Counterion Effects in $[Ru(bpy)_3](X)_2$-Photocatalyzed Energy Transfer Reactions

Juliette Zanzi,[a] Zachary Pastorel,[b] Carine Duhayon,[a] Elise Lognon,[c] Christophe Coudret,[d] Antonio Monari,[c] Isabelle M. Dixon,*[e] Yves Canac,*[a] Michael Smietana*[b] and Olivier Baslé*[a]

[a]*LCC-CNRS, Université de Toulouse, CNRS, UPS, Toulouse, France*

[b]*Institut des Biomolécules Max Mousseron, Université de Montpellier, CNRS, ENSCM, Montpellier, France*

[c]*ITODYS, Université Paris Cité and CNRS, F-75006 Paris, France*

[d]*Université de Toulouse, UPS, Institut de Chimie de Toulouse, FR2599, 118 Route de Narbonne, F-31062 Toulouse, France*

[e]*LCPQ, Université de Toulouse, CNRS, Université Toulouse III - Paul Sabatier, 118 route de Narbonne, F-31062 Toulouse, France*

ABSTRACT: Photocatalysis that uses the energy of light to promote chemical transformations by exploiting the reactivity of excited state molecules is at the hearth of a virtuous dynamic within the chemical community. Visible light metal-based photosensitizers are most prominent in organic synthesis, thanks to their versatile ligand structure tunability allowing to adjust photocatalytic properties towards specific applications. Nevertheless, a large majority of these photocatalysts are cationic species whose counterion effects remains underestimated and overlooked. In this report, we show that modification of the X counterions constitutive of $[Ru(bpy)_3](X)_2$ photocatalysts modulates their catalytic activities in intermolecular [2+2] cycloaddition reactions operating through triplet-triplet energy transfer (TTEnT). Particularly noteworthy is the dramatic impact observed in low-dielectric constant solvent over the excited state quenching coefficient, which varies by two orders of magnitude depending on whether X is a large weakly bound ($BAr^F{}_4^-$) or a tightly bound anion (TsO^-). In addition, the counterion identity also greatly affects the photophysical properties of the cationic ruthenium complex, with $[Ru(bpy)_3](BAr^F{}_4)_2$ exhibiting the shortest ^{3}MLCT excited state lifetime, highest excited state energy and photostability, enabling remarkably enhanced performance (up to >1000 TON at low 500 ppm catalyst loading) in TTEnT photocatalysis. These findings supported by density functional theory-based molecular modeling demonstrate that counterions have a critical role in modulating cationic transition metal-based photocatalyst potency, a parameter that should be taken into consideration also when developing energy transfer processes.

INTRODUCTION

Photocatalysis under visible light irradiation has witnessed tremendous successes in recent years in a wide variety of applications,[1] and has notably initiated a paradigm shift in organic synthesis.[2] Within the design of efficient molecular photocatalytic systems, the architecture of metal-based photocatalysts can easily be tuned to efficiently achieve the desired photochemical properties required to promote specific transformations.[3] Thanks to their strong absorption in the visible region, rapid intersystem crossing, and long-lived triplet excited state, polypyridyl ruthenium(II) and cyclometalated iridium(III) complexes are the most widely explored classes of photoredox catalysts and photosensitizers.[4] The common method to manipulate excited state properties of such photocatalysts relies on modifying the ligand scaffold around the metal center: a *modus operandi* which has so far provided access to a plethora of photocatalysts covering a broad range of redox potentials and triplet excited state energies.[5,6] Importantly, a large majority of these Ir(III) and Ru(II) complexes, but also of emerging earth-abundant transition metal-based luminescent complexes,[7] are cationic species associated with counterion(s) to ensure charge balance of the molecular photocatalysts. Nevertheless, while photocatalytic reaction cycles are usually modeled considering the properties of the sole organometallic cation, the true nature of the photocatalyst is that of a salt in which the anions could play a role beyond the one of mere spectators.[8] In this respect, recent reports have brought to light the influence of counterion identity on the efficiency of single electron transfer (SET) processes operated by cationic iridium and ruthenium photoredox catalysts (Figure 1A).[9] However, despite the fact that both SET and energy transfer (EnT) photocatalyses rely on the triplet metal-to-ligand charge-transfer (^{3}MLCT) excited state of the photocatalyst, counterion effects on EnT processes have not been demonstrated yet. In triplet-triplet energy transfer (TTEnT) processes, a burgeoning field of research with a wide range of synthetic applications,[10] the photocatalyst is excited by the direct absorption of visible light and subsequently transfers its triplet excited state energy to a desired substrate, opening otherwise elusive relaxation channels and photoreactive routes, leading to original chemical transformations.[11] Herein, we show that modification of the X counterions in the emblematic $[Ru(bpy)_3](X)_2$ complex series has a dramatic impact on their photophysical properties and photocatalytic activity in TTEnT. Specifically, we demonstrate that large non-coordinating tetrakis[3,5-bis(trifluoromethyl) phenyl]borate ($BAr^F{}_4^-$) anions are highly beneficial giving rise to much higher photocatalytic activity (up to >1000 TON) in intermolecular [2+2] cycloaddition reactions of olefins to yield cyclobutanes (Figure 1B).[12] We rationalize the superior photocatalytic performance of $[Ru(bpy)_3](BAr^F{}_4)_2$ over its congeners on the basis of experimental and density functional theory (DFT) studies that indicate an increase of both the energy of the ^{3}MLCT and the metal-centered (^{3}MC) excited state, which are respectively responsible for a larger energy transfer driving force and an

improved photostability. Our results provide fundamental insights into how counterions impact cationic transition metal-based photocatalyst potency in triplet energy transfer-activated reactions, a topic of great interest but surprisingly overlooked so far in the literature.

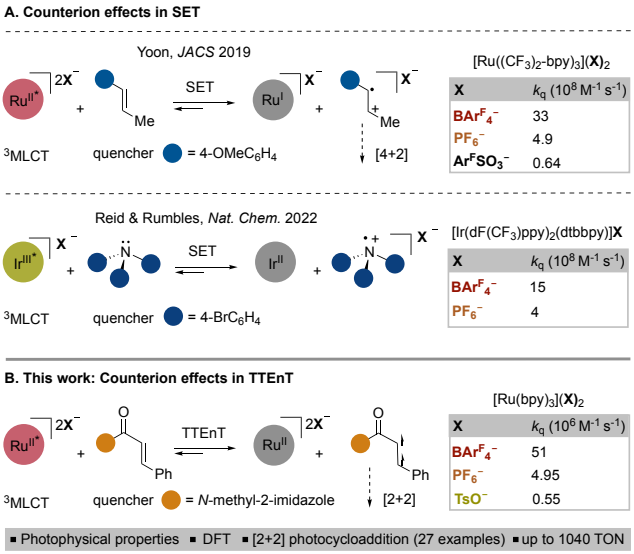


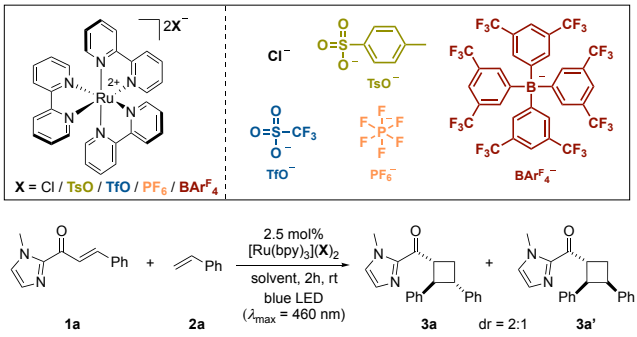
Figure 1. Counterion dependence in photochemical processes. (A) Previously reported counterion effects on single electron transfer (SET). (B) Counterion effects in $[Ru(bpy)_3]X_2$-catalyzed intermolecular [2+2] cycloaddition enabled by energy transfer. Bimolecular quenching rate coefficient (k_q). 3,5-bis(trifluoromethyl)benzenesulfonate ($Ar^FSO_3^-$). Turnover number (TON).

RESULTS AND DISCUSSION

As a model reaction to evaluate the impact of counterion effect in TTEnT photocatalysis, we selected the [2+2]-cycloaddition of α,β-unsaturated 2-acyl imidazole **1a** with styrene **2a** using $[Ru(bpy)_3](X)_2$ as triplet sensitizer under visible light irradiation. The choice of the reaction has been made in light of recent studies on "triplet activation" of **1a**,[13] in particular that of Yoon and coworkers,[13b] who showed that the presence of trifluoromethanesulfonic acid as a Brønsted acid co-catalyst is strictly necessary to efficiently promote this $[Ru(bpy)_3]^{2+}$-catalyzed [2+2] photocycloaddition in acetonitrile. The effect was justified by the fact that the protonated acyl-imidazole substrate is a better energy acceptor for the triplet excited state of $[Ru(bpy)_3]Cl_2$*. We therefore assumed that by modifying the counterions of $[Ru(bpy)_3](X)_2$ photosensitizers and assessing their ability to promote the aforementioned [2+2] photocycloaddition in the absence of any additive, we could acquire the first evidence of counterion effects in TTEnT photocatalysis. The results of the initial screening of a series of Ru(II) photosensitizers for the intermolecular [2+2] cycloaddition between **1a** and **2a** are summarized in Table 1. Under the standard conditions, using 2.5 mol% of $[Ru(bpy)_3](X)_2$ as sensitizer under blue light irradiation, the rates of the photocycloaddition performed in either acetonitrile or dichloromethane as solvent were compared after 2 h of reaction. $[Ru(bpy)_3]Cl_2$ proved incompetent as a photosensitizer to promote the desired transformation, affording a very low 5% yield of the cyclobutane products in acetonitrile and displaying a complete lack of catalytic activity in dichloromethane; and was therefore left out for the rest of the experimental study.[13b,14]

On the other hand, using *p*-toluenesulfonate (TsO^-), trifluoromethanesulfonate (TfO^-), hexafluorophosphate (PF_6^-) or $BAr^F{}_4^-$ anions proved beneficial, and similar yields (~30%) of the desired cyclobutane products were obtained in acetonitrile after 2h. More interestingly, a striking difference in reaction rates was observed in dichloromethane depending on the counterion identity. While up to 43% yield of the [2+2]-cycloaddition products was obtained after 2h of irradiation using the $[Ru(bpy)_3](BAr^F{}_4)_2$ photocatalyst, the reaction proceeded to 26% yield with $[Ru(bpy)_3](PF_6)_2$, 22% with $[Ru(bpy)_3](OTf)_2$ and only 8% yield with $[Ru(bpy)_3](OTs)_2$.

Table 1. Initial screening of counterion effects on EnT[a]



X =	Cl	TsO	TfO	PF_6	$BAr^F{}_4$
Yield[b] (in CH_3CN)	5	31	30	30	29
Yield[b] (in CH_2Cl_2)	nd[c]	8	22	26	43

[a]Reaction conditions: **1a** (0.1 mmol), **2a** (1 mmol), $[Ru(bpy)_3]X_2$ ($2.5.10^{-3}$ mmol), degassed solvent (2 mL), blue LED irradiation (λ_{max} = 460 nm) for 2 hours under Ar. [b]Yield in % was determined by ^1H NMR using 1,3,5-trimethylbenzene as an internal standard. [c]Not detected by ^1H NMR.

In order to gain a better understanding of the parameters influencing the catalytic performances of this series of $[Ru(bpy)_3](X)_2$ photosensitizers, their optical properties were recorded in both acetonitrile and dichloromethane solutions. In acetonitrile, characterized by relatively high dielectric constant (ε_r ~37) impeding ion association, the properties of the ground state and $[Ru(bpy)_3]^{2+}$* excited state of solvated $[Ru(bpy)_3](X)_2$ salts should not be significantly affected by the nature of the counterion. Indeed, all complexes exhibited a similar metal-to-ligand charge-transfer (MLCT) absorption band at λ_{max} = 451 nm with similar molar extinction coefficients (1500 $M.cm^{-1}$)[15], and demonstrated superimposable emission spectra with λ_{em} = 605 nm (Figure 2A). In dichloromethane (ε_r ~ 9), on the other hand, while the impact of counterion identity on absorption spectra was modest with only a slight increase of molar extinction coefficients from the more coordinating TsO^- anion to the least coordinating $BAr^F{}_4^-$,[16] differences were more significant in emission spectroscopy (Figure 2B). [17] First, the emission of the TsO^- complex is characterized by a maximum at 604 nm/2.053 eV, then the TfO^- and PF_6^- complexes showed a slight hypsochromic shift (λ_{em} = 598 nm/2.073 eV and 596 nm/2.080 eV, respectively) and finally the complex with the largest and least coordinating $BAr^F{}_4^-$ ions exhibited the most blue-shifted emission at λ_{em} = 575 nm/2.156 eV, up by

29 nm/0.103 eV. This relative ordering was satisfactorily reproduced by DFT ΔSCF calculations (Table S23). In dichloromethane of low dielectric constant, the emission energy of ion pairs thus seems to correlate with the charge density of the counterions. In fact, in the MLCT state, the transferred electron is localized on a single bipyridine ligand inducing a dipole moment in the $[(bpy^{\bullet-})Ru^{III}(bpy)_2]^{2+*}$ excited state of C_2 symmetry,[18] which can be stabilized by anion-dipole interaction with the paired anions positioned away from the formally reduced ligand (Figure 2C). The extent of this stabilization is inversely correlated to the counterion size.[19] The blue shifted emission of $[Ru(bpy)_3](BAr^F{}_4)_2$ reflects its increased ^{3}MLCT energy, which is 2.4 kcal/mol higher than the triplet energy of $[Ru(bpy)_3](TsO)_2$, that could in part account for the improved performance of the $BAr^F{}_4^-$ photocatalyst by increasing the driving force for energy transfer from its triplet state (E_T = 49.7 kcal/mol)[20] to the acyl-imidazole substrate **1a** (E_T = 47.6 kcal/mol).[13b] The GS-^{3}MLCT gaps for $[Ru(bpy)_3](X)_2$ complexes have been computed as the difference in their electronic energies at their respective equilibrium geometries in dichloromethane, as reported in Table S22. They follow a trend that perfectly reflects the experimental observations: the gap goes from 46.6 kcal/mol for $[Ru(bpy)_3](BAr^F{}_4)_2$ to 45.4 kcal/mol for $[Ru(bpy)_3](PF_6)_2$, 44.4 kcal/mol for $[Ru(bpy)_3](TfO)_2$ and 44.1 kcal/mol for $[Ru(bpy)_3](TsO)_2$. These values correspond to the electrostatically favorable adjacent arrangement of the counterions, *i.e.* counterions in neighbouring quadrants and remote from the formally reduced bpy ligand, as opposed to an unfavorable opposite arrangement where counterions would be diametrically opposed with respect to the metal centre and with one counterion in the immediate vicinity of the reduced bpy ligand ($\Delta E_{adj/opp}$ = 0.6–3.7 kcal/mol) (Figure 1D). Noteworthy is the ^{3}MLCT state of $[Ru(bpy)_3](BAr^F{}_4)_2$ with an adjacent arrangement of the counterions that is slightly different from the others. Due to their bulkiness and specific geometry, the $BAr^F{}_4^-$ counterions are pinching both neutral bpy ligands in a clip-like configuration (Figure 1E & SI for related discussions and additional visual representations).

To gain further insight into the influence of counterion on energy transfer performance, we conducted Stern-Volmer experiments of intermolecular deactivation of the triplet excited states of ruthenium complexes by an increasing concentration of substrate **1a** as quencher.[21] First, in agreement with the very similar catalytic activities observed in acetonitrile (*vide supra,* Table 1), all excited complexes demonstrated the same ability to be quenched by **1a** with a Stern-Volmer rate constant (K_{sv}) of ~8.1 M^{-1} (Figure 3A). In sharp contrast, Stern-Volmer plots measured in dichloromethane evidenced large differences depending on the identity of the counterions (Figure 3B). The quenching rate of the photoexcited catalysts by the substrate increased by a factor of twenty-five from the TsO^- to the $BAr^F{}_4^-$ counterion, also in line with the markedly superior

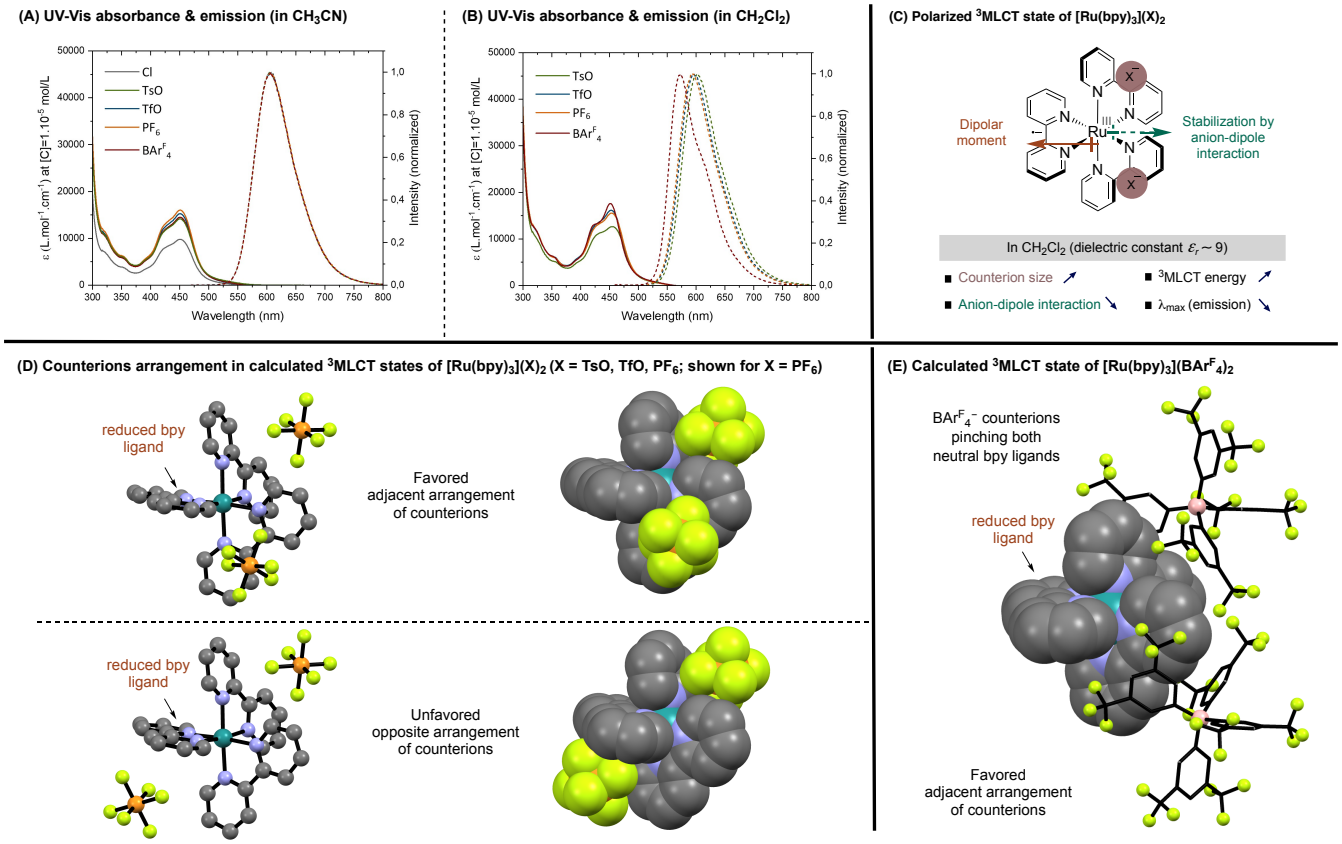


Figure 2. Counterion effect on absorption (solid lines) and emission (dashed lines, λ_{ex} = 450 nm) properties of $[Ru(bpy)_3](X)_2$ in acetonitrile (A) and in dichloromethane (B) at 25 °C. (C) ^{3}MLCT state of $[Ru(bpy)_3](X)_2$. (D) Opposite and adjacent arrangements of counterions in calculated ^{3}MLCT states of $[Ru(bpy)_3](X)_2$ shown for X = PF_6 in ball and stick (left) and space-filling (right) models. (E) Configuration of counterions in calculated ^{3}MLCT state of $[Ru(bpy)_3](BAr^F{}_4)_2$.

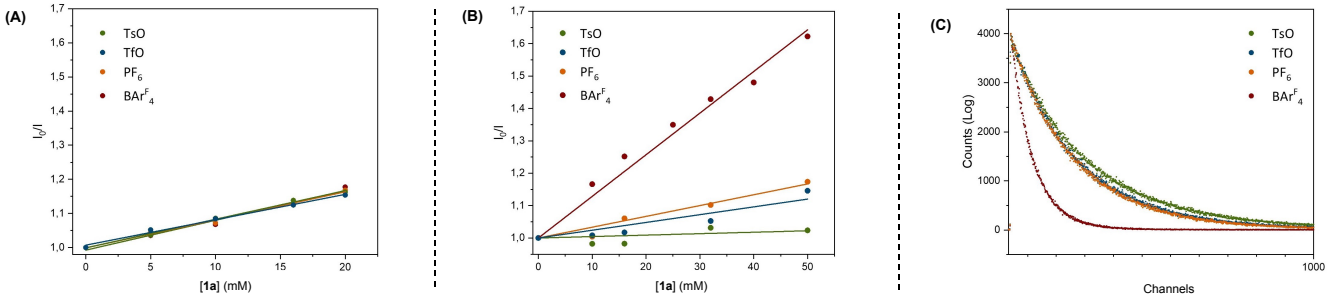


Figure 3. Stern-Volmer plots for excited state quenching of $[Ru(bpy)_3]X_2$ by **1a** in acetonitrile (**A**) and in dichloromethane (**B**). Steady state life time of $[Ru(bpy)_3]X_2$ in CH_2Cl_2 (**C**).

Table 2. Counterion dependence of the optical properties and quenching constants of $[Ru(bpy)_3]X_2$ in dichloromethane[a]

X^-	ε ($M^{-1}.cm^{-1}$)	λ_{em} (nm)	K_{sv} (M^{-1})	τ_0 (ns)[b]	k_q (10^6 M^{-1} s^{-1})[c]	k_r (10^4 s^{-1})[d]	k_{nr} (10^6 s^{-1})[e]	Φ_{lum}
TsO^-	12450	604	0.45	804	0.55	8.71	1.16	0.070
TfO^-	15340	598	2.40	694	3.46	7.92	1.36	0.055
PF_6^-	16020	596	3.34	675	4.95	7.41	1.41	0.050
$BArF^-$	17530	575	11.8	231	51.08	9.96	4.23	0.023

[a]Photophysical data were obtained in dry, deaerated CH_2Cl_2 at 293 K. ε is the molar extinction coefficient at the lowest-energy absorption band maximum, and λ_{em} is the wavelength of the emission band maximum.[b]λ_{exc} = 455 nm. [c]$k_q = K_{sv}/\tau_0$. [d]$k_r = \Phi_{lum}/\tau_0$. [e]$k_{nr} = (1 - \Phi_{lum})/\tau_0$.

photocatalytic performance of the latter. Then, their ^{3}MLCT excited state lifetimes (τ_o) were measured in dichloromethane and once again a significant influence of the counterions was established. The $BAr^F{}_4^-$ complex evidenced the shortest excited state lifetime (τ_o = 231 ns), which is almost four times shorter than that of the TsO^- complex (τ_o = 804 ns). This behavior is usually rationalized by a reduced energy gap between the ^{3}MLCT excited state and low-lying metal-centered (^{3}MC) excited states of $[Ru(bpy)_3]^{2+}$, which facilitates the ^{3}MLCT-^{3}MC internal conversion enabling more efficient nonradiative deactivation pathways,[22,23] as reflected in the decreased luminescence quantum yield (Φ_{lum}, Table 2). The better performance of $[Ru(bpy)_3](BAr^F{}_4)_2$ photocatalyst is all the more remarkable as its excited state lifetime is significantly reduced, which decreases the probability of encountering the substrate for productive energy transfer. The actual quenching rate coefficient (k_q) for this series of $[Ru(bpy)_3](X)_2$ photosensitizers could be accessed thanks to the Stern-Volmer relationship ($K_{sv} = k_q\ \tau_o$) and remarkably, a decrease by two orders of magnitude between the large and non-coordinating $BAr^F{}_4^-$ and the small coordinating TsO^- was determined, highlighting the impressive superiority of $[Ru(bpy)_3](BAr^F{}_4)_2$ to efficiently transfer its triplet energy to the *α,β*-unsaturated 2-acyl imidazole substrate **1a**. This reactivity could be explained by the higher tendency of the bulky $BAr^F{}_4^-$ counterions to dissociate from their cation to form solvent-separated ion pairs,[9b,24] thus unshielding the $[Ru(bpy)_3]^{2+}$ cation in its triplet-excited state and facilitating collisions with the substrate for effective triplet-triplet energy transfer.

Another parameter that plays an important role in photocatalysis, is the photostability of catalysts.[22] Nevertheless, $[Ru(bpy)_3]^{2+}$ complexes are recognized for their relatively low photostability in non-polar solvents such as dichloromethane. In fact, photodecomposition of $[Ru(bpy)_3]^{2+}$ is known to operate through photodissociation of a bipyridine (bpy) ligand, a phenomenon that becomes significant in solvent of low dielectric constant with strongly coordinating anions that can substitute the bpy ligand leading to photocatalytically inactive $Ru(bpy)_2X_2$ complexes such as $Ru(bpy)_2Cl_2$.[14] Therefore, the photostability of our series of $[Ru(bpy)_3]X_2$ complexes was investigated by monitoring the luminescence intensity of CH_2Cl_2 solutions irradiated with blue LEDs (λ_{max} = 460 nm). The decrease of luminescence intensity (I/I_0) as a function of irradiation time is presented in Figure 4. As expected, $[Ru(bpy)_3](TsO)_2$ with the most coordinating counterion in the series evidenced the lowest photostability, leading to almost complete decomposition (>85%) after only 10 minutes of irradiation. A significantly lower decomposition rate was measured for $[Ru(bpy)_3](TfO)_2$ and $[Ru(bpy)_3](PF_6)_2$, which maintained 25% and 40% of their initial luminescence after 1h of irradiation, respectively. Interestingly, $[Ru(bpy)_3](BAr^F{}_4)_2$ exhibited the highest photostability being roughly twice as robust as $[Ru(bpy)_3](PF_6)_2$. The superior photostability of $[Ru(bpy)_3](BAr^F{}_4)_2$ compared to $[Ru(bpy)_3](PF_6)_2$, both involving non-coordinating ions, is however remarkable given its increased ^{3}MLCT state energy. DFT calculations were performed to optimize the lowest MLCT and MC states for the PF_6 and $BAr^F{}_4$ complexes (and the bare cation as a reference, SI, since all our previous calculations on $[Ru(bpy)_3]^{2+}$ were conducted using MeCN as implicit solvent,).[25] Indeed, several

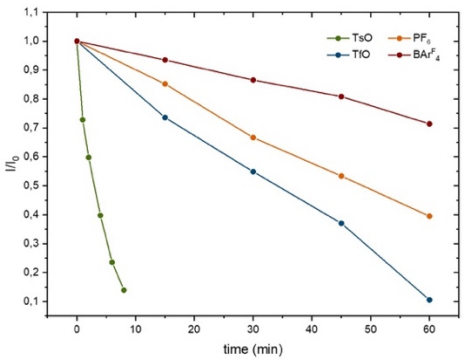


Figure 4. Photostability of $[Ru(bpy)_3]X_2$ complexes in dichloromethane (8 mM) at 25 °C under 50W blue LED (λ_{max} = 460 nm) irradiation.

types of MC states may be invoked, whose electronic structures differ in the nature of the populated d_{σ^*} orbital. As a global picture, when populating a d_{z^2}-like orbital, two major Ru–N elongations will occur involving two pyridines from *separate* ligands in tris(bidentate) complexes such as $Ru(bpy)_3^{2+}$, leading to what we termed MC_{trans} states. On the contrary, if a $d_{x^2-y^2}$-like orbital is populated, three or four major elongations will occur, two of which involving *the same* ligand, leading to MC_{cis} states. For this reason, photoinstability issues due to ligand loss have been suggested to involve preferably $d_{x^2-y^2}$-based MC_{cis} states, while nonradiative deactivation has been suggested to proceed preferably through d_{z^2}-based MC_{trans} states (Figure 5). The calculated Jablonski diagram (Figure 6) shows that MC_{cis} states (red dots) are higher in energy and better separated from MC_{trans} states (green dashes) for the BAr^F_4 complex compared to the PF_6 analogue, in line with the higher photostability of the former. Besides, since the ^{3}MLCT lifetime is long enough to allow relaxation of the anions,[9b] the globally lowest state, labelled as ^{3}MLCT-adj and presenting adjacent anions, is the one which should be considered as the most populated. The gap between ^{3}MLCT-adj and $^3MC_{trans}$-adj is smaller for $[Ru(bpy)_3](BAr^F_4)_2$ (5.3 kcal/mol) than for $[Ru(bpy)_3](PF_6)_2$ (7.0 kcal/mol), in agreement with the shorter MLCT lifetime of the BArF complex (*vide supra* Figure 3C & Table 2).

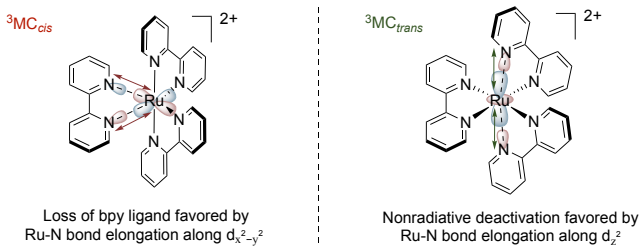


Figure 5: Schematic representation of ^{3}MC states of $[Ru(bpy)_3]^{2+}$ with elongation along either $d_{x^2-y^2}$ ($^3MC_{cis}$, left) or d_{z^2} ($^3MC_{trans}$, right).

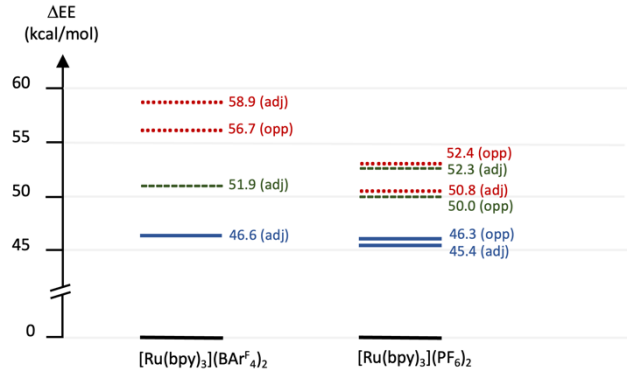
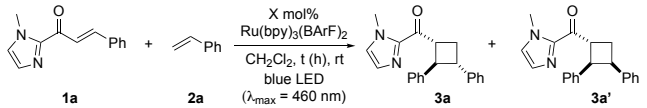


Figure 6: Jablonski diagram schematizing the relative energies of optimized GS (black), ^{3}MLCT (blue lines), $^3MC_{cis}$ (red dots) and $^3MC_{trans}$ (green dashes) states for $[Ru(bpy)_3](BAr^F_4)_2$ and $[Ru(bpy)_3](PF_6)_2$ complexes. Multiple states of the same color indicate various anion arrangements, such as adjacent (adj) and opposite (opp) (see text and SI).

After demonstrating the high photostability of the BAr^F_4 catalyst and its excellent ability to transfer its triplet energy to substrate **1a** in dichloromethane, we optimized the conditions for the [2+2] photocycloaddition reaction in the presence of styrene **2a** as a partner. First, using 2.5 mol% of $[Ru(bpy)_3](BAr^F_4)_2$ as photocatalyst and by increasing the reaction concentration from 0.05 to 0.1 M, the rate of the cycloaddition after 2h of irradiation under blue LEDs (λ_{max} = 460 nm) improved to reach 56% yield of the desired product (Table 3, entries 1 & 2). Interestingly, a decrease of the photocatalyst loading from 2.5 mol% to 1 mol% (Table 3, entry 3) and 0.5 mol% (Table 3, entry 4) did not significantly affect the productivity since very similar results were obtained after 2h of reaction, allowing to reach a turnover number (TON) above 100 in the latter case. The optimized conditions used a low photocatalyst loading of 0.5 mol% and a reaction concentration of 0.2M (Table 3, entry 6), which allowed full conversion within 16h at room temperature, and 72% yield of the desired cycloaddition products (Table 3, entry 6). Finally, by further decreasing its catalytic loading and increasing the reaction concentration, we were able to highlight the outstanding catalytic performance of the BAr^F_4 photosensitizer, which achieved an impressive TON of up to 1040 at a very low loading of 500 ppm (Table 3, entry 7, and SI).

Table 3. Optimization using $[Ru(bpy)_3](BAr^F_4)_2$ as catalyst[a]

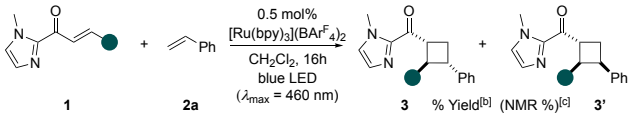


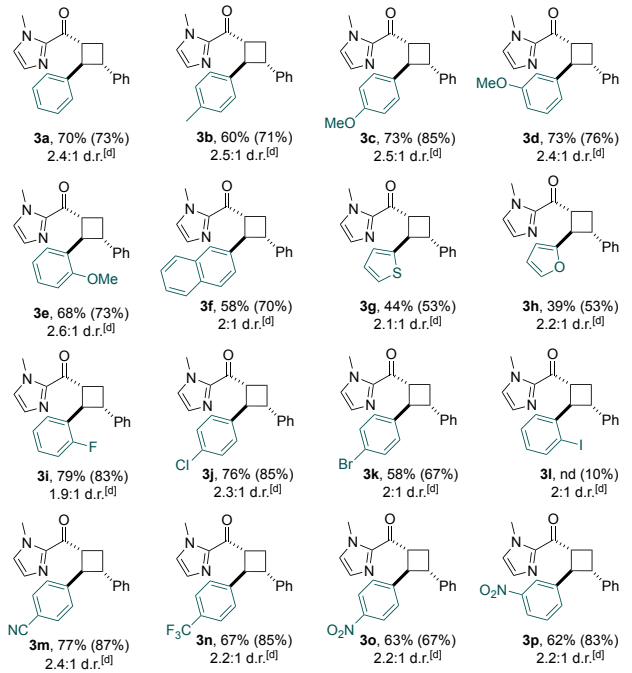
entry	$[Ru(bpy)_3](BAr^F_4)_2$ (mol %)	[**1a**] (mol.L^{-1})	Time (h)	Yield (%)[b]	TON[c]
1	2.5	0.05	2	43	17
2	2.5	0.1	2	56	22
3	1	0.1	2	56	54
4	0.5	0.1	2	54	108
5	0.5	0.2	2	70	140
6	0.5	0.2	16	72	144
7	0.05	1.0	16	52	1040

[a]Reaction conditions: **1a** (0.1 mmol), **2a** (1 mmol), $[Ru(bpy)_3](BAr^F_4)_2$ (0.05 to 2.5x10^{-3} mmol), degassed CH_2Cl_2 (0.25 to 2 mL), blue LED irradiation (λ_{max} = 460 nm) for 2 to 16 hours under Ar. [b]Yield in % of (**3a**+**3a'**) was determined by ^{1}H NMR using 1,3,5-trimethylbenzene as an internal standard.[c]Turnover number.

With the optimized conditions in hand, we then investigated the scope of the intermolecular [2+2] cycloaddition reaction by first varying the electronic and steric properties of the *β*-aryl substituent of the *α*,*β*-unsaturated 2-acyl *N*-methylimidazole **1** (Table 4). Aryl groups with an electron donating substituent were well tolerated affording the desired products (**3b**-**3e**) in good isolated yields. A slight decrease in yield was nevertheless observed when the steric hindrance increased from a methoxy substituent in the *para*-position (**3c**) to a *meta*-(**3d**) and *ortho*-position (**3e**). Substrate **1f** bearing the bulky 2-naphthyl group also efficiently engaged in the [2+2] cycloaddition with styrene to afford the product **3f** in 58% isolated yield. Thienyl and furanyl groups were well tolerated, allowing the formation of products **3g** and **3h** in reasonable yields. With the exception of the *o*-iodophenyl group (**3l**) which induced a very sluggish reaction, substrates bearing an electron-deficient fluoro-, chloro-, bromo-, cyano-, trifluoromethyl, or nitro-aryl group (**3i**-**3p**) were all amenable to efficient cycloaddition catalyzed by the $[Ru(bpy)_3](BAr^F_4)_2$ photocatalyst. We then investigated

Table 4. Scope of *α,β*-unsaturated 2-acyl *N*-methylimidazole[a]





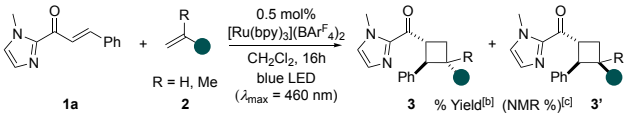
[a]Reaction conditions: **1** (0.2 mmol), **2a** (2 mmol), $[Ru(bpy)_3](BAr^F_4)_2$ ($1x10^{-3}$ mmol), degassed CH_2Cl_2 (1 mL), blue LED irradiation (λ_{max} = 460 nm) for 16 hours under Ar. [b]Isolated yield of **3** and **3'**. [c]Yield in % of **3** and **3'** determined by ^1H NMR using 1,3,5-trimethylbenzene as an internal standard. [d]Diastereomeric ratio (**3**:**3'**) determined by ^1H NMR analysis.

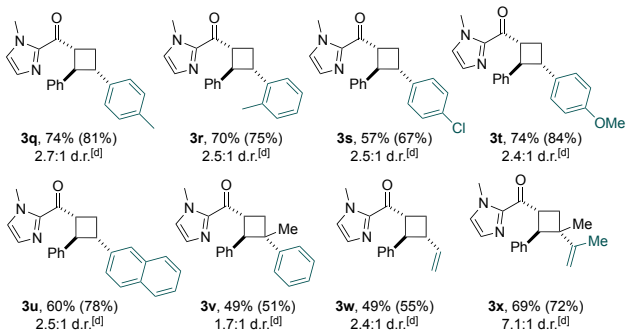
the scope of olefin partner and the results are shown in Table 5. Aryl-substituted electron- rich and electron poor styrenes, including sterically congested (**2r**), were valuable substrates to afford the desired products (**3q**-**3u**) in good yields. Gratifyingly, the presence of a methyl group at the α-position of styrene (**2v**) did not hamper the reactivity. Butadiene and 2,3-dimethylbutadiene are also compatible with the [2+2] cycloaddition, allowing the synthesis of cyclobutanes **3w** and **3x** in good yields and with improved 7:1 selectivity in favor of the trans-trans diastereomer in the latter case. Moreover, the method is not limited to *α,β*-unsaturated 2-acyl *N*-methylimidazole **1** since the *N*-methylimidazole moiety can be replaced by other heteroaromatic groups, such as *N*-phenylimidazole (**4a**), or pyridine (**4b**) with no significant impact on the catalytic performance as demonstrated by the isolation of produts (**5a**) and (**5b**) in 72% and 63% yields, respectively (Table 6). Interestingly, energy transfer from the triplet state energy of $[Ru(bpy)_3](BAr^F_4)_2$ to chalcone is also operative in CH_2Cl_2, allowing cycloaddition with styrene to produce the corresponding intermolecular cycloadduct **5c** in a reasonable 44% isolated yield.

CONCLUSION

In this work, we reveal counterion effects in triplet-triplet energy transfer processes catalyzed by a cationic transition-metal based photosensitizer in low polarity solvent. We observe

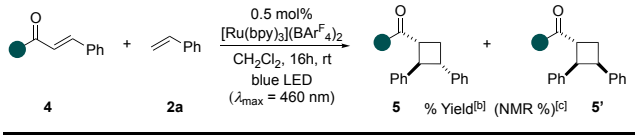
Table 5. Scope of styrene partner[a]

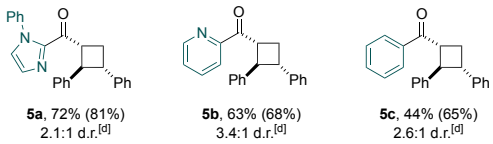




[a]Reaction conditions: **1a** (0.2 mmol), **2** (2 mmol), $[Ru(bpy)_3](BAr^F_4)_2$ ($1x10^{-3}$ mmol), degassed CH_2Cl_2 (1 mL), blue LED irradiation (λ_{max} = 460 nm) for 16 hours under Ar. [b]Isolated yield of **3** and **3'**. [c]Yield in % of **3** and **3'** determined by ^1H NMR using 1,3,5-trimethylbenzene as an internal standard. [d]Diastereomeric ratio (**3**:**3'**) determined by ^1H NMR analysis.

Table 6. Scope of *α,β*-unsaturated ketones[a]





[a]Reaction conditions: **1** (0.2 mmol), **2a** (2 mmol), $[Ru(bpy)_3](BAr^F_4)_2$ ($1x10^{-3}$ mmol), degassed CH_2Cl_2 (1 mL), blue LED irradiation (λ_{max} = 460 nm) for 16 hours under Ar. [b]Isolated yield of **5** and **5'**. [c]Yield in % of **5** and **5'** determined by ^1H NMR using 1,3,5-trimethylbenzene as an internal standard. [d]Diastereomeric ratio (**5**:**5'**) determined by ^1H NMR analysis.

a strong impact of the counterion identity on the excited state properties of $[Ru(bpy)_3](X)_2$ photocatalysts and on their capacity to transfer triplet energy to cinnamoyl-type substrates. The limited stabilization by anion-dipole interaction of the $[Ru(bpy)_3]^{2+*}$ excited state in the presence of large $BArF^-$ anions in dichloromethane results in a ^{3}MLCT excited state of reduced lifetime but of higher energy, offering an increased driving force for the energy transfer process. In comparison, the efficiency of the energy transfer drops by two orders of magnitude in the presence of small anions like TsO^-. The non-coordinating $BAr^F_4{}^-$ anion is also beneficial for the photostability of the cationic photocatalyst by disfavoring the loss of a bpy ligand, which is a predominant decomposition pathway in solvents of low dielectric constant such as dichloromethane in the presence of coordinating/nucleophilic anions. This modulation is due to the fact that $BAr^F_4{}^-$ anions favor the population of non-dissociative $^3MC_{trans}$ states over dissociative $^3MC_{cis}$ states. Thanks to the excellent photostability and high efficiency in triplet-triplet energy transfer of the $[Ru(bpy)_3](BAr^F_4)_2$ photocatalyst in dichloromethane, excellent

photocatalytic activities (up to >1000 TON) could be achieved in intermolecular [2+2] cycloaddition reactions of broad scope with photocatalyst loading as low as 500 ppm. These results clearly demonstrate that the nature of counterion makes a major contribution to cationic photocatalyst potency, a parameter so far overlooked in the literature regarding photocatalyzed transformations and especially reactions enabled by triplet energy transfer.

AUTHOR INFORMATION

Corresponding Authors

Isabelle M. Dixon – *LCPQ, Université de Toulouse, CNRS, Université Toulouse III - Paul Sabatier, 118 route de Narbonne, F-31062 Toulouse, France; Email: dixon@irsamc.ups-tlse.fr*

Yves Canac – *LCC-CNRS, Université de Toulouse, CNRS, UPS, Toulouse, France; Email: yves.canac@lcc-toulouse.fr*

Michael Smietana – *Institut des Biomolécules Max Mousseron, Montpellier, Université de Montpellier, CNRS, ENSCM, France; Email: michael.smietana@umontpellier.fr*

Olivier Baslé – *LCC-CNRS, Université de Toulouse, CNRS, UPS, Toulouse, France; Email: olivier.basle@lcc-toulouse.fr*

Authors

Juliette Zanzi – *LCC-CNRS, Université de Toulouse, CNRS, UPS, Toulouse, France*

Zachary Pastorel – *Institut des Biomolécules Max Mousseron, Montpellier, Université de Montpellier, CNRS, ENSCM, France*

Carine Duhayon – *LCC-CNRS, Université de Toulouse, CNRS, UPS, Toulouse, France*

Elise Lognon – *ITODYS, Université Paris Cité and CNRS, F-75006 Paris, France*

Christophe Coudret – *SOFTMAT, Université de Toulouse, UPS, Institut de Chimie de Toulouse, FR2599, 118 Route de Narbonne, F-31062 Toulouse, France*

Antonio Monari – *ITODYS, Université Paris Cité and CNRS, F-75006 Paris, France*

Notes

There are no conflicts to declare

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