Cyclic arrays of chlorophyll molecules occur widely in photosynthetic light-harvesting systems, such as those in purple bacteria.\[1\] Synthetic rings of porphyrins are valuable models for understanding the photophysics of these light-harvesting arrays.\[2\] They are also studied as multitopic receptors for molecule recognition,\[3\] and as expanded annulenes for exploring nanoscale aromaticity.\[4\] Many large porphyrin-based macrocycles have been synthesized using oligopyridine ligands as non-covalent templates.\[5\] Here we investigate the use of a covalent template for the synthesis of an 18-porphyrin nanoring. The main advantage of a covalently attached template is that it allows coupling reactions to be carried out under conditions that would cause a non-covalent template to dissociate. Previously, covalent templates have been used to synthesize a wide variety of macrocycles,\[6\] but to the best of our knowledge, they have not been used to prepare cyclic porphyrin oligomers.

This work is part of a project directed towards the investigation of 5,15-linked porphyrin nanorings 5,15-PN and edge-fused porphyrin nanobelts EF-PN (Figure 1). There are no previous reports of macrocycles of the type 5,15-PN, although 5,10-linked cyclic porphyrin oligomers have been synthesized.\[7\] Recently, we achieved the synthesis of a 5,15-linked 24-porphyrin nanoring with a single butadiyne bridge, using non-covalent oligopyridine templates.\[8\] However, it has not yet been possible to synthesize fully 5,15-linked porphyrin nanorings using that strategy, which is why we turned to covalent templates.

Osuka and coworkers pioneered the synthesis of linear 5,15-linked porphyrin oligomers, consisting of up to 1024 porphyrin units.\[9\] These oligomers are not π-conjugated, due to the severe twist between neighboring porphyrins, but there is strong exciton coupling along the chain, leading to ultra-fast excited-state energy migration,\[10\] which makes it interesting to investigate the photophysics of 5,15-PNs. Fully 5,15-linked porphyrin nanorings are also potential precursors to edge-fused porphyrin nanobelts EF-PN. Molecular belts of these type have not yet been synthesized, although they were mentioned in a theoretical study.\[11\] They are fascinating targets because the corresponding linear porphyrin tapes display exceptional electronic delocalization, leading to optical electronic transitions in the IR\[12\] and single-molecule conductance that is almost independent of length.\[13\]

**Figure 1.** Generalized structures of a fully 5,15-linked porphyrin nanoring 5,15-PN and an edge-fused porphyrin nanobelt EF-PN. (N is the number of porphyrin units in the ring and Ar is an aryl solubilizing group.)
We used a six-armed covalent oligophenylene template to direct the formation of the 5,15-linked 18-porphyrin macrocycle 1, by oxidative coupling of the non-cyclic hexamer of trimers 2 (Scheme 1). These compounds have a mixture of zinc(II) and nickel(II) metalation to facilitate the synthesis: preliminary studies showed that meso-bromo nickel(II) porphyrins undergo more efficient Suzuki coupling, whereas oxidative meso-meso coupling occurs most easily with zinc(II) porphyrins. The most innovative step in this synthesis is perhaps the construction of 2 by trimerization of the H-shaped porphyrin hexamer 3. This trimerization strategy has the advantage that it generates 2 without forming closely related species that would be difficult to separate.

Suzuki-Miyaura coupling of dibromo nickel(II) porphyrin 4 with two equivalents of zinc porphyrin boronic ester 5, followed by Miyaura borylation of the para-chlorophenyl substituent, gave porphyrin trimer 6 (70% yield over two steps). Coupling of 6 with dibromide 7 generated porphyrin hexamer 3 in 50% yield. Cobalt-catalyzed trimerization of tolan 3 gave the porphyrin 18-mer 2, which was isolated in 25% yield after purification by recycling gel-permeation chromatography (GPC). Previously, this type of trimerization reaction has been used to synthesize radial porphyrin hexamers, but this is the first time it has been applied to create a molecule in this size regime (molecular weight of 2: 21.7 kDa). Our optimized reaction conditions for conversion of 3 to 2 require the use of one equivalent of Co$_2$(CO)$_8$ and a high concentration of 3 in toluene (35 mM) in a sealed tube at 140 °C for 16 hours. The high solubility of 3 and 2 in toluene, conferred by the octyloxy, tri-n-hexylsilyl and octyl sidechains, is crucial for the success of this reaction. The $^1$H NMR spectra of 3 and 2 are similar (Figure 2), indicating unhindered fast rotation of individual subunits on the NMR timescale. $^{13}$C NMR spectroscopy clearly shows disappearance of the alkyne signal at $\delta = 89.8$ ppm on conversion of 3 to 2, while MALDI TOF mass spectrometry shows the expected three-fold increase in molecular weight.
Meso-meso coupling and fusion of all the zinc(II) porphyrin units in 2 was carried out to give nanoring 1, without isolating the intermediate meso-meso-linked ring, using bis(trifluoroacetoxy)iodobenzene (PIFA) as the oxidant.\(^{[14,16]}\) The analytical GPC trace of the turquoise product from cyclization of 2 with PIFA showed a main fraction with the same retention time as 2, indicating a similar molecular size. However, the electronic properties manifested in the UV-vis-NIR absorption spectra are completely different (Figure 3). The most red-shifted absorption band of 1 is located at 1055 nm, in a spectral window characteristic of edge-fused Zn\(^{11}\)-Zn\(^{11}\) porphyrin dimers.\(^{[15]}\) In comparison, absorption bands of both tolan 3 and octadecamer 2 do not extend beyond 600 nm. Moreover, the absorption spectra match well with those of previously reported linear mixed-metal Ni\(^{11}\) monomer-Zn\(^{11}\) fused dimer hybrids (see Figure S50).\(^{[14]}\)

The identity of nanoring 1 was confirmed by NMR spectroscopy (Figure 2c) and mass spectrometry. The \(^1\)H NMR spectrum shows the absence of a meso proton (\(\delta = 10.2\) ppm), while retaining the symmetry of 2. Most of the resonances in the \(^1\)H NMR spectrum of 1 were assigned using 2D COSY and NOESY spectra (Figures S46–48). Insights into the 3D shapes of 1 and 2 were provided by scanning tunnelling microscopy (STM) characterization of molecules deposited on an Au(111) surface, held under vacuum conditions, by electrospray deposition from solutions in toluene/methanol (Figure 4a–c). Images for 1 are approximately circular: circumference 12 ± 1 nm; long axis 4.1 ± 0.3 nm; short axis 3.7 ± 0.3 nm. These dimensions compared well with Ni::Ni diameter of 4.7 nm from semiempirical tight binding xTB calculations (Figure 4d). The central template is visible inside the ring in the STM images (Figure 4a). Images of 2 are similar to those of 1, but the molecules are slightly larger and more elliptical (circumference 15 ± 1 nm; long axis 5.5 ± 0.3 nm; short axis 4.3 ± 0.2 nm), and the template component is less clearly visible, which probably reflects the greater flexibility of 2 (Figure 4c). XPS spectra of samples of 1 and 2 on a gold surface confirmed the presence of zinc and nickel, and show a single nitrogen chemical environment, in agreement with the presence of metalated porphyrin species. Elemental composition ratios for Zn:Ni and N:Ni (2:1 and 6:1, respectively – Figure S56) agree with the expected stoichiometries of 1 and 2.

In conclusion, we have demonstrated an efficient synthesis of the first example of a partially fused, directly 5,15-linked porphyrin nanoring, without any linkers between the adjacent porphyrin units. The new spoked nanoring was characterized both in solution and on surface by STM. This work prepares the way for
the synthesis of similar covalently templated 5,15-PN nanorings using a template with cleavable links (e.g. esters), so that the template can be removed and the whole belt can be fused to give an edge-fused nanobelt EF-PN.

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Conflict of Interest

The authors declare no conflict of interest.

Data Availability Statement

The data that support the findings of this study are available in the supplementary material of this article.

Keywords: porphyrin • nanoring • template • aromatic compounds • oxidative coupling

Cobalt-catalyzed alkyne trimerization followed by oxidating coupling and fusion of zinc porphyrin units was used to construct a porphyrin nanoring around a central covalent spoke. The size and shape of these nanorings (diameter: 4.7 nm) was confirmed by scanning tunnelling microscopy (STM). A combination of zinc(II) and nickel(II) porphyrin units was used to increase the efficiency of the synthesis and control the positions of oxidative fusion.

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