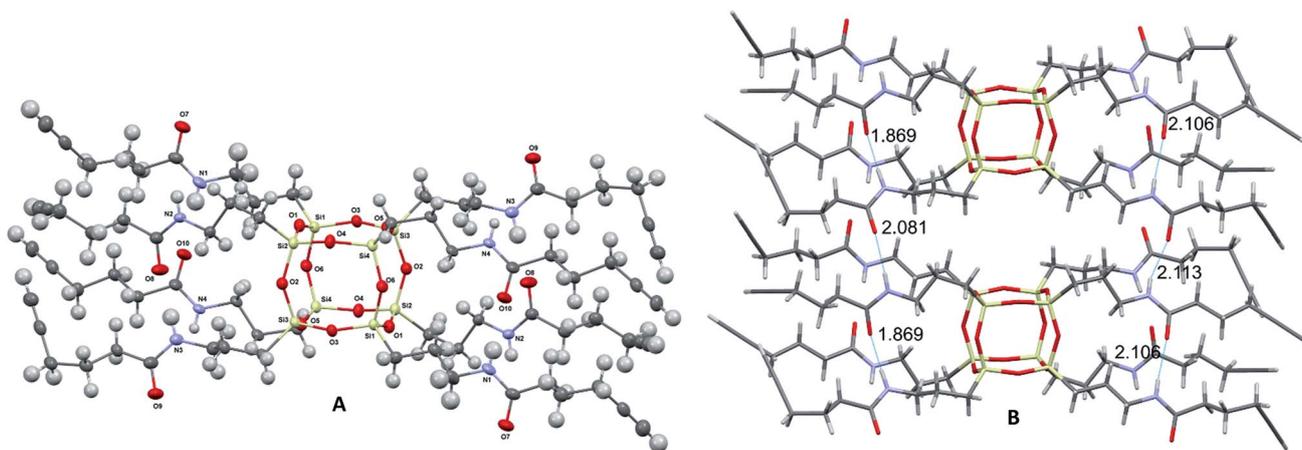


Fig. 2 Molecular structure of 2 grown fragment (thermal ellipsoids drawn with 50% probability) (A) and the font intra- and inter- H-bonding distances (B).

presents a particularly versatile route which provides a facile and convenient way to functionalise a cubic silsesquioxane core with biomolecules that are more readily available as their azido derivative than their alkyne derivative.

Compound 2 was prepared in one step from commercially available materials; octa(3-aminopropyl)octasilsesquioxane (1) and 5-hexynoic acid (Scheme 1), in 82–90% yield. Product 2 was isolated and purified by column chromatography, followed by characterisation using standard techniques (see ESI†). The crystal structure determined by X-ray crystallography (Fig. 2) suggests that intra- and inter-molecular hydrogen bonding between the arms were a fundamental driving force for the formation of a well-defined crystal structure.‡

The length of intramolecular nitrogen–hydrogen (N⋯H) bonds varies between 2.09(3) and 2.12(3) Å, whereas for an intermolecular bond the distance is 1.87(3) Å.

The completion of the cycloaddition reaction was confirmed by MALDI-TOF and the reaction progress was monitored by

observing the disappearance of the azide asymmetric stretch at  $2093\text{ cm}^{-1}$  and the triple bond  $\text{C}\equiv\text{C}$  asymmetric stretch of  $\text{T}_g$ -[propylhex-5-ynamide]<sub>8</sub> (2) at  $2100\text{ cm}^{-1}$  by FT-IR spectroscopy together with monitoring the disappearance in the  $^{13}\text{C}$ -NMR spectrum of the two peaks (89.20 and 76.56 ppm) representing the triple bond of 2.

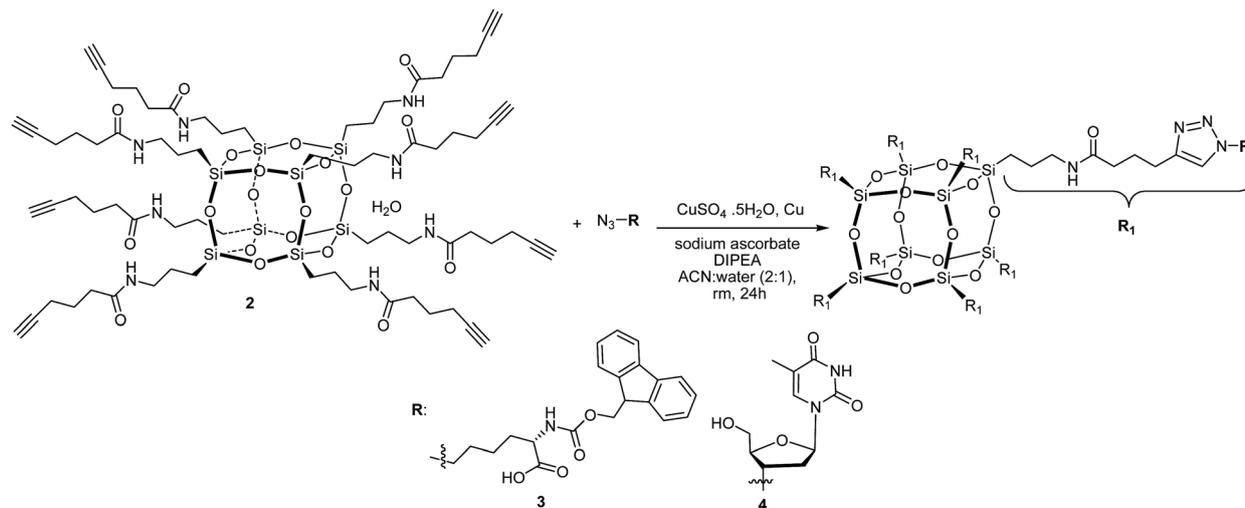
Compounds 3 and 4 have been analysed and characterised using NMR ( $^1\text{H}$ ,  $^{13}\text{C}$  and  $^{29}\text{Si}$ ) spectroscopy, infrared and MALDI-TOF mass spectrometry in positive ion mode with a DHB matrix.

Trastoy *et al.*<sup>32</sup> have reported an efficient preparation of highly functionalised cubic-octameric POSS frameworks by click chemistry and the highest yield (96%) was obtained with the  $\text{CuSO}_4\cdot\text{H}_2\text{O}$ /sodium ascorbate precatalyst system<sup>20</sup> using a biphasic organic solvent/water mixture at room temperature for 24 hours. We have used these reaction conditions for the functionalisation of the octa-alkyne-terminated POSS with azido-*N*-Fmoc-norleucine and 3'-azido-3'-deoxythymidine (Scheme 2).

The MALDI-TOF MS of compound 3 and 4 revealed that the octa-alkyne-terminated POSS has been fully functionalised with azido-*N*-Fmoc-norleucine for 3 and 3'-azido-3'-deoxythymidine for 4. The molecular ion peak of 3 observed at found 4787 Da is attributed to  $[\text{M} + \text{H}]^+$  and 4 observed at 3835.3 Da is attributed to  $[\text{M} + \text{Cu}]^+$ .

‡ Crystal data of compound 2:  $\text{C}_{72}\text{H}_{112}\text{N}_8\text{O}_{20}\text{Si}_8$  ( $M = 1634.41\text{ g mol}^{-1}$ ): triclinic, space group  $P\bar{1}$  (no. 2),  $a = 9.6202(3)\text{ Å}$ ,  $b = 14.1254(3)\text{ Å}$ ,  $c = 17.6565(6)\text{ Å}$ ,  $\alpha = 71.392(2)^\circ$ ,  $\beta = 74.675(3)^\circ$ ,  $\gamma = 70.560(2)^\circ$ ,  $V = 2110.47(12)\text{ Å}^3$ ,  $Z = 1$ ,  $T = 100.15\text{ K}$ ,  $\mu(\text{Mo K}\alpha) = 0.198\text{ mm}^{-1}$ ,  $D_{\text{calc}} = 1.286\text{ g cm}^{-3}$ , 28 211 reflections measured ( $6.088^\circ \leq 2\theta \leq 50.054^\circ$ ), 7434 unique ( $R_{\text{int}} = 0.0375$ ,  $R_{\text{sigma}} = 0.0368$ ) which were used in all calculations. The final  $R_1$  was 0.0459 ( $I > 2\sigma(I)$ ) and  $wR_2$  was 0.1280 (all data).





Scheme 2 Synthesis of 3 and 4 using CuAAC approach.

## Conclusions

In this study we have described a novel, efficient method for the synthesis of 3D radially symmetrical biomolecule-POSS hybrids. We have developed a one-step synthesis of 2 from commercially available octakis(3-aminopropyl)octa-silsesquioxane (1) with high yield (82–90%). The X-ray crystal structure shows that compound 2 exhibits plane-to-plane stacking with an intra- and inter-molecular hydrogen bond network. The octa-alkyne-terminated POSS was efficiently and regioselectively octa-functionalised with two azido-R species (where R are Fmoc-Leu and thymidine) by copper(I)-catalysed 1,3-dipolar azide-alkyne cycloaddition (CuAAC) under biphasic conditions. This led to new hybrid biofunctional nanocages 3 and 4 in high yield. This new strategy of functionalisation of terminated alkyne-POSS *via* CuAAC opens many possibilities for the efficient and controlled assembly of new hybrid biomaterials with a high degree of symmetry and with carefully tailored functional properties.

## Acknowledgements

We thank the EPSRC National Mass Spectrometry Service Centre (NMSSC) at Swansea and MEDAC Ltd. of Brunel University for elemental analysis. We thank the EPSRC UK National Crystallography Service at the University of Southampton for the collection of the crystallographic data.

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