Colour tuning by the ring roundabout: [Ir(C^N)₂(N^N)]⁺ emitters with sulfonyl-substituted cyclometallating ligands†

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A series of cationic bis-cyclometallated iridium(III) complexes [Ir(C^N)₂(N^N)]⁺ is reported. Cyclometallating C^N ligands are based on 2-phenylpyridine with electron-withdrawing sulfonyl substituents in the phenyl ring: 2-(4-methylsulfonylphenyl)pyridine (H2pzpy) and 2-(3-methylsulfonyl-1H-pyrazol-1-yl)pyridine (dmmpzpy) are used as electron-rich ancillary N^N ligands. The complexes have been fully characterized and the single crystal structure of [Ir2(dmmpzpy)][PF6]·MeCN has been determined. Depending on the position of the methylsulfonyl group, the complexes are green or blue emitters with vibrationally structured emission maxima at 491, 523 nm for [Ir2(Hpzpy)][PF6] or 463, 493 nm for [Ir2(N^N)[PF6] in MeCN solution. The marked vibrational structure and the absence of a rigidochromic shift, together with theoretical predictions based on density functional theory calculations, confirm the 3LC nature of the emitting triplet state. All four complexes have relatively high photoluminescence quantum yields in de-aerated solution (53 to 77%). On going from solution to powder samples, the emission is red-shifted and the quantum yields are considerably lower (~11%). The complexes were tested in light-emitting electrochemical cells (LECs) achieving maximum luminances of 141 cd m⁻² when operated at 100 A m⁻² using pulsed current driving conditions.

Introduction

Cationic iridium(III) complexes of the type [Ir(C^N)₂(N^N)]⁺, in which C^N is a cyclometallated ligand and N^N is a diimine ligand, are well-established emissive materials in light-emitting electrochemical cells (LECs). Prototype LECs incorporated conjugated polymers in the active layer, but the development of LECs with emissive ionic transition-metal complexes (iTMCs) has led to single-component devices. The application of a bias across the device causes the ions in the active layer to migrate towards the electrodes where their accumulation results in a drop in potential near the electrode surface. The formation of ionic junctions lowers the barrier for electron and hole injection and, as a consequence, the device is independent of the work function of the electrode material. Hence, it is not a prerequisite that LECs use low work-function metals, as is necessary in organic light-emitting diodes (OLEDs), and air-stable electrodes (e.g. Al) are employed in LECs. This has a beneficial consequence: the rigorous exclusion of air, which is essential in the fabrication of OLEDs, is unnecessary in LECs. Another advantage of LECs over OLEDs is their far simpler architecture and the fact that they can be assembled using solution processing.

Despite the fabrication advantages of LECs, there is a lack of wide bandgap iTMCs to produce the blue iTMC-LECs needed to combine with orange emitters for white light. In [Ir(C^N)₂(N^N)]⁺ complexes, the HOMO contains contributions from the Ir(III)dπ orbitals and phenyl π-orbitals of the cyclometallated C^N ligands, and the LUMO is usually localized on the N^N ligand. The localized character of the HOMO–LUMO manifold lends itself perfectly to colour tuning the emission of [Ir(C^N)₂(N^N)]⁺ complexes by introducing N^N ligands that destabilize the LUMO and/or C^N ligands that stabilize the HOMO. In 2006, Nazeeruddin et al. showed that [Ir(ppy)₂(4,4’-Me₂N)₂bpy]⁺ (Hppy = 2-phenylpyridine, 4,4’-bis(dimethylamino)-2,2’-bipyridine) was an efficient blue-green emitter, and pointed to the potential use of 1-phenylpyrazole or fluorinated 2-phenylpyridine ligands to stabilize the HOMO of [Ir(C^N)₂(N^N)]⁺. This approach to blue emission was
demonstrated in 2005 by Thompson and coworkers in a landmark paper. The quest for blue emitters continues, with utilization of the structure–property ‘rules’, leading to the replacement of 1-phenylpyrazolyl- by 2-methyl-5-phenyl-2H-tetrazolyl-functionalized C’N ligands. The number and position of fluoro groups in Hppy have only a small influence on the photophysical properties of [Ir(C’N)2(4,4’-Bu2bpy)]+ complexes (4,4’-di-tert-butyl-2,2’-bipyridine) although increasing the number of fluorine atoms leads to shorter-lived LECs. Indeed, the use of fluorinated C’N ligands in iridium(m) complexes in OLEDs has been questioned due to the instability of the Cphenyl–F bond; hence, fluorine-free blue [Ir(C’N)2(N’N)]+ emitters are desirable. It must be mentioned that blue photoluminescence (PL) in solution, thin-film or powder, is no guarantee of blue electroluminescence (EL) under device conditions, and significant shifts in emission maxima on going from PL to EL are common.

Following studies of blue emitting complexes for OLEDs, Evariste et al. demonstrated the use of cyclometallated methyl- and methoxy-substituted 2,3’-bipyridine ligands as the C’N domain in blue-emitting [Ir(C’N)2(4,4’-Bu2bpy)] complexes. An alternative approach is to replace the fluoro-substituents in the C’N ligands by electron-withdrawing sulfone groups, a design strategy previously demonstrated in OLEDs, and one that we have applied in [Ir(C’N)2(N’N)] complexes (Scheme 1).

However, the complexes in Scheme 1 are green emitters and we consider here further modification to achieve a shift in emission to higher energies. An appealing strategy is the use of sulfone-functionalized C’N ligands and derivatives of 1-phenylpyrazole as the N’N ligand, combining the effects of the electron-withdrawing sulfone groups with those of the electron-rich pyrazole ring, leading to larger HOMO–LUMO gaps in [Ir(C’N)2(N’N)] complexes. Stereoisomerism in [Ir(C’N)3] emitters in OLEDs is important for the enhancement of phosphorescence quantum yield and electrochemical stability, with the fac-isomer being superior to the mer-isomer, and we have now investigated the effect of positional isomerization in the sulfone-functionalized ligands as it is known that the emission behaviour of some iridium complexes is sensitive to the positions of the substituents in the C’N ligands. Density functional theory (DFT) calculations have shown that photophysical properties of cationic [Ir(ppy)2(P’P)] complexes vary with the position of functionalization of the [ppy] ligands.

To further blue-shift the emission, we have synthesized a series of complexes combining sulfone-substituted H1 and H2 and pyrazolopyridine ancillary ligands (Scheme 2). The complex [Ir(ppy)2(pzpy)][PF6] previously prepared by He et al. is used as a reference compound for the comparison of the photophysical and electrochemical properties of the new complexes (Scheme 3), which are further analysed through DFT calculations. The electroluminescent properties of the complexes have been investigated in LEC devices operated under pulsed driving conditions.

Scheme 1 Previously reported [Ir(C’N)2(N’N)]+ complexes containing sulfone-functionalized C’N ligands.

Scheme 2 Structures of cyclometallating ligands 2-(4-methyl-sulfonylphenyl)pyridine (H1) and 2-[3-methylsulfonylphenyl]pyridine (H2) and ancillary ligands 2-(1H-pyrazol-1-yl)pyridine (pzpy) and 2-(3,5-dimethyl-1H-pyrazol-1-yl)pyridine (dmzpy).

Scheme 3 Structures of [Ir(ppy)2(pzpy)][PF6] and methylsulfonyl-substituted complexes [Ir(C’N)2(N’N)][PF6] with C’N = [1]− or [2]−, and N’N = 2-(1H-pyrazol-1-yl)pyridine (pzpy) or 2-(3,5-dimethyl-1H-pyrazol-1-yl)pyridine (dmzpy).
Experimental

General experimental, ligand synthesis and IR spectroscopic data for the complexes are given in the ESI†

Synthesis

[Ir(2)2Cl]2. Compound H2 (105 mg, 0.450 mmol) was dissolved in 2-ethoxyethanol (3 mL). The solution was purged with N2, then [Ir(cod)Cl]2 (77.2 mg, 0.115 mmol) was added. The mixture was heated at reflux for 22 h. After letting the mixture cool to room temperature, the precipitate was filtered off, washed with 2-ethoxyethanol, H2O and EtOH and dried under vacuum to yield [Ir(2)2Cl]2 as a yellow powder (131 mg, 0.0946 mmol, 84.0%) which was used for subsequent reactions without further purification. 1H NMR (500 MHz, CDCl3) δ/ppm 9.19 (dd, J = 5.7, 1.5, 0.7 Hz, 4H, Hδ), 8.11 (dd, J = 8.5, 1.4, 0.7 Hz, 4H, Hβ2), 8.05 (d, J = 2.0 Hz, 4H, Hδ), 7.96 (dd, J = 8.1, 1.5, 0.7 Hz, 4H, Hβ3, Hδ), 7.09 (dd, J = 8.3, 2.0 Hz, 4H, Hβ3, Hβ4), 6.98 (dd, J = 7.3, 5.5, 1.4 Hz, 4H, Hβ3, Hβ4), 6.65 (d, J = 8.3 Hz, 4H, Hα), 2.91 (s, 12H, Hδ), 16H, [Ir(C^N)2(MeCN)2]+ (calc. 657.1), 698.1 [Ir(C^N)2(MeCN)]+ (calc. 698.1) [Ir(C^N)]2(MeCN)]+ (calc. 739.1).

General procedure for the synthesis of iridium(III) complexes. The iridium(III) dimer and the ancillary ligand were suspended in MeOH (15 mL) in a vial and heated at 120 °C for 1 h in a microwave reactor (14 bar). The resulting yellow solution was filtered through cotton and concentrated under reduced pressure. The residue was dissolved in MeOH, an excess of solid NH4PF6 was added and the resulting suspension was stirred for 5 min at room temperature. The yellow precipitate was removed by filtration and redissolved in CH2Cl2/CH3CN. The solvent was removed under reduced pressure, the crude product was purified by column chromatography (silica) and the solvent removed again. The residue was dissolved in CH2Cl2 or CH3CN, precipitated with EtO and left in a refrigerator overnight. The resulting precipitate was removed by filtration, washed with MeOH and then EtO and dried under vacuum.

[Ir(1)(pzpy)][PF6]. [Ir(1)Cl]2 (125 mg, 0.0903 mmol) and pzpy (27.0 mg, 0.186 mmol). Purification by column chromatography (silica, CH2Cl2 changing to CHCl3–5% MeOH) and precipitation from a CH3CN solution. [Ir(1)(pzpy)][PF6] was isolated as a yellow solid (169 mg, 0.179 mmol, 99.0%). 1H NMR (500 MHz, CD3CN) δ/ppm 8.70 (d, J = 3.1 Hz, 1H, Hα3), 8.25–8.21 (overlapping m, 2H, Hβ1–2), 8.19 (dd, J = 8.5, 7.6, 1.6 Hz, 1H, Hβ3), 8.10 (dt, J = 8.5, 1.0 Hz, 1H, Hβ3), 8.05–7.97 (overlapping m, 4H, Hα1–2, Hβ1–2, Hβ3, Hδ), 7.85–7.82 (overlapping m, 2H, Hβ), 7.75 (dd, J = 5.5, 1.7, 0.7 Hz, 1H, Hβ3, Hδ), 7.58 (dd, J = 8.2, 1.9 Hz, 1H, Hβ4), 7.56 (dd, J = 8.2, 1.9 Hz, 1H, Hα), 7.36 (dd, J = 7.6, 5.6, 1.1 Hz, 1H, Hβ3), 7.32 (d, J = 2.1 Hz, 1H, Hα), 7.30–7.22 (overlapping m, 2H, Hβ5–6), 6.77 (dd, J = 3.1, 2.1 Hz, 1H, Hβ4), 6.71 (d, J = 1.9 Hz, 1H, Hα3), 6.70 (d, J = 1.9 Hz, 1H, Hα4), 2.88 (s, 3H, Hβ1–2), 2.88 (s, 3H, Hβ1–2). 11C[1H] NMR (126 MHz, CD2CN) δ/ppm 166.2 (C1), 161.6 (C2), 151.2 (C3), 150.9 (C4), 150.5 (C5), 150.4 (C6), 150.39 (C7), 150.38 (C8), 149.4 (C9), 146.8 (C10), 145.0 (C11), 142.8 (C12), 142.2 (C13), 141.9 (C14), 140.5 (C15), 140.4 (C16), 133.3 (C17), 129.8 (C18), 129.7 (C19), 126.34 (C20), 126.32 (C21), 126.21 (C22), 126.17 (C23), 126.0 (C24), 122.7 (C25), 122.54 (C26), 122.51 (C27), 122.48 (C28), 114.5 (C29), 114.3 (C30), 44.2 (C31), 44.2 (C32), 44.1 (C33), 44.1 (C34), 44.1 (C35), 44.1 (C36), 44.1 (C37), 44.1 (C38), 44.1 (C39), 44.1 (C40), 44.1 (C41), 44.1 (C42), 44.1 (C43), 44.1 (C44), 44.1 (C45).
(C Sm), 125.8 (C D), 124.1 (C A), 123.7 (C C), 121.89 (C B/D), 121.86 (C A/D), 114.5 (C E), 112.3 (C F), 44.6 (C A-SO2Me-C4-SO2Me).

UV/Vis (CH3CN, 1.0 \times 10^{-5} \text{mol dm}^{-3}) / \text{nm} (\epsilon/\text{dm}^3 \text{mol}^{-1} \text{cm}^{-1}) = 253 (67000), 280 (44000), 375 sh (50000). Emission (CH3CN, 1.0 \times 10^{-5} \text{mol dm}^{-3}, \lambda_{exc} = 259 \text{ nm}) \lambda_{em}^{max} = 463, 493 \text{ nm}. ESI-MS m/z 805 [M-PF6]\text{]} (calc. 802.4). Found C 39.62, H 3.20, N 7.51; C32H27F4IrN5O4PS2 \text{H2O} requires C 39.15, H 3.20, N 7.51%.

\textbf{[Ir(2)2(dmpzpy)]PF6,} [Ir(2)Cl]2 (143 mg, 0.103 mmol) and dmpzpy (34.6 mg, 0.200 mmol) purification by column chromatography (silica, CH2Cl2 changing to CH2Cl2-MeOH). The sulfone-substituted cyclometallating ligand H3N- SO2Me, 2.83 (overlapping m, 3H, HB5+C5+E5), 2.82 (overlapping s, 3H, HA4-SO2Me), 2.81 (s, 3H, HB4+D4+B6), 2.74 (dd, J = 5.5, 1.6, 0.8 Hz, 1H, HE6), 2.76 (dd, J = 8.1, 2.0 Hz, 1H, HE4), 2.73-2.74 (overlapping m, 3H, H8-SO2Me), 2.72 (dd, J = 7.4, 5.8, 1.4 Hz, 1H, H6\text{SO2Me}), 2.69 (d, J = 8.1 Hz, 1H, H5\text{F5-Me}), 2.63 (d, J = 8.1 Hz, 1H, H4\text{F4-Me}), 2.64 s (1H, H3\text{F3-Me}), 3.03 s (3H, H3\text{F3-Me}).

13C{1H} NMR (126 MHz, CD3CN) /ppm 8.30 \text{ (CD2)}, 166.4 (CD2), 166.0 (CB2), 158.6 (CA4), 158.1 (CZ4), 156.6 (CZ3), 151.2 (CZ2), 151.1 (CB6), 150.7 (CD6), 150.4 (CB4), 147.0 (CZ3), 146.4 (CA4), 146.3 (CA3), 142.5 (CZ4), 140.4 (CB3), 140.2 (CA5), 136.9 (CA3), 136.3 (CA5), 133.6 (CZ6), 133.0 (CA5), 129.1 (CA2), 128.4 (CB5), 126.0 (CB5), 125.7 (D5), 125.1 (CA5), 124.1 (CA5), 123.9 (CA5), 123.8 (CB3), 121.9 (D3), 115.3 (CZ3), 114.2 (CZ4), 44.5 (C4-Ar-SO2Me-C4-SO2Me), 44.5 (C4-Ar-SO2Me-C4-SO2Me), 15.6 (CF3-Me), 13.8 (CF3-Me).

UV/Vis (CH3CN, 1.0 \times 10^{-5} \text{mol dm}^{-3}) / \text{nm} (\epsilon/\text{dm}^3 \text{mol}^{-1} \text{cm}^{-1}) = 254 (66000), 280 sh (41000), 375 sh (43000). Emission (CH3CN, 1.0 \times 10^{-5} \text{mol dm}^{-3}, \lambda_{exc} = 259 \text{ nm}) \lambda_{em}^{max} = 463, 493 \text{ nm}. ESI-MS m/z 830 [M-PF6]\text{]} (calc. 830.1). Found C 41.89, H 3.54, N 7.18; C34H31F6IrN5O4PS2 \text{H2O} requires C 41.89, H 3.20, N 7.18%.

Crystallography

Single crystal data were collected on a Bruker APEX-II diffractometer; data reduction, solution and refinement used SHELXTL. Structure analysis used Mercury v. 3.5.1.38

\textbf{[Ir(2)2(dmpzpy)]PF6,} MeCN. \text{C}34\text{H}28\text{F}4\text{IrN}6\text{O}4\text{PS}2, \text{M} = 1009.97, yellow plate, monoclinic, space group \text{C}2/c, \text{a} = 25.335(3), \text{b} = 12.8383(13), \text{c} = 24.8213(4), \beta = 109.7554(4), \text{U} = 7598.2(8) \text{A}^2, Z = 8, D_x = 1.766 \text{Mg m}^{-3}, \mu(\text{Cu-Kz}) = 8.920 \text{mm}^{-1}, T = 123.5 \text{K}. Total 50748 reflections, 6985 unique, \text{R}_{int} = 0.049. Refinement of 6361 reflections (505 parameters) with \text{I} > 2\sigma (\text{I}) converged at final \text{R}_1 = 0.0239 (\text{R}_1 \text{ all data} = 0.0251), \text{wR}_2 = 0.0257 (\text{wR}_2 \text{ all data} = 0.0282), \text{gof} = 1.0886.

Computational details

DFT calculations were carried out with the D.01 revision of the Gaussian 09 program package\textsuperscript{39} using Becke’s three-parameter B3LYP exchange-correlation functional\textsuperscript{40,41} together with the 6-31G** basis set for C, H, N, S and O\textsuperscript{42} and the “double-\zeta” quality LANL2DZ basis set for Ir.\textsuperscript{43} The geometries of the singlet ground state (S0) and of the lowest-energy triplet state (T1) were fully optimized without imposing any symmetry restriction. The geometry of T1 was calculated at the spin-unrestricted UB3LYP level with a spin multiplicity of three. Phosphorescence emission energies were estimated as the vertical difference between the energy of the minimum of the lowest-energy triplet state and the energy of S0 at the T1 optimized geometry. All the calculations were performed in the presence of the solvent (acetonitrile). Solvent effects were considered within the self-consistent reaction field (SCRF) theory using the polarized continuum model (PCM) approach.\textsuperscript{44-46} The calculation of the energy of S0 at the T1 geometry was performed as an equilibrium single-point calculation with respect to the solvent reaction field/ solute electronic density polarization process. Time-dependent DFT (TD-DFT)\textsuperscript{47-49} calculations of the lowest lying 15 triplets were performed in the presence of the solvent at the lowest-energy triplet geometry optimized for the ground state.

Device preparation and characterization

Electroluminescent LEC devices were made as follows. First, glass substrates with patterned ITO contact (Naranjo Substrates) were cleaned by sonication with soap, deionized water, isopropanol and an UV-O3 lamp for 20 min. Then, an 80 nm layer of poly(3,4-ethylenedioxythiophene):polystyrene sulfonate (PEDOT:PSS) (CLEVIOS™ P VP Al 4083, aqueous dispersion, 1.3–1.7% solid content, Heraeus) was spin-coated on the ITO substrates to improve the reproducibility of the devices and to prevent the formation of pinholes. After this layer deposition, a 100 nm transparent film of the iTMC and the ionic liquid (IL) 1-butylium-3-methylidiazolium hexafluorophosphate ([Bmim][PF6]) (Sigma-Aldrich) in a 4 : 1 molar ratio were spin-coated from a 20 mg mL\textsuperscript{-1} acetonitrile solution. The devices were transferred into an inert atmosphere glovebox (<0.1 ppm O2 and H2O, M. Braun). The Al electrode (70 nm) was thermally vapour-deposited using a shadow mask under a vacuum (<1 \times 10\textsuperscript{-6} mbar) with an Edwards Auto500 evaporator integrated in the glovebox. The final device configuration was ITO/PEDOT:PSS/iTMC:IL/Al. The device characteristics were measured by applying a pulsed current (average current density 100 A m\textsuperscript{-2}, 50% duty cycle, 1 kHz, block wave) and monitoring the current and luminance versus time by a True Colour Sensor MAZeT (MTCSICT Sensor) with a Botest OLT OLED Lifetime-Test System. The electroluminescence spectra were measured using an Avantes AvaSpec-2048 Fiber Optic Spectrometer.

Results and discussion

Ligand synthesis and characterization

The sulfone-substituted cyclometallating ligand H2 was prepared in three steps in an analogous manner to the synthesis of 2-(4-methylsulfonyl)phenylpyridine (H1),\textsuperscript{11} starting from 2-bromopyridine and 3-fluorophenylboronic acid. The first step was a Suzuki coupling in EtOH/H2O with PdCl2,\textsuperscript{28} followed by a nucleophilic aromatic substitution with NaSMe to yield the corresponding thioether derivative. Oxidation of sulfur using \text{H2O2}/Na2WO4 in MeOH to give H2 (Scheme 4) was performed.
Synthesis and characterization of [Ir(C^N)2(N^N)][PF6] complexes

Complexes [Ir(C^N)2(N^N)][PF6] (C^N = [1]^- and [2]^2-; N^N = pzpy and dmpzpy) were synthesized from the corresponding dimers and ancillary ligands pzpy and dmpzpy in MeOH in a microwave reactor (Scheme 6). Subsequent anion exchange yielded the products as yellow powders in good yields.

In the ESI mass spectra of [Ir(C^N)2(N^N)][PF6] (C^N = [1]^- and [2]^2-, N^N = pzpy and dmpzpy), the base peak in each case corresponded to [M–PF6]^+ and exhibited a characteristic iridium isotope pattern. ^1H and ^13C NMR spectra were assigned using COSY, NOESY, HMQC and HMBC methods and were consistent with the structures shown in Scheme 6. The ^1H NMR spectrum of complex [Ir(2)2(pzpy)][PF6] is shown in Fig. 1 as a representative example. The cyclometallating ligands are inequivalent due to the asymmetric pzpy ligand; the most characteristic signals are those for H^46 and H^56 which occur at lowest frequency. The inequivalent cyclometallating ligands were distinguished by NOESY cross peaks between protons H^46 and H^56, and between protons H^55/H^55-Me and H^66 and/or H^56 (see Scheme 6 for numbering).

Crystal structure of [Ir(2)2(dmpzpy)][PF6]·MeCN

X-Ray quality crystals of [Ir(2)2(dmpzpy)][PF6]·MeCN were grown by overlaying a MeCN solution with Et3O. Structural determination confirmed the expected octahedral environment of the iridium(III) centre and the presence of three chelating ligands. The cation is necessarily chiral and the asymmetric unit contains the A-[Ir(2)2(dmpzpy)]^2+ cation shown in Fig. 2; important bond parameters are given in the figure caption. The compound crystallizes in the monoclinic space group C2/c and the unit cell therefore contains both

Scheme 5 Synthetic route to [Ir(2)2Cl2]. Conditions: (i) [Ir(cod)Cl]2, 2-ethoxyethanol, N2, overnight, reflux. Numbering scheme is for NMR spectroscopic assignments.

Scheme 4 Synthetic route to H1 and H2. Conditions: (i) 2-bromopyridine, PdCl2, K2CO3, EtOH/H2O 1 : 1, overnight, room temperature or reflux; (ii) NaSMe, N-methyl-2-pyrroliodine, N2, microwave reactor, 1 h, 80 °C or 120 °C; (iii) H2O2, Na2WO4·2H2O, MeOH, overnight, room temperature. Numbering scheme is for NMR spectroscopic assignments.

under mild conditions. High yields were achieved in all three steps. Compound H2 was characterized using multinuclear NMR spectroscopy, mass spectrometry, infrared spectroscopy and elemental analysis. The base peak in the LC-ESI mass spectrum corresponded to [M + H]^+ with an isotope pattern matching that calculated. ^1H and ^13C NMR signals were assigned using COSY, NOESY, HMQC and HMBC methods. Pyrazolylpyridine ancillary ligands were synthesized according to literature.34
A preliminary structure of $\text{[Ir}(1)_2\text{(dmpzpy)}][\text{PF}_6]$ was determined with crystals grown from a MeCN solution of the complex overlaid with Et$_2$O. This revealed the octahedral metal centre as well as the presence of the three bidentate ligands, confirming the expected structure of the complex.

**Photophysical properties**

Absorption spectra of $\text{[Ir}(C^N)\text{N}(N^N)][\text{PF}_6]$ ($C^N = [1]^-$ and $[2]^-$, $N^N = \text{pzpy}$ and dmpzpy) were recorded in MeCN solutions (Fig. 4). The spectra are similar and the most intense absorption bands lie in the UV region and are assigned to spin-allowed, ligand-centred $\pi^* \rightarrow \pi$ transitions. Between 350 and 450 nm, less intense bands arise from $1\text{MLCT}$ and $1\text{LLCT}$ transitions.$^3$ The lower energy absorption tails of $\text{[Ir}(1)_2(N^N)][\text{PF}_6]$ extend further into the visible than those of $\text{[Ir}(2)_2(N^N)][\text{PF}_6]^-$, with broad, but weak, maxima at 390 and 420 nm (Fig. 4).

The photoluminescence spectra of the complexes were recorded in MeCN solution both at room temperature and at 77 K and are shown in Fig. 5; emission maxima are summarized in Table 1. The emission profile is independent of the excitation wavelength. In complexes of the type $\text{[Ir}(C^N)\text{N}(N^N)]^-$, emission occurs from the lowest-lying triplet state (T$_1$), which comprises contributions from metal-to-ligand/ligand-to-ligand charge-transfer transitions.

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**Figure 1** 500 MHz $^1$H NMR spectrum of complex $\text{[Ir}(2)_2\text{(pzpy)}][\text{PF}_6]$ in CD$_2$CN with signal assignments. See Scheme 6 for proton labelling. Scale: $\delta$/ppm.

**Figure 2** Structure of the $\text{[Ir}(2)_2\text{(dmpzpy)}]^+$ cation in $\text{[Ir}(2)_2\text{(dmpzpy)}][\text{PF}_6]$: MeCN; ellipsoids plotted at 40% probability level and H atoms omitted for clarity. Selected bond lengths and angles: $\text{Ir}_1$–$\text{N}_1 = 2.144(2)$, $\text{Ir}_1$–$\text{N}_2 = 2.126(2)$, $\text{S}_1$–$\text{C}_21 = 1.763(3)$, $\text{S}_1$–$\text{O}_1 = 1.441(2)$, $\text{S}_2$–$\text{O}_3 = 1.442(2)$, $\text{S}_2$–$\text{O}_4 = 1.437(2)$, $\text{C}_22$–$\text{S}_1 = 1.769(3)$, $\text{N}_2$–$\text{N}_3 = 1.385(3)$ Å, $\text{N}_1$–$\text{Ir}_1$–$\text{N}_3 = 75.53(8)$, $\text{N}_4$–$\text{Ir}_1$–$\text{C}_21 = 80.34(9)$, $\text{N}_5$–$\text{Ir}_1$–$\text{C}_33 = 80.67(9)$, $\text{N}_4$–$\text{Ir}_1$–$\text{N}_5 = 75.53(8)$, $\text{N}_1$–$\text{Ir}_1$–$\text{C}_21 = 178.07(9)$, $\text{N}_3$–$\text{Ir}_1$–$\text{C}_33 = 171.26(9)$, $\text{O}_1$–$\text{S}_1$–$\text{O}_2 = 119.32(14)$, $\text{O}_3$–$\text{S}_2$–$\text{O}_4 = 118.32(12)$ Å.

**Figure 3** Packing diagram for $\text{[Ir}(2)_2\text{(dmpzpy)}][\text{PF}_6]$: MeCN showing sheets of cations in the bc-plane separated by $\text{[PF}_6^-$ anions.

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enantiomers. Both the $\text{[PF}_6^-$ anion and MeCN solvent molecule are ordered. Analyses of the structures of aryl-alkyl sulfones$^{34}$ and diaryl sulfones$^{54,55}$ demonstrate the presence of intra- and intermolecular $\text{CH}_\text{aryl}$⋯$\text{OS}$ hydrogen bonds, and we have observed this feature both in the structures of the ligand $\text{H}_1$ and the iridium dimer $\text{[Ir}(1)_2\text{Cl}_2]$. In the $\text{[Ir}(2)_2\text{(dmpzpy)}]^+$ cation, the sulfone groups are twisted with respect to the phenyl rings to which they are bonded with torsion angles of $\text{O}_1$–$\text{S}_1$–$\text{C}_18$–$\text{C}_17 = 27.4(3)$, $\text{O}_2$–$\text{S}_1$–$\text{C}_18$–$\text{C}_19 = -22.4(3)$, $\text{O}_3$–$\text{S}_2$–$\text{C}_30$–$\text{C}_31 = 15.0(3)$ and $\text{O}_4$–$\text{S}_2$–$\text{C}_30$–$\text{C}_29 = -35.1(2)^\circ$ giving rise to short $\text{O}_\text{sulfone}$⋯$\text{HC}_\text{phenyl}$ contacts in the range 2.6–2.7 Å. Intermolecular $\text{O}_\text{sulfone}$⋯$\text{HC}_\text{arene}$ and $\text{O}_\text{sulfone}$⋯$\text{HC}_\text{methyl}$ interactions are dominant packing interactions along with $\text{PF}_\text{anion}$⋯$\text{HC}$ contacts. The $\text{[Ir}(2)_2\text{(dmpzpy)}]^+$ cations pack into sheets that are coparallel with the $bc$-plane (Fig. 3).
MeCN solution Powder samples

At 298 K At 77 K At 298 K

Complex $\lambda_{\text{em}}^\text{max}$ [nm] $\varphi$ [µs] PLQY$^a$ [%] $k_r$ [10$^5$ s$^{-1}$] $k_{\text{nr}}$ [10$^5$ s$^{-1}$] $\lambda_{\text{em}}^\text{max}$ [nm] $\lambda_{\text{em}}^\text{max}$ [nm] $\varphi$ [µs] PLQY$^a$ [%]

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<th>$\varphi$ [µs]</th>
<th>PLQY$^a$ [%]</th>
<th>$k_r$ [10$^5$ s$^{-1}$]</th>
<th>$k_{\text{nr}}$ [10$^5$ s$^{-1}$]</th>
<th>$\lambda_{\text{em}}^\text{max}$ [nm]</th>
<th>$\lambda_{\text{em}}^\text{max}$ [nm]</th>
<th>$\varphi$ [µs]</th>
<th>PLQY$^a$ [%]</th>
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<td>[Ir(2)$_2$(pzpy)][PF$_6$]</td>
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<td>72</td>
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<td>1.2</td>
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<tr>
<td>[Ir(2)$_2$(dmpzpy)][PF$_6$]</td>
<td>463, 493</td>
<td>2.27</td>
<td>53</td>
<td>2.3</td>
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<td>459, 494, 523</td>
<td>491</td>
<td>0.510 (94%), 1.33 (6%)</td>
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</table>

$^a$ $\lambda_{\text{exc}} = 261$ nm for [Ir(1)$_2$(N$^\text{N}$N)][PF$_6$], 259 nm for [Ir(2)$_2$(N$^\text{N}$N)][PF$_6$].

$^b$ $\lambda_{\text{exc}} = 280$ nm.

$^c$ $\lambda_{\text{exc}} = 320$ nm.
higher than for their non-substituted analogues, but the difference is less significant than in the previously reported examples.\textsuperscript{35} When solutions of $[\text{Ir}(\text{C}^N\text{N})_2(\text{N}^N\text{N})][\text{PF}_6]$ ($\text{C}^N = [1^-]$ and $[2^-]$), $\text{N}^N = \text{pzpy}$ and dmpzpy) are not de-aerated, quantum yields decrease significantly to between 2.9 and 5.1%, consistent with oxygen quenching.

Excited-state lifetimes ($\tau$) in de-aerated MeCN solution are of the order of a few microseconds for all the complexes. $[\text{Ir}(\text{pzpy})_2(\text{pzpy})][\text{PF}_6]$ and $[\text{Ir}(\text{bpy})_2(\text{bpy})][\text{PF}_6]$\textsuperscript{36} have lifetimes of 2.3 $\mu$s, slightly longer than $\tau$ of $[\text{Ir}(\text{pppy})_2(\text{pppy})][\text{PF}_6]$ ($1.56 \mu$s).\textsuperscript{4} The complexes $[\text{Ir}(1)(\text{N}^N\text{N})][\text{PF}_6]$ exhibit longer excited state lifetimes of around 5 $\mu$s. For $[\text{Ir}(1)(\text{N}^N\text{N})][\text{PF}_6]$ and $[\text{Ir}(2)(\text{dmpzpy})][\text{PF}_6]$, the calculated radiative decay rate constants ($k_r$) are between $1.3 \times 10^5$ and $2.3 \times 10^5$ s$^{-1}$; these values compare to a $k_r$ value of $1.5 \times 10^5$ s$^{-1}$ for $[\text{Ir}(\text{pppy})_2(\text{pppy})][\text{PF}_6]$.\textsuperscript{8} Only $[\text{Ir}(2)(\text{pzpy})][\text{PF}_6]$ has a higher $k_r$ of $3.1 \times 10^5$ s$^{-1}$, comparable to $[\text{Ir}(1)(\text{bpy})][\text{PF}_6]$ with a value of $3.2 \times 10^5$ s$^{-1}$.\textsuperscript{11} A higher $k_r$ indicates a lower $^3\text{LC}$ character of the emissive state.\textsuperscript{7,8}

Excitation of powder samples of $[\text{Ir}(\text{C}^N\text{N})_2(\text{N}^N\text{N})][\text{PF}_6]$ ($\text{C}^N = [1^-]$ and $[2^-]$, $\text{N}^N = \text{pzpy}$ and dmpzpy) results in the emission spectra shown in Fig. 6, and PL data are summarized in Table 1. With respect to the solution spectra (Fig. 5), the solid-state emission maxima are red-shifted by about 30 nm and vibrational structure is less pronounced, suggesting strong intermolecular interactions in the solid state.\textsuperscript{9} Such a red-shift of the emission maximum on going from the solution to the solid seems to be common for blue or blue-green emitting iridium complexes. The extent of the red-shift can, however, vary between $\leq 15$ nm\textsuperscript{8,28,58,59} and $\geq 33$ nm.\textsuperscript{29,60,61}

In our previously reported sulfone-substituted $[\text{Ir}(\text{C}^N\text{N})_2(\text{bpy})][\text{PF}_6]$ complexes, red-shifts in the emission maxima of $\approx 40$ nm were observed on going from solution to solid. In contrast, the yellow emitting fluoroine- and thioether-containing $[\text{Ir}(\text{C}^N\text{N})_2(\text{bpy})][\text{PF}_6]$ in this series showed blue-shifts of 10–19 nm from solution to powder.\textsuperscript{13} Excited-state lifetimes (determined using second-order fits) in the solid state are much smaller than in MeCN solution. In addition to the dramatically decreased PLQYs of powdered samples ($\leq 11\%$), this suggests strong quenching of the excited state in powder samples.

### Electrochemical properties

Cyclic voltammetric data for $[\text{Ir}(\text{C}^N\text{N})_2(\text{N}^N\text{N})][\text{PF}_6]$ ($\text{C}^N = [1^-]$ and $[2^-]$, $\text{N}^N = \text{pzpy}$ and dmpzpy) are given in Table 2. All complexes show reversible oxidation processes. However, the first reduction is only reversible for $[\text{Ir}(1)(\text{N}^N\text{N})][\text{PF}_6]$. The HOMO is located on the cyclometallating ligand as well as the iridium(m) centre.\textsuperscript{5} Therefore, the similarity between $E^{*}_{1/2}$ for $[\text{Ir}(1)(\text{N}^N\text{N})][\text{PF}_6]$ and $[\text{Ir}(1)(\text{bpy})][\text{PF}_6]$ (+1.19, +1.20 and +1.18 V, respectively, with respect to Fc/Fc$^+$) and for $[\text{Ir}(2)(\text{N}^N\text{N})][\text{PF}_6]$ (+1.26 and +1.27 V) is as expected. The oxidation of each of $[\text{Ir}(2)(\text{N}^N\text{N})][\text{PF}_6]$ ($\text{N}^N = \text{pzpy}$ and dmpzpy) is shifted to slightly higher potential, consistent with stabilization of the HOMO. Compared to a value of $-1.72$ V for $[\text{Ir}(1)(\text{bpy})][\text{PF}_6]$,\textsuperscript{47} the first reduction occurs at lower potential (between $-2.00$ and $-2.14$ V). This is consistent with the replacement of bpy by the more electron-rich pyrazolylpyridine ligands leading to a destabilization of the LUMO. The larger electrochemical bandgap for complexes $[\text{Ir}(2)(\text{N}^N\text{N})][\text{PF}_6]$ is consistent with the blue-shift of the emission compared to that of $[\text{Ir}(1)(\text{N}^N\text{N})][\text{PF}_6]$.

### DFT/TD-DFT calculations

To gain a deeper insight into the electrochemical and photophysical properties of complexes $[\text{Ir}(\text{C}^N\text{N})_2(\text{N}^N\text{N})][\text{PF}_6]$ ($\text{C}^N = [1^-]$ and $[2^-]$, $\text{N}^N = \text{pzpy}$ and dmpzpy), a combined DFT/TDDFT theoretical investigation was undertaken at the B3LYP/(6-31G** + LANL2DZ) level in the presence of solvent (acetonitrile). A similar study was recently performed for $[\text{Ir}(\text{ppy})_2(\text{ppy})]^+$,\textsuperscript{62} which is used here as a reference for comparison purposes.

Calculations predict a near-octahedral structure for the complexes in their ground electronic state ($S_0$). The ancillary ligand is mostly planar in the pzpy complexes, the inter-annular N–N–C–N dihedral angle having a value of 3.4$^\circ$. The presence of the methyl group in 5-position of the coordinated dmpzpy ligand introduces some steric hindrance and the N–N–C–N angle increases to 11.5$^\circ$. This loss of planarity does not significantly affect the Ir-N$_{pz}$ and Ir-N$_{pz}$ coordination distances with the ancillary ligand, that change from 2.23 and 2.19 Å in the complexes with $\text{N}^N = \text{pzpy}$ to 2.21 and 2.21 Å in those with $\text{N}^N = \text{dmmpzpy}$, respectively.

Fig. 7 compares the energy and electron density contours calculated for the highest occupied (HOMO) and lowest-occupied molecular orbitals (LUMO and LUMO+1) of $[\text{Ir}(\text{ppy})_2(\text{ppy})]^+$ with those obtained for $[\text{Ir}(\text{C}^N\text{N})_2(\text{N}^N\text{N})]^+$ ($\text{C}^N = [1^-]$ and $[2^-]$) and $\text{N}^N = \text{pzpy}$ and dmpzpy). The introduction of the sulfone groups leads to a stabilization of the HOMO, which is located, as expected, on the Ir atom and the phenyl rings of the cyclometallating ligands. The stabilization is slightly larger (0.08 eV) for the complexes with $\text{C}^N = [2^-]$ because of the higher participation of the carbon in 4-position to which the sulfone group is attached. The attachment of methyl groups to the N–N ligand has no relevant effect on the energy of the HOMO because this ligand does not contribute to the orbital. These trends explain the higher oxidation potentials measured for $[\text{Ir}(2)(\text{N}^N\text{N})][\text{PF}_6]$ (+1.27 V) compared with $[\text{Ir}(1)(\text{N}^N\text{N})][\text{PF}_6]$ (+1.20 V) and the negligible changes observed when substituting the pzpy ligand by dmpzpy (Table 2). The oxidation potential

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**Fig. 6** Photoluminescence spectra of complexes $[\text{Ir}(\text{C}^N\text{N})_2(\text{N}^N\text{N})][\text{PF}_6]$ ($\text{C}^N = [1^-]$ and $[2^-]$, $\text{N}^N = \text{pzpy}$ and dmpzpy) in the solid state (powder). Excitation wavelength: 280 nm.
Table 2: Electrochemical data of complexes $\text{[Ir(C^N)2(N^N)] PF_6}$ (C^N = [1]^− and [2]^−, N^N = pzpy and dmpzpy) in MeCN solution referenced to Fc/Fc\textsuperscript{+} with 0.1 M $\text{[Bu$_4$N]+}$ as supporting electrolyte and a scan rate of 0.1 V s\textsuperscript{-1} (ir = irreversible, qr = quasi-reversible).

<table>
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<tr>
<th>Complex</th>
<th>$E^{\text{ox}}_{1/2}$ [V]</th>
<th>$E^{\text{red}}_{1/2}$ [V]</th>
<th>$\Delta E_{1/2}$ \textsuperscript{a} [V]</th>
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<tbody>
<tr>
<td>$\text{[Ir(1)(pzpy)] PF_6}$</td>
<td>+1.19</td>
<td>−2.02, −2.26\textsuperscript{ir}, −2.68\textsuperscript{ir}</td>
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<td>$\text{[Ir(1)(dmmpzpy)] PF_6}$</td>
<td>+1.20</td>
<td>−2.00, −2.27\textsuperscript{ir}, −2.42\textsuperscript{ir}, −2.63\textsuperscript{ir}</td>
<td>3.20</td>
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<tr>
<td>$\text{[Ir(2)(pzpy)] PF_6}$</td>
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<tr>
<td>$\text{[Ir(2)(dmmpzpy)] PF_6}$</td>
<td>+1.27</td>
<td>−2.14\textsuperscript{ir}, −2.37\textsuperscript{ir}, −2.68\textsuperscript{ir}</td>
<td>3.41</td>
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</table>

\textsuperscript{a} $\Delta E_{1/2} = E^{\text{ox}}_{1/2} - E^{\text{red}}_{1/2}$

The combination of these effects leads to HOMO–LUMO energy gaps that increase along the series $\text{[Ir(pzpy)$_2$(pzpy)]}$: 3.66 eV < $\text{[Ir(1)(pzpy)]}$: 3.86 eV < $\text{[Ir(2)(pzpy)]}$: 4.02 eV < $\text{[Ir(2)(dmmpzpy)]}$: 4.08 eV. This trend correctly reproduces the evolution of the electrochemical gap (3.07 V < 3.21 V < 3.37 V < 3.41 V, respectively, Table 2) and the blue-shifted emission of complexes $\text{[Ir(2)(N^N)]}$ with respect to $\text{[Ir(1)(N^N)]}$ (Fig. 5). Indeed, if a direct correlation is established between the HOMO–LUMO gap and the emission energy, all the complexes should be expected to be bluer than $\text{[Ir(pzpy)$_2$(pzpy)]}$. This is however the case since the emission of $\text{[Ir(1)(pzpy)]}$ and $\text{[Ir(1)(dmmpzpy)]}$ ($\lambda_{\text{em}}^{\text{max}} = 491, 523$ nm, Table 1) is red-shifted compared to $\text{[Ir(pzpy)$_2$(pzpy)]}$ ($\lambda_{\text{em}}^{\text{max}} = 475, 503$ nm),\textsuperscript{b} whereas that of $\text{[Ir(2)(pzpy)]}$ and $\text{[Ir(2)(dmmpzpy)]}$ ($\lambda_{\text{em}}^{\text{max}} = 463, 493$ nm) is blue-shifted.

To disentangle this apparent contradiction between theory and experiment and to obtain more information about the emitting state, the geometry of the lowest-energy triplet excited state $T_1$ was optimized using the spin-unrestricted UB3LYP approach. Fig. 8a summarizes the adiabatic energy difference between $S_0$ and $T_1$ ($\Delta E$) and the emission energy ($E_{\text{em}}$) estimated as the vertical energy difference between $T_1$ and $S_0$ at the optimized minimum-energy geometry of $T_1$. Fig. 8b shows the unpaired-electron spin density distributions calculated for the lowest-energy triplet of complexes $\text{[Ir(1)(pzpy)]}$ and $\text{[Ir(2)(pzpy)]}$. Almost identical spin-density plots are obtained for complexes with N^N = dmmpzpy and for $\text{[Ir(bpy)$_2$(pzpy)]}$. Therefore, independently of the nature of the LUMO, the spin-density distribution calculated for $T_1$ is mainly centred on one of the cyclometallating ligands (~1.65 unpaired electrons) with some contribution from the iodinum atom (~0.29e). This indicates that the lowest-energy triplet state has a predominant LC nature with some MLCT character in accord with the structured shape of the emission band observed experimentally.

In agreement with experiment (Table 1), calculations find almost identical emission energies for complexes with C^N = [1]^− ($E_{\text{em}} \sim 2.25$ eV) and C^N = [2]^− ($E_{\text{em}} \sim 2.41$ eV). The emission energy is not affected by the addition of methyl groups to the ancillary ligand because the emission is centred on the C^N ligands (Fig. 8). Compared with $\text{[Ir(pzpy)$_2$(pzpy)]}$ ($E_{\text{em}} = 2.35$ eV), the calculated emission energies correctly reproduce the red-shifted and blue-shifted emission observed experimentally for complexes $\text{[Ir(1)(N^N)]}$ and $\text{[Ir(2)(N^N)]}$, respectively.

Time dependent DFT (TD-DFT) calculations confirm the $^3$LC nature of the lowest-lying triplet (Table S1). The first triplet
The state results from the excitation from the HOMO to the first virtual orbital located on the C^N ligands, i.e. the LUMO for \([\text{Ir}(1)_2(\text{pzpy})]^{+}\), \([\text{Ir}(1)_2(\text{dmpzpy})]^{+}\), and \([\text{Ir}(2)_2(\text{pzpy})]^{+}\), and the LUMO + 1 for \([\text{Ir}(pppy)_2(\text{pzpy})]^{+}\) and \([\text{Ir}(2)_2(\text{pzpy})]^{+}\). A second 3LC triplet is found very close in energy (~0.05 eV) due to the excitation from the HOMO to the second virtual MO centred on the C^N ligands. The first 3MLCT state is predicted to be higher than the 1LC state by an energy ranging from 0.2 eV in \([\text{Ir}(pppy)_2(\text{pzpy})]^{+}\) to 0.6 eV in \([\text{Ir}(1)_2(\text{dmpzpy})]^{+}\).

**Electroluminescence**

Complexes \([\text{Ir}(\text{C}^\text{N})_2(\text{N}^\text{N})][\text{PF}_6]\) \((\text{C}^\text{N} = \text{[1]}^{+} \text{ and } 2^{+}, \text{N}^\text{N} = \text{pzpy and dmpzpy})\) were tested in LEC devices, which were operated using a pulsed current driving mode with a frequency of 1 kHz and a duty cycle of 50%. The LEC performance parameters measured using an average current density of 100 A m\(^{-2}\) are given in Table 3. Under these conditions, all the devices showed a typical LEC behaviour (Fig. S1–S4†). The voltage needed to maintain the established current density starts at a high value due to the initially high injection barriers and rapidly drops as a result of ion migration towards the electrodes forming an electric double-layer that reduces the injection barriers. The time needed to reach the maximum luminance \((t_{\text{on}})\) is relatively short ranging from 30 to 78 seconds. The relatively fast response matches with the rather poor stability of the devices with lifetimes \((t_{1/2})\) of a few minutes (Table 3). The results are comparable to those obtained for other blue-green LECs reported in the literature, for which a limited lifetime was observed especially when operated using a pulsed driving mode. In these cases, a relatively fast \(t_{\text{on}}\) was also measured which is consistent with the values reported here.

The maximum luminance achieved under pulsed current density of 100 A m\(^{-2}\) was 141 cd m\(^{-2}\) for the LEC prepared using \([\text{Ir}(\text{pzpy})_2][\text{PF}_6]\). The device performance is therefore not very high due to the moderate efficiencies. The trends obtained for the power conversion efficiency (PCE) and the external quantum efficiency (EQE) follow the photoluminescence quantum yields measured for thin films of the complexes with the same composition of the active layer in the device (Fig. S5†). Complex \([\text{Ir}(\text{pzpy})_2][\text{PF}_6]\) with the lowest efficiency values (Table 3) show the smallest PLQY (6.9%), and the other three complexes with comparable higher efficiencies have PLQY values ranging from 13.0 to 16.9%.

**Table 3** Performance parameters obtained for ITO/PEDOT:PSS/active layer/Al LEC devices by applying a block-wave pulsed current of 100 A m\(^{-2}\) at a frequency of 1 kHz and duty cycles of 50%. Active layer: \([\text{Ir}(\text{C}^\text{N})_2(\text{N}^\text{N})][\text{PF}_6] : [\text{Bmim}][\text{PF}_6] 4 : 1\) molar ratio \([\text{Bmim}][\text{PF}_6] = 1\)-butyl-3-methylimidazolium hexafluorophosphate

<table>
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<tr>
<th>Complex cation</th>
<th>(t_{\text{on}}) (a) [s]</th>
<th>(\text{Lum}_{\text{max}}) (b) [cd m(^{-2})]</th>
<th>(t_{1/2}) (c) [min]</th>
<th>PCE (d) [lm W(^{-1})]</th>
<th>EQE (d) [%]</th>
<th>(\lambda^\text{max}) (d) [nm]</th>
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<td>([\text{Ir}(1)_2(\text{pzpy})]^{+})</td>
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<tr>
<td>([\text{Ir}(1)_2(\text{dmpzpy})]^{+})</td>
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<td>125</td>
<td>6.2</td>
<td>0.6</td>
<td>0.4</td>
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<tr>
<td>([\text{Ir}(2)_2(\text{pzpy})]^{+})</td>
<td>42</td>
<td>49</td>
<td>3.5</td>
<td>0.2</td>
<td>0.2</td>
<td>466, 496, 547</td>
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<tr>
<td>([\text{Ir}(2)_2(\text{dmpzpy})]^{+})</td>
<td>30</td>
<td>129</td>
<td>2.4</td>
<td>0.5</td>
<td>0.4</td>
<td>466, 496, 547</td>
</tr>
</tbody>
</table>

\(a\) Time to reach the maximum luminance. \(b\) Maximum luminance reached. \(c\) Time to reach one-half of the maximum luminance. \(d\) Power conversion efficiency (PCE). \(e\) External quantum efficiency (EQE).

**Fig. 8** (a) Schematic diagram showing the adiabatic energy difference \((\Delta E)\) between \(S_0\) and \(T_1\) and the emission energy \((E_{\text{em}})\) from \(T_1\) computed for \([\text{Ir}(\text{pzpy})_2(\text{pzpy})]^{+}\) and for complexes \([\text{Ir}(\text{C}^\text{N})_2(\text{N}^\text{N})][\text{PF}_6]\) \((\text{C}^\text{N} = \text{[1]}^{+} \text{ and } 2^{+}, \text{N}^\text{N} = \text{pzpy and dmpzpy})\). (b) Unpaired-electron spin density contours (0.002 au) calculated for the fully relaxed \(T_1\) state of \([\text{Ir}(1)_2(\text{pzpy})]^{+}\) and \([\text{Ir}(2)_2(\text{pzpy})]^{+}\).

**Fig. 9** Electroluminescence spectra of LEC devices containing complexes \([\text{Ir}(\text{C}^\text{N})_2(\text{N}^\text{N})][\text{PF}_6]\) \((\text{C}^\text{N} = \text{[1]}^{+} \text{ and } 2^{+}, \text{N}^\text{N} = \text{pzpy and dmpzpy})\) driven using a block-wave pulsed current of 100 A m\(^{-2}\) at a frequency of 1 kHz and duty cycles of 50%.
Fig. 9 shows the electroluminescent (EL) spectra recorded for the LECs prepared. All the LECs exhibit structured electroluminescent emission bands coherent with the LC character of the emissive state. The position of the peaks forming the EL band are similar to the PL spectra depicted in Fig. 5 (compare Tables 1 and 3), but their relative intensity drastically changes. For complexes [Ir(2)(N-N)][PF6], the EL spectra show bluer weak emission peaks at 466 and 496 nm and the maximum emission is at 546 nm (Fig. 9). For complexes [Ir(1)(N-N)][PF6], an intense peak is observed at 492 nm but the maximum emission appears around 530 nm. The relative intensity of the EL peaks determines that the LEC devices exhibit a green colour for all the four complexes.

Conclusions

A series of [Ir(C\(^{-}\))\(_{2}(N-N)\)]\(^{+}\) complexes with 2-phenylpyridine-based cyclometallating and pyrazolopyridine ancillary ligands has been synthesized. Two different cyclometallating ligands, H1 and H2, with a sulfonyl substituent in either the 3- or 4-position of the phenyl ring of the ppy domain were combined. Complexes were designed to blue-shift the emission of the [Ir(C\(^{-}\))\(_{2}(N-N)\)]\(^{+}\) complex. In practice, blue or green emissions were observed, the colour depending on the sulfonyl substituent position in the cyclometallating ligand.

The complexes were characterized by spectroscopic and mass spectrometric methods, and the single crystal structure of [Ir(2)(dmpzpy)][PF6] in MeCN solution leads to structured emission spectra with maxima in the green region for complexes with 2-(4-methylsulfonylphenyl)pyridine and 2-(3,5-dimethyl-1H-pyrazol-1-yl)pyridine (dmpzpy). These ligand combinations were designed to blue-shift the emission of the [Ir(C\(^{-}\))\(_{2}(N-N)\)]\(^{+}\) complex. The marked vibrational structure and the absence of a rigidochromic shift in the low temperature spectra indicate significant 3LC contributions to the emissive state. Theoretical calculations showed that the sulfonyl substituents strongly stabilize the orbitals located over the cyclometallating ligands and confirm the 3LC nature of the lowest-lying triplet state. PLQY values in de-aerated MeCN solution are quite high for all four complexes (between 53 and 77%), whereas the PLQY values of powdered samples are considerably lower than in solution (≤11%), suggesting strong quenching effects due to intermolecular interactions. Unfortunately, all four complexes performed poorly when tested in LEC configuration.

Methyl groups on the pyrazolopyridine ligand did not have a significant effect on the properties of the complexes since both the emission and electrochemical properties are mainly determined by the cyclometallating ligands. On the other hand, changing the substitution position of the sulfonyl group on the cyclometallating ligand produced a shift of ≈30 nm in the emission maximum. This shift is reproduced in the theoretical study and is due to the different effect that sulfonyl substitution has on the frontier molecular orbitals depending on the substitution position. We are currently investigating the effect of substituential isomerism on the cyclometallating ligand in relation to colour tuning in more detail.

Acknowledgements

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Notes and references