Design of porphyrin-based conjugated microporous polymers with enhanced singlet oxygen productivity†

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Novel non-toxic materials with antimicrobial surfaces are needed for medicinal applications. Potential alternatives for bacterial inactivation include materials that produce singlet oxygen, $\text{O}_2(^1\Delta_g)$, a short-lived, highly oxidative, and cytotoxic species. We synthesized a promising group of materials, porphyrin-based conjugated microporous polymers (CMPs), which effectively generate $\text{O}_2(^1\Delta_g)$ under visible light irradiation. CMPs were rationally designed and synthesized to maximize $\text{O}_2(^1\Delta_g)$ production. A strategy based on three-dimensional frameworks allowed the immediate environment of the porphyrin units to be tuned and their structure–property relationships to be elucidated. We investigated the photophysical and photochemical properties of the frameworks and compared them with the properties of porphyrin-based CMPs and metal–organic frameworks. In general, the $\text{O}_2(^1\Delta_g)$ production activity of the CMPs correlated with neither the surface area nor with the pore volume. The novel CMPs displayed high $\text{O}_2(^1\Delta_g)$ production, were stable in organic solvents, and did not undergo measurable photobleaching.

Introduction

In recent decades, great effort has been devoted to the development of antimicrobial materials that prevent the formation of bacterial colonies or directly kill bacteria upon contact. A wide range of strategies has been employed including the use of silver ions, antibiotics, or nanostructured metal oxides, such as TiO$_2$ and ZnO, combined with UV irradiation. Growing resistance against antibiotics and other chemotherapeutics has led to a search for antimicrobial treatments to which pathogenic microorganisms will not be easily able to develop resistance. A very promising alternative is photodynamic inactivation based on the formation of reactive oxygen species (ROS), including singlet oxygen, $\text{O}_2(^1\Delta_g)$, a short-lived, highly oxidative, and cytotoxic species, generated in situ via energy transfer from an excited molecule of a photosensitizer to an oxygen molecule. The development of nanomaterials producing $\text{O}_2(^1\Delta_g)$ has blossomed as a field of great interest because of the ease with which their functionality is both rapidly switched on and off by light and without introducing other reactants into the system. Novel non-toxic materials displaying antimicrobial properties under visible light irradiation are needed for medicinal applications. In this respect, porphyrin-based polymers that effectively generate $\text{O}_2(^1\Delta_g)$ represent a promising group of materials.

Porous polymers are a fast growing class of materials because of their high surface areas, tunable structures, good diffusion properties, and variable chemical functionality. Among others, purely organic structures such as covalent organic frameworks (COFs) and conjugated microporous polymers (CMPs) are receiving increased attention. In COFs, the building blocks are connected by reversibly formed bonds such as boroxine, boronyl ester, triazine, or imine, and therefore, the resulting materials are crystalline. On the other hand, their chemical stability can be lower than that of CMPs produced by ethynyl cyclotrimerization, metathesis, Suzuki, or Sonogashira polycondensations.

CMPs have a three dimensional structure created by condensation polymerization with one of the two monomers bearing more than two reactive end groups. Commonly, building blocks of porous polymers are rigid molecules with a well-defined geometry allowing a rational design of networks with the desired size and topology of pores. Thus, the use of molecules containing strictly sp$^3$ hybridized carbon atoms, such as 1,3,5-trisubstituted benzenes or 5,10,15,20-tetrasubstituted porphyrins, leads to planar networks stacked by $\pi–\pi$ interactions (abbreviated as 2D CMPs), whereas molecules with sp$^2$ carbon or silicon atoms (e.g., tetraphenyl methane, tetraphenyl silane, or adamantane) enable the preparation of...
materials with a three-dimensional covalently bound network (abbreviated as 3D CMPs).

Porphyrin molecules are commonly used building blocks for the construction of porous polymers because of their rigidity, planar geometry, and chemical stability. In addition, porphyrin units add synergistic functionalities resulting in materials suitable for gas storage and separation, sorption, optoelectronic devices, and electrocatalytic and catalytic applications. Porphyrins are also among the dyes that can be excited by visible light to excited singlet and triplet states in high quantum yields. The triplet states interact with molecular oxygen via energy transfer resulting in the formation of O$_2$(^3$\Delta_g$).

In recent years, a number of composite materials containing porphyrins for antimicrobial applications have been developed. The photosensitizing properties of porphyrin molecules are strongly dependent on surrounding environment because commonly occurring $\pi-\pi$ porphyrin stacking diminishes the photosensitizing properties. Therefore, tailoring properties for a specific purpose requires a high degree of control over the porphyrin environment. This is difficult to achieve in amorphous materials such as silica-based composites or nanowires. For these reasons, crystalline materials with an organized arrangement of porphyrin units have been extensively investigated, such as layered metal hydroxides and metal-organic frameworks (MOFs); however, these materials often possess limited chemical stability in aqueous environments. CMPs represent a unique combination of the stability of the polymeric structures bonded by C-C bonds with added structural variability and control.

In this work, we synthesized novel CMPs constructed from porphyrin building blocks suitable for the design of antimicrobial materials. To elucidate their structure–property relationship, we investigated their photophysical and photocatalytic properties and compared these properties with the properties of two known CMPs with the planar configuration of the building blocks and porphyrin-based MOFs. The newly prepared CMPs displayed markedly higher O$_2$(^3$\Delta_g$) production than 2D CMPs and MOFs, were stable in organic solvents, and did not undergo measurable photobleaching.

**Experimental**

**Materials**

Tetraphenyl methane (Manchester Organics), Br$_2$, BuLi (2.5 M in hexanes), B(O-iPr)$_3$, Pd(PPh$_3$)$_4$, palladium acetylacetone (Pd(acac)$_2$), pyrrole, benzaldehyde, piconol, trifluoroacetic acid, 2,3-dichloro-5,6-dicyano-1,4-benzoquinone (DDQ), benzene-1,4-diboronic acid (2,3-dichloro-5,6-dicyano-1,4-benzoquinone (DDQ), benzene-1,4-diboronic acid (II), Et$_3$N, MgSO$_4$ (all Sigma-Aldrich), 5,10,15,20-tetrakis(4-bromophenyl)porphyrin (8, Frontier Scientific Europe, Ltd., UK), 5,15-dibromo-10,20-diphenylporphyrin (10, PorphyChem, France), NaOH, HCl, acetone, diethyl ether, hexane (mixture of isomers), CH$_2$Cl$_2$ (all Lachner, Czech Republic), NaCl, NaHCO$_3$, K$_2$CO$_3$ (all Lachema, Czech Republic), tetrahydrofuran (THF), methanol (MeOH), and 1,4-dioxane (all Penta, Czech Republic) were used as purchased. For the Suzuki coupling reaction, 1,4-dioxane was dried with Na/Pb$_2$CO and distilled under N$_2$. HPLC grade acetonitrile (ACN) (Chromasolv Plus, Sigma-Aldrich) was used for all photophysical measurements.

**Instrumental methods**

The porosity and adsorption properties were probed by measuring nitrogen adsorption isotherms at the boiling point of liquid nitrogen (ca. 77 K) with a Micromeritics ASAP 2010 apparatus. Prior to adsorption experiments, the samples were evacuated at 80 °C for at least 24 hours. Because of the complex character of CMPs, the adsorption isotherms were analysed by several techniques including the BET method, Broekhoff-de Boer (BdB) t-plots, and DFT/NLDFT methods for various pore shapes and interactions potentials (as provided by the Micromeritics software), with the aim to achieve consistent texture parameters.

$^1$H and $^{13}$C NMR spectroscopy were performed on a Varian Mercury 400 Plus Instrument ($^1$H, 399.98 MHz; $^{13}$C, 100.59 MHz) at 298 K. The chemical shifts (δ/ppm) are given relative to the residual $^1$H signal of the deuterated solvent. Thermal analyses were carried out using a Setaram SETSYS Evolution-16-MS instrument coupled with a mass spectroscopy system. The measurements were performed in a synthetic air atmosphere (99.999%, Messer) with 30 mL min$^{-1}$ flow rate and a heating rate of 5 °C min$^{-1}$. Fourier transform infrared spectra (FTIR) were recorded with a Nicolet NEXUS 670-FT spectrometer in KBr pellets. The elemental analysis of CHN was performed using a standard combustion technique.

The UV/Vis absorption spectra of CMP dispersions in acetonitrile were recorded on a Perkin Elmer Lambda 35. Solid samples were recorded on the same instrument using a Labsphere RSA-PE-20 integration sphere accessory. The luminescence properties were monitored on a Fluorolog 3 spectrometer equipped with a cooled TBX-05-C photon detection module (Horiba Jobin Yvon) or with a Hamamatsu H10330-45 photomultiplier for measurement of the O$_2$(^3$\Delta_g$) emission spectra. The fluorescence lifetime measurements were performed using a laser-diode excitation at 405 nm (NanoLED-405LH, pulse width 750 ps, repetition rate 1 MHz). The emission was recorded at fluorescence maxima at 660 and 720 nm using a cooled TBX-05-C photon detection module in a time-correlated single-photon counting regime. The decay fits were made with exponential functions using the iterative reconvolution procedure of the DAS6 software (v. 6.4, Horiba Jobin Yvon, 2009).

**Photosensitization activity of CMPs**

The photosensitizing ability of CMPs was assessed using the reaction of photoproduced O$_2$(^3$\Delta_g$) with 9,10-diphenylanthracene leading to the corresponding endoperoxide. The course of the reaction was followed by UV/Vis spectroscopy as it is indicated by a decreasing absorption of 9,10-diphenylanthracene between 230 and 410 nm. In each experiment, 1.0 mg of CMP was dispersed in 20 mL of 10$^{-4}$ M 9,10-diphenylanthracene in acetonitrile and placed in a quartz cuvette. After sonication for 10 s, the dispersion was stirred with a magnetic bar and continuously irradiated by a 300 W Xe lamp (ozone free,
General procedure for preparation of Pd(II) porphyrins

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General procedure for preparation of Pd(II) porphyrins

Porphyrin and Pd(acac)2 were dissolved in toluene, and the mixture was refluxed at 110 °C for 24 h. The reaction mixture was filtered, and the solvent was evaporated. The product was triturated with hexane to remove excess Pd(acac)2 and the residual solvent. The product purity was checked by NMR and luminescence emission spectroscopy. If the sample still contained non-metallated porphyrin, then the procedure was repeated.

Preparation of Pd(II)-5,15-bis(4-bromophenyl)-10,20-diphenylporphyrin

(7). Porphyrin 6 (232 mg; 0.3 mmol), Pd(acac)2 (110 mg; 0.36 mmol) and toluene (200 mL) were used.

1H NMR (CDCl3): δ 7.75 (d, 6H, 3JHH = 7.6 Hz); 7.87 (d, 4H, 3JHH = 6.8 Hz); 8.03 (d, 4H, 3JHH = 8.4 Hz); 8.16 (d, 4H, 3JHH = 6.4 Hz); 8.81 (dt, 8H).

Preparation of Pd(II)-5,10,15,20-tetrakis(4-bromophenyl)porphyrin

(9). Porphyrin 8 (279 mg; 0.3 mmol), Pd(acac)2 (110 mg; 0.36 mmol) and toluene (300 mL) were used.

1H NMR (CDCl3): δ 7.88 (d, 8H, 3JHH = 8.0 Hz); 8.02 (d, 8H, 3JHH = 8.0 Hz); 8.80 (s, 8H).

General procedure for CMP polymerization

A Schlenk tube charged with the corresponding porphyrin, linker, and K2CO3 (0.4 M solution) was purged three times with argon. Then, a solution of Pd(PPh3)4 catalyst in dry 1,4-dioxane was added, and the reaction mixture was degassed in three freeze–pump–thaw cycles. The amounts of reactants, catalyst, and dioxane are given in Table 1. The reaction mixture was stirred at 100 °C for 24 h. The resulting precipitate was collected by filtration and thoroughly washed with water, MeOH, acetone, and THF. The solid product was purified in a Soxhlet extractor with THF, MeOH, and acetone for at least 24 h with each solvent.

Results and discussion

Syntheses and textural properties

Scheme 1 depicts the synthetic strategy for a series of CMPs. Polymerization of substituted porphyrins with linkers was performed using Suzuki–Miyaura cross-coupling reactions forming robust porous polymeric structures connected via C–C bonds. The material robustness allowed their purification by Soxhlet extraction, which ensured the complete removal of unreacted porphyrins. The prepared CMPs were insoluble solids that can be well dispersed in organic solvents such as acetonitrile or dimethylformamide; the hydrophobicity of the CMPs prevented them from use water for solution processing.

Polymers 3D-TPP, 3D-PdTPP, and 3D-DPP were designed by a combination of well-defined rigid 4-connected tetrahedral and 2-connected linear shaped building units to form a diamond-like structure with accessible porphyrin centres in pores (Fig. 1). The size of the cavities was modulated using differently sized building units while maintaining the porous structure. As anticipated, the reaction between two equivalents of trans-disubstituted porphyrin building blocks and one
equivalent of tetraphenyl methane linkers in 1,4-dioxane catalysed by Pd(PPh₃)₄ resulted in formation of 3D CMPs in high yields. The syntheses and properties of these materials are reported for the first time. 3D-DPP was constructed using 5,15-dibromo-10,20-diphenylporphyrin (10) to obtain a structure with narrow pores due to a close arrangement of the porphyrin units that are separated only by tertiary substituted carbon atoms breaking their conjugation in the polymer backbone. To enlarge the distance between the porphyrin units, tetraphenyl porphyrins 5 or 7 were the building blocks for 3D-TPP and 3D-PdTPP, respectively. Pd(II) in 3D-PdTPP brings an additional photofunctionality to the material.

For comparison purposes, two 2D CMPs were synthesized using a method derived from recently published procedures. 27 2D-TPP was built using tetrasubstituted tetraphenyl porphyrin 8 connected by 1,4-phenylene linkers 11, and the analogous 2D-PdTPP was prepared using Pd(II) porphyrin 9 instead of 8. Based on the idealized structures of the 2D CMP units, the porphyrin layers are closely stacked, forming pores oriented perpendicularly to the porphyrin plane and thus limiting the accessibility of the porphyrin centres (Fig. 1).

To optimize the O_{2}(1Δg) productivity investigated in the following experiments, Pd(II) porphyrins were used for the preparation of 2D-PdTPP and 3D-PdTPP (Scheme 1). Apart from the altered photophysical properties, tetracoordinated Pd(II) cation imposes a planar conformation on the porphyrin unit, thus increasing the rigidity of the CMPs structure.

The CMPs were characterized by various analytical techniques including ¹³C MAS-NMR, FTIR, thermogravimetric analysis, and elemental analysis, and their textural properties were assessed by the analysis of N₂ adsorption isotherms. The ¹³C MAS-NMR spectra (Fig. S1†) indicated the signals of tertiary and quaternary pyrrole carbon atoms in the range of 118–121 and 152–155 ppm, respectively. The intensive peaks at 141 ppm were attributed to both quaternary phenylene and meso-carbon atoms of the porphyrin rings. The most intensive peak at 127 ppm can be ascribed to tertiary phenylene carbon atoms. In addition, 3D CMP materials showed peaks of quaternary carbon and tertiary phenylene carbon atoms of tetraphenyl methane at 65 and 131 ppm, respectively. These results suggest that the spectra of CMPs are in agreement with their expected network structure as presented in Fig. 1. The FTIR spectra of CMPs exhibited vibrational peaks characteristic for porphyrins: stretching aromatic C–H vibrations in the range of 3022–3054 cm⁻¹, a sharp peak at 794–798 cm⁻¹ corresponding to pyrrole ring vibration, and an N–H stretching vibration at 3313 cm⁻¹ that disappeared in palladium analogues 2D-PdTPP and 3D-PdTPP (Fig. S2†).

Thermogravimetric analyses of 3D CMPs made of metal-free porphyrins revealed thermal stability up to 300 °C (Fig. S3 and S4†). The most thermally stable was 2D-TPP decomposing from approximately 350 °C (Fig. S5†). On the other hand, Pd-porphyrin based CMPs decomposed from approximately 250 °C, which could be due to palladium-catalysed oxidation (Fig. S6 and S7†). A partial mass loss below decomposition temperature was due to the evaporation of residual solvents. The decomposition of the samples was indicated by the coincident evolution of the gaseous products CO₂, H₂O, and NO₂. Elemental analyses of CMPs showed that the weight percentages of C, H, and N were slightly lower than the...
can be related to the imposed planarity of the Pd(II) porphyrin units.

### Photophysical and photochemical properties

The microporosity of CMP materials (Table 2) indicated good accessibility of the excited porphyrin units to oxygen, which is a fundamental prerequisite for the CMPs to act as efficient photosensitizers of \( \text{O}_2(1\Delta_g) \). In general, the processes after the excitation of porphyrin units involve the decay of the excited singlet states, \( 1\text{P}^* \), via fluorescence (eqn 1)), non-radiative vibrational relaxation, and intersystem crossing leading to the low-lying triplet states, \( 3\text{P} \) (eqn 2)). The triplet states are deactivated spontaneously via phosphorescence emission (eqn 3)), non-radiative relaxation, or by energy transfer with molecular oxygen in the ground state to form \( \text{O}_2(1\Delta_g) \) (eqn 4)).

\[
1\text{P}^* \rightarrow 1\text{P} + hr_P \\
1\text{P}^* \rightarrow 3\text{P} \\
3\text{P} \rightarrow 1\text{P} + hr_P \\
3\text{P} + \text{O}_2(1\Delta_g) \rightarrow 1\text{P} + \text{O}_2(1\Delta_g) \\
\text{O}_2(1\Delta_g) \rightarrow \text{O}_2(1\Delta_g) + hr_P
\]

The UV/Vis absorption spectra of CMP dispersions in acetonitrile displayed the characteristic Soret and Q bands of the porphyrin units (Fig. S10 and S11†). In comparison with the sharp Soret bands of monomeric porphyrins such as 5,10,15,20-tetraphenylporphyrin at 413 nm, embedding the porphyrin units into the polymer backbone led to a broadening and a shift to 440–450 nm with a shoulder at approximately 415 nm. The Q bands were red-shifted, with a largest shift for 2D CMPs indicating the most extensive interaction between the porphyrin units in these materials. The spectral features are characteristic of porphyrin polymers. The Q(0,0) and Q(0,1) fluorescence bands were broadened and slightly shifted to 658–666 and 720–721 nm, respectively, when compared with the corresponding bands at 647 and 714 nm measured for 5,10,15,20-

### Table 2  Textural parameters and fluorescence lifetimes of the investigated CMPs in acetonitrile dispersions at room temperature

<table>
<thead>
<tr>
<th>CMP</th>
<th>( S_{\text{BET}} )$/m^2 \text{~g}^{-1}$</th>
<th>( (P/P_0)_{\text{cal}} )</th>
<th>( V_{\text{pore}} )/cm$^3 \text{~g}^{-1}$</th>
<th>Pore width/nm</th>
<th>( \tau_f )/ns</th>
</tr>
</thead>
<tbody>
<tr>
<td>2D-TPP</td>
<td>443</td>
<td>0.05–0.10</td>
<td>0.18</td>
<td>2.0</td>
<td>0.5 (82), 7.2 (18)</td>
</tr>
<tr>
<td>2D-PdTPP</td>
<td>37</td>
<td>0.05–0.20</td>
<td>—</td>
<td>Nonporous</td>
<td>—</td>
</tr>
<tr>
<td>3D-TPP</td>
<td>353</td>
<td>0.05–0.25</td>
<td>0.20</td>
<td>2.1</td>
<td>0.5 (98), 4.1 (2)</td>
</tr>
<tr>
<td>3D-PdTPP</td>
<td>112</td>
<td>0.05–0.25</td>
<td>0.043</td>
<td>1.4</td>
<td>—</td>
</tr>
<tr>
<td>3D-DPP</td>
<td>624</td>
<td>0.05–0.10</td>
<td>0.29</td>
<td>&lt; 1.8</td>
<td>~0.2 (92), 7.3 (8)</td>
</tr>
</tbody>
</table>

\( S_{\text{BET}} \) stands for the BET surface area calculated within the range of relative pressures \( (P/P_0)_{\text{cal}} \) ensuring the validity of the BET equation; \( V_{\text{pore}} \) is the pore volume determined by the BdB \( t \)-plot method; pore width was determined by the NLDFT method (Fig. S9); \( \tau_f \) are the fluorescence lifetimes (amplitude fractions in % in parentheses).
tetraphenylporphyrin in acetonitrile. In addition, the intensity of the Q(0,1) band increased at the expense of the Q(0,0) band (Fig. 2) and fluorescence lifetimes were considerably shortened (Table 2) as follows from comparison of the average fluorescence lifetimes of 1.7, 0.8, and 0.6 ns for 2D-TPP, 3D-DPP, and 3D-TPP, respectively, with the single lifetime of 5,10,15,20-tetraphenylporphyrin in acetonitrile being 9.2 ns. For completeness, the fluorescence excitation spectra of all measured samples matched the corresponding absorption spectra supporting the identity of the fluorescent components.

The dynamics of the triplet states directly predetermine the efficacy of O$_2$(^1D_g) photosensitization according to eqn (4). Because 2D-PdTPP and 3D-PdTPP contained phosphorescent Pd-porphyrin units, the dynamics of their triplet states were probed by luminescence spectroscopy. The emission spectra of 3D-PdTPP were considerably affected by oxygen (Fig. 3). While the fluorescence bands at 575 and 613 nm remained unchanged, the phosphorescence bands at 710 and approximately 785 nm were quenched considerably in the presence of oxygen. Phosphorescence decay curves recorded in the absence of oxygen were best analysed by a biexponential function giving an average lifetime of approximately 120 µs. This lifetime is in the same order of magnitude as the lifetimes of monomeric Pd-porphyrins (e.g., Pd(n)-5,10,15,20-tetrakis(4-sulfonatophenyl) porphyrin has a lifetime of 297 µs in water). In contrast, the emission of 2D-PdTPP dispersions was only slightly affected by oxygen (Fig. S12†), which is in accordance with the nonporous character of this material with closely stacked porphyrin units leading to self-quenching of the excited states.

The above results document that the photophysical properties of the porphyrin units were affected by porphyrin–porphyrin interactions within a polymeric backbone and/or between adjacent backbones. The formation of O$_2$(^1D_g) was indirectly indicated only for 3D-PdTPP on the basis of phosphorescence quenching by oxygen according to eqn (4) (Fig. 3). To prove the formation of O$_2$(^1D_g), we measured the O$_2$(^1D_g) luminescence emission of the solid samples in the near infrared region (eqn (5)). As anticipated, both palladium-containing 2D-PdTPP and 3D-PdTPP had a luminescence band at 1272 nm, the characteristic of O$_2$(^1D_g) (Fig. S13†). The high intensity of the luminescence band of 3D-PdTPP suggested that this CMP is a better producer of O$_2$(^1D_g) than 2D-PdTPP, in accordance with the phosphorescence results. The other CMPs exhibited only tiny signals of O$_2$(^1D_g) presence.

To document the photosensitizing properties of CMPs in more detail, we performed luminescence measurements in acetonitrile dispersions. The photoproduction of O$_2$(^1D_g) by 3D-PdTPP was proven by the appearance of the corresponding luminescence band and by a strong dependence of the peak intensity on oxygen concentrations (Fig. 4). Clearly, no O$_2$(^1D_g) was produced in the absence of oxygen. Similar behaviour was observed for 2D-PdTPP and 3D-TPP dispersions (Fig. S14 and S15†); however, the intensities of photosensitized O$_2$(^1D_g) were...
much smaller when compared with the activity of 3D-PdTPP. In contrast, 3D-DPP (Fig. S16†) and 2D-TPP (Fig. 5) dispersions produced \( \text{O}_2(1^3\Delta_g) \) even after long-term purging of the dispersions with argon. These results indicate that molecular oxygen is quite difficult to remove from the microporous structure of both CMPs, such that these polymeric networks retain oxygen even in an argon atmosphere. As mentioned, the 3D-DPP network has large pore volumes with quite narrow pores (<1.8 nm) (Table 2). The pore width of 2D-TPP is 2.0 nm.

CMPs, composed of building blocks with a photosensitization functionality, can be good candidates for novel \( \text{O}_2(1^3\Delta_g) \) photosensitizers. The porphyrin units themselves used in this work are good \( \text{O}_2(1^3\Delta_g) \) sensitizers. However, when the porphyrin units become a part of the porous polymer network, it is difficult to accurately access the effectiveness of the formation of the porphyrin triplet states (eqn (2)), and, consequently, the effectiveness of \( \text{O}_2(1^3\Delta_g) \) generation (eqn (4)). This is because the porphyrin units tend to undergo π–π stacking interactions that lead to fast competitive non-radiative relaxations of the excited states that diminish \( \text{O}_2(1^3\Delta_g) \) productivity. For comparison, benzothiadiazole- and zinc phthalocyanine-based CMPs were identified as solid-state photosensitizers of \( \text{O}_2(1^3\Delta_g) \) under irradiation by visible and 700 nm light on the bases of indirect experiments. Similarly, the selective oxidation of thioanisole into the corresponding sulfoxide under visible light irradiation, occurring in the presence of BODIPY-based CMP, was attributed to a \( \text{O}_2(1^3\Delta_g) \) reaction channel. In contrast to these works, the presented results here prove directly the formation of triplet states and \( \text{O}_2(1^3\Delta_g) \), and therefore unambiguously indicate that the porphyrin-based CMPs can be considered as promising \( \text{O}_2(1^3\Delta_g) \) photosensitizers.

**Photosensitization activity**

Driven by the motivation to utilize the porphyrin-based CMPs as solid-state photosensitizers of \( \text{O}_2(1^3\Delta_g) \), we investigated the \( \text{O}_2(1^3\Delta_g) \) production efficiency in a reaction of 9,10-diphenylanthracene to its corresponding endoperoxide (Scheme 2). The reaction kinetics were followed by a concentration decrease of 9,10-diphenylanthracene recorded using UV/Vis spectroscopy. Blank experiments excluded the substrate sorption by the CMPs and the substrate reaction upon irradiation in the absence of the CMPs. As presented in Fig. 6, the kinetics strongly depend on the porphyrin arrangement in the CMPs. Two samples, 3D-PdTPP and 3D-TPP, exhibited the most effective photosensitizing activity, whereas 2D-PdTPP and 3D-DPP were the least photosensitizing materials. Based on these results, it is evident that the activity of the CMPs to produce \( \text{O}_2(1^3\Delta_g) \) available to react with the substrate can be correlated with neither the BET surface area nor with the corresponding pore volume, as these characteristics are quite low in the most effective 3D-PdTPP compared with the other CMPs (Table 2). In general, \( \text{O}_2(1^3\Delta_g) \) produced by a CMP particle can be formed at the external surface and/or in the bulk including the pores, and therefore the reactions of \( \text{O}_2(1^3\Delta_g) \) with a target species, commonly located at the surface, in pores, or in the close surroundings of a particle, depend on the ability of \( \text{O}_2(1^3\Delta_g) \) to reach these locations. This originates from the fact that \( \text{O}_2(1^3\Delta_g) \) has a relatively short lifetime, limiting its oxidation power to the proximity of a site where it was formed. This reasoning suggests that materials with good accessibility of the porphyrin units to

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**Fig. 5** Luminescence emission spectra of photosensitized \( \text{O}_2(1^3\Delta_g) \) produced in acetonitrile dispersions of 2D-TPP in oxygen (a), air (b), and argon atmosphere (c) upon excitation at 420 nm.

**Fig. 6** Reaction of photosensitized \( \text{O}_2(1^3\Delta_g) \) produced by CMP and MOF dispersions in acetonitrile with 9,10-diphenylanthracene according to Scheme 2.
oxygen and a low degree of porphyrin–porphyrin interactions have high photosensitizing activities, which is probably the case with 3D-PdTPP and 3D-TPP. On the other hand, the very low activity, especially of 3D-DPP, can be due to a close arrangement of the porphyrin units resulting in the quenching of the porphyrin triplet states.

To broaden the examination of porphyrin polymers, we also investigated the photosensitizing activity of porphyrin containing water-stable MOFs; namely, we synthesized PCN-222, MOF-525, and Al-TPPC MOF. The amount of MOFs in the experiments was adjusted to the same content of porphyrin units as used in the CMPs. Even though the porphyrin units in the MOFs do not stack together and the MOFs display significantly higher porosity than the CMPs (BET surface area: 2300, 2600, and 1400 m² g⁻¹ for PCN-222, MOF-525, and Al-TPPC MOF, respectively), the endoperoxide formation was rather low when compared with the activity of the CMPs. Having established the photosensitizing activities of a series of porphyrin-based materials, it is clear that structural and chemical composition differences of the immediate porphyrin environment cause significant variations in the O₂(1Δg) production.

To prove the stability and repeatable applicability of CMP photosensitizers, the most active 3D-PdTPP was separated after the complete reaction and reused in a new run. As presented in Fig. S17, the photosensitizer preserved its activity and can be repeatedly used. To analyse the effects of oxygen concentration, some irradiation experiments were performed in an argon atmosphere. Both 2D-PdTPP and 2D-TPP decreased the 9,10-diphenylanthracene concentration down to 54.7 ± 0.9 and 40.3 ± 1.1%, respectively, after 3 hours of irradiation in air. In agreement with the excitation scheme (eqn (1)–(4)), the photosensitizing activity of 2D-PdTPP completely diminished in the absence of oxygen. In contrast, the usage of 2D-TPP in an argon-saturated dispersion decreased the substrate concentration down to 88.5%. This finding is in agreement with the behaviour of 2D-TPP described above (Fig. 5) and confirms that oxygen in 2D-TPP is confined in the microporous structure of the polymer network and cannot be easily removed as in the case of 2D-PdTPP. This can be advantageous in systems where the concentration of oxygen is fluctuating with a constant need for O₂(1Δg) generation.

Conclusions

The presented materials are a platform for the fabrication of heterogeneous photosensitzers functioning under visible light. They are photostable, and their nature allows for simple separation and reuseing. We prepared a series of new porphyrin-containing 3D CMPs with a diamond-like topology and compared their photophysical properties with those of corresponding 2D CMPs and water-stable MOFs. It is evident that the activity of CMPs in the O₂(1Δg)-based reactions cannot be directly correlated with their BET surface area nor with their pore volume, but strongly correlates with the immediate porphyrin environment. When porphyrins are horizontally stacked, as in the 2D CMPs, or are separated by a short organic linker (3D-DPP), most of the absorbed light energy is dissipated by non-radiative processes. Enlarging the porphyrin separation, as in 3D-PdTPP and 3D-TPP, leads to effective solid photosensitzers that display a significantly higher activity. Compared with the MOFs, the CMPs offer greater control over the porphyrin environment and superior stability by replacing susceptible coordination bonds with robust covalent bonds; therefore, the CMPs are more suitable and more efficient photosensitzers.

The CMP materials represent a substantial step forward in the fabrication of heterogeneous photosensitzers acting under visible light. They are photo- and chemically stable, and their nature allows for recycling and utilization in composites for the construction of antimicrobial coatings under visible light. Ongoing research is focused on the development of composites by fixing porous CMP nanoparticles on polymeric surfaces and dispersing them as fillers into continuous polymeric matrices.

Acknowledgements

This work was supported by the Czech Science Foundation (No. 16–15020S). J. H. gratefully acknowledges the Grant Agency of the Charles University in Prague (No. 252216) for financial support.

Notes and references