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Effects of Methyl Substitution in Ruthenium Tris(2-pyridylmethyl)amine Photocaging Groups for Nitriles

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Abstract

Four complexes of the general formula $[\text{Ru}(\text{L})(\text{CH}_3\text{CN})_2](\text{PF}_6)_2$, $[\text{L} = \text{TPA}$ (**5**), MeTPA (**6**), Me_2TPA (**7**), and Me_3TPA (**8**)] [$\text{TPA} = \text{tris}[(\text{pyridin-2-yl})\text{methyl}]\text{amine}$, where methyl groups were introduced consecutively onto the 6-position of py donors of TPA, were prepared and characterized by various spectroscopic techniques and mass spectrometry. While **5** and **8** were isolated as single stereoisomers, **6** and **7** were isolated as mixtures of stereoisomers in 2:1 and 1.5:1 ratios, respectively. Steric effects on ground state stability and thermal and photochemical reactivities were studied for all four complexes using ^1H NMR and electronic absorption spectroscopies and computational studies. These studies confirmed that the addition of steric bulk accelerates photochemical and thermal nitrile release.

Graphical abstract

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Supporting Information

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Crystallographic information file for **7a** (CIF)

General considerations, DFT results and photochemical studies, X-ray crystallographic analysis, spectral data for **6**, **7**, and **8**, cartesian coordinates, frequencies (in cm^{-1}) of the calculated structures, electronic structure information for the ^3MC states of **5** and **6**, and TD-DFT for the two isomers of $[\text{Ru}(\text{TPA})(\text{CH}_3\text{CN})(\text{Cl})]^+$ (PDF)

Notes

The authors declare no competing financial interest.

INTRODUCTION

Compounds that undergo photochemical cleavage reactions have important applications in materials science,¹ chemistry, and biology.² The binding of compounds to photolabile protecting groups, also known as photocaging, gives researchers the ability to achieve spatial and temporal control over release of an active agent using light. For decades, chemists have employed organic protecting groups as photocages.^{3,4} More recently, metal complexes have become an important class of photocaging groups.⁵ Metal complexes hold several advantages over their organic counterparts. An important aspect is that their photochemistry can be tuned over a broad range of the visible spectrum by manipulating the ancillary ligands.^{6,7} Metal complexes also bind to functional groups that cannot be caged with organic fragments, including nitrogen-containing heterocycles,^{8–15} thioethers^{16–18} and nitriles.^{19–28} Thus, metal complexes offer an orthogonal approach to organic caging methods.

Nitriles are a robust pharmacophore found in many biologically active compounds, including over 30 drugs currently used in the clinic.²⁹ Despite their prevalence in biological tools and drugs, nitriles are a functional group that to date cannot be protected with an organic fragment. Thus, metal complexes are the only option for photocaging nitriles, and represent an attractive target for further development. Seminal studies established that the caging group Ru(bpy)₂ (bpy = 2,2'-bipyridine) can be used to cage 5-cyanouracil (5CNU), a cytotoxic agent that inhibits pyrimidine catabolism in vivo.²⁰ Later work showed that [Ru(tpy)(5CNU)₃]²⁺ (tpy = 2,2':6',2''-terpyridine) releases the same agent in cervical cancer cells when irradiated with visible light.²³ In addition, the Ru(bpy)₂ photocaging group was applied to a series of nitrile-based protease inhibitors, initiating enzyme inhibition against purified cysteine cathepsins only upon photoactivation, as well as cathepsin activities in lysates and live cells.^{21,24,30,31}

Pioneering work in neuroscience demonstrated that ruthenium complexes can be used to cage neurotransmitters without causing toxicity.^{8,32–34} Since then, most efforts in developing ruthenium-based photocaging groups focused on planar, heteroaromatic ligands similar to bpy, where ancillary ligands are typically bi- or tridentate possess denticities of three or below. We recently reported that a ruthenium fragment based on the tetradentate ligand tris[(pyridin-2-yl)methyl]-amine (TPA, Figure 1) is an effective photocaging group for nitriles.²⁵ Even though the Ru(TPA) motif had been investigated in photochemical molecular machines and switches,^{35–38} oxidation^{39–49} and hydrogenation⁵⁰ catalysts, DNA metallointercalators,⁵¹ and for proton-coupled electron transfer properties,^{52–56} its behavior as a photocaging group had only been investigated for release of nitric oxide.⁵⁷ Gratifyingly, Ru(TPA) showed promising activity as a caging group for nitriles, including stability in buffer and high selectivity for enzyme inhibition under dark versus light conditions. We also disclosed a solid-phase method that can be used to synthesize and screen derivatives of TPA as ligands for ruthenium caging groups to rapidly assess effects of the ancillary ligand on tuning spectral properties and photoreactivity for nitrile release.²⁷

Beyond tuning the identity of the donor atom or increasing conjugation, steric effects are known to control photochemical reactivity in ruthenium complexes. The introduction of steric bulk is used to distort the octahedral field, lower the bond dissociation energy and

bring dissociative triplet metal-centered (^3MC) states closer in energy to triplet metal-to-ligand charge transfer ($^3\text{MLCT}$) states, which are generated by photoexcitation.¹⁶ Thus, more efficient photodissociation can occur in distorted complexes. For example, sterically encumbered analogs of bpy, such as 6,6'-dimethyl-2,2'-bipyridine, undergo efficient release from ruthenium complexes upon irradiation with visible light, which opens coordination site for DNA binding.⁵⁸ More recently, steric effects have also been used to increase the quantum yields of monodentate pyridine release from ruthenium caging groups.^{12,59} Given the success already demonstrated in these reports, we hypothesized that introducing steric bulk could prove favorable in controlling the photoreactivity of Ru(TPA) complexes. However, steric effects on photodissociation have not been investigated systematically in these complexes, including studies geared toward controlling efficiency of ligand release.

In this Article, we report a systematic study to determine the role of steric effects in Ru(TPA) and related complexes used to cage the model nitrile CH_3CN . Four complexes were synthesized and characterized structurally and photochemically, where methyl groups were introduced consecutively onto the 6-position of the pyridine donors. In addition, calculations were used to gain insight into ground state stabilities as well as the thermal and photochemical reactivities. These studies confirm that steric effects can facilitate nitrile release, both from the ground state and from excited states accessed by photoexcitation.

EXPERIMENTAL SECTION

General Considerations

NMR spectra were recorded on a Varian FT-NMR Mercury-400 MHz spectrometer. Mass spectra were recorded on a Time-of-Flight Micromass LCT Premier XE Spectrometer. IR spectra were recorded on a Nicolet FT-IR spectrophotometer (KBr pellet). UV-vis spectra were recorded on a Varian Cary 50 spectrophotometer. All reactions were performed under ambient atmosphere unless otherwise noted. Anaerobic reactions were performed by purging the reaction solutions with Ar or N_2 . Ligands **1–4**⁶⁰ and complex $[\text{Ru}(\text{TPA})(\text{CH}_3\text{CN})_2](\text{PF}_6)_2$ (**5**)²⁵ were synthesized according to literature procedures.

Synthesis of Ru Complexes

[Ru(MeTPA)(CH₃CN)₂](PF₆)₂ (6**)**—A pressure flask was charged with *cis*- $[\text{RuCl}_2(\text{Me}_2\text{SO})_4]$ (30 mg, 0.062 mmol), MeTPA ligand (19 mg, 0.062 mmol) and MeOH (6 mL). The solution was deoxygenated by bubbling Ar through a submerged needle for 15 min, sealed and refluxed at 70 °C for 4 h under inert atmosphere during which time the reaction color changed from pale yellow to dark yellow. The reaction mixture was cooled to RT and concentrated in vacuo to give a dark yellow solid. The crude reaction mixture containing $[\text{Ru}(\text{MeTPA})(\text{DMSO})\text{Cl}]\text{Cl}$ (34 mg, 0.062 mmol), as 2:1 mixture of stereoisomers, was dissolved in a 1:1 mixture of H_2O and CH_3CN (6 mL) under argon atmosphere and the resulting solution was refluxed at 80 °C for 16 h under inert atmosphere. Ice cold water (5 mL) was added to the reaction mixture followed by a saturated solution of aqueous NH_4PF_6 (2 mL), resulting in a pale yellow precipitate that was isolated by centrifugation, washed with ice-cold H_2O and dried under reduced pressure to get the title complex as a yellow solid (45 mg, 0.058 mmol) obtained as a 2:1 mixture of two isomers in

94% overall yield. ^1H NMR (400 MHz $\text{C}_3\text{D}_6\text{O}$): δ 9.37 (d, 2H, $J = 5.7$ Hz), 9.07 (d, 2H, $J = 5.7$ Hz), 8.90 (d, 2H, $J = 5.7$ Hz), 7.95–7.86 (m, 5H), 7.82 (t, 2H, $J = 7.7$ Hz), 7.75–7.65 (m, 4H), 7.62 (d, 2H, $J = 5.7$ Hz), 7.55 (t, 1H, $J = 7.7$ Hz), 7.51–7.43 (m, 4H), 7.42–7.35 (m, 6H), 7.26 (d, 2H, $J = 7.7$ Hz), 7.04 (d, 1H, $J = 7.7$ Hz), 5.44–5.36 (m, 6H), 5.26–5.18 (m, 6H), 5.00–4.83 (m, 6H), 3.13 (s, 3H), 2.97 (s, 6H), 2.88 (s, 6H), 2.86 (s, 3H), 2.55 (s, 3H), 2.54 (s, 6H). IR (KBr) ν_{max} (cm^{-1}): 3854, 3839, 3801, 3750, 3675, 3649, 3421, 2934, 2853, 2279, 1734, 1707, 1610, 1575, 1508, 1460, 1386, 1312, 1226, 1167, 1039, 995, 839, 788, 772, 739. ESMS calcd for $\text{C}_{23}\text{H}_{26}\text{N}_6\text{Ru}$ (M^{+2}): 244, found 244. UV–vis (acetonitrile): $\lambda_{\text{max}} = 370$ ($\epsilon = 10\,000\text{ M}^{-1}\text{cm}^{-1}$). Anal. Calcd for $\text{C}_{23}\text{H}_{26}\text{F}_{12}\text{N}_6\text{P}_2\text{Ru}$ ($6 \cdot 2.5\text{ H}_2\text{O}$): C, 33.59; H, 3.80; N, 10.22. Found: C, 33.35; H, 3.41; N, 10.49.

[Ru(Me₂TPA)(CH₃CN)₂](PF₆)₂ (7)—Complex **7** was synthesized using the same method as with **6**, starting from Me₂TPA (33 mg, 0.103 mmol). After 2 steps, **7** was obtained as a 1.5:1 mixture of two isomers in 89% overall yield (73 mg, 0.092 mmol). ^1H NMR (400 MHz $\text{C}_3\text{D}_6\text{O}$): δ 9.61 (d, 1.5H, $J = 5.6$ Hz), 9.03 (d, 1H, $J = 5.6$ Hz), 7.89–7.84 (m, 2H), 7.79–7.71 (m, 5H), 7.62–7.55 (m, 2H), 7.51 (d, 1H, $J = 7.6$ Hz), 7.46 (d, 3H, $J = 7.6$ Hz), 7.43–7.39 (m, 2.5H), 7.32–7.29 (m, 5H), 7.26 (d, 1H, $J = 7.6$ Hz), 7.10 (d, 1H, $J = 7.6$ Hz), 5.49–5.39 (m, 5H), 5.22–5.17 (m, 5H), 4.94 (d, 1H, $J = 17.9$ Hz), 4.90 (s, 3H), 4.76 (d, 1H, $J = 17.9$ Hz), 3.18 (s, 3H), 2.97 (s, 3H), 2.92 (s, 7.5H), 2.90 (s, 9H), 2.59 (s, 4.5H), 2.55 (s, 3H). IR (KBr) ν_{max} (cm^{-1}): 3854, 3839, 3737, 3676, 3402, 2940, 2281, 1992, 1735, 1716, 1610, 1574, 1508, 1461, 1438, 1386, 1354, 1271, 1233, 1171, 1096, 1037, 998, 970, 839, 789, 770, 739, 644, 611. ESMS calcd for $\text{C}_{24}\text{H}_{28}\text{N}_6\text{Ru}$ (M^{+2}): 251, found 251. UV–vis: λ_{max} (acetonitrile) = 365 ($\epsilon = 9700\text{ M}^{-1}\text{cm}^{-1}$). Anal. Calcd for $\text{C}_{24}\text{H}_{28}\text{F}_{12}\text{N}_6\text{P}_2\text{Ru}$: C, 36.42; H, 3.57; N, 10.62. Found: C, 36.46; H, 3.58; N, 10.38.

[Ru(Me₃TPA)(CH₃CN)₂](PF₆)₂ (8)—A pressure flask was charged with *cis*-[RuCl₂(Me₂SO)₄] (30 mg, 0.062 mmol) and ligand **4** (21 mg, 0.062 mmol) in dry EtOH (5 mL) was deoxygenated by bubbling Ar through a submerged needle for 10 min in a sealable tube. The reaction mixture was sealed and refluxed for 18 h at 90 °C under Ar atmosphere. The reaction mixture was cooled to RT and concentrated in vacuo, giving a mixture of [Ru(Me₃TPA)Cl(Me₂SO)]Cl complexes, as judged by ^1H NMR and UV–vis spectroscopic analysis of the crude reaction mixture. The crude mixture was dissolved in a 1:1 mixture of H₂O and CH₃CN (5 mL) and the resulting solution was heated under argon atmosphere to reflux at 80 °C for 18 h. After the mixture was cooled to RT, Ar was flowed over the mixture for 1 h to expel acetonitrile. The reaction mixture was transferred to a solution of NH₄PF₆ at 0 °C (15 mL), resulting in formation of a pale yellow precipitate that was isolated by centrifugation. The pellet was washed several times with cold water. After drying in vacuo, complex **8** was isolated as a yellow solid (42 mg, 84%). Crystals suitable for X-ray crystallographic analysis were obtained by diffusing Et₂O into a solution of **8** in MeCN: mp = 220 °C (decomp). ^1H NMR (400 MHz $\text{C}_3\text{D}_6\text{O}$): δ 7.79 (t, 2H, $J = 7.7$ Hz), 7.58 (t, 1H, $J = 7.7$ Hz), 7.45 (d, 2H, $J = 7.7$ Hz), 7.35 (d, 2H, $J = 7.7$ Hz), 7.29 (d, 1H, $J = 7.7$ Hz), 7.11 (d, 1H, $J = 7.7$ Hz), 5.49 (part of an AB system, 2H, $J_{\text{AB}} = 15.6$ Hz), 5.17 (part of an AB system, 2H, $J_{\text{AB}} = 15.6$ Hz), 4.82 (s, 2H), 3.35 (s, 3H), 2.97 (s, 6H), 2.94 (s, 3H), 2.60 (s, 3H). IR (KBr) ν_{max} (cm^{-1}): 3903, 3854, 3839, 3821, 3802, 3751, 3735, 3677, 3567, 3261, 3005, 2941, 2361, 1772, 1699, 1611, 1575, 1541, 1522, 1508, 1473, 1458, 1396, 1355,

1238, 1222, 1174, 1098, 1034, 992, 969, 838, 790, 739, 644, 612. ESMS calcd for $C_{25}H_{30}N_6Ru$ (M^{+2}): 258, found 258. UV-vis: λ_{max} (acetonitrile) = 360 ($\epsilon = 5600 M^{-1} cm^{-1}$). Anal. Calcd for $C_{25}H_{30}F_{12}N_6P_2Ru$: C, 37.28; H, 3.75; N, 10.43. Found: C, 37.51; H, 3.84; N, 10.38.

Photochemistry

Photolysis and ligand exchange quantum yield experiments were performed with a 150 W Xe arc lamp (USHIO) contained in a MilliArc lamp housing unit and powered by an LPS-220 power supply equipped with an LPS-221 igniter (PTI). A 395 nm long pass filter (CVI Melles Griot) was used for the photolysis experiments, and a 400 nm bandpass filter (Thorlabs) with a 335 nm long-pass filter were used together for the ligand exchange quantum yield experiments. The samples were dissolved in either CH_2Cl_2 with 10 mM Bu_4NCl or H_2O with <5% acetone. Potassium tris(ferrioxalate) was used as the chemical actinometer to determine the photon flux of the lamp with $\lambda_{irr} = 400$ nm (2.1×10^{-8} mol photons/min).⁶¹ For NMR studies, solutions of **5** and **6** in CD_3CN (3 mM) were dried with 4 Å molecular sieves. After filtration, solutions were irradiated in NMR tubes with a long wave UV TLC lamp (8W, 365 nm) at a distance of 13 cm from the lamp to the tubes, and 1H NMR spectra were recorded at 1 h intervals from $t = 0$ –5 h. To analyze the photoproducts of **6** by NMR, the complex was dissolved in CD_3OD to ~3 mM and irradiated (150 W Xe arc lamp, $\lambda_{irr} = 395$ nm selected using a long-pass filter). The 1H NMR spectra were collected using a Bruker 400 MHz DPX spectrophotometer. The resulting photoproduct was isolated by drying the sample in the NMR tube with air to remove the CD_3OD solvent, then CD_3CN was added to the dried sample in the NMR tube and the solution was kept in the dark by wrapping the tube in aluminum foil, and the NMR spectrum was collected after 16 h. To analyze the chloride-bound samples, **6** was irradiated in CD_3OD as described above, then ~10 mg of $CaCl_2$ was added to the solution. The sample was wrapped in aluminum foil and kept in the dark overnight to allow the exchange of Ru-bound CD_3OD for Cl^- , and the NMR spectrum was collected after 16 h.

Computational Details

Electronic structure calculations were carried out with the Gaussian09 software package⁶² using the BP86 density functional.^{63,64} The SDD basis set and effective core potential^{65,66} were used for Ru atom and the 6-31G(d) basis set^{67,68} was used for the other atoms. Solvation effects in acetonitrile were incorporated by using the implicit SMD solvation model⁶⁹ and were included during structure optimization. All of the optimized structures were confirmed as minima by harmonic vibrational frequency calculations and the converged wave functions were tested for the SCF stability. The triplet metal-centered states (3MC) were fully optimized using the SCF method. The 3MC states have been modeled as five-coordinate Ru complexes. The identities of the 3MC electronic configurations were confirmed by S^2 expectation values $\langle S^2 \rangle$ and by plots of the spin density (isovalue = 0.004 au, visualized using GaussView⁷⁰). TD-DFT calculations^{71,72} were performed in dichloromethane solvent with the same density functional and basis sets using ground state geometries to characterize the photoproduct from complex **5**. The electronic transitions were checked by visualizing the canonical orbitals (isovalue = 0.05 au) using GaussView. To explore the potential energy surfaces for CH_3CN dissociation from Ru complexes, relaxed

potential energy surface scans were performed by stretching the Ru–NCCH₃ bond and optimizing the remaining coordinates.

RESULTS AND DISCUSSION

Synthesis

Complexes for our study were synthesized starting from the known tetradentate ligands tris[(pyridin-2-yl)methyl]amine (TPA, **1**), [(6-methylpyridin-2-yl)methyl]bis-[(pyridin-2-yl)methyl]amine (MeTPA, **2**), bis[(6-methylpyridin-2-yl)methyl][(pyridin-2-yl)methyl]amine (Me₂TPA, **3**), and tris[(6-methylpyridin-2-yl)methyl]amine (Me₃TPA, **4**).⁶⁰ Treating ligands **1–3** with 1 equiv of *cis*-[RuCl₂(Me₂SO)₄] in MeOH at 70 °C for 4 h, followed by concentration resulted in the metalation and afforded mixtures of [Ru(L)Cl(Me₂SO)]Cl complexes, where L = **1–3**.⁷³ Heating each of these mixtures of complexes in 1:1 CH₃CN/H₂O at 80 °C for 16 h, followed by precipitation with NH₄PF₆ gave [Ru(TPA)(CH₃CN)₂](PF₆)₂ (**5**), [Ru(MeTPA)(CH₃CN)₂](PF₆)₂ (**6**), and [Ru(Me₂TPA)-(CH₃CN)₂](PF₆)₂ (**7**) as yellow solids, in yields ranging from 48% to 94%. Complex **5** was isolated as a single isomer, whereas complexes **6** and **7** were obtained as 2:1 and 1.5:1 mixtures of stereoisomers, respectively. Structural assignments for major and minor isomers of **6** and **7** are provided below. Applying the same two-step procedure to the ligand Me₃TPA resulted in <20% conversion, as judged by ¹H NMR spectroscopic analysis, with the remainder of the material being unreacted ligand **4**. Further inspection of each step confirmed that metalation of **4** with *cis*-[RuCl₂(Me₂SO)₄] was significantly slower than with ligands **1–3**, and thus would require more forcing conditions. Heating ligand **4** at 90 °C with 1 equiv of *cis*-[RuCl₂(Me₂SO)₄] in EtOH for 18 h led to higher conversion to [Ru(Me₃TPA)Cl(Me₂SO)]Cl, which proceeded readily to form [Ru(Me₃TPA)(CH₃CN)₂](PF₆)₂ (**8**) upon heating in 1:1 CH₃CN/H₂O at 80 °C for 18 h, followed by precipitation with NH₄PF₆. Using these optimized conditions, complex **8** was isolated in 84% overall yield in 2 steps from **4**.

X-ray Crystallographic Studies

Complexes **5** and **8** were characterized by X-ray crystallography. Diffusion of Et₂O into a solution of **5** or **8** in CH₃CN furnished small yellow blocks suitable for X-ray crystallographic analysis. Parameters of data collections are described in Table 1. Tables 2 and 3 describe selected bond distances and bond angles, respectively.

Complex **5** crystallized in the triclinic space group $P\bar{1}$ with $Z = 4$.⁷⁴ Two nearly identical dications of [Ru(TPA)(CH₃CN)₂]²⁺ are located in the asymmetric unit. Because of their similarity, structural parameters for only one of the two dications are described herein; parameters for the second dication can be found in the Supporting Information. The coordination environment around the ruthenium center in the dication [Ru(TPA)(CH₃CN)₂]²⁺ (Figure 2A) possesses a distorted octahedral geometry, with bond angles ranging from 81.7(2)° to 99.5(2)°. The two bound CH₃CN ligands are in different chemical environments, with one CH₃CN group *cis* and one *trans* to the basic nitrogen donor N3. Despite this fact, the Ru1–N1 and Ru1–N6 bond distances to nitrile are identical within error at 2.031(5) and 2.037(5) Å, respectively. Ruthenium–nitrogen bond distances to the three

pyridine donors and one basic nitrogen (N2–N5) range only slightly from 2.053(4) to 2.071(4) Å.

Complex **8** crystallized in the monoclinic space group $C12/c1$ with $Z = 8$. Data for the two dications in the asymmetric unit were similar, thus data for only one of the two are presented in Tables 2 and 3 and Figure 2B. Similar to **5**, the dication $[\text{Ru}(\text{Me}_3\text{TPA})(\text{CH}_3\text{CN})_2]^{2+}$ of **8** possesses a distorted octahedral geometry. Most of the bond angles in **8** are in good agreement with angles observed in **5**, varying by only three degrees, with the exception of angles involved with the nitrile donors. As shown in Figure 2B, introduction of the methyl groups into the ligand results in a severe distortion of the N1–Ru1–N5 bond angle to $104.9(1)^\circ$, as compared with the same three atoms of **5**, which showed a bond angle of only $93.9(2)^\circ$. This distortion results in a contraction of the bond angle between the two nitrile donors in **8**, with N1–Ru1–N6 equal to $79.9(1)^\circ$, as compared to $88.8(2)^\circ$ for **5**. As opposed to **5**, bond distances to the two nitriles of **8** are not identical within error. The Ru1–N1 distance of $2.064(2)$ Å of **8** is considerably longer than the Ru1–N6 distance of $2.028(2)$ Å, presumably because of steric crowding by the three methyl groups present on the ligand Me_3TPA . As expected, ruthenium bond distances to the three pyridine N-donors in **8** (N2, N4 and N5) are also considerably longer than those observed for **5**, ranging from $2.125(3)$ to $2.130(2)$ Å.

Electronic Absorption Data

Complexes **5–7** show maxima in acetone, DMSO, and CH_3CN ranging from 365 to 380 nm, consistent with metal-to-ligand charge transfer (MLCT) bands. The spectra of these complexes are largely independent of solvent, consistent with the $[\text{Ru}(\text{L})-(\text{CH}_3\text{CN})_2]^{2+}$ dications remaining intact in weakly and strongly coordinating solvents, where $\text{L} = \mathbf{1–3}$. In contrast, spectra for **8** become broader in acetone and DMSO relative to CH_3CN , with increased absorbance at wavelengths >400 nm, consistent with dynamic behavior between $[\text{Ru}(\text{Me}_3\text{TPA})(\text{CH}_3\text{CN})_2]^{2+}$ and solvent. Exchange of bound CH_3CN groups with solvents was explored further with ^1H NMR spectroscopic analysis.

^1H NMR Spectroscopic Data

^1H NMR spectroscopic analysis was used to gain further insight into the structures of complexes **5–8** in solution, and to measure ratios of isomers for compounds **6** and **7**. ^1H NMR spectroscopic analysis of complex **5** in acetone- $d_6/\text{D}_2\text{O}$ (9:1) indicated the presence of two distinct CH_3CN ligands, with singlets at 2.88 and 2.47 ppm, consistent with the structure shown in Figure 2A with one CH_3CN group *cis* and one CH_3CN *trans* to the basic nitrogen of TPA. The CH_3CN group containing N6 is assigned as the farthest upfield signal at 2.47 ppm, based on NOESY data.²⁵ This structural assignment is further supported by the fact that an upfield shift is expected for this CH_3 group, due to shielding effects of the two *cis*-pyridine rings' π -systems. Doublets at 9.18 and 8.83 ppm, integrating for 1 and 2 protons, respectively, are assigned as resonances belonging to the 6-pyridyl C–H protons of TPA based on their downfield nature and 5.9 Hz coupling constants.⁷⁵ These data are consistent with $[\text{Ru}(\text{TPA})-(\text{CH}_3\text{CN})_2]^{2+}$ containing a plane of symmetry where two pyridyl groups are equivalent (see pyridine rings containing N2 and N4 donors, Figure 2A). In addition, two doublets and one singlet, each integrating for two protons, are observed

between 5.25 and 4.86 ppm. ^1H COSY analysis indicates coupling between doublets at 5.25 and 5.15 ppm. With coupling constants of 15.6 Hz observed by ^1H NMR spectroscopy, which are in the range for geminal coupling, these protons are assigned as two CH_2 units, related by the plane of symmetry, each possessing diastereotopic protons. The last singlet present at 4.86 ppm is assigned to a CH_2 unit with equivalent protons. In this case, the plane of symmetry passing through the pyridine and two MeCN ligands bisects the two protons, making them equivalent by symmetry so geminal coupling is not observed.

Spectral data from **5** were used to assign the structures of major and minor isomers for **6** and **7**, based on the aforementioned considerations of symmetry. Ruthenium complexes derived from MeTPA were isolated as a 2:1 mixture of stereoisomers, complexes **6a** and **6b**, respectively. The spectrum of **6a** and **6b** in acetone- d_6 shows three doublets that are furthest downfield from 9.4 to 8.8 ppm, which are assigned as 6-pyridyl C–H protons based on arguments presented with **5**. The major isomer **6a** contains pyridine donors that are nonequivalent by symmetry. Thus, two signals are observed. In the minor isomer **6b**, the two pyridyl donors are equivalent by symmetry, thus only one signal at 9.05 ppm is observed. In addition, the methyl group present in **6a** breaks the plane of symmetry observed with **5**, making each of the three CH_2 units nonequivalent, giving **6a** a total of 6 diastereotopic protons that are evident in the spectrum from 5.5 to 4.7 ppm. Complexes derived from Me_2TPA were isolated as a 1.5:1 mixture, complexes **7a** and **7b**, respectively. The major isomer **7a** shows a 6-pyridyl C–H resonance at 9.6 ppm, whereas the minor isomer **7b** shows a resonance at 9.05 ppm. Two doublets and a singlet are observed as major peaks in the region from 5.5 to 4.6 ppm, consistent with the plane of symmetry present in **7a**. Signals for the six diastereotopic protons belonging to the CH_2 units of **7b** are observed as minor peaks. In both cases, placing the 6-methylpyridyl group in the axial position leads to the least stable isomer. These product ratios are in good agreement with DFT calculations that measured relative thermodynamic stabilities for **6a/b** and **7a/b** (vide infra).

^1H NMR spectra were recorded for the Me_3TPA complex **8** in multiple solvents. In acetone- d_6 , complex **8** displays many of the same features present in **5**, including 3-, 4-, and 5-pyridyl C–H protons present in a 2:1 ratio from 7.8 to 7.05 ppm, as well as two doublets and a singlet present from 5.5 to 4.7 ppm, assigned to the CH_2 units. These data are consistent with the plane of symmetry present in **8**. Resonances assigned to the 6-ethylpyridyl groups are located at 3.35 and 2.97 ppm, integrating for 3 and 6 protons, respectively. Resonances assigned to the bound CH_3CN groups occur at 2.94 and 2.60 ppm, with one upfield and one downfield signal, the same as observed with **5**. Interestingly, the signal for the downfield CH_3CN group is not evident in the spectrum of **8** recorded in CD_3CN . Immediately upon dissolution of **8** in CD_3CN , the resonance for the downfield N1 CH_3CN group disappears and a strong peak at 2.05 ppm is observed, consistent with free CH_3CN . The signal for the upfield N6 CH_3CN group at 2.4 ppm remains unchanged and does not decrease in intensity, even after storing in CD_3CN solution for 18 h in the dark. These data are consistent with one CH_3CN ligand (see N1, Figure 2B) undergoing facile exchange in coordinating solvents, whereas in weakly coordinating solvents, such as acetone- d_6 , the N1 CH_3CN group remains bound.

Photochemical Studies

Photochemical studies were carried out with **5** and **6** in CH₂Cl₂ with 10 mM Bu₄NCl and in aqueous solution. Control experiments using electronic absorption spectroscopy confirm that **5** and **6** are stable in the dark for over 1 h, whereas **7** and **8** show spectral changes over the same time period (Figures S4–6). These data are consistent with **7** and **8** showing a higher rate of thermal ligand exchange than **5** and **6**, making photochemical release data for **7** and **8** difficult to investigate. Therefore, photochemical studies with **7** and **8** were not undertaken. The photoinduced reactions of **5** and **6** in CH₂Cl₂ with 10 mM Bu₄NCl and in H₂O (<5% acetone) were monitored by electronic absorption spectroscopy and the resulting spectra as a function of irradiation time are shown in Figures 4A and 4B, respectively (λ_{irr} = 395 nm). The exchange of CH₃CN for Cl[−] or H₂O in **5** occurs with a decrease in the absorption feature at 375 nm and a concomitant increase in the peak at 475 nm with an isosbestic point at 398 nm in the Cl[−] system (Figure 4A), and an increase in the peak at 397 nm with an isosbestic point at 384 nm is observed in H₂O (Figure 4C). The quantum yield for this ligand exchange in **5** was measured as 0.016(1) in CH₂Cl₂ with 10 mM Bu₄NCl and 0.0090(2) in H₂O (λ_{irr} = 400 nm). Similar irradiation of complex **6**, which exists in two isomers (**6a** and **6b**, Figure 5A), exhibits analogous spectral changes but no isosbestic point is observed in the electronic absorption spectra upon irradiation due to either the formation of more than one photoproduct or different rates of reaction of the two isomers (Figures 4B and 4D). The presence of two photoactive isomers of **6** in solution precludes measurement of an accurate quantum yield of ligand dissociation; however, an apparent quantum yield of 0.041(4) was determined by monitoring the decrease in absorbance of the reactant peak at 370 nm at early irradiation times (0–15 min) in CH₂Cl₂ with 10 mM Bu₄NCl, and a value of 0.011(2) was measured in H₂O using the same procedure (λ_{irr} = 400 nm). This increase in efficiency for **6a/b** relative to **5** by addition of a methyl substituent is consistent with a stabilized ³MC state due to the addition of steric bulk, which distorts the pseudo-octahedral geometry around the Ru(II) center; this effect has been demonstrated in a variety of related Ru(II)-diimine complexes.^{76,77}

To unequivocally gain structural information on the photoproducts, samples of **5** and **6a/b** were photolyzed in CD₃CN with λ_{irr} = 365 nm and monitored by ¹H NMR spectroscopy. In addition to determination of the products following CH₃CN ligand exchange, these experiments also provide evidence that no other photochemical pathways are operative in CD₃CN (Figures S1–2). When a CH₃CN ligand is exchanged for CD₃CN, the electronic and chemical structures of the complex do not change, such that the signals for the protons of all remaining bound ligands on the complex remain unchanged. In this case, if no other photochemistry is taking place, then the only changes observed are the concomitant loss of the signal of the bound CH₃CN ligand and the growth of that associated with free CH₃CN. The labeling scheme for the CH₃CN ligands and a plot of the integration of the CH₃CN protons as a function of irradiation time are provided in Figure 5.

The sample for **5** consisted of a single isomer, whereas **6** was composed of the two isomers **6a** and **6b** in a 2:1 ratio, respectively. As expected from previous work with **5**, the CH₃CN ligand coordinated *trans* to a pyridine, labeled as N6 (Figure 5A), is exchanged for CD₃CN much more facile exchange than the N1 CH₃CN ligand (Figure 5B). For **6a/b** (Figure 5C),

the N6 acetonitrile ligands exchanged on a similar time scale to that observed with **5**. However, the N1 CH₃CN ligand for **6a/b** underwent more rapid exchange than that observed for **5**. This finding indicates that the placement of the methyl group on the TPA ligand generally increases the photolability of the CH₃CN ligands.

The irradiation of the 2:1 mixture of **6a/6b** in weakly coordinating CD₃OD resulted in unexpected isomerization that provides an inversion of the isomer ratio. ¹H NMR spectra of a solution of **6a/6b** in CD₃OD were recorded as a function of irradiation time (λ_{irr} 395 nm; Figure S7). Prior to photolysis, three doublets appear furthest downfield at 8.72 and 9.20 ppm, corresponding to the protons in the 6-position of the two inequivalent pyridyl rings in **6a**, and at 8.91 ppm, corresponding to the protons in the 6-position of the two equivalent pyridyl rings in **6b** (Figure S7A). The 2:1 **6a:6b** mixture of isomers is evident by the equal integrations of these three doublets. Upon irradiation, as free CH₃CN appears along with the loss of bound CH₃CN (Figure S7B), the three downfield doublets decrease in intensity while two new small doublets at 8.92 and 9.08 ppm and one large doublet at 9.11 ppm grow as a function of irradiation time (Figure S7A). The ¹H NMR spectrum of the photoproduct is consistent with [Ru(MeTPA)(CH₃CN)(CD₃OD)]²⁺ in two different isomers, **6a-CD₃OD** or **6b-CD₃OD**, whose structures are depicted in Figure 6. The two small doublets, corresponding to **6a-CD₃OD**, integrate to one proton each, while the large doublet, corresponding the **6b-CD₃OD**, integrates to 4 protons; these integrations translate to a 1:2 ratio of **6a-CD₃OD** and **6b-CD₃OD**, respectively. It may be concluded from the change in distribution of isomers that a pyridyl arm of the MeTPA ligand is labile and is able to replace the weakly coordinated CD₃OD ligand, as shown in Figure 6. Conversely, this process does not occur when the sample is irradiated in CD₃CN, discussed above, because CD₃CN is a much stronger ligand than CD₃OD, such that it is not displaced by the pyridine arm of MeTPA.

To confirm the inversion of isomer distribution, the irradiated sample containing **6a-CD₃OD** and **6b-CD₃OD** was dried by removing the CD₃OD solvent, and the product was dissolved in CD₃CN. The CD₃CN solvent displaced the weaker CD₃OD ligand to provide [Ru(MeTPA)(CH₃CN)-(CD₃CN)]²⁺ (**6a-CD₃CN** and **6b-CD₃CN**, Figure 6), which are expected to have identical chemical shifts in the aromatic region of the ¹H NMR spectrum compared to **6a** and **6b**, but with different integrations as determined by the isomer ratio. These two spectra are compared in Figure S8, showing the initial 2:1 mixture of **6a** and **6b** in CD₃CN and the 1:2 mixture of **6a-CD₃CN** and **6b-CD₃CN** products.

Analysis of the chloride-substituted [Ru(MeTPA)(CH₃CN)-Cl]⁺ product (**6a-Cl/6b-Cl**) by ¹H NMR was achieved by irradiation of **6a/6b** in CD₃OD with λ 395 nm until the CH₃CN *trans* to py/Me-py was fully replaced by CD₃OD, followed by addition of excess CaCl₂, a CD₃OD-soluble source of Cl⁻ ions. The solution was allowed to react in the dark overnight, and the resulting NMR spectrum shows formation of **6a-Cl** and **6b-Cl** in a 1:2 isomer ratio, respectively (Figure S9). The two small doublets at 8.76 and 9.16 ppm correspond to the protons in the 6-position of the inequivalent pyridyl groups in **6a-Cl**, and the large doublet at 8.95 ppm corresponds to the protons in the 6-position of the two equivalent pyridyl groups in **6b-Cl**. The scheme depicting the photochemical release of

CH₃CN, rearrangement of the MeTPA ligand, and coordination of chloride is provided in Figure 7.

COMPUTATIONAL CHEMISTRY

Calculated Structures and Energies

Calculations were carried out to gain insight into the structure and thermal reactivities of **5**–**8**. As illustrated in Tables 2 and 3, the DFT-calculated structures are in good agreement with the X-ray crystallographic data for complexes **5** and **8**. However, the agreement is not quantitatively accurate plausibly because of the lack of crystal packing effect in the calculated structures. Calculations were performed in order to understand the relative stabilities of the isomers of complexes **6** and **7** (Table S1, Figure S3). Compound **6a** is found to be lower in energy ($E = 3.1$ kcal/mol) than its isomer **6b**, and **7a** is calculated to be about 1 kcal/mol more stable than **7b**. These results are consistent with the observations from the NMR data, which show **6a** and **7a** are the major isomers for complexes **6** and **7**, respectively. Inspection of the molecular geometries shows that in complex **6a**, the methyl group on the N2-pyridine moiety has one *cis*-pyridine group at the N5 position while the methyl group in **6b** has two *cis*-pyridine groups at the N2 and N4 positions. Consequently, the methyl moiety in the former complex is embedded in an environment that is sterically less crowded than that in the latter compound. In addition, as displayed in Figure 8, there is an unfavorable steric interaction between the methyl moiety and the acetonitrile ligand at the N1 position in complex **6b**. The acetonitrile ligand bends away from the methyl group to relieve this repulsion and, as a result, the angle N6–Ru–N1 is contracted to about 83.6° giving rise to a distorted octahedral geometry of the complex. Complex **7b** containing two methyl-pyridine moieties *cis* to each other is found to be less stable than **7a** having two methyl-pyridine moieties *trans* to each other (Figure S3).

Thermal Dissociation Properties

As observed by ¹H NMR spectroscopy, the CH₃CN ligand at the N1 position in complex **8** is found to be labile as it dissociates immediately upon dissolution of the complex in CD₃CN in the absence of light. In contrast, no CH₃CN dissociation was found for complex **5** under similar conditions. Relaxed potential energy scans were performed for **5** and **8** to gain insights into these different dissociation behaviors. The Ru–NCCH₃ bonds were elongated at N1 and N6 positions for each of the complexes with an increment of 0.1 Å in each step while the rest of the molecule was reoptimized. As illustrated in Figure 9, the dissociation of the CH₃CN ligand at the N1 position in complex **8** is associated with an estimated energy barrier of about 12 kcal/mol. The estimated barriers for the dissociation events of N6-CH₃CN in complex **8** and the two CH₃CN ligands in **5** are found to be higher, ranging from 20 to 21 kcal/mol. These results are consistent with ¹H NMR spectroscopic data, which showed that complex **8** undergoes rapid dissociation of the N1 CH₃CN ligand. In contrast, the N6 CH₃CN ligand of **8** and the N1 and N6 CH₃CN groups of **5** show no measurable dissociation under ambient conditions, as judged by ¹H NMR spectroscopic analysis.

Differences in thermal dissociation behavior between **5** and **8** can be explained by considering their respective ground state structures. The N1 CH₃CN ligand in **8** is sterically

encumbered between the three methyl-substituted pyridine moieties while the absence of the methyl groups on the pyridine moieties in **5** gives rise to a less sterically crowded environment for the N1 CH₃CN ligand. In order to relieve the steric congestion the N1–Ru–N6 angle decreases by 3.5° from **5** to **8**, and the Ru–N1–C(CH₃CN) bond angle decreases from 177.3° in **5** to 170.6° in **8**. Consequently, the Ru–NCCH₃ interaction in **8** is weakened and the N1 CH₃CN ligand becomes more susceptible to dissociation compared to that in complex **5**. Dissociation of the N1 CH₃CN ligand in **8** relieves the steric repulsion present between the two CH₃CN ligands. The Ru–N–C(CH₃CN) bond angle at the N1 position in **8** is calculated to deviate significantly from the linear arrangement when compared to all other Ru–N–C(CH₃CN) bond angles (177–178°) of complexes **5** and **8**. As a result, the release of the N1 CH₃CN ligand is calculated to be facile in **8** with an estimated activation barrier of about 12 kcal/mol, which is the lowest among the barriers of all CH₃CN dissociations in **5** and **8**.

The relaxed potential energy scans were also performed for the major and minor isomers of complexes **6** and **7** to gain further insights into their dissociation behaviors. The Ru–NCCH₃ bonds were elongated at N1 and N6 positions for each of the complexes with an increment of 0.1 Å in each step while the rest of the molecule was reoptimized (Figure S10). In both complexes the energy barrier for dissociation of the N1–CH₃CN ligand is the lowest in the case of minor isomer, 16.7 kcal/mol for **6b** and 15.1 kcal/mol for **7b**, whereas in **6a** and **7a** the barriers are somewhat higher, 19.0 and 17.2 kcal/mol, respectively. In both minor isomers **6b** and **7b**, methyl groups of the axial pyridyl donors distort the CH₃CN ligand at the N1 position, decreasing the N6–Ru–N1 bond angle, and making the CH₃CN ligand more labile. Taken together, these results indicate that addition of steric bulk around the N1 CH₃CN groups can be used to tune dissociation energies between 12 and 22 kcal/mol for **5**–**8**.

Photochemical Dissociation Properties

DFT calculations were used to gain further insight into the excited state behaviors of **5**, **6a**, and **6b**. Excited state energies were not examined for **7a/b** and **8**, because of their higher rates of thermal ligand exchange. In our previous study we used SCF calculations to characterize the ³MLCT states for the Ru complexes with quinoline and bipyridine acceptor ligands.²⁸ In our present study the Ru complexes have pyridine ligands which are known to be poorer acceptors than the quinoline and bipyridine ligands. We were unable to optimize the ³MLCT states for the Ru-pyridine complexes with SCF calculations. After many attempts, we concluded that these ³MLCT states are high in energy and without stable minima, causing the calculations to converge to lower energy ³MC states. The six coordinate ³MC states can lose a CH₃CN ligand from N1 or N6 to yield more stable five coordinate complexes that are 45–50 kcal/mol above the *S*₀ ground state for **5** (see Figure 10). Loss of CH₃CN from N6 leads to ³MC₁, a trigonal bipyramidal complex (with N3–Ru–N1 as the axis) that is 5 kcal/mol more stable than the loss of CH₃CN from N1 which forms ³MC₂, a square pyramidal structure (with Ru–N3 as the axis).²⁸ These results are in good agreement with previous observations,²⁵ which showed that photochemical dissociation of the N6 CH₃CN is significantly more facile than the N1 CH₃CN for Ru-quinoline and Ru-bipyridine complexes.²⁸ The N6 CH₃CN was found to be more labile than

the N1 CH₃CN in the triplet excited states because of a favorable orbital mixing between the aromatic ligand π^* and the Ru $d\sigma^*$ orbitals that characterize the Ru–N6 CH₃CN dissociation. By contrast, the N1 CH₃CN is more labile than the N6 CH₃CN in the ground states for **6**–**8** because of the steric repulsion by the methyl groups.

The ³MC₁ and ³MC₂ states of **6a** are 43.7 and 46.9 kcal/mol higher in energy, respectively, than the ground state **6a**, while the ³MC₁ and ³MC₂ states of **6b** are 40.6 and 44.6 kcal/mol higher, respectively, than the ground state **6b** (see Figure 11). Inspection of the ground state geometries of **6a** and **6b** finds that the steric repulsion present between the methyl group on the N5 pyridine and the N1 CH₃CN in **6b** distorts the octahedral field of the complex relative to that of **6a**. The Ru–N1–C(CH₃CN) and N6–Ru–N1 angles in the ground state of **6b** are 170.5° and 83.6°, respectively, while the same bonds in **6a** show angles of 174.2° and 86.9°, respectively. As a result, the ground state of **6b** is 3.1 kcal/mol higher in energy than that of **6a**. In both **6a** and **6b**, the ³MC₁ states resulting from the loss of the N6 CH₃CN ligands are lower in energy than the ³MC₂ states resulting from the loss of the N1 CH₃CN ligands. This agrees well with the observation that the N6 CH₃CN undergoes exchange with CD₃CN with shorter irradiation times than the N1 CH₃CN for both the complexes. Moreover, the ³MC₁ states for **6a** and **6b** are lower in energy than the ³MC₁ state of **5** (43.7 and 40.6 kcal/mol vs 45.5 kcal/mol) and the ³MC₂ states for **6a** and **6b** are lower in energy than the ³MC₂ state of **5** (46.9 and 44.6 kcal/mol vs 50.4 kcal/mol) when compared to their respective ground states. These data are consistent with the 2.6-fold increase in the quantum yield of ligand dissociation observed for **6a** and **6b** relative to **5**. Thus, both steric interactions destabilizing the ground state **6a** and **6b**, and electronic interactions stabilizing the ³MC₁ excited state relative to the ³MC₂ state contribute to the greater quantum yield for selective photochemical nitrile release when **6a** and **6b** are compared to **5**.

CONCLUSIONS

In conclusion, we report the synthesis and characterization of a series of Ru-caged nitrile complexes derived from the ligand TPA. Steric effects on ground state stability and thermal and photochemical reactivity were studied by the consecutive introduction of methyl groups onto the 6-positions of the pyridyl donors of **5**. Overall, the data indicate that a delicate balance exists between introducing steric bulk to accelerate photochemical dissociation of the nitrile donors vs accelerating thermal dissociation in the dark, which compromises the effectiveness of the caging group. Introduction of one methyl group into **5** leads to a mixture of stereoisomers (**6a** and **6b**) with roughly 2.6-fold higher quantum efficiency for photochemical nitrile release than **5**. Adding more methyl substituents, as in the case of **7a/b** and **8**, leads to steric crowding and higher rates of nitrile release in the dark. Computational results, supported by experimental data, indicate that barriers for thermal nitrile exchange can be tuned by over 10 kcal/mol by introducing steric crowding around the nitrile donors. Taken together, these results show that nitrile release can be tuned in Ru(TPA) complexes, which may aid in the design of new caging groups.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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1) TPA ($R_1, R_2, R_3 = H$)

2) MeTPA ($R_1 = Me, R_2, R_3 = H$)

3) Me₂TPA ($R_1, R_2 = Me, R_3 = H$)

4) Me₃TPA ($R_1, R_2, R_3 = Me$)

Figure 1.

Structures of the tetradentate ligands TPA (1), MeTPA (2), Me₂TPA (3), and Me₃TPA (4).

A **B** **Figure 2.**

ORTEP diagrams of the dication $[\text{Ru}(\text{TPA})(\text{MeCN})_2]^{2+}$ from compound **5** (A)²⁵ and $[\text{Ru}(\text{Me}_3\text{TPA})(\text{MeCN})_2]^{2+}$ from compound **8** (B). Thermal ellipsoids are shown at 50% probability. Hydrogen atoms are omitted for clarity.

Figure 3.
Assigned structures for major and minor isomers of **6** and **7**.

Figure 4.

Electronic absorption spectra of **5** (A) and **6** (B) irradiated for 0–45 min with $\lambda_{\text{irr}} = 395$ nm in CH_2Cl_2 with 10 mM Bu_4NCl , and for **5** (C) and **6** (D) irradiated for 0–25 min with $\lambda_{\text{irr}} = 395$ nm in H_2O (<5% acetone).

Figure 5.

(A) Methyl proton labeling scheme for **5** and **6a–b**. Plots of integration vs irradiation time in CD_3CN with $\lambda_{\text{irr}} = 365 \text{ nm}$ determined by ^1H NMR for (B) **5** (3 mM), Ru–N1 (red filled circle), Ru–N6 (green filled triangle), free CH_3CN (black filled square), and (C) **6a/b** (3 mM), Ru–N1 (**6a**) (red filled circle), Ru–N1 (**6b**) (blue filled diamond), Ru–N6 (**6a**) and Ru–N6 (**6b**) (green filled triangle), free CH_3CN (black filled square).

Figure 6.

Scheme representing the photodissociation of CH_3CN in **6a** and **6b** in CD_3OD followed by removing the CD_3OD solvent and redissolving in CD_3CN .

Figure 7.

Schematic representation of the photosubstitution of CH₃CN in complexes 6a and 6b in CD₃OD followed by addition of Cl⁻.

Figure 8. Calculated optimized structures of **6a** and **6b**. The acetonitrile ligand at the N1 position in **6b** bends away from the methyl group of the N5-pyridine moiety and it leads to a distorted octahedral geometry with the N6–Ru–N1 angle squeezed to 83.6°.

Figure 9. Relaxed potential energy scans of Ru–N1 and Ru–N6 acetonitrile ligands in complexes **5** (Ru–N1, green filled triangle; Ru–N6, black filled square) and **8** (Ru–N1, blue filled diamond; Ru–N6, red filled circle). The Ru–N bond is elongated by 0.1 Å in each step and the rest of the molecular geometry is reoptimized.

Figure 10.

Calculated energies of the ^3MC states relative to the ground state S_0 for $[\text{Ru}(\text{TPA})(\text{CH}_3\text{CN})_2]^{2+}$ **5**. Each ^3MC state is five-coordinate and has completely lost one acetonitrile ligand. The energies have been calculated as $\Delta E(\text{kcal/mol}) = E_{^3\text{MC}} + E_{\text{CH}_3\text{CN}} - E_{S_0}$.

Figure 11.

Calculated energies of the ^3MC states of $[\text{Ru}(\text{MeTPA})(\text{CH}_3\text{CN})_2]^{2+}$ isomers, **6a** and **6b**, relative to their respective ground states S_0 . Each ^3MC state is five-coordinate and has completely lost one acetonitrile ligand. The energies have been calculated as

$$\Delta E(\text{kcal/mol}) = E_{^3\text{MC}} + E_{\text{CH}_3\text{CN}} - E_{S_0}.$$

Scheme 1. Synthesis of [Ru(L)(MeCN)](PF₆)₂ Complexes, Where L = TPA (5), MeTPA (6), Me₂TPA (7), and Me₃TPA (8)^a

^aConditions: (a) (1) *cis*-[RuCl₂(Me₂SO)₄] (1 equiv), MeOH, 70 °C, 4 h, (2) CH₃CN/H₂O (1:1) 80 °C, 16 h; (b) (1) *cis*-[RuCl₂(Me₂SO)₄] (1 equiv), EtOH, 90 °C, 18 h, (2) CH₃CN/H₂O (1:1) 80 °C, 18 h.

Table 1X-ray Crystallographic Data for 5²⁵ and 8

	[Ru(TPA)(CH₃CN)₂] (PF₆)₂ (5)	[Ru(Me₃TPA)(CH₃CN)₂] (PF₆)₂ (8)
empirical formula	C ₄₉ H ₅₀ F ₅₇ N ₁₅ P ₄ Ru ₂	C ₂₇ H ₃₃ F ₁₂ N _{6.50} P ₂ Ru
formula weight	1651.12	839.61
crystal system	triclinic	monoclinic
space group	<i>P</i> $\bar{1}$	<i>C</i> 2/ <i>c</i> 1
<i>a</i> (Å)	12.1129(8)	12.5538(11)
<i>b</i> (Å)	12.2928(8)	12.7608(11)
<i>c</i> (Å)	21.8490(15)	39.841(4)
α (deg)	94.436(3)	90
β (deg)	97.000(3)°	93.501(3)
γ (deg)	91.324(3)°	90
<i>V</i> (Å ³)	3217.7(4)	6370.5(10)
<i>Z</i>	4	8
final R indices [<i>I</i> > 2 σ (<i>I</i>)]	R1 = 0.0579, wR2 = 0.1853	R1 = 0.0530, wR2 = 0.1239
R indices (all data)	R1 = 0.0771, wR2 = 0.1952	R1 = 0.0543, wR2 = 0.1245

Table 2A Comparison of X-ray Data and DFT Calculations for Selected Bond Distances (Å) for 5²⁵ and 8

bond	complex 5		complex 8	
	X-ray data	DFT calcd	X-ray data	DFT calcd
Ru1–N1	2.031(5)	2.007	2.064(2)	2.017
Ru1–N2	2.062(4)	2.087	2.128(3)	2.158
Ru1–N3	2.053(4)	2.095	2.055(2)	2.093
Ru1–N4	2.071(4)	2.085	2.125(3)	2.162
Ru1–N5	2.056(4)	2.064	2.130(2)	2.156
Ru1–N6	2.037(5)	2.020	2.028(2)	1.999

Table 3Comparison of X-ray Data and DFT Calculations for Selected Bond Angles (deg) for 5²⁵ and 8

bond angle	complex 5		complex 8	
	X-ray data	DFT calcd	X-ray data	DFT calcd
N1–Ru1–N2	96.4(2)	98.3	96.1(1)	100.3
N1–Ru1–N4	99.5(2)	98.3	101.6(1)	97.9
N1–Ru1–N5	93.9(2)	95.3	104.9(1)	101.7
N1–Ru1–N6	88.8(2)	87.3	79.9(1)	83.8
N2–Ru1–N3	81.7(2)	81.5	80.4(1)	82.7
N2–Ru1–N5	90.4(2)	90.3	89.9(1)	85.9
N2–Ru1–N6	89.1(2)	90.0	91.7(1)	89.5
N3–Ru1–N4	82.6(2)	82.0	81.9(1)	78.9
N3–Ru1–N5	82.6(2)	82.6	83.4(1)	82.0
N3–Ru1–N6	94.8(2)	94.8	92.0(1)	92.8
N4–Ru1–N5	90.0(2)	88.2	87.4(1)	94.8
N4–Ru1–N6	89.7(2)	90.8	89.5(1)	88.1