Room-temperature NIR phosphorescence of new iridium (III) complexes with ligands derived from benzoquinoxaline¹

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Abstract: A new series of new iridium (III) complexes (**1–5**) bearing ligands derived from benzoquinoxaline were designed and synthesized. X-ray structural analyses of **1** reveal a distorted octahedral geometry around the Ir atom in which the pyrazolate chelate is located opposite to the cis-oriented carbon donor atoms of benzoquinoxaline, while the benzoquinoxaline ligands adopt an eclipse configuration and their coordinated nitrogen atoms and carbon adopt trans- and cis-orientation, respectively. Complexes **1–5** exhibit moderate NIR phosphorescence with peak maxima located at around 910–930 nm. As supported by the TDDFT approach, the transition mainly involves benzoquinoxaline $^3\pi-\pi^*$ intraligand charge transfer (ILCT) and metal (Ir) to benzoquinoxaline charge transfer (MLCT) of which the spectroscopy and dynamics of relaxation have been thoroughly investigated. The relatively weak NIR emission can be tentatively rationalized by the low energy gap of which the radiationless deactivation may be governed by nearly temperature-independent, weak-bonding motions in combination with a minor channel incorporating small torsional motions associated with phenyl ring in the benzoquinoxaline sites.

Key words: phosphorescence, NIR, iridium, benzoquinoxaline, isoquinoline, bipyridine, pyrazolate, acetylacetonate.

Résumé : On a préparé une nouvelle série de complexes de l'iridium(III) (**1–5**) portant des ligands dérivés de la benzoquinoxaline. L'analyse des données structurales du composé **1**, obtenues par diffraction des rayons X, révèle la présence d'une géométrie octaédrique déformée autour de l'atome d'iridium dans laquelle le chélate pyrazolate se trouve à l'opposé des atomes de carbone donneurs de la benzoquinoxaline orientés en cis, alors que les ligands benzoquinoxaline adoptent une conformation éclipse et que leurs atomes d'azote coordonnés adoptent respectivement des orientations cis et trans. Les complexes **1–5** présentent de la phosphorescence modérée dans le proche infrarouge (PIR) avec un pic maximal situé autour de 910 à 930 nm. Tel que le prédit l'approche TDDFT, la transition implique principalement un transfert de charge interligand (TCIL) $^3\pi-\pi^*$ de la benzoquinoxaline et un transfert de charge du métal (iridium) à la benzoquinoxaline (TCML) dont on a étudié à fond la spectroscopie et la dynamique de relaxation. L'émission relativement faible dans le proche infrarouge pourrait être rationalisée en fonction d'une discontinuité de faible énergie de laquelle la désactivation sans rayonnement pourrait être orientée par des mouvements de faible liaison, pratiquement indépendants de la température, en combinaison avec une voie mineure qui incorporerait de faibles mouvements de torsion du noyau phényle dans les sites benzoquinoxalines.

Mots clés : phosphorescence, PIR, iridium, benzoquinoxaline, isoquinoléine, bipyridine, pyrazolate, acétylacétonate.

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This paper is dedicated to Professor Arthur J. Carty; he has been our mentor, collaborator, and friend for the past 15 years.

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Introduction

Transition metal complexes, which exhibit strong emission in fluid solution at room temperature, have been extensively investigated owing to their fundamental interest and key roles in the design of novel systems for solar energy conversion (1), telecommunication and other optical technologies (2), as well as fabrication of high efficiency phosphorescent organic light emitting devices (OLEDs) (3). So far, most of the reported luminescent metal complexes emit in the visible region, while studies of emission properties in the NIR region are relatively rare. One reason lies in that decreasing the energy gap toward NIR may trigger a much faster radiationless decay channel because of the interaction between ground and emitting states, i.e., the so-called energy gap law, giving rise to poor quantum efficiencies (4).

The first class of complexes showing notable NIR luminescence should be credited to the lanthanide complexes incorporating the Nd(III), Yb(III), and Er(III) ions (5). Sharp emission signals for lanthanide complexes are expected simply because the 4f orbitals in the lanthanide elements are buried deep within the atom and would not be seriously coupled with the ligand vibration motions, as opposed to the broad d–d transitions of the transition metals. As a second class, the Pd porphyrinate types of complexes are documented as exhibiting moderate NIR phosphorescence in the fluid state at room temperature, a property which is desirable for application as oxygen sensors for biomedical applications *in vivo* (6). The third class consists of the polypyridyl complexes involving the Re(I), Ru(II), and Os(II) metal ions with the d⁶ electronic configuration (7). Their systematic synthesis has remained as the outstanding issue in view of applications. These complexes typically absorb appreciably in the high energy region of the spectra, accompanied by the MLCT absorptions at lower energy, as well as a certain degree of contribution from direct ⁵MLCT absorption, a result of the heavy atom enhanced spin-orbit coupling. Accordingly, NIR phosphorescent emissions are achievable for systems with extended electronic delocalization between the metal and ligands in both ground and excited states, making them important as light harvesting sensitizers for solar cell applications.

As an extension of our contribution to the third-row metal phosphorescent materials, we have embarked on the development of Ir(III)-based metal complexes showing detectable luminescence in the NIR spectral region. The key designing feature is to utilize a diphenyl substituted benzoquinoline ligand that allows formation of a highly extended π conjugation. The choice of Ir as the central atom renders the advantages of linking the benzoquinoline ligand through facile orthometalation, affording a desirable metal–chelate bonding interaction and a presumably rigid molecular skeleton. Along this line, the emission gaps are expected to shift significantly toward the lower energy, i.e., the NIR region, without drastic reduction to the corresponding emission efficiency.

Experimental section

General experiments

The pyrazole ligands, 3-trifluoromethyl-5-(2-pyridyl)pyrazole (fppz)H and 1-(5-*tert*-butyl-2H-pyrazol-3-yl)isoquino-

line (bpiq)H, were prepared according to the literature methods (8). Commercially available chemicals were used without further purification. All reactions were monitored by TLC with Merck silica gel plates (0.20 mm with fluorescent indicator UV₂₅₄). Compounds were visualized with 254 or 365 nm UV irradiation. Column chromatography was carried out using silica gel purchased 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. ¹H and ¹³C NMR spectra were recorded on a Varian INOVA-500 instrument; chemical shifts are quoted with respect to the internal standard tetramethylsilane for ¹H and ¹³C NMR data. Elemental analysis was carried out with a Heraeus CHN-O rapid elementary analyzer.

Cyclic voltammetry (CV)

CV measurements were performed using a BAS 100 B/W electrochemical analyzer. The oxidation and reduction measurements were recorded, respectively, in anhydrous CH₂Cl₂ and anhydrous THF containing 0.1 mol/L TBAPF₆ as the supporting electrolyte, at a scan rate of 100 mV s⁻¹. The potentials were measured against a Ag/Ag⁺ (0.01 mol/L AgNO₃) reference electrode with ferrocene as the internal standard.

Preparation of 2,3-diphenyl benzo[g]quinoxaline

An anhydrous ethanol solution (30 mL) of 2,3-naphthalenediamine (1.0 g 6.33 mmol) and benzil (1.34 g, 6.33 mmol) was heated to reflux for 5 h. Then the solution was gradually cooled to room temperature. The solid deposited from solution was then filtered, washed with a few drops of ethanol, and dried in vacuum overnight to afford 1.8 g of yellow-green 2,3-diphenyl benzo[g]quinoxaline (5.42 mmol, 86%), denoted as (dpbq)H. The yellow-colored derivative 2,3-di-*p*-tolyl benzo[g]quinoxaline, (dtbq)H, was obtained from 4,4'-dimethylbenzil in a similar fashion.

Spectral data for (dpbq)H

¹H NMR (400 MHz, CDCl₃, 298 K) δ : 8.72 (s, 2H), 8.07 (dd, 2H, $J_{\text{HH}} = 6.4, 2.8$ Hz), 7.56 (d, 4H, $J_{\text{HH}} = 7.2$ Hz), 7.52 (dd, 2H, $J_{\text{HH}} = 6.4, 2.8$ Hz), 7.38–7.32 (m, 6H). MS (EI) m/z : 332 [M⁺]. Anal. calcd. for C₂₄H₁₆N₂: N 8.43, C 86.72, H 4.85; found: N 8.48, C 86.63, H 5.05.

Spectral data for (dtbq)H

¹H NMR (400 MHz, CDCl₃, 298 K) δ : 8.69 (s, 2H), 8.09–8.06 (m, 2H), 7.54–7.51 (m, 2H), 7.46 (d, 4H, $J_{\text{HH}} = 10$ Hz), 7.15 (d, 4H, $J_{\text{HH}} = 10$ Hz), 2.37 (s, 6H). MS (EI) m/z : 360 [M⁺]. Anal. calcd. for C₂₆H₂₀N₂: N 7.77, C 86.64, H 5.59; found: N 7.41, C 86.65, H 5.26.

Preparation of [(dpbq)₂IrCl]₂

The chloride-bridged complex [(dpbq)₂IrCl]₂ was synthesized according to the method reported by Nonoyama (9), involving treatment of IrCl₃·3H₂O (1.0 g, 2.84 mmol) with (dpbq)H (2.83 g, 8.5 mmol) in 2-methoxyethanol (30 mL) for 24 h. Then the solution was treated with water (30 mL) to induce precipitation of a dark brown solid. The solid was then filtered, washed with methanol, and dried under vacuum (1.8 g, 1.22 mmol, 71%).

Preparation of [(dtbq)₂IrCl]₂

On the basis of a procedure similar to that described for [(dpbq)₂IrCl]₂, the *p*-tolyl-substituted complex [(dtbq)₂IrCl]₂ was synthesized in ~75% yield.

Preparation of [(dpbq)₂Ir(fppz)] (1)

A mixture of [(dpbq)₂IrCl]₂ (300 mg, 0.168 mmol), 3-trifluoromethyl-5-(2-pyridyl)pyrazole (fppzH, 80 mg, 0.38 mmol), and Na₂CO₃ (100 mg, 0.94 mmol) in 2-methoxyethanol (20 mL) was heated to reflux for 4 h. An excess of deionized water was added after cooling the solution to room temperature (RT). The precipitate was collected by filtration and washed with methanol (10 mL), followed by diethyl ether (10 mL). Black crystals of [(dpbq)₂Ir(fppz)] (1) were obtained from a mixture of CH₂Cl₂ and methanol at RT (210 mg, 0.19 mmol, 58%).

Spectral data for 1

¹H NMR (400 MHz, CDCl₃, 298 K) δ: 8.77 (s, 1H), 8.64 (d, 1H, *J*_{HH} = 5.6 Hz), 8.54 (s, 1H), 8.49 (s, 1H), 8.05 (s, 1H), 7.94–7.88 (m, 4H), 7.75 (d, 1H, *J*_{HH} = 8.0 Hz), 7.72–7.61 (m, 5H), 7.61–7.55 (m, 3H), 7.45–7.38 (m, 4H), 7.35 (t, 1H, *J*_{HH} = 6.0 Hz), 7.27–7.32 (m, 2H), 7.25–7.20 (m, 3H), 6.78 (d, 1H, *J*_{HH} = 7.6 Hz), 6.72 (t, 1H, *J*_{HH} = 8.0 Hz), 6.65 (t, 1H, *J*_{HH} = 7.6 Hz), 6.60 (t, 1H, *J*_{HH} = 7.2 Hz), 6.50–6.47 (m, 2H), 6.36 (d, 1H, *J*_{HH} = 7.2 Hz). MS (FAB, ¹⁹³Ir) *m/z*: 1067 [M⁺]. Anal. calcd. for C₅₇H₃₅F₃IrN₇: N 9.19, C 64.15, H 3.31; found: N 9.09, C 64.21, H 3.62.

Preparation of [(dpbq)₂Ir(bpiq)] (2)

Procedures identical to that of 1 were followed, using 200 mg of [(dpbq)₂IrCl]₂ (0.112 mmol), 71 mg of (bpiq)H (0.28 mmol), and 60 mg of Na₂CO₃ (0.57 mmol) in 20 mL of 2-methoxyethanol. Black crystals of [(dpbq)₂Ir(bpiq)] (2) were obtained from a mixture of CH₂Cl₂ and hexane at RT (74 mg, 0.085 mmol, 33%).

Spectral data for 2

¹H NMR (400 MHz, CDCl₃, 298 K) δ: 9.00 (s, 1H), 8.50–8.43 (m, 5H), 8.30 (s, 1H), 7.97 (b, 2H), 7.87 (d, 2H, *J*_{HH} = 8.0 Hz), 7.82 (d, 1H, *J*_{HH} = 8.0 Hz), 7.72 (d, 1H, *J*_{HH} = 8.0 Hz), 7.69–7.58 (m, 7H), 7.52 (t, 1H, *J*_{HH} = 7.2 Hz), 7.44–7.28 (m, 4H), 7.26–7.18 (m, 5H), 6.74–6.66 (m, 2H), 6.64–6.53 (m, 3H), 6.45 (t, 1H, *J*_{HH} = 7.6 Hz), 6.31 (d, 1H, *J*_{HH} = 8.0 Hz), 1.37 (s, 9H). MS (FAB, ¹⁹³Ir) *m/z*: 1105 [M⁺]. Anal. calcd. for C₆₄H₄₆F₃IrN₇: N 8.87, C 69.54, H 4.19; found: N 8.79, C 69.48, H 4.56.

Preparation of [(dpbq)₂Ir(bpy)] (3)

A mixture of [(dpbq)₂IrCl]₂ (200 mg, 0.11 mmol), 2,2'-bipyridine (bpy, 45 mg, 0.29 mmol), and AgBF₄ (100 mg, 0.51 mmol) in CH₂Cl₂ (25 mL) was stirred at RT for 3 h. The solution was filtered and the filtrate was evaporated to give the crude product. Black crystals of [(dpbq)₂Ir(bpy)][BF₄] (3) were obtained from a mixture of CH₂Cl₂ and hexane at RT (110 mg, 0.09 mmol, 41%).

Spectral data for 3

¹H NMR (400 MHz, CDCl₃, 298 K) δ: 8.71 (d, 2H, *J*_{HH} = 4.8 Hz), 8.60 (s, 2H), 8.55 (d, 2H, *J*_{HH} = 8.8 Hz), 8.20 (t, 2H, *J*_{HH} = 7.6 Hz), 7.96 (d, 2H, *J*_{HH} = 8.0 Hz), 7.86–7.80

(m, 8H), 7.68–7.64 (m, 8H), 7.49 (t, 2H, *J*_{HH} = 3.6 Hz), 7.40 (t, 2H, *J*_{HH} = 7.6 Hz), 7.03 (d, 2H, *J*_{HH} = 8.0 Hz), 6.75 (t, 2H, *J*_{HH} = 7.6 Hz), 6.63 (t, 2H, *J*_{HH} = 7.2 Hz), 6.54 (d, 2H, *J*_{HH} = 7.6 Hz). MS (FAB, ¹⁹³Ir) *m/z*: 1011 [M⁺]. Anal. calcd. for C₅₈H₃₈F₄IrN₆: N 7.65, C 63.45, H 3.49; found: N 7.71, C 63.25, H 3.57.

Preparation of [(dtbq)₂Ir(fppz)] (4)

Procedures identical with that of 1 were followed, using 200 mg of [(dtbq)₂IrCl]₂ (0.106 mmol), 113 mg of (fppz)H (0.53 mmol), and 100 mg of Na₂CO₃ (0.94 mmol) in 20 mL of 2-methoxyethanol. Black crystals of [(dtbq)₂Ir(fppz)] (4) were obtained from a mixture of CH₂Cl₂ and methanol at RT (120 mg, 0.107 mmol, 50%).

Spectral data for 4

¹H NMR (400 MHz, CDCl₃, 298 K) δ: 8.69 (s, 1H), 8.57 (d, 1H, *J*_{HH} = 8.0 Hz), 8.49 (s, 1H), 8.44 (s, 1H), 7.98 (s, 1H), 7.91 (d, 1H, *J*_{HH} = 8.0 Hz), 7.86 (d, 1H, *J*_{HH} = 8.0 Hz), 7.81 (d, 2H, *J*_{HH} = 5.2 Hz), 7.72 (d, 1H, *J*_{HH} = 8.0 Hz), 7.54 (t, 1H, *J*_{HH} = 1.2 Hz), 7.50–7.41 (m, 5H), 7.40 (t, 3H, *J*_{HH} = 1.2 Hz), 7.31–7.24 (m, 4H), 7.20–7.12 (m, 3H), 6.63 (s, 1H), 6.57 (d, 1H, *J*_{HH} = 8.0 Hz), 6.48 (d, 1H, *J*_{HH} = 8.0 Hz), 6.44 (s, 1H), 6.20 (s, 1H), 2.54 (s, 3H), 2.51 (s, 3H), 1.86 (s, 3H), 1.79 (s, 3H). MS (FAB, ¹⁹³Ir) *m/z*: 1123 [M⁺]. Anal. calcd. for C₆₁H₄₃F₃IrN₇: N 8.73, C 65.23, H 3.86; found: N 8.64, C 64.89, H 3.81.

Preparation of [(dtbq)₂Ir(acac)] (5)

A mixture of [(dtbq)₂IrCl]₂ (200 mg, 0.106 mmol), acetylacetone (acacH, 300 mg, 3.0 mmol), and Na₂CO₃ (100 mg, 0.94 mmol) in CH₂Cl₂ (20 mL) was stirred at RT for 24 h. After stopping the reaction, the solution was washed with water (3 × 50 mL), dried over MgSO₄, and concentrated to dryness. Black crystals of [(dtbq)₂Ir(acac)] (5) were recrystallized from CH₂Cl₂ and hexane at RT (90 mg, 0.89 mmol, 42%).

Spectral data for 5

¹H NMR (400 MHz, CDCl₃, 298 K) δ: 8.84 (s, 2H), 8.65 (s, 2H), 8.03 (d, 2H, *J*_{HH} = 8.0 Hz), 7.94–7.89 (m, 6H), 7.49–7.42 (m, 8H), 7.06 (d, 2H, *J*_{HH} = 8.4 Hz), 6.41 (d, 2H, *J*_{HH} = 8.4 Hz), 6.32 (s, 2H), 4.63 (s, 1H), 2.52 (s, 6H), 1.79 (s, 6H), 1.61 (s, 6H). MS (FAB, ¹⁹³Ir) *m/z*: 1010 [M⁺]. Anal. calcd. for C₅₇H₄₅IrN₄O₂: N 5.55, C 67.77, H 4.49; found: N 5.51, C 67.63, H 4.83.

X-ray structural analysis

Single crystal X-ray diffraction data were measured on a Bruker Smart CCD diffractometer using (Mo Kα) radiation (λ = 0.710 73 Å). The data collection was executed using the SMART program. Cell refinement and data reduction were made with the SAINT program. The structure was determined using the SHELXTL/PC program and refined using full-matrix least-squares. All non-hydrogen atoms were refined anisotropically, whereas hydrogen atoms were placed at the calculated positions and included in the final stage of refinements with fixed parameters. The crystallographic refinement parameters for complexes 1 and 3 are summarized in Table 1, and selected bond distances and angles of these complexes are listed in Tables 2 and 3, respectively.

Table 1. Crystal data and refinement parameters for complexes **1** and **3**.

Compound	1	3
Formula	C ₅₇ H ₃₅ F ₃ N ₇ Ir·CH ₂ Cl ₂	C ₅₈ H ₃₈ N ₆ Ir·BF ₄ ·CH ₂ Cl ₂
Molecular weight	1 152.05	1 182.88
Crystal system	Triclinic	Triclinic
Space group	<i>P</i> -1	<i>P</i> -1
Crystal size (mm ³)	0.4×0.15×0.12	0.18×0.15×0.05
<i>a</i> (Å)	11.9568(4)	16.3626(6)
<i>b</i> (Å)	12.0739(5)	17.2949(6)
<i>c</i> (Å)	17.1667(7)	19.5430(7)
α (°)	74.820(1)	73.631(1)
β (°)	82.535(1)	72.197(1)
γ (°)	84.788(1)	70.003(1)
<i>V</i> (Å ³)	2 367.4(2)	4 850.1(3)
<i>Z</i>	2	4
ρ_{calc} (g cm ⁻³)	1.616	1.620
Temp (K)	295(2)	150(1)
μ (mm ⁻¹)	2.994	2.927
Reflections collected	31 690	64 361
Independent reflections	10 878 (<i>R</i> (int) = 0.0419)	22 242 (<i>R</i> (int) = 0.0680)
<i>R</i> _F , <i>R</i> _w (<i>F</i> ²) (all data)	0.0445, 0.0928	0.0873, 0.1475
<i>R</i> _F , <i>R</i> _w (<i>F</i> ²) [<i>I</i> > 2σ(<i>I</i>)]	0.0360, 0.0860	0.0606, 0.1319
GOF	1.043	1.144

Note: $R_F = \sum |F_o - F_c| / \sum F_o$ and $R_w(F^2) = [\sum w(F_o^2 - F_c^2)^2 / \sum w(F_o^4)]^{1/2}$.

Table 2. Selected bond lengths (Å) and angles (°) for complex **1**.

Bond lengths (Å)	
Ir—N(1)	2.201(3)
Ir—N(4)	2.065(3)
Ir—C(10)	1.990(4)
C(25)···N(2)	3.013(7)
C(49)···N(1)	3.165(6)
Ir—N(2)	2.116(3)
Ir—N(6)	2.047(3)
Ir—C(34)	1.991(4)
C(25)···N(3)	3.381(6)
C(49)···C(1)	3.439(6)
Bond angles (°)	
N(1)—Ir—N(2)	75.02(13)
N(6)—Ir—C(34)	79.47(15)
N(2)—Ir—C(10)	174.19(14)
N(4)—Ir—C(10)	79.21(15)
N(1)—Ir—C(34)	170.85(14)
N(4)—Ir—N(6)	178.16(12)

Table 3. Selected bond lengths (Å) and angles (°) for complex **3**.

Bond lengths (Å)	
Ir(1)—N(1)	2.062(6)
Ir(1)—N(5)	2.171(6)
Ir(1)—C(1)	1.992(7)
C(23)···N(6)	2.995(8)
C(47)···N(5)	3.146(9)
Ir(1)—N(3)	2.076(5)
Ir(1)—N(6)	2.160(6)
Ir(1)—C(25)	1.999(6)
C(23)···C(58)	3.336(8)
C(47)···C(49)	3.438(8)
Bond angles (°)	
N(1)—Ir(1)—C(1)	79.4(3)
N(5)—Ir(1)—N(6)	75.2(2)
N(6)—Ir(1)—C(1)	171.7(2)
N(3)—Ir(1)—C(25)	79.3(2)
N(1)—Ir(1)—N(3)	173.7(2)
N(5)—Ir(1)—C(25)	174.9(2)

The crystallographic data for complexes **1** and **3** (excluding structure factors) have been deposited.⁵

Spectral and dynamic measurement

Steady-state absorption and emission spectra were re-

corded with a Hitachi (U-3310) spectrophotometer. The NIR emission spectra were acquired by exciting the sample solution under a rectangular excitation configuration using an Ar ion laser (488 or 514 nm, Innova 90, Coherent Inc., Santa Clara, California) and the continuous wave was chopped by

⁵Supplementary data for this article are available on the journal Web site (<http://canjchem.nrc.ca>) or may be purchased from the Depository of Unpublished Data, Document Delivery, CISTI, National Research Council Canada, Ottawa, ON K1A 0R6, Canada. DUD 5002. For more information on obtaining material refer to http://cisti-icist.nrc-cnrc.gc.ca/irm/unpub_e.shtml. CCDC 276875 and 276876 contain the crystallographic data for this manuscript. These data can be obtained, free of charge, via <http://www.ccdc.cam.ac.uk/conts/retrieving.html> (Or from the Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax +44 1223 336033; or deposit@ccdc.cam.ac.uk).

a mechanic chopper operated at 200 Hz. To minimize the re-absorption effect, the excitation beam was focused as closely as possible to the front wall of the quartz cell. The NIR emission was measured by the photomultiplier tube (Hamamatsu R5509-72, operated at $-80\text{ }^{\circ}\text{C}$) coupled with a Lock-in amplifier (SR830, Stanford Research Systems, Sunnyvale, California) and a monochromator (SpectraPro-275, Acton Research Corporation, Acton, Massachusetts) in that the grating was blazed at 1200 nm. IR125 (Exciton Inc., Dayton, Ohio) with $\Phi \sim 0.11$ ($\lambda_{\text{max}} \sim 925\text{ nm}$) in DMSO solution was used as a standard to calculate the emission quantum yield (10). For the NIR lifetime measurement, the sample was excited by a 532 nm pulse (2nd harmonic of Nd:YAG, 6 ns). The emission trace was detected by a photomultiplier tube (Hamamatsu R5509-72) operated at $-80\text{ }^{\circ}\text{C}$ and recorded by a sampling oscilloscope (TDS 3012, Tektronix, Beaverton, Oregon). An average of 100 shots was taken in the lifetime measurement, unless specified elsewhere.

Computational methodology

Calculations on the electronic ground state of complexes **1** and **3** were carried out using B3LYP density functional theory (11, 12). A double- ζ quality basis set consisting of Hay and Wadt's effective core potentials (LANL2DZ) (13) was employed for the Ir atom and a 6-31G* (14) basis for the H, C, N, and F atoms. A relativistic effective core potential (ECP) replaced the inner core electrons of Ir(III), leaving the outer core ($5s^25p^6$) electrons and the $5d^6$ valence electrons. Time-dependent DFT (TDDFT) calculations (15) using the B3LYP functional were then performed based on the structural optimized geometries. Typically, the lowest 10 triplet and 10 singlet roots of the nonhermitian eigenvalue equations were obtained to determine the vertical excitation energies. Oscillator strengths were deduced from the dipole transition matrix elements (for singlet states only). The ground-state B3LYP and excited-state TDDFT calculations were carried out using Gaussian03 (16).

Results and discussion

Syntheses and structural characterization

The preparation of highly conjugated ligand chromophores is necessary to achieve the NIR-emitting iridium complexes. As depicted in Scheme 1, the required ligands can be synthesized by treatment of 2,3-naphthalenediamine and dibenzil to produce 2,3-diphenyl- or di-*p*-tolyl-substituted benzo[*g*]quinoxaline in good yields. This synthetic approach follows the literature procedure utilized for the analogous diphenylquinoxaline, and is best prepared from related *o*-phenylenediamine and the corresponding α -diketone in refluxing ethanol (17).

Subsequent reactions of $\text{IrCl}_3 \cdot n\text{H}_2\text{O}$ with (dpbq)H or (dtbq)H in refluxing ethoxyethanol afforded chloride-bridged dimers $[(\text{dpbq})_2\text{IrCl}]_2$ or $[(\text{dtbq})_2\text{IrCl}]_2$, respectively. These dimer complexes can easily react with the pyrazole chelate ligand (fppz)H in the presence of Na_2CO_3 to afford black-colored complexes $[(\text{dpbq})_2\text{Ir}(\text{fppz})]$ (**1**) and $[(\text{dtbq})_2\text{Ir}(\text{fppz})]$ (**4**), where (fppz)H = 3-trifluoromethyl-5-(2-pyridyl)pyrazole. Similar reactions towards other bidentate chelates such as 1-(5-*tert*-butyl-2H-pyrazol-3-yl)isoquinoline ((bpiq)H), 2,2'-

Scheme 1.

Scheme 2.

bipyridine (bpy), and acetylacetone (acacH) yielded the corresponding Ir(III) complexes, $[(\text{dpbq})_2\text{Ir}(\text{bpiq})]$ (**2**), $[(\text{dpbq})_2\text{Ir}(\text{bpy})][\text{BF}_4]$ (**3**), and $[(\text{dtbq})_2\text{Ir}(\text{acac})]$ (**5**), respectively (see Scheme 2).

All iridium complexes were easily soluble in organic solvents such as CH_2Cl_2 , THF, and acetone, and the corresponding solutions showed negligible decomposition for a period of over 24 h, upon exposure to air. Routine characterizations were conducted using microanalysis, mass spectrometry, and ^1H NMR spectroscopy. Moreover, the fppz complex **1** was further characterized by single crystal X-ray diffraction analysis. Its ORTEP diagram is depicted in Fig. 1, while crucial bond distances and angles are listed in Table 1.

As depicted in Fig. 1, complex **1** reveals a distorted octahedral geometry around the Ir atom, consisting of two cyclometalated phenylbenzoquinoxaline fragments and one (2-pyridyl) pyrazolate ligand. The unique pyrazolate chelate is located opposite to the cis-oriented carbon donor atoms of benzoquinoxaline, while the benzoquinoxaline ligands adopt an eclipse configuration and their coordinated nitrogen atoms (N(4) and N(6)) and carbon atoms (C(10) and C(34)) adopt trans and cis orientation, respectively. This coordinative arrangement is akin to those of the parent phenyl pyridine ligands observed for the chloride-bridged dimer

Fig. 1. (a) ORTEP diagram of **1** with thermal ellipsoids shown at the 30% probability level; fluorine atoms of the CF₃ substituent were omitted for clarity. (b) Same ORTEP diagram with an emphasis on the spatial distortion of dpbq ligands.

complex [(ppy)₂Ir(μ-Cl)]₂ (**18**), the diketonate complex (ppy)₂Ir(acac) (**19**), and even the pyrazolate derivatives, suggesting that the incoming fppz ligand occupies the positions of the bridging chloride ligands in its starting material, [(dpbp)₂IrCl]₂. Moreover, the uncoordinated and cyclometalated phenyl groups show twisting angles of 40°–54°, as well as unusually large deformation angles of 23°–28° with respect to the middle benzoquinoxaline fragments. The deformation angles reveal an unfavorable steric interaction between the chelating ligands and can be gauged by the least-squares calculation using planes defined by the cyclometalated phenyl group and the middle C₆ hexagon of the benzoquinoxaline. It appears to us that the nonbonding repulsion between benzoquinoxaline and pyridyl pyrazolate ligands is the main driving force causing such a large distortion at the cyclometalated phenyl groups, as most of the interligand C...C and C...N nonbonding contacts listed in Table 2 are shorter than a limited value of ca. 3.4 Å, i.e., the sum of the van der Waals radii of the carbon atoms.

For a further comparison, the X-ray structural study on the cationic dipyrindine complex **3** was also examined. Comparison of the ORTEP diagram (Fig. 2) and the metric parameters (Table 3) with those of **1** shows no significant variation on the overall molecular architecture, except for the formation of an approximate C₂ rotational axis that bisects the bipyridine ligand and passes through the central Ir metal atom. This increasing molecular symmetry is consistent with the simplification of the ¹H NMR spectral pattern observed in solution at RT (see the Experimental section).

Electrochemistry

The electrochemical properties of these Ir complexes were investigated by CV using ferrocene as the internal standard, and the respective redox data are listed in Table 4. During the anodic scan, all iridium complexes exhibited a reversible oxidation wave with potentials in the region 0.50–0.77 V. It

is believed that the oxidation process mainly occurred at the metal center and, in part, with a slight contribution from the surrounding chelate ligands (**20**). Thus, the formal oxidation potential was strongly dependent on the electronic environment of the Ir(III) cation, i.e., a better electron donor would shift the oxidation potential to a less positive value. This hypothesis was verified by the increased π conjugation of the anionic bpiq ligand in **2**, which decreased the oxidative potential from 0.77 (in **1**) to 0.61 V (in **2**), while the attachment of the 2,2'-bipyridine ligand in **3** led to the complete suppression of the metal oxidation signal owing to its cationic nature. Moreover, replacement of the phenyl substituents with tolyl substituents of benzoquinoxaline also slightly altered the oxidation potential, as supported by the fact that the oxidation potential of **4** is cathodically shifted to 0.70 V in comparison with that of the parent complex **1**. Owing to the presence of an extra methyl substituent, this result could possibly be attributed to a slightly better σ-donor ability of the cyclometalated tolyl group.

Upon switching to the cathodic sweep in THF, two reversible reduction processes, with potentials ranging from –1.33 to –1.96 V, were detected for all complexes (**1**–**5**). We assigned these signals to a sequential reduction of the cyclometalated benzoquinoxaline fragments. For dtbq complex **4**, the first reduction peak is shifted to –1.65 V in comparison with that of the dpbq counterpart **1** (–1.54 V) because of the electron-releasing effect of the *para*-methyl substituents. Conversely, complex **3** showed the lowest reduction potentials, owing to its cationic nature, while complexes **2** and **5** gave the highest reduction potential among the whole series. The results indicate that the third ancillary ligand also significantly affects the redox properties of these Ir complexes, possibly via their enhanced donor capability as well as the extended π conjugation. Qualitatively, this observation is consistent with both the photophysical properties and theoretical predictions elaborated as follows.

Fig. 2. (a) ORTEP diagram of **3** with thermal ellipsoids shown at the 50% probability level. (b) Same ORTEP diagram with an emphasis on the spatial distortion of dpbq ligands.

Table 4. One-electron redox potentials of complexes **1–5**.

Complex	$E_{1/2}^{\text{ox}}$ (V) ^a	$E_{1/2}^{\text{red}}$ (V) ^a
1	0.77 [100] ^b	−1.54 [110], −1.81 [100]
2	0.61 [80]	−1.64 [100], −1.91 [100]
3	—	−1.31 [80], −1.61 [80]
4	0.70 [100]	−1.65 [110], −1.91 [120]
5	0.50 [90]	−1.73 [90], −1.96 [100]

^aPotential values referenced vs. Fc/Fc⁺.

^bData in square bracket denotes the $|E_{\text{pa}} - E_{\text{pc}}|$ values in mV.

Photophysical properties

Figure 3 shows the absorption and emission spectra of complexes **1–5** in CH₂Cl₂. As supported by CV measurements as well as the frontier orbital analyses (see the following section), it is reasonable to assign the first S₀–S₁ transition to absorption peaks around 500–600 nm ($\epsilon \sim 5 \times 10^3$ (mol/L)^{−1} cm^{−1}) to a metal to ligand (benzoquinoxaline part) charge transfer (MLCT) overlapping with the interligand phenyl to benzoquinoxaline transition. Due to a large molecular framework, the composition of higher electronic states is complicated, so that any corresponding assignments for complexes **1–5** may not be meaningful. Furthermore, the Ir-metal-enhanced spin-orbit coupling should result in nonnegligible absorptivity for the triplet transition, as indicated by a corresponding small absorption hump around 650–700 nm. However, absorption bands ascribed to the triplet manifolds could not be distinctly resolved mainly owing to their hiding inside the UV region of the strong singlet transitions. Detailed discussion regarding the transition in the triplet manifold will be given in the Theoretical approach section.

Figure 3 also depicts the emission spectra of complexes **1–5** in CH₂Cl₂. The spectral feature, quantum efficiency and relaxation dynamics (vide infra) of complexes **1–5** show concentration independence up to 10^{−3} mol/L, indicating a

Fig. 3. The absorption and emission spectra of complexes **1–5** in CH₂Cl₂. The peak intensities were normalized. $\lambda_{\text{ex}} = 514$ nm. The arrow indicates the peak for ¹O₂ ¹Δg(v' = 0) → Σ_g[−](v'' = 0) 1276 nm emission.

negligible aggregation effect in fluid solution. To reveal any differences in the spectral profile, the emission spectra were normalized at the peak wavelength. In comparison, for all complexes studied, only a slight variation was observed in view of the emission spectral feature and peak position. As listed in Table 5, the relaxation dynamics of **1–5** are O₂ concentration dependent. For example, the decay time of complex **1** was measured to be 286 ns in the degassed CH₂Cl₂, while it was decreased to 200 ns upon aeration. In the O₂ dependent study, in which the O₂ concentration was monitored by its vapor pressure filled in the vacuum line and deduced by Henry's law, a straight Stern–Volmer plot could be formulated (not shown here), and an oxygen-quenching rate constant of 1.7×10^9 (mol/L)^{−1} s^{−1} was deduced, which is nearly one-ninth of the diffusion-controlled rate of 2.0×10^{10} (mol/L)^{−1} s^{−1} calculated from the Stokes–Einstein equa-

Table 5. Photophysical properties of complexes **1–5** in CH₂Cl₂ at room temperature.

Complex	λ_{abs} (nm) (ϵ , 10 ³ (mol/L) ⁻¹ cm ⁻¹)	λ_{em} (nm)	Φ_{aer} ($\times 10^3$) ^a	τ_{aer} (ns)	τ_{deg} (ns)
1	535 (7), 491 (10), 414 (29)	916	6.48	200	286
2	575 (5), 464 (10), 414 (27)	922	9.53	195	276
3	540 (5), 484 (9), 412 (26)	922	3.59	245	403
4	541 (9), 491 (13), 422 (40)	916	10.2	209	302
5	570 (7), 508 (10), 421 (31)	904	21.8	216	290

^aQuantum efficiencies were measured in aerated CH₂Cl₂.**Fig. 4.** HOMO and LUMO of complex **1** involved in the low-lying transition. **Table 6.** The calculated lowest excited singlet and triplet states of complex **1**.

State	Assignment	λ (nm)	E (eV)	f
S ₁	HOMO → LUMO (+95%)	618.3	2.01	0.0067
T ₁	HOMO-1 → LUMO (+42%)	743.6	1.67	~0
	HOMO → LUMO (+37%)			
	HOMO-1 → LUMO+1 (23%)			
	HOMO-2 → LUMO (11%)			

tion in CH₂Cl₂. The result is consistent with the O₂-quenching triplet state according to the theory of electron-exchange type (Dexter type) of energy transfer expressed as (21):

The overall spin must be conserved upon forming a collisional complex. Accordingly, the possibility of each collision producing ¹O₂ is statistically one-ninth. Perhaps the strongest support for the O₂ quenching is given by the observation of ¹O₂ (¹Δ_g (v'' = 0) → ¹Σ_g⁺ (v' = 0)) emission with a peak wavelength at ~1276 nm (22) for complexes **1–5** in aerated CH₂Cl₂ (see the arrow in Fig. 3). The results, in combination with the rather small radiative decay rate of <10⁶ s⁻¹, lead us to unambiguously assign the dominant emission in complexes **1–5** to be of phosphorescence character. The spectra show a distinctly 0–0 vibronic onset of the transition, a profile that normally appears in a ³π–π* transition of aromatic chromophores with high symmetry. The results pro-

vide a clue that the observed phosphorescence possesses, in part, a ³π–π* character plausibly originating from a ligand-center chromophore of long conjugation such as benzoquinoxaline. Nevertheless, the involvement of other transitions such as ³MLCT is also possible. Furthermore, the observation of rather small changes in spectral features, as well as peak positions, implies that frontier orbitals responsible for the T₁–S₀ transition may be mainly derived from the same origin for complexes **1–5**. It is thus of fundamental importance to gain more detailed insights into the transition properties of the phosphorescence based on the theoretical approaches elaborated as follows.

Theoretical approach

Theoretical confirmation of the photophysical properties of complex **1** was provided by the DFT calculations. In this study, the crystallographic data for **1** was used as the framework for the extended Hückel formalism, and the calculation

was performed via a CACAO program (see the Experimental section). Figure 4 depicts the features of the HOMO to HOMO-3 and LUMO to LUMO+3 frontier orbitals mainly involved in the lower-lying transition, while the descriptions and the energy gaps of each transition are listed in Table 6. The validity of the TDDFT approach can be readily perceived by the calculated $T_1 \rightarrow S_0$ emission peak at 745 nm, which is very close to the 0-0 onset peak (~ 785 nm) observed experimentally. Based on the results (see Fig. 4 and Table 6), the electron densities of the HOMO are mainly located on the Ir(III) metal center and, to a small extent, the cyclometalated phenyl ligand, while a large portion of the electron densities of HOMO-1 and HOMO-2 is attributed to the π orbitals of benzoquinoxaline overlapping with certain d_π orbitals of the Ir(III) atom. Conversely, the electron densities of LUMO and LUMO+1 are distributed mostly on the benzoquinoxaline moiety. The $S_0 \rightarrow S_1$ and $S_0 \rightarrow S_2$ transitions mainly incorporate HOMO $\rightarrow$ LUMO (+95%) and HOMO $\rightarrow$ LUMO+1 (+93%), respectively, and thus are dominated by MLCT (d_π (Ir) $\rightarrow \pi^*$ (benzoquinoxaline)) overlapping with π - π^* ILCT (mainly cyclometalated phenyl site (π) $\rightarrow$ benzoquinoxaline pyridine site (π^*)) in character. As supported by the data shown in Table 6, the higher excited states in the singlet manifold, $S_0 \rightarrow S_n$ ($n > 3$), are relatively much more complicated, the results of which make a definitive frontier orbital assignment meaningless. In contrast, the lowest electronic transition in the triplet manifold, i.e., $S_0 \rightarrow T_1$, mainly incorporates the ligand center benzoquinoxaline $^3\pi$ - π^* transition and ^{3} MLCT (d_π (Ir) $\rightarrow \pi^*$ (benzoquinoxaline)).

The contribution of the $^3\pi$ - π^* transition is manifested by the low-temperature (77 K) approach to complex **1**. Figure 5 shows the luminescence spectrum of **1** in a 77 K CH_2Cl_2 matrix. Two salient remarks can be pointed out promptly according the spectral layout. First, in comparison to that measured at 298 K, the enhancement of the vibronic progression at 77 K is obvious, showing the notable population switching from perhaps ^{3} MLCT/ $^3\pi$ - π mixing character to the $^3\pi$ - π dominating state under this condition. Nevertheless, one cannot eliminate the possibility that the increase in vibronic resolution upon cooling may just indicate an increase in rigidity of the molecule, making the vibrational levels more identical to one another, as well as the disappearance of hot bands. Secondly, the phosphorescence intensity at 77 K is increased ~ 2.5 -fold with respect to that at 298 K. Similar structural phosphorescence with 2- to ~ 3 -fold increases in intensity was observed for complexes **2-5** in 77 K CH_2Cl_2 . Thus, temperature-dependent radiationless deactivation pathways may only play a minor role to the relaxation dynamics. Moreover, as listed in Table 5, substitution at non-benzoquinoxaline sites causes neither spectral change nor large alternation of the quantum efficiency. The latter implies that the large amplitude vibration or any torsional motion at non-benzoquinoxaline sites, which generally quenches emission in the NIR region, are not actively involved in the deactivation process. This viewpoint can be rationalized by the negligible contribution of non-benzoquinoxaline ligands to both HOMO (or HOMO-1 and HOMO-2) and LUMO (or LUMO+1). Since complexes **1-5** all bear aryl substituents at benzoquinoxaline sites, it is plausible that the torsional motion of phenyl (or tolyl) substituents, the relaxation dynam-

Fig. 5. The luminescence spectrum of **1** at room temperature (■) and in the 77 K CH_2Cl_2 matrix (Δ). Note that both spectra were analyzed using the same absorbance and excitation wavelength (514 nm). Thus, the intensity ratio can be directly estimated from the spectral integration.

ics of which should be temperature dependent, partly induces the radiationless transition. However, one ought to realize that because of the much smaller NIR energy gap, excitation that causes any weakening of the bond strength may result in crossing of the potential energy surface between T_1 and S_0 states and eventually trigger the rapid radiationless deactivation. In the case of complexes **1-5**, these nearly temperature-independent, weak vibrational motions play a key role in accounting for the dominant radiationless deactivation.

Conclusion

In summary, we report the syntheses and characterization of a new series of NIR Ir(III) complexes (**1-5**). These complexes exhibit moderate NIR phosphorescence with peak wavelengths around 910–920 nm at RT. As supported by the TDDFT approach, the transition mainly involves benzoisoquinoline $^3\pi$ - π^* intraligand charge transfer and metal (Ir) to benzoisoquinoline charge transfer. The relatively weak NIR emission can be tentatively rationalized by the low energy gap of which the radiationless deactivation may be governed by the large amplitude vibration and, in part, the phenyl torsional motions at the benzoquinoxaline sites. Knowing that current interest in Ir(III) types of phosphorescence emitter is focused on the Ir(III)-type OLED, to switch gears in research orientation, future applications of the titled NIR complexes and (or) relevant derivatives may be motivated by the strategy of using these complexes in the fabrication of photovoltaic devices (23). Focus on this approach is in progress.

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