

Highly Non-Planar Dendritic Porphyrin for pH Sensing: Observation of Porphyrin Monocation

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Metal-free porphyrin-dendrimers provide a convenient platform for the construction of membrane-impermeable ratiometric probes for pH measurements in compartmentalized biological systems. In all previously reported molecules, electrostatic stabilization (shielding) of the core porphyrin by peripheral negative charges (carboxylates) was required to shift the intrinsically low porphyrin protonation pK_a 's into the physiological pH range (pH 6–8). However, binding of metal cations (e.g., K^+ , Na^+ , Ca^{2+} , Mg^{2+}) by the carboxylate groups on the dendrimer could affect the protonation behavior of such probes in biological environments. Here we present a dendritic pH nanoprobe based on a highly non-planar tetraaryl-tetracyclohexenoporphyrin (Ar_4TCHP), whose intrinsic protonation pK_a 's are significantly higher than those of regular tetraarylporphyrins, thereby eliminating the need for electrostatic core shielding. The porphyrin was modified with eight Newkome-type dendrons and PEGylated at the periphery, rendering a neutral water-soluble probe (TCHpH), suitable for measurements in the physiological pH range. The protonation of TCHpH could be followed by absorption (e.g., $\epsilon_{\text{Soret}}(\text{dication}) \sim 270,000 \text{ M}^{-1} \text{ cm}^{-1}$) or by fluorescence. Unlike most tetraarylporphyrins, TCHpH is protonated in two distinct steps (pK_a 's 7.8 and 6.0). In the region between the pK_a 's, an intermediate species with a well-defined spectroscopic signature, presumably a TCHpH monocation, could be observed in the mixture. The performance of TCHpH was evaluated by pH gradient measurements in large unilamellar vesicles. The probe was retained inside the vesicles and did not pass through and/or interact with vesicle membranes, proving useful for quantification of proton transport across phospholipid bilayers. To interpret the protonation behavior of TCHpH we developed a model relating structural changes on the porphyrin macrocycle upon protonation to its basicity. The model was validated by density functional theory (DFT) calculations performed on a planar and non-planar porphyrin, making it possible to rationalize higher protonation pK_a 's of non-planar porphyrins as well as the easier observation of their monocations.

Introduction

Over the past decade, porphyrin-dendrimers have been continuously attracting attention in numerous areas of chemistry, including light harvesting, electron transfer, host–guest chemistry, photodynamic therapy, and design of biological sensors and imaging agents.¹ Dendritic encapsulation² offers a convenient method for isolation of the porphyrin moiety from components of biological systems, improving solubility and enabling control over the probe distribution in complex biological environments.

As molecular probes, porphyrin-dendrimers have proven particularly useful for imaging of molecular oxygen³ as well as membrane-impermeable pH probes for compartmentalized biological systems.⁴ The latter application makes use of the fact that protonation of the two imine nitrogens in the core of the porphyrin macrocycle leads to dramatic changes of its optical spectra. Yet both protonated and deprotonated

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forms retain strong absorption and in many cases substantial fluorescence, enabling ratiometric pH measurements. Compared to other macromolecular pH indicators,⁵ porphyrins offer the advantage of very high extinction coefficients (reaching up to $\sim 400,000 \text{ M}^{-1} \text{ cm}^{-1}$ in the Soret-band region), thereby ensuring high signal-to-noise ratios (SNR) and high measurement accuracy. Once encapsulated inside dendrimers with hydrophilic peripheral groups, porphyrins become highly water-soluble and unable to pass through and/or interact with lipid bilayers—a valuable property for pH measurements in multicompartments systems.^{4b,6}

Intrinsic protonation pK_a 's of tetraarylporphyrins⁷ are usually below the physiological pH range (pH 6–8). However, there are several ways by which the proton affinity of the porphyrin macrocycle can be increased, such as substitution with electron-donor groups,⁷ structural deformation⁸ and electrostatic core stabilization.^{9,4} In the previously designed dendritic probes, electrostatic stabilization (shielding) of porphyrin cations by peripheral negative charges (carboxylates) was utilized to tune the protonation pK_a 's into the physiological pH range.⁴ Some of these probes proved instrumental in functional studies;^{4b,6,10} however, dependence of the protonation state of the sensor on the ionization of the peripheral layer made these probes sensitive to environmental effects. For example, binding of divalent metal cations (Ca^{2+} , Mg^{2+} , etc.) by the carboxylate groups and/or alterations in K^+ or Na^+ concentrations influenced the protonation state of the porphyrin, skewing the measurements.

A preferred way to tune the probe's pK_a would be to modulate the intrinsic proton affinity of the porphyrin. Such modulation can be achieved by altering the geometry of the porphyrin macrocycle, since it is well-known that non-planar deformations can dramatically affect porphyrin basicity.⁸ Guided by our previous observation¹¹ that protonation pK_a 's of strongly saddled tetraaryltetracyclohexenoporphyrins (Ar_4TCHP) are several pH units higher than those of regular tetraarylporphyrins, we envisioned using Ar_4TCHPs as cores of pH probes. Below we describe design, synthesis, and properties of a dendritic probe TCHpH, based on Ar_4TCHP , suitable for pH measurements by absorption and/or fluorescence in physiological pH range. TCHpH is an addition to the class of highly non-planar porphyrin-dendrimers, of which there are presently only a few examples.^{3b,c,e,4b,12} The probe was found useful in measurements of proton gradients in microcompartment systems.

A remarkable feature of TCHpH is the existence of a pH range in which the porphyrin monocation is readily observable in mixtures of the protonated forms. This result is of independent interest, since in the overwhelming majority of cases the two protonations of the pyrroline nitrogens of the porphyrin macrocycle are inseparable,¹³ and with rare exceptions^{14–18} porphyrin monocations (or monoacids) are extremely elusive species. To explain this and other properties of TCHpH we developed a model relating acid–base properties of porphyrins to their structural features. The model was validated by DFT calculations, making it possible to rationalize higher protonation pK_a 's of TCHpH and other non-planar porphyrins as well as the presence of their monocations.

Experimental Section

General Information. All solvents and reagents were obtained from commercial sources and used as received. Tetrahydroisoindeole ethyl ester (pyrrole ester) (**1**)¹⁹ and di-*n*-butyl 5-formylisophthalate (aromatic aldehyde)²⁰ were synthesized as described previously. Tris[2-(*tert*-butoxycarbonyl)ethoxymethyl]-methylamine (Gen 1 Newkome-type dendron) was synthesized according to the published method.²¹ Thin-layer chromatography was performed on aluminum-supported silica gel plates (Aldrich). Column chromatography was performed on Selecto silica gel (Fisher). Preparative size exclusion chromatography (SEC) was performed on S-X1 beads (Bio-Rad), using tetrahydrofuran (THF) as a mobile phase. ¹H and ¹³C NMR spectra were recorded on a Bruker DPX-400 spectrometer. The mass spectra were obtained on a MALDI-TOF Voyager-DE RP Bio-Spectrometry workstation, using α -cyano-4-hydroxycinnamic acid as the matrix.

Quartz fluorometric cells (Starna Cells, Inc., 1 cm optical path length) were used in both UV–vis and fluorescence experiments. Optical absorption spectra were recorded on a Perkin-Elmer Lambda 40 UV–vis spectrophotometer and/or an Avantes AvaSpec-2048 fiberoptic spectrometer with a Tungsten Halogen light-source, connected to an in-house constructed titration system.⁶ Steady-state fluorescence measurements were performed on a FS-900 spectrofluorometer (Edinburgh Instruments). The fluorescence quantum yields were determined relative to the fluorescence of Rhodamine 6G ($\phi_f = 0.95$ in EtOH)²² or tetraphenylporphyrin ($\phi_f = 0.11$ in deox. C_6H_6).²³ Time-resolved fluorescence measurements were performed at the Ultrafast Optical Processes Laboratory at the University of Pennsylvania. Dynamic light scattering (DLS) measurements were performed on a Zetasizer Nano-S instrument (Malvern Instruments).

For details of the DFT calculations see Supporting Information. Normal-mode Structural Decomposition (NSD) analysis²⁴

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was performed using an online Java applet available at the Web site of Prof. J. A. Shelnutt, <http://jasheln.unm.edu/jasheln/content/NSD/NSDEngine/start.htm> (Supporting Information, S6).

Synthesis. Tetraaryltetracyclohexenoporphyrin Octabutyl Ester (3). A mixture of pyrrole ester (**1**) (300 mg, 1.55 mmol) and potassium hydroxide (~85%, 140 mg, 2.55 mmol) in ethylene glycol (9 mL) was refluxed under Ar for 1 h. The mixture was cooled to 0 °C and CH₂Cl₂ (20 mL) was added. The organic phase was washed with water and with brine. The product, pyrrole **2**, was extracted with CH₂Cl₂. The solution was washed with water and brine, dried over Na₂SO₄, and the solvent was evaporated in vacuum. The residue was purified by column chromatography (Selecto silica gel/CH₂Cl₂). The resulting pyrrole **2** was dissolved in CH₂Cl₂ (150 mL). The flask was protected from ambient light and flushed with Ar, after which dibutyl-5-formylisophthalate (430 mg, 1.41 mmol) was added in one portion. The solution was stirred for 10 min, after which BF₃·Et₂O (40 mg, 0.28 mmol) was added, and the mixture was stirred at room temperature (r.t.) for 2 h. DDQ (350 mg, 1.55 mmol) was added, and the mixture was stirred overnight. The resulting green solution was washed with 10% aq Na₂SO₃, 10% aq Na₂CO₃, 5% aq HCl, and finally with brine. The organic layer was dried over Na₂SO₄ and reduced in volume by rotary evaporation. The green colored residue was purified by column chromatography on silica gel using CH₂Cl₂/EtOAc (85:15 by volume) as a mobile phase. After evaporation of the solvent, and drying in vacuum, porphyrin **3** was isolated as a green-brown solid. Yield: 230 mg, 10% (based on **1**). UV-vis (CH₂Cl₂-TFA, 9:1) λ_{max} (log ϵ) 464 (5.29), 611 (4.08), 671 nm (4.29); ¹H NMR (CDCl₃) δ 8.40–8.37 (m, 12H), 4.90 (s, 16H), 2.64–2.44 (m, 32H), 1.86–1.84 (m, 16H), 1.68–1.65 (m, 16H), 0.89 (s, 24H), 0.31 (br s, 2H); ¹³C NMR (CDCl₃) δ 159.6, 140.6, 135.0, 131.1, 127.7, 124.1, 123.7, 88.8, 56.1, 39.1, 28.9, 24.0, 15.6, 12.3; MALDI, m/z for C₁₀₀H₁₁₈N₄O₁₆ calcd 1630.8, found 1630.8.

Tetraaryltetracyclohexenoporphyrin Octacarboxylic Acid (4). Porphyrin **3** (46 mg, 0.028 mmol) was dissolved in THF (6 mL) at 0 °C, and two-three pellets of NaOH and MeOH (~0.5 mL) were added to the mixture. The mixture was left to stir at r.t. overnight, after which it was concentrated in vacuum, and several drops of water were added to dissolve the brown precipitate. The solution was stirred for 1 h at r.t., and THF was removed by rotary evaporation. The solution was acidified with conc. HCl. The precipitate was isolated by centrifugation and washed with water three times to give **4** as a dark green powder. Yield: 33 mg, 94%. ¹H NMR (DMSO-*d*₆) δ 8.33–8.05 (m, 12H), 1.45–0.94 (m, 32H), 0.39 (br s, 2H); MALDI, m/z for C₆₈H₅₄N₄O₁₆ calcd 1182.3, found 1182.7.

Porphyrin-Dendrimer Poly-¹Bu-ester (5). Porphyrin **4** (52 mg, 0.044 mmol) and HBTU (250 mg, 0.66 mmol) were dissolved in dry NMP (25 mL). DIPEA (520 mg, 4.05 mmol) was added to the solution at r. t., immediately followed by the addition of the Newkome G1 dendron (334 mg, 0.66 mmol). The resulting mixture was stirred for 4 days at r.t., after which it was poured into aq. NaCl (3%). The resulting precipitate was collected by centrifugation, washed with water by suspension/centrifugation cycles (3 × 30 mL), and dried in vacuum. The green-brown viscous solid was redissolved in THF and purified by SEC (Bio-Rad S-X1 Beads, 200–400 mesh, THF) to yield the porphyrin-dendrimer **5**. MALDI, m/z for C₂₆₈H₄₁₄N₁₂O₈₀ calcd 5080.9, found 5080.3. The MALDI spectrum revealed that the resulting product contained ~15% of the porphyrin modified with 7 Newkome dendrons.

Porphyrin-Dendrimer Polycarboxylic Acid (6). To a well-stirred solution of **5** (198 mg, 0.039 mmol) in CH₂Cl₂ (12 mL) was added TFA (10 mL) dropwise at 0 °C over a period of 20 min. The solution was allowed to react for 2 h, after which the solvent and the excess TFA were evaporated in vacuum at r.t. The residual TFA was removed by redissolving the precipitate in

CH₂Cl₂ (10 mL) and evaporating it in vacuum. This procedure was repeated several times, leaving polyacid **6** as a dark green viscous material. Yield: 142 mg, 98%. MALDI, m/z for C₁₇₂H₂₂₂N₁₂O₈₀ calcd 3735.4, found 3735.4. The MALDI spectrum revealed that the resulting product contained ~15% of the porphyrin modified with 7 Newkome dendrons.

Porphyrin-Dendrimer Poly-PEG-ester (TCHpH). Polyacid **6** (129 mg, 0.035 mmol) was dissolved in CH₂Cl₂ (10 mL), monomethoxyoligoethyleneglycol (Av. MW 750, 3 mL) was added, and the solution was left under stirring for 15 min at r.t., after which it was cooled on an ice bath. *sym*-Collidine (8–10 drops), HOBT (1.18 g, 8.75 mmol), and DCC (1.81 g, 8.75 mmol) were added to the mixture, and it was left to react under stirring for 4 days at r.t. Ice-cold 0.05 M aq. HCl (50 mL) was added to the mixture to quench the reaction, and the mixture was stirred for 30 min to decompose the unreacted DCC. The mixture was filtered through a Whatman 55 filter paper. The filtrate was extracted with CH₂Cl₂, and the solvent was removed in vacuum. The product was purified by size-exclusion column chromatography (Bio-Rad S-X1 beads, 200–400 mesh, THF) to yield the target TCHpH probe as a green viscous solid. Yield: 396 mg, 53%.

pH Titrations. Bulk titrations of TCHpH were performed at 22 °C using 50 mM potassium phosphate buffer or 25 mM HEPES buffer. The pH was adjusted by addition of HCl, H₂SO₄, or KOH. K₂SO₄, Ca(OAc)₂, and Mg(OAc)₂ were used to test the influence of metal ion concentrations. In the titrations by absorption, absorbances of solutions at Soret peak maxima were kept below 1.0 OD. In titrations by fluorescence, absorbances at the excitation wavelengths were kept below 0.1 OD.

The titration curves were constructed by plotting absorbances at selected wavelengths, or, in the case of fluorescence titrations, integrated fluorescence versus pH. In ratiometric titrations, ratios of the integrated fluorescence spectra upon excitation at selected wavelengths (e.g., at the Soret peaks corresponding to the free-base and dication forms) were plotted versus pH. The titration data were fitted to the Henderson–Hasselbalch-type equation derived for the case of two sequential protonations (Supporting Information, S2):

$$S_{\lambda}(\text{pH}) = \frac{P_3 \times 10^{-2\text{pH}} + P_2 \times 10^{-\text{pH}} + P_1}{10^{-2\text{pH}} + P_4 \times 10^{-\text{pH}} + P_5} \quad (1)$$

where $S_{\lambda}(\text{pH})$ designates the absorbance at a selected wavelength λ and fitting parameters P_1 – P_5 are the functions of the equilibrium constant(s), probe concentration, and molar extinction coefficients of the free-base, monocation, and dication. In eq 1, $P_4 = K_4$ and $P_5 = K_3K_4$ (Supporting Information, S2). Similar treatment was used previously to extract protonation constants K_3 and K_4 of other water-soluble porphyrins.²⁵ Fitting was performed using Microcal Origin software (Origin 7.0).

Liposome Experiments. Large unilamellar liposomes containing TCHpH were prepared from palmitoyl-oleoyl phosphatidylcholine (Avanti Polar Lipids) according to the protocol described previously.⁶ The buffer was potassium phosphate (20 mM, pH 7) with addition of EDTA (250 μ M) and K₂SO₄ to bring the total K⁺ concentration to 50 mM. This procedure is designed specifically for the preparation of proteoliposomes, and, therefore, it includes the step of partial liposome solubilization after their initial formation. The solubilization was performed using octylglucoside (Avanti Polar Lipids) as a detergent, which was subsequently removed by adsorption on hydrophobic beads (Bio-Beads, SM-2, Bio-Rad). TCHpH was added to the mixture prior to the step of detergent removal to a total concentration of 4.67 mg/mL. After vesicle formation, the probe not enclosed into liposomes was removed by centrifugation of the liposomes at 40,000 rpm for 2 h, discarding the supernatant

and resuspending the formed pellet in a fresh buffer (6 mL). This procedure was repeated three times. On the basis of the DLS measurements, the liposomes were characterized by homogeneous size distributions with the average diameter of ~ 200 nm.

For calibration of the signal from the encapsulated probe, gramicidin (Sigma) solution in dimethylsulfoxide (DMSO) was added to liposome suspensions to make up the final concentration of $\sim 1 \mu\text{g/mL}$. The bulk pH was titrated, and the absorption spectra were corrected for the scattering background arising from the turbidity of the liposome suspension. The normalized quantity R_{fb} , used for fitting in these experiments, was determined using absorbances at 430 nm (on the blue edge of the Soret band of the free-base) and 459 nm (λ_{max} of the Soret band of the dication):

$$R_{fb} = \frac{S_{430} - S_{459} \times a}{S_{430} \times b + S_{459} \times a} \quad (2)$$

where constants a and b account for the difference in the extinction coefficients of differently protonated forms and their spectral overlap.

The pH-jump experiment (Figure 5) was performed using a suspension of liposomes in a mixture of Mes buffer (20 mM), potassium phosphate buffer (20 mM), and EDTA (250 μM). The pH in the extraliposomal (bulk) solution was set to 4.75. The spectra were recorded every 5 s during ~ 16 min, after which gramicidin in DMSO was added to make a final concentration of $1 \mu\text{g/mL}$.

The passive proton permeability of the liposome bilayer was calculated from the difference in pH between times t_1 (pH₁) and t_2 (pH₂) during the initial period prior to the addition of gramicidin (Figure 5), taking into account the capacity of the buffer inside the liposomes (20 mM, $\text{p}K_a$ 6.87):

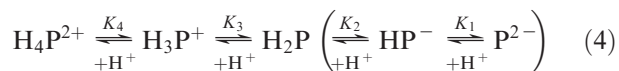
$$\Delta[\text{H}^+] = \frac{20 \times 10^{(6.87 - \text{pH}_2)}}{1 + 10^{(6.87 - \text{pH}_2)}} - \frac{20 \times 10^{(6.87 - \text{pH}_1)}}{1 + 10^{(6.87 - \text{pH}_1)}} \quad (3)$$

Expression 3 gives the change in the proton concentration $[\text{H}^+]$ (in mM) inside the liposomes between times t_1 and t_2 . When multiplied by the total internal liposome volume, this quantity gives the total amount of protons that traversed the phospholipid membrane.

Adhesion of TCHpH to polystyrene beads (SM-2, Bio-Rad) was tested by adding the beads (100 mg) to a solution of the probe in the buffer (1 mL, 8.5 mg/mL of TCHpH, pH 7.2), incubating the suspension under stirring for 1 h, adding more beads (50 mg) and continuing stirring for additional 1 h.

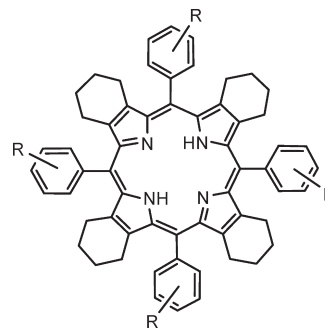
Results and Discussion

Choice of the Porphyrin Core and Probe Synthesis. Free-base porphyrins (H_2P) are able to bind two protons by the core imine nitrogens with formation of *mono*- (H_3P^+) and *di*- (H_4P^{2+}) cations. Traditionally, these two proton dissociation reactions are described by equilibrium constants K_3 and K_4 , while constants K_1 and K_2 are reserved for formation of porphyrin anions.⁷



The choice of Ar_4TCHP (Chart 1) as a core for the dendritic probe was motivated by our previous measurements of $\text{p}K_a$'s ($\text{p}K_3$ and $\text{p}K_4$) of water-soluble Ar_4TCHPs ¹¹ and subsequent studies of Ar_4TCHP photophysics.²⁶ It is well-known that $\text{p}K_a$'s of non-planar porphyrins are

Chart 1. Structure of Ar_4TCHP



typically higher than $\text{p}K_a$'s of their planar counterparts.⁸ Ar_4TCHPs belong to the class of highly non-planar porphyrins; and in addition, their proton affinities are increased because of the σ -electron-donor effect of the cyclohexeno-substituents in the β -pyrrolic positions. As a result, $\text{p}K_a$'s of Ar_4TCHPs may become too high, that is, higher than the physiological pH range. To counterbalance the effect of the cyclohexeno-substituents we introduced electron-withdrawing carboxyl groups into the *meta*-positions of the *meso*-aryl rings, which simultaneously provided anchor points for linking the solubilizing dendrons.

The synthesis of probe TCHpH (Scheme 1) followed the path developed previously for the synthesis of phosphorescent Pt and Pd porphyrin dendrimers.^{3c} Pyrrole **2** with fused cyclohexene rings was obtained by decarboxylation of pyrrole-ester **1**, which was synthesized by the modified Barton–Zard method, as reported previously.¹⁹ Octabutoxycarbonyltetracyclohexenoporphyrin **3** was synthesized by the Lindsey method²⁷ and converted into the corresponding octacarboxylic acid **4** by way of base-mediated hydrolysis. The eight carboxylic groups on **4** were modified with Newkome-type dendrons using HBTU/DIPEA coupling chemistry, and the resulting porphyrin-dendrimer **5** was purified by SEC on polystyrene beads using THF as a mobile phase. The product was found to contain 10–15% of the porphyrin modified with seven Newkome dendrons, which is quite typical for modification of porphyrins with many anchor points.

Newkome-type ester-amide dendrons²⁸ were chosen because they possess a trifurcated branching junction, thereby producing 24 peripheral functionalities already at the first generation. Newkome-type dendrons are easily accessible²⁹ and have been used previously for constructions of porphyrin-dendrimers.^{30,4b} The peripheral *tert*-butyl ester groups on **5** were cleaved by TFA, giving polyacid **6**, whose carboxyl groups were subsequently esterified with monomethoxyoligo(ethylene glycol)

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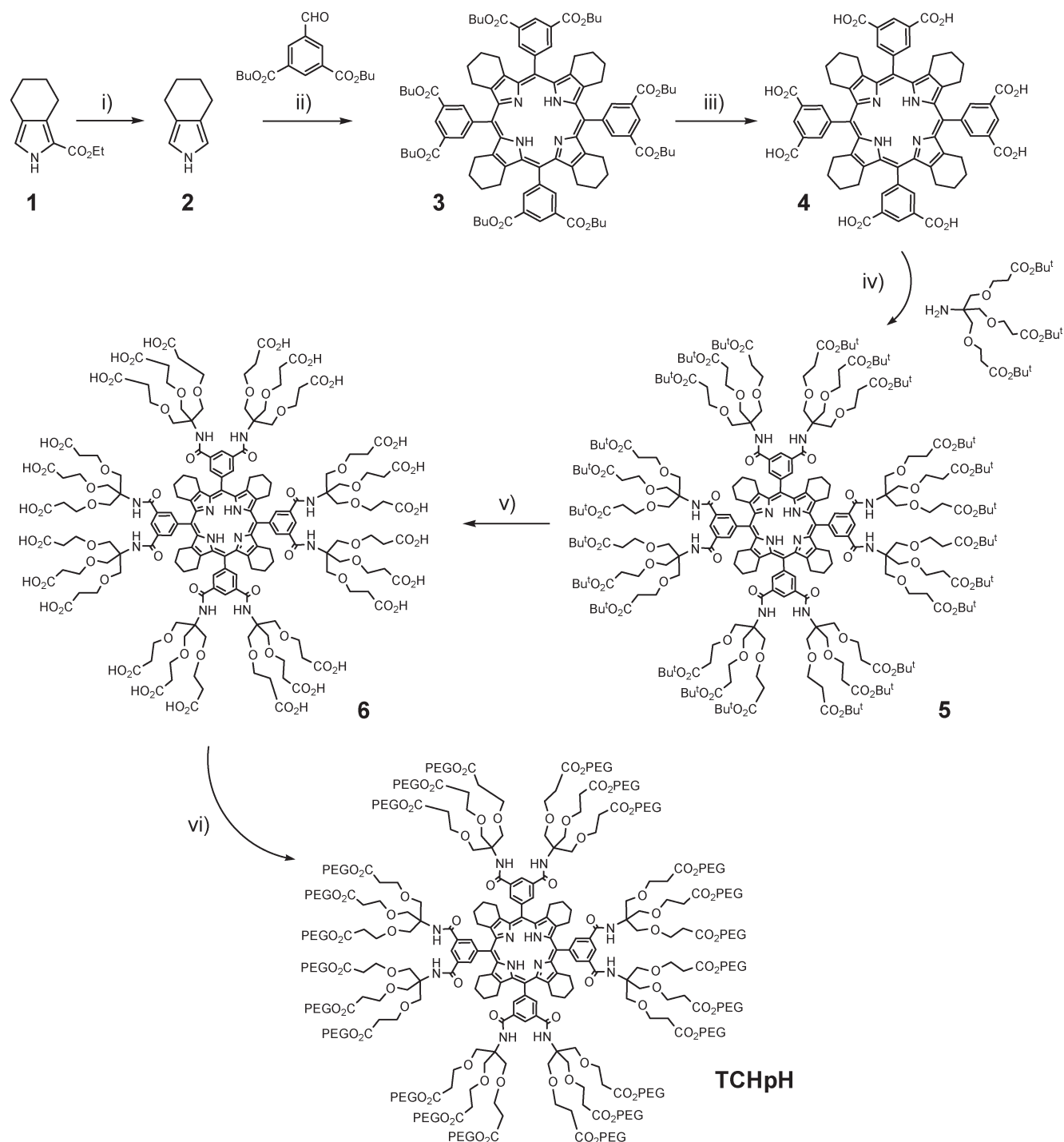
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Scheme 1. Synthesis of Probe TCHpH



Reagents and conditions: (i) KOH, HOCH₂CH₂OH, reflux, 1 h; (ii) (a) BF₃·Et₂O, CH₂Cl₂, r.t., 2 h; (b) DDQ, r.t., 12 h, 10% for two stages; (iii) NaOH, THF, r.t., 12 h, 94%; (iv) HBTU, DIPEA, NMP, r.t., 4 days, 86%; (v) TFA, 0 °C, 20 min, 98%; (vi) PEG750, DCC, HOBT, *Sym*-collidine, CH₂Cl₂, r.t., 4 days, 53%. Abbreviations: NMP = *N*-methylpyrrolidone; HBTU = 2-(1*H*-benzotriazol-1-yl)-1,1,3,3-tetramethyluronium hexafluorophosphate; DIPEA = *N,N*-diisopropylethylamine; TFA = Tri fluoroacetic acid; PEG750 = oligoethyleneglycol monomethyl ether, Av. MW 750; DCC = *N,N*-dicyclohexylcarbodiimide; HOBT = 1-hydroxybenzotriazole.

(Av. MW 750) using the DCC/HOBT method.^{30a,b} The final PEGylated porphyrin-dendrimer (TCHpH) was purified twice by SEC on polystyrene beads. TCHpH was isolated as a dark green viscous material, exhibiting very high aqueous solubility.

Spectroscopic Properties of TCHpH. The changes in the optical absorption spectra of TCHpH upon changes in

pH are shown in Figure 1. The free-base probe (pH 10.5) has its Soret band ($S_0 \rightarrow S_2$) at 442 nm (Figure 1b) and shows the characteristic four-band pattern in the Q-band ($S_0 \rightarrow S_1$) region ($Q_y(1,0)$, $Q_y(0,0)$, $Q_x(1,0)$, $Q_x(0,0)$) (Figure 1c), which is typical for tetraarylporphyrins.³¹ All

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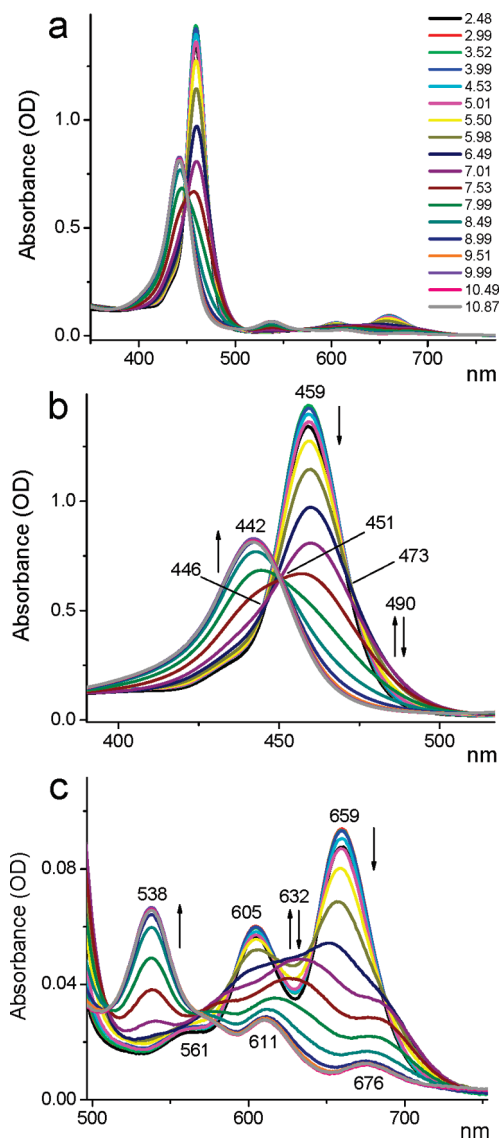


Figure 1. Changes in the absorption spectra of probe TCHpH upon changes in pH (pH is shown in the legend, upper graph). Buffer: phosphate, 50 mM. (a) Full spectrum; (b) Soret band region; (c) Q-band region. Arrows indicate the direction of the intensity changes with increase in pH.

bands are broadened and red-shifted compared to TPP, which is consistent with the high non-planarity and conformational flexibility of the Ar_4TCHP skeleton.^{26,32,33} The red-most absorption ($\text{Q}_x(0,0)$) has its maximum at 676 nm.

The spectrum of the fully protonated species (porphyrin dication, pH 4.0) exhibits a narrower and higher intensity Soret band ($\lambda_{\text{max}} = 459$ nm), which is 838 cm^{-1} lower in energy than the peak of the free-base Soret band (Figure 1b). This is a much smaller shift compared to what is typically encountered in planar tetraarylporphyrins.^{33d,e} For example, the corresponding difference between the Soret peaks of the free-base and dication of water-soluble probe Glu3, based on tetracarboxyphenylporphyrin, is 1317 cm^{-1} .⁶

Because of the higher symmetry of the dication, the four bands of the free-base merge into the single $\text{Q}(1,0)\text{-Q}(0,0)$ pattern ($\lambda_{\text{max}} = 605\text{ nm}, 659\text{ nm}$), shifted blue relative to the lower energy $\text{Q}_x(0,0)$ band of the free-base. This behavior is different from tetraarylporphyrins, where the protonation is accompanied by large red shifts, which occur because of the induced strong non-planar deformations.^{33d}

The extinction coefficients were measured for the dication of porphyrin **3** in $\text{CH}_2\text{Cl}_2/\text{TFA}$ at 22°C ($\epsilon_{\text{Soret}} = 197,000\text{ M}^{-1}\text{ cm}^{-1}$, $\lambda_{\text{max}}(\text{Soret}) = 464\text{ nm}$). It was assumed that for the same transitions the oscillator strengths of the dendritic porphyrin (TCHpH probe) in water and of the parent porphyrin **3** in CH_2Cl_2 would be similar. The extinction coefficients of TCHpH in an aqueous buffer were calculated to be $\epsilon_{459}(\text{dication}) = 272,000\text{ M}^{-1}\text{ cm}^{-1}$ and $\epsilon_{442}(\text{free-base}) = 163,000\text{ M}^{-1}\text{ cm}^{-1}$. These values are somewhat lower than the extinction coefficients of the previously reported porphyrin-based pH probes,^{4,6} but still are more than adequate for analytical applications.

Examination of the absorption changes during protonation revealed the presence of several isosbestic points. For example, in the Soret band region two such points (446 and 473 nm) occur in the pH range 3.0–6.5 and one (451 nm) in the range 7.5–10.0, suggesting two distinct protonation processes. To evaluate the dissociation pK_a 's, absorbances at selected wavelengths were plotted as a function of pH and fitted to eq 1, which describes two sequential protonations (Figure 2) (Supporting Information, S2).

The two wavelengths were chosen to be 435 nm, the blue edge of the Soret peak of the free-base, and 459 nm, λ_{max} of the Soret peak of the dication. Independent least-squares fitting of the two data sets revealed nearly the same pK_a 's: $\text{pK}_3 = 7.79$ and $\text{pK}_4 = 6.03$ (Figure 2a). As expected for non-planar porphyrins, these values are shifted significantly toward higher pH compared to pK_a 's of regular tetraarylporphyrins.⁷ The optical spectra were found to be practically unaffected by the choice of the buffer (for example, HEPES (25 mM) vs 50 mM phosphate) in the physiological pH range (Supporting Information, S9).

For the purpose of practical pH determination it is useful to use ratiometric curves, whose midpoint values (pH_{mid}) can be tuned by changing measurement wavelengths (Supporting Information, S3). An example of such curves is shown in Figure 2b. Ratios of absorbances at 435 and 459 nm were plotted against pH and fitted to Boltzmann-type sigmoidal curves (Supporting Information, S5). The midpoints of the two fits, $\text{pH}_{\text{mid}} = 6.64$ and $\text{pH}_{\text{mid}} = 8.15$, make these curves optimally suited for measurements in two overlapping, but different pH ranges, about 4.5–8.5 and 6.5–10.

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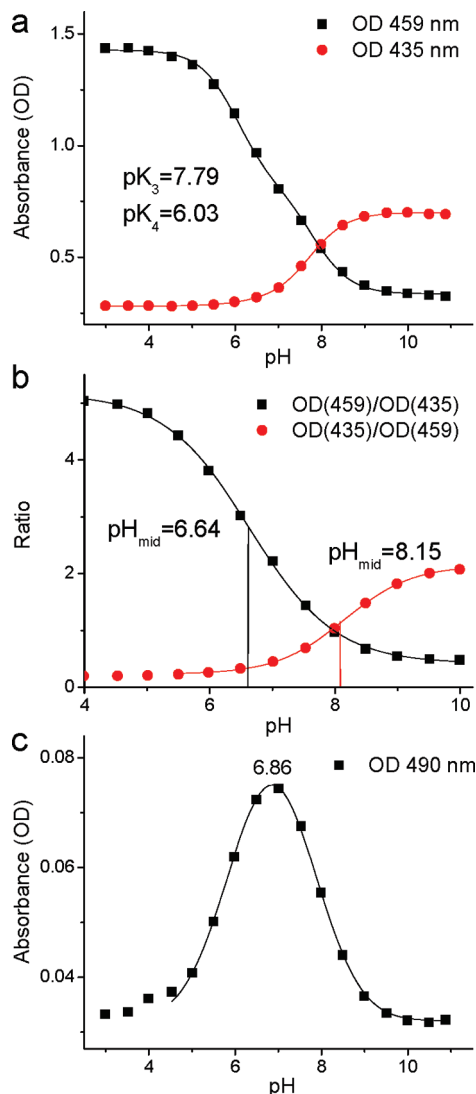


Figure 2. Changes in absorbances of TCHpH with pH: (a) OD at 435 nm and OD at 459 nm and their fits to eq 1; (b) ratios of absorbances and their fits to the Boltzmann type sigmoidal curves (S3–S5); (c) OD at 490 nm and its fit to a Gaussian form ($pH_{max} = 6.86$).

A remarkable feature of the titrations shown in Figure 1b is that a distinct spectral pattern emerges and then disappears again with a monotonic increase in pH. For example, the absorbance at ~ 632 nm rises and then falls back, and the absorbance at the red edge of the Soret band (490 nm) undergoes similar changes, reaching the maximum around pH 6.9 (Figure 2c). This behavior indicates the presence of an intermediate species, which was tentatively assigned to the porphyrin monocation.

To obtain the spectral signature of the monocation, the absorption spectrum recorded at pH 7.01 was fitted with a linear combination of the spectra of the free-base and the dication (recorded at pH 10.0 and 3.5, respectively) and the sum of parameters, each representing the absorbance at its own individual wavelength (see Supporting Information, S12). The fitting was performed by the linear least-squares, implemented as a positively constrained quadratic optimization algorithm.³⁴

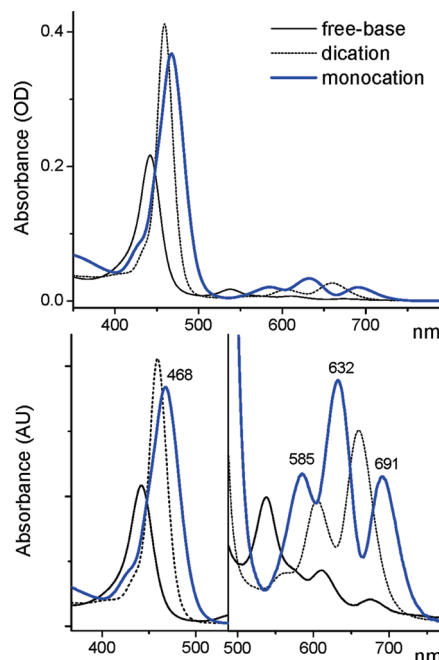


Figure 3. Optical absorption spectra of the three absorbing species: TCHpH monocation (blue), free-base and dication. The spectrum of the monocation was obtained by deconvolution of the experimental spectrum recorded at pH 7.01.

The result is depicted in Figure 3, together with the weighted spectra of the dication and the free-base. The original experimental spectrum (the sum of all three components) can be seen in Figure 1 (also see Supporting Information, Figure S6). Very similar results were obtained by fitting the spectra of the probe recorded at other pH values between 6.5 and 7.5.

The spectrum of the monocation is distinctly different from the spectra of the free-base and dicationic forms. It exhibits a red shift of the Soret band versus the Soret bands of both the free-base (by 1257 cm^{-1}) and the dication (by 419 cm^{-1}). The Soret band of the monocation is broadened and, in fact, it may well be a superposition of two or more overlapping bands resulting from splitting because of the monocation asymmetry.

In the Q-band region, the absorption spectrum of the monocation consists of three peaks: 585, 632, and 691 nm. A similar three-band pattern was observed previously for the monocation of octaethylporphyrin (H_3OEP^+), stabilized by a large transition-metal based anionic complex.¹⁴ Since the monocation is an asymmetrical species, it is expected to possess two orthogonal transition dipoles, Q_x and Q_y , just like the parent free-base;^{31,33e} however, the Q_x , Q_y transitions of the monocation may be closer in energy, and their $Q_y(0,0)$ and $Q_x(1,0)$ components may overlap, giving rise to the observed three-band pattern. The lowest energy $Q_x(0,0)$ band of the monocation ($\lambda_{max} = 691\text{ nm}$) is red-shifted by 321 cm^{-1} relative to the $Q_x(0,0)$ band of the free-base, which may be due to the monocation's non-planar distortion.^{33d}

The fluorescence quantum yield of the parent porphyrin 3 (dication) in CH_2Cl_2 /TFA was found to be 0.0050 ± 0.0007 when measured against fluorescence of Rhodamine 6G in EtOH ($\phi_f = 0.95$)²² and 0.010 ± 0.001 when measured against fluorescence of H_2TPP in deoxygenated

(34) Shrager, R. I. *Commun. ACM* **1972**, *15*, 41.

benzene ($\phi_f = 0.11$).^{23,35} This low yield is a result of rapid internal conversion of the S_1 state, a process well documented for non-planar porphyrins,³³ including Ar₄TCHPs.²⁶ The fluorescence quantum yield of the TCHpH dication was found to be approximately twice as high when measured in aqueous buffer ($\phi_f = 0.012 \pm 0.0008$, measured against Rhodamine 6G), which suggests lower conformational flexibility of the porphyrin skeleton and could in part result from the modification of the porphyrin with bulky dendritic substituents. The fluorescence decay of the dication in aqueous buffer (Supporting Information, S16) could be approximated by a single-exponential function with the time constant of 0.59 ns, which resembles S_1 lifetimes of other distorted porphyrins.³³ The fluorescence quantum yield of TCHpH free-base (pH 10.2) is about two times lower than that of the dication ($\phi_f = 0.006 \pm 0.0008$, measured against Rhodamine 6G).

In spite of its low quantum yield, fluorescence of TCHpH still can be useful as a pH indicator, especially for bulk ensemble measurements. Changes in the emission spectra with varying pH are shown in Figure 4. Fluorescence was excited at either 442 nm (λ_{max} (Soret) of the free-base) or at 459 nm (λ_{max} (Soret) of the dication), while the absorbances were kept below 0.1 OD.

Similar to other non-planar porphyrins, TCHpH exhibits broad featureless emission spectra, whose changes upon protonation follow the changes in absorption (Figure 1). Since the emission signal is weak, the most practical method for pH measurements would be dual-wavelength excitation with collection of the entire integrated fluorescence output. For example, a ratiometric curve could be constructed by plotting the ratio of the integrated fluorescence intensities upon excitation at the Soret peaks of the free-base and dication forms (Figure 4, inset), showing suitability of the fluorescent signal for physiological pH measurements.

Attempts to isolate the excitation spectrum of the monocation by taking measurements of a sample equilibrated at pH 6.9 resulted in spectra with maxima varying with the emission wavelength, but all the spectra were superpositions of all three emitting species (free-base, monocation and dication). Similarly, emission spectra obtained at different excitation wavelengths showed shifts in the emission maxima (Supporting Information, S15). Further, a series of emission spectra were recorded at different pH values for the excitation at 490 nm (Figure 4c), that is, at the red-shoulder of the monocation absorption (Figure 3). Near pH 7.0, the fluorescence has its maximum at 715 nm, which is significantly red-shifted relative to the emission peaks of the dication (Figure 4b). The emission at pH 7.0 originates mostly from the porphyrin monocation.

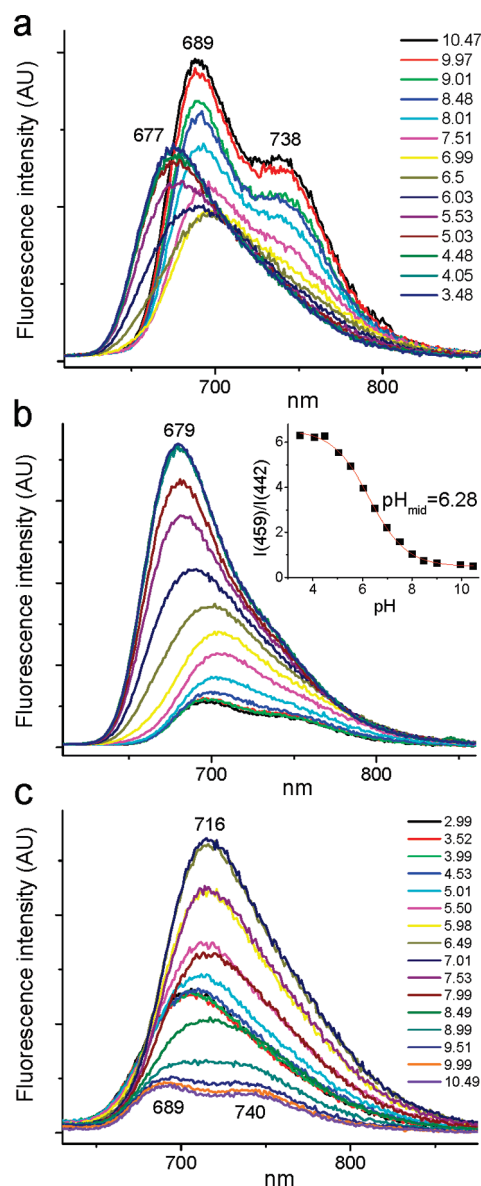


Figure 4. Changes in fluorescence spectra of TCHpH upon changes in pH (pH is shown in the legend, upper graph). Buffer: phosphate, 50 mM. (a) $\lambda_{\text{ex}} = 442$ nm; (b) $\lambda_{\text{ex}} = 459$ nm; (c) $\lambda_{\text{ex}} = 490$ nm. Inset: ratio of integrated fluorescence intensities versus pH: $R = I(459)/I(442)$, where $I(459)$, fluorescence spectrum integrated from 600 to 860 nm, $\lambda_{\text{ex}} = 459$ nm; $I(442)$, fluorescence spectrum integrated from 700 to 860 nm, $\lambda_{\text{ex}} = 442$ nm.

Probe Performance in Microcompartment Systems - Liposome Experiments. To evaluate the suitability of TCHpH for measurements in microcompartment biological systems, we first investigated the dependence of the probe read-out on the concentration of metal cations. Titrations of TCHpH were performed at several K^+ concentrations, showing no apparent difference in the calibration plots (Supporting Information, S10). In contrast, carboxyl-terminated dendritic probes, for example, Glu,³ were shown to be sensitive to ionic strength.⁶ Between pH 6 and 8 in HEPES buffer (25 mM) the spectra of TCHpH did not show any visible changes upon addition of up to 28 mM of Ca or Mg acetates (Supporting Information, S10). However, at lower pH values (e.g., pH 5.0) absorbances at the Soret bands' maxima somewhat decreased at higher cation concentrations.

(35) The discrepancy between the quantum yields measured against the two common standards is a result of overestimated quantum yield for the reference fluorescence of H₂TPP. The quantum yield of $\phi_f(\text{H}_2\text{TPP}) = 0.11$ was measured relative to the fluorescence of chlorophyll *b* in benzene ($\phi_f = 0.12$), as reported by Weber and Teale (Weber, G.; Teale, F. W. J. *Trans. Faraday Soc.* **1957**, 53, 646). In the same paper, Weber and Teale report the quantum yield $\phi_f = 0.97$ for Rhodamine B in EtOH. As was shown later (Karstens, T.; Kobs, K. J. *Phys. Chem.* **1980**, 84, 1871) this value was significantly overestimated. If measured against the fluorescence of H₂TPP in deoxygenated benzene, assuming $\phi_f(\text{H}_2\text{TPP}) = 0.11$, the fluorescence of Rhodamine 6G in EtOH has the quantum yield of 1.9.

A useful quality of dendritic probes for proton translocation measurements is the lack of adsorption to hydrophobic beads, which are typically used in preparation of proteoliposomes. The adsorption of TCHpH was found to be somewhat higher than that of polycarboxylated probes,⁶ but significantly below that of commercial pH probes. For example, after 90 min of incubation with polystyrene beads, 89% of TCHpH remained in solution, whereas 72% of commercial macromolecular probe SNARF-dextran (10 kDa) remained in solution under the same conditions.

The performance of TCHpH in pH gradient measurements was evaluated by incorporating the probe into large unilamellar vesicles made of phosphatidyl choline. The vesicles were prepared by sonication, extrusion, solubilization by detergent and detergent removal using beads, followed by the removal of the bulk probe by ultracentrifugation.⁶ The average liposome diameter was 200 nm, as determined by dynamic light scattering. The concentration of the probe inside vesicles was kept high, 0.22 mM, to ensure high signal-to-noise ratios in absorbance measurements. To calibrate the pH response of the encapsulated probe, the liposomes were treated with a non-specific ionophore gramicidin. Gramicidin forms channels in phospholipid bilayers, making them permeable for small ions, including protons,³⁶ but impermeable to large molecules, such as TCHpH. The absorption spectra of the probe encapsulated inside the liposomes were collected at different pH values, and a calibration curve was constructed (Figure 5a) by fitting the normalized absorbance (R_{pH}) (eq 2) to a sigmoidal curve. This curve was found to be nearly identical to the one obtained in bulk titrations, showing that the liposomal entrapment of the probe and its high local concentration have no effect on the calibration parameters.

A pH gradient across the phospholipid bilayer was created by mixing one part of the liposomes' suspension in a buffer at pH 7.0 with nine parts of a buffer at pH 4.75 (Figure 5). Mixing of the two buffers brought the extraliposomal pH to 5.3, while the intraliposomal pH right after mixing was 7.0. The probe was encapsulated inside the liposomes. Its absorption spectra were recorded continuously, and absorbances at selected wavelengths were converted into pH values using the calibration curve shown in Figure 5a.

During the initial period, the pH inside the liposomes (pH_{in}) showed a slow drift, reflecting passive transport of protons across the phospholipid bilayer. The internal liposome volume was calculated to be about 1.5% of the total volume of the suspension. Taking into account the average liposome size and the buffer concentration, the change in pH_{in} in the absence of gramicidin (Figure 5b) could be converted into the average passive permeability of the liposome bilayer, giving the value of ~ 30 protons/ $\text{s} \cdot \mu\text{m}^2$ under the pH gradient of 1.7. This permeability agrees well with previous estimations.⁶

After approximately 16 min, gramicidin was added, instantly forming pores and permitting free proton flow along the concentration gradient, that is, from the outside into the liposomes. As a result, pH_{in} abruptly dropped, reflecting fast equilibration of the external and internal volumes. This experiment demonstrates that TCHpH is a

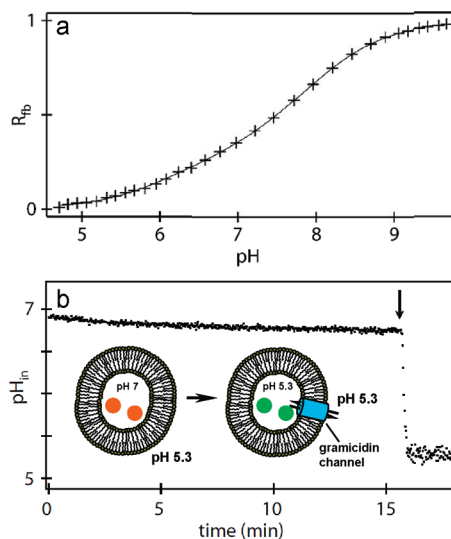


Figure 5. Probe inside phospholipid vesicles. (a) pH titration plot obtained by measuring absorption spectra of the probe inside liposomes with channels made by gramicidin. (b) pH-jump experiment. pH_{in} – pH inside liposomes, as measured by absorbance, converted into pH values using the calibration curve in the upper graph. Gramicidin added after 16 min (arrow), resulting in an abrupt decrease in pH_{in} because of the rapid mixing of internal and external volumes.

membrane-impermeable probe, capable of reporting pH in internal liposomal compartments.

Acid–Base Properties of TCHpH - Structural Analysis.

In view of the observed behavior of TCHpH it is interesting and relevant to discuss acid–base properties of non-planar porphyrins. Traditionally, higher protonation $\text{p}K_{\text{a}}$'s ($\text{p}K_3$ and $\text{p}K_4$ in eq 4) of non-planar porphyrins are attributed to easier accessibility of the core nitrogens to protons in distorted macrocycles.³⁷ However, easy accessibility does not explain the relative thermodynamic stability of protonated forms, but rather implies faster protonation reactions. The rate of protonation may or may not correlate with the protonation free-energy ΔG_{p} ,³⁸ and hence the question still needs to be answered: why are non-planar porphyrins more basic than planar porphyrins, that is, $\Delta G_{\text{p}}(\text{non-planar}) < \Delta G_{\text{p}}(\text{planar})$?

In our discussion below we focus on electronic effects. Therefore, instead of using ΔG_{p} (Gibbs free energy of protonation), we consider only internal electronic energy ΔE_{p} (also see ref 45). We divide ΔE_{p} into two parts, the intrinsic proton affinity ΔE_{intr} and the structural term ΔE_{st} .¹¹

$$\Delta E_{\text{p}} = \Delta E_{\text{intr}} + \Delta E_{\text{st}} \quad (5)$$

It is well-known that protonation of porphyrins leads to their non-planar, typically saddling, deformations.³⁷ With only rare exceptions,^{39,40} porphyrin cations are known to

(37) (a) Stone, A.; Fleisher, E. B. *J. Am. Chem. Soc.* **1968**, *90*, 2735. (b) Pasternack, R. F.; Sutin, N.; Turner, D. H. *J. Am. Chem. Soc.* **1976**, *98*, 1908. (c) Cheng, B.; Munro, O. Q.; Marques, H. M.; Scheidt, W. R. *J. Am. Chem. Soc.* **1997**, *119*, 10732.

(38) We define ΔG_{p} as the free energy of porphyrin protonation, which is the opposite of proton dissociation, associated with ΔG_{d} : $\Delta G_{\text{prot}} = -\Delta G_{\text{d}}$, $\Delta G_{\text{d}} \sim \text{p}K_{\text{a}}$. For the two step reaction: $K_{\text{a}} = K_1 K_2$ (or $K_3 K_4$, as in eq 4) and $\Delta G_{\text{p}} = \Delta G_{\text{p}}(1) + \Delta G_{\text{p}}(2)$.

(39) Cetinkaya, E.; Johnson, A. W.; Lappert, M. F.; McLaughlin, G. M.; Muir, K. W. *J. Chem. Soc., Dalton. Trans.* **1974**, 1236–1243.

(40) Senge, M. O.; Forsyth, T.; Nguyen, L. T.; Smith, K. M. *Angew. Chem., Int. Ed. Engl.* **1994**, *33*, 2485–2487.

be strongly non-planar species.^{37a,c,41} We define ΔE_{intr} as the energy of the protonation of a free-base porphyrin, whose geometry matches that of the corresponding cation (i.e., mono- or dication). ΔE_{intr} encompasses all the electronic effects of the substituents as they affect the proton affinity. Thus, electron-donating substituents make ΔE_{intr} more negative, favoring protonation, while electron-withdrawing substituents do just the opposite. Importantly, the effect of a particular substituent on ΔE_{intr} may depend on the structure of the porphyrin. For example, electron-withdrawing and electron-donating *meso*-aryl groups would have a stronger effect on ΔE_{intr} if they would be more aligned and thus better conjugated with the tetrapyrrole macrocycle.⁴²

The structural term ΔE_{st} in turn consists of the two parts: $\Delta E_{\text{st}} = \Delta E_{\text{st}}(d) + \Delta E_{\text{st}}(c)$. $\Delta E_{\text{st}}(d)$ represents the energetic cost of the protonation-induced distortion of the macrocycle relative to its free-base conformation. Since non-planar distortion destabilizes the macrocycle,⁴² it disfavors the protonation ($\Delta E_{\text{st}}(d) > 0$). The second term, $\Delta E_{\text{st}}(c)$, is related specifically to *meso*-aryl-substituted porphyrins, for which it has been shown⁴² that a saddling distortion is coupled to the rotation of the *meso*-aryl groups relative to the macrocycle mean plane. This rotation increases the conjugation between the *meso*-aryls and the macrocycle π -system, stabilizing porphyrin cations⁴³ and thus favoring protonation ($\Delta E_{\text{st}}(c) < 0$). Notably, *meso*-aryl substituents always prefer to align with the macrocycle plane; however, starting from a certain angle their rotation begins to force the macrocycle out of planarity, thereby costing energy. The fact that tetraphenylporphyrins, such as tetraphenylporphyrin (TPP), remain planar as free-bases suggests that the benefit of the conjugation ($\Delta E_{\text{st}}(c)$) is not large enough to overcome the structural penalty associated with distortion, that is, $|\Delta E_{\text{st}}(c)| < |\Delta E_{\text{st}}(d)|$.⁴⁴

Considering the above formalism, it is possible now to contemplate the porphyrin protonation reaction as a sequential process (Figure 6). At first, the free-base (fb) is distorted to match the geometry of the monocation, forming the hypothetical species fb-mc. This process raises the energy by the structural penalty term $\Delta E_{\text{st}}(1)$. A proton is then added to fb-mc, producing the monocation (mc) and lowering the energy by $\Delta E_{\text{intr}}(1)$. In the second step, the monocation is first distorted to match the geometry of the dication (dc), producing the hypothetical

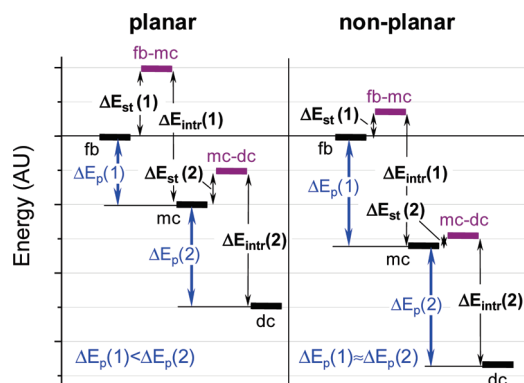


Figure 6. Energy diagram depicting two sequential protonations of planar (left) and non-planar (right) porphyrins.

species mc-dc and raising the energy by $\Delta E_{\text{st}}(2)$, followed by the addition of the second proton and release of $\Delta E_{\text{intr}}(2)$.

Assuming the intrinsic proton affinities ΔE_{intr} (for example $\Delta E_{\text{intr}}(1)$) are equal for the planar and non-planar porphyrins under comparison, it is now easy to see why the protonation of non-planar macrocycles is more energetically favorable. The attachment of protons to already non-planar free-bases requires less extra-distortion, and, therefore, it is associated with a smaller structural penalty term $\Delta E_{\text{st}}(d)$. At the same time, the component associated with the energetic benefit of increased conjugation ($\Delta E_{\text{st}}(c)$), which in principle may be larger for planar porphyrins, is still smaller in its absolute value than $\Delta E_{\text{st}}(d)$ (see above). Thus, we first of all can conclude that all other terms being equal, higher basicity of non-planar porphyrins is due to the lesser energetic penalty associated with protonation-induced distortion of the macrocycle: $\Delta E_{\text{st}}(\text{planar}) > \Delta E_{\text{st}}(\text{non-planar})$.

The second conclusion from the diagram in Figure 6 is related to the existence of the monocation of TCHpH, which is an interesting and unusual property. In the overwhelming majority of cases two sequential protonations of porphyrins are inseparable, and with only rare exceptions^{14–18} porphyrin monocations (monoacids) are extremely elusive species. To rationalize this behavior we recall that normally the attachment of the first proton to a neutral species disfavors the second protonation because of the electrostatic repulsion between the monocation and the second proton, that is, $\Delta E_p(2) > \Delta E_p(1)$. This leads to a difference in protonation pK_a 's and makes detection of the monocations feasible in the pH range between these pK_a 's. However, in porphyrins the situation is different presumably because of the cooperativity effect. In general, cooperativity entails a mechanism by which the first protonation assists the second protonation, thereby reducing the difference between $\Delta E_p(1)$ and $\Delta E_p(2)$ and, in extreme cases, making $\Delta E_p(2)$ more negative than $\Delta E_p(1)$, that is, $\Delta E_p(2) < \Delta E_p(1)$. The latter situation has been computationally demonstrated, for example, for the case of TPP,¹⁷ and it is shown in Figure 6. In general, the smaller the difference $[\Delta E_p(2) - \Delta E_p(1)]$, the less monocation is present in the mixture under equilibrium.

The cooperative enhancement of the second protonation in porphyrins can be explained by the structural effect: the protonation of the first imine nitrogen

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(44) For example, as calculated in ref 42 for TPP, bending of the free-base to match the saddling angle of the dication ($\varphi = 30^\circ$), while keeping the *meso*-phenyl rings as in their free-base orientation (tilt angle $\theta = 60^\circ$), is associated with the energetic penalty $\Delta E_{\text{st}}(d) \sim 13$ –14 kcal/mol. At the same time, rotation of the phenyls from $\theta = 60^\circ$ to $\theta = 30^\circ$, to match the orientation in the dication, brings back $\Delta E_{\text{st}}(c)$ of only ~ 4 –5 kcal/mol, making up the total structural penalty term $\Delta E_{\text{st}} \sim 9$ kcal/mol.

(45) ΔE_p 's were calculated as gas-phase uncorrected electronic free energies. Use of ΔE_p 's facilitates comparison between the energies of equilibrium structures (free-bases, monocations, and dications) and the energies of hypothetical non-equilibrium structures (fb-mc and mc-dc in Figure 6), for which thermochemical data could not be obtained and Gibbs free energies could not be calculated.

induces already strong enough deviation of the macrocycle from planarity, so that the attachment of the second proton requires much less structural deformation, reducing the associated structural penalty term $\Delta E_{st}(2)$. This scenario is illustrated in Figure 6. The intrinsic proton affinities for the first and second protonations, $\Delta E_{intr}(1)$ and $\Delta E_{intr}(2)$, were considered equal to more clearly show that the structural effect alone may be responsible for more negative $\Delta E_p(2)$ than $\Delta E_p(1)$. Most relevant to our discussion is the fact that a decrease in $\Delta E_{st}(1)$ and $\Delta E_{st}(2)$, which occurs in non-planar porphyrins, leads to an increase in the difference [$\Delta E_p(2) - \Delta E_p(1)$]. As a result, cooperativity decreases, and the monocation becomes more readily observable.

Because of electrostatic and other reasons, intrinsic proton affinities $\Delta E_{intr}(1)$ and $\Delta E_{intr}(2)$ in real systems are most certainly not the same. For example, the values of ΔE_{intr} may be affected significantly by counterions. The effect of counterions on the detectability of porphyrin monocations has been discussed in the literature.¹⁷ Nonetheless, the above conclusions regarding the structural effect should hold true as long as ΔE_{intr} themselves are not strongly influenced by planar deformations. The limitations of this assumption will have to be evaluated separately, but the calculations performed so far on model porphyrins (see below) suggest that ΔE_{intr} 's indeed depend only slightly on the degree of non-planar deformations.

To apply the described structural model to interpret the experimentally observed properties of TCHpH, we performed calculations on two model systems: planar tetraphenylporphyrin (TPP), which has been modeled previously,^{17,42,43} and highly non-planar Ph₄TCHP (Chart 1, R = H), which is a close analogue of the core in TCHpH. The σ -electron donor effect of tetracyclohexeno-rings in Ph₄TCHP could be estimated by comparing proton affinities of unsubstituted porphine (P) and *meso*-unsubstituted tetracyclohexenoporphyrin (TCHP), since in the absence of *meso*-aryl substituents tetracyclohexeno-groups do not induce structural deformations (Supporting Information, S6). Unsubstituted *meso*-phenyl groups were chosen in this model study because regardless of their orientation with respect to the macrocycle, orthogonally or at an angle, they presumably do not significantly affect intrinsic proton affinities (ΔE_{intr}). Of course, the electron-withdrawing carboxyl groups in *meso*-aryl rings of TCHpH decrease its intrinsic proton affinity compared to that of Ph₄TCHP. However, this effect is not structural, and it is apparently much weaker than the structural effect, since TCHpH still exhibits higher pK_a 's than the majority of tetraarylporphyrins.⁷

The calculations were performed by the DFT methods using B3LYP/6-31G(d) model chemistry (Supporting Information, S6). All structures were analyzed by the Normal Mode Structural Decomposition (NSD) method,²⁴ which provides a convenient tool for quantification of porphyrin distortions. The difference in the total mean out-of-plane displacements (D_{oop}), as calculated by the NSD program, between free-base and protonated forms (ΔD_{oop}) was used as a measure of the macrocycle distortion induced by protonation (Supporting Information, S6).

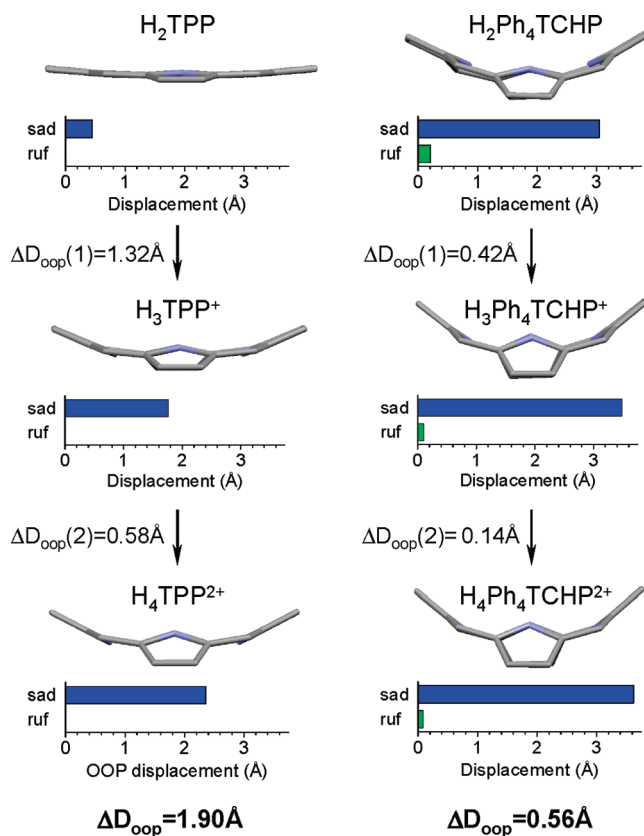


Figure 7. Structural changes upon protonation of TPP (left) and Ph₄TCHP (right). Side views of the tetrapyrrole macrocycles are shown. Structures of cations were calculated for the corresponding chloride salts. Charts graphically depict the degrees of two main out-of-plane (*oop*) displacements: saddling (*sad*) and ruffling (*ruf*). ΔD_{oop} designates the difference between the mean square *oop* displacements (sum of all non-planar modes) among differently protonated forms.²⁴

The results of the calculations are summarized in Figure 7 and Table 1.

It can be clearly seen that the overall protonation-induced deformation (Figure 7), going from free-bases to dications, is significantly larger for TPP ($\Delta D_{oop} = 1.90$ Å) than for Ph₄TCHP ($\Delta D_{oop} = 0.56$ Å); and that the magnitudes of the deformations, which are in all cases dominated by saddling, correlate with the calculated proton affinities (ΔE_p).⁴⁵ Total ΔE_p of Ph₄TCHP, for the two protonation steps combined, is ~ 18.8 kcal/mol more negative than that of TPP (Supporting Information, S6).⁴⁶ On the basis of the comparison between porphine (P) and unsubstituted TCHP, only ~ 5.5 kcal/mol of this difference is due to the σ -donor effect of the cyclohexeno substituents in Ph₄TCHP. The remaining ~ 13.3 kcal/mol is mostly accounted for by the difference between the structural terms: $|\Delta E_{st}(\text{TPP}) - \Delta E_{st}(\text{Ph}_4\text{TCHP})| \sim 10.2$ kcal/mol, suggesting that the decrease in the structural penalty indeed plays the central role in defining high proton affinity of Ph₄TCHP and presumably of other non-planar porphyrins.

(46) The gas-phase Gibbs free-energies calculated for the first (K_3) and second (K_4) protonations of TPP at 298 K, 17.8 and -54.6 kJ mol⁻¹ K⁻¹, respectively, were found to be higher than the earlier reported values, 1.5 and -66.7 kJ mol⁻¹ K⁻¹.¹⁷ The discrepancy is probably due to the difference in the model chemistries used in these two calculations.

Table 1. Energies^a (kcal mol⁻¹) for Gas Phase Protonations of TPP and Ph₄TCHP^b

	TPP	Ph ₄ TCHP	Ph ₄ TCHP _{corr} ^c
$\Delta E_p(1)$	-10.5848	-23.0291	-20.6710
$\Delta E_p(2)$	-26.8487	-33.1956	-29.9947
ΔE_p	-37.4335	-56.2247	-50.6657
$[\Delta E_p(2) - \Delta E_p(1)]$	-16.2639	-10.1665	-9.3237
$\Delta E_{st}(1)$	24.9494	16.4256	16.4256
$\Delta E_{st}(2)$	8.4672	6.8071	6.8071
$[\Delta E_{st}(2) - \Delta E_{st}(1)]$	-16.4822	-9.6185	-9.6185
$\Delta E_{intr}(1)$	-35.5342	-39.4547	-37.0966
$\Delta E_{intr}(2)$	-37.0966	-40.0027	-36.8018

^a All energies shown are derived from the pure uncorrected electronic energies of the corresponding components (see Supporting Information for details). ^b Cl⁻ was used as counterion in calculations of mono- and dications. ^c Energies in this column were corrected by the differences in the corresponding proton affinities of TCHP and P to account for the σ -donor effect of tetracyclohexenogroups in Ph₄TCHP.

As expected, after correction for the σ -donor effect of the cyclohexeno substituents in Ph₄TCHP, the intrinsic proton affinities ΔE_{intr} of TPP and Ph₄TCHP are indeed very similar. This indicates that the degree of non-planar distortion and the orientation of unsubstituted *meso*-phenyls with respect to the macrocycle, which are different in fb-mc and mc-dc structures of TPP and Ph₄TCHP, have only a minor effect on the intrinsic proton affinities. In other words, the intrinsic proton affinity of the porphyrin macrocycle is only weakly affected by its distortion.

The calculations confirmed that the overwhelming part of the non-planar deformation occurs with the first protonation, and that the absolute values of the distortions (ΔD_{oop} 's) in both protonation stages are significantly higher in the case of TPP (Figure 7). The associated structural terms $\Delta E_{st}(1)$ and $\Delta E_{st}(2)$ (Table 1) follow these trends, making the difference between the free energies of the first and second protonations $[\Delta E_p(2) - \Delta E_p(1)]$ substantially lower in the case of TPP, resembling the situation shown in Figure 6. This enables us to conclude that the relative ease of detectability of the monocation of TCHpH is due to the high non-planarity of its core porphyrin. Notably, one of the recent reports on porphyrin monocations also describes highly non-planar dodecasubstituted porphyrin.¹⁸

Several spectroscopic properties of TCHpH can also be rationalized using the described above structural model. First, we note that the red shift of the Soret band of TCHpH is smaller than the corresponding shifts typically observed for regular porphyrins. This is consistent with a smaller extra-distortion of the non-planar core of TCHpH induced by protonation, taking into account that non-planar deformations cause red shifts in the spectra of porphyrins.^{32,33} Second, the red-most Q-band of the TCHpH dication is shifted blue relative to the Q_x(0,0) transition of the free-base, unlike in regular porphyrins where the corresponding shift occurs in the opposite direction. Symmetrization of the porphyrin upon protonation causes its Q_x and Q_y bands to coalesce, and in the case of regular porphyrins, strong non-planar distortion causes the resulting band to move beyond the Q_x(0,0) band of the free-base. In the case of TCHpH the distortion is much smaller, and the band is red-shifted significantly less.

Calculations performed by the ZINDO/S method (Supporting Information, S8),⁴⁷ using the structures shown in Figure 7, were able to qualitatively reproduce the relative shifts of the bands of Ph₄TCHP dication versus the free-base, positions of Q_x and Q_y bands of the monocation versus those of the free-base, and the merging of the Q_x and Q_y transitions into the single Q-band of the dication. However, both the Soret and the Q-bands of the dication were predicted to be red-shifted relative to the bands of the free-base and the monocation, which does not agree with the experiment. Higher level calculations, such as performed recently on monocations of other tetraarylporphyrins,¹⁷ will be required to fully delineate the spectral features of TCHpH.

Conclusions

The newly developed dendritic probe TCHpH is a macro-molecular porphyrin-based pH-sensitive dye suitable for measurements in microcompartmentalized systems, including dynamic measurements of proton gradients. The probe consists of a polyfunctionalized non-planar porphyrin, modified with eight trifurcated Newkome-type dendrons. The latter together with peripheral PEG residues serve to increase the probe's solubility, control its size, and prevent its interactions with components of the measurement system. In the present version of the probe, PEG residues with an average molecular weight of 750 Da were used; however, larger probes can be easily constructed by modifying terminal groups on the dendrimer with larger methoxypolyethyleneglycol or methoxypolyethyleneglycol amine residues, which are readily available. Such alterations will not change the protonation pK_a's, but may appear instrumental in optimizing the probe's distribution in microcompartmentalized systems.

To rationalize acid–base properties of TCHpH we further developed a proposed earlier¹¹ structural model that relates electronic effects of macrocycle deformation to porphyrin basicity. The predictions of this model were validated by DFT calculations on a planar and non-planar porphyrin, a structural analogue of the core of TCHpH. One relevant prediction is the relative ease of observations of monocations of strongly out-of-plane distorted porphyrins, which was experimentally confirmed in the case of TCHpH. The unique monocation spectrum was identified in the spectra of the mixture of variously protonated forms by spectral deconvolution and could be partly interpreted in terms of the classic porphyrin electronic structure model.

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Supporting Information Available: Characterization data for the new compounds, derivation of expressions for fitting the spectroscopic data, details of DFT calculations, results of NSD analyses, additional spectroscopic data. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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