Bottom-Up Hierarchical Self-Assembly of Chiral Porphyrins through Coordination and Hydrogen Bonds

Cristina Oliveras-González,† Florent Di Meo,‡ Arántzazu González-Campo,† David Beljonne,§ Patrick Norman,‡ Maite Simón-Sorbed,† Mathieu Linares,‡ and David B. Amabilino*†

†Institut de Ciència de Materials de Barcelona (ICMAB-CSIC), Campus Universitari de Bellaterra, 08193 Cerdanyola del Vallès, Catalonia, Spain
‡Department of Physics, Chemistry and Biology (IFM), Linköping University, SE-58 583 Linköping, Sweden
§Laboratory of Chemistry for Novel Materials, Mons University, Place du Parc, Mons B-9000, Belgium
∥School of Chemistry, The University of Nottingham, University Park, Nottingham NG7 2RD, United Kingdom

Supporting Information

ABSTRACT: A series of chiral synthetic compounds is reported that shows intricate but specific hierarchical assembly because of varying positions of coordination and hydrogen bonds. The evolution of the aggregates (followed by absorption spectroscopy and temperature-dependent circular dichroism studies in solution) reveal the influence of the proportion of stereogenic centers in the side groups connected to the chromophore ring in their optical activity and the important role of pyridyl groups in the self-assembly of these chiral macrocycles. The optical activity spans 2 orders of magnitude depending on composition and constitution. Two of the aggregates show very high optical activity even though the isolated chromophores barely give a circular dichroism signal. Molecular modeling of the aggregates, starting from the pyridine—zinc(II) porphyrin interaction and working up, and calculation of the circular dichroism signal confirm the origin of this optical activity as the chiral supramolecular organization of the molecules. The aggregates show a broad absorption range, between approximately 390 and 475 nm for the transitions associated with the Soret region alone, that spans wavelengths far more than the isolated chromophore. The supramolecular assemblies of the metalloporphyrins in solution were deposited onto highly oriented pyrolytic graphite in order to study their hierarchy in assembly by atomic force microscopy. Zero and one-dimensional aggregates were observed, and a clear dependence on deposition temperature was shown, indicating that the hierarchical assembly took place largely in solution. Moreover, scanning electron microscopy images of porphyrins and metalloporphyrins precipitated under out-of-equilibrium conditions showed the dependence of the number and position of chiral amide groups in the formation of a fibrillar nanomaterial. The combination of coordination and hydrogen bonding in the complicated assembly of these molecules—where there is a clear hierarchy for zinc(II)—pyridyl interaction followed by hydrogen-bonding between amide groups, and then van der Waals interactions—paves the way for the preparation of molecular materials with multiple chromophore environments.

INTRODUCTION

Porphyrins and their derivatives based on a tetrapyrrolic macrocycles linked by methine bridges are natural chromophores1 that are essential for the photosynthetic pathway2 in electron transport and light harvesting processes.3 This property is shown exemplarily by chlorophyll a4,5 or the chlorosomes of green-sulfur bacteria, which mainly comprises aggregated bacteriochlorophyll.6,7 These macrocycles have chiral environments arising from self-organization through noncovalent interactions. In all of these systems, a metal ion is coordinated in the core of the chromophore, and this feature plays an important role in their self-assembly and function. To understand the role of metal ions and self-organization of porphyrins, enzymes and light-harvesting systems mimicking the natural pigments have been synthesized and studied, confirming the importance of the supramolecular structure on the light harvesting ability.8–12

Self-assembled chiral systems based on synthetic porphyrins present a broad range of potential applications in areas such as photovoltaic cells,13 sensors,14 optoelectronics15 and nanoelectronic devices16 and also nonlinear optical materials.17 The number and position of the stereogenic centers in molecules united by covalent or noncovalent bonds directly affect the chiral transfer from the molecular level up to the supramolecular aggregates that they form in solution and consequently their optical activity can be modified.18–20

Received: August 28, 2015
Published: November 23, 2015

DOI: 10.1021/jacs.5b08081
J. Am. Chem. Soc. 2015, 137, 15795−15808
The self-assembly of porphyrins in supramolecular stacked systems can be controlled by a combination of noncovalent interactions such as hydrogen-bonding, van der Waals interactions and π−π-interactions between the porphyrin cores. The presence of exo-facing coordinating ligands and any metal ion coordinated at the center of the porphyrin ring also influence their self-assembly.21−23 The ability of metalloporphyrins to coordinate with axial ligands containing nitrogen, oxygen or sulfur atoms gives a great variety of self-assembled superstructures, with facility to tune their optical and electronic properties, as is the case of many zinc(II) metalloporphyrins that form stable supramolecules in nonpolar media when they self-assemble through coordinating ligands.24,25 The coordination of zinc porphyrins with axial nitrogen derivative ligands, such as imidazolyl or pyridyl groups afford the formation of well-organized macromolecules in the form of cyclic structures,26−29 or linear polymers30,31 and oligomers.32−34 The coordination between zinc(II) and pyridyl ligands has a relatively large association constant (∼10^3 M^−1)35,36 and usually does not disturb the photoexcited state of the metalloporphyrin.37,38

Zinc(II) porphyrins have a strong tendency to form pentacoordinate aggregates with a square pyramidal geometry, although the hexacoordination of this class of metalloporphyrins have also been reported in solid state structures.39−42 The control of the fifth coordination site of metalloporphyrins with nitrogen ligands is an important parameter in order to study the kinetic and thermodynamic effects in the self-assembly process of these compounds.43,44

Bearing in mind this wealth of former knowledge, compounds Zn-(R,R,R)-1, Zn-(R,R)-2, Zn-(R,R)-3 and Zn-(R)-4 have been designed and synthesized to probe hierarchical assembly in solution using different noncovalent interactions at each stage. Each compound differs from the other in the number and position of stereogenic centers—and associated hydrogen bond-forming unit—and pyridyl groups (Figure 1). Alkyl chains provide solubility for the systems in solution as well as adhesion to each other and nonpolar surfaces through van der Waals interactions,45 and pyridyl groups allow self-assembly through coordination with the zinc(II) ion in the core of porphyrinic rings.

For these compounds, two different coordinating atoms can in principle bind the metal ion, the pyridyl nitrogen atom or the carbonyl oxygen atom. Furthermore, hydrogen-bonding between amide groups can be the main noncovalent interactions to form an aggregate.18,21 However, more than one interaction can be participating in the self-assembly of these chromophores: Figure 2 shows the stronger potential interactions that need to be considered.

In this work the influence of the proportion and position of the 4-pyridyl and 4-phenyl rings bearing the stereogenic center in a lactamide moiety on the aggregation of the molecules in solution and upon precipitation was studied. In solution, absorption and circular dichroism (CD) spectroscopy—a chiroptical tool that allows precise monitoring of aggregation—gave information about the complex assembly and relative orientations of molecules in aggregates.27,31,46−48 In conjunction, simulations provided insight into the nature of the hierarchical assembly that led to the very high optical activity. The morphology of the aggregates was investigated by atomic force microscopy (AFM) when they were transferred from solution to surface and by scanning electron microscopy (SEM) to analyze their morphology in the solid state when precipitated far from equilibrium conditions.

■ RESULTS AND DISCUSSION

Synthesis and Characterization. The metalloporphyrin derivatives Zn-(R,R,R)-1, Zn-(R,R)-2, Zn-(R,R)-3 and Zn-(R)-4 were prepared following the synthetic route detailed in the
Supporting Information (Scheme S1). Briefly, the free base chiral porphyrin intermediates were prepared in one step—that required their separation by column chromatography—from the condensation of \((R)-\text{methyl 2-} (4\text{-formylphenoxy})\text{-propanoate})^{52} \text{and 4-formylpyridine with pyrrole in refluxing proponic acid in air. Habitual analytical and spectroscopic techniques were used to characterize each one. The constitutional isomers containing two stereogenic centers were distinguished by the multiplicity of the resonances of the hydrogen atoms of the pyrrole group in their nuclear magnetic resonance (NMR) spectra. In the case of \((R,R)-2 \text{ a double } \text{ doublet resonance (AB system) was observed. The NMR of the other isomer ((R,R)-3) showed an AB system with two singlets located between the doublets. The amimation of the ester group in these porphyrins at reflux in toluene with the presence of an excess of octadecylamine afforded the free base porphyrin derivatives (R,R)-1, (R,R)-2, (R,R)-3 and (R)-4. A decrease in solubility with increase number of pyridyl groups attached to the porphyrin made us consider a new route to obtain the porphyrin derivatives. The strategy to prepare the free base porphyrin derivatives \((R,R,R)-1, (R,R)-2, (R,R)-3 \text{ and } (R)-4 \text{ resolved the problem of low solubility of the intermediate reactant porphyrins was via an imine derived from the lactate. Condensation of the imine and 4-formylpyridine with pyrrole in refluxing proponic acid in air gave the desired free-base porphyrin derivatives directly, although yields were similar to the more conventional route. The insertion of zinc(II) into the porphyrin core was done by reaction of the acetate metal salt with free-base porphyrin derivative in a refluxing mixture of CH₂CH₂/CH₃OH to give the desired metallocompounds Zn-(R,R,R)-1, Zn-(R,R)-2, Zn-(R,R)-3 and Zn-(R)-4. The \(^1\)H NMR spectra of all the metalloporphyrins in CDCl₃ showed broad resonances at room temperature because of equilibria between aggregated states that occur at a rate comparable to the time scale of the spectroscopy. When the NMR was run at lower temperature for metalloporphyrin Zn-(R,R,R)-1 clear peaks were observed, and will be discussed in detail in the next section. IR and mass spectroscopy allow the identification of the desired compounds. The UV-visible absorption spectra of metalloporphyrins in CHCl₃ were the typical ones for these compounds, showing a Soret band at around 420 nm and two Q-bands at approximately 550 and 650 nm.

Self-Assembly of Porphyrins and Metalloporphyrins in Solution. The porphyrins and metalloporphyrins synthesized here were prepared to be able to self-assemble in solution by different noncovalent interactions such as hydrogen bonding, \(\pi-\pi\) stacking interactions directly related with the \(\pi\)-conjugated skeleton or by coordination with the metal ion in the core of porphyrin. \(^{50} \text{UV-visible absorption spectroscopy is a good technique to study the aggregation of the chromophores because the electronic } \pi-\pi^* \text{ transitions typical for these compounds are extremely sensitive to the aggregation state. The electronic absorption spectra depend on the exocyclic functionalization and the coordinated metal ion; they are well explained by the Gouterman four-orbital model.}^{51}

In general, free-base porphyrins and metalloporphyrins present an intense band (the Soret band) at around 420 nm with an extinction coefficient over \(10^5 \text{ mol/L cm}^{-1}\) was the focus of our studies. For some metalloporphyrins the Soret band can be shifted to lower wavelength because of the metal in the relative energy transitions of the chromophores ring. \(^{27} \text{Closed-shell ions (d0–d10) like zinc(II) have a weak effect on the } \pi-\pi^* \text{ gap, meaning that the Soret band does not move significantly with respect to the free-base porphyrin.}^{52–54} \text{ This band provides us useful information for the study of aggregation of our compounds. Two general types of stacked aggregates exist: } H \text{-aggregates, where porphyrins overlap face-to-face and present a blue (hypsochromic) shift respect to the Soret band of the monomer, and } J \text{-aggregates, where a red (bathochromic) shift of the Soret band is observed because of staircase-type packing with partial overlapping of the porphyrins.}^{55}

The absorption measurements of free-base porphyrins and the corresponding metalloporphyrins were initially carried out in chloroform at room temperature. The spectra of the free-base porphyrins showed a Soret band centered at 419 nm and for metalloporphyrins at 421 nm, indicating that in this solvent molecules behave as isolated chromophores surrounded by solvent. \(^{56–58} \text{ The absorption spectra were also recorded in methycyclohexane (with 3% chloroform to ensure solubility) for the same compounds and it was observed that the free-base porphyrins do not show any significant change in the position of the bands (Figure S1) indicating no aggregation of the chromophores and therefore insignificant hydrogen bonding under these conditions.}^{58} \text{ The absorption spectra of the metalloporphyrins show both slight blue-shifted (hypsochromic effect) Soret bands and new bands at longer wavelengths that might correspond to } J \text{-aggregates which are caused by the exciton coupling between porphyrins (Figure 3).}^{59} \text{ The spectral patterns are different for each metalloporphyrin indicating a distinct assembly for each, leading to unique structures. The spectra for compounds Zn-(R,R,R)-1 and Zn-(R,R)-3 show a shoulder at aaround 435 nm that has been assigned previously to the coordination of pyridyl groups with the zinc(II) ion of the neighboring metalloporphyrin.}^{60,61} \text{ The other two metalloporphyrins Zn-(R,R)-2 and Zn-(R)-4 show more red-shifted bands with a maxima at 450 nm, suggesting a higher number of electronic transitions due to formation of a larger aggregate or even another kind of } J \text{-aggregate because of the different position of the peripheral groups. The development of the equilibria in the self-assembly of each metalloporphyrin was studied at different temperatures by UV-visible absorption and CD spectroscopies, which gave information about the orientation that the chromophores adopt with respect to one another as the intensity of the optical

![Figure 3. UV-visible absorption spectrum of compounds Zn-(R,R,R)-1, Zn-(R,R)-2, Zn-(R,R)-3 and Zn-(R)-4 at room temperature in methycyclohexane/3% CHCl3 (5 μM).](https://zenodo.org/doi/10.1021/jacs.5b08081)
signals is directly proportional to the interactions between electronic transitions.\textsuperscript{62}

Solutions of compounds Zn-(1–4) in methycyclohexane/3% CHCl\textsubscript{3} were prepared (at a concentration 5 \(\mu\)M), and CD spectra were recorded in the range of temperatures from 338 to 263 K after a heating and cooling cycle to ensure that equilibrium conditions applied.

The temperature dependence of the CD spectra of compound Zn-(\(R, R, R\))-1 shows two clearly distinct states of self-assembly. A weak Cotton effect arises at the position of the band associated with the pyridyl-Zn(II) porphyrin complex, suggesting a perpendicular disposition of the two metalloporphyrins and a thermodynamically stable aggregate in the range of temperatures from 300.5 to 338 K (Figure 4, red lines). The isodichroic points at 415 and 443 nm in the CD spectra suggest equilibria between two species, either monomer and cyclic tetramer or monomer and linear aggregate. Furthermore, in this range of temperature, a clear isosbestic point at 425 nm appears in the absorption spectra, indicating a coexistence of monomer metallocorphyrin with an absorption maximum at 419.5 nm and the cyclic tetramer (the most stable aggregate in dilute solutions).\textsuperscript{35,63} The evolution of the aggregate suggesting equilibria between the discrete aggregate and the larger superstructure.

The evolution of the tetramer aggregate with temperature is shown in the plots of band intensity versus temperature (Figure 5). The bands at 438 nm in the CD spectra and 431.5 nm in the corresponding absorption spectra reach a maximum at around 300 K and then an abrupt decay of these bands is apparent, while the 451 nm band starts to increase below this critical temperature. This behavior suggests that the cyclic tetramer aggregates to a larger superstructure at lower temperatures, as corroborated by the presence of an isosbestic point at 441 nm.\textsuperscript{66}

For compound Zn-(\(R, R, R\))-2 the evolution from monomer to large aggregate does not pass through a specific intermediate. The formation of a discrete structure is not evident when the aggregation is followed by CD or absorption spectroscopy (Figure 6).\textsuperscript{67} Two intense nonsymmetrical bisignate CD signals were observed, one with positive maximum at 421 nm and the other one corresponding to a supramolecular arrangement at longer wavelength with maximum at 449 nm. The CD spectra and corresponding absorption spectra of Zn-(\(R, R, R\))-2 were simpler than those of metalloporphyrin Zn-(\(R, R, R, R\))-1 presumably because of the higher symmetry in the former compound. The presence of two pyridyl groups in Zn-(\(R, R, R\))-2 is apparently an important factor for linear growth in this compound, which must involve cooperative assembly in which both pyridyl-zinc(II) coordination and hydrogen bonding, as the CD pattern for the aggregate is remarkably similar to the larger aggregates observed for Zn-(\(R, R, R, R\))-1, pointing to a similar chiral arrangement of the porphyrin moieties in the aggregate.

Three isodichroic points are evident in the CD spectra, at approximately 415 nm, 435 and 453 nm. The isodichroic points at 415 and 443 nm in the CD spectra suggest equilibria between two species, either monomer and cyclic tetramer or monomer and linear aggregate. Furthermore, in this range of temperature, a clear isosbestic point at 425 nm appears in the absorption spectra, indicating a coexistence of monomer metallocorphyrin with an absorption maximum at 419.5 nm and the cyclic tetramer (the most stable aggregate in dilute solutions).\textsuperscript{35,63} The evolution of the aggregate suggests equilibria between the discrete aggregate and the larger superstructure.

The evolution of the tetramer aggregate with temperature is shown in the plots of band intensity versus temperature (Figure 5). The bands at 438 nm in the CD spectra and 431.5 nm in the corresponding absorption spectra reach a maximum at around 300 K and then an abrupt decay of these bands is apparent, while the 451 nm band starts to increase below this critical temperature. This behavior suggests that the cyclic tetramer aggregates to a larger superstructure at lower temperatures, as corroborated by the presence of an isosbestic point at 441 nm.\textsuperscript{66}

For compound Zn-(\(R, R, R\))-2 the evolution from monomer to large aggregate does not pass through a specific intermediate. The formation of a discrete structure is not evident when the aggregation is followed by CD or absorption spectroscopy (Figure 6).\textsuperscript{67} Two intense nonsymmetrical bisignate CD signals were observed, one with positive maximum at 421 nm and the other one corresponding to a supramolecular arrangement at longer wavelength with maximum at 449 nm. The CD spectra and corresponding absorption spectra of Zn-(\(R, R, R\))-2 were simpler than those of metalloporphyrin Zn-(\(R, R, R, R\))-1 presumably because of the higher symmetry in the former compound. The presence of two pyridyl groups in Zn-(\(R, R, R\))-2 is apparently an important factor for linear growth in this compound, which must involve cooperative assembly in which both pyridyl-zinc(II) coordination and hydrogen bonding, as the CD pattern for the aggregate is remarkably similar to the larger aggregates observed for Zn-(\(R, R, R, R\))-1, pointing to a similar chiral arrangement of the porphyrin moieties in the aggregate.

Three isodichroic points are evident in the CD spectra, at approximately 415 nm, 435 and 453 nm. The isodichroic points at 415 and 443 nm in the CD spectra suggest equilibria between two species, either monomer and cyclic tetramer or monomer and linear aggregate. Furthermore, in this range of temperature, a clear isosbestic point at 425 nm appears in the absorption spectra, indicating a coexistence of monomer metallocorphyrin with an absorption maximum at 419.5 nm and the cyclic tetramer (the most stable aggregate in dilute solutions).\textsuperscript{35,63} The evolution of the aggregate suggests equilibria between the discrete aggregate and the larger superstructure.

The evolution of the tetramer aggregate with temperature is shown in the plots of band intensity versus temperature (Figure 5). The bands at 438 nm in the CD spectra and 431.5 nm in the corresponding absorption spectra reach a maximum at around 300 K and then an abrupt decay of these bands is apparent, while the 451 nm band starts to increase below this critical temperature. This behavior suggests that the cyclic tetramer aggregates to a larger superstructure at lower temperatures, as corroborated by the presence of an isosbestic point at 441 nm.\textsuperscript{66}
different species are in equilibrium in the temperature range
studied. The absence of a band at 435 nm in the absorption
spectra could indicate that the species involved in the
equilibrium are the monomer metallocompound and a larger
aggregate. An isosbestic point at 437 nm is present in the
absorption spectra corroborating the existence of an equili-
brum between monomer and aggregate. The remarkable point
of this spectrum is the presence of two bands at 448 and 453
nm, clearly observable at and above room temperature. As the
temperature is decreased these bands collapse into one. A
possible explanation of this behavior is the formation of a J-
aggregate whose excited state energies become closer as it gets
longer resulting in a band with a shoulder. Even at the higher
temperatures studied (up to 338 K), a supramolecular aggregate is present in solution as indicated by the high intensity in the CD spectra. This feature shows that metalloporphyrin Zn-(R,R)-2 presents a much greater tendency to aggregate than Zn-(R,R,R)-1, although the CD spectra infer a somewhat similar arrangement of the chromophores within the aggregates. Therefore, it is believed that the tetramer formed by the pyridyl coordination giving pentacoordinate zinc(II) ions is present in the aggregates formed by Zn-(R,R)-2.

The behavior in solution of compound Zn-(R,R)-3 was completely different to either compound Zn-(R,R,R)-1 or Zn-(R,R)-2. The difference with compound Zn-(R,R)-2 is particularly striking, because both have two chiral amide groups and two pyridyl substituents in the periphery of the metalloporphyrin core. This contrasting behavior indicates the importance of the position of the groups on the porphyrin ring for self-assembly. The CD spectra of Zn-(R,R)-3 (Figure 7) show two bisignate signals of low intensity. This pattern is similar to that of compound Zn-(R,R,R)-1 above 300 K when it is present as a cyclic tetramer. Therefore, this feature indicates that for Zn-(R,R)-3 the main aggregate form in the whole range of temperature studied is the cyclic tetramer and there is no subsequent aggregation through secondary interactions between tetramers. In the corresponding absorption spectra only a broad band at 435 nm is seen, similar to that of Zn-(R,R,R)-1 and attributed to a discrete aggregate. Moreover, the broadness of the band might corroborate the formation of a cyclic aggregate.

Compound Zn-(R)-4 with only one stereogenic center in its structure and three pyridyl groups showed contrasting spectra compared to the other compounds, but principally because the solubility of the compound is so low in the solvent used. Nonetheless, the appearance of the CD spectrum reflects an environment of the molecules (Figure S2) that is different to the homologues.

The results of the temperature-dependent studies of the metalloporphyrins clearly demonstrate the importance of the position and number of pyridyl groups and chiral amide in the periphery of the macrocycle. In the case of metalloporphyrin Zn-(R,R,R)-1 two distinct assembly regimes arise. Above 300 K a tetramer structure is formed and an elongation process takes place at lower temperatures. A linear growth for metalloporphyrins Zn-(R,R)-2 and Zn-(R)-4 appears likely according to the changes in the CD and absorption spectra with temperature. On the other hand, the results for Zn-(R,R)-3 suggest the cyclization of the aggregate, but the lack of effective hydrogen bonding negates further aggregation.

Mathematical models for the aggregation curves from CD intensity at distinct wavelengths versus temperature were applied in order to determine precisely the self-assembly processes taking place in each Zn (II) porphyrin. Two main models are associated with the self-assembly processes in this kind of aggregation. The cooperative mechanism has two regimes, the (costly) formation of nuclei with critical size which
acts as an activation step followed by a favorable elongation process. On the other hand, the isodesmic model is governed by a gradual increase in the number of aggregate species with the same energy increment at each addition of monomer. To analyze the temperature-dependent data from CD spectra of each metalloporphyrin, both models were applied and the best fits identified the most suitable model in each case (Figure S3–S4).

The self-assembly of Zn-(R,R,R)-1 clearly shows two distinct states in its assembly in the CD and absorption spectra, suggesting that a nucleation–growth process take place (Figure 8). Fitting reproduced very well the shape of the observed curves. The formation of a tetramer aggregate follows a sigmoidal shape as a function of temperature which is indicative of isodesmic self-assembly and acts as nucleation structure. The following elongation regime appears as the temperature decreases below 300 K; the nuclei evolve to a larger aggregate following a cooperative process.

Given the restrictions in solvent and concentration a direct comparison between CD and NMR data was not possible. However, a concentrated solution (20 mM) in a more competitive solvent was used. The comparison of free-base porphyrin (R,R,R)-1 1H NMR spectrum and at that of Zn-(R,R,R)-1 in chloroform-d at -45 °C (where it was possible to distinguish clearly all the peaks of the molecule) shows a clear shift of the pyridyl protons to high field. In the Zn(II) compound they resonate at 6.32 and 6.07 ppm for the H β-pyridyl and 2.58 and 1.97 ppm for the Hα-pyridyl, while for the free-base porphyrin they appear as a doublets at 8.88 and 8.17 ppm hidden by the pyrrole peak and the aromatic peak, respectively. A splitting of the signals is observed in the Zn(II) compound, mostly the aromatic protons and pyridyl protons close to the porphyrin ring because the two faces of porphyrin are not equivalent due to the pyridyl-zinc(II) ion coordination (Figure 9).73–76

The complexity of the 1H NMR spectrum of metalloporphyrin Zn-(R,R,R)-1 required 2D 1H NMR studies in order to identify correctly all the protons of the molecule. A ROESY (Rotating Frame Overhauser effect spectroscopy) experiment allowed full assignment of the signals and indicated those protons in close proximity (Figure S8). Figure 10 shows the proposed tetramer structure for Zn-(R,R,R)-1 with arrows indicating the correlation between protons extracted from the ROESY studies.

Modeling of the Metalloporphyrin and Pyridine-Metalloporphyrin Systems. Theoretical models of a Zn-
(R,R,R)-1 and its pyridine complex were constructed by replacement of octadecyl with ethyl chains since alkyl side chains are not relevant to the Soret band of ECD spectra. First-principles quantum mechanical calculations on the models were carried out at the ωB97X-D/6-31+G(d,p)/LANL2DZ/ωB97X-D/6-31G(d)/LANL2DZ level of theory. The free and complexed metallorporphyrin have planar tetracoordinate and square-pyramidal pentacoordinate geometry, respectively. Both singlet and triplet states were considered for both systems. Singlet states are significantly more stable than triplets, with singlet–triplet splittings being −30.9 and −26.1 kcal mol⁻¹ for the free metallorporphyrin and the pyridine complex, respectively. Pyridine complexation is highly favorable, showing ΔE_{association} = −20.1 kcal mol⁻¹.

Pyridine complexation to the metallorporphyrin leads to slight conformational changes of the tetrapyrole core. Bond distances between the metal ion center and the tetrapyrole nitrogen-atoms are slightly increased owing to the presence of the new electron-donor pyridine ligand. For instance, typical N_{tetrapyrrole}–Zn bond distances are respectively 2.06 and 2.09 Å in the metallorporphyrin and its complex (Table S1), and the zinc atom is moved out of the tetrapyrole core plane as seen experimentally in this type of complex.68,79 Passing from planar tetracoordinate to square pyramidal pentacoordinate Zn-(II) also leads to a redistribution of charges. The pyridine ligand plays a significant electron-donor role, decreasing the positive charge of the zinc atom from 1.185e for the free metalloporphyrin to 1.038e for the pyridine complex, as well as increasing charges of pyridine and tetrapyrole N atoms (Table S2). An important electron exchange feature is observed in the pyridine-zinc bond suggesting a strong dative nature. The π-back-donation between 3d orbitals of Zn-atom and π-orbital of pyridine is not important (verified by analyzing bond distances of the pyridine ligand that are not affected with respect to the standalone pyridine). The π-back-donation is thus restricted to the Zn-tetrapyrole core. Thereby, electrostatic interactions between pyridine and zinc are expected to play an important role in such association. The nitrogen-atom charge of standalone pyridine is very similar to that of the moiety in Zn-(R,R,R)-1, confirming the relevance of this model for the Zn-pentacoordinate complexation.

The highest occupied and the lowest unoccupied molecular orbitals (HOMO and LUMO) are destabilized by approximately 0.3 and 0.2 eV upon pyridine complexation leading to a smaller HOMO–LUMO gap by around 0.1 eV. HOMO and LUMO distributions are rather similar between the free metallorporphyrin and the pyridine complex confirming the experimentally suggested π–π* gap (Figure S15). Only a very weak change is expected in energy transitions of the tetrapyrole chromophore while bound to pyridine (i.e., in the Soret band). Such complexation would lead to a red-shift of the high energy band as observed experimentally.

Self-Assembly Leading to Proposed Tetramer Structure. NPT molecular dynamics (MD) simulations were performed at room temperature to establish the self-assembly of the ethyl chain equivalent of Zn-(R,R,R)-1, starting with the proposed tetramer structure to test its existence and stability. For sake of clarity, each sequential metallorporphyrin (i.e., monomer) of a tetramer square are numbered 1, 2, 3, and 4 and 1 → 2, 2 → 3, 3 → 4 and 4 → 1 are metallorporphyrin pairs connected by N_{pyrrole}–Zn bonds. The tetramer is actually stable throughout the simulation and pentacoordinate mode is maintained with an average N_{pyrrole}–Zn bond distance of 2.35 Å as well as an average N_{pyrrole}–Zn–N_{pyrrole} angle of approximately 91°. The zinc(II) atom is slightly more in the plane than in the quantum mechanical model, although this very slight bond length difference is not expected to influence dramatically the superstructure, especially given the flexibility in the bonding scenario. The N_{pyrrole}–Zn bond distances are shorter than N_{pyrrole}–Zn ones leading to a certain flexibility between metalloporphyrins (N_{Zn}–Zn_{N} bond distances and N_{pyrrole}–Zn–N_{pyrrole} angles ranges are respectively [2.1, 2.6] Å and [75, 100]°, Figure 11a and b). The meso-substituted pyridyl moieties are twisted to the tetrapyrole core with either right or left chirality (Figure 11c). No preference between right- and left-handed deviations of the pyridyl dihedral angle is observed. The deformations of zinc atoms out of the tetrapyrole cores obtained from MD simulations are slightly smaller than those observed in the QM calculations, possibly because of the deformation of the tetrapyrole core in the tetramer. Indeed, tetrapyrole cores tend to lose their planarity in the tetramer. For example, improper dihedral angles formed by each N atom of tetrapyrole cores exhibits distribution from −10° to 10°.

The global tetramer structure can be defined by the square formed by each zinc-atom, with edge Zn–Zn distances of around 9.7 Å. A square shape is confirmed by the observation of diagonal Zn–Zn distances to be 13.7 Å, which is in perfect agreement with the square diagonal equation. The dot products between vectors defined from Zn to N_{pyrrole} atoms were also calculated (Table S3), and they show that adjacent metalloporphyrins remain orthogonal with respect to each other along the MD trajectory whereas diagonal metalloporphyrins keep an antiparallel orientation.

In spite of the importance of the Zn–N_{pyrrole} bonds in the tetramer, other noncovalent interactions also play a crucial role in these aggregates. The total noncovalent interaction energy in a tetramer was calculated to be −94.7 kcal mol⁻¹. This energy can be decomposed into noncovalent dimer energy between each metalloporphyrin. The main contribution to noncovalent interactions in the tetramer arises from adjacent metal-
loporphyrin dimers (i.e., $1 \rightarrow 2$, $2 \rightarrow 3$, $3 \rightarrow 4$ and $4 \rightarrow 1$). There is an equal contribution of van der Waals and Coulombic interactions (average Coulombic and van der Waals dimer energy interactions are 11.7 and 11.9 kcal mol$^{-1}$ respectively). It is worth noting that electrostatic interactions are less dominant probably due to the solvation in the very nonpolar solvent. The perpendicular T-shape arrangement of adjacent metalloporphyrins underlines the importance of dispersive $\sigma-\pi$ interactions between the pyridyl groups and the tetrapyrrole cores. Cofacial antiparallel metalloporphyrins (i.e., $1 \rightarrow 3$ and $2 \rightarrow 4$) are weakly interacting since the respective intermolecular distances are at the upper limit of such interactions. Average Coulombic and van der Waals interaction energies are $-0.3$ and $-1.0$ kcal mol$^{-1}$, respectively.

**Self-Assembly of the Tetramer Structures.** The key parameter in the calculation of the elongation step is the elucidation of the association mode between each tetramer. Since, the process takes place at temperatures below 300 K experimentally, weak noncovalent interactions are the driving force. Amide groups in the metalloporphyrin side chains are expected to play a crucial role as hydrogen-bonding between tetramer nuclei. Likewise, $\pi$-delocalization over the tetrapyrrole cores could lead to $\pi-\pi$ interactions in tetramer aggregates. Any association mode between tetramer aggregates must balance these interactions. In theory, three binding modes can be proposed (Figure 12).

![Figure 12. Proposed self-assembly modes of the tetramer metalloporphyrin complex. Tetramer square nuclei are depicted from the top view (porphyrins perpendicular to the plane of the drawing).](Image)

In the top view, two types of side chain can be defined in each tetramer, lateral and top/under side chains are respectively coplanar and orthogonal to the drawing plane.

Mode A may be defined by an almost face-to-face arrangement of tetramers that have hydrogen bonds between lateral side chains. Mode B and C slide one tetramer upon another allowing hydrogen-bonding between lateral and top-side chains. It must be stressed that mode B allows two hydrogen bonds while mode C allows four hydrogen bonds to be made. In the latter, one side chain of one tetramer is between two top side chains of another allowing the central amide group to be bonded to one electron-acceptor and one electron-donor respectively, thanks to NH and C=O moieties, thus satisfying all its hydrogen-bonding potential.

Particular attention has been paid to mode A since several submodes of this arrangement are possible according to the position of the side chains involved in hydrogen-bond networks (Figure 13). Ladder and pyramidal shape are defined when both side chains of one tetramer are both on the top (bottom) of the side chains of the other tetramer square. They differ only when at least three tetramers are self-assembled. In the latter, the middle tetramer side chains are below both neighbor ones while the former exhibits a kind of staircase shape. On the other hand, helical structures may be obtained when one side chain is on the top of the other tetramer side chain while the other side chain of the former tetramer is under. Both $P$- and $M$-helices were considered according to the tetrapyrrole planes rotation along helical axis (Figure 13).

Dodecamer structures (trimers of tetramer squares) were built following the several modes of association and were minimized at the molecular mechanics (MM) level of theory. Again, the ethyl chain equivalent of Zn-(R,R,R)-1 was used to optimize computational time in these very large and complex superstructures, and therefore allow a reasonable exploration of the supramolecular possibilities in the structure. The binding modes considered as the strongest here—through metal coordination and hydrogen bonding—are not apparently influenced significantly by the weak interactions between alkyl side chains, which are located at the periphery of the self-assembled structure. However, this simplification should not be taken as a general result since in other systems lateral alkyl chains might play a crucial role in the self-assembly process, and might influence in the subsequent hierarchical levels of assembly. The use of energies to select the most stable conformation is challenging because of (i) the small difference in relative energies per molecule ranging from 0.0 to 13.0 kcal mol$^{-1}$ and (ii) the considered level of theory. However, such results associated with theoretical and experimental CD spectra permit trends about self-assembly modes to be gleaned. Mode C seems to be unlikely to occur owing to steric hindrance of alkyl side chains leading to the least stable structure. Interestingly, the pyramidal mode A may also be discarded while ladder mode A is one of the most plausible at first sight. Finally, both $P$- and $M$-helices also appear possible, the latter being more stable in vacuo.

**Toward the Elucidation of Second-Order Self-Assembly Using Semiempirical CD Calculations.** Experimental CD in the Soret region below 300 K is mainly characterized by very high optical activity with respect to higher temperature spectra. We stress that the CD signals arises from induced chirality of tetrapyrrole cores by self-assembly (and not from intrinsic molecular chirality). As suggested experimentally, higher temperature CD signals should be considered as tetramers. The low intensity CD signals within a range from $-5$ to $5$ mdeg suggests that the chiral deformation of the tetrapyrrole core is significant but small. Furthermore, the theoretical CD spectra of the monomer corroborate that even though the metalloporphyrin is chiral, it does not present a significant optical activity (Figure 14). This low signal can be attributed to the pyridyl moiety binding to the zinc(II) metal ion. It must be stressed that a similar environment-induced chirality has already been observed in metalloporphyrins with, e.g., DNA G-quadruplexes. However, the CD signal at low temperature shows that the tetrapyrrole core chirality is strongly affected by hierarchical assembly. This might be attributed to large chiral excitonic coupling between two tetrapyrrole cores that belong to different tetramer structures. To be effective, exciton coupling requires a relatively short distance between tetrapyrrole cores. Therefore, mode B and mode C should be discarded as in this self-assembly mode the tetrapyrrole cores are neither facing each other nor close (average distance being ca. 18 Å).
Ladder and helix arrangements appear to be the only relevant modes for the tetramer aggregation because they exhibit cofacial arrangements of the tetrapyrrole cores and short intercore distances that would lead to efficient exciton coupling. Moreover, these association modes are the only ones comprising both hydrogen-bonding and π−π interactions between tetramers. As mode B and C have been discarded, semiempirical excitonic calculations of CD spectra were carried out only on the four assemblies of mode A (Figure 14). Despite the limited potential energy surface exploration available, the high optical activity is reproduced only with helical shapes. Intertetramer cofacial tetrapyrrole cores are twisted with respect to each other leading to a strong chiral exciton coupling. According to clockwise or anticlockwise twist, P- and M-helix CD spectra are mirror images. It must be stressed that the high optical activity is not observed in the ladder arrangement since excitonic metalloporphyrins are not twisted. That should thus discard ladder shape as a relevant association mode.

Self-Assembly Studies of Porphyrins and Their Metalloporphyrins on a Surface. So as to probe the hierarchically grown structure of the aggregates of the porphyrins more microscopically, solutions used for spectroscopic studies were deposited on highly oriented pyrolytic graphite (HOPG) by casting and the solvent was allowed to evaporate. The material was investigated with intermittent mode atomic force microscopy (AFM). As was observed in solution, the self-assembly was dependent on temperature, so the organization of the supramolecular aggregate of metallocompounds on the surface can also be affected by this external factor. Molecule–surface interactions are an important factor to take into account to successfully adsorb the aggregates from solution onto a solid support in a controlled way. The metallocompounds under observation here have a long alkyl chains in the side positions that can help the adsorption of the aggregates due to the van der Waals interaction with the hydrophobic surface. Metalloporphyrin solutions (used in CD) were drop cast onto HOPG surface at room temperature and let the solvent evaporate. Topographic images of Zn-(R,R,R)-1 showed disc-like objects distributed evenly over the surface after evaporation of the solvent. The measured heights of the discs were around 2.5 nm, that must correspond to a standing up porphyrins (with their planes perpendicular to the surface), otherwise a coordinating pyridyl group of aporphyrin flat on the surface could not interact with another to produce the cycle for steric reasons (Figure S16). The ill-defined shape is presumably because the alkyl chains atop the structure are free to slump in no preferred direction on or beside the aggregate.

The globular stacks present on surface presumably come directly from the solution, because the molecule–molecule interaction is apparently stronger than the molecule–surface forces. These results were in agreement with the observation in solution where tetrameric structures were formed from 300 K upward for this compound.

Figure 15 shows a statistic study about the dimensions of the globular objects from a solution of metalloporphyrin Zn-(R,R,R)-1 where it can be seen that most of the particles present diameter between 20 to 30 nm (from 100 particles).
This behavior could be attributed to different numbers of tetramers close to one another because of the hydrogen-bonding between amide groups of different tetramer aggregates that lead to larger structures at lower temperature.

In order to corroborate that the aggregates formed in solution were transferred to the surface suffering a minimum effect of the adsorption, the metalloporphyrin solution was cooled down to 0 °C and then drop-cast onto HOPG. Two different regions of the surface were observed by AFM with different behavior of the aggregates (Figure 16). The image in Figure 16a shows big planar blocks with a height around 12 nm (see profile) and small rods of 3 nm height around them. The blocks have well-defined edges characteristic of very small crystals (the biggest blocks are around 700 nm long). Figure 16b shows a region with a fibrillar organization with vertical dimensions between 1.5 to 2 nm, that could correspond to the porphyrin size width. This behavior coincides with the evolution of the aggregates in solution because, even though the fibers are not well-defined, they are clearly distinct from the discs seen for the same sample when deposited at room temperature.

Topographic images of a solution of Zn-(R,R,R)-1 deposited at room temperature onto HOPG showed a mixture of spherical particles with heights around 10 nm and a fibrillar network with heights around 3 nm. When the same solution was cooled down at 0 °C and cast onto graphite (HOPG) a fiber network was observed as the dominant self-assembled structure. The profile showed a 3 nm height for the fibers and a 1.5 nm height for an underlying structure possibly arising from a non-aggregated metalloporphyrin, always present in solution, that physisorbs to the HOPG on evaporation and aligns with its axes (Figure S17). A close up image implied helicity in the fiber formation, clearly observable in the corresponding phase image (Figure 17). This helicity indicates a clear chiral transfer from monomer to aggregate that was observed in the high intensity CD signal.

While the AFM images of assemblies of Zn-(R,R)-2 deposited solutions at low temperature matched well with the observation of long aggregates in circular dichroism, the room temperature solution transferred onto HOPG did not show significant large aggregates. This difference between solution and surface studies could be noted for the lability of the tetramer that evolves toward the fibrillar structure. As mentioned before, the transfer of aggregates from solution to surface can be affected by different factors such as molecule—
surface interaction and the solvent used. The faster evaporation of the solvent at room temperature might favor the observation of the discrete aggregate that was not possible to detected by spectroscopic techniques.

The statistical study of the room temperature deposited Zn-\((R,R)\)-2 in one area where round particles predominates as aggregate showed a narrow dimensions distribution of the particles presenting diameters between 14 to 26 nm as a result of interactions between different tetramers aggregates (Figure S18). The objects are smaller than those formed by the monopyridyl homologue, possibly because of mass depletion to the fibers that are also present on the surface.

Topographic images of Zn-\((R,R)\)-3 showed a globular structure as a main morphology on graphite, with heights between 3 and 6 nm when the solution were adsorbed at room temperature, while a bigger globular structures where observed when it was deposited at 0 °C (Figure S19). Statistical analyses of the dimensions of the small particles in the AFM images at room temperature for metalloporphyrin Zn-\((R,R)\)-3 showed that the major number of the small globular stacks have a diameter between 14 to 20 nm, although they have a great tendency to form a bigger aggregates as the topographic images showed (Figure S20). Therefore, little specificity in assembly is observed, and no significant fibrillar morphology was seen.

Metalloporphyrin Zn-\((R)\)-4 showed an apparent monolayer organization on the surface aligned with the substrate symmetry axes and with a height lower than 0.5 nm in their topographic images, corresponding to flat-adsorbed molecules. Moreover at room temperature spheres were appreciated that presented height of 2.5 and 1 nm (Figure S21). The dimensions for the spherical structure of Zn-\((R)\)-4 when the solution was drop-cast onto graphite surface at room temperature were around 14 to 20 nm in diameter (Figure S22).

At a yet larger scale, the solid morphology of free-base porphyrins and the corresponding metalloporphyrins were analyzed by scanning electron microscopy (SEM) after precipitation in far from equilibrium conditions from CHCl₃/EtOH solution (Figure 18).

As observed in the solution studies, free-base porphyrins \((R,R,R)\)-1, \((R,R)\)-2, \((R,R)\)-3, \((R)\)-4 did not show any kind of aggregate, but under precipitation the same porphyrins showed a kind of fibrillar organization.

Porphyrins \((R,R,R)\)-1 and \((R,R)\)-2 revealed a much clearer fibrillar organization in their SEM images (Figure 18a and b) than porphyrins \((R,R)\)-3 and \((R)\)-4 (Figure 18c and d) where a more compact solids were observed. Especially for porphyrin \((R)\)-4 the lack of alkyl chains might be responsible for this effect. The main force for the arrangement of free-base porphyrins is the hydrogen-bonding between amide groups. However, porphyrins \((R,R)\)-2 and \((R,R)\)-3, both with two chiral amide groups and two pyridyl groups connected in the meso position of the ring, showed a different morphology that is an
indicative that the constitution of the chromophore ring related with the position of the groups has a direct effect in the way of precipitation, and this is not only attributed to the aggregation in solution.

The solid hierarchy organization for metalloporphyrins observed in the SEM images revealed how Zn-(R,R,R)-1 and Zn-(R,R,R)-2 precipitated in a fibrillar manner (Figure 18a and b), whereas for Zn-(R,R)-3 and Zn-(R)-4 a more chaotic solid was observed after precipitation (Figure 18c and d). Both Zn-(R,R,R)-1 and Zn-(R,R,R)-2 show a thicker fibrillar structure compared with the corresponding free base porphyrins, showing how the hierarchy of organization is propagated from the smallest unit and strongest interaction (the zinc(II)-pyridyl) up to the solid where van der Waals forces must dominate.

CONCLUSIONS

It has been shown that a family of chiral zinc(II) porphyrins can exhibit hierarchical self-assembly with a complicated combination of supramolecular environments that give rise to high optical activity over a wide range of wavelength—from 390 to 475 nm—from a single chromophoric unit. The zinc(II) interaction with pyridyl moieties is the dominant interaction, but hydrox bonding and van der Waals interactions are responsible for the massive increase in optical activity that is observed.

Molecular modeling has confirmed the hierarchy in the self-assembly of the compound bearing one pyridyl group and that weaker interactions between side-chains is a viable way of inducing high optical activity in the second step of the self-assembly of this compound. The second step is well differentiated from the first because of the difference in binding energy of the hydrogen bonds compared with the zinc(II) to pyridyl coordination bond. The optical activity of this compound was important for the interpretation of the other self-assembled structures, particularly for Zn-(R,R)-2.

The compound Zn-(R,R)-2 incorporating two pyridyl groups at the 12 and 6 o’clock positions of the porphyrin core is the system that aggregates most strongly. The optical activity of the aggregate indicates that interporphyrin stacking through weak interactions is very important as it is in the compound with one pyridyl group. However, the concerted nature of the assembly involving coordination and hydrogen bonds in this case suggests an arrangement where the pyridyl group not bonded to zinc(II) can interact with the amide function of a neighboring tetramer. Therefore, the assembly of these compounds is extremely sensitive to molecular constitution, as demonstrated by the other zinc porphyrin compounds with either two pyridyl groups at 12 and 3 o’clock or three pyridyl groups, where only small aggregates are observed with no obvious specific assembly pattern.

Importantly, the deposition of these aggregates onto graphite reflects the state of assembly in solution. When the compound incorporating one pyridyl ring was deposited at room temperature only small closed aggregates were observed, while a lower temperature deposition from solution larger scale fibrous structures were seen by AFM.

The effect of constitution on assembly in solution is carried through to higher-level aggregates formed under far from equilibrium conditions. The morphology of the materials shows fibers and lamellae with features that are unique in each case, and very different from those of the corresponding free base porphyrins. This result emphasizes the potential of hierarchical assembly approaches not only because there exist strong interactions facilitate aggregation, but also since the influence of weaker interactions can be enhanced by them.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/jacs.5b08081.

Full synthetic experimental procedures and characterization of compounds, additional absorption and CD spectra and titration curves, 2D NMR spectrum, IR spectra of all the porphyrins, computational details, additional AFM images and analysis. (PDF)

Dodecamer XYZ coordinates. (ZIP)

AUTHOR INFORMATION

Corresponding Author

david.amabilino@nottingham.ac.uk

Notes

The authors declare no competing financial interest.

ACKNOWLEDGMENTS

This work was supported by MINECO (Spain, Project CTQ2010-16339 and MAT2013-47869-C4). M.L. acknowledges SeRC (Swedish e-Science Research Center) for funding. P.N. and F.D.M. thank the Swedish Research Council (Grant No. 621-2014-4646). Authors P.N., F.D.M. and M.L. acknowledge SNIC (Swedish National Infrastructure for Computing) for providing computer resources (snic001-12-192). D.B. is a FNRS Research Director.

REFERENCES

(1) Babu, S. S.; Bonifazi, D. ChemPlusChem 2014, 79, 895–906.
(2) Renger, T.; Schlodder, E. ChemPhysChem 2010, 11, 1141–1153.
(3) Ohtani, Y.; Shimada, T.; Takagi, S. J. Phys. Chem. C 2015, 119, 18896–18902.
(4) Lo, P.-C.; Leng, X.; Ng, D. K. P. Coord. Chem. Rev. 2007, 251, 2334–2353.
(5) Hu, J.-S.; Guo, Y.-G.; Liang, H.-P.; Wan, L.-J.; Jiang, L. J. Am. Chem. Soc. 2005, 127, 17090–17095.
(6) Nishimori, R.; Tamiaki, H.; Kashiwara, S.; Saga, Y. Supramol. Chem. 2015, 27, 28–36.
(7) Van Rossum, V.-J.; Steensgaard, D. B.; Mulder, F. M.; Boender, G. J.; Schaffner, K.; Holzwarth, A. R.; de Groot, H. J. M. Biochemistry 2001, 40, 1587–1595.
(8) Taamaki, H.; Ariki, N.; Yasuda, S.; Miyatake, T.; Oba, T. Tetrahedron 2014, 70, 9768–9775.
(9) Balaban, T. S.; Linke-Schaelzel, M.; Bhise, A. D.; Vanthuyne, N.; Roussel, C.; Anson, C. E.; Bath, G.; Eichöfer, A.; Foster, K.; Garab, G.; Gliemann, H.; Goddard, R.; Javorfi, T.; Powell, A. K.; Rösner, H.; Schimmel, T. Chem. - Eur. J. 2005, 11, 2267–2275.
(10) Balaban, T. S.; Bhise, A. D.; Bringmann, G.; Bürgk, J.; Chappaz-Gillot, C.; Eichöfer, A.; Fenske, D.; Götz, D. C. G.; Kräuter, A.; Miroguchi, T.; Mössinger, D.; Rösner, H.; Roussel, C.; Schraut, M.; Tamiaki, H.; Vanthuyne, N. J. Am. Chem. Soc. 2009, 131, 14480–14492.
(11) Cnatonetti, V.; Monti, D.; Venanzi, M.; Bombelli, C.; Ceccacci, F.; Mancini, G. Tetrahedron: Asymmetry 2004, 15, 1969–1977.
(12) Jin, H. G.; Balaban, M. C.; Chevalier-Michaud, S.; Rigazzea, M.; Balaban, T. S. Chem. Commun. 2015, 51, 11884–11887.
(13) Li, L.-L.; Diao, E.W.-G. Chem. Soc. Rev. 2013, 42, 291–304.
(14) Ding, Y. B.; Tang, Y. Y.; Zhu, W. H.; Xie, Y. S. Chem. Soc. Rev. 2015, 44, 1101–1112.
(82) Rubio-Magnieto, J.; Di Meo, F.; Lo, M.; Delcourt, C.; Clement, S.; Norman, P.; Richeter, S.; Linares, M.; Surin, M. *Org. Biomol. Chem.* 2015, 13, 2453–2463.

(83) Jonkheijm, P.; Hoeben, F. J. M.; Kleppinger, R.; Van Herrikhuyzen, J.; Schenning, A. P. H. J.; Meijer, E. W. *J. Am. Chem. Soc.* 2003, 125, 15941–15949.

(84) Koepf, M.; Wytako, J. A.; Bucher, J.-P.; Weiss, J. *J. Am. Chem. Soc.* 2008, 130, 9994–10001.

(85) Rauch, V.; Wytako, J. A.; Takahashi, M.; Kikkawa, Y.; Kanesato, M.; Weiss, J. *Org. Lett.* 2012, 14, 1998–2001.

(86) Iavicoli, P.; Simón-Sorbed, M.; Amabilino, D. B. *New J. Chem.* 2009, 33, 358–36.