In Search of High-Performance Platinum(II) Phosphorescent Materials for the Fabrication of Red Electroluminescent Devices


The rational design and syntheses of Pt(iqdz)_2 (1) and Pt(pyzdz)_2 (2) bearing isoquinolinyl indazole (iqdz)H or pyridyl indazole (pydz)H groups on the coordinating ligands are reported. Single-crystal X-ray diffraction studies of 1 reveal a square planar geometry, in which two iqdiz ligands adopt a trans-configuration. Short N···H contacts (~2.21 Å) are detected between the ortho-hydrogen atom of isoquinoline and the adjacent N atom of the indazole fragment, making the overall molecular geometry analogous to that of the platinum(ii) porphyrinato complexes. The lowest absorption bands for both complexes reveal strong state mixings between singlet and triplet (metal-to-ligand charge transfer and intra-ligand) manifolds. This, in combination with the introduction of a camphor-like structure to avoid the stacking effect, leads to phosphorescence with unprecedented brightness both in solution and in the solid state. Organic light-emitting diode (OLED) devices fabricated using 1 as a dopant emitter have been achieved in a multilayer configuration. The results constitute the first highly efficient Pt(ii)-based red OLED.

1. Introduction

Owing to their potential to harness the energies of both the singlet and triplet excitons after charge recombination, transition-metal-based phosphorescent materials have recently received considerable attention for fabricating organic light-emitting diodes (OLEDs). The main advantages are due to the heavy-atom-induced singlet-to-triplet intersystem crossing as well as the large enhancement of radiative decay rate from the resulting triplet manifolds. In this regard, numerous attempts have been made to exploit third-row transition metal complexes as dopant emitters for OLED fabrication, among which quite a few Pt(II), Os(II), and Ir(III) complexes have been reported to exhibit highly efficient device performances. Despite these developments, attempts to further expand the potential of the square planar Pt(II) complexes, in which the central metal ion possesses a higher atomic number than Os(II) and Ir(III), for efficient OLED applications has encountered many intrinsic obstacles. For example, the POEP (where H_2OEP = octaethylporphyrin) type of emitter commonly has a ligand-based phosphorescence with lifetimes as long as 30–50 µs, so that saturation of emissive sites and a rapid drop in device efficiency at high drive current is observed. Another factor contributing to the poor device efficiency is the planar molecular configuration of many Pt(II) complexes, which leads to stacking and the formation of aggregates or excimers. To realize the potential of Pt(II) materials for application in high-efficiency OLEDs, the rational design of Pt(II) complexes aimed at the reduction of the phosphorescence radiative lifetime and the prevention of stacking behavior are critical. In this paper, we report the design and synthesis of a new series of emissive Pt(II) complexes, in which the associated ligand chromophores possess a bulky, rigid camphor-like architecture to effectively suppress the aggregation effect. Moreover, drastic reduction of the phosphorescence radiative lifetime to several microseconds has also been achieved due to the strong singlet–triplet state mixing. Consequently, highly efficient Pt(II) OLEDs operating at 610-630 nm have been successfully prepared for the first time, in which the dopant (Pt(II) complex) concentration, owing to the reduced aggregation effect, can be substantially increased to maximize performance.

2. Results and Discussion

2.1. Synthesis and Characterization

As depicted in Scheme 1, the required camphor-derivatized chelating ligand was synthesized in two steps employing a direct condensation of ethyl 1-isoquinolinonecarboxylate and (1R,2R)-camphor, followed by treatment with hydrazine hy-
drate in refluxing ethanol to induce formation of the indazole skeleton. \(^{(7)}\) Subsequent treatment with \(\text{K}_2\text{PtCl}_4\) in a mixture of water and ethanol gave a dark-red Pt complex denoted as \(\text{Pt}(\text{iqdz})_2\) (1) in good yield. A similar method, but with ethyl 1-isoquinolinecarboxylate replaced by picolinate, yielded the yellowish green platinum complex \(\text{Pt}(\text{pydz})_2\) (2). Both Pt complexes were highly soluble in organic solvents and have been characterized using various spectroscopic methods (see Experimental section). Their spectroscopic data are in good agreement with the expected square planar Pt\(^{IV}\) complexes coordinated with two indazole chelates. Complex 1 was further examined by single-crystal X-ray diffraction analysis to establish its molecular structure.

Figure 1 depicts the ORTEP diagram of 1, along with that of a second, symmetry-related molecule shown in pale gray. An overlap between the isoquinoline fragments of these two proximal complexes is clearly noted with a moderate interplanar spacing of 3.64 Å. All methyl-substituted bridges on the camphor moieties point in the same direction due to the selection of a chiral reagent, (1R)-(+) camphor, for the original ligand preparation. The Pt–Pt separation was calculated to be 6.857(7) Å, which is much too long to expect any significant Pt–Pt interaction such as those observed in dimers of cyclometallated Pt\(^{II}\) complexes (3.15–3.76 Å).\(^{(8)}\) Moreover, the Pt atom is bonded to two chelating iqdz ligands in a trans-arrangement. The Pt–N\(_{\text{mda}}\) distances of 1.976 and 2.005 Å in 1 are slightly shorter than the Pt–N\(_{\text{quim}}\) dative bonds (2.011 and 2.047 Å), which are within the range reported for those of [Pt(α-ditopic)Cl\(_2\)] (1.992–2.015 Å) and related complexes.\(^{(9)}\) Moreover, short N–H contacts (~2.21 Å) are also detected between the ortho-hydrogen atom of isoquinoline and the adjacent N atom of the indazole fragments, forming H(1)···N(6) and H(21)···N(3) dual hydrogen bonds, which are further supported by the observation of an unusual downfield \(^1\)H NMR chemical shift of δ 10.93. The proposed intramolecular H-bonding interactions are reminiscent of those reported for pyridyl pyrazolate Os complexes as well as cobaloxime complexes.\(^{(10)}\)

It is believed that this unusual H-bonding pattern, to a certain extent, provides a strong driving force to stabilize the observed trans-geometry. In contrast to our discovery, the X-ray structural analyses of related Pt\(^{IV}\) complexes such as Pt(thppy)\(_2\), where thppy is 2-(2’-thienyl) pyridine, revealed that ligated thiopeine and pyridine fragments are oriented exclusively in a cis-configuration, despite the large distortion and deformation of square planar geometry that occurred at the junction of the ligands.\(^{(11)}\) This could be due to the exceedingly strong trans-effect exerted by the carbon donor atom of the ligated thiopeine.

### 2.2. Photophysical Measurements

As shown in Figure 2, although the energy gaps are quite different, similar spectral features are observed for complexes 1 and 2, consisting of a weak, broad band located in the long-wavelength region, accompanied by a vibronic progression feature in the short-wavelength region that is commonly assigned to the singlet π–π* intra-ligand transitions (\(^1\)IL). The low-frequency absorption bands have a relatively small extinction coefficient (1: 5240 M\(^{-1}\) cm\(^{-1}\) at 502 nm; 2: 4690 M\(^{-1}\) cm\(^{-1}\) at 457 nm) and are tentatively assigned to the transition incorporating a state mixing between singlet and triplet metal–ligand charge transfer (\(^1\)MLCT and \(^3\)MLCT) and, to a certain extent, the intra-ligand triplet state (\(^3\)IL). Support for this viewpoint is first provided by a distinct shoulder at 530 nm (ε ~ 4080 M\(^{-1}\) cm\(^{-1}\)) resolved in the absorption spectrum of complex 1, which can be tentatively assigned to the lowest lying \(^3\)MLCT band (see below). The close energetics and absorptivity between the \(^1\)MLCT and \(^3\)MLCT bands suggest that the \(^3\)MLCT transition, induced by spin–orbit coupling and the
proximal energy levels with respect to \( ^1 \text{MLCT} \), is greatly enhanced and becomes partially allowed. This novel spectral feature is in accord with data for other recently published Pt\(^{II} \) complexes bearing bis(phenoxo)diimine auxiliaries capable of tetradentate bonding.\(^{[13]} \) Although we have not explicitly examined and discussed the details of the singlet–triplet mixing, its presence is unambiguously confirmed by the obvious overlap between this UV-vis absorption band and the leading edge of the corresponding emission profile.

As for complex 2, the \( ^1 \text{MLCT} \) and \( ^3 \text{MLCT} \) states are so close that an asymmetric band was observed rather than the well-resolved dual absorption profile. Complex 1 exhibits an intensive emission with a maximum at 635 nm (quantum yield \( \Phi = 0.81 \); lifetime \( \tau = 5.34 \mu s \)) in degassed \( \text{CH}_2\text{Cl}_2 \). The oxygen quenching rate of \( 1.78 \times 10^9 \text{M}^{-1}\text{s}^{-1} \) for the emission in \( \text{CH}_2\text{Cl}_2 \), in combination with its spectral mirror image with respect to the lowest absorption profile, leads us to conclude that the emission mainly originates from a triplet manifold. Similarly, complex 2 also exhibits strong phosphorescence with a peak wavelength at 553 nm (\( \Phi = 0.64 \); \( \tau = 3.63 \mu s \)), which is comparable to those observed in the cyclometalated dipyridylbenzene Pt complexes.\(^{[13]} \) The observed radiative lifetimes for 1 and 2 in \( \text{CH}_2\text{Cl}_2 \) are relatively long for a pure \( ^3 \text{MLCT} \) emission from complexes incorporating a central heavy atom like Pt\(^{II} \) and a \( ^1 \text{MLCT} \) state in proximity. Accordingly, we tentatively propose that there exists, in part, a further state mixing with the \( ^3 \text{IL} \) manifold.\(^{[14]} \) Firm support for this viewpoint is given by the unusually broad, structureless emission feature, with a full width at half maximum (FWHM) of 108 nm and 109 nm for 1 and 2, respectively, in \( \text{CH}_2\text{Cl}_2 \) at room temperature (RT), while upon cooling to 77 K the emission reveals distinctive vibronic-like features with peak wavelengths at 580, 614 (520), 656 (552), and 698 (596) nm for complex 1 (2) (see Fig. 3 and Table 1). Although not well resolved, similar structural features were also observed for both complexes in the solid state at RT (see Fig. 2). The spectral progression of >1000 cm\(^{-1} \) for each successive peak cannot be rationalized by the much smaller d-level splitting in a square planar coordination, but is akin to that corresponding to the vibrational modes (1270–1300 cm\(^{-1} \)) of aromatic terpyridyl ligands.\(^{[15]} \) Alternatively, it may be plausible that the broad FWHM, together with the vibronic structure features in a 77 K \( \text{CH}_2\text{Cl}_2 \) solution as well as in the solid state at RT, arises from a state mixing between \( ^3 \text{MLCT} \) and \( ^3 \text{IL} \).\(^{[16]} \) In a central planar configuration such as 1 and 2, the strong mixing of these two transitions essentially requires covalent interaction of the relevant d-orbitals and the ligand \( \pi \)-system, in which MLCT [\( d_{x^2-y^2} \rightarrow \pi^* \)] transitions should be the most likely candidates.\(^{[17]} \)

Due to the planar geometry of the central Pt\(^{II} \) atom possessing a dsp\(^2 \) configuration, one has to consider the possible stacking effects for both 1 and 2. We thus carried out a concentration-dependent absorption/emission study in an attempt to resolve this issue. Upon varying the sample concentrations from \( 3.45 \times 10^{-3} \text{M} \) to \( 1.07 \times 10^{-2} \text{M} \), both absorption and emission spectra remain unchanged for 1 and 2, indicating that the stacking effect, if any, is too small to affect any associated photophysical behavior. It is thus reasonable to conclude that the introduction of a bulky camphor-derived group on the indazole fragment drastically increases the steric hindrance and hence suppresses aggregation. Supplementary support for this viewpoint is provided by the solid-state emission spectra in that both complexes (1: \( \lambda_{\text{em}} = 638 \text{ nm} \); 2: \( \lambda_{\text{em}} = 550 \text{ nm} \), see Fig. 2)

Figure 2. UV-vis absorption and emission spectra of 1 (2.68 \times 10^{-5} \text{ M}, filled circles joined by solid lines) and 2 (2.14 \times 10^{-5} \text{ M}, triangles joined by solid lines) in \( \text{CH}_2\text{Cl}_2 \) at RT. Note that the normalized emission spectra were acquired under degassed conditions. The dotted lines denote the corresponding solid-state emission obtained from a thin-film sample at RT.

Figure 3. The phosphorescence spectra of 1 (circles) and 2 (triangles) in the 77 K solid \( \text{CH}_2\text{Cl}_2 \) matrices. Insets: The phosphorescence decay profiles of complexes 1 and 2.
Table 1. Photophysical properties of complexes 1 and 2 in degassed CH₂Cl₂ at RT.

<table>
<thead>
<tr>
<th></th>
<th>( \lambda_{\text{abs}} , [\text{nm}] )</th>
<th>( \lambda_{\text{em}} , [\text{nm}] )</th>
<th>( \Phi )</th>
<th>( \tau , [\text{ns}] )</th>
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</thead>
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<tr>
<td>Pt(qdz)₂ (1)</td>
<td>502 (5240)</td>
<td>635</td>
<td>0.81</td>
<td>5.34</td>
</tr>
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<td></td>
<td></td>
<td>(610, 638, 688) [a]</td>
<td>0.20 [a]</td>
<td>1.10 [a,c]</td>
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<tr>
<td></td>
<td></td>
<td>(580, 614, 656, 698) [b]</td>
<td>3.0 [b]</td>
<td></td>
</tr>
<tr>
<td>Pt(pdz)₂ (2)</td>
<td>457 (4690)</td>
<td>553</td>
<td>0.64</td>
<td>3.63</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(528, 550, 581) [a]</td>
<td>0.15 [a]</td>
<td>3.38 [a,c]</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(520, 552, 596) [b]</td>
<td>2.78 [b]</td>
<td></td>
</tr>
</tbody>
</table>

[a] The solid-state emission obtained from a thin-film sample at RT. [b] The phosphorescent emission recorded in frozen CH₂Cl₂ matrices at 77 K. [c] Lifetime is an average value from a two-component fit.

manifest negligible spectral shifts from the corresponding emission maximum in solution, providing unambiguous evidence for the negligible stacking interaction in both complexes. It is noted that the solid-film photoluminescence (PL) is narrower than the solution PL. In addition, the PL spectra in the solid film show a slight blue shift for 2 and red shift for 1 in comparison with their corresponding emission in solution. If the solid state of a complex lacks strong intermolecular interaction such as hydrogen bonding, \( \pi \) stacking, etc., a slight blue shift and narrowing for the solid-film PL relative to that of the solution PL may be expected, and can be attributed to a “medium effect”. For complex 2 in solution, the stronger interaction from the solvent (e.g., CH₂Cl₂) makes the emission broader, whereas, due to the lack of \( \pi \) stacking, complex 2 is more or less frozen and inhibited from having a closer interaction with itself. In comparison, the additional fused benzene of isoquinoline in complex 1 introduces a weak but perhaps non-negligible \( \pi \) interaction in the solid film, resulting in a slightly red-shifted emission. Nevertheless, from solution to solid, the shift of peak wavelength is rather small for 1, indicating that the intermolecular interaction cannot be large. This viewpoint can be supported from X-ray single-crystal analysis, in which a rather long Pt–Pt distance has been resolved for 1 (see above).

Table 1 lists detailed spectroscopic and dynamics data for 1 and 2 in solution and for single crystals. Despite the much lower energy gap with respect to that of the pyridyl counterpart 2, significant enhancement of the luminescent quantum yield (\( \Phi \sim 0.81 \)) is observed in 1, accompanied by a longer lifetime (5.34 us) in degassed CH₂Cl₂. These results seem to contradict the energy-gap law, in which the theory pertaining to non-radiative decay concludes that the non-radiative deactivation should increase with decreasing energy gap of the transition.[18] We thus tentatively propose that the remarkable but unusual luminescence behavior in complex 1 is due to the highly conjugated \( \pi \) systems in indazole coupled with the nearby isoquinoline fragments. One possible strategy to suppress the active vibrational modes, such as ring stretching and bending of the acceptor ligand that commonly dominates the deactivation of the MLCT excited states, is to use a ligand with a rigid o-framework. In addition, upon excitation, the electron occupation of the lowest \( \pi^* \) acceptor orbital results in increases of the C–C and C–N bond distances, inducing non-radiative transition due to the loose bonding effect.[19] Since the Franck–Condon factor for non-radiative transitions is qualitatively proportional to the square of the bonding displacement,[20] enlarging the \( \pi \) conjugation should reduce the distortion of the ligand framework due to the smaller changes in average distance between ground and excited states. Accordingly, the non-radiative decay rate in 1 is expected to be relatively small to compete with the red phosphorescence, giving rise to an exceptionally high emission quantum yield. For comparison, Nazeruddin et al. have recently reported near unity quantum yields for blue, green, and yellow emission by meticulous selection of the ligand on the \( \text{Ir}^{III} \) system for strong ligand field strength, which increased the energy gaps between triplet emitting states and the nearby deactivating metal centered (MC) level.[21] Other factors leading to the high emission yield for 1 or 2 are also possible. These include a) Pt\( \text{II} \) metal ion intrinsically possessing a relatively large d-orbital splitting, b) isoquinolinyl indazole (or pyridyl indazole) with a fairly strong ligand field inducing a larger gap between the MC states and the lowest unoccupied molecular orbital (LUMO) of the ligands, and c) close-lying \( \pi^*\pi \) and MLCT states together with the heavy atom effect enhancing the spin–orbit coupling. A comprehensive understanding of the relaxation mechanisms might have to rely on future theoretical approaches, which we are currently exploring.

2.3. OLED Fabrication

Due to its high phosphorescence quantum efficiency in the red, multilayer device of the configuration ITO/NPB (40 nm)/CBP 1 (30 nm)/BCP 1 (10 nm)/Alq₃ (30 nm)/LiF (1 nm)/Al (150 nm) were prepared, with doping concentrations of 1 varying from 6%, 12%, 20%, 50% to a neat film. Very bright red emission was observed for all the concentrations prepared, including the one with a pure layer of the platinum complex. The current–voltage (\( I–V \)) curves, plotted in Figure 4, show a trend of increasing current density with increasing concentration of 1. The results may imply that the phosphorescent sites serve as charge trapping sites. The electroluminescence (EL) spectra originated only from the complex.

![Figure 4. I–V characteristics of OLED devices based on complex 1 for various dopant concentrations.](http://www.afm-journal.de)
plex in all cases, but with a small red shift of the EL spectra $\lambda_{\text{max}}$ with increasing dopant concentration from 610 nm for the 6% dopant concentration to 630 nm for the neat film (Fig. 5). The FWHM of the EL spectrum also increased slightly (from 76 nm to 92 nm) over the same range of doping concentrations. Comparing Figures 2 and 5, the EL from the device is slightly blue shifted from PL of the solid film. One possible origin for the shift is from the microcavity effect, which is caused by the interference between the forward-traveling light and the light reflected from the metal electrode.\textsuperscript{[22]}

![Figure 5. Electroluminescence spectra of OLED devices based on complex 1 for various dopant concentrations.](image)

All devices showed a turn-on voltage as low as 4.0 V. Although the devices still exhibited a similar dropping trend with increasing current (Fig. 6), as is the case for most phosphorescence-based devices, the performance characteristics are nevertheless very encouraging. For the device doped with 12% 1 driven at a current of 100 mA, a brightness of 10,677 cd/m\(^2\) was achieved with an external quantum efficiency of ~7%, a luminance efficiency of ~11 cd/A and a power efficiency of 3.3 lm/W. The results also exhibited a decreasing trend with increasing concentration of the platinum dopant. However, it is noteworthy that, even using a pure film of 1 as the emission layer, a brightness of 2653 cd/m\(^2\) and an external quantum efficiency of 2.46%, luminance efficiency of 2.65 cd/A and power efficiency of 0.93 lm/W can be achieved. The relatively high efficiency of the device even in the neat complex is tentatively attributed to the unusually short radiative lifetime that avoids the triplet–triplet annihilation. Table 2 summarizes the performance data for various concentrations studied. The achievement of high luminescence efficiency can be attributed to much shorter phosphorescence radiative lifetime in combination with a rationally designed structure that greatly suppresses the aggregation effect. It should be noted that rigid steric blockers such as a pinene functionality incorporated into an octahedral phenyl pyridine $\text{H}^{11}$ complex\textsuperscript{[23]} have been reported to effectively reduce self-quenching of the phosphorescent dopant. While improving the device luminescence efficiency, this prior innovation based on Ir(ppy)$_3$ and its pinene-derivatized complexes exhibits green emission rather than the much-needed saturated red emission.

### 3. Conclusion

In summary, we report detailed syntheses and luminescence properties of red-light- (1) and green-light- (2) emitting $\text{Pt}^{II}$ metal complexes using a bulky ligand incorporating a camphor-derived moiety. These complexes exhibit high emission quantum yields, short phosphorescence radiative lifetimes in the range of several microseconds and, more importantly, a much lower tendency to aggregation than previously reported $\text{Pt}^{II}$ porphyrinato or $\beta$-diketonato complexes for electroluminescent applications.\textsuperscript{[2]} Remarkable improvement of the device performance has been achieved at higher dopant concentrations or even in a pure emission layer, constituting for the first time a highly efficient $\text{Pt}^{II}$-based OLED in the red.

### 4. Experimental

**General Procedures:** All reactions were performed under nitrogen. Solvents were distilled from appropriate drying agents prior to use. Commercially available reagents were used without further purification unless otherwise stated. All reactions were monitored by thin-layer chromatography (TLC) with Merck pre-coated glass plates (0.20 mm with fluorescent indicator UV$_{365}$). Compounds were visualized with UV light irradiation at 254 nm and 365 nm. Flash column chromatography was carried out using silica gel from Merck (230–400 mesh). Mass spectra were obtained on a JEOL SX-102A instrument operating in electron impact (EI) or fast atom bombardment (FAB) mode. $^1$H and $^13$C NMR spectra were recorded on a Bruker-400 or INOVA-500 instrument; chemical shifts are quoted with respect to the internal standard tetramethylsilane for $^1$H and $^13$C NMR data. Elemental analysis was carried out with a Heraeus CHN-O Rapid Elementary Analyzer.

**Spectroscopic and Dynamic Measurements:** Steady-state absorption and emission spectra were recorded on a Hitachi (U-3310) spectrophotometer and an Edinburgh (FS920) fluorimeter, respectively. Both wavelength-dependent excitation and emission response of the fluorimeter were calibrated. A configuration of front-face excitation was used to measure the emission of the solid sample, in which the cell was made
by assembling two edge-polished quartz plates with various Teflon spacers. A combination of appropriate filters was used to avoid interference from the scattering light. Lifetime studies were performed by an Edinburgh F900000 photon-counting system with a hydrogen-filled lamp or a nitrogen lamp as the excitation source. Data were analyzed using a nonlinear least squares procedure in combination with an iterative convolution method. The emission decays were analyzed by the sum of exponential functions, which allows partial removal of the instrument time broadening and consequently renders a temporal resolution of ~200 ps.

To determine the photoluminescence quantum yield in solution, samples were degassed by three freeze–pump–thaw cycles under vigorous stirring conditions. 4-(Dicyanomethylene)-2-methyl-6-(p-dimethylaminostyryl)-4H-pyran (DCM, $\lambda_{\text{em}} = 615$ nm, Exciton, Inc.) in methanol was used as a reference, assuming a quantum yield of 0.43 with 430 nm excitation [24]. An integrating sphere (Labsphere) was applied to measure the quantum yield in the solid state, in which the solid-sample film was prepared via either spin-coating or vapor-deposition methods and was excited by a 514 nm (complex 1) or 457 nm (complex 2) Ar$^+$ laser line. The resulting luminescence was led to an intensified charge-coupled detector for subsequent quantum yield analyses to obtain the PL quantum yield in the solid state, the emission was collected via an integrating sphere, and the quantum yield was calculated according to a reported method [25].

Synthesis of 4,8,8-Trimethyl-3-isoxazolin-1-yl-4,5,6,7-tetrahydro-2H-4,7-methano-indazole, (iqdz)$_2$: To a stirred mixture of NaH (0.26 g, 10.8 mmol) and tetrahydrofuran (THF, 10 mL) at 0 °C was added a solution of (1H)-cyclophosphorin (1.64 g, 10.8 mmol) in THF for a period of 30 min. The temperature of the reaction mixture was slowly increased to RT and stirring was continued for about 30 min. The solution was heated to 60 °C, and ethyl 1-isoxazolinocarboxylate (1.7 g, 8.5 mmol) in THF was added slowly and refluxed for about 3 h. After this period, the reaction mixture was cooled to 0 °C and quenched with dilute HCl to pH 8–9. Then it was extracted with ethyl acetate (2 × 100 mL), and the extracts were washed with brine and water, dried over anhydrous MgSO$_4$ and concentrated in vacuo to give a yellow oil (2.2 g). Hydrazine monohydrate (4.2 mL, 86.0 mmol) in EtOH was added dropwise to a refluxing solution of the above oil (2.2 g), without further purification, in EtOH (30 mL). After the mixture was refluxed for 12 h, the solvent was removed under vacuum. The residue obtained was dissolved in ethyl acetate and washed with water, dried over anhydrous MgSO$_4$, and concentrated again. The residue obtained was passed through a silica gel column using mixtures of hexane and ethyl acetate as eluents to give (iqdz)$_2$ as colorless crystals (1.4 g, 65 %).

**Table 2.** Performance characteristics for the electrophosphorescence devices based on complex 1.

<table>
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<tr>
<th>Doping conc. [%]</th>
<th>Brightness $B_{\text{max}}$ [cd/m$^2$]</th>
<th>$\tau_{\text{tint}}$ [µs]</th>
<th>Lumin. eff. $\eta$ [%]</th>
<th>Power eff. $\eta_{\text{power}}$ [W/W]</th>
<th>$V_{\text{opt}}$ [V]</th>
<th>$V_{\text{max}}$ [V]</th>
<th>$\lambda_{\text{max}}$ [nm]</th>
<th>CIE (x, y)</th>
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<tr>
<td>6</td>
<td>3451 [a]</td>
<td>10.6</td>
<td>17.4</td>
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<td>8.61</td>
<td>61/76</td>
<td>0.61, 0.38</td>
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<tr>
<td>12</td>
<td>10846 [a]</td>
<td>6.6</td>
<td>10.92</td>
<td>3.24</td>
<td>10.61</td>
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<tr>
<td>33193 [15 (c)]</td>
<td>14.9 (5.5)</td>
<td>24.75 (5.5)</td>
<td>14.86 (5.0)</td>
<td></td>
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<td>20</td>
<td>2378 [a]</td>
<td>8.2</td>
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<td>63/92, 0.64, 0.35</td>
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<td>10733 [15 (c)]</td>
<td>3.91 (4.5)</td>
<td>4.22 (4.5)</td>
<td>2.95 (4.5)</td>
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</table>

[a] Values collected at 20 mA/cm$^2$. [b] Values collected at 100 mA/cm$^2$. [c] Maximum values of the devices. Values in parentheses are the voltages at which they were obtained.

**Spectral Data of Complex 1**: emission peak at 617.1 nm, $\lambda_{\text{max}} = 617.1$ nm, excitation peak at 585.5 nm, $\lambda_{\text{max}} = 585.5$ nm, and the full width at half maximum (FWHM) is 64.0 nm.

**Spectral Data of Complex 2**: emission peak at 617.2 nm, $\lambda_{\text{max}} = 617.2$ nm, excitation peak at 585.5 nm, $\lambda_{\text{max}} = 585.5$ nm, and the full width at half maximum (FWHM) is 64.0 nm.
The crystallographic data of this complex (excluding structure factors) have been deposited at the Cambridge Crystallographic Data Centre with the deposition number CCDC-238230. The data can be obtained free of charge on application to CCDC, 12 Union Road, Cambridge CB2 1EZ, UK (fax: (+44) 1223 336 033; e-mail: deposit@ccdc.cam.ac.uk).

**OLED Fabrication:** Charge transporting materials such as NPB (4,4′-bis[N-(1-naphthyl)-N-phenylamino]biphenyl) and Alq3 (tris[8-hydroxyquinolinato]aluminum) [11], as well as the host material C60 (4,4′,4′′-tri(N-carbazolyl)-1,1′,1″-biphenyl) were synthesized according to literature procedures [26], and were sublimed twice through a temperature-gradient sublimation system before use. BCP (2,9-dimethyl-4,7-di-phenyl-1,10-phenanthroline) was obtained from Aldrich. Patterned ITO-coated glass substrates (sheet resistance \( \leq 30 \Omega \) in a square with an effective individual device area of 3.14 mm²) were cleaned by sonication in a detergent solution, water, and ethanol, separately, and then dried by a flow of nitrogen. The substrates were further treated with oxygen plasma for 5 min before being loaded into the vacuum chamber. Various organic layers were deposited sequentially at a rate of 0.1-0.3 Å/minute under a pressure of 2 x 10⁻⁸ torr in an UvClay Cyrogenic deposition system. Phosphorescent dopants were co-evaporated with C60 via two independent sources. A thin layer of LiF (1 nm) and a thick layer of Al (150 nm) followed as the cathode. The current-voltage-illumination of the devices was measured in ambient conditions with a Keithley 2400 Source meter and a Newport 1835C Optical meter equipped with an 818ST silicon photodiode. The EL spectrum was obtained using a Hitachi F4500 spectrophotometer. The active area of the device was 3.14 mm² and that of the silicon photodiode was 100 mm². The device was placed close to the photodiode such that all the forward light went to the photodiode. The external quantum efficiency was calculated according to the method described before [27]. The luminous flux \( \mathcal{L} \) is defined [28] by

\[
P_\mathcal{L} = K_{\text{lum}} P_{\text{sc}} \mathcal{L}(\lambda) d\lambda
\]

where \( K_{\text{lum}} \) is the maximum luminous efficiency (683 lm/W), \( P_{\text{sc}} \) is the spectral distribution of radiant flux \( \mathcal{L}(\lambda) \), \( P_{\text{sc}} \) is the relative photopic luminance efficiency function; the luminance \( \text{cd/m}^2 \) is defined by \( P_{\text{sc}}/\Delta \text{A} \), where \( \Delta \text{A} \) is the device area; the luminous efficiency \( \text{cd/A} \) is defined by \( P_{\text{sc}}/\Delta \text{A} \), where \( I \) is the current; power efficiency is defined as \( P_{\text{sc}}/V \), where \( V \) is the applied voltage.

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