Effects of Hydrogen Bonding on Internal Conversion of GFP-like Chromophores. II. The meta-Amino Systems

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Supporting Information

ABSTRACT: To rationalize the efficient quenching of the fluorescence and the Z → E photoisomerization of m-ABDI, the meta-amino analogue of the green fluorescent protein (GFP) chromophore, in protic solvents, the femtosecond time-resolved fluorescence and transient infrared (TRIR) spectra of m-ABDI in CD₃CN, CH₃OH, and CD₃OD are determined. For solutions in CD₃CN, the fluorescence decay lifetime is ~7.9 ns and IR absorption lines near 1513, 1531, 1557, and 1613 cm⁻¹ of m-ABDI in its electronically excited state were observed with a decay time >5 ns. For solutions in CH₃OH, the fluorescence decay is double exponential with time constants of ~16 and 62 ps. In addition to IR absorption lines of m-ABDI in its electronically excited state with a decay time of ~16 ps, new features near 1513, 1532, 1554, and 1592 cm⁻¹ were observed to have a rise time of ~19 ps and a decay constant of ~58 ps, indicating formation of an intermediate. The assignments for the IR spectra of the ground and excited states were assisted with DFT and TDDFT calculations, respectively. We conclude that the torsion of the exocyclic C=C bond (the τ torsion) is responsible for the nonradiative decay of electronically excited m-ABDI in CD₃CN. However, in CH₃OH and CD₃OD, the solute–solvent hydrogen bonding (SSH) interactions diminish significantly the barrier of the τ torsion and induce a new pathway that competes successfully with the τ torsion, consistent with the efficient fluorescence quenching and the diminished yield for Z → E photoisomerization. The new pathway is likely associated with excited-state proton transfer (ESPT) from the solvent to m-ABDI, particularly the carbonyl group, and generates an intermediate (ESPT*) that is weakly fluorescent.

INTRODUCTION

The phenomenon of enhanced photochemical reactivity for meta- vs para-substituted systems is often termed the “meta effect.”¹⁻³ Zimmerman devised this term to account for the solvolytic activity of a number of meta-disubstituted benzenes in their excited states enhanced relative to the corresponding para-disubstituted benzenes, which is of the reverse for the ground-state reactivity.¹ The meta effect has been demonstrated with other photochemical properties.⁴⁻⁵ One particular example is the meta-amino effect on the enhancement of fluorescence quantum efficiency (Φᵢ) of trans-stilbenes; in cyclohexane, Φᵢ (0.78) of trans-3-aminostilbene is 15 times that (0.05) of trans-4-aminostilbene.⁶ The meta-amino effect arises from an increased barrier height for the torsion of the C=C bond in the lowest singlet excited state so that the fluorescence decay becomes kinetically favored.

The great utility of the green fluorescent protein (GFP) as a biomarker⁷⁻¹⁰ has inspired much research on GFP-like chromophores both to understand the structure–property relationship and for applications as fluorescent probes¹¹ or light-emitting materials.¹²⁻¹³ It is well-documented that the parent GFP chromophore, p-hydroxybenzylidenedimethylimadazolinone (p-HBDI), is nearly nonemissive (Φᵢ < 10⁻³) in nonviscous solutions at ambient temperature.¹⁴⁻¹⁵ The ultrafast fluorescence quenching is associated with the torsion of the exocyclic C=C bond (the τ torsion).¹⁶⁻¹⁸ Effective suppression of this τ torsion is thus a prerequisite to creation of GFP-like chromophores of large Φᵢ. Whereas this can be readily achieved with covalent linking of the two aromatic rings that constrains the τ torsion,¹⁹⁻²⁰ most unconstrained GFP-like chromophores are weakly fluorescent with Φᵢ lower than 0.10.²¹⁻²⁵

One of our laboratories recently reported that the meta-amino effect on enhanced fluorescence applies also to the GFP chromophore. The meta-amino derivative m-ABDI (Chart 1) exhibits Φᵢ as large as 0.34 in hexane, which is more than 100 times that of the para isomer p-ABDI (Φᵢ < 10⁻³).¹⁶ In contrast, the fluorescence is nearly quenched and the quantum yield of Z → E isomerization (Φₓₓ) is greatly diminished in protic solvents, such as CH₃OH, indicating a significant solute–
solvent hydrogen bonding (SSHB) effect on the decay dynamics. To characterize the fluorescence-quenching SSHB effect, we have measured the femtosecond fluorescence decay and time-resolved infrared (TRIR) spectra of both p-ABDI and m-ABDI in selected solvents. In the preceding paper, the excited-state dynamics of p-ABDI were discussed;26 we report here the results for m-ABDI. We conclude that the nonradiative decay of m-ABDI in aprotic solvents is dominated by the τ torsion, whereas in protic solvents the SSHB opens new channels for internal conversion (IC). According to calculations of the time-dependent density functional theory (TDDFT) on the IR absorption of m-ABDI in the excited state, the IC channel might be coupled with a proton transfer from the solvent molecule to the carbonyl oxygen of m-ABDI in the excited state. The SSHB effect on the τ torsion is also discussed.

### Experimental Section
The synthesis and X-ray crystal structure of m-ABDI have been reported.16 The steady-state and femtosecond time-resolved instruments and experimental methods were the same as those described for p-ABDI in the preceding paper.26 The 1H NMR spectra of both the Z and E isomers of m-ABDI in DMSO-d6 and HPLC chromatograms of m-ABDI before and after irradiation are shown in Figure S1 (Supporting Information).

#### Picosecond Time-Resolved Measurements.
Picosecond temporally resolved spectra were obtained with a time-correlated single-photon counting (TCSPC) system (Fluotime 200, PicoQuant). The excitation source was provided by a 400 nm pulsed-diode laser (LDH-400, PicoQuant), which was controlled by a pulsed laser driver (PDL800-D, PicoQuant). The excitation laser was focused onto the sample holder by a lens. A lens collected fluorescence emitted from the sample at a right angle. An iris was used to attenuate the intensity of the detected signal, and the polarization of the detected fluorescence relative to the excitation laser pulse was set at 54° (magic angle) with a polarizer. A subtractive double monochromator (9030DS, Sciencetech) compensated the GVD of fluorescence and selected the detection wavelength. A multichannel plate photomultiplier (R3809U, Hamamatsu) was connected to a computer with a TCSPC-module (SPC-630, Becker and Hickl) card for data acquisition.

#### Data Analysis of TRIR Spectra.
For the quantitative analysis of TRIR data for m-ABDI in selected solvents, we performed the global and target analysis by using the GLOTARAN software package.27,28 The aim of the analysis is to obtain a model-based description of the time-resolved data in terms of a small number of precisely estimated rate constants and difference spectra so as to elucidate a photoinduced process in complex systems. The global analysis performs a simultaneous analysis using a model of sequential reactions. In this program, all TRIR spectra are collected in a matrix, which was globally fitted by applying a sequential kinetic scheme with increasing lifetimes. In this work, the time-resolved difference spectra were fitted with two kinetic models. A parallel model

\[
A \rightarrow B \rightarrow C
\]

of independently decaying components yielded decay associated difference spectrum (DADS), which represent the decay of spectra associated with varied species with distinct lifetimes upon photoexcitation. The sequential model

\[
A \rightarrow B \rightarrow C \rightarrow \text{target}
\]

and found that converged results could be obtained only when \(k_1\) or \(k_2\) is fixed, and the resultant \(k_1 + k_2\) is identical to the \(k_1\) in the sequential model

\[
A \rightarrow B \rightarrow C
\]

#### Calculations.
All the calculations were carried out using the Gaussian 09 package.27,28 The electronic ground- and excited-state geometry optimizations adopted the density functional theory (DFT) and time-dependent density functional theory (TDDFT) methods, respectively. Becke’s three-parameter hybrid exchange functional and the Lee–Yang–Parr correlation functional (B3LYP)30,31 were utilized in the calculation with a 6-31G** basis set. Vibrational frequencies were evaluated for the optimized geometries at the same level to ensure that the structure obtained was a minimum on the potential energy surface. The polarizable continuum model (PCM)32 was employed for the bulk solvent effects with the solvents CH3CN (dielectric constant, \(\varepsilon = 35.688\)) and CH3OH (dielectric constant, \(\varepsilon = 32.613\)). The geometric optimizations of the hydrogen-bonded m-ABDI–CH3OH complexes were considered here as one and two methanol molecules with m-ABDI in the ground and excited states. The IR intensities were determined from the gradients of the dipole moment. All the calculated vibrational frequencies are scaled by a factor of 0.96 to correct the anharmonicity effects.33–35 To predict the absorption maxima, vertical excitation energies were calculated at the ground-state geometry with the time-dependent DFT (TDDFT) method at the B3LYP/6-31G** level.

### Results and Discussion

#### Steady-State Electronic Spectra.
The UV–vis absorption and emission spectra of m-ABDI in hexane, THF, CH3CN, and CH3OH are shown in Figure 1. The absorption spectra depend only slightly on the solvent. For example, the absorption maximum (\(\lambda_{max}\)) of m-ABDI was bathochromically shifted by ~400 cm\(^{-1}\) when hexane was replaced by CH3OH, but the corresponding solvatochromic shift was ~1900 cm\(^{-1}\) for the para isomer p-ABDI.26 The small solvent effect in the para systems is attributed to the inherently weaker mesomeric interactions between the amino donor and the imidazolinedione acceptor. In contrast, the fluorescence maximum (\(\lambda_{fmax}\)) exhibits a large solvatochromic shift (5431 cm\(^{-1}\)) on proceeding from hexane to CH3OH, indicating that the lowest excited state (S1) is highly polarized. For comparison, the corresponding...
solvatofluorochromic shift for p-ABDI is 2466 cm⁻¹. Evidently, the excited state of m-ABDI possesses a larger extent of charge separation than that of p-ABDI, which is again due to weaker mesomeric interactions between the donor and acceptor groups after photoinduced charge transfer. It should also be noted that the spectrum in CH₂OH shows dual fluorescence. The short-wavelength band near 495 nm is similar to that in hexane and the intensity is much weaker than the long-wavelength band at 677 nm. According to the femtosecond fluorescence and IR transients (vide infra), the short- and long-wavelength emissions result from m-ABDI and a protonated intermediate, respectively, in the excited state.

The electronic structure of m-ABDI in the gas phase with the geometry optimized at the B3LYP/6-31G* level was investigated with TDDFT calculations at the TD-B3LYP/6-31G** level. The benzylideneimidazolinone moiety is planar in the optimized structure, consistent with the reported X-ray crystal structure for m-ABDI, in which the dihedral angle defined by the phenyl and imidazolinone rings is only 5.1°. Two distinct conformers, syn and anti, depend on the relative orientation of the NH₂ and C=O groups. The anti form is more stable than the syn form by ~0.5 kcal mol⁻¹ in the gas phase (Table S1, Supporting Information), but only the syn form was observed in the solid state according to the X-ray crystal structure. The calculated absorption wavelength (λ_max,cal) and oscillator strength (f) agree satisfactorily with the observed spectra (Table S2, Supporting Information). The lowest energy transition S₀ → S₁ near 395 nm is weak (f < 0.1), and the more intense transitions are at higher energies with f being 0.30 and 0.20, respectively, for the nearly degenerate S₀ → S₂ and S₁ → S₁ transitions near 342 nm. The HOMO → LUMO is the major configuration of the S₁ state. As shown in Figure 2, the probability of finding electrons in the amino group is negligible in the LUMO but significant in the HOMO. The S₃ state thus possesses a charge-transfer character. In contrast, the S₂ and S₃ states are more localized in the benzylideneimidazolinone moiety. The large solvatofluorochromic shift from hexane to CH₂CN is consistent with a charge-transfer S₁ state.

The prediction of a low-intensity transition to the charge-transfer S₁ state and a more intense one to the localized S₂ state for m-ABDI is reminiscent of trans-3-aminostilbene (m-AS) and its derivative trans-3-((N,N-dimethylamino)stilbene (m-DS). The S₁→S₃ energy gaps estimated from the absorption spectra in methycyclohexane are 3162 and 4317 cm⁻¹ for m-AS and m-DS, respectively. The increase in the S₁→S₃ gap on going from m-AS to m-DS reflects the enhanced charge-transfer character of the chromophores. Along with the increase in the S₁→S₃ gap is a decrease in the intensity of the S₀ → S₁ absorption band, indicating the occurrence of intensity borrowing from the higher excited states of more allowed transition. This phenomenon was also observed for m-ABDI and its dimethylamino derivative m-DMABDI. Whereas there is no distinct band but a small tailing that corresponds to the weak S₀ → S₁ transition in the absorption spectra of m-ABDI, a low-lying shoulder near 420 nm that is distinct from the intense band near 350 nm is present for m-DMABDI. It is anticipated that the S₁→S₂ gap could be further enlarged in the excited state because of the enhanced charge-transfer character of the S₁ state, and thus, the rate of the S₁ → S₀ emission is reduced. This argument is supported by the long fluorescence lifetimes for all these meta systems (vide infra).

Quantum Yields. The Φᵢ values of m-ABDI in hexane (0.34), THF (0.28), CH₂CN (0.16), and CH₂OH (<10⁻⁸) at 296 K have been reported. Compared with other unconstrained GFP-like chromophores, the Φᵢ of m-ABDI in the aprotic solvents is unusually high as a result of the meta-amino effect. The Φᵢ decreases with increasing solvent polarity from hexane to CH₂CN. However, the behavior is different in CH₂OH, in which the fluorescence is nearly quenched. The Φ_ZE values of m-ABDI in hexane (0.37), THF (0.35), CH₂CN (0.45), and CH₂OH (0.08) were also reported. We have previously proposed that the parameter Φᵢ + 2Φ_ZE or Φᵢ + 2Φ_EZ is a useful probe to study the decay channels of alkenes of E-Z-photoisomerization activity. This proposition stems from the one-bond-flip (OBF) mechanism of photoisomerization, in which IC occurs at the perpendicular state with an ~90° torsion of the C=C bond so that the probability of forming the E and Z isomers is ~50% each. This condition corresponds to a quantum yield 2Φ_ZE and 2Φ_EZ for the C=C torsion in the Z and E isomers, respectively. For m-ABDI, the moderate magnitude of Φᵢ and that Φᵢ + 2Φ_ZE ≈ 1.0 in aprotic solvents indicate that fluorescence and the τ torsion are responsible for the deactivation of the excited state. The increase in Φ_ZE at the expense of Φᵢ in more polar solvents reveals a solvent effect on the barrier of τ torsion, which is decreased in more polar solvents. In CH₂OH, the Φᵢ becomes negligible and 2Φ_ZE is much less than 1.0. This condition indicates the presence of a new nonradiative decay channel that is induced by the SSHB interactions; such an indication is supported by the time-resolved fluorescence and IR spectra (vide infra).

Steady-State IR Spectra. The IR spectra of m-ABDI in its electronic ground state in CD₃CN, CH₂OH, and CD₃OD in the spectral region 1500–1750 cm⁻¹ are shown in Figure 3; the intensity is normalized to absorption near 1650 cm⁻¹. The spectra can be deconvoluted into six major vibrational bands (Figure S2, Supporting Information), and the wavenumbers...
and relative amplitudes of these bands are summarized in Table 1, with their assignments according to DFT calculations at the B3LYP/6-31G* level of theory (Figure S3, Supporting Information). The calculated wavenumbers and relative intensities agree satisfactorily with the observed spectra. The intense band near 1650 cm\(^{-1}\) corresponds to the exocyclic C=C bond stretching coupled with the C=N stretching and the NH\(_2\) bending. The band near 1570 cm\(^{-1}\) is associated with the phenyl stretching coupled with the exocyclic C=C and C=N stretching modes. The bands at 1585 and 1600 cm\(^{-1}\) correlate with the phenyl stretching coupled with the C=N stretching and NH\(_2\) bending modes. The shoulder at 1630 cm\(^{-1}\) in CD\(_3\)CN is ascribed to the NH\(_2\) bending, but the corresponding band in CH\(_3\)OH and CD\(_3\)OD might be overlapped with the band at \(\sim\)1650 cm\(^{-1}\). Finally, the broad band at \(\sim\)1710 cm\(^{-1}\) is associated with the C=O stretching mode. This band splits into two bands near 1700 and 1725 cm\(^{-1}\) in both CH\(_3\)OH and CD\(_3\)OD as a consequence of SSHB interactions. A splitting of the C=O absorption band in protic solvents was also observed for \(\text{p-ABDI}^2\) and other organic dyes with carbonyl groups.\(^4\)\(^2\)\(^4\)\(^3\)

**Time-Resolved Fluorescence Profiles.** Fluorescence lifetimes (\(\tau_f\)) of \(\text{m-ABDI}\) in CD\(_3\)CN and CH\(_3\)OH at 296 K were measured with time-correlated single photon counting (TCSPC). The fluorescence decay profile of \(\text{m-ABDI}\) in CD\(_3\)CN is fitted well with a single exponential function,

<table>
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<th>mode</th>
<th>CD(_3)CN</th>
<th>CH(_3)OH</th>
<th>CD(_3)OD</th>
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<td>1568</td>
<td>1558</td>
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<tr>
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<td>(19)</td>
<td>(28)</td>
<td>(28)</td>
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<td>1582</td>
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<td>(34)</td>
<td>(28)</td>
<td>(33)</td>
<td>(33)</td>
<td>(11)</td>
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<td>1630</td>
<td>u.r.(^c)</td>
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<td>(100)</td>
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<tr>
<td>(22)</td>
<td>(22)</td>
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<td>(22)</td>
<td>(22)</td>
<td>C=C = C=C + C=N (S) + NH(_2) (B)</td>
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</table>

Relative intensities normalized to the absorption band near 1645 cm\(^{-1}\) are listed in parentheses.\(^4\)Vibrational wavenumbers calculated with B3LYP/6-31G* in the gas phase are scaled by 0.96.\(^5\)u.r. = unresolved bands.\(^4\)Additional vibrational band split from \(\nu_{68}\) due to SSHB.
corresponding to a lifetime of 7.9 ns with excitation at 400 nm and detection at 600 nm (Figure S8, Supporting Information). The fluorescence decay of m-ABDI in CH$_3$OH was, however, much faster and unattainable with this technique because of the instrument response time 150 ps of TCSPC. Such a solvent effect on $\tau_1$ is consistent with the relative $\Phi_f$ determined in the two solvents.

To resolve the excited-state dynamics of m-ABDI in CH$_3$OH, the temporal profiles were investigated further with femtosecond fluorescence up-conversion. Figure 5 shows the fluorescence transients at a series of emission wavelengths from 500 to 725 nm. The fluorescence transients exhibit a rapid decay feature in the short-wavelength region and a rising feature to a plateau in the long-wavelength region, which is characteristic of solvation dynamics for an organic solute in a polar solvent. A plot of fluorescence intensity at 725 nm as a function of time (Figure S9, Supporting Information) shows a rise that exhibits a maximum near 10 ps; at 725 nm, the solvation dynamics has little effect on the temporal profile. A satisfactory fit of the plot requires three rising components (0.1 ± 0.2, 0.8 ± 0.3, and 5.2 ± 1.0 ps) and one decay component (55 ± 2 ps).

The wavelength-dependent fluorescence transients of m-ABDI in CH$_3$OH were converted to a time-resolved frequency plot (Figure S10, Supporting Information) according to a standard procedure.$^{45-48}$ The emission maximum at 508 nm (19667 cm$^{-1}$) in CH$_3$OH at time zero is near that at 495 nm (20202 cm$^{-1}$) in hexane. The emission maximum shifted to 700 nm (14500 cm$^{-1}$) after ~12 ps. The solvation dynamics can be described by three decay components (0.15 ± 0.01, 1.26 ± 0.02, and 8.8 ± 0.1 ps), which are consistent with the results (multiple exponential decays with the average response time coefficient 5.0 ps) reported for Coumarin 153 in CH$_3$OH.$^{49}$ We constructed the time-resolved emission spectra (TRES) accordingly, and the results are shown in Figure S11 (Supporting Information). The TRES show the spectral shift toward low-energy region from 0.2 to 31.5 ps. After 30 ps, the spectra of the TRES show no significant change and the decays of the spectral intensities were mainly due to population loss in the excited state. We also estimated the total intensity by fitting the fluorescence spectra to an equation with the parameters width, height, and asymmetry factor.$^{50}$ The estimated total intensity involves a large uncertainty in the first 2 ps (Figure 6).

The fluorescence decays rapidly for 2–10 ps. Because of the large uncertainties in the initial stage, we fitted data after 10 ps to a biexponential decay with time constants of 16 and 62 ps. The biexponential decays were tentatively assigned to two emissive species with different excited-state lifetimes detailed in the last section. Because of the complexity of the fluorescence transient spectra involving solvation, it is difficult to deconvolute the TRES shown in Figure S11 (Supporting Information) into two components as two emissive species in the excited state, but the trend of spectral shift can be easily identified.

**Time-Resolved Infrared Spectra.** The two-dimensional contour plots of the time-resolved infrared (TRIR) difference spectra in the spectral ranges 1500–1740 cm$^{-1}$ (in CD$_3$CN and CH$_3$OH) and 1360–1740 cm$^{-1}$ (CD$_3$OD) and in the time domains 0–1300 ps (CD$_3$CN), 0–300 ps (CH$_3$OH), and 0–800 ps (CD$_3$OD) for m-ABDI excited at 400 nm are shown in Figure 7a; positive values indicate formation, and negative values indicate depletion. The inverted ground-state IR absorption spectra are also shown in green in Figure 7a for comparison. The spectral region was extended to 1360 cm$^{-1}$ for samples in CD$_3$OD because of less interference from the solvent. The difference spectra recorded at 0.5, 23, 103, 503, and 1403 ps in CD$_3$CN and at 0.5, 5.0, 30, 100, and 300 ps in CH$_3$OH and CD$_3$OD are shown in Figure 7b. In all three solvents, the ground-state absorption bands near 1650 and 1700 cm$^{-1}$ are diminished (negative signals) upon irradiation. In addition, several positive features of transient absorption (TA), which depend on the solvent, appear in the region 1630–1500 cm$^{-1}$. Figure 7c shows the temporal profiles of TA signals at selected wavenumbers.

We performed global analysis$^{28,51}$ on the time-resolved difference spectra observed upon photoexcitation of m-ABDI in each solvent. For m-ABDI in CD$_3$CN, the data can be fitted with a single exponential decay and the DADS is shown in Figure 8a, with the associated decay curve presented in Figure 8b; the ground-state IR absorption spectra (green) are presented for comparison. The negative features are due to ground-state depletion, and the positive features are due to absorption of a new species (or state) upon photoexcitation. The ground-state recovery and the decay of new features were
fitted satisfactorily with a single exponential decay constant of 5.0 ns; this time constant might be smaller than the actual value because we only probed the absorption with a maximal delay of 1.5 ns. The decay constant of $\tau > 5.0$ ns for the new features is consistent with the observed fluorescence lifetime of 7.9 ns. For m-ABDI in CH$_3$OH, the observed spectra were fitted satisfactorily only with a sequential model $\text{A} \xrightarrow{k_1} \text{B} \xrightarrow{k_2} \text{C}$ with $k_1 = 15.5 \pm 0.3$ ps and $k_2 = 57.7 \pm 0.3$ ps. The same sequential model was also applicable to m-ABDI in CD$_3$OD, and time constants of $k_1 = 16.0 \pm 0.1$ ps and $k_2 = 247 \pm 1$ ps were obtained. This is conceivable, as we observed a clear rise and decay for the feature near 1594 cm$^{-1}$ (Figure 7c) and an isosbestic-like point near 1598 cm$^{-1}$ in CH$_3$OH and near 1605 cm$^{-1}$ in CH$_3$OD.
Table 2. Experimental and Calculated Wavenumbers (cm\(^{-1}\)) and Relative Intensities of the Transient Excited-State Vibration Bands in the Region 1450–1750 cm\(^{-1}\) for m-ABDI and m-ABDI–OH\(^{+}\) in CD\(_3\)CN, CH\(_3\)OH, and CD\(_3\)OD

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<th>Mode</th>
<th>Species A</th>
<th>Species B</th>
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<td>1528 ± 1 (100)</td>
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\(\nu_{65}\) in CD\(_3\)OD (Figure 7b). The EADS obtained for m-ABDI in CH\(_3\)OH and CD\(_3\)OD are shown in Figure 8c and f, with associated evolution curves shown in Figure 8d and f, respectively. In both solvents, the positive features of the black curves (species A) exhibit a single exponential decay of time constant \(\sim 16\) ps. The positive features of the red curves (species B) are characteristic of an intermediate with a rise time constant of \(\sim 16\) ps in both CH\(_3\)OH and CD\(_3\)OD and a decay constant of \(\sim 58\) ps in CH\(_3\)OH and \(\sim 247\) ps in CD\(_3\)OD.

The observed positive features of the black curves (species A) in Figure 8a, c, and e were deconvoluted, and the data are shown in Table 2. The deconvoluted spectra of these features in the three solvents are shown in Figure S12 (Supporting Information). In CD\(_3\)CN, four bands at 1513 ± 13, 1531 ± 5, 1557 ± 1, and 1613 ± 1 cm\(^{-1}\) with relative intensities S1:100:74:27 were observed. The deconvoluted bands in CH\(_3\)OH are at 1512 ± 4, 1531 ± 3, 1560 ± 2, and 1608 ± 2 cm\(^{-1}\) with relative intensities 33:100:103:38. Similar bands and relative intensities (in parentheses) at 1515 ± 3 (37), 1530 ± 2 (100), 1557 ± 1 (100), and 1615 ± 1 (32) cm\(^{-1}\) were observed in CD\(_3\)OD; additional features at 1414 ± 2, 1434 ± 1, 1471 ± 2, and 1500 ± 9 were also observed because of less interference by the solvent in this spectral region.

To characterize the nature of the observed bands for species A, we performed TD-DFT optimizations on the first electronically excited state of m-ABDI in CH\(_3\)CN and CH\(_3\)OH at the B3LYP/6-31G* level with the incorporation of the influence of the solvent by applying a polarizable-continuum model (PCM). The calculated results (Table 2 and Figure 9a and b) agree satisfactorily with the observed new features in terms of wavenumbers and relative intensities. The observed features at 1513, 1531, 1557, and 1613 cm\(^{-1}\) correspond to 1507, 1512, 1557, and 1624 cm\(^{-1}\) predicted by TDDFT. We thus conclude that these new features are due to IR absorption of the S\(_1\) state of m-ABDI. Accordingly, the observed TA band at 1613 cm\(^{-1}\) in CD\(_3\)CN is assignable to the C=O stretching coupled with the NH\(_2\) bending mode (calculated at 1624 cm\(^{-1}\)) of the S\(_1\) state (Figures S13 and S14, Supporting Information). That the wavenumbers of the C=O stretching mode are smaller in the excited state than in the ground state (1710 cm\(^{-1}\)) is attributed to the decreased double-bond character of the carbonyl in S\(_1\).

We deconvoluted the absorption features of the intermediate observed in CH\(_3\)OH (red curves in Figure 8c) into four bands at 1513 ± 4, 1532 ± 2, 1554 ± 3, and 1592 ± 2 cm\(^{-1}\), with relative intensities S2:100:87:28. Similar bands at 1512 ± 4, 1528 ± 1, 1553 ± 1, and 1596 ± 1 cm\(^{-1}\) were observed for m-ABDI in CD\(_3\)OD; additional features at 1415 ± 3, 1433 ± 1, 1475 ± 2, and 1501 ± 6 cm\(^{-1}\) were also observed because of less interference by the solvent in this spectral region. In general, the spectrum of this intermediate species is similar to that of the S\(_1\) state of m-ABDI except some spectral shifts; the band near 1608 cm\(^{-1}\) disappeared, and the band near 1592 cm\(^{-1}\) is enhanced and becomes isolated. It should be noted that the intensity ratios of overlapped bands for the electronically excited state (black curve in Figure 8) and the intermediate (red curve in Figure 8) might not be as reliable as the isolated bands.

To characterize the nature of these TA bands of the intermediate (species B), we performed TD-DFT optimizations on the S\(_1\) state of m-ABDI with the SSHB modes shown in Figure 4 and that of protonated species m-ABDI–NH\(^{+}\) and m-ABDI–OH\(^{+}\) (Chart 2), which can be formed via excited-state proton transfer (ESPT). These optimized excited-state SSHB systems are shown in Figure S15 (Supporting Information).
and the vibration modes are depicted in Figures S16–S20 (Supporting Information). A comparison of the TDDFT-predicted and the observed absorption features of the intermediate are depicted in Figure 9c–g. As the hydrogen-bonded species shown in Figure 4 represent only some possible SSHB forms for m-ABDI in CH$_3$OH and each observed TA signal might result from more than one SSHB form, a definitive assignment of the carrier of these bands is difficult. Nevertheless, the calculations provide some indication of a possible origin of the observed TA signals of the intermediate. Regarding the new species, the two candidates m-ABDI−OH’ and m-ABDI−NH’ have predicted bands similar to observations, with a unique band near 1592 cm$^{-1}$. However, the formation of m-ABDI−NH’ is less likely than that of m-ABDI−OH’ because an intense band near 1648 cm$^{-1}$ that corresponds to the C=O stretching mode was predicted for the former but unobserved (Figure 9f). Recall that the C=O stretching in S$_1$ is at 1608 cm$^{-1}$ (Figure 9b) and this band disappears upon the formation of the intermediate. The unique feature near 1592 cm$^{-1}$ might correspond to the phenyl and the exocyclic C=C stretching modes of m-ABDI−OH’ (Table 2). The excited state m-ABDI−OH’ is likely formed via ESPT from CH$_3$OH to m-ABDI.

**Excited-State Relaxation Mechanism.** The observation that $\Phi_f + 2\Phi_{\text{re}} \approx 1.0$ for m-ABDI in aprotic solvents$^{16}$ conforms to the OBF mechanism for $E/Z$-photoisomerization of alkenes, in which the IC occurs from a perpendicular $^1p^*$ state and the probability of partition to the $E$ and $Z$ isomers is equal. The fluorescence decay time $\sim 7.9$ ns for m-ABDI in CD$_3$CN is about 10,000 times that for $p$-ABDI (0.3–1.0 ps), which indicates that the $\tau$ torsion in the excited state encounters a significant barrier. This phenomenon is similar to that observed for meta- and para-aminostilbenes, which was dubbed as the meta-amino effect.$^{4–6}$ In analogy to the cases of trans-aminostilbenes,6,38–40,52,53 a large $\tau$-torsion barrier for m-ABDI might result from a lower energy for the $Z^*$ vs $p^*$ state. Compared to the $Z^*$ and $p^*$ states of $p$-ABDI, the energy of the $Z^*$ state should be lowered and that of the $p^*$ state should be raised in the case of m-ABDI because of a stronger charge-transfer character for the $Z^*$ and less mesomeric stabilization of the $p^*$ state by the amino group. As estimated with the fluorescence maxima (578 vs 469 nm) in CH$_3$CN, the $Z^*$ state of m-ABDI is lower by 11.5 kcal mol$^{-1}$ than that of $p$-ABDI. The dark $^1p^*$ state does not provide an emission signal for direct evaluation of the energy difference of the $^1p^*$ state in m-ABDI vs $p$-ABDI. It was reported to be $\sim 7$ kcal mol$^{-1}$ higher for the $^1p^*$ state of trans-3-aminostilbene than that of trans-4-aminostilbene.$^6$ A similar size might be expected for the case of m-ABDI vs $p$-ABDI. Although the relative energies of the $Z^*$ and $p^*$ states of $p$-ABDI are unknown and the $Z^* \rightarrow p^*$ process should be exergonic, the simultaneous stabilization of $Z^*$ and destabilization of $p^*$ on going from $p$-ABDI to m-ABDI with a total energy change near 19 kcal mol$^{-1}$ should render the $Z^* \rightarrow p^*$ process of m-ABDI endergonic with a significant barrier. In addition to a slower rate for the $\tau$ torsion, another contribution to the longer fluorescence lifetime of m-ABDI vs $p$-ABDI is the less allowed S$_1 \rightarrow$ S$_0$ transition (vide supra). On the basis of $\Phi_f$ (0.16) and $\tau_f$ (7.9 ns), the time constants of the radiative process ($\tau_f/\Phi_f$) and of the nonradiative relaxation through the $\tau$ torsion ($\tau_f/(1-\Phi_f)$) of m-ABDI in CH$_3$CN are $\sim 50$ and $\sim 10$ ns, respectively (Figure 10a).

The rapid fluorescence quenching ($\sim 16$ and 62 ps) and the small $\Phi_{\text{re}}$ (0.08) for m-ABDI in CH$_3$OH indicate that the nonradiative decay in CH$_3$OH differs from that in CD$_3$CN and is strongly affected by the SSHB interactions. A simplified scheme for the decay dynamics is shown in Figure 10b, where the HB in the labels denotes the presence of hydrogen bonding with the solvent molecules. The presence of an intermediate excited state (ESPT*) that is formed via ESPT is proposed on the basis of the observation of a rise time of $\sim 16$ ps and a decay time of $\sim 58$ ps for the species B in the global analysis of TRIR spectra. An alternative explanation of the intermediate excited state is a HB-associated TICT state.$^{9,54,55}$ However, the dynamics of a HB-associated TICT state are generally more
ESPT process. It should also be noted that the driving force for the excited-state dynamics of m-ABDI differs significantly from that of the para isomer p-ABDI, for which ESPT does not occur and for which only the PES of the τ torsion is perturbed. According to the scenario of Figure 10b, the absence of ESPT for p-ABDI could be attributed to the ultrafast decay (~1 ps) of the excited state via the τ torsion so that the ESPT process cannot compete with the τ torsion. The meta-amine effect in m-ABDI raises the torsion barrier and slows the τ torsion. The time constant 100 ps for the τ torsion of m-ABDI in CH$_3$OH is much larger than that (19 ps) for the ESPT process. It should also be noted that the driving force for the ESPT process is inherently larger in m-ABDI vs p-ABDI, as the former possesses a stronger charge-transfer character and thus a better hydrogen-bond acceptor of the carbonyl oxygen. The rapid fluorescence intensity decay for the initial 10 ps (Figure 6) deserves a comment. Both HB-Z$^*$ and ESPT$^*$ are fluorescent, and their decay coefficients ~16 and ~62 ps match well with the decay constants ~16 and 58 ps derived from global fitting of the TRIR spectra. The rapid fluorescence decay in the first 10 ps has no corresponding component in the TRIR spectra, indicating that this decay does not involve HB-Z$^*$ or ESPT$^*$. The rapid decay occurred during the period when solvent-induced vibrational relaxation and solvation were taking place so that interpretation of this relaxation process becomes complex. Nevertheless, it is possible that the molecular conformation and electronic character of m-ABDI in the excited state was significantly affected by SSHB in a way that the optical transition probability is much lower for the relaxed electronic excited state (HB-Z$^*$) than for the Franck–Condon state (FC) to account for the ~60% decrease in total fluorescence intensities in the first 10 ps. Another possibility is that the FC state with higher energy might overcome the τ torsion barrier more readily than the HB-Z$^*$ state. However, as is the situation of the SSHB effect on the PES along the τ torsion coordinate for p-ABDI, the τ torsion is aborted in an early stage (i.e., at small τ torsion angles), so that the observed Z → E isomerization quantum yield (0.08) is much lower than a total quantum yield of ~0.60 for the τ torsion. In this case, the rate for HB-Z$^*$ → ESPT$^*$ might be just ~16 ps and the rate of HB-Z$^*$ → HB-1p$^*$ could be much greater than 100 ps, with the

Figure 10. Schematic representation of the proposed excited-state relaxation dynamics for m-ABDI in (a) CH$_3$CN and (b) CH$_3$OH. The HB and PT in the labels denote hydrogen-bonded and proton transfer, respectively.
total decay of HB-Z* ~16 ps. Further experiments are needed to confirm such a possibility.

Finally, it deserves a comparison of the meta-NH2 and the meta—OH effect58,59 on the excited-state dynamics of GFP chromophores. The meta-OH isomer m-HBDI was shown to display fluorescence decay times up to 30 times longer than p-HBDI does in CH3CN. Such a meta-OH effect is much less significant as compared to the meta-NH2 effect, which induces a 10^4 times increase in the fluorescence lifetime. The smaller meta-OH vs meta-NH2 effect can be attributed to a smaller increase in the barrier for the τ torsion. We reasoned that the hydroxyl group is a weaker donor than the amino group so that the stabilization of the Z* state and the destabilization of the p* state on going from p-HBDI to m-HBDI are to a much weaker extent than the meta-NH2 effect. Similar behavior was found for the comparison of trans-3-hydroxy stilbenes60-62 and trans-3-aminostilbenes.63,64 For anionic m-HBDI, the stronger oxid (O−) donor does lead to a feature of band splitting in the electronic absorption spectrum similar to the case of the meta-NH2 effect.59 This phenomenon would in principle significantly raise the τ-torsion barrier as is the meta-NH2 effect. However, the solvent condition of CH3OH/H2O (1/9) and pH 11.5 required for the formation of anionic m-HBDI in the ground state also introduces SSSH interactions and ultrafast ESPT processes, which counterbalances the substituent effect.

**CONCLUSION**

The excited decay dynamics for m-ABDI in CD3CN, CH3OH, and CD3OD have been investigated with femtosecond fluorescence and transient IR absorption spectroscopy. In CD3CN, the decays are mainly through the fluorescence and the exocyclic C=C bond (τ) torsion. Because of a slow rate (10 ns) of the τ torsion, the observed Φτ (0.16) is exceptionally large among the unconstrained GFP chromophores known in nonviscous solvents. In CH3OH, the participation of SSSH strongly affects the decay mechanism, as evidenced by the small Φτ (0.08) and Φτ (10^-3). There exists a fluorescent intermediate state (ESPT*), which was not observed in the case of the para isomer p-ABDI. The ESPT* is likely due to ESPT from CH3OH to the carbonyl oxygen of m-ABDI. The SSSH interactions might also couple with the τ torsion to decrease the barrier and/or to initiate the ESPT. That the SSSH effects on the decay dynamics are larger for m-ABDI than for p-ABDI is attributed to a stronger charge-transfer character and a larger barrier for the τ torsion of the S state. These results indicate the importance of not only SSSH but also the location of substituents on the excited decay dynamics of a GFP-like chromophore. These results provide useful information for structural engineering toward highly fluorescent GFP chromophores.

**ASSOCIATED CONTENT**

1. Supporting Information

Deconvoluted IR spectrum of m-ABDI in selected solvents, fluorescence decay profiles, time-dependent frequencies, fitting of decay profiles, TDDFT-optimized geometries of hydrogen-bonded complexes, TDDFT-calculated infrared absorption, EADS from target analysis and TRES of m-ABDI in CH3OH, table of optimized structures and relative energies of m-ABDI and associated complexes, tables of calculated vibrational modes, complete ref 29, and Cartesian coordinates of m-ABDI and its hydrogen-bonded complexes in their electronically excited and ground states. This material is available free of charge via the Internet at http://pubs.acs.org.

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**Notes**

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