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Selective Excited-State Dynamics in a Unique Set of Rationally Designed Ni Porphyrins

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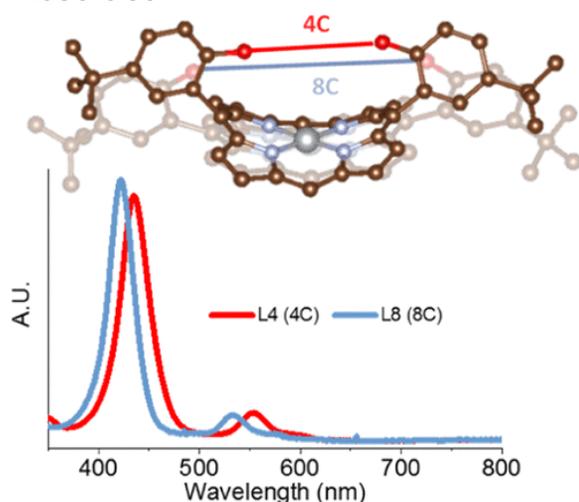
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Abstract



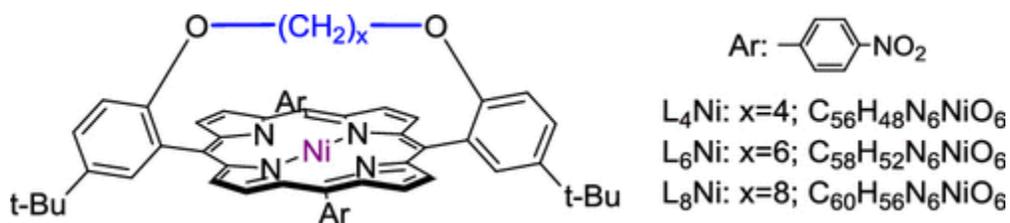
In this work, we report the design and photophysical properties of a unique class of Ni porphyrins, in which the *tert*-butyl benzene substituents at the meso positions of the macrocycle were tethered by ethers with alkyl linkers. This not only results in the permanently locked *ruf* distortion of the macrocycle but also enables the engineering of the degree of distortion through varying the length of alkyl linkers, which addressed the complication of uncertainty in the specific structural distortions that has long plagued the porphyrin photophysical community. Using advanced time-resolved optical and X-ray absorption spectroscopy, we observed tunability in the excited-state relaxation pathway depending on the degree of distortion and characterized the associated transient intermediate structure. These findings provide a new avenue to afford accessibility to a wide range of excited-state properties in nonplanar porphyrins.

Introduction

Metalloporphyrins represent versatile model systems for various biophysical processes ranging from photosynthesis, oxygen transport, and oxygen activation.(1–3) The active research on these systems in the past decades has significantly advanced the understanding of their structure and photochemical properties, which in turn led to the discovery of many new applications including photodynamic therapy,(4,5) light emitting diodes,(6,7) and solar energy conversion.(8–10) Previous studies on their electronic and photophysical

properties demonstrated that the macrocycle ring in metalloporphyrins is highly flexible and often undergoes various deformations from planarity due to functionalization with bulky peripheral substituents, which dramatically alters their chemical and photophysical properties from their planar analogues.(11,12) For example, nonplanar distortion due to either *ruf* or *sad* deformation can tune the redox potential,(13–15) axial binding affinity of center metal,(14,16,17) basicity of N atoms, and so forth,(18,19) the properties that directly dictate their biological function for proteins with the porphyrin cofactor. Substituent-induced nonplanar distortion can also significantly impact the ground-state optical properties and energetic ordering of excited states (ESs), leading to tunable relaxation dynamics,(12,18,20–30) which are the key photophysical properties that determine the light harvesting and charge transport function of photosynthetic pigments in proteins and solar energy conversion.

While these previous studies have certainly provided important insights into the correlation of electronic and chemical properties of metalloporphyrins with macrocycle conformational distortion, the distortion was achieved via substitution of the meso or β positions of the porphyrin periphery with steric crowding substituents.(11,31,32) Under these conditions, different conformations can be adopted by the macrocycle and perturbed by external environment, leaving significant uncertainty in probing their structural features and photophysical properties. Moreover, the assignment of *sad*, *ruf*, or other distortions largely rely on calculations or crystal structures, which are certainly dynamic in the solution phase where photophysical studies are performed. As a result, it is highly needed to have a permanently locked model system for which the distortion is known to be one of the fixed distortions such that the effects of the specific distortion on the chemistry and photochemical behavior can be unambiguously determined and distinguished from the effects of other types of distortions. In this work, we report the design and photophysical studies of a unique class of Ni porphyrins (Scheme 1), in which the meso positions of the macrocycle were tethered by ethers that are attached to the *t*-butyl benzene substitutes.(33) Using advanced time-resolved optical and X-ray absorption spectroscopy (XAS), we observed tunability in the ES relaxation pathway dependent on the degree of distortion and also determined the associated transient intermediate structure.



Scheme 1. Design Principle of Permanently Distorted Ni Porphyrins

Experimental Section

Synthesis

Strapped nickel porphyrin L8-Ni was prepared according to previously published standard methods.(33) 2,2'-[Octane-1,8-diylbis(oxy)]di(5-*tert*-butyl)benzaldehyde (4.66 g, 10 mmol) and bis(2-pyrrolyl)-4-nitrophenylmethane (5.34 g, 20 mmol) were added to a boiling propionic/butyric (2:1, v/v) acid mixture (200 mL) in small portions within 5 min. Ni(OAc)₂·4H₂O (~2.7 g, 11 mmol) was added to the solution after 15 min under reflux; the mixture was further stirred for 45 min, and then, the mixed acids were removed under reduced pressure. The solid residue was extracted three times with dichloromethane. The extracting solution was washed five times with water and dried with sodium sulfate. After removal of the solvent, the residue was separated by silica gel column chromatography with dichloromethane/petroleum ether (1:2, v/v) to give purple solid L8-Ni 1.27 g. Yield: 12.5%. UV–vis (chloroform, 293 K): λ_{max} = 421, 510, 578 nm. HR MS (ESI): found for C₆₀H₅₆N₆NiO₆ [M]⁺, 1014.3616; calcd, 1014.3609. All reagent grade reactants and solvents were used as received from chemical suppliers.

Characterization

UV–visible spectroscopy was performed using an HP Agilent 8453 spectrometer. The high-resolution mass spectrum was obtained by using a Varian QFT-MALDI mass spectrometer. Crystal structure determination by X-ray diffraction was performed by using a Bruker SMART 1000 charge-coupled device (CCD) diffractometer equipped with a normal focus, 2.4 kW sealed tube X-ray source, using monochromatic Mo K α ($\lambda = 0.71073$ Å) radiation. A suitable single crystal of each compound was glued to a thin glass fiber with superglue. The SMART program package was used to determine the unit cell parameters and for data collection. The structure was solved by direct methods and were refined on F^2 by the full-matrix least-squares method by using the SHELXT-97 program. The positional parameters for the Ni, N, and O atoms were obtained by direct methods for the compound. All hydrogen atoms were placed geometrically and were refined by riding on their appropriate O and C atoms. The final positions of all nonhydrogen atoms were refined anisotropically. Crystals of L8-Ni were obtained by solvent diffusion in chloroform and methanol.

Optical Transient Absorption

Optical transient absorption (OTA) measurements were performed using a Spectra-Physics Solstice Ti:sapphire amplifier [120 ps full width at half-maximum (fwhm), 800 nm, 1 kHz]. Pump pulses were generated with a TOPAS OPA using 75% of the Ti:sapphire output and were chopped at 500 Hz for transient absorption measurement. A portion of the remaining 800 nm output was directed into a Helios transient absorption spectrometer (Ultrafast Systems). Time delay between the pump and probe is achieved by a delay stage on the probe line (800 nm). The probe is then focused onto a sapphire crystal for white light generation (430–780 nm). Samples were prepared in a screw-top 2 mm quartz cuvette with PTFE-lined septa. The concentration was 0.1 mM in toluene, purged with N₂ before measurement. The sample was stirred during measurement using a magnetic stirrer and PTFE-coated stir bar. UV–visible spectroscopy was used to confirm that laser exposure did not result in sample decomposition. Soret band or Q-band excitations were performed, respectively, using 420 or 530 nm pulses with 0.3 μ J/pulse.

Nanosecond transient absorption measurements were performed using an Edinburgh Instruments LP900 flash photolysis spectrometer. The pump pulse was generated by an Nd:YAG (Spectra-Physics, INDI-10) with the third harmonic (355 nm) output. The 355 nm output was converted to 525 nm using an optical parametric oscillator (Spectra-Physics). The probe source is a pulsed 450 W xenon arc lamp, which was detected before and after laser exposure by the integrated CCD detector.

X-ray Transient Absorption Spectroscopy

X-ray transient absorption spectroscopy was performed at the Advanced Photon Source at Argonne National Laboratory in beamline 11-ID-D. The storage ring was under the standard operating mode, with 80 ps fwhm X-ray pulses that were used as the probe. The 100 fs 400 nm pump pulse (535 mW) was the second harmonic output of an Nd:YLF Ti:sapphire regenerative amplified laser at 10 kHz. The sample (1 mM in toluene) was placed in an N₂-purged double-walled flask cooled to 5 °C and was continuously flowed through a stainless steel tube using a peristaltic pump to produce a 550 μ M sample jet. The jet is positioned between two avalanche photodiodes that collect the X-ray fluorescence signals. Soller slits with Co filters were positioned between the sample and photodiode detectors. Delay scans were collected at the peak of the negative XTA difference signal, which corresponded to the on-edge 1s–p₂ feature, the largest feature in the difference spectrum. The full-energy range-extended X-ray absorption fine structure (EXAFS) spectra were then collected at the pump delay that produced the largest difference signal, approximately 50 ps for all samples. Because the fwhm of X-ray pulses are 80 ps, the difference spectrum observed is insensitive to pump delay, as XTA can only probe the cooled MC state in Ni porphyrins.

X-ray Absorption Data Analysis

XTA produced two X-ray absorption spectra: laser-on and laser-off spectra. The difference signal was generated by laser-on minus laser-off. First, the ground-state (laser-off) spectra were fit using the Artemis module of the Demeter software package. The free energy force field (FEFF) models were constructed using a single molecule with atomic coordinates directly taken from the CIF files. In order to accurately quantify the first-shell Ni–N distance, second-shell and multiple-scattering paths were included into the model. The ground-state spectra in XTA experiments were used for steady-state FEFF analysis. The k -range used for the Fourier transform into R -space was approximately 1.7–8.6 for all data sets.

Once satisfactory fits were obtained for the ground-state (laser-off) spectra, the laser-on spectra were fit. Because the excitation pulse does not excite all of the molecules in the X-ray beam path, the laser-on spectrum is actually a combination of both excited and nonexcited molecules. To account for this, two amplitude-weighted FEFF models were used to fit the data, where the amplitude is the ES fraction. The first model was the same model used for the ground-state molecule, with fixed Δ_{enot} , N , R , and σ^2 . The second model consisted of the same paths; however, R and σ^2 were allowed to vary to account for their new values in excited molecules. The ES fraction was fixed during this fitting. Various ES fractions were used to determine a reasonable value (Figure S6) by following the R -factor of the resulting fit. It was determined that an ES fraction of 0.3 best represented the set of porphyrins investigated and was used to determine ES parameters. Finally, the modeled ES R -space spectrum was generated by a single FEFF model with ES values for R and σ^2 with 100% amplitude weighting.

Results and Discussion

As shown in Scheme 1 and Figure S1, the alkyl linker tends to pull two side rings that connected with the *tert*-butyl benzene group, resulting in a locked *ruf* distortion of the macrocycle. Moreover, the degree of distortion can be systematically tuned by varying the length of the ether-tethered alkyl strap group enabling an unambiguous study on the correlation of their conformational distortion with photophysical properties. As shown in Scheme 1, the length of the alkyl group was varied to include 4 C atoms (L4), 6 C atoms (L6), and 8 C atoms (L8). The synthetic details and characterization of permanently distorted Ni porphyrins (L4 and L6) have been reported previously.⁽³³⁾ L8 was prepared in this work to further decrease the degree of distortion, and its synthesis is outlined in the Experimental Section. The ruffling effect should be more pronounced as the chain length reduces from L8 to L4. The crystal structures confirmed this trend (Figure S1 and Table S1), namely, the macrocyclic distortion effectively induces a contraction of the N_4 core, where the core diameter (L_{NN}) decreases from 3.89 Å in L8, to 3.81 Å in L6, and 3.78 Å in L4. Such a result is related to the compression of size (L_{CC}) between the tethered meso carbon atoms from 6.72 to 6.50, and 6.34 Å, respectively.

The electronic and local structure of the Ni porphyrin analogues were investigated by XAS via the Ni K edge. The X-ray absorption near-edge structure (XANES) of each sample is compared in Figure 1a. Two distinct features are observed on the rising edge, including a sharp feature at ~ 8.338 keV, corresponding to the $1s-4p_z$ transition and the white line transition at the absorption edge (~ 8.360 keV), corresponding to the $1s-4p_{x,y}$ transition.^(12,25,34) The whole energy spectra show blue shifts above-edge, suggesting the decrease of the Ni–N bond length from L8 to L4. To gain quantitative insights into the nature of these features, the EXAFS spectra (Figure S2) of these samples were analyzed using FEFF fitting with the Demeter software package.⁽³⁵⁾ Figure 1b shows the Fourier-transformed R -space spectra along with their best fits. It can be clearly seen that significant decreases of Ni–N first shell bond lengths are observed with decreasing tether length/increasing degree of distortion, indicating that distortion results in closer Ni–N distance, consistent with XANES results. These values are slightly shorter than the average distances observed in the X-ray crystal structure (L8, 1.944 Å; L6, 1.910 Å; and L4, 1.892 Å), which is likely explained by the measurement environment

and conditions, as XAS was performed in the solution phase, while the X-ray crystal structures were obtained in the solid state. The full FEFF fitting parameters are given in Tables S2–S4.

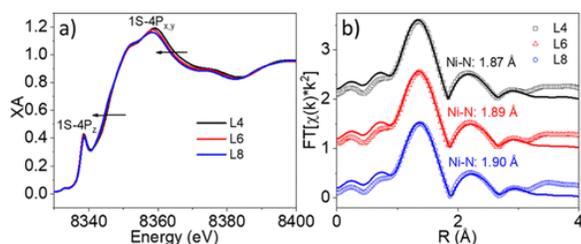


Figure 1. (a) XANES spectra for L4–L8 at the Ni K edge. (b) FEFF fitting results of the EXAFS data with the obtained first shell bond distances with uncertainty of approximately ± 0.02 Å. Data are shown as open points and best fits as solid lines.

Figure 2a shows the UV–visible absorption spectra of Ni porphyrins, which are featured by an intense band in the UV region (Soret band, S_0 – S_2 transition) and relatively weaker bands in the visible region of the spectrum (Q-band, S_0 – S_1 transitions).^(11,21,36) A clear redshift trend is observed from L8–L4 as a function of distortion of the porphyrin macrocycle. From the molecular orbital diagrams,⁽³⁶⁾ the lowest energy-allowed transitions consist of occupied $a_{1u}(\pi)$ and $a_{2u}(\pi)$ initial states and unoccupied $e_g(\pi^*)$ final states. Upon ruffling of the porphyrin macrocycle, redshifts in the UV–visible spectra were observed due to destabilization of the $a_{1u}(\pi)$ and $a_{2u}(\pi)$ molecular orbitals.^(11,20) Therefore, the trend observed in L8–L4 demonstrates that systematic distortion results in shifting of the Soret band (Figure 2a) and Q-band (inset) transition energies, consistent with literature reports.

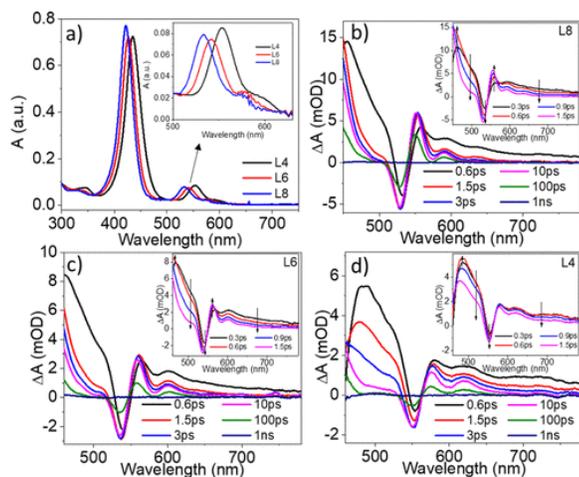


Figure 2. UV–visible spectra (a) for Ni-porphyrin analogues with the inset showing the expanded Q-band region. OTA spectra after 400 nm excitation of the S_0 – S_2 transition for L8 (b), L6 (c), and L4 (d). Insets of (b–d) display the early time spectra and highlighted spectral evolution on this timescale.

Femtosecond OTA spectroscopy was performed to investigate the ES dynamics of these Ni porphyrins. Figure 2b–d shows the OTA spectra for L8, L6, and L4, respectively, following 400 nm excitation which triggers Soret band transition. The initial spectrum of the least-distorted sample L8 (0.3 ps, inset of Figure 2b) is featured by a positive band in the 460–500 nm region, broad absorption at the 550–780 nm region, and a bleach band centered at ~ 530 nm that can be assigned to the ground-state bleach (GSB) of the Q-band. Distinct spectral evolutions were observed for these features within 2 ps, including the decay of the positive feature at 460–500 nm, the growth of the Q-band GSB, decay of the 550–780 nm absorption, and the formation of a positive feature at 553 nm. The presence of a clear isosbestic point at 562 nm between the latter two features suggest these evolutions correspond to the same relaxation process. Because similar spectral

evolution was observed in the transient absorption spectra following Q-band excitation (Figure S3), this evolution cannot be attributed to the relaxation of S_2-S_1 . In addition, the absence of stimulated emission following the Soret band excitation excludes the contribution from the S_1 state. These results, which are in good agreement with previous reports that S_2-S_1 and $S_1-(3d_x^2-y^2, 3d_z^2)$ (denoted as (d,d)) relaxation is ultrafast and beyond our instrument resolution of 300 fs, suggest that the early time spectral evolution can be attributed to intramolecular vibrational relaxation (IVR) from the hot (d,d) state.(22–24,27,37–39) Next, the OTA spectra show two more distinct spectral evolutions, including (1) the whole spectrum showing slight blueshift which is accompanied by relatively slow decay within 2–20 ps and (2) the simultaneous decay of all transient features without spectral shift >20 ps. While the spectral evolution for the latter can be attributed to the returning of transient species to the GS, we assign the former to the vibrational cooling (VC) process from a hot (d,d) state to cooled (d,d) configuration.(34–36) L6 (Figure 2c) appears to show similar dynamics to L8, significantly slower spectral changes on both the VC and IVR timescales in the more distorted L4 analogue (Figure 2d), as well as a shorter ES lifetime.

To quantitatively understand these relaxation dynamics and elucidate trends as a function of structural distortion, we fit the decay kinetics at the described spectral regions with multiexponential rise/decay functions. The excited state absorption (ESA) in Figure 3a is described predominantly by slow processes relative to the ESA in Figure 3b. From the Q-band bleach (Figure 3c), it is evident that the early-time growth on a linear scale is followed by at least two distinct decay components as revealed by log scaling after the break. Based on both the kinetic and spectral observations, a triexponential model was employed to quantitatively describe the ES dynamics, with fitting parameters shown in Table 1. For nearly all traces, τ_1 , corresponding to an intramolecular vibrational redistribution (IVR) process, is 0.5 ps for L6 and L8 and 1.33 ps for L4, although for L6 and L8, τ_1 corresponds to a rise in the 562 and 553 nm traces, respectively. The Q-band bleach also rises for all samples with τ_1 , in agreement with the spectral observations. After IVR, the kinetics for L4 are markedly different from that of L6 and L8 with a trend at some wavelengths, most clearly demonstrated in the positive feature on the red-side of the Q-band bleach (Figure 3d). At this feature, τ_2 which represents VC is fit with a growth component in L8 while L6 and L4 decay in this region. This process is accompanied by a growth in the Q-band bleach in L4, while L6 and L8 show Q-band decay for τ_2 . Finally, τ_3 , which corresponds to the return of the cooled (d,d) state to GS, shows a clear trend with respect to structural distortion, where L8 has the longest lifetime and L4 has the shortest. The ESA at 690 nm (Figure 3e) closely resembles the decay at the probe wavelengths shown in Figure 3b. A scheme showing the fit model is presented in Figure 3f. The Q-band excitation data could be fit by the same model (Figure S4 and Table S5), implying that the same relaxation processes are observed regardless of excitation wavelength.

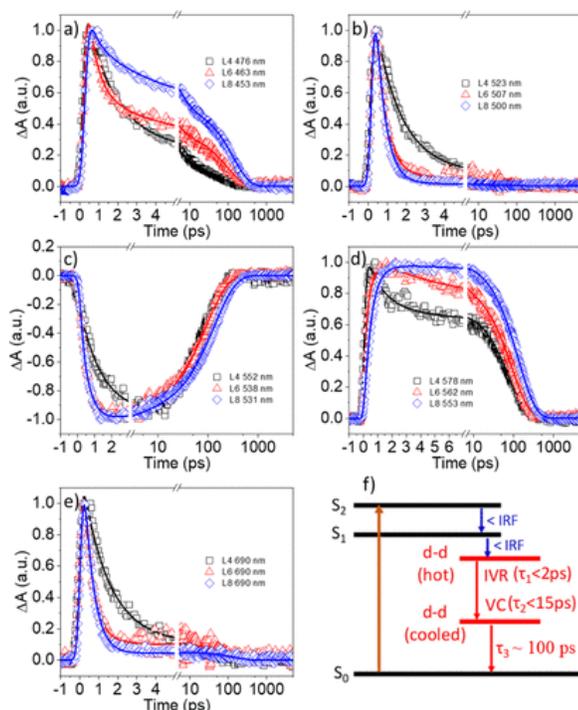


Figure 3. Kinetic traces representing the broad positive feature in OTA spectra on the blue-side of Q-band bleach (a,b), Q-band bleach (c), derivative-shaped positive feature (d), and broad vis–NIR feature probed at 690 nm (e). (f) Scheme showing the three-component model used to fit OTA data.

Table 1. OTA Kinetic Fit Parameters, Soret Excitation

sample	probe (nm)	τ_1 (ps)	A_1 , %	τ_2 (ps)	A_2 , %	τ_3 (ps)	A_3 , %
L8	453	0.5	14.6	4.05	39.8	133	45.6
	500		98.8		0.54		0.66
	531		42.7		-4.11		-53.1
	553		-54.0		-2.23		43.5
	690		96.0		1.07		2.93
L6	463	0.5	56.6	3.14	20.8	91.5	22.6
	507		94.5		3.60		1.90
	538		47.3		-3.28		-49.6
	562		-33.7		12.5		53.8
	690		93.7		0.10		6.2
L4	476	1.33	68.6	8.2	22.4	57.96	9.00
	523		88.3		9.08		2.62
	552		44.60		5.70		-49.70
	578		33.80		4.40		61.80
	690		87.0		8.36		4.64

The differences observed in OTA imply a fundamental difference in the ES relaxation cascades as a function of structural distortion. For L4, both τ_1 and τ_2 are longer but τ_3 is shorter than those in L8 and L6, which directly implies that the more distorted (d,d) state could prevent the IVR and IC relaxation while facilitate the relaxation from the cooled (d,d) state to GS. This dependence of the ES deactivation pathway on the degree of distortion is

unprecedented for Ni porphyrins. One explanation for the slower IVR and VC processes is related to the relaxation to the cooled d–d state being radiationless and depositing ca. $10\,000\text{ cm}^{-1}$ of energy into macrocycle vibrational modes. However, the highly distorted L4 porphyrin is most restricted to *ruf* modes and, therefore, has less vibrational degrees of freedom than L6 and L8.

The highly altered photodynamics for the most highly distorted L4 porphyrin might suggest that the final Ni d–d state ($3d_x^2-y^2$, $3d_z^2$) that forms via vibrational relaxation through the macrocycle modes(26,40) could be different in nature for L4 compared to L6 and L8. To investigate the nature of the vibrationally cooled state, X-ray transient absorption was performed at the Ni K-edge, which has previously been used to demonstrate the Ni d–d ($3d_x^2-y^2$, $3d_z^2$) character of the final state.(24) In the current study, the fwhm of synchrotron pulses used for time-resolved measurements is approximately 80 ps, which can only probe the MC state. XTA experiments were performed using 400 nm laser pump and Ni K-edge X-ray probe (Figure S5). The laser-off spectra constitute the GS spectra, while the laser-on spectra represent a combination of ES molecules excited by the laser pulse and GS molecules that were not excited. Therefore, an ES fraction was incorporated into the FEFF fitting model in order to extract ES information (Figure S6).

FEFF fitting was performed using the optimized ES fraction of 0.3, meaning that the laser excitation pulse excites 30% of molecules in the X-ray beam path while 70% of molecules remain in their ground state (Figure S7). The resulting fits are shown in Figure 4a–c along with the modeled ES spectrum. From comparison of the peak positions in the first shell representing Ni–N distance, it is qualitatively observable that macrocycle core expansion about the Ni metal center is observed in the ES, in agreement with previous reports.(23,24,27,28,40) The fit values for R_{ES} indicate that first shell expansion is similar for these samples with expansion occurring from $1.87 \rightarrow 1.94\text{ \AA}$ for L4 (Figure 4a), from $1.89 \rightarrow 1.95\text{ \AA}$ for L6 (Figure 4b), and from $1.90 \rightarrow 1.96\text{ \AA}$ for L8 (Figure 4c). The lifetime of this Ni d–d ($3d_x^2-y^2$, $3d_z^2$) ES tracked by delay scans (Figure 4d, fit parameters in Table S6) for each sample shows a good agreement with the long-lived state in OTA measurements (τ_3), further supporting that this is the cooled (d,d) state and also confirming that the Ni centered d–d state deactivation is faster in the most distorted analogue.

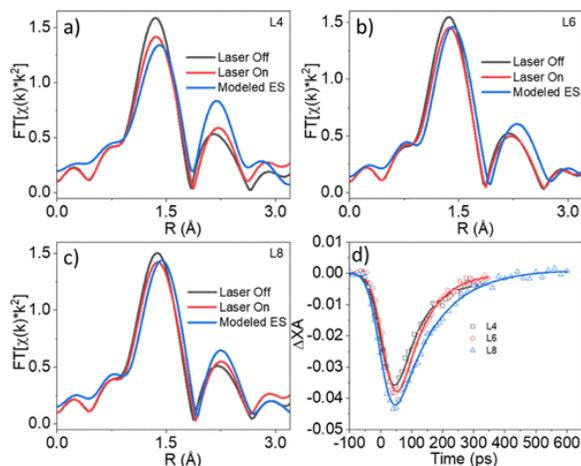


Figure 4. *R*-space data for XTA experiments. Laser-off, laser-on, and modeled ES spectra in *R*-space for L4 (a), L6 (b), and L8 (c). The ES Ni–N distance is shown in each figure. Delay scans at the maximum negative difference signal for L4–L8 and best fits (d).

One of the most interesting findings of this work is that increased distortion leads to faster deactivation of the Ni d–d ($3d_x^2-y^2$, $3d_z^2$) ES. The cooled Ni d–d state involving the expanded core macrocycle involves an interplay between macrocycle and metal-centered states. It has been previously shown that ruffling influences the Ni d–d energetics.(27) Specifically, it was shown for a sterically distorted porphyrin species that the $3d_x^2-y^2$ orbital

mixed with the porphyrin π orbitals and resulted in a small degree of charge transfer from the macrocycle to metal. The observation that a larger degree of permanent distortion leads to faster ES decay in this study implies the involvement of the *ruf* mode of the macrocycle in relaxation of the Ni d–d ES. These effects could be a combination of dynamic, including nonadiabatic effects, or static effects.

Interestingly, distortion leads to faster ES deactivation also for free-base analogues in the triplet manifold (Figures S8 and S9). It can also be observed that the ES features in free-base ns-OTA and Ni-metallated fs-OTA data are significantly broadened as a function of distortion (Figure S10) despite the most distorted macrocycle being the most restricted and having the fewest vibrational degrees of freedom. Therefore, the contribution of dynamic interconversion with out-of-plane modes other than *ruf* can be disregarded. We suggest that anharmonic coupling of various in-plane modes and the *ruf* mode is enhanced due to distortion, leading to broadening of the ES features in both the Ni metallated and free-base cases, and to enhanced ES decay.

Conclusions

In summary, we report the design and photophysical properties of a unique class of Ni porphyrins permanently locked in *ruf* distortion resulting from the tethered meso positions of the macrocycle by ethers that are attached to the *tert*-butyl benzene substitutes. We show that by varying the length of ether chains, the degree of distortion can be systematically controlled. Using the combination of advanced OTA and XTA spectroscopy, which allows us to directly probe the structural dynamics of Ni porphyrins, we demonstrate that ES relaxation dynamics can be highly tuned by the degree of distortion. The IVR and VC processes are slowed significantly in permanently distorted porphyrins due to reduced vibrational degrees of freedom. Furthermore, it was shown that the lifetime of the final Ni d–d state formed in the relaxation cascade is dependent on the degree of distortion, where higher distortion leads to enhanced ES decay. These results suggest that these permanently distorted porphyrins are a useful scaffold for investigating macrocycle-metal ES dynamics, implying vast utility in the porphyrin community.

The authors declare no competing financial interest.

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