Reversible oxygen addition on a triplet sensitizer molecule: protection from excited state depopulation

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We demonstrate that photoactivated oxygen addition to diphenylanthracene moieties can be used as a tool for protection of porphyrin’s phosphorescence against oxygen quenching. Phosphorescent palladium(ii) tetrabenzozoporphyrin, covalently linked to four diphenylanthracene moieties, was synthesized and studied. Upon irradiation with ambient light or red laser in solution in air, addition of oxygen and formation of the corresponding endoperoxides were observed. Heating of the irradiated samples afforded the parent porphyrin material.

Introduction

Organic chromophores able to form triplet excited states upon absorption of light (triplet sensitizers) are used in various fields, e.g. electroluminescence,1 bioimaging and molecular sensing,2 photocatalytic organic reactions,3 and triplet–triplet annihilation photon upconversion (TTA-UC).4 Organic materials based on triplet sensitizers promise to make revolutionary transformations in solar energy conversion technologies by improving the light-harvesting of amorphous silicon solar cells5 and extending the infrared limit of oxygenic photosynthesis.6 However, compared to fluorophores, applications for triplet sensitizers are much less developed considering a substantially higher sensitivity of triplet states towards non-emissive deactivation processes. The most common process which leads to the loss of triplet excited state population is related to the presence of molecular oxygen in the corresponding samples. It involves a triplet energy transfer between an excited chromophore and oxygen, giving rise to singlet oxygen and the ground state of the chromophore.7

An efficient protection against oxygen quenching is essential for applications which include triplet excited state formation. Two general approaches for the protection are known: (1) passive protection, based on decreasing oxygen permeability e.g. by means of encapsulation in polymer films8 or nano- and micro-dendrimers9 and (2) active protection – by applying oxygen barrier properties leads to a substantial decrease of the TTA-UC efficiency, because in solid state environments the local mobility of the chromophores involved a significant decrease.14 Until recently the direct incorporation of dyes into polymers was not enough to provide the imperative requirements for effective and sustainable annihilation upconversion, that is, high local mobility and exhaustive oxygen protection of the UC-chromophores. Recently we succeeded in developing organic polyphosphate oxygen protection matrices15 and cellulose bioinspired oxygen protection films.16 However, these approaches either require a change in the sample’s architecture or affect the photophysical and chemical properties of a triplet sensitizer.

Herein we report a new strategy for protection of triplet excited state depopulation by quenching, relying on chemical modification of a triplet sensitizer molecule. It is based on binding the molecular oxygen, present in a sample, to specially designed structural subunits which do not affect the photophysical properties and allow the triplet sensitizer to act in an undisturbed manner further. This protection strategy is of sacrificial character, and is time-limited depending on the integral photon flux, applied to the sample. However, the starting triplet

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sensitizer material can be fully regenerated afterwards through oxygen release upon moderate heating. Furthermore, protective groups do not bind oxygen in its ground (triplet) state and the corresponding material is stable towards photooxidation in the course of synthetic procedures and purification. The protection is active only against singlet oxygen, thus only when sensitizer triplet states are formed. Therefore, the presence of an additional stimulus, i.e. optical excitation of the sensitizer, triggers the protection process.

In our recent work on the synthesis of tetraanthraporphyrins, we observed a photoactivated addition of up to four oxygen molecules per porphyrin molecule, leading to the formation of the corresponding endoperoxides. We proposed that such an addition can serve as a tool for preventing oxygen from quenching of porphyrin phosphorescence. However, in the case of tetraanthraporphyrins, the addition of oxygen leads to a drastic change in the optical properties, particularly to a blue-shift of absorption and emission bands by 200 nm due to a partial loss of conjugation in the π-system. Here we suggested to introduce anthracene subunits in meso-positions of the porphyrin in order to avoid alteration of the optical properties.

The concept is illustrated in Fig. 1 using an example of Pd-porphyrin bearing four anthracene subunits in meso-positions of the macrocycle. Pd-porphyrins are known to possess high intersystem crossing coefficient values and are widely used as triplet sensitizers. Alternatively, anthracenes are capable of reversible binding of singlet oxygen species. Upon selective excitation of the porphyrin chromophore, a triplet excited state is formed (step 1). In the presence of the molecular oxygen a quenching process takes place, leading to the ground state of the sensitizer and singlet oxygen (step 2). The key point is step 3 where binding of the singlet oxygen to the attached hydrocarbon via the Diels–Alder-type process takes place. We proposed that complete binding to all of four hydrocarbon subunits can be achieved (shown as step 4 in Scheme 1) and the corresponding (O₂)₄ adduct can be obtained, otherwise adducts with a variable number of bound oxygens are produced. Consumption of oxygen results in partial or complete termination of the quenching process that provides emissive relaxation of newly formed triplet states of the Pd-porphyrin core. Due to the reversibility of the oxygen addition to anthracenes, the starting material can further be regenerated by means of heating (step 5).

Although many polyaromatic hydrocarbons, e.g. naphthalenes, 2-pyridone or rubrenes, are known to bind oxygen in a reversible manner, anthracenes are most suitable to be used for oxygen protection of the phosphorescence. First, the absorption bands of the corresponding anthracenes do not intersect with porphyrin absorption that allows monitoring of the protection by absorption spectroscopy. Second, anthracene endoperoxides decompose only upon heating above 100 °C whereas naphthalene and 2-pyridone endoperoxides release oxygen even at room temperature. Finally, anthracene provides a scaffold that allows straightforward generation of suitable substitution patterns.

The introduction of such type of triplet sensitizer into the arsenal of materials used in the relevant areas of research would open new opportunities for the control of the corresponding photophysical processes. Particularly, it provides an alternative way to oxygen protection of phosphorescent samples or to develop new molecular “singlet oxygen reservoirs” for oxygen storage and thermal release. In this paper we report the
synthesis and properties of phosphorescent palladium(II) porphyrin, bearing substituted anthracenes in meso-positions, which is capable of photoactivated oxygen addition. Anthracene groups preserve the phosphorescence properties of porphyrin even in the oxygen rich environment. Furthermore, thermal-mediated reversibility of the oxygen binding was demonstrated.

### Results and discussion

#### Synthesis

Our attempts towards the synthesis of the target molecule were firstly focused on β-unsubstituted porphyrin with anthracene groups directly attached in meso-positions.

According to literature data, the conventional method of porphyrin chemistry – Lewis acid catalyzed condensation of pyrrole with the corresponding 10-phenyl-anthracene-9-carboxaldehyde 1 – results in very low preparative yields. An alternative method based on pyrrole-carbinol tetramerization is being commonly applied for such type of substrates. Following this approach, the starting 9-phenylanthracene 2 was brominated with Br₂ to give 9-bromo-10-phenylanthracene 3. It was then subjected to the reaction with n-butyllithium to give the corresponding organolithium compound 4 which was further introduced into the reaction with pyrrole-2-carboxaldehyde (Scheme 1). The obtained pyrrole-carbinol 5 was used for acid-catalyzed tetramerization and delivered the corresponding porphyrin 6 with reasonable yield (15%). Subsequent insertion of palladium was achieved by reflux in benzonitrile. Unfortunately the obtained product 6-Pd was found to possess too low solubility for adequate characterization and study of oxygen addition.

Generally, for large conjugated aromatic systems, solubility is ensured by introduction of suitable side chains into the aromatic moiety, e.g. alkyl and alkoxy groups. However, the introduction of electron-donating groups on anthracenes is known to have a pronounced effect on the rates of oxygen binding and decomposition of the endoperoxide. In some cases it results in the irreversibility of the oxygen addition. Thus in order to optimize the sensitizer structure we decided to modify the porphyrin core instead of the anthracene subunit. It was shown that β-substitution of the porphyrin macrocycle, especially along with introduction of meso-aryl substituents leads to a strong distortion of the macrocycle due to steric repulsion. This in turn results in an improved solubility. Following this approach we decided to introduce two additional structural features: (1) a phenylene bridge between porphyrin and anthracene subunits and (2) annelated cyclohexane rings at β-positions of the porphyrin macrocycle. Such a modification also allows for shifting of absorption and emission bands into the red or far red region, the latter by means of aromatization of annelated rings.

9-Bromo-10-phenylanthracene 3 was subjected to Suzuki-coupling with formylboronic acid giving the corresponding aldehyde 8. Its condensation with 4,5,6,7-tetrahydrosioindole 9 provided the corresponding porphyrin 10 (Scheme 2), which was then metallated with bis(benzonitrile)palladium(II) in boiling benzonitrile. Further aromatization of porphyrin 10-Pd into the corresponding tetrabenzo-porphyrin 11-Pd was achieved by reflux with an excess of DDQ in toluene. Complete aromatization of all four annelated rings was achieved without significant by-product formation. Porphyrin 11-Pd was found to be well-soluble in the common solvent (chlorinated hydrocarbons, THF, toluene) and was unambiguously characterized by NMR and mass spectroscopy (see ESI†).

#### Oxygen addition and release

As is shown in Fig. 2, porphyrin 10-Pd possesses characteristic absorption in the region of 300–400 nm, which correspond to the anthracene subunits. When a solution of 10-Pd was kept under daylight for several days or irradiated with the green line of the HeNe laser (λ = 543 nm, broad beam), disappearance of the anthracene absorption was observed (Fig. 2, red line) giving instead a shapeless background absorption. On the other hand, spectral features corresponding to the porphyrin macrocycle – Soret (430 nm) and Q-bands (540 and 570 nm) – remained unchanged. Light irradiation of the same solution prepared and sealed in a glove-box (<1 ppm oxygen) showed no change in absorption.

The mass spectrum of the irradiated solution showed a single peak with a mass of 2072 Da, corresponding to an adduct of 10-Pd with four oxygen molecules (Scheme 3).
A similar process with compound 11-Pd was monitored by UV-Vis spectroscopy. Its solution (4 × 10^{-5} M) was irradiated with the red line of HeNe laser (λ = 633 nm, broad beam) at an intensity of only 3 mW cm^{-2}. The laser beam was set to cover the whole cuvette. The complete disappearance of the anthracene absorption in the region of 300–400 nm was observed within 9 h (Fig. 3). Mass spectra of the obtained samples showed a series of (M+n×32) peaks indicating the formation of intermediate products bearing 1, 2, 3 or 4 oxygen molecules attached. At 3 mW cm^{-2} the addition reaction was complete in 9 h as is evidenced by the absence of spectral change and a sole peak of 2056 Da in the mass spectrum corresponding to the adduct with four oxygen molecules (Scheme 4). Exposure of the solution for longer times of irradiation (up to 40 h) did not result in the change in absorption spectra or origin of other peaks in the mass spectrum.

Fig. 2 Absorption spectrum of 10-Pd and its change after light irradiation of air-saturated toluene solution (green-line HeNe laser, 3 μW cm^{-2}, 10 h) (A); the corresponding mass spectra of 10-Pd (B, black trace) and the product of irradiation (B, red trace).

Fig. 3 Change in absorption upon irradiation of 4 × 10^{-5} M toluene solution of 11-Pd using 633 nm laser (3 μW cm^{-2}) (A); the corresponding change in mass spectra (B): 11-Pd (black trace), after 5 h of irradiation (blue trace), and after 10 h of irradiation (yellow trace). Note: saturation of porphyrin absorption is due to high concentration of the solution taken for the experiment, essential for monitoring the relatively weak anthracene absorption.

Scheme 3 Photosensitized oxygen addition to porphyrin 10-Pd.

Scheme 4 Photosensitized oxygen addition to porphyrin 11-Pd.
To further support the idea of photosensitized oxygen addition on anthracene moieties in 11-Pd we performed a similar experiment using a mixture of 9,10-diphenylanthracene and palladium tetrabenzytenzoporphyrin in a molar ratio of 4 : 1. Under the same conditions (concentration, laser intensity) a similar spectral transformation was observed by UV-Vis spectroscopy on a timescale of 6 h (see ESI† Fig. S14).

The release of bound oxygen molecules was achieved when solution of the adduct 13-Pd was concentrated to a solid and then heated at 100–110 °C in vacuum (1 mbar) for 3–5 h. A complete recovery of anthracene absorption (with respect to 11-Pd) as well as a perfect overlap with the original spectrum of 11-Pd were observed. The mass spectrum showed a 1928 Da mass peak (11-Pd) as well as a perfect overlap with the original spectrum of 11-Pd). The sample of 11-Pd was isolated. In order to investigate their emission properties a series of 11-Pd solutions were illuminated (633 nm, 3 μW cm⁻²) at different time intervals (1–9 h). The obtained samples were concentrated to solids, transferred into the glove-box (<1 ppm of oxygen) and dissolved in toluene to prepare a set of solutions of equal concentration for phosphorescence measurement. No significant difference in the phosphorescence intensity between starting 11-Pd and irradiated samples was observed (Fig. 5). Only a moderate decrease of the emission intensity (approximately 25%) takes place upon addition of oxygen molecules to all four anthrancenes. This is probably due to a decrease in the phosphorescence quantum yield upon increasing molecule’s complexity usually resulting in an increase of non-radiative decay rates.

The fact that adducts with a different number of bound oxygen molecules possess similar emission properties enables us to use the photo-activated oxygen addition to enhance the phosphorescence signal intensity of oxygen contaminated (e.g. as a result of leaking) samples. Under laser irradiation during a short time (minutes or less, depending on excitation intensity), oxygen molecules are bound to the sensitizer, performing local and real-time “deoxygenation” of the sample. Due to the decrease of the quencher concentration, the phosphorescence efficiency increases.

To demonstrate such an effect, a solution of 11-Pd was prepared and sealed in an atmosphere containing 100 ppm of oxygen. The phosphorescence spectrum registered has shown a peak intensity of 2160 cps (Fig. 6, black line). Then the cuvette was placed under a laser beam with 250 μW cm⁻² intensity for a period of 10 min. The phosphorescence spectrum registered after irradiation has shown 100% increase of the peak intensity (Fig. 6, red line). Obviously, the increase of the signal intensity is due to the binding of residual amounts of oxygen present in the sample.

Moreover, an enhancement of the phosphorescence originating from porphyrin 11-Pd can be achieved in a local area (spot diameter d = 400 μm) of excitation without preliminary irradiation of the whole sample. In the course of the measurement, the same laser beam being used for the excitation of phosphorescence can...
system will be reported in due course.

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a few seconds, if the excitation intensity is lower, the necessary
time is substantially longer (Fig. 7, the black line). The experi-
mental atmosphere was contaminated by 100 ppm of oxygen.

Owing to the deoxygenation ability, molecules such as 11-Pd

can become promising “self-healing” sensitizers for the pro-
cess of TTA-UC, which is known to crucially depend on the

content. In order to confirm the suitability of 11-Pd as a

sensitizer for TTA-UC, it was mixed with perylene as an emitter

in an inert atmosphere (<1 ppm of oxygen) and then used for
the generation of the upconversion fluorescence. Upon the red
excitation at 638 nm, along with some residual phosphores-
cence of the sensitizer, a blue upconverted emission of perylene
in the region 430–550 nm with the reasonable quantum yield
(~3%) was observed (Fig. 8). A corresponding study devoted to
the effect of oxygen on the upconversion efficiency of the
system will be reported in due course.

Conclusions

While in many studies devoted to phosphorescence and triplet
excited state dynamics the problem of oxygen quenching is being
solved by physical or chemical protection of the sample from
oxygen contamination, the development of “self-protective” triplet
sensitizer molecules represents an almost unexplored approach.

In this work, palladium(II) porphyrin bearing four anthracene
subunits in meso-positions was synthesized. Optimization of struc-
tural features delivered a material with high solubility and strong
near-IR absorption. It was found to possess oxygen-binding proper-
ties upon irradiation with either daylight or lasers. There was no
effect on the absorption features of porphyrin, and only a weak
decrease of phosphorescence intensity was observed during addition
of oxygen. Such properties enable us to perform partial or complete
doxygenation of oxygen-contaminated samples, thus enhancing
their phosphorescence intensity. Two regimes of deoxygenation were
explored: a conventional one – oxygen scavenging of the whole
sample, and second – local, and real-time oxygen scavenging,
suitable for sensing applications. The starting sensitizer material
can be regenerated by means of heating in vacuum. A palette of
observed properties makes the material promising for application as
phosphorescent sensing probes and a self-protecting sensitizer for
the process of TTA-UC.

Although such a protection strategy is limited by the capacity of
the sensitizer to bind no more than four oxygen molecules, it may
have a potential in applications which demand very low oxygen
concentration levels. For example, most of the materials comprising
the OLEDs suffer from degradation effects due to the presence of
environmental oxygen and water, as both compounds can penetrate
into the device. Much research and effort have been put into
fabrication methods and appropriate device encapsulation to help
mitigate these environmental effects. Thus the demonstrated
oxidation protection strategy on a molecular level might become
complementary to those already developed.

Experimental

4,5,6,7-Tetrahydroisoindole and pyrrole-2-carboxaldehyde were prepared according to published synthetic protocols.
9-Phenylantracone, 4-formylphenyboronic acid, bis(benzonitrile) palladium(u) chloride, and DDQ, were purchased from Sigma-Aldrich. The handling of all air/water sensitive materials was carried out using standard high vacuum techniques. All solvents and reagents were obtained from commercial sources and used as received. Extra dry DMF, toluene and THF were purchased from Sigma-Aldrich. Triethylamine and DCM were distilled from CaH₂. Where mixtures of solvents were used, ratios are reported by volume. Column chromatography was carried out on silica gel 60 at normal pressure. NMR spectra were recorded onBruker DPX 250, Bruker AC300 NMR and Bruker Avance 500 spectrometers, with the solvent proton or carbon signal as an internal standard. Elemental analysis was carried out using a Foss Heraeus Vario EL. Electronic absorption spectra were recorded on a PerkinElmer Lambda 25 instrument. MALDI-TOF spectra were obtained on a Bruker Reflex spectrometer III instrument using dithranol as a matrix. HR ESI spectrometry was performed on a QToF Ultima 3 Fa. Melting points were determined on a Büchi hot stage apparatus and are uncorrected. The emission spectra and phosphorescence lifetime of substances were measured using our home-built spectrometer.33 For the excitation and irradiation of studied materials a temperature stabilized diode laser with λ = 638 nm (QL63H5S, average power 20 mW) was used. In order to irradiate the samples in the quartz cuvettes (2 × 1 × 1 cm), the laser beam size was increased using a negative lens, resulting in the average intensity of 5 µW cm⁻² on the entrance surface of the cuvette for 638 nm laser. The suppression of the excitation radiation during phosphorescence spectrum measurements was done using a notch filter NF 03-633U-25 or NF03-532E-25 (Semrock Inc.).

4-(10-Phenyl-anthracen-9-yl)-benzaldehyde 8

The title compound was prepared following a modified literature procedure.34 9-Phenylantracone (3.5 g, 13.8 mmol) was dissolved in acetic acid (150 mL). The solution was heated to 65 °C under nitrogen, and bromine (2.2 g, 13.8 mmol) was added dropwise over a period of 10 min. The reaction mixture was allowed to cool to room temperature and evaporated in vacuum. The residual solid was recrystallized from ethanol to give 4-(10-phenyl-9-bromoanthracene (4.59 g, 95%) as yellow crystals with m.p. 154–155 °C (lit. 153–155 °C). ¹H NMR (250 MHz, CD₂Cl₂) δ 8.64–8.57 (m, 2H), 7.69–7.55 (m, 7H), 7.44–7.35 (m, 4H).

Porphyrin 6

Butyllithium (1.6 M solution in hexane, 3.1 mL, 5 mmol) was added dropwise to a mixture of 10-phenyl-9-bromoanthracene (1.5 g, 4.5 mmol) in diethyl ether (30 mL) and the mixture was stirred at room temperature for 30 min. Then a solution of pyrrole-2-carboxaldehyde (0.285 g, 3 mmol) in diethyl ether (5 mL) was added, the resulting solution was stirred for 1 h and poured into water (100 mL). The organic phase was separated, washed with water (3 × 10 mL) and evaporated in vacuum. The residue was then dissolved in a mixture of toluene (15 mL) and propionic acid (5 mL). The reaction mixture was stirred for 3 h at 100 °C then allowed to cool to room temperature. Solvents were evaporated in vacuum, the residue was then redissolved in CHCl₃ and passed through a filter with silica. The first red-brown fraction was collected and the solvents were removed. The crude product was recrystallized from DCM–methanol to give porphyrin 6 (0.148 g, 15% with respect to pyrrole-2-carboxaldehyde) as a red-brown solid. ¹H NMR (300 MHz, CD₂Cl₂) δ 8.40–8.34 (m, 8H), 7.96 (d, J = 8.9 Hz, 8H), 7.78–7.65 (m, 20H), 7.40 (ddd, J = 8.9, 6.0, 1.6 Hz, 8H), 7.29–7.17 (m, 16H). ¹³C NMR (75 MHz, CD₂Cl₂) δ 147.63, 141.88, 139.16, 135.74, 132.58, 131.87, 130.50, 130.22, 129.22, 128.57, 127.97, 127.77, 127.64, 125.99, 125.87, 119.50. UV/vis (toluene) λmax (log ε): 430 (5.54), 519 (4.58), 551 (4.05), 592 (4.09), 649 (3.73). MALDI-TOF: m/z found 1319.4076, calcd for [M⁺] C₁₀₀H₆₂N₄ 1319.5008.

Porphyrin 10

4,5,6,7-Tetrahydrosoindole (0.3 g, 2.48 mmol) was dissolved in CH₂Cl₂ (250 mL) freshly distilled from CaH₂, and 4-(10-phenyl-anthracen-9-yl)-benzaldehyde (0.888 g, 2.48 mmol) was added. The mixture was stirred under nitrogen for 10 min in the dark at room temperature. BF₃·Et₂O (0.035 g, 0.248 mmol) was added in one portion, and the mixture was stirred for an additional 2 h. DDQ (0.422 g, 1.86 mmol) was added followed by additional stirring for 2 h in aqueous Na₂SO₄, dried over Na₂SO₄ and concentrated in vacuum. The residue was purified on a silica gel column (eluent CH₂Cl₂, then CH₂Cl₂–AcOH, green band collected). Additional purification by repetitive precipitation from CH₂Cl₂–AcOH (10:1) with hexane delivered the product (0.279 g, 23%) as a dark-green powder. ¹H NMR (250 MHz, CD₂Cl₂) δ 10.20 (s, 1H), 8.15 (d, J = 8.1 Hz, 2H), 7.76–7.68 (m, 4H), 7.66–7.57 (m, 5H), 7.50 (ddd, J = 7.6, 1.7 Hz, 2H), 7.42–7.33 (m, 4H).

4-(10-Phenyl-anthracen-9-yl)-benzaldehyde 8

The title compound was prepared following a modified literature procedure.35 A mixture of 10-phenyl-9-bromoanthracene (2.5 g, 7.5 mmol), 4-formylphenyboronic acid (1.35 g, 9 mmol), tetrakis(triphenylphosphine)palladium (0.433 g, 0.375 mmol), K₂CO₃ (2.48 g, 18 mmol), benzene (100 mL), ethanol (20 mL) and water (40 mL) was refluxed under nitrogen for 24 h. The organic phase was separated, washed with brine, and dried over Na₂SO₄. The solvent was evaporated in vacuum and the remaining solid was purified on a silica gel column using the EtOAc–pentane (1:5) mixture as an eluent. Fractions containing the product (determined by TLC) were concentrated to give the aldehyde (2.36 g, 88%) as yellow crystals with m.p. 234–236 °C. ¹H NMR (250 MHz, CD₂Cl₂) δ 128.57, 127.97, 127.77, 127.64, 125.99, 125.87, 119.50. UV/vis (toluene) λmax (log ε): 400 (5.07), 430 (5.54), 519 (4.58), 551 (4.05), 592 (4.09), 649 (3.73). MALDI-TOF: m/z found 1319.4076, calcd for [M⁺] C₁₀₀H₆₂N₄ 1319.5008.
Porphyrin 10-Pd

Bis(benzonitrile)palladium(u) chloride (0.078 g, 0.204 mmol) and triethylamine (0.101 g, 1 mmol) were added to a solution of porphyrin 10 (0.2 g, 0.102 mmol) in benzonitrile (10 mL), and the mixture was heated at 180 °C for 1–2 h. The conversion was monitored by UV-vis spectroscopy (solvant CH2Cl2-TFA) and was considered complete after the absorption band of the product (0.06 g, 60%) as blue-green crystals. 1H NMR (300 MHz, CD2Cl4) δ 7.76 (d, J = 7.4 Hz, 8H), 7.86 (d, J = 8.8 Hz, 8H), 7.86–7.51 (m, 52H). 13C NMR (126 MHz, CD2Cl4) δ 141.95, 141.16, 140.53, 139.08, 138.87, 137.66, 137.31, 136.92, 134.32, 131.50, 130.08, 128.87, 128.26, 128.05, 127.38, 127.09, 126.82, 126.15, 124.94, 124.20, 118.96, 29.42, 23.87. UV/vis (toluene): 340 (4.29), 357 (4.37), 376 (4.48), 430 (5.11), 538 (4.21), 572 (4.09). HRMS (ESI): m/z found 1927.5749, calcd for [M+] C140H84N4Pd 1927.5764.

Porphyrin 11-Pd

Porphyrin 10-Pd (0.1 g, 0.051 mmol) was dissolved in THF (10 mL), DDQ (0.187 g, 0.82 mmol) was added, and the mixture was refluxed for 20–40 min. During refluxing the color changed from red-brown to deep green. The mixture was allowed to cool, diluted with CH2Cl2, washed with 10% aqueous solution of Na2SO3, with brine, and dried over Na2SO4. The solvent was removed in a vacuum, and the remaining solid was purified on a silica gel column using CH2Cl2 as an eluent. The first dark-green fraction was collected. The solvent was evaporated and the residue was recrystallized from CH2Cl2–Et2O to give the product (0.06 g, 60%) as blue-green crystals. 1H NMR (300 MHz, CD2Cl4) δ 8.48 (d, J = 7.4 Hz, 8H), 8.68 (d, J = 8.4 Hz, 8H), 7.94 (d, J = 8.8 Hz, 8H), 7.86–7.51 (m, 52H). 13C NMR (126 MHz, CD2Cl4) δ 141.03, 139.63, 139.75, 138.35, 138.24, 137.65, 136.37, 133.99, 133.17, 132.52, 131.30, 129.97, 129.70, 128.46, 127.57, 127.35, 126.70, 125.60, 125.23, 124.42, 120.19, 117.93. UV/vis λmax(toluene)/nm (log ε): 340 (4.29), 357 (4.4), 375 (4.53), 395 (4.54), 446 (5.33), 628 (4.89). HRMS (ESI): m/z found 1943.7042, calcd for [M+] C140H100N4Pd 1943.7016.

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Notes and references


4 (a) A. Turshatov, D. Busko, S. Baluschev, T. Miteva and K. Landfester, New J. Phys., 2011, 10, 083035; (b) C. Wohnhaas, A. Turshatov, V. Mailaender, S. Lorenz,


27 (a) N. K. S. Davis, M. Pawlacki and H. L. Anderson, *Org. Lett.*, 2008, 10, 3945; (b) N. K. S. Davis, A. L. Thompson and...


