Synthesis of New Porphyrin Trimers via Buchwald–Hartwig Amination Reaction

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Two types of the novel porphyrin trimers linked by amines were synthesized by the Buchwald–Hartwig amination reaction of zinc complex of 5-(4-bromophenyl)-β-octaalkylporphyrin with zinc 5,15-di(4-aminophenyl)-10,20-dimesitylporphyrin, and zinc 5,15-diaryl-β-octaalkylporphyrins, where aryl groups were functionalized with piperazine or diaza-18-crown-6 ether. One type of trimers was bridged by one nitrogen atom of diphenylamine and another type of trimers was linked through two different nitrogen atoms of the cyclic diamines. Each type of the trimers has required specific catalyst composition varying in the ligands for catalytic palladium complex: BINAP and DavePhos correspondingly. The highest 75 % yield was achieved for the piperazine linked trimer. UV-Vis spectroscopic investigations of the synthesized compounds were carried out and showed that the observed spectra of the trimers consist of the spectra sum of their porphyrin components. Thus the trimers have shown negligible interchromophore interaction in both ground and excited states. The geometry of the trimer bridged by diphenylamine was optimized by molecular mechanics MM+ method showing angled structure with the distance between centers of the macrocycles of 16.86 Å. The interaction of the chromophores should be weak on such a distance, and there is no conjugation of electron systems of chromophores due to the orthogonal orientation of the bridging benzene rings relative to the tetrpyrrole plane. These results of the geometry calculations well correspond to the spectroscopic observations.

Keywords: Porphyrin trimer, oligoporphyrin, interchromophore interaction, aminoporphyrin, Buchwald-Hartwig amination, cyclic diamine, diazacrown ether.

Синтез новых порфириновых тримеров с помощью реакции аминирования Бухвальда–Хартвига

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Два типа новых порфириновых тримеров с аминовыми мостиками были синтезированы с помощью реакции аминирования Бухвальда-Хартвига между цинковыми комплексами 5-(4-бромфенил)-β-октаэтилпорфирин и 5,15-ди(4-аминофенил)-10,20-димезитилпорфирин и 5,15-диарил-β-октаэтилпорфиринов, в которых арильные группы функционализированы пиперазином или диаза-18-краун-6-эфиром. Один тип тримеров использован в качестве связывающего мостика один атом азота дифениламина, а другой тип тримеров связан через два различных атома азота циклических диаминов. Для получения каждого типа тримеров требуется специфический состав катализатора, отличающегося используемыми лигандами для каталитического палладиевого комплекса: BINAP и DavePhos, соответственно. Наиболее высокий выход 75 %

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Introduction

Multiporphyrins are of increasing interest due to their special electronic properties arising from the interaction of aromatic systems of the tetrapyrole macrocycles. Such interactions give rise to the new properties inherent in this ensemble which are not the simple sum of components. The nature of the interactions depends on a relative spatial positioning, geometry, rigidity and type of the bridges connecting porphyrinic cycles. Conjugation conductive groups are of special interest. Expanded π-conjugation leads to unusual electronic properties applicable in molecular electronic devices,[1–4] photon tunnels,[5] multispin molecules with magnetic properties,[6] devices for the information storage,[7] nonlinear optical (NLO) materials,[8,9] and accompanied to that increased long-wave absorbance is important in sensitizers for photodynamic therapy (PDT)[10] and solar cells.[11]

Numerous oligoporphyrins with a large variety of connecting fragments were reported.[12] Among them just a few types of relatively short linkers were explored. The shorter distance between tetrapyrole aromatic systems means the higher degree of their interaction. But the distance is only one of the numerous parameters which can tune the interchromophore interaction. Mutual arrangement, angles between tetrapyrole planes, nature of the bridges linking porphyrins play an important role.[13] The strongest interaction was with π-conjugation conducting linkers. These types of linkers are easy to insert via the well developed catalytic cross-coupling methodology such as Heck, Suzuki, and Sonogashira reactions.[12,14–16] Among them acetylene, phenylene, and their combinations were frequently used as linkers in oligoporphyrins. Acetylene groups were used as the most effective conductors of π-conjugation.[13,14] The very short distance and high conjugation lead to the strong interaction which practically combine all porphyrins to the linear layout. All these relatively short linkers have allowed the rapid and efficient energy transfer in excited state despite the lack of conjugation.[14,17,18] The Heck and Stille reactions allow to obtain angeld and rigid oligoporphyrins with the ethylene and diphenylethylene linkers.[12,19] Another type of linking methodology via 1,2,3-triazole was performed by catalytic 1,3-dipolar cycloaddition which is so called ‘click’ reaction.[20] 1,4-Diphenyl-1,2,3-triazole group is a rigid angled linker providing effective electronic communication of the porphyrin units in the excited state which was investigated by femtosecond spectroscopy clearly demonstrating very interesting coherent phenomenon.[21]

Among various linkers amino group is of special interest as it is capable to coordinate metal cations, imparting porphyrinic molecules additional bonding centers. It is necessary to emphasize, that metalloporphyrins with amino groups contain two opposite types of bonding centers: electrophilic metallocenter and nucleophilic nitrogen center. Such combination provides these molecules with ability to self-assembly, and enriches design opportunities of supramolecular ensembles, amplifies receptor properties. Besides by means of amine groups as strong donor substituents it is possible to influence electronic density of porphyrin macrocycle and, consequently, photophysical characteristics. meso-Diarylamino substituted porphyrins were shown to be the best sensitizers for the dye-sensitized solar cells.[22] Among oligoporphyrins one of the highest efficiencies were achieved with porphyrin dimers linked by aminophenyl groups attached to 1,3,5-triazole heterocycle.[23] Earlier we have investigated synthesis of the bisporphyrins linked by diphenylamine and various diamine linkers and have showed their abilities for supramolecular assembly. In the given work we have applied the earlier developed approaches to a synthesis of new porphyrin trimers linked with diphenylamine group and cyclic diamines.

Ключевые слова: Тример порфирина, олигопорфирина, взаимодействие хромофоров, аминопорфирина, аминирование по Бухвальду-Хартвигу, циклический диамин, диазакраун-эфир.
Experimental

Materials. 1,4-Dioxane was distilled and dried over sodium under atmosphere of Ar. CH₂Cl₂ was distilled over CaH₂, CHCl₃ was distilled over P₂O₅, and was freed of acids by stirring over K₂CO₃. Methanol was distilled over magnesium turnings. The starting materials were generally used as received (Aldrich). 5-(4-Bromophenyl)-2,3,7,12,18,28,32-hexamethyl-13,17-dithranol matrix. UV-Visible spectra were recorded on a Cary-100 UV-Visible Spectrophotometer. All reactions were performed under argon at room temperature and referenced to the residual protons of solvent (CDCl₃). δH ppm: 8.85 (4Н, d, J = 7.69 Hz, H-2); 7.79 (4H, d, J = 7.78 Hz, Ar-H); 7.41 (4H, s, Ar-H); 7.05 (4H, d, J = 7.78 Hz, Ar-H); 4.06 (4H, br s, NH₂); 2.61 (6H, s, CH₃); 1.86 (12H, s, CH₂); UV-Vis (CH₂Cl₂) λ nm (lg ε): 424 (5.6), 551 (4.34), 572 (3.93), 593 (3.73). Mass spectrum (MALDI TOF) m/z: 790.28 M⁺ [M+3H]⁺ for [C₁₉H₁₈N₂Zn]⁺.

Trimer ZnP(PhPip)-(ZnPPh₂). ZnP(PhPip)₂ (37.8 mg, 0.037 mmol), ZnPPhBr (55 mg, 0.073 mmol), Pd(dbta)₂ (3.3 mg, 0.01 mol (5.64·10⁻⁴ mmol, 8 mol %)), DavePhos (4.6 mg, NaOBut (9.8 mg) and dioxane (1 ml) were placed in a dried, resposable round-bottom flask with magnetic stirrer, which was backfilled with argon, sealed and heated to reflux with stirring for 24 hrs. The reaction mixture was then cooled to room temperature, filtered, and the solvent was evaporated in vacuo. The crude product was purified by column chromatography on silica gel with gradient eluent dichloromethane/methanol 100:0.5–100:2. Yield: 65.5 mg (75 %). 1H NMR (600 MHz, CDCl₃, 298 K) δ ppm: 10.19–10.11 (8H, m, 15.20, 26.50, 22.79, 21.67, 15.77, 14.16, 12.18, 11.63). UV-Vis (CH₂Cl₂) λ nm (lg ε): 406 (5.18), 424 (4.8), 536 (4.00), 570 (4.02). Mass spectrum (MALDI TOF) m/z: 2137.09 [M+3H]⁺ for [C₁₉H₁₈N₂Zn]⁺.

Results and Discussion

The synthesis of porphyrin trimers bridged by amino group was performed via palladium catalyzed Buchwald-Hartwig amination reaction between p-aminophenyl and p-bromophenyl meso-substituted porphyrin components where amino group replaces bromine atom thus linking two porphyrins together through nitrogen atom of amino group. Recently we have reported on the synthesis of bisporphyrins[34] bridged by the tertiary diarylkylamine group formed via two sequential couplings of two bromophenylporphyrin molecules with the primary alkylamine. In the current work two primary arylamine groups of the bis(aminophenyl)porphyrin have reacted with two bromophenylporphyrin molecules to give a trimer with...
a secondary diarylamine bridging group. In principle such trimers can be synthesized via two alternative ways (Figure 1): 1) interaction of one bis(aminophenyl)porphyrin building block with two mono(bromophenyl)porphyrin blocks; 2) reaction of one bis(bromophenyl)porphyrin building block with two mono(aminophenyl)porphyrin blocks. The choice was determined by the ease of the precursors synthesis. Both symmetrical 5,15-functionalized porphyrins are easily available by the [2+2] McDonald condensation of dipyrromethane with substituted benzaldehyde. The hardest is the synthesis of unsymmetrical monosubstituted porphyrin. Synthesis of the mono(aminophenyl)porphyrin via mixed condensation has given low yield with hardly separable byproducts (Supporting Information, Figure 1S) whereas the synthesis of meso-mononor(bromophenyl) porphyrin was developed\(^{[28,29]}\) by the directional way with high yield through condensation of the arylaldehyde with β-octaalkyl substituted a,c-biladiene\(^{[30]}\) obtained from the corresponding β-substituted 2-formylpyrrole and dipyrromethane (Supporting Information, Figure 2S).

Thus 5,15-bis(aminophenyl)porphyrin was synthesized as follows: [2+2] McDonald condensation of meso-mesityldipyrromethane\(^{[31]}\) with of p-nitrobenzaldehyde has given 5,15-bis(p-nitropheno)porphyrin. Mesityl substitution was used as sterically hindered mesityl groups were shown to inhibit side reactions of scrambling in a synthesis of unsymmetrically meso substituted porphyrins.\(^{[32]}\) Then without isolation nitroporphyrin porphyrin was reduced with \(\text{SnCl}_2\cdot2\text{H}_2\text{O/}\text{HCl}\) in chloroform/acetic acid following treatment with aqueous ammonia to give \(\text{H}_2\text{PMe}_3\cdot(\text{PhNH}_2)\)\(^{[29]}\) with 25 % yield over two stages accordingly. It is known that free base porphyrins are not convenient for homogenus transition metal catalyzed reactions as they can coordinate catalytic metal ions removing them from the catalytic cycle thus inhibiting the reaction. Therefore the free base porphyrins were converted to their zinc complexes by reaction with an excess zinc acetate monohydrate in boiling chloroform for two hours (Supporting Information, Figure 1S). 5-(4-Bromophenyl)-2,3,7,8,12,18-hexamethyl-13,17-di(α-pentyl)porphyrin (ZnPPhBr) was prepared (Supporting Information, Figure 2S) as we have earlier reported.\(^{[30]}\) Obtained porphyrins are convenient due to their good solubility in nonpolar solvents provided by β-alkyl substituents in bromoporphyrin and mesityl groups in aminoporphyrins. Different substitution in the tetrapyrrole cores of the porphyrin building blocks is helpful to discriminate porphyrin units in a trimer. Besides, such heterogeneity is useful for the subsequent investigation of processes of energy transfer between tetrapyrrolic macrocycles.

The coupling reaction between aminophenylporphyrin and bromophenylporphyrin was carried out with a palladium catalyst formed from 10 mol % \(\text{Pd(dba)}_2\) in a combination with 20 mol % of a diphasine ligand (±)-2,2′-bis(diphenylphosphino)-1,1′-binaphthyl ((rac)-BI-NAP) with 1,4 eq. of NaOBF\(_4\) as a base in a boiling dioxane under argon during 24 hrs. The choice of catalyst and conditions was based on the results of our previous work of amination of bromophenylporphyrins with various amines.\(^{[24,25,31,33]}\) The reaction between 5,15-bis(4-aminophenyl)porphyrin ZnPMe\(_3\cdot(\text{PhNH}_2)\)_2 and 2 equivalents of 5-(4-bromophenyl)porphyrin ZnPPhBr has led to arylation of both amino groups yielding 21 % of porphyrin trimer ZnPMe\(_3\cdot(\text{PhNH}_2)\)\(_2\)-(ZnPPh)\(_2\) (Figure 2).

Two other trimers were synthesized from porphyrin functionalized with cyclic amines such as piperazine and diaza-18-crown-6 ether. Reaction of one equivalent ZnP(PhPip), (bis-piperazine substituted zinc porphyrin) with two equivalents of ZnPPhBr in presence of monophosphine ligand 2-(dicyclohexylphosphino)-2′-(N,N-dimethy lamino) biphenyl (DavePhos) has led to the formation of trimer ZnP(PhPip)\(_2\)-(ZnPPh), where three porphyrin fragments are linked with two piperazine bridges (Figure 3) in an excellent 75 % yield. This is more than twice higher compared to the yield of the corresponding dimer earlier obtained from the monopiperidine substituted porphyrin.\(^{[24]}\) The trimer with diazacrown ether as linker was considerably harder to obtain compared to the analogous dimer.\(^{[64]}\) Interaction of the porphyrin with two diaza-18-crown-6 ether fragments ZnP(PhDiaza18C6), and bromoporphy rin ZnPPhBr has given the trimer ZnP(PhDiaza18C6)\(_2\)-(ZnPPh), in 27 % yield which is almost three times lower than for the piperidine linker (Figure 4).

The structure of the porphyrin trimer ZnPMe\(_3\cdot(\text{PhNH}_2)\)\(_2\)-(ZnPPh), was confirmed by \(^1\)H NMR. Resonances of protons at β- and meso-positions of the tetrapyrrolic macrocycle and benzene rings of the ZnPMe\(_3\cdot(\text{PhNH}_2)\)\(_2\)-(ZnPPh), were shifted downfield due to the mutual deshielding interaction

![Figure 1](image1.png)

**Figure 1.** Schemes of synthesis of porphyrin trimers from amino and bromo porphyrin building blocks.
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**Figure 2.** Synthesis of the porphyrin trimer.

**Figure 3.** Synthesis of the porphyrin trimer linked by piperazine.

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of the porphyrin aromatic systems for ∆δ=0.3 (0.24), 0.24 (0.14), 0.15–0.69 ppm correspondingly (Support, Table 1S). But phenyl protons at β-position (Support, Figure 3S) were slightly upshifted due to the donor effect of nitrogen atom (instead of bromine). Thus the phenyl protons resulted from bromo component were discriminated shifting in opposite directions. The methyl groups of mesityl substituents were upshifted for 0.68 and 0.32 ppm. The proton shifts in both dimers were not so unambiguous. Shifting of β-protons of dimers was insignificant (~0.1 ppm). Shifts of β-protons of amino bridged dimer (ZnPMe₃,Ph)₂NH were downfield and dimer without amino group (ZnPMe₃,Ph)₂ had an upshift. But the shifts of the protons of bridging benzene rings of amino component were considerable and downfield for all compounds. The oligomers linked by cyclic diamines contained one type of β-alkylsubstituted tetrapyrole rings. The meso-protons of the oligomers were shifted downfield whereas phenyl protons were not shifted unidirectionally (Support, Table 2). The CH₃ protons of the cyclic diamines were shifted upfield for the piperazine bridged trimer and downfield for diazacrown ether bridged oligomers.

The electronic spectra of the trimer ZnPMe₃,(PhNH)₂- (ZnPPh)₂ have revealed the splitting of the Soret band to two components with λₙm=424 nm and λₙs=406 nm (Table 1). Their intensity ratio is 2:1 which corresponds to the ratio of the two different types of the porphyrin components (one central and two peripheral), and the positions of the Soret bands are also the same as in starting porphyrin building blocks ZnPMe₃,(PhNH)₂, and ZnPPhBr. Thus the observed spectra of the trimer could be just a simple sum of the spectra of two types of porphyrin components. The difference in the positions of the adsorption bands results from the different substitution of the tetrapyrole macrocycles: one has β-alkyl substituents and 3 free meso-positions, and another has free β-positions and meso-mesityl substituents.

![Figure 4. Synthesis of the porphyrin trimer linked by diazacrown ether.](image)

<table>
<thead>
<tr>
<th>Compound</th>
<th>λₙm, nm (lgε)</th>
<th>Soret band</th>
<th>Q bands</th>
</tr>
</thead>
<tbody>
<tr>
<td>H₂PPhBr</td>
<td>404(5.29)</td>
<td>503(4.21), 537(3.92), 571(3.88), 624(3.57)</td>
<td></td>
</tr>
<tr>
<td>ZnPPhBr</td>
<td>406(5.56)</td>
<td>536(3.98), 572(3.95)</td>
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<tr>
<td>ZnPMe₃,(PhNH)₂</td>
<td>424(5.6)</td>
<td>551(4.34), 593(3.73)</td>
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<tr>
<td>ZnPMe₃,(PhNH)₂,(ZnPPh)₂</td>
<td>406(5.18)</td>
<td>536(4.00), 570(4.02)</td>
<td></td>
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<tr>
<td>ZnP(PhPip)</td>
<td>413(5.02)</td>
<td>542(3.87), 574(3.6)</td>
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<tr>
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<td>407(4.77)</td>
<td>536(3.54), 571(3.45)</td>
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<tr>
<td>ZnP(PhDiaza18C6)</td>
<td>418(4.57)</td>
<td>545(3.43), 577(3.01)</td>
<td></td>
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<tr>
<td>ZnP(PhDiaza18C6),(ZnPPh)₂</td>
<td>408(4.74)</td>
<td>537(3.59), 571(3.45)</td>
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<tr>
<td>ZnP(PhDiaza18C6),-ZnPPh</td>
<td>406(5.59)</td>
<td>536(4.44), 571(4.35)</td>
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</tbody>
</table>

It is known that the splitting of the Soret band in the multiporphyrins can be a consequence of the excitation coupling[35,36] and reported as characteristic for meso-meso linked porphyrins.[37] The geometry of the trimer was optimized by molecular mechanics MM’ method (Figure 5). The distance between centers of the macrocycles is 16.86 Å. The interaction of the chromophores is weak on such distance, and there is no conjugation of electron systems of chromophores due to the orthogonal orientation of the bridging benzene rings relatively tetrapyrole plane. Based on the optimized geometry the calculated value (made by point-dipole approximation)[38] of the splitting for the trimer (~6 nm) turned out to be three times less than difference between centers of the Soret bands of the trimer porphyrin components (18 nm). But the conformational freedom of the

Table 1. Electronic spectra of the porphyrin substrates and coupling products.
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trimer could lead to the decreased exciton coupling in the other conformers and broad shape of the two overlapping Soret bands probably didn’t allow to reveal the possible splitting.

The distances between centers of the macrocycles in the trimers bridged by piperazine and diazacrown ether are considerably larger. Besides these bridges are even more conformationally flexible. Thus the splitting of the Soret band in these cases should be completely negligible.

Conclusions

The novel porphyrin trimer compounds were synthesized by Buchwald-Hartwig amination reaction. Yields of the reaction are highly dependent on the substrates. The higher 75 % yield was achieved for piperazine linked trimer. UV-Vis spectroscopic investigations did not reveal interchromophore interaction between porphyrin components in both types of the synthesized porphyrin trimers which can be explained by the conformational freedom of the linkers allowing the variety of relative positions of the monomer components in the trimer. Obtained trimers are interesting as they are linked by the groups capable to coordination with metal cations with the possible formation of the supramolecular assemblies. Especially cyclic piperazine and diazacrown ether linkers are strong chelating ligands. The coordination of metal cations with the trimers could lead to the changes of the mutual position of the porphyrin components and correspondingly to the changes of their electronic communication giving possibilities to control the photophysical properties of the porphyrin trimers. The investigation of the supramolecular coordination of the trimers and their photophysical properties is on the way and will be reported in upcoming publications.

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