Hyperporphyrin effects extended into a J-aggregate supramolecular structure in water†

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The relationship between the acid–base chemistry and the supramolecular behavior of 5-(4-aminophenyl)-10,15,20-tris(4-sulfonatophenyl) porphyrin in acidic water is reported. A new species exhibiting a prominently red-shifted Q-absorption band at 742 nm is described which is in accordance with a hyperporphyrin-type spectrum of a J-aggregate in water. UV-vis spectroscopy and peak force microscopy reveal that depending on the pH value of the medium the porphyrin self-assembles into two structurally different mesophases which can be reversibly interconverted at will.

Self-organized materials obtained from the self-assembly of monomers that form J-aggregates† arranged in a regular fashion are nowadays promising candidates with potential applications in energy and electron transfer processes in nanostructured materials. In this regard, supramolecular homoassociates of water-soluble porphyrins have received much attention.

Inspired both by the hyperporphyrin aggregates described for the first time by Mons `u Scolaro et al.3 using hydroxy-substituted porphyrins and by the more recent report by Wamser et al.4 on the acid–base chemistry of amino-substituted hyperporphyrins in organic media, we set forth to investigate if both approaches could converge using the water-soluble 5-(4-aminophenyl)-10,15,20-tris(4-sulfonatophenyl)porphyrin (TPPS3NH2, Scheme 1) as a suitable monomeric building block with the aim to compare its supramolecular behavior with that of the well-known 5,10,15,20-tetrakis(4-sulfonatophenyl) porphyrin (TPPS4).5 The results of our investigation are presented in what follows.

The spectrophotometric titration‡ of TPPS3NH2 (species of Fig. 1) could not be conveniently performed in buffered solutions at different pH values owing to an enhanced propensity of this porphyrin to aggregate in solutions of high ionic strength (see below). Instead, a 1.4 × 10⁻⁶ M aqueous solution of the free base form was titrated with sulfuric acid without any interference from the presence of aggregated species (ESI†). In spite of sharp isosbestic points in the spectrophotometric titration (ESI†), only a rough average value for (pKₐ₁ + pKₐ₂)/2 = 5.06 ± 0.02 corresponding to the transition from the free base porphyrin to the diprotonated form (protonation at the inner pyrroleninic nitrogen atoms, Fig. 1) could be measured. This was caused by the unusually high presence of the monoprotonated species‡ as a consequence of its stabilization originated by the electronic conjugation between the protonated central nitrogen atom the peripheral unprotonated amino group.‡ However, although the presence of the lateral amino group does indeed seem to stabilize the monoprotonated form as reported,‡ it does not have a substantial effect on the basicity of the central

![Scheme 1 Chemical structure of TPPS3NH2 and its schematic representation used in this communication.](image-url)
core of the porphyrin ring in water. It had already been unambiguously established that the inner pyrrolic nitrogen atoms of the macrocycle are more basic than the peripheral meso-4-aminophenyl group, as we could corroborate during the titration of TPPS$_3$NH$_2$. Then, the pK$_a$ value determined for the meso-anilinium cation was found to be: 2.84 ± 0.02, corresponding to a basicity more than two orders of magnitude lower than that of the central nitrogen atoms. As a consequence, solutions of the monomeric free-base, diprotonated, and triprotonated forms can all be adequately obtained so that their UV-vis spectra can be recorded as individual species (Fig. 1 and Table 1). The spectrum of the free base shows the expected absorption bands for a monosubstituted meso-4-amino-phenylporphyrin. Upon diprotonation of the inner nitrogen atoms a distinctive hyperporphyrin spectrum showing a split Soret band together with an unusually red-shifted Q-band is very much in line with the typical spectrum of those of tetrasulfonated porphyrins. Further protonation at the anilinic amino group results again in the typical absorption of the diacidic form of a porphyrin protonated at both inner pyrrolic nitrogen atoms without any hyperporphyrin features in the spectrum.

We then studied how the protonation state of the peripheral amino group affected the supramolecular behavior of the porphyrin and the electronic properties of the aggregates, in comparison to the currently well-known example of TPPS$_4$. In relation to its tetraysulfonated counterpart, TPPS$_3$NH$_2$ was found to have an increased propensity to homomassociate. Already at micromolar concentrations of the porphyrin, an order of magnitude lower than for TPPS$_4$ (ESI†), aggregates could be detected by UV-vis spectroscopy, both in aqueous HCl 0.1 M (diprotonated form, NH$_2^-$) and in AcONa/AcOH 0.1 M buffer at pH = 4.0 (diprotonated form, NH$_2$). The UV-vis spectra of these aggregated species are presented on the inset of Fig. 1. The absorption spectrum of the aggregates of the triprotonated form [λ$_{\text{max}}$: 418 nm (H-agg.); 486 nm (J-agg.)] in aqueous HCl 0.1 M corresponding to the aggregates of the triprotonated zwitterionic form; the presence of residual monomeric units is evident from the shoulder at 433 nm. Further protonation of the amino group results in a most significant red-shift of the Q-band displaced to 742 nm. We attribute these changes in the J-aggregate absorption band to a hyperporphyrin effect caused by the electronic conjugation of the lone pair of the basic peripheral amino group with the exciton band of the aggregate in analogy to what is observed for the corresponding diprotonated monomeric species. It is also significant that when the amino group is not protonated, even

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### Table 1: UV-vis absorption data [λ$_{\text{max}}$ (ε)] of the monomeric species of TPPS$_3$NH$_2$ corresponding to the spectra in Fig. 1

|                | Free base | Diprotonated form | Triprotonated form |
|----------------|-----------|-------------------|-------------------|
| 415 (320.000)  | 423 (135.000) | 434 (362.000)     |
| 519 (13.400)   | 436 (126.400) | 593 (5.700)       |
| 560 (9.100)    | 610sh (6.900) | 645 (35.000)      |
| 579 (7.900)    | 669 (33.900)  |                   |
| 639 (4.100)    |           |                   |
at porphyrin concentrations as low as 10 μM the presence of the monomer is not detected. Yet another difference between both types of aggregates is that the triprotonated form (−NH₃⁺) easily flocculates in the solution while the diprotonated form (−NH₂⁺) does not. In addition, both different aggregates (−NH₂⁺/−NH₋) can be reversibly interconverted merely by changing the pH of the solution once the mesophases are already formed (ESI†). Notice, however, that increasing the acidity of the medium to the extent that the sulfonato groups become protonated (pH ≪ 0) results, as expected, in the total disappearance of the aggregates.

The acid–base reversibility between both aggregates is also evidenced by peak force microscopy (PFM) measurements (Fig. 2). The mesophases deposited onto highly ordered pyrolytic graphite from a solution that shows an electronic spectrum with (Fig. 2). The mesophases deposited onto highly ordered pyrolytic graphite evidenced by peak force microscopy (PFM) measurements (ESI†). Notice, however, that increasing the acidity of the medium to the extent that the sulfonato groups become protonated (pH ≪ 0) results, as expected, in the total disappearance of the aggregates.

The acid–base reversibility between both aggregates is also evidenced by peak force microscopy (PFM) measurements (Fig. 2). The mesophases deposited onto highly ordered pyrolytic graphite from a solution that shows an electronic spectrum with the hyperporphyrin absorption bands are very regular (aqueous HCl solution of TPPS₃NH₂ at pH = 4.0 obtained from the addition of a concentrated solution of the sodium salt of the free base porphyrin over the acid and left to aggregate for 24 hours leading to a solution with the expected UV-vis spectrum). In this case, wide plate-like structures consisting of monolayered 2-D aggregates appear on the graphite surface showing regions in which the flat structure is partially folded into a bilayer (3.7 nm of height) in analogy to what has been reported for other porphyrins bearing hydrophobic meso-substituents. When an analogous solution of TPPS₃NH₂ is similarly prepared but at lower pH values so that the amino group is protonated (aqueous HCl solution, pH = 1.3, with a UV-vis spectrum showing the predominance of the aggregates of the triprotonated form), only particles of different heights corresponding to multilayered structured can be detected (ESI†). The reversible interconversion between both types of aggregates was also corroborated by PFM: when a typical solution of TPPS₃NH₂ at pH = 4.0 showing the folded mesophases is further acidified to pH = 1.0 with HCl, the folded structures do disappear, smaller aggregates form and flocculate, and again only irregularly shaped multilayered structures can be detected on the HOPG surface.

It is now rather well-established how the number and relative position of the anionic sulfonato groups in the para-position of the meso-phenyl groups of a protonated tetraphenylporphyrin determine the shape of their self-assembled mesophases and, hence, some of their properties like chirality expression at the mesoscopic scale. The results presented herein based on the tunable amino group (neutral vs. positively charged) at the periphery of the macrocycle show that not only the electronic properties of the J-aggregates can be effectively controlled at will, but also provide further evidence on how the supramolecular architectures of the aggregates can indeed be now predicted to a reasonable extent. The experimental results reported above can be reasonably interpreted as depicted in Scheme 2. Inasmuch as there are two opposite meso-4-sulfonatophenyl groups in TPPS₃ NH₂, the porphyrin is able to form aggregates via the same basic supramolecular pattern than TPPS₄. In a first stage, oligomeric aggregates of the porphyrin further self-assemble into a two dimensional sheet-like structure consisting of J- and H-aggregates by the interaction of two opposite sulfonato groups of the same molecule with the positively charged central pyrrolic nitrogen atoms of adjacent units, together with π–π interactions and hydrophobic effects. Thereupon, the out-of-plane 5-(4-aminophenyl) and 15-(4-sulfonatophenyl) groups can give rise to different stereoisomers of the porphyrin aggregates depending on the tacticity of these lateral groups in relation to the 2-D supramolecular sheet. Depending on the protonation state of the amino group, the conspicuously different growth of the particles that leads to two distinctive mesophases can be rationalized as follows: when at lower pH values the peripheral amino groups are protonated, multilayered structures can easily grow owing to interlayer coulombic interactions (Scheme 2); this fact, together with the total charge compensation in the zwitterionic nature of the triprotonated

Fig. 2 PFM images (topographical map) of TPPS₃NH₂ aggregates deposited on highly ordered pyrolytic graphite (HOPG) from an 8.8 × 10⁻⁵ M aqueous solution of the porphyrin at pH = 4.0 (HCl). The inset shows the height analysis of the section of the spiral structure folded so as to form bilayered regions.

Scheme 2 Schematic representation of the different 3D packing motifs of the 2D sheet-like supramolecular structure of the TPPS₃NH₂ aggregates depending on the protonation state of the peripheral amino group.
species accounts for the increased tendency of the porphyrin to flocculate. The bilayered structures, however, can be exclusively self-assembled when the amino groups are in their basic neutral form and a large enough region of the monolayer rearranges itself into an isotactic pattern with all the 4-aminophenyl groups on the same side of the 2-D structure. In this way, the supramolecular structure as a whole can gain additional thermodynamical stabilization owing to these supplementary interlayer hydrophobic and aromatic π–π interactions provided by folding.

In summary, we report a new type of J-aggregate in water that exhibits a pH-activatable hyperporphyrin type absorption spectrum obtained from self-assembly of a meso-sulfonatophenyl-substituted porphyrin. In addition we have shown how the origin of the unusually red-shifted absorption Q-band of the J-aggregate is unambiguously correlated to the protonation state of the peripheral p-aminophenyl group of TPPS₃NH₂. In this way, the electronic characteristics of the J-aggregates can be reversibly tuned depending on the acidity of the media. This fundamental result may be of potential applicability for the design of novel pH-switchable self-assembled functional materials which optical properties in the near-infrared are based on a programmed response of the porphyrinic J-aggregates.

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