A strong steric hindrance effect on ground state, excited state, and charge separated state properties of a CuI-diimine complex captured by X-ray transient absorption spectroscopy


Total number of authors: 11

Published in:
Dalton Transactions

Link to article, DOI:
10.1039/c4dt02046d

Publication date:
2014

Document Version
Publisher's PDF, also known as Version of record

Link back to DTU Orbit

Citation (APA):
L. X. Chen et al.,
A strong steric hindrance effect on ground state, excited state, and charge separated state properties of a Cu(I)-diimine complex captured by X-ray transient absorption spectroscopy.
A strong steric hindrance effect on ground state, excited state, and charge separated state properties of a CuI-diimine complex captured by X-ray transient absorption spectroscopy


Photophysical and structural properties of a CuI diimine complex with very strong steric hindrance, [CuI(dppS)2]+ (dppS = 2,9-diphenyl-1,10-phenanthroline disulfonic acid disodium salt), are investigated by optical and X-ray transient absorption (OTA and XTA) spectroscopy. The bulky phenylsulfonic acid groups at 2,9 positions of phenanthroline ligands force the ground state and the metal-to-ligand charge-transfer (MLCT) excited state to adopt a flattened pseudo-tetrahedral coordination geometry in which the solvent access to the copper center is completely blocked. We analyzed the MLCT state dynamics and structures as well as those of the charge separated state resulting from the interfacial electron injection from the MLCT state to TiO2 nanoparticles (NPs). The OTA results show the absence of the sub-picosecond component previously assigned as the time constant for flattening, while the two observed time constants are assigned to a relatively slow intersystem crossing (ISC) rate (~13.8 ps) and a decay rate (100 ns) of the [CuI(dppS)2]+ MLCT state in water. These results correlate well with the XTA studies that resolved a flattened tetrahedral Cu(I) coordination geometry in the ground state. Probing the 3MLCT state structure with XTA establishes that the 3MLCT state has the same oxidation state as the copper center in [CuI(dppS)2]2+ and the Cu–N distance is reduced by 0.06 Å compared to that of the ground state, accompanied by a rotation of phenyl rings located at 2,9 positions of phenanthroline. The structural dynamics of the photoinduced charge transfer process in the [CuI(dppS)2]+/TiO2 hybrid is also investigated, which suggests a more restricted environment for the complex upon binding to TiO2 NPs. Moreover, the Cu–N bond length of the oxidized state of [CuI(dppS)2]+ after electron injection to TiO2 NPs shortens by 0.05 Å compared to that in the ground state. The interpretation of these observed structural changes associated with excited and charge separated states will be discussed. These results not only set an example for applying XTA in capturing the intermediate structure of metal complex/semiconductor NP hybrids but also provide guidance for designing efficient CuI diimine complexes with optimized structures for application in solar-to-electricity conversion.

Introduction

CuI diimine complexes, with similar light absorption to those of quintessential ruthenium complexes, have attracted much attention because of their potential applications in solar energy conversion1–5 chemical sensing6,7 and molecular devices.8,9 Although these complexes have been studied for decades, studies in the past few years have focused on their functions that are aligned with those performed by Ru(II) polypyridyl complexes in solar energy conversion. CuI diimine complexes have strong potential to replace the less abundant Ru(II) complexes used in solar energy conversion, because in addition to their similar photophysical properties of the MLCT (metal-to-ligand charge-transfer) state to that of Ru(II) complexes, CuI diimine complexes are more environmentally and economically viable.10–15 Furthermore, the MLCT state of CuI diimine complexes, due to its electron transfer from the copper center to the ligand, yields a 3d electronic configuration that is susceptible to the Jahn–Teller distortion to transform from a...
pseudo-tetrahedral geometry to $D_{2d}$ symmetry to a flattened tetrahedral geometry with $D_2$ symmetry. Such excited state structural movements have been explored for functions of molecular machines.\textsuperscript{12} Therefore, it is important to gain detailed knowledge on structural and environmental control of the excited state properties of the Cu\textsuperscript{I} diimine complexes before they are employed in different applications in solar energy conversion.

Perhaps the most interesting aspect of these complexes is their structure and environment dependent photophysical properties, such as the MLCT state lifetimes, intersystem crossing rates and luminescence quantum yields. Also, the MLCT state of such complexes undergoes a transformation in the coordination geometry due to the different preferences of the electronic configurations of 3d\textsuperscript{10} Cu(\textit{i}) in the ground state and 3d\textsuperscript{9} Cu(\textit{ii}) in the MLCT state.\textsuperscript{16} As summarized in a recent review,\textsuperscript{17} the MLCT state properties of a series of Cu\textsuperscript{I} bis-phenanthroline based complexes can be altered via different steric hindrance exerted by the substituents at the 2,9 positions of the phenanthroline ligands that can modulate the angle between the two phenanthroline ligand planes from 90°, such as in the complex with 2,9-di-tert-butyl-1,10-phenanthroline ligands with an orthogonal tetrahedral coordinating geometry, to $\sim$68°, a “flattened” tetrahedral geometry, such as in less hindered species that are capable of undergoing the flattening.\textsuperscript{4,8,13,17–24} It has been observed that when the two phenanthroline ligand planes are forced to be orthogonal, the intersystem crossing (ISC) from the 1MLCT to 3MLCT state occurs on a timescale of a few picoseconds, while the same process takes 10–15 ps when the two ligand planes are not orthogonal,\textsuperscript{19,23–25} i.e. a flattened tetrahedral geometry. According to the calculations by Siddique \textit{et al.},\textsuperscript{24} the spin–orbit coupling coefficients are lower for the molecular orbitals stabilized by a flattened coordinating geometry due to the changes in the symmetry from $D_{2d}$ to $D_2$, resulting in a relatively slow ISC rate. In addition, the 3MLCT excited state lifetimes and emission quantum yields are also sensitive to the capability of an “exciplex” formation with a solvent molecule, no matter whether the solvent is considered as coordinating or non-coordinating with the copper center. It has been shown that the presence of bulky groups at 2,9 positions can block solvent access to the Cu center to prevent the formation of an “exciplex”, lowering the energy gap between the MLCT and the ground states and accelerating the ground state recovery, and thus result in a longer MLCT lifetime and a higher emission quantum yield.\textsuperscript{14,15,18,26–29} These studies suggest that the excited state dynamics of Cu\textsuperscript{I} diimine complexes could be controlled by the dihedral angle between the two phenanthroline ligand planes as well as the solvent accessibility to the Cu center.

Previously, we have investigated the structural influence on the excited state dynamics in an extreme case of a completely locked tetrahedral coordination geometry due to the sterically-bulky tert-butyl groups in bis(2,9-di-tert-butyl-1,10-phenanthroline)-copper(\textit{i}), [Cu\textsubscript{I}(dtbp)\textsubscript{2}]\textsuperscript{+}.\textsuperscript{19} We showed that [Cu\textsubscript{I}(dtbp)\textsubscript{2}]\textsuperscript{+} has a three microsecond 3MLCT lifetime, no sub-picosecond component that was previously assigned as the Jahn–Teller distortion of flattening of the pseudo-tetrahedral coordination geometry and a few picosecond intersystem crossing time constant. Because sufficient lifetimes and energetics are prerequisites for many photochemical processes as well as the sensitivity of these Cu\textsuperscript{I} excited states to their structures, we want to investigate whether the 1MLCT and 3MLCT state lifetimes can be simultaneously prolonged in the Cu\textsuperscript{I} phenanthroline complexes with locked flattened geometry in both the ground and excited states. Furthermore, if these excited state lifetimes are extended by structural tuning, we may achieve sufficient efficiency for interfacial charge injection from these complexes to semiconductor nanoparticles in the dye-sensitized solar cells.\textsuperscript{2,17,25}

In this report, we examine the structural influence on the excited state dynamics of [Cu\textsubscript{I}(dppS)\textsubscript{2}]\textsuperscript{+} (dppS = 2,9-diphenyl-1,10-phenanthroline disulfonic acid disodium salt), which represents another extreme case in which the complex is almost locked into the flattened coordination geometry. In our recent paper,\textsuperscript{25} we have reported an efficient electron injection process directly from the 1MLCT state of [Cu\textsubscript{I}(dppS)\textsubscript{2}]\textsuperscript{+} to TiO\textsubscript{2} NPs followed by an ultraslow back electron transfer process. This observation has been attributed to the already flattened ground state geometry of [Cu\textsubscript{I}(dppS)\textsubscript{2}]\textsuperscript{+}, which prolongs the ISC time to enable ET from the 3MLCT state. While the successful electron injection has been studied in great detail, the quantitative structural analysis of the MLCT state as well as the charge separated state in this Cu(\textit{i}) diimine complex have not been reported. Here, our investigation is focused on the detailed structural dynamics of these transient states of the [Cu\textsubscript{I}(dppS)\textsubscript{2}]\textsuperscript{+} complex in the excited state in H\textsubscript{2}O and the charge separated state formed after electron injection to TiO\textsubscript{2} NPs as well as their correlation with the ground state structure. Optical transient absorption (OTA) spectroscopy is used to determine excited MLCT state kinetics by monitoring the MLCT excited state absorption of [Cu\textsubscript{I}(dppS)\textsubscript{2}]\textsuperscript{+} in the visible region. An analogous method in the X-ray regime, X-ray transient absorption (XTA) spectroscopy, is used to track electron configurations and the corresponding molecular structural changes of [Cu\textsubscript{I}(dppS)\textsubscript{2}]\textsuperscript{+} by measuring the transient X-ray absorption near edge structure (XANES) and the extended X-ray absorption fine structure (EXAFS).\textsuperscript{17,19,22,23,25,30–34} We would like to answer the following questions: (1) what are the structures of the MLCT state and charge separated state structures; (2) what are the structural changes of the complex when it binds to TiO\textsubscript{2} nanoparticles; and (3) what can we learn from our study on Cu\textsuperscript{I} diimine complex design for its applications in solar energy conversion? Understanding the molecular structures and dynamics of both photoexcited states and charge separated intermediates in the current system will not only provide guidance by revealing their practical applications in dye sensitized solar cells but also an insight into the nature of photochemical reactions.

Results and discussion

Ground state photophysical properties of [Cu\textsubscript{I}(dppS)\textsubscript{2}]\textsuperscript{+} in H\textsubscript{2}O

The UV-vis absorption spectrum of [Cu\textsubscript{I}(dppS)\textsubscript{2}]\textsuperscript{+} in H\textsubscript{2}O (blue) is presented in Fig. 1b. The spectra of [Cu\textsubscript{I}(phen)\textsubscript{2}]\textsuperscript{+} (black)
and $[\text{Cu}^1(\text{dppS})_2]^+$ (red) in dichloromethane are also shown in Fig. 1b for comparison. Analogous to those of $[\text{Cu}^1(\text{phen})_2]^+$ and $[\text{Cu}^1(\text{dpp})_2]^+$, the spectrum of $[\text{Cu}^1(\text{dppS})_2]^+$ is typically characterized by an intense main absorption peak in the 420–470 nm region and a shoulder feature at lower energy, both of which have been assigned to MLCT bands based on previous studies$^{17,19,22,25,34,35}$ with the latter assigned to the MLCT transition from the flattened coordination geometry. Apparently, with the increasing steric hindrance at the 2,9 positions of phenanthroline groups, i.e. $[\text{Cu}^1(\text{phen})_2]^+ < [\text{Cu}^1(\text{dpp})_2]^+ < [\text{Cu}^1(\text{dppS})_2]^+$ (Fig. 1a), the maximum position of the main absorption peak shifts to blue, accompanied by the increasing relative intensity of the low-energy shoulder to the main absorption peak, indicating the correlation between the steric hindrance and the MLCT transitions. According to the DFT calculations by Siddique et al.,$^{24}$ both excitation energies and oscillator strengths of MLCT transitions in Cu$^1$ phenanthroline complexes, corresponding to the transitions between the HOMO and LUMO orbitals, are strongly dependent on the dihedral angles of the two phenanthroline planes. When the dihedral angle of the two phenanthroline planes is 90°, the two transition energies are nearly degenerate, therefore only the main peak in the 440–460 nm region is expected in the absorption spectrum, which is represented by the extreme case of $[\text{Cu}^1(\text{dtbp})_2]^+$ as observed in our previous studies.$^{34}$ When the dihedral angle deviates from 90° through a flattening distortion from $D_{2d}$ to $D_2$, the two MLCT transition energies split, leading to a red-shift of the low-energy shoulder and a blue shift of the main MLCT absorption band. This is clearly seen in Fig. 1b, where the main absorption band shows a blue shift and the shoulder shows a red shift in the most sterically hindered $[\text{Cu}^1(\text{dppS})_2]^+$. As shown here as well as in our previous study,$^{14}$ the relative intensity of the low-energy shoulder indicates the degree of dynamic flattening distortion in the ground state likely through low frequency rocking modes of the pseudo-tetrahedral coordination geometry in $D_{2d}$ symmetry,$^{35}$ with an increased shoulder intensity correlated with a larger flattening distortion. Therefore, the highest relative intensity shoulders in the spectrum of $[\text{Cu}^1(\text{dppS})_2]^+$ compared to those of $[\text{Cu}^1(\text{phen})_2]^+$ and $[\text{Cu}^1(\text{dpp})_2]^+$ suggested the largest degree of flattening as well as other accompanying distortions. These results all support that $[\text{Cu}^1(\text{dppS})_2]^+$ with bulky groups at 2,9 positions of the phenanthroline groups adapts a likely flattened geometry in the ground state, which is verified from our XANES results to be discussed later.

**Excited state dynamics of the $[\text{Cu}^1(\text{dppS})_2]^+$ complex in H$_2$O**

Fig. 2a shows the femtosecond transient absorption (fs-TA) spectra of the $[\text{Cu}^1(\text{dppS})_2]^+$ complex in H$_2$O in the visible region after 415 nm excitation. The entire spectra exhibit a broad positive absorption, and can be assigned to the MLCT state absorption. The spectral evolution for $[\text{Cu}^1(\text{dppS})_2]^+$ in H$_2$O was dominated by ISC from the $^1$MLCT to the $^3$MLCT state at the early times (<25 ps) and intrinsic $^3$MLCT state decay at later times (>25 ps). Fig. 2b shows the kinetics trace of the $[\text{Cu}^1(\text{dppS})_2]^+$ complex in H$_2$O at 570 nm. The kinetic trace fits a dual-exponential function which reveals two decay time
constants, \( \tau_1 = 13.7 \text{ ps} \) and \( \tau_2 \gg 3 \text{ ns} \). \( \tau_1 \) is consistent with the ISC time constant previously observed in \([\text{CuI}(\text{dpp})_2]\) \(^{22}\) (dpp = 2,9-diphenyl-1,10-phenanthroline) and in \([\text{CuI}(\text{dmp})_2]\) \(^{20,21,23}\) (dmp = 2,9-dimethyl-1,10-phenanthroline) after the excited state flattening distortion,\(^{20,21,23,24}\) and thus can be attributed to the ISC from the \( ^1\text{MLCT} \) to the \( ^3\text{MLCT} \) excited state. Compared to \([\text{CuI}(\text{tbp})_2]\) with two phenanthroline ligand planes locked to orthogonal orientation with each other, the ISC rate in the \([\text{CuI}(\text{dppS})_2]\) complex is several times slower. Meanwhile, this complex does not have a sub-ps component in its TA kinetics corresponding to the flattening as in \([\text{CuI}(\text{dmp})_2]\) \(^{20,21,23}\) and structural reorganization as in \([\text{CuI}(\text{dpp})_2]\) \(^{22}\), indicating that its severe steric hindrance prevents the Jahn–Teller distortion in the excited state, which will be further discussed in the XTA results. As discussed in previous studies, the prolonged ISC time constant is likely an indication of a flattened tetrahedral coordination geometry for the Cu(I) center in the ground state due to the strong steric hindrance of the two sulfonated phenyl ligands at the 2,9 positions of phenanthroline. Consequently, the spin–orbit coupling is significantly weakened and hence results in a slower ISC process and a prolonged \( ^1\text{MLCT} \) lifetime of 13.7 ps. The long decay time constant (>3 ns) can be accurately determined by nanosecond absorption measurements (ns-TA) and assigned to the \( ^3\text{MLCT} \) state lifetime. The ns-TA spectra of the \([\text{CuI}(\text{dppS})_2]\) complex in H\(_2\)O after 420 nm excitation (Fig. 2c) show the dominating excited state absorption in the whole spectral window, similar to the results in the fs-TA measurement (Fig. 2a). The kinetics monitored at the probe wavelength of 570 nm (Fig. 2d) yields a \( ^3\text{MLCT} \) lifetime of \( \sim 100 \text{ ns} \), similar to that of \([\text{CuI}(\text{dpp})_2]\) in ethanol or acetonitrile.\(^{23}\) In contrast, the \( ^3\text{MLCT} \) lifetime of \([\text{CuI}(\text{phen})_2]\) is only a few picoseconds even in non-coordination solvents, dichloromethane,\(^{14}\) and that of \([\text{CuI}(\text{dmp})_2]\) is \(< 2 \text{ ns} \) in coordination solvents.\(^{23}\) Therefore, the prolonged \( ^3\text{MLCT} \) lifetime of \([\text{CuI}(\text{dppS})_2]\) suggests a complete shielding of the Cu center from the solvent, preventing the formation of an “exciplex” with the coordination solvent molecules. As mentioned earlier, the sub-ps rising component attributed to the excited state flattening dynamics in other Cu complexes\(^{20,23}\) was not observed in this system. These observations further support the assumption that the ground state of the \([\text{CuI}(\text{dppS})_2]\) \(^3\) complex is already locked in a “flattened” geometry, and hence no further flattening occurs in the excited state. These results together indicate that the \( ^3\text{MLCT} \) excited state of the \([\text{CuI}(\text{dppS})_2]\) complex formed after excitation converts to the \( ^3\text{MLCT} \) excited state without undergoing Jahn–Teller distortion, followed by \( ^3\text{MLCT} \) state decay with a 100 ns time constant.

### Structural dynamics of the \([\text{CuI}(\text{dppS})_2]\) \(^3\) complex in H\(_2\)O

X-ray absorption spectroscopy (XAS) was used to directly extract the structures of \([\text{CuI}(\text{dppS})_2]\) in H\(_2\)O for the ground and excited states as shown in Fig. 3. The Cu K-edge XANES (X-ray absorption near edge structure) spectrum for the ground state of \([\text{CuI}(\text{dppS})_2]\) \(^3\) (black plot, Fig. 3a) recorded before the laser pulse arrives shows a rather pronounced peak in the middle of the transition edge attributed to the \( 1s \rightarrow 4p_z \) transition at 8.986 keV as assigned for the first row transition metal coordination environments from comprehensive studies of different model complexes,\(^{36}\) and has been used to directly probe the dihedral angle of the two phenanthroline ligand planes of the Cu(I) complex in the solution.\(^{36}\) Such a capability is enabled by both experimental and theoretical studies showing that the intensity of \( 1s \rightarrow 4p_z \) edge features varies significantly for different Cu(I) complexes with various coordination geometries.\(^{36}\) The \( 1s \rightarrow 4p_z \) peak intensity is associated with the vacancy of the \( 4p_z \) orbital of the first row transition metal complexes, which changes with the coordination geometry. Hence, the more prominent \( 1s \rightarrow 4p_z \) peak intensity observed in \([\text{CuI}(\text{dppS})_2]\) \(^3\) than in the \([\text{CuI}(\text{phen})_2]\) \(^3\) complex\(^{34}\) suggests a more “flattened” tetrahedral coordination geometry for the copper center in the ground state, which is consistent with the MLCT transition analysis based on its UV-visible absorption spectrum.

X-ray transient absorption (XTA) spectroscopy\(^{32,37–41}\) is used to study the excited state structural dynamics of \([\text{CuI}(\text{dppS})_2]\) \(^3\) in H\(_2\)O. The XANES spectrum of \([\text{CuI}(\text{dppS})_2]\) \(^3\) in H\(_2\)O with laser excitation is shown in Fig. 3a (red plot). The laser-on spectrum is recorded at 150 ps after the laser pulse pump, corresponding to the \( ^3\text{MLCT} \) state. Due to the presence of both ground state and excited state molecules in the system, the laser-on XANES spectrum represents an algebraic sum of both the MLCT excited and ground states at 150 ps time delay between the laser pump and the X-ray probe pulses. Therefore, the MLCT state XANES spectrum needs to be extracted by subtracting the contribution of the remaining ground state from the total spectrum. Based on the energy distinction of the shoulder feature for Cu(I) species at 8.986 keV, the excited state spectrum (blue curve) was obtained by subtracting 55% of the ground state from the laser-on spectrum until the shoulder feature at 8.986 keV disappears. Similar to the previous studies on Cu(I) dimine complexes,\(^{17,22,25,30,31,14}\) the following spectral changes in the MLCT state compared to that of the ground state were observed: (1) the transition edge energy shifts to 3 eV higher, suggesting the higher positive charges in the copper center; (2) the shoulder feature at 8.986 keV, representing a 1s-to-4p\(_z\) transition disappears, typically observed for
transformation from CuI to CuII; (3) a pre-edge feature at 8.979 keV appears which is attributed to a 1s–3d transition that was only present for the CuII (d9). These results for the ground and MLCT states for the [CuI(dppS)2]+ complex are in good agreement with previously reported spectra for Cu diimine complexes and suggest a Cu(II) center in the MLCT state.

Fig. 3b compares the XAFS spectra in R-space after Fourier transformation for the ground state (black open dot) and laser illuminated [CuI(dppS)2]+ [red solid plot]. The Fourier transforms of these XAFS spectra were processed within the k-range of 2.6–7.6 Å⁻¹ with a k³ weighting. The peaks in these spectra present a radial distribution of Cu-neighboring atom distances in the complex without the phase corrections. The first peak, the Cu–N scattering paths, shifts to a smaller R for the laser illuminated [CuI(dppS)2]+ compared to that in the ground state. The magnitudes as well as the peak positions in the R-range of 2–3.5 Å for laser illuminated [CuI(dppS)2]+ reveal differences from that of the ground state [CuI(dppS)2]⁺. The peaks in this region correspond to the Cu–C distances with one distance fixed at its corresponding

Table 1  XAFS fitting parametersa of [CuI(dppS)2]⁺ in H2O

<table>
<thead>
<tr>
<th>Vector</th>
<th>N</th>
<th>R (Å)</th>
<th>σ² (Å²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cu–N (GS)b</td>
<td>4</td>
<td>2.00</td>
<td>0.009</td>
</tr>
<tr>
<td>Cu–N (ES)b</td>
<td>4</td>
<td>1.94</td>
<td>0.006</td>
</tr>
<tr>
<td>Cu–C1 (GS)</td>
<td>4</td>
<td>2.68</td>
<td>0.002</td>
</tr>
<tr>
<td>Cu–C1 (ES)</td>
<td>4</td>
<td>2.65</td>
<td>0.001</td>
</tr>
<tr>
<td>Cu–C2 (GS)</td>
<td>4</td>
<td>2.90</td>
<td>0.002</td>
</tr>
<tr>
<td>Cu–C2 (ES)</td>
<td>4</td>
<td>2.85</td>
<td>0.010</td>
</tr>
<tr>
<td>Cu–C3 (GS)</td>
<td>4</td>
<td>3.18</td>
<td>0.001</td>
</tr>
<tr>
<td>Cu–C3 (ES)</td>
<td>4</td>
<td>3.00</td>
<td>0.001</td>
</tr>
<tr>
<td>Cu–C4 (GS)</td>
<td>4</td>
<td>3.32</td>
<td>0.002</td>
</tr>
<tr>
<td>Cu–C4 (ES)</td>
<td>4</td>
<td>3.25</td>
<td>0.001</td>
</tr>
</tbody>
</table>

a The fitting precision for the distance is ±0.01 Å, and 0.5 for N, the coordination number. b GS: ground state. ES: excited state.

Cu–ligand distance obtained from ground state spectrum fitting with 55% fraction to account for the contribution of the ground state molecules to the laser illuminated spectrum. Using this approach, the laser illuminated spectrum can be adequately fit with an average Cu–N distance of 1.94 Å for [CuI(dppS)2]⁺ in the MLCT excited state, which is shortened by 0.06 Å relative to the ground state [CuI(dppS)2]⁺. In addition, the Cu–C1 and Cu–C2 bond distances (indicated in Scheme 1) are 0.03 and 0.05 Å shorter, respectively, in the MLCT state than in the ground state. This distance change is expected because C1 and C2 atoms in the phenanthroline are directly connected to the four Cu-ligating N atoms. It is interesting to note that a larger shortening of the Cu–C3 bond distance 0.18 Å is observed, while the Cu–C4 bond distance reveals a relatively small change with 0.07 Å shorter than in the ground state. This change suggests that the larger change of the Cu–C3 bond distance compared to the Cu–C4 distance may be due to the phenyl ring rotation, shortened by 0.06 Å relative to the ground state [CuI(dppS)2]⁺.

In addition to the solution [CuI(dppS)2]⁺ MLCT excited state structure, we also investigated the geometry change of the complex upon its attachment onto TiO2 nanoparticles (NPs). Fig. 5a compares the XANES spectra of [CuI(dppS)2]⁺ in solution and on the TiO2 nanoparticle surface. In both cases, the 1s–4pₓ transition energy remains the same at 8.986 keV, while the intensity of the latter is lowered, accompanied by a simultaneous increase of the white line intensity at 8.997 keV. Because the correlations between the 1s–4pₓ transition intensity and the angle between the ligand planes have been established in comparing the intensity of this feature between the tetrahedral [CuI(dmp)2]⁺ and flattened [CuI(dpp)2]⁺ in our

![Fig. 4 XAFS spectra (open dot) and the best fit (solid line) of [CuI(dppS)2]⁺ in H2O. (a) R-space, ground state, (b) k-space, ground state, (c) R-space, excited state, (d) k-space, excited state.](image)

![Scheme 1 Atom centers used in XAFS data fitting.](image)
previous studies,\textsuperscript{22} the reduced intensity can be attributed to a possible increase of the angle between the two phenanthroline ligand planes (or less flattened). It is known that the oxidation of Cu\textsuperscript{I} to Cu\textsuperscript{II} in [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+} is not only featured by the reduced 1s–4p\textsubscript{z} transition, but also featured by the edge shift to higher energy, the emergence of the 1s–3d pre-edge feature at 8.979 keV, as well as the reduced band at 8.996 keV. None of these additional features were observed in the XANES spectrum of the [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+}/TiO\textsubscript{2} hybrid, suggesting that the formation of [Cu\textsuperscript{II}(dppS)\textsubscript{2}]\textsuperscript{2+} in the [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+}/TiO\textsubscript{2} hybrid can be excluded. These results indicate that a less flattened geometry of [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+} was observed in the [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+}/TiO\textsubscript{2} hybrid.

A comparison of the XAFS spectra of [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+} in H\textsubscript{2}O and on TiO\textsubscript{2} NPs in R-space after Fourier transformation is presented in Fig. 5b. Compared to [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+} in H\textsubscript{2}O, the Cu–N peak for the [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+}/TiO\textsubscript{2} hybrid showed enhanced intensity. The increased peak amplitude, in general, can be attributed to an increase of the coordination number or a narrower conformation distribution due to restricted environments, i.e. smaller Debye–Waller factor. In this case, due to the two bulky sulfonated phenyl groups in the 2,9 positions of phenanthroline which blocks the solvent access, it is unlikely that ligation can occur in the hybrid. Therefore, we attributed the increased Cu–N peak amplitude to the more restricted environment caused by binding to TiO\textsubscript{2} nanoparticles which further restricted the dynamic movement of the ligands. This result can be confirmed by comparing the Debye–Waller factor of the Cu–N bond in the [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+}/TiO\textsubscript{2} hybrid sample with [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+}. The best fits of XAFS spectra of [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+}/TiO\textsubscript{2} at R and k-space are shown in Fig. 6 and their fitting results are listed in Table 2. A notably reduced Debye–Waller factor of the Cu–N bond was observed in the [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+}/TiO\textsubscript{2} hybrid, which is consistent with the assignment of a more restricted environment of [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+} after attaching to TiO\textsubscript{2} NPs. All of these results suggest that the complex adopted a less flattened geometry after linking to TiO\textsubscript{2} NPs.

We have previously demonstrated efficient electron injection from the MLCT state of [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+} to TiO\textsubscript{2} nanoparticles using a combined EPR, TA, and XTA spectroscopy characterization.\textsuperscript{25} Electron injection to TiO\textsubscript{2} NPs was unambiguously determined by probing the formation of the electron injection product, \textit{i.e.} oxidized state [Cu\textsuperscript{II}(dppS)\textsubscript{2}]\textsuperscript{2+}, monitoring the XANES region of the spectrum collected at 9 ns when the MLCT state population in the hybrid can be neglected. In order to extract the intermediate structure of [Cu\textsuperscript{II}(dppS)\textsubscript{2}]\textsuperscript{2+} in the [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+}/TiO\textsubscript{2} hybrid associated with the electron transfer process in the hybrid, here we analyzed its XAFS spectrum taken at 9 ns delay, which represents [Cu\textsuperscript{II}(dppS)\textsubscript{2}]\textsuperscript{2+} as a result of the oxidized [Cu\textsuperscript{II}(dppS)\textsubscript{2}]\textsuperscript{2+} after the electron injection. The XAFS spectra for [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+} in the [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+}/TiO\textsubscript{2} hybrid at R- and k-space after Fourier transformation as well as their best fits are presented in Fig. 7. The structural parameters for the distance from the copper center to neighboring shells from the XAFS analysis are also listed in Table 2. Analogous to the previous fitting approach, the laser initiated spectrum was fitted using a two bond model with one distance fixed at the same parameters for ground state fitting. In this

\begin{table}[h]
\centering
\caption{XAFS fitting parameters of [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+}/TiO\textsubscript{2}}
\begin{tabular}{llll}
\hline
Vector & \(N\) & \(R\) (Å) & \(\sigma^2\) (Å\textsuperscript{2}) \\
\hline
Cu–N (GS) & 4 & 1.99 & 0.006 \\
(OS) & 4 & 1.94 & 0.009 \\
Cu–C\textsubscript{1}(GS) & 4 & 2.48 & 0.001 \\
(OS) & 4 & 2.44 & 0.007 \\
Cu–C\textsubscript{2}(GS) & 4 & 2.74 & 0.005 \\
(OS) & 4 & 2.72 & 0.004 \\
Cu–C\textsubscript{3} (GS) & 4 & 3.31 & 0.001 \\
(OS) & 4 & 3.47 & 0.010 \\
Cu–C\textsubscript{4} (GS) & 4 & 3.55 & 0.002 \\
(OS) & 4 & 3.58 & 0.001 \\
\hline
\end{tabular}
\textsuperscript{a} OS: oxidized state.
\end{table}

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{fig7}
\caption{Comparison of the XANES (a) and the Fourier-transformed XAFS (b) spectra between [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+} in H\textsubscript{2}O (black) and [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+}/TiO\textsubscript{2} (red) in the ground state.}
\end{figure}

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{fig6}
\caption{Comparison of the XAFS spectra in R- (a) and k-space (b) for [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+}/TiO\textsubscript{2} in the ground state.}
\end{figure}

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{fig5}
\caption{Comparison of the XAFS spectra in R- (a) and k-space (b) for [Cu\textsuperscript{I}(dppS)\textsubscript{2}]\textsuperscript{+}/TiO\textsubscript{2} taken at 9 ns after laser excitation (75% ground state and 25% oxidized state).}
\end{figure}
case, the fraction for the ground state of \([\text{Cu}^I(\text{dppS})_2]^+\) is 75% to account for the contribution from the remaining ground state to the laser initiated spectrum. From the fitting results, we observed that the Cu–N bond distance in the oxidized state of \([\text{Cu}^I(\text{dppS})_2]^+\) in the hybrid is decreased by 0.05 Å compared to that in the ground state. Similar to the results observed for \([\text{Cu}^I(\text{dppS})_2]^+\) in H₂O, Cu–C₁ and Cu–C₂ bond distances also decrease with the Cu–N distance, which is expected because C₁ and C₂ are directly connected to the four Cu-ligating N atoms in the phenanthroline rings. In comparison with the MLCT state structure, the Cu in the phenanthroline rings. In comparison with the MLCT state, the Cu–N distance, which is expected because C₁ and C₂ are directly connected to the four Cu-ligating N atoms in the phenanthroline rings. In comparison with the MLCT state structure, the Cu–N distance, which is expected because C₁ and C₂ are directly connected to the four Cu-ligating N atoms in the phenanthroline rings. In comparison with the MLCT state structure, the Cu–N distance, which is expected because C₁ and C₂ are directly connected to the four Cu-ligating N atoms in the phenanthroline rings. In comparison with the MLCT state structure, the Cu–N distance, which is expected because C₁ and C₂ are directly connected to the four Cu-ligating N atoms in the phenanthroline rings. In comparison with the MLCT state structure, the Cu–N distance, which is expected because C₁ and C₂ are directly connected to the four Cu-ligating N atoms in the phenanthroline rings.

**Conclusions**

Our structural and dynamics studies on the \([\text{Cu}^I(\text{dppS})_2]^+\) complex in water and in a constrained \([\text{Cu}^I(\text{dppS})_2]^+\)/TiO₂ hybrid revealed a flattened coordination geometry in both environments for the complex, which ensured a slower ISC rate that allows the interfacial electron injection to TiO₂ from a \(^1\)MLCT state. The structural difference around the copper
center at the interface with the TiO2 nanoparticle from that in water suggested a highly constrained environment which causes distortion of the \( [\text{CuI(dppS)}_2]^+ \) structure via binding. Also observed are the structural differences in the "MLCT state compared to the ground state with shortening of the Cu–N distance due to the stronger electrostatic interactions between the metal center and the ligand in solution. However, the structural change between the oxidized intermediate \( [\text{Cu}^{\text{n}}(\text{dppS})_2]^+/\text{TiO}_2 \) and its ground state is significantly less due to the constraints in the environment. The capability of obtaining structural details of an environment dependent photoinduced interfacial charge transfer system provides opportunities to theoretically model these systems to gain insight into the transient structure–function correlation on the time scale of the charge separation, and structural guidance in material design.

Acknowledgements

We acknowledge the support from the U. S. Department of Energy, Office of Science, Office of Basic Energy Sciences, under contract no. DE-AC02-06CH11357. The use of the Advanced Photon Source at Argonne National Laboratory was supported by the U. S. Department of Energy, Office of Science, Office of Basic Energy Sciences, under contract no. DE-AC02-06CH11357. The authors would like to thank Drs Klaus Attenkofer (now at NSLS-II, Brookhaven National Laboratory), Guy Jennings, and Mr Charles Kurtz of the Advanced Photon Source for their contributions to the Beamline 11ID-D facility at the APS. The instrumentation support (to LXC and others) from the US Department of Energy for purchasing lasers, detectors and other related equipment enabling the initiation and upgradation of XTA experiments at Beamline 11IDD is greatly appreciated. K. Haldrup gratefully acknowledges support from the Carlsberg and Villum Foundations.

Notes and references