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■ Host–Guest Systems

Allosteric Control of Naphthalene Diimide Encapsulation and Electron Transfer in Porphyrin Containers: Photophysical Studies and Molecular Dynamics Simulation

Laura Zanetti-Polzi,^[a] Ryan Djemili,^[b] Stéphanie Durot,^[b] Valérie Heitz,^{*,[b]} Isabella Daidone,^{*,[c]} and Barbara Ventura^{*,[d]}**Abstract:** ■■■ Please provide academic title for each author

■■■ The complexation processes of N,N'-dibutyl-1,4,5,8-naphthalene diimide (NDI) into two types of π -electron-rich molecular containers consisting of two Zn^{II}-porphyrins connected by four flexible linkers of two different lengths, were characterized by means of absorption and emission spectroscopies and molecular dynamics simulation. Notably, the addition of NDI leads to a strong quenching of the fluorescence of both cages only when they are in an open conformation suitable for guest encapsulation, a situation triggered by silver(I) ions binding to the lateral triazoles. Molecular dynamics simulations confirm the fast binding of NDI, likely assisted by NDI-silver(I) interactions. Upon NDI complexation, the two porphyrin macrocycles get closer, with an

optimized face to face orientation, suggesting an induced-fit mechanism through π - π interactions with the NDI aromatic cycle. Ultrafast transient absorption experiments allowed to identify the process of quenching of the Zn-porphyrin fluorescence as an efficient photoinduced electron transfer reaction between the cage porphyrin and the included NDI guest. The process occurs on fast and ultrafast time scales in the two complexes (1.5 ps and ≤ 300 fs) leading to a short-lived charge separated state (charge recombination lifetimes in the order of 30–40 ps). The combined computational and experimental approach used here is able to furnish a reliable model of the NDI-cage complexation mechanism and of the corresponding electron transfer reaction, attesting the allosteric control of both processes by the silver(I) ions.

Introduction

Reversible encapsulation of guest molecules inside the cavity of three-dimensional structures is a phenomenon of great interest in supramolecular chemistry. It enables the guest to be isolated from the bulk medium and to establish new interac-

tions with its host which affect their mutual physical and chemical properties.^[1] The new properties that emerge from this close association have opened the way to applications in many areas of chemical, biological and material sciences. Encapsulation is often a critical step to access to more complex processes where the guest plays the role of the substrate or the catalyst in chemical transformations, the active drug in drug-delivery systems or the functional molecular unit in self-healing materials.^[2] Therefore, the stability and lability of the inclusion complex in a given environment has to be controlled by a fine electronic and geometrical match between the host and the guest which is not easily transposable from one environment to another one.^[3] Interestingly, guest complexation can also be controlled with external stimuli, such as photons,^[4] electrons^[5] or chemical entities,^[6] that interact with specific sites of the host, remote from the binding sites of the guest and induce large and reversible modifications of the host structure. Such control is reminiscent of the allosteric control used by many biological systems such as proteins, enzymes and molecular machines.^[7] Inclusion complexes are also well suited for achieving efficient photoinduced electron transfer, a key process in natural photosynthesis. The primary photoinduced electron transfer steps occurring in the Photosystem II involve a photoexcited dimer belonging to the family of porphyrin derivatives and a series of electron acceptors all maintained in a specific spatial organization through non covalent

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interactions in a protein host.^[8] Notably, the last electron acceptor, a quinone, once doubly reduced and protonated loses its affinity for its binding pocket, leaves the protein and is replaced by a fresh quinone.^[9] The modified affinity of the quinone in its reduced and protonated states for its host and its reversible binding is at the origin of the developed proton-motive force and the subsequently generated chemical energy. Regarding the importance of these processes, photoinduced electron transfer between porphyrins and electron acceptors in non-covalent hybrids has been investigated in different biomimetic inclusion complexes,^[10] in mechanically linked structures in the context of molecular machines^[11] and supramolecular optoelectronic devices.^[12] Nevertheless, limited examples have reported on an allosteric control of photoinduced electron transfer.^[10c, 11a,b]

In this vein, we were interested in studying the inclusion complex of *N,N'*-dibutyl-1,4,5,8-naphthalene diimide (**NDI**) into π -electron-rich hosts possibly giving rise to photoinduced electron transfer processes. In particular, we studied the inclusion of **NDI** in molecular cages, that we call **1** and **2**, consisting of two Zn^{II}-porphyrins connected by four flexible linkers of different lengths, endowed with triazole binding sites (Figure 1). 1,4,5,8-Naphthalene diimide, as a versatile aromatic π -acceptor, was previously combined with various π -rich donors including porphyrins in the context of supramolecular host-guest systems.^[13] Previous studies performed with the molecular container **1** endowed with the "short" linkers revealed that it behaves as an allosteric receptor, able to encapsulate **NDI** only once the container was in an open form, obtained upon silver(I) ions binding to the lateral triazoles.^[14] Here, combined spectroscopic and computational studies are reported to shed light on the binding affinity of **NDI**, on its dynamic encapsulation induced by the opening of containers **1** and **2** upon silver(I) ions binding and on the photoinduced electron transfer process taking place within the assembly. The structural char-

acterization of the complexation process is obtained by multiple, explicit solvent, molecular dynamics (MD) simulations and the photophysics of the inclusion complexes is investigated by means of ultrafast transient absorption spectroscopy. Binding of silver(I) ions to the lateral linkers is shown to be the event that controls and drives **NDI** inclusion and thus the occurrence of the photoinduced electron transfer reaction between the Zn-porphyrins of the host and the **NDI** guest.

Results and Discussion

Photophysical characterization and evaluation of guest binding in diluted solutions

The photophysical properties of cages **1** and **2** and of their silver(I) complexes $[\text{Ag}_4(\mathbf{1})](\text{OTf})_4$ and $[\text{Ag}_4(\mathbf{2})](\text{OTf})_4$ in $\text{CH}_2\text{Cl}_2/\text{MeOH}$ (90:10), a solvent mixture that could ensure good solubility of the porphyrin cages and of the silver salt $\text{Ag}(\text{OTf})$, were previously reported.^[15] A complete characterization of the absorption and emission properties of the guest **NDI** in the same solvent mixture has been performed, and the results are summarized in the ESI (Figure S1 and Table S1). In diluted solutions ($C \leq 10^{-6} \text{ M}$) **NDI** shows the typical absorption and emission features of the monomeric species: a structured absorption in the UV region, with $\epsilon_{380 \text{ nm}} = 2.62 \times 10^4 \text{ M}^{-1} \text{ cm}^{-1}$, and a weak fluorescence ($\phi_f = 0.0017$) with a vibronic structure mirroring the absorption spectrum.^[16] Upon increasing the concentration, additional absorption features in the range 450–600 nm and a broad emission band peaking at 566 nm appear; a similar red-shifted emission has been reported for excimers or excited dimers of naphthalene diimides in concentrated solutions or packed structures.^[16a, 17] In our system this secondary emission could be clearly characterized in concentrated solutions (10^{-4} M onwards) where the excitation spectrum collected at 600 nm matches the absorption bands present in the 450–

Figure 1. Chemical structures of the hosts **1** and **2**, their silver(I) complexes and of the **NDI** guest.

600 nm region (Figure S1), evidencing that it arises from a ground state dimer or higher order aggregate (indicated generally as NDI_n).

Moving towards the analysis of the binding between the bis-porphyrinic cages **1** and **2** and the **NDI** guest, we examined the photophysical properties of the cages, in their closed conformation, once treated with an excess of **NDI**. Absorption and emission spectra registered for mixtures of cage (ca. 2×10^{-7} M) and **NDI** (ca. 1×10^{-4} M, ≥ 500 equiv) are reported in Figures S2 and S3 for cages **1** and **2**, respectively. It is evident that there are no changes in the absorption features of the cages in the mixtures, while a slight reduction of emission intensity, that is, 3% for **1** and 7% for **2**, is observed. The latter is paralleled by measured excited state lifetimes of 1.64 ns (vs. 1.71 ns) and 1.54 ns (vs. 1.64 ns) for **1** and **2** in the mixtures (vs. cage alone),^[15] respectively. Though small, this quenching can be considered as a result of a bimolecular interaction between the cage and **NDI** molecules, and reaction rate constants of $2.7 \times 10^{11} \text{ M}^{-1} \text{ s}^{-1}$ for **1** and $4.4 \times 10^{11} \text{ M}^{-1} \text{ s}^{-1}$ for **2** could be derived, values one order of magnitude higher than the diffusion controlled rate constant ($k_{\text{diff, CH}_2\text{Cl}_2} = 1.5 \times 10^{10} \text{ M}^{-1} \text{ s}^{-1}$ at 293 K).^[18] This outcome seems to indicate that short-lived encounter complexes, where the cages interact with external **NDI** molecules, are operative in the reduction of the porphyrins fluorescence for less than 10%. Formation of such encounter complexes is also predicted by molecular dynamics simulations (see below).

The process of inclusion of the guest **NDI** inside the two zinc-porphyrin cages, once their cavity is enlarged by the binding of Ag^+ to the peripheral triazole ligands, has been investigated in diluted solutions by means of absorption and fluorescence spectroscopy. Prior to the addition of **NDI**, the Ag^+ complexes $[\text{Ag}_4(\text{1})](\text{OTf})_4$ and $[\text{Ag}_4(\text{2})](\text{OTf})_4$ were prepared by adding $\text{Ag}(\text{OTf})$ to $\text{CH}_2\text{Cl}_2/\text{MeOH}$ (90:10) solutions of the cages, as previously described.^[15,19]

Addition of **NDI** (0–463 equivalents) to a solution of $[\text{Ag}_4(\text{1})](\text{OTf})_4$ (1.8×10^{-7} M) leads to a decrease in absorption of the Soret band, as shown in Figure 2A, testifying an electronic interaction between the Zn-porphyrins of the cage and the guest. Fluorescence titration experiments were performed

upon excitation at 434 nm, isosbestic of the absorption spectra. In order to clearly analyse the trend of the emission intensity of the open cage within the titration, each emission spectrum has been subtracted by the contribution of NDI_n , collected from a “blank” solution of **NDI** prepared at the same concentration as in each titration point (see Figure S4). This was made necessary since, even if the absorbance of the formed NDI_n at 434 nm is very low (< 0.0010) in the range $0\text{--}8.5 \times 10^{-5}$ M of **NDI** used for the titration, its emission is intense enough ($\phi = 0.15$, Table S1) to show up and sum to the fluorescence of the porphyrin cage (Figure S4). The emission spectra thus derived ($\Delta I(\lambda)$), reported in Figure 2B, show a decrease of the emission intensity of $[\text{Ag}_4(\text{1})]^{4+}$ upon addition of **NDI**. The observed changes clearly indicate the formation of an inclusion complex between the $[\text{Ag}_4(\text{1})]^{4+}$ open cage and **NDI**.

In order to derive the association constant for the process of **NDI** complexation, emission titration data have been analysed both by Reactlab Equilibria^[20] and Hypspec^[21] software (the best fittings were obtained using a 1:1 complexation model) and by fitting the data with a specific equation related to a 1:1 complex formation (see Equation (1) in the Experimental Section). A couple of examples of fittings of the emission data treated with Hypspec software are reported in Figure S5. An average association constant of $(4150 \pm 39) \text{ M}^{-1}$ is derived for the formation of the $\text{NDI} @ [\text{Ag}_4(\text{1})]^{4+}$ complex. It can be noticed that a lower value of $K_a = (95 \pm 20) \text{ M}^{-1}$ was found for the formation of $\text{NDI} @ [\text{Ag}_4(\text{1})]^{4+}$ in CD_2Cl_2 , through ^1H NMR titration of $[\text{Ag}_4(\text{1})](\text{BARF})_4$ with **NDI**.^[14] Some considerations can be made to explain the discrepancy of the two constants. The present value might be overestimated by the contribution given by bimolecular external quenching occurring when the **NDI** concentration increases. Moreover, absorption and emission titrations were conducted in a low complex concentration range (about 20–30% of formed complex at the end of the titration) due to the necessity to work in optical conditions and the fittings were limited to an initial close to linear dependence range (Figure S5). Also, the formation of NDI_n species could contribute to lower the real monomeric **NDI** concentration compared to the expected one, affecting the mathematical treatment. The latter should have only a slight influence, by

Figure 2. A) Absorption spectra of $\text{CH}_2\text{Cl}_2/\text{MeOH}$ (9:1) solutions of cage $[\text{Ag}_4(\text{1})]^{4+}$ (1.8×10^{-7} M) with increasing amounts of **NDI** (0–463 equiv). B) Differential (see text) emission spectra of the same solutions as in A; excitation at 434 nm.

considering self-association constants of $1\text{--}2\text{ M}^{-1}$ reported for a naphthalene diimide in solvents of various polarity.^[22] Finally, one should consider that both the solvent and the complex counterions were different in this study with respect to the previous one,^[14] plausibly leading to a different value of the association constant. A solvophobic effect can, in fact, be operative in the present case where the more polar solvent mixture favours **NDI** encapsulation.^[23]

The same titration experiments were performed with $[\text{Ag}_4(\mathbf{2})](\text{OTf})_4$. Absorption and emission changes observed for a $2.0 \times 10^{-7}\text{ M}$ solution of $[\text{Ag}_4(\mathbf{2})]^{4+}$ titrated with **NDI** in the range 0–463 equivalents, are reported in Figure S6. The emission trend was followed upon excitation at 425 nm, isosbestic of the absorption titration, and by correcting the spectra for the emission of **NDI**_n (see Figure S7), as in the previous case. The observed changes in both absorption and emission of $[\text{Ag}_4(\mathbf{2})]^{4+}$ are similar to those observed for $[\text{Ag}_4(\mathbf{1})]^{4+}$, pointing to the same type of complexation. The titration data have been mathematically analyzed by means of Reactlab Equilibria and Hypspec software, with a 1:1 complexation model, and by using Equation (1) in the Experimental section. Examples of trends of emission intensities at selected wavelengths, analysed with Hypspec software, are reported in Figure S8. The derived average association constant for the formation of the **NDI**@ $[\text{Ag}_4(\mathbf{2})]^{4+}$ complex is $(4018 \pm 47\text{ M}^{-1})$. Also in this case, we cannot exclude an overestimation of the association constant for the reasons discussed above.

Overall, the spectral changes observed in the absorption titrations and the stronger quenching observed in the fluorescence experiments once the cages **1** and **2** are open with sil-

ver(I), endorse an assembly process where the **NDI** guest is steadily included inside the cavity of the cage. In order to gain precious information on type and geometry of the host/guest interactions at the basis of the observed phenomena, detailed computational investigation has been performed (see below).

Computational results

Two 100-ns long MD simulations of silver(I)-complexed cage **2**, $[\text{Ag}_4(\mathbf{2})]^{4+}$, in the presence of four **NDI** molecules are performed in $\text{CH}_2\text{Cl}_2/\text{MeOH}$ (90:10) at 300 K reproducing the experimental conditions (see the Methods section for a justification of this choice). Two starting configurations for the four **NDI** molecules are chosen. In the first one, the four **NDI** molecules are placed at a distance from the center of mass of the two Zn-porphyrins varying from $\approx 1\text{ nm}$ to $\approx 2\text{ nm}$ and with different orientations between the **NDI** plane and the porphyrin planes (see Figure 3A).

In this configuration the **NDI** molecules are not forced to interact with the $[\text{Ag}_4(\mathbf{2})]^{4+}$ cage in a specific way. However, at such distances both the **NDI** molecules and the cage do not behave as freely diffusing molecules in solution, as an interaction among them is already present.^[24] Thus, this choice allows both to follow an unbiased interaction among the five molecules and to speed up a possible complexation between the **NDI** molecules and the cage, bypassing the time-limiting step of the purely diffusive encountering of the solvated molecules. The analysis of the MD simulation starting from this configuration highlighted the presence of transient interactions between the **NDI** molecules and the Ag^+ ions coordinated to the

Figure 3. Top (left) and side (right) view of the first (A) and second (B) starting configurations for the silver(I)-complexed cage **2** and the four **NDI** molecules.

linkers (see below). Thus, a second starting configuration is tested in which the **NDI** molecules are placed perpendicularly to the two porphyrin planes in-between the four linkers (i.e., with each **NDI** molecule at a distance of ≈ 0.5 nm from a silver(I) ion of one linker) with an average distance between the center of mass of the **NDI** and the center of mass of the two Zn atoms of ≈ 1 nm (see Figure 3B). In this second starting configuration the interaction between each **NDI** molecule and a silver(I) ion is favored, in order to investigate the possible

role of such interaction in the formation of the **NDI**@ $[\text{Ag}_4(2)]^{4+}$ complex.

In both simulations, one **NDI** molecule spontaneously enters the cage after a few nanoseconds forming a complex that remains stable for the rest of the simulation. In the MD simulation starting from the first configuration, an **NDI** molecule (hereafter called **NDI2**) enters the cage after ≈ 6 ns (see Figure 4A). After an initial diffusion (lasting ≈ 2.5 ns) during which the distance between the cage and the **NDI2** centers of mass

Figure 4. A, E) Time evolution of the distance between the center of mass of an **NDI** molecule (**NDI2** in the MD starting from the first configuration, panel A, and **NDI4** in the MD starting from the second configuration, panel E) and the center of mass of the two Zn-porphyrins. The vertical dashed line highlights the time at which the **NDI** molecules enter the cage (≈ 6 ns and ≈ 2.2 ns for the first and second starting configurations, respectively). Representative structures of the relative position of the **NDI** molecules and the $[\text{Ag}_4(2)]^{4+}$ cage are reported. B, F) Normalized distribution of the distance between the center of mass of **NDI2/NDI4** and each of the two Zn atoms of the porphyrin rings after **NDI** enters the $[\text{Ag}_4(2)]^{4+}$ cage (6–100 ns and 2.2–100 ns for the first (B) and second (F) starting configurations, respectively). C, G) Time evolution of the distance between the two Zn atoms of the porphyrin rings. D, H) Normalized distribution of the distance between the two Zn atoms of the porphyrin rings before (0–6 ns and 0–2.2 ns for the first (D), and second (H), starting configuration, respectively, dashed lines) and after (6–100 ns and 2.2–100 ns for the first (D) and second (H) starting configuration, respectively, solid lines) **NDI** enters the $[\text{Ag}_4(2)]^{4+}$ cage.

reaches ≈ 2.7 nm, **NDI2** approaches the cage, interacting with the Ag^+ ion of one of the linkers. During the ≈ 2.5 ns in which **NDI2** interacts with the silver(I) ion, the distance between the Ag^+ ion and the center of mass of the **NDI** molecule is maintained at ≈ 0.5 nm, suggesting the presence of a cation- π interaction between the silver(I) ion and the **NDI**. Then, **NDI2** starts entering the cage. After further ≈ 1 ns the entering process is completed and **NDI2** is in the center of the cage, essentially parallel to the porphyrin planes (see below). In the MD simulation starting from the second configuration, an **NDI** molecule (**NDI4**) enters the cage after ≈ 2.2 ns (see Figure 4E). In the first ≈ 1 ns of the simulation **NDI4** interacts with the silver(I) ion, with an average distance between the Ag^+ ion and the center of mass of **NDI4** of ≈ 0.55 nm, analogously to what observed in the MD simulation starting from the first configuration.

The results of both MD simulations suggest that the interaction with the Ag^+ ion might be relevant for the complexation between the **NDI** and the $[\text{Ag}_4(2)]^{4+}$ cage, maintaining the **NDI** molecule between the Zn-porphyrin planes until the proper orientation to enter the cage cavity is found.

It can be noted from Figure 4B and F that in the stable complex, the **NDI** molecule is in the center of the cage. In fact, the normalized distributions of the distance between the center of mass of **NDI2/NDI4** and each of the two Zn^{II} metal centers of the porphyrin rings are essentially indistinguishable and peaked at the same distance (0.39 nm). Such a distance is only slightly higher than a typical π - π interaction distance (0.35 nm),^[23,25] suggesting a suboptimal electronic interaction between the **NDI** and the porphyrin rings, in agreement with the experimental binding constant measured by UV-visible and NMR titrations.^[14]

In our previous work,^[15] MD simulations showed that in the complexed $[\text{Ag}_4(2)]^{4+}$ cage the two Zn-porphyrins are highly rigid and adopt, in solution, a cofacial position with an average distance between the Zn^{II} centers of mass of ≈ 0.85 nm. Upon complexation with the **NDI** molecule a reduction of the distance between the two porphyrin macrocycles is observed

(see Figure 4C, D, G, and H). In fact, the distance between the two Zn^{II} metal ions of the porphyrins decreases from ≈ 0.85 nm to ≈ 0.78 nm upon complexation, suggesting a sort of induced-fit mechanism.

The **NDI**-cage complex was further investigated by analyzing the relative orientation of the three planes (i.e., the two porphyrin planes and the **NDI** plane). In what follows, the results obtained from the MD simulations starting from the two different conformations, being analogous, are reported together. To assess the effect of the formation of the complex on the geometry of the $[\text{Ag}_4(2)]^{4+}$ cage, the angles between each of the Zn-porphyrin plane and the vector connecting the two centers of mass (θ_1 , θ_2) are analyzed before and after the formation of the complex (see Figure 5A). It can be observed that, upon complexation with the **NDI** molecule, the porphyrins of the $[\text{Ag}_4(2)]^{4+}$ cage adopt a more cofacial conformation, 82° being the most populated angle with a sharper distribution when the **NDI** molecule is located inside the cage and 75° being the most populated one and with a broader distribution, when the **NDI** molecule is outside the cage. The two porphyrin planes are also mainly parallel, as the bi-dimensional distribution of θ_1 and θ_2 shows a strong linear correlation (see Figure S9A). The 75° value found for θ_1 and θ_2 when the **NDI** is outside the cage is in agreement with what was observed during a previously performed 100 ns-long MD simulation of the same cage in the absence of **NDI** molecules, in which the most populated angle was 78° .^[15]

The angles between the **NDI** plane and the vectors connecting the centers of mass of each porphyrin plane and of the **NDI** molecule (ϕ_1 and γ_1), as well as the angles between the Zn-porphyrin planes and the same vectors (ϕ_2 and γ_2) were also analyzed. Their normalized distribution (Figure 5B) shows that the **NDI** plane is also almost cofacial to the two porphyrin planes. It can be noted that the most populated values of ϕ_1 and γ_1 are slightly lower than the ones of ϕ_2 and γ_2 (78° vs. 82°), but such a difference is negligible and the three planes are essentially parallel and cofacial.

Figure 5. A) Normalized distribution of θ_1 and θ_2 before (dashed line) and after (solid line) the complexation of the $[\text{Ag}_4(2)]^{4+}$ cage with the **NDI** molecule. B) Normalized distribution of ϕ_1 and ϕ_2 , γ_1 and γ_2 when the **NDI**-cage complex is formed.

The angle defining the rotation of the porphyrin planes one with respect to the other (β) and the angles between the **NDI** plane and each porphyrin plane (β_1 and β_2) were also analyzed along the MD simulations (see Figure S10). In agreement with what previously found,^[15] such angles display small variations along the simulation, the most populated β angle being 26° and the most populated β_1 and β_2 angles being 14° (Figure S10).

The behavior of the three **NDI** molecules that are not complexed with the $[\text{Ag}_4(2)]^{4+}$ cage was also analyzed during the MD simulation by calculating the distance between the center of mass of each **NDI** molecule and the center of mass of the $[\text{Ag}_4(2)]^{4+}$ cage. The results obtained from the MD simulation starting from the first configuration are reported in Figure 6 and the ones obtained from the MD simulation starting from the second configuration are reported in Figure S11. It can be observed in the figures that the three **NDI** molecules that remain outside the cage interact with it by forming short-lived complexes (i.e. encounter complexes) in which the **NDI** molecules interact both with the Ag^+ ions and with the Zn-porphyrins (see representative structures in Figure 6). In addition, interactions among the three **NDI** molecules can be also observed (see Figure S12).

Photoinduced processes within the inclusion complexes

As revealed by spectrofluorimetric titration experiments, formation of the complexes **NDI**@ $[\text{Ag}_4(1)]^{4+}$ and **NDI**@ $[\text{Ag}_4(2)]^{4+}$ causes a quenching of the fluorescence of the $[\text{Ag}_4(1)]^{4+}$ and $[\text{Ag}_4(2)]^{4+}$ cages. This quenching can be ascribed to a fast photo-

induced process occurring between the units of the host and the included guest, depopulating the porphyrins singlet excited states. An energy transfer process from the cage porphyrin to the guest can be excluded from thermodynamic reasons. Indeed, the lowest singlet excited state of **NDI** is higher in energy than the cage Zn-porphyrin singlet excited state (an energy value of 3.20 eV for ^1NDI can be derived from the maximum of fluorescence emission, Figure S1, and the value of 2.06 eV for $^1[\text{Ag}_4(1)]^{4+}$ and $^1[\text{Ag}_4(2)]^{4+}$ has been previously reported^[15]). Conversely, a LUMO-LUMO electron-transfer leading to a charge-separated (CS) state with a Zn-porphyrin radical cation and a **NDI** radical anion is a feasible process. By considering a reported Zn-porphyrin oxidation potential of + 0.70 V versus SCE in CH_2Cl_2 ^[26] and a **NDI** reduction potential of -0.65 V in the same solvent (converted from -1.10 V vs. Fc/Fc^+ ^[16d,27]), in fact, the energy of the CS state can be approximately estimated at 1.4 eV, thus thermodynamically accessible from the cage Zn-porphyrin singlet excited state. Moreover, the process is strongly favoured by the close proximity of the porphyrin planes and the **NDI** π system in the fixed sandwich geometry of the complexes ($d_{\text{porphyrin-NDI}} \leq 0.4$ nm, from MD simulations).

In order to gain better insights into the photophysics of the systems, pump-probe transient absorption experiments with femtosecond resolution were performed. Higher concentration conditions were used to provide an appropriate absorbance of the solutions at 560 nm, the excitation wavelength chosen to selectively populate the Zn-porphyrin singlet and to avoid photodegradation of the silver(I)-cages. Solutions of cages **1** and **2**, ca. 6×10^{-6} M, have been treated with $\text{Ag}(\text{OTf})$ to obtain

Figure 6. Time evolution of the distance between the center of mass of each **NDI** molecule and the center of mass of the two Zn-porphyrins. Representative structures of the relative position of the **NDI** molecules and the $[\text{Ag}_4(2)]^{4+}$ cage are reported.

the open conformation and then added with **NDI** to have a sizeable amount of complex in solution (50% of complexed cage). **NDI** concentrations of 2.1×10^{-3} M and 3.6×10^{-3} M were used for $[\text{Ag}_4(1)]^{4+}$ and $[\text{Ag}_4(2)]^{4+}$, respectively. Absorption and emission spectra of the mixtures, compared with those of the open cages, are reported in Figures S13 and S14; it can be seen that the trend in the decrease of the Soret band upon complexation with **NDI** is confirmed, accompanied by a slight red-shift of the same band and the formation of a shoulder at ca. 430 nm. The ca. 50% quenching measured for the porphyrin fluorescence in the mixtures attested the amount of formed complex.

Transient absorption measurements were performed on the host–guest mixtures and on solutions of bare cages and of silver(I)-complexed cages, at the same concentrations, for comparison purposes. Figure 7 shows the time evolution of the transient absorption spectra of **1**, $[\text{Ag}_4(1)]^{4+}$ and the mixture containing **NDI**@ $[\text{Ag}_4(1)]^{4+}$, upon excitation at 560 nm. In cage **1**, the end-of-pulse spectrum presents a positive signal with maximum at ca. 470 nm superimposed with the ground state bleaching of the Q-bands, at 560 and 600 nm, and stimulated emission at 606 and 660 nm. These absorption features, typical of a Zn-porphyrin singlet excited state, evolve towards a spectrum with maximum at 476 nm, that can be safely attributed to triplet absorption. A clear isosbestic point at 504 nm and the lifetime of 1.7 ns found for signals decays and growths (see Figure S15A for the kinetic analysis at representative wavelengths) leads to identify the process as $\text{S}_1 \rightarrow \text{T}_1$ intersystem crossing. In the silver(I)-complexed cage, the picture is very similar, except for a prompt decay (13 ps, pre-exponential weight $\leq 20\%$) of the singlet signal that precedes the intersystem crossing evolution (Figure S15B). This fast event can be ascribed to static quenching between the porphyrin units forced to stay in a face-to-face conformation in the Ag^{I} -complexed cage or to a specific interaction with the Ag^+ ions.^[28] The picture for the mixture containing **NDI**@ $[\text{Ag}_4(1)]^{4+}$ is markedly different (Figure 7C). A signal rise of 1.5 ps is observed in the range 470–650 nm, then the transient spectrum decays with a lifetime of ca. 30 ps and, on longer time scales, a kinetics of 1700 ps for the decay of the singlet signal and rise of the triplet spectrum, is detected (Figure S15C). Although weak, the spectral features of the species that forms in 1.5 ps and decays in 30 ps can be safely assigned to the CS state P^+-NDI^- (with P: cage porphyrin). In fact, the Zn-porphyrin cation is known to have a broad absorption between 600 and 700 nm^[29] whereas the **NDI** radical anion has characteristic absorption bands at ca. 475 (most intense), 610, 680 and 760 nm.^[16c,30] It can be noticed that the amount of formed triplet is approximately half that observed for the $[\text{Ag}_4(1)]^{4+}$ solution, indicating that the **NDI**-free cages undergo the normal intersystem crossing path, whereas the ones complexed with **NDI** experience fast charge separation followed by charge recombination to the ground state.

The same analysis has been performed for **2**, $[\text{Ag}_4(2)](\text{OTf})_4$ and the mixture containing **NDI**@ $[\text{Ag}_4(2)](\text{OTf})_4$; the results are presented in Figures S16 and S17. Features similar to the previous case are observed for the transient absorption evolution of

Figure 7. Time evolution of the transient absorbance of $\text{CH}_2\text{Cl}_2/\text{MeOH}$ (9:1) solutions of A) cage **1** (5.8×10^{-6} M), B) cage $[\text{Ag}_4(1)]^{4+}$ (5.8×10^{-6} M) and C) mixture of the latter with **NDI** 2.1×10^{-3} M; excitation at 560 nm ($A_{560} = 0.050$, 0.2 cm optical path). The spectral region around 560 nm is covered by scattering of the excitation beam.

2 and its silver(I)-complex (Figure S16A and B): intersystem crossing occurs with a time constant of 1.6 ns in **2**, and this process is preceded by a fast decay of ca. 4 ps for $[\text{Ag}_4(2)]^{4+}$ (Figure S17A and B). In the mixture, at a difference with the previous case, the rise of the absorption bands of the radicals

is not observed, and the end-of-pulse spectrum already contains weak CS features superimposed with the singlet absorption of the guest-free cage (Figure S16C). This implies that charge separation occurs within the time resolution of the system, that is, 300 fs. This spectrum then evolves with a decay of 38 ps and, on longer time scales, the triplet features appear as a result of triplet population in the free cages present in solution (Figures S16C and S17C).

Overall the data indicate that a photoinduced host–guest electron transfer process occurs in both complexes upon excitation of the porphyrin components on fast and ultrafast scales (1.5 ps and ≤ 300 fs for $\text{NDI}[\text{Ag}_4(1)]^{4+}$ and $\text{NDI}[\text{Ag}_4(2)]^{4+}$, respectively). The faster rate for the latter is likely due to the shorter distance between the Zn-porphyrin and **NDI**, allowed by the longer flexible linkers of cage **2**.^[15] Charge recombination occurs with a similar kinetics in the two systems, with lifetime of the CS state of ca. 30–40 ps (an energy level diagram and the observed rate constants are reported in Scheme 1). These values reflect the high efficiency of both processes, likely due to the very short distance between donor and acceptor units in the formed inclusion complexes, as evidenced by the MD simulation and in accordance with Marcus electron transfer theory.^[31]

Conclusions

A thorough study of the encapsulation of **NDI** in Zn-porphyrin molecular cages endowed with triazoles-including connectors of different lengths have been performed by means of absorption and emission spectroscopies and molecular dynamics sim-

ulations. Whereas the photophysical properties of the cages are overall not affected by the presence a large excess of **NDI**, addition of silver(I) ions triggers **NDI** encapsulation and a strong quenching of the cage Zn-porphyrin emission. Molecular dynamics simulation shows that the stable host–guest complex forms within a few nanoseconds, probably mediated by favorable cation– π interactions between the Ag^+ ions and **NDI** molecules. Silver(I) binding controls the overall disposition of the Zn-porphyrins in the cages in a face to face geometry that finely adjust upon **NDI** encapsulation to optimize the π donor– π acceptor interactions, as shown by the simulation. In these Ag^+ -triggered **NDI** inclusion complexes a photoinduced electron transfer process thus occurs with almost unitary efficiency from the excited Zn-porphyrin components to **NDI**, in 1.5 ps for $\text{NDI}[\text{Ag}_4(1)](\text{OTf})_4$ and ultrafast scales (≤ 300 fs) for $\text{NDI}[\text{Ag}_4(2)](\text{OTf})_4$. Following charge separation, charge recombination between the oxidized Zn-porphyrin and the reduced **NDI** in both inclusion complexes occurs in tens of picoseconds. The present study demonstrates the Ag^+ ions allosteric control on the encapsulation of the guest and, consequently, on the host–guest photoinduced electron transfer and on the photophysical properties of the cages. Such control opens the way to develop fine-tuned sensors, signaling receptors or photoreactors.

Experimental Section

Absorption and emission spectroscopy: Spectroscopic grade solvents from Merck were used as received. Silver trifluoromethanesulfonate ($\text{Ag}(\text{OTf})$), from Sigma–Aldrich, was stored under argon in a sealed vial in dark and dry conditions. Freshly prepared $\text{Ag}(\text{OTf})$ solutions were always used. Solutions of cage **2** were filtered with 0.2 μm membrane filters (Minisart RC4 filters) prior to each measurement, to avoid aggregation issues.

Absorption and emission spectra were recorded with a PerkinElmer Lambda 650 spectrophotometer and a Edinburgh FLS920 fluorimeter, equipped with a Peltier-cooled Hamamatsu R928 PMT (280–850 nm), respectively. The reported emission spectra are uncorrected for the phototube response. Titration experiments were performed by incremental addition of micro aliquots of stock solutions of **NDI** ($8 \times 10^{-4} \text{ M} \times 10^{-3} \text{ M}$) to a solution of silver(I)-complexed cage (ca. $2 \times 10^{-7} \text{ M}$). The final volume added was kept below 10% of the total volume, to avoid dilution of the porphyrin. The experiments have been conducted avoiding light exposure of the solutions.

Titration data have been analyzed with Reactlab Equilibria and Hypspec softwares and according to Equation (1), derived for a 1:1 host–guest stoichiometry:

$$Obs = Obs_0 + (\Delta Obs_{max} / 2 S_0) \times \{K_d + X + S_0 - [(K_d + X + S_0)^2 - 4 X S_0]^{1/2}\} \quad (1)$$

where Obs_0 is the signal (absorbance or emission intensity) of the free $[\text{Ag}_4(1)]^{4+}$, ΔObs ($Obs - Obs_0$) is the change in the optical signal with respect to that of the free $[\text{Ag}_4(1)]^{4+}$, S_0 is the constant concentration of $[\text{Ag}_4(1)]^{4+}$, X is the variable concentration of **NDI**, ΔObs_{max} is the maximum variation of the observed signal between non-complexed and complexed states and $K_d = 1/K_a$ is the dissociation constant of the 1:1 complex.^[10c,k,33]

Scheme 1. Energy level diagram and derived kinetics of the photoinduced processes in the inclusion complexes $\text{NDI}[\text{Ag}_4(1)]^{4+}$ (red) and $\text{NDI}[\text{Ag}_4(2)]^{4+}$ (blue). P stands for cage porphyrin. The singlet and triplet energy levels are taken from data of the present paper and from literature.^[15,32]

Emission lifetimes were determined by using an IBH time-correlated single-photon counting apparatus with excitation at 560 nm for porphyrin cages and at 331 nm and 465 nm for **NDI** and **NDI_n**, respectively.

Pump-probe transient absorption measurements were performed with an Ultrafast Systems HELIOS (HE-VIS-NIR) femtosecond transient absorption spectrometer by using, as excitation source, a Newport Spectra Physics Solstice-F-1K-230 V laser system,^[34] combined with a TOPAS Prime (TPR-TOPAS-F) optical parametric amplifier (pulse width: 100 fs, 1 kHz repetition rate, selected output wavelength: 560 nm). The overall time resolution of the system is 300 fs. Air-equilibrated solutions in 0.2 cm optical path cells were analyzed under continuous stirring. To reduce photo-degradation, the pump energy on the sample was reduced to 2 μJ/pulse. Surface Explorer V4 software from Ultrafast Systems was used for data acquisition and analysis. The 3D data surfaces were corrected for the chirp of the probe pulse prior to analysis.

Estimated errors are 10% on transient absorbance lifetimes, 10% for luminescence lifetimes, 10% for molar absorption coefficients and 10% on quantum yields. The reductions in emission intensity and lifetime observed for the “closed” cages treated with **NDI** can be considered reliable even if they fall within instrumental errors, due to the maintenance of the same experimental conditions and to the consistency of the decrease determined with two different techniques.

Molecular dynamics simulations: MD simulations of silver(I)-complexed cage **2**, [Ag₄(**2**)⁴⁺], in the presence of 4 **NDI** molecules are performed in CH₂Cl₂/MeOH (90:10) to mimic the experimental conditions at the end of the titrations with **NDI**, in which the number of equivalents of **NDI** is around 400. Despite the high number of equivalents, the experimental **NDI** concentration is very low ($\approx 10^{-5}$ – 10^{-6} M). Given the limited size of the simulation box (compared to what would be needed to reproduce the very low experimental concentration), to find a compromise between **NDI** concentration and number of equivalents and to avoid over-crowding of the simulation box, we had to drastically reduce the number of **NDI** molecules from around 400 to 4. The initial structure of the silver(I)-complexed cage **2** was taken from a previous MD simulation in the same solvent.^[15]

MD simulations are performed with the GROMACS software package using the GROMOS force field. The previously derived GROMOS force field parameters and atomic charges for the silver(I)-complexed cage **2** are used.^[15] The GROMOS force field parameters and atomic charges for the **NDI** molecules are obtained from the Automated Topology Builder (ATB).^[35]

The silver(I)-complexed cage **2** and the four **NDI** molecules are placed in a dodecahedral box large enough to contain the molecule and at least 1.0 nm of solvent on all sides with a proper number of CH₂Cl₂ molecules to reproduce the density of CH₂Cl₂ at 300 K and 1 bar (1.33 g cm⁻³). A proper number of CH₂Cl₂ molecules was then substituted with the corresponding number of MeOH molecules to reproduce the experimental CH₂Cl₂/MeOH (90:10) proportion. The force field parameters for CH₂Cl₂ and MeOH are taken from the GROMACS topology database.

The initial structure of the silver(I)-complexed cage **2** is placed in the center of the box and two starting configurations for the four **NDI** molecules are chosen. In the first one, the four **NDI** molecules are placed below the cage cavity with randomly selected orientations between the **NDI** plane and the porphyrin planes and with distances between the center of mass of the **NDI** and the center of mass of the two Zn atoms varying from ≈ 1 nm to ≈ 2 nm (average distance ≈ 1.7 nm, see Figure 3 A). In the second starting configura-

tion the **NDI** molecules are placed perpendicularly to the two porphyrin planes in-between the four linkers with an average distance between the center of mass of the **NDI** and the center of mass of the two Zn atoms of ≈ 1 nm (see Figure 3 B).

Simulations are carried out in the NVT (i.e., with constant number of molecules, volume and temperature) ensemble at a constant temperature of 300 K using the velocity rescaling temperature coupling.^[36] The LINCS algorithm^[37] is used to constrain bond lengths and a time step of 2 fs for numerical integration of the equations of motion is used. The particle mesh Ewald method^[38] is used for the calculation of the long-range interactions and a cut-off of 1.1 nm is used. After a solute optimization and a subsequent solvent relaxation, each system is gradually heated from 50 to 300 K using short MD simulations. The trajectories were then propagated for 100 ns for each system. Coordinates are saved at every 1 ps.

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Conflict of interest

The authors declare no conflict of interest.

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FULL PAPER

■ Host–Guest Systems

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■■■ – ■■■

Allosteric Control of Naphthalene
Diimide Encapsulation and Electron
Transfer in Porphyrin Containers:
Photophysical Studies and Molecular
Dynamics Simulation

Ag⁺ ions as chemical stimuli to control
naphthalene diimide encapsulation and

photoinduced electron transfer in por-
phyrin containers.

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