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

Mikhail A. Filatov,⁎a,b Ernesta Heinrich,a Dmitry Busko,a Iliyana Z. Ilieva,c Katharina Landfester*a and Stanislav Baluschevac,d

We demonstrate that photoactivated oxygen addition to diphenylanthracene moities can be used as a tool for protection of porphyrin’s phosphorescence against oxygen quenching. Phosphorescent palladium(ii) tetrabenzo porphyrin, 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 electronic 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-9 and micro-carriers,10 or incorporation into supramolecular complexes11 or dendrimers12 and (2) active protection – by applying oxygen scavenging species.13 The problem of triplet excited state quenching is exceptionally important in the case of the TTA-UC process. Both the sensitizer and emitter ensembles can transfer triplet energy to the molecular oxygen and be depopulated, leading to the aging of samples and loss of quantum efficiency. Straightforward embedding of UC-active substances into inert polymer films with high 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
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 \( \pi \)-system. Here we suggested to introduce anthracene subunits in \( \text{meso} \)-positions of the porphyrin in order to avoid alteration of the optical properties.

The concept is illustrated in Fig. 1 using an example of \( \text{Pd-porphyrin} \) bearing four anthracene subunits in \( \text{meso} \)-positions of the macrocycle. \( \text{Pd-porphyrins} \) 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 \( (\text{O}_2)_4 \) 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 \( \text{Pd-porphyrin core} \). 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-tetrahydroisoindole 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 tetrabenzoporphyrin 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 × 10⁻² M) was irradiated with the red line of HeNe laser (λ = 633 nm, broad beam) at an intensity of only 3 μW 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 μW 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.
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 tetraphenyltetrabenzoquinophyrin 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 parent peak. Thus a completely reversible oxygenation–deoxygination process takes place for 11-Pd.

It should be mentioned that the decomposition temperature (and time) of the adduct depends on the substitution pattern of the aromatic subunits. In our case the parameters of oxygen release match those previously described for 9,10-diphenylanthracene decomposition. However, modification of the starting materials would allow varying decomposition parameters. Particularly, it can be made irreversible (by means of introducing one or two alkoxy-groups into the parent anthracene derivative). Alternatively, the temperature of endoperoxide decomposition can be adjusted to 30–40 °C (by using naphthalene derivatives as oxygen traps, instead of anthracenes).

**Optical properties: protection of phosphorescence against oxygen quenching**

Absorption and emission spectra of 11-Pd are shown in Fig. 4. Its optical properties are very similar to those of tetraphenyltetrabenzoquinophyrin-Pd (see ESI†). The sample of 11-Pd prepared in a glove-box (containing <1 ppm oxygen) exhibits strong phosphorescence centered at 796 nm with a quantum yield of 0.18 (with respect to Ph4TBPPd in toluene (0.21)). No phosphorescence was observed in the sample open to air. The triplet excited state lifetime was measured to be 217 μs (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.

the effect of oxygen on the upconversion efficiency of the sensitizer, a blue upconverted emission of perylene excitation at 638 nm, along with some residual phosphorescence of the sensitizer for TTA-UC, it was mixed with perylene as an emitter in an inert atmosphere (toluene, 4 \( \times \) 10\(^{-5}\) M, 633 nm, 250 \( \mu \)W cm\(^{-2}\)) (red line).

A new material was synthesized, palladium(II) porphyrin bearing four anthracene subunits in meso-positions, and using 6 \( \times \) 10\(^{-5}\) M (toluene) perylene as an emitter. Quantum yield of UC at an excitation intensity of \( \sim 1\) W cm\(^{-2}\) is 3%. Inset: a photo of the upconversion sample being irradiated with 638 nm laser.

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 structural features delivered a material with high solubility and strong near-IR absorption. It was found to possess oxygen-binding properties 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 deoxygenation 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 oxygen protection strategy on a molecular level might become complementary to those already developed.

Experimental

4,5,6,7-Tetrahydroisoindole\(^{31}\) and pyrrole-2-carboxaldehyde\(^{32}\) were prepared according to published synthetic protocols.
9-Phenylanthracene, 4-formylphenylboronic 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 on Bruker 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. Elemental analysis was carried out using a Foss Heraeus Vario EL. 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 ie = 638 nm (QL63H5S, average power 20 mW) was used. In order to irradiate the samples in the quartz cuvettes (2 x 1 x 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.).

10-Phenyl-9-bromoanthracene 3

The title compound was prepared following a modified literature procedure.34 9-Phenylanthracene (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 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 x 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, 138.53, 138.39, 137.31, 136.83, 135.96, 132.47, 131.91, 130.62, 130.46, 129.10, 128.23, 127.76, 127.30, 126.13, 125.80, 118.72, 113.55, 103.63, 96.81, 78.01. MALDI-TOF: m/z found 1319.4076, caled for [M+H]+ C140H103N4 1319.4076.

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-formylphenylboronic 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, C2D2Cl2) 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 (dd, J = 7.6, 1.7 Hz, 2H), 7.42–7.33 (m, 4H).

Porphyrin 10

4,5,6,7-Tetrahydroisoindole (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-EtO (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 Na2SO3, 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 8.88 (d, J = 7.9 Hz, 2H), 8.10–8.01 (m, 4H), 7.84–7.77 (m, 2H), 7.73–7.40 (m, 9H), 3.09 (d, J = 17.5 Hz, 8H), 2.44 (d, J = 17.6 Hz, 8H), 2.03 (d, J = 5.4 Hz, 8H), 1.63–1.35 (m, 14H). 13C NMR (75 MHz, CD2Cl2) δ 144.65, 141.27, 139.55, 138.53, 138.39, 137.31, 136.83, 135.96, 132.47, 131.91, 130.62, 130.46, 129.10, 128.23, 127.76, 127.30, 126.13, 125.80, 118.72, 113.55, 112.26. UV/vis (toluene) λmax (log ε): 357 (4.64), 375 (4.79), 396 (4.81), 470 (5.45), 617 (4.06), 675 (4.45). MALDI-TOF: m/z found 1840.9465, caled for [M+H]+ C140H103N4 1840.8216.
Porphyran 10-Pd

Bis(benzo[tr]thiophene)chlorodium[α] chloride (0.078 g, 0.204 mmol) and triethylamine (0.101 g, 1 mmol) were added to a solution of porphyran 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 (solvent CH2Cl2-TFA) and was considered complete after the absorption band of the dication at 470 nm disappeared. The mixture was allowed to cool then evaporated in vacuum to dryness. The residue was diluted with CH2Cl2, filtered through a thin layer of Celite to remove Pd black, and the solvent was evaporated. The product was purified by chromatography on silica gel using CH2Cl2 as eluent (dark red band collected). The solvent was evaporated and the residue was either recrystallized from CH2Cl2–ether to give porphyran 10-Pd (0.151 g, 76%) as a dark-brown powder.

1H NMR (300 MHz, C2D2Cl4) δ 8.48 (d, J = 7.4 Hz, 8H), 8.18 (d, J = 8.6 Hz, 8H), 7.88 (dd, J = 17.7, 8.1 Hz, 16H), 7.74–7.43 (m, 36H), 2.95 (s, 16H), 1.94 (s, 16H). 13C NMR (126 MHz, C2D2Cl4) δ 141.95, 141.16, 140.53, 139.08, 138.87, 137.66, 137.31, 136.92, 136.38, 135.90, 130.59, 130.08, 128.87, 128.26, 128.15, 128.05, 127.38, 127.09, 126.82, 126.15, 124.94, 124.00, 118.96, 29.42, 23.87. UV/vis (toluene) max (410-800 nm) (log ε): 340 (4.29), 357 (4.53), 395 (4.54), 446 (5.33), 628 (4.89). HRMS (ESI): m/z found 1927.5749, calcd for [M+] C140H84N4Pd 1927.5764.

Porphyran 11-Pd

Porphyran 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 (FP7-MCA-609305) for the financial support.

Notes and references


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