

Modulating the luminescence of an iridium(III) complex incorporating a di(2-picolyl)anilino-appended bipyridine ligand with Zn^{2+} cations†

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A novel iridium complex incorporating a di(2-picolyl)anilino-appended bipyridine ligand was synthesized and its optical properties studied. The presence of Zn^{2+} ions specifically perturbs the excited state, giving rise to a blue-shifted absorption and emission, and a shorter luminescence lifetime.

Cyclometallated iridium(III) complexes are well known for their rich photochemical and photophysical properties.¹ These complexes, with low-lying triplet excited states and lifetimes of around a microsecond, behave as a promising class of phosphorescent dyes in optoelectronic devices² and as new luminescent sensors for analytes,³ including protons,⁴ ions⁵ and biomolecules.⁶ Among them, cationic N^*N -diimine complexes $[\text{Ir}(N^*C\text{-ppy})_2(N^*N)]^+$ (where Hppy is 2-phenylpyridine and N^*N represents a diimine, such as bpy or phen) are particularly attractive.^{7,8} They are readily synthesized, and the facile functionalization of diimine ligands provides access to diverse emissive states, allowing fine tuning of their optical properties.

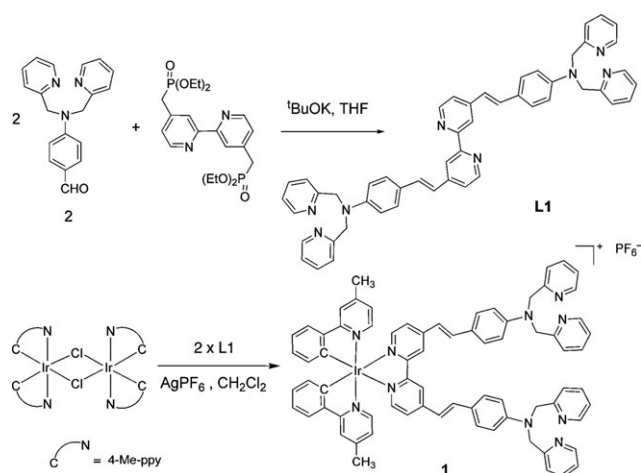
We have previously shown that 2-phenylpyridines incorporating π -conjugated substituents are good building blocks for the construction of luminophores such as neutral, bis-^{9a} and tris-cyclometalated^{9b} 4-(4-(donor)styryl-2-phenylpyridine) iridium(III) and platinum(II)^{9c} complexes, the interesting phosphorescent properties of which are governed by strong triplet intraligand transitions, promoted by an admixture of MLCT character $\{d/\pi \rightarrow \pi^*\}$.

With our ongoing interest in the design of Ir(III) complexes, we have turned our attention to cationic bipyridine complexes incorporating a styryl group substituted with a di(2-picolyl)amino

(DPA) group (Scheme 1). The affinity of the nitrogen-based DPA group towards metal ions, such as Ni^{2+} , Zn^{2+} and Cd^{2+} cations, has previously been demonstrated.^{10,11} We anticipated that attachment of this receptor to an Ir-coordinated bipyridine ligand would allow modulation of the electronic structure upon coordination of metal cations at the DPA site by attenuating the electron donating ability of the amino group; the emission properties should be perturbed as a result.

We report herein the synthesis, characterization and photophysical properties of the Ir complex $[\text{Ir}(N^*C\text{-4-Me-ppy})_2(N^*N\text{-bpy-CH=CH-C}_6\text{H}_4\text{-DPA})][\text{PF}_6]$ (**1**) and the changes to its optical properties upon binding Ni^{2+} , Zn^{2+} and Cd^{2+} . The addition of Zn^{2+} gives a unique response, including a significant emission wavelength shift and a change in luminescence lifetime.

Target bipyridine **L1** was readily prepared in 88% yield by means of a Horner–Wadsworth–Emmons condensation between a bisphosphonate-bipyridine derivative¹² and the appropriate benzaldehyde amine, **2** (Scheme 1). The latter compound was obtained by following a reported procedure for the substitution of aniline by two equivalents of 2-chloromethylpyridine, followed by oxidation with POCl_3 in DMF to introduce the aldehyde functionality.^{10e} The ^1H and ^{13}C NMR spectra of **L1** were in agreement with the proposed structure. Complex **1** was subsequently prepared by the reaction of **L1** with Ir dimer $[\text{Ir}(N^*C\text{-4-Me-ppy})_2(\mu\text{-Cl})_2]$ in CH_2Cl_2 in the presence of AgPF_6 (Scheme 1). It was isolated as orange-red



Scheme 1 Preparation of **L1** and **1**.

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† Electronic supplementary information (ESI) available: Full experimental procedures, photophysical data and UV-vis titrations, details of the X-ray structure of **1**. CCDC reference number 718415. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/b9nj00515c.

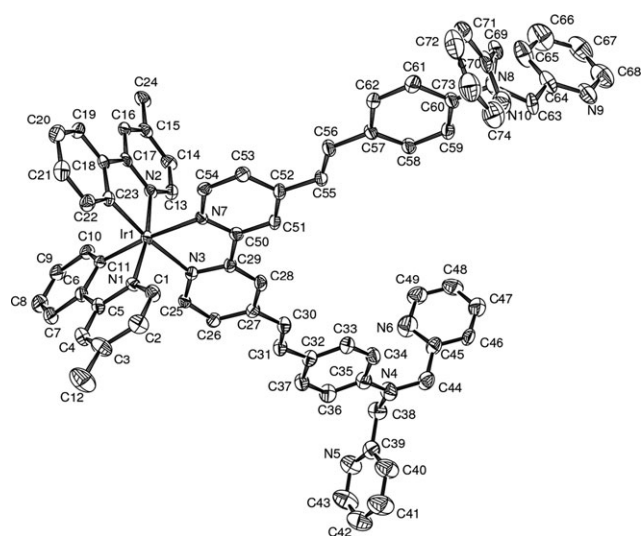


Fig. 1 An ORTEP plot of **1** with thermal ellipsoids at the 50% probability level; $T = 120$ K. Hydrogen atoms are omitted for clarity.

microcrystals in 70% yield after recrystallization from CH_2Cl_2 –pentane; the structure determined by X-ray diffraction is shown in Fig. 1. The initial *E* configuration of the two $\text{C}=\text{C}$ double bonds was retained upon metal complexation. However, upon UV irradiation, the $\text{C}=\text{C}$ bonds of **1** underwent a slow *E*–*Z* isomerization process, as revealed by ^1H NMR spectroscopy.

The electronic absorption and luminescence data for **1** and **L1** are summarized in Table S1.† The absorption spectrum of **1** displays high energy bands in the UV region (250–340 nm) assigned to intraligand IL ($\pi \rightarrow \pi^*$) (bpy and ppy) transitions (Fig. 2). An intense low energy absorption band centered at 449 nm ($\epsilon = 27\,500 \text{ dm}^3 \text{ mol}^{-1} \text{ cm}^{-1}$) probably arises from the influence of the electron donating nature of the amino group. The effect of the amino substituent is also seen in the behavior of the corresponding uncoordinated ligand **L1**, which displays a low energy absorption band ($\lambda_{\text{max}} = 385 \text{ nm}$). The 64 nm red shift in the IL band of **L1** upon complexation to iridium(III) (highlighted in Figure S1†) can be attributed to the enhanced π -acceptor character of the bpy moiety upon coordination, lowering the LUMO energy level.¹³ Based on the spectra of complexes such as $[\text{Ir}(\text{N}^+\text{C-ppy})_2(\text{N}^+\text{N-bpy})]^+$,^{7,8} one or more bands due to MLCT transitions [$d\pi(\text{Ir}) \rightarrow \pi^*(\text{diimine and ppy})$] would be anticipated over the range 330–400 nm; in the present complex, they are superimposed by the tail of the low energy band associated with the aminostyryl unit.

Uncoordinated ligand **L1** is fluorescent in solution at room temperature (RT) ($\lambda_{\text{max}} = 486 \text{ nm}$ in CH_2Cl_2). In contrast, complex **1** is non-emissive in solution at RT. Simpler iridium(III) complexes, such as $[\text{Ir}(\text{N}^+\text{C-ppy})_2(\text{N}^+\text{N-bpy})]^+$, display luminescence from the lowest-lying triplet state under these conditions, promoted by spin–orbit coupling. The absence of RT phosphorescence in the present case suggests the introduction of a non-radiative decay pathway that is not open to simpler derivatives and/or a radiative rate constant, k_r , that is unusually low; this is confirmed by its behavior at 77 K. Under these conditions, the complex emits brightly, displaying a highly structured spectrum deep into the red

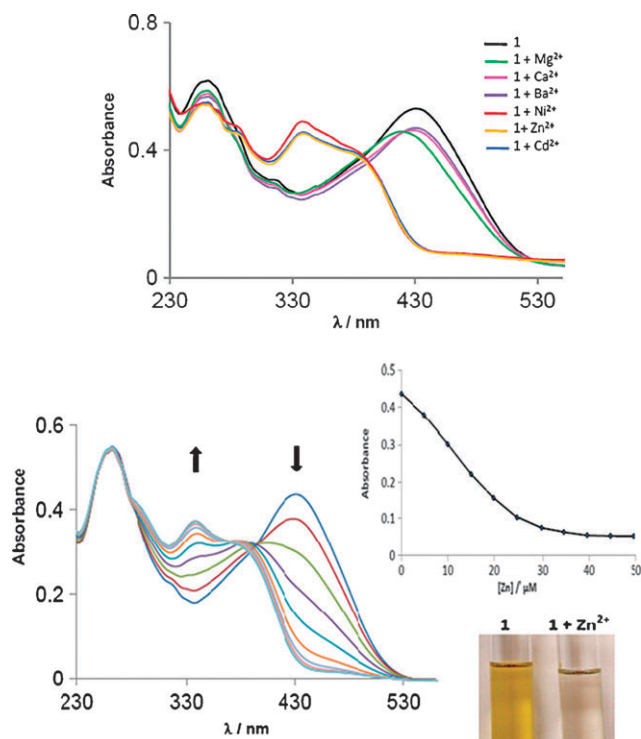


Fig. 2 Top: UV-vis absorption spectral changes of **1** (conc. = $1 \times 10^{-5} \text{ M}$ in CH_3CN) after the addition of 50 equivalents of Ba^{2+} , Ca^{2+} , Mg^{2+} , Cd^{2+} , Zn^{2+} and Ni^{2+} ions. Bottom: UV-vis absorption spectral changes of **1** upon addition of incremental (0.5 equivalent) amounts of $\text{Zn}(\text{ClO}_4)_2$. The inset shows a plot of absorbance against $[\text{Zn}^{2+}]$ monitored at 430 nm.

region ($\lambda_{\text{max}}^{0-0} = 640 \text{ nm}$, $\Delta\nu = 1400 \text{ cm}^{-1}$) (Fig. 3). The lifetime of 67 μs is exceptionally long, confirming that k_r is indeed low. The values are in stark contrast to those of, for example, $[\text{Ir}(\text{N}^+\text{C-ppy})_2(\text{N}^+\text{N-4,4'-Me}_2\text{-bpy})]^+$ ($\lambda_{\text{em}} = 506 \text{ nm}$ at 77 K, $\tau = 4.83 \mu\text{s}$),^{7d} where the emission is attributed to an MLCT state. Evidently, in complex **1**, the emission emanates from a state of ^3IL character localized on the styryl-substituted bpy ligand, with relatively low metal participation; triplet excited states, in which the metal plays only a minor role, are expected to have longer radiative lifetimes owing to less efficient spin–orbit coupling pathways. The triplet radiative rate constant is probably so low that, at RT, the excited state is essentially fully deactivated by the faster *E*–*Z* isomerization process, accounting for the lack of emission. In free ligand **L1**, in contrast, the fluorescence is fast enough to be competitive with this process.

The behavior of **1** in the presence of various metal cations was studied (Ba^{2+} , Ca^{2+} , Mg^{2+} , Ni^{2+} , Zn^{2+} and Cd^{2+} as their perchlorate salts in CH_3CN). The absorption spectrum was dramatically modified upon adding Ni^{2+} , Zn^{2+} and Cd^{2+} (Fig. 2). The solution color visibly changed from orange to colorless, the low energy absorption band ($\lambda_{\text{max}} 430 \text{ nm}$ in CH_3CN) decreased and a new band at $\sim 330 \text{ nm}$ concomitantly evolved (Fig. 2 and Fig. S3†). In contrast, no significant changes were observed with Ba^{2+} , Ca^{2+} and Mg^{2+} .

The blue shift of the absorption band with Ni^{2+} , Zn^{2+} and Cd^{2+} is intuitively consistent with the binding of the metal ion

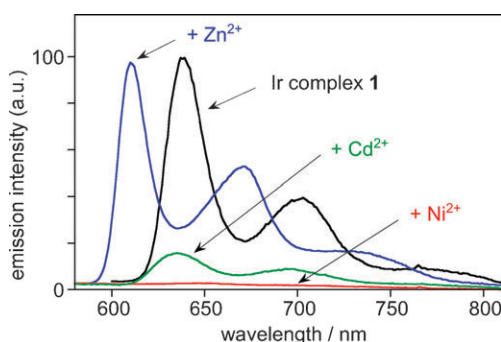


Fig. 3 Luminescence spectral changes (77 K) of **1** (conc. = 3×10^{-6} M) in ethanol–methanol (4 : 1, v/v) upon the addition of $M(\text{ClO}_4)_2$ ($M = \text{Ni}, \text{Zn}, \text{Cd}$) (conc. = 6×10^{-5} M); $\lambda_{\text{ex}} = 385$ nm (isosbestic point).

at the DPA amine nitrogen, which would reduce its donating ability, lowering the energy of the highest-occupied molecular orbital. The stoichiometric ratio of the species formed between complex **1** and M^{2+} was estimated to be 1 : 2 in accordance with the presence of two DPA receptors. An isosbestic point at 385 nm is consistent with a two-species equilibrium ($\mathbf{1} + 2M^{2+} \rightarrow \mathbf{1} \cdot (M^{2+})_2$). The following binding constants, $\log K_s$, were determined: $\text{Ni}^{2+} = 7.01 \pm 0.03$, $\text{Cd}^{2+} = 6.56 \pm 0.61$ and $\text{Zn}^{2+} = 5.46 \pm 0.12$.

The effect of the same set of metal ions on the luminescence of **1** at 77 K was also investigated. The emission ($\lambda_{\text{max}} = 640$ nm) was completely quenched upon adding Ni^{2+} ions, whereas Cd^{2+} lead only to partial quenching and a scarcely perceptible blue shift (3 nm) (Fig. 3). Most strikingly, the coordination of Zn^{2+} ions induced a unique response: a blue-shifted emission ($\lambda_{\text{max}}^{0-0} = 610$ nm) of the intensity comparable to **1** (for excitation at the isosbestic point at 385 nm) with a lifetime of 34 μs was observed, indicative of the formation of a new emissive species. In the presence of a sub-stoichiometric quantity of Zn^{2+} , the emission profiles of the two species—the ion-free complex **1** and the Zn^{2+} -coordinated complex—were independently observable at 77 K (Fig. S5†). Two distinct sets of vibrational bands were clear, and the distinctly different decay curves registered at 610 and 637 nm, respectively, are in line with those of the fully bound and unbound species. Interestingly, the addition of H^+ ions (from trifluoroacetic acid) lead to related but much smaller effects on the emission: a blue shift of 12 nm and a small decrease in the lifetime to 50 μs .

As for the absorption, the emission blue shift can be interpreted in terms of a stabilisation of the highest occupied molecular orbitals as the N lone pair becomes bound to the Zn^{2+} ion. That selectivity for Cd^{2+} is observed only in emission and not in absorption presumably reflects a subtle difference in the electronic distribution in the triplet and singlet excited states. The selective blue shift induced by Zn^{2+} implies that, in the excited state, the more charge-dense Zn^{2+} ion interacts more strongly with the amino group of the DPA unit than does Cd^{2+} . The softer nature of the Cd^{2+} ion probably means that its predominant mode of binding to the DPA in the excited state is *via* the picolyl units.

In summary, complex **1** displays an unusual phenomenon of selective zinc-dependent wavelength- and lifetime-modulation of its emission.

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Experimental

Preparation of **1**

Chloride-bridged dimer $[\text{Ir}(\text{N}^4\text{-C-4-Me-ppy})_2(\mu\text{-Cl})_2]$ (100 mg, 0.09 mmol), bipyridine derivative **L1** (140 mg, 0.19 mmol) and AgPF_6 (60 mg, 0.24 mmol) were mixed in CH_2Cl_2 (10 mL). The reaction mixture was stirred under Ar for 2 h. The solution was then concentrated to dryness and the product extracted with CH_2Cl_2 (3×5 mL). Crystallization from a CH_2Cl_2 –diethyl ether mixture gave an orange-red powder (186 mg, 70%). ^1H NMR (500 MHz, CDCl_3): 8.68 (s, 2H, $\text{H}_3\text{-bpy}$), 8.61 (d, 4H, $\text{H}_6\text{-Py}^*$, $^3J = 4.4$ Hz), 7.70 (s, 2H, $\text{H}_6\text{-bpy}$), 7.65 (m, 2H, $\text{H}_3\text{-Ph}$, 4H, $\text{H}_3\text{-Py}^*$ and 2H, $\text{H}_3\text{-Py}$), 7.48 (d, 4H, C_6H_4 , $^3J = 8.9$ Hz), 7.44 (d, 2H, $\text{H}_6\text{-Py}$, $^3J = 6.0$ Hz), 7.41 (d, 2H, $=\text{CH}_8$, $^3J = 16.2$ Hz), 7.27 (m, 2H, $\text{H}_5\text{-bpy}$), 7.25 (d, 4H, $\text{H}_4\text{-Py}^*$, $^3J = 7.9$ Hz), 7.20 (dd, 4H, $\text{H}_5\text{-Py}^*$, $^3J = 6.8$ Hz, $^3J = 5.03$ Hz), 7.11 (d, 2H, $=\text{CH}_7$, $^3J = 16.2$ Hz), 7.01 (td, 2H, $\text{H}_4\text{-Ph}$, $^3J = 7.5$ Hz, $^4J = 1.0$ Hz), 6.90 (td, 2H, $\text{H}_5\text{-Ph}$, $^3J = 7.5$ Hz, $^4J = 1.2$ Hz), 6.84 (dd, 2H, $\text{H}_5\text{-Py}$, $^3J = 6.0$ Hz, $^4J = 1.4$ Hz), 6.69 (d, 4H, C_6H_4 , $^3J = 8.8$ Hz), 6.34 (dd, 2H, $\text{H}_6\text{-Ph}$, $^3J = 7.5$ Hz, $^4J = 0.8$ Hz), 4.86 (s, 8H, CH_2) and 2.49 (s, 6H, CH_3). $^{13}\text{C}\{^1\text{H}\}$ NMR (75 MHz, CDCl_3): 167.23 ($\text{C}_2\text{-py}$), 158.13 ($\text{C}_2\text{-py}^*$), 156.05 ($\text{C}_2\text{-bpy}$), 151.67 ($\text{C}_1\text{-Ph}$), 149.74 ($\text{C}_6\text{-py}^*$), 149.53 ($\text{C}_4\text{-py}$), 149.38 (C_9), 149.13 ($\text{C}_4\text{-bpy}$), 149.05 ($\text{C}_6\text{-bpy}$), 147.90 ($\text{C}_6\text{-py}$), 143.64 ($\text{C}_2\text{-Ph}$), 137.19 (C_8), 136.89 ($\text{C}_4\text{-py}^*$), 131.84 ($\text{C}_6\text{-Ph}$), 130.32 ($\text{C}_5\text{-Ph}$), 129.65 (C_{10}), 124.90 (C_{12}), 124.36 ($\text{C}_3\text{-Ph}$), 124.26 ($\text{C}_5\text{-py}$), 122.44 ($\text{C}_5\text{-bpy}$), 122.17 ($\text{C}_5\text{-py}^*$ and $\text{C}_3\text{-bpy}$), 122.08 ($\text{C}_4\text{-Ph}$), 120.80 ($\text{C}_3\text{-py}^*$), 120.06 ($\text{C}_3\text{-py}$), 119.70 (C_7), 112.54 (C_{11}), 56.98 (CH_2) and 21.29 (CH_3). Elemental analysis: $\text{C}_{74}\text{H}_{62}\text{F}_6\text{N}_{10}\text{IrP} \cdot 0.5\text{CH}_2\text{Cl}_2$ calc. C, 60.83, H, 4.32, N, 9.52. Found: C, 60.20, H, 4.36, N, 9.18%. HRMS: m/z calc. for $\text{C}_{74}\text{H}_{62}\text{N}_{10}\text{Ir}$: 1283.47883, found: 1283.4793.

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