Synthesis, crystal structure, photophysical property and metal ion-binding behavior of a cyclometalated platinum(II) terpyridylacetylide with efficient π-conjugation degree†

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A cyclometalated Pt(II) acetylide derivative bearing a carbazole donor and a terpyridine (TPY) receptor has been successfully synthesized and characterized. X-ray crystallography shows its monoclinic crystal system with the P2(1)/c space group and the three-component co-planarity molecular structure. This efficient π-conjugation system displays an intense low-energy absorption band (ε = 1.51 × 10^4 dm^3 mol^-1 cm^-1) at λ_{max} = 442 nm and a strong phosphorescence emission (Φ = 0.045) at λ_{max} = 596 nm in an air-saturated CH2Cl2 solution at room temperature, resulting from the dπPt → π*(C=N=N) metal-to-ligand charge transfer (MLCT) mixing with the π(C≡C–Ar) → π*(C=N=N) ligand-to-ligand charge transfer (LLCT) transition. With the introduction of the TPY receptor, this complex possesses ca. 1.6-fold luminescence-enhancing response for Zn^{2+} and Cd^{2+} but dramatic luminescence quenching response toward other common transition metal cations. Consecutive titrations exhibit that the added metal ions bond to its free TPY receptor with 1 : 2 stoichiometry to form the heterotrinuclear (Pt–M–Pt) complex. The Fe^{2+}-quenched and Zn^{2+}-enhanced luminescence behaviors with the maximum PET efficiency of 92.4% and 62.7%, respectively, strongly suggest the presence of the intra-molecular energy transfer between the [M(TPY)_3]^{2+} core and the terminal phosphorescent Pt(II) complex units.

Introduction

Luminescent chemosensors with high sensitivity and selectivity have been widely studied for their attractive potential in environmental and bio-medical applications.1,2 The “Chromophore–Spacer–Receptor” model has been generally designed, and the transition metal complexes with electronic transition in visible/near-IR region, high quantum efficiency and long lifetime have been extensively employed as fluorophores instead of the organic counterparts.3 Owing to their intriguing spectroscopic and luminescence properties, square-planar platinum(II) polypyridyl complexes imparted with metal-binding units, such as P- and N-donor crown ether pendants,4 S-benzo-15-crown-5,5 4-ethylbenzo-15-crown-5,6 5,17-diethynyl-25,27-dimethoxy-calix[4]crown-5,7 bipyridylacetylides,8 and terpyridylacetylides,9 have been reported in such investigations. Since the strong ligand field effect of the cyclometalated carbon raises the energy of the d–d states to reduce the non-radiation decay probability, cyclometalated platinum(II) complexes with tridentate phenyl-substituted pyridine [(N=C=N) motif]10–13 and bipyridine [(C=N=N) motif]14–22 as well as the related diphenyl-substituted pyridine [(C=N=C) motif]23,24 ligands are good emitters with high quantum yields at room temperature, which are expected to be used as better candidates in chemosensors.25,26 Cyclometalated [(C=N=N)Pt(µ)] σ-alkynyl complex with a benzo-15-azacrown-5 receptor27 and a bis(2-picolyl)aniline (DPA) receptor28 have been applied in ion-binding studies. Our group reported a cyclometalated [(C=N=N)Pt(µ)] terpyridylacetylide with a rich electron-donating bis-arylamine moiety, which exhibits the H’-triggered, Zn^{2+}-enhanced and OH’-quenched phosphorescence properties and an unexpected Zn^{2+}-selective luminescence chemosensor behavior in acid solution.29 However, owing to the lack of a clear molecular structure, the structure–property relationship is not well established for this class of multi-component compounds.

Herein, we design a novel three-component molecular system (Chart 1, complex 3), in which the [(C=N=N)Pt(µ)] acetylide core and the pendant TPY unit still serve as a planar fluorophore and a general receptor for transition metal ions, respectively, while the carbazole (CBZ) moiety serves as a rigid...
and higher energy-donating component. Its synthesis, crystal structure, photophysical properties, and transition metal ion-binding luminescent behaviors are mainly discussed with complex 2 (Scheme 1) as the reference compound.

**Experimental**

**Materials and general procedures**

The commercially available reagents 3,6-di-tert-butylcarbazole, 18-crown-6, triethylamine (Et3N), phenylacetylene (C6H5C≡CH), anhydrous K2CO3, CuI, and K2Ptl4 were of analytical grade and used as received without further purification. Reactions were carried out under an argon atmosphere in an oven-dried glassware using standard Schlenk techniques.

The solvents used for synthesis were also of analytical grade and distilled under an argon atmosphere from molten sodium. Dichloromethane was distilled before using it as a solvent for photophysical and electrochemical determinations of the sample. All metal salts, such as Zn(ClO4)2, Cd(ClO4)2, Fe(ClO4)2, NaClO4, KCIO4, Ca(CH3COO)2, Mg(CH3COO)2, Mn(CH3COO)2, Co(CH3COO)2, Ni(CH3COO)2, Cu(CH3COO)2, and Pb(CH3COO)2, were of analytical reagent grade and used as supplied to prepare the titrant CH3OH solutions containing 3.5 × 10−3 mol dm−3 metal ions. Tetrabutylammonium perchlorate ([Bu4N]ClO4), the supporting electrolyte for electrochemical studies, was prepared by the metathesis of tetrabutylammonium bromide and perchloric acid. The electrolyte was recrystallized from hot pentanol/ethyl acetate solution twice to give colorless crystals and dried in a vacuum oven. (Caution: perchlorate salts are potentially explosive and should be handled with care and in small amounts.)

Syntheses of 4-[(p-bromophenyl)-6-phenyl-2,2′-bipyridine and 4′-(4-ethylphenyl)-2,2′:6′,2′-terpyridine have been described previously.29

**Synthesis of 4-[(p-[9-(3,6-di-tert-butylcarbazolyl)]-phenyl-6-phenyl-2,2′-bipyridine (HL)**

A mixture of 3,6-di-tert-butylcarbazole (3.91 g, 14.0 mmol), 4-(p-bromophenyl)-6-phenyl-2,2′-bipyridine (3.87 g, 10.0 mmol), anhydrous K2CO3 (4.18 g, 30.0 mmol), CuI (0.12 g, 0.6 mmol), and 18-crown-6 (0.28 g, 1.1 mmol) was transferred into a flask, purged with argon, and then heated to 185 °C under stirring for 24 h. After the reaction mixture was cooled to ambient temperature, CH2Cl2 and H2O were added. The aqueous phase was discarded, and the organic phase was washed with distilled H2O and then dried over anhydrous Na2SO4. The evaporated residue was applied to a chromatography column (neutral Al2O3) and eluted with CH2Cl2-petroleum ether (2:1). The pure product was obtained by recrystallization from its MeOH/CH2Cl2 solution as a white crystal in a yield of 78%.1H NMR (400 MHz, CDCl3): δ 8.76–8.73 (m, 3H), 8.26 (d, J = 7.6 Hz, 2H), 8.16 (s, 2H), 8.09 (s, 1H), 8.05 (d, J = 8.0 Hz, 2H), 7.90 (t, J = 7.6 Hz, 1H), 7.73 (d, J = 8.2 Hz, 2H), 7.56 (t, J = 7.4 Hz, 2H), 7.51–7.43 (m, 5H), 7.38 (t, J = 6.0 Hz, 1H, Ar), 1.52 (s, 18H, –CH2).13C {1H} NMR (100 MHz, CDCl3): δ 157.4, 156.3, 156.2, 149.4, 149.0, 143.2, 139.4, 139.0, 138.9, 137.1, 137.0, 129.2, 128.8, 128.7, 127.1, 127.0, 124.0, 123.8, 123.6, 121.7, 118.4, 117.5, 116.4, 109.2 (Ar), 34.8 (–C(CH3)3), 32.0 (–CH3); elemental analysis calc (%): for C33H28N4: C, 81.55; H, 6.18; N, 12.27.

**Synthesis of 1**

A mixture of HL (0.29 g, 0.5 mmol), K2Ptl4 (0.21 g, 0.5 mmol) and glacial acetic acid (50 mL) was refluxed for 12 h under an argon atmosphere in the absence of light. The reaction mixture was then cooled to room temperature and filtered. The obtained solid was recrystallized from MeOH/CH2Cl2 solution to form the desired product as an orange crystal in a yield of 92%.1H NMR (300 MHz, DMSO-d6): δ 8.92 (d, J = 4.8 Hz, 1H), 8.78 (d, J = 7.8 Hz, 1H), 8.63 (s, 1H), 8.41–8.33 (m, 6H), 7.92 (t, J = 6.6 Hz, 2H), 7.87 (d, J = 8.4 Hz, 2H), 7.54–7.42 (m, 5H), 7.13 (t, J = 8.4 Hz, 2H, Ar), 1.44 (s, 18H, –CH3). Elemental analysis calc (%): for C42H38ClN3Pt: C, 61.87; H, 4.70; N, 5.15.

**Synthesis of 2**

A mixture of rectification 3 (0.24 g, 0.3 mmol), C4H9C≡CH (0.10 g, 1.0 mmol), Et3N (5 mL) and CuI (5.0 mg, 0.03 mmol) in degassed CH2Cl2 (50 mL) was stirred for 24 h under an argon atmosphere at room temperature in the absence of light. The reaction mixture was then evaporated to dryness under reduced pressure. The crude product was purified by flash chromatography (neutral Al2O3, CH2Cl2 as eluent) and recrystallized from CH2Cl2/MeOH solution to form the desired product as a red crystal in a yield of 84%.1H NMR (400 MHz, DMSO-d6): δ 9.09 (d, J = 4.4 Hz, 1H), 8.80 (d, J = 8.0 Hz, 1H), 8.71 (s, 1H), 8.46 (s, 1H), 8.42–8.38 (m, 3H), 8.35 (s, 2H), 7.92–7.87 (m, 4H), 7.78 (d, J = 7.6 Hz, 1H), 7.53 (d, J = 8.8 Hz, 2H), 7.45 (d, J = 8.4 Hz, 2H), 7.40 (d, J = 7.2 Hz, 2H), 7.30 (t, J = 7.8 Hz, 2H), 7.19–7.09 (m, 3H, Ar), 1.44 (s, 18H, –CH3). Elemental analysis calc (%): for C50H43N3Pt: C, 68.17; H, 4.70; N, 5.15; found: C, 62.03; H, 4.83; N, 4.98.

**Synthesis of 3**

To a stirred solution of 1 (0.14 g, 0.17 mmol) and 4′-(4-ethylphenyl)-2,2′:6′,2′-terpyridine (0.07 g, 0.18 mmol) in degassed CH2Cl2 (20 mL), CuI (5.4 mg, 0.03 mmol) and Et3N (4 mL) were successively added. The mixture was stirred for 24 h under an argon atmosphere at room temperature in the absence of light and then evaporated to dryness under reduced pressure. The crude product was purified by flash chromatography (neutral
General procedure of consecutive titrations

The titrand solutions of 3 (100 mL) were freshly prepared in CH2Cl2 before the consecutive titration experiment. Absorption and emission spectra were recorded simultaneously after each addition (10 µL) of the corresponding titrant solution. Each titration step was stopped after no more changes were observed in the absorption and emission properties. The selectivity of 3 toward different metal cations was monitored by the parallel PL measurements and emission properties. The selectivity of 3 toward Fe2+ and Zn2+ were evaluated from their fluorescence titration data using the Stern–Volmer expression and the linear Benesi–Hildebrand equation, respectively.

Determination of the association constant (K)

The binding constants of 3 toward Fe2+ and Zn2+ from the absorption titration data were calculated using the linear Benesi–Hildebrand equation. Moreover, the binding constants of 3 toward Fe2+ and Zn2+ were evaluated from their fluorescence titration data using the Stern–Volmer expression and the linear Benesi–Hildebrand equation, respectively.

Determination of the limit of detection (LOD)

The limits of detection for 3 of Fe2+ and Zn2+ ions were calculated using the following equation:

\[ \text{LOD} = 3 \sigma / S \]

where \( \sigma \) is the standard deviation of the observed absorption and emission data, and \( S \) is the slope of the calibration curve.

Results and discussion

Synthesis and characterization

The synthesis of ligand HL and complexes 1-3 explored in this work is shown in Scheme 1. Furthermore, 4-(p-bromophenyl)-2,2′-bipyridine was obtained using Kröhnke’s method, and 4-[[p-[3,6-di-tert-butyl]carbazolyl]-phenyl-2,2′-bipyridine (HL) was synthesized by the high-temperature Ullmann condensation between 3,6-di-tert-butylcarbazole and 4-(p-bromophenyl)-6-phenyl-2,2′-bipyridine, followed by chromatographic purification. The cyclometalated Pt[II] chloride was synthesized by refluxing HL and K2PtCl4 in glacial acetic acid for 12 h. Finally, reaction of 1 with phenylacetylene and 4-(4-ethynylphenyl)-2,2′-6′,2″-terpyridine by Sonogashira’s method and then purification by flash chromatography and recrystallization resulted in complexes 2 and 3 as red crystals in good yields, respectively. Both of them have good solubility in CH2Cl2 and were characterized by 1H NMR, elemental analysis, and/or MALDI-TOF MS. In addition, their structures were revealed by X-ray crystallography.

Crystal structures

The single crystals of complexes 2 and 3 (CCDC: 991876 and 991877) were grown by slowly evaporating the solvent from their concentrated CH2Cl2/MeOH solutions, and the crystal data and crystal structure parameters are listed in Table S1.† Their crystallographic geometry of the Pt atom is a distorted square planar configuration with the C(1)–Pt(1)–N(2) and N(1)–Pt(1)–C(35) angles of 160.4(18) and 175.8(18)° for 2 and 160.5(2) and 178.1(2)° for 3. The bond distances of Pt(1)–C(1), Pt(1)–N(1) and Pt(1)–N(2) are 2.016(5), 1.991(4) and 2.111(4) Å for 2 and 1.993(6), 1.981(5) and 2.096(5) Å for 3, which are comparable to those of previously reported analogs.26-39 The C(1)–C(6)–C(7)–N(1) and N(1)–C(11)–C(12)–N(2) torsion angles of 1.91 and 1.60° for 2 and 4.34 and 7.23° for 3 suggest that the configuration of (C=N=N) ligand is nearly planar. The dihedral angles between the phenyl ring at 4-position of the (C=N=N) ligand and the plane of the [(C=N=N)Pt] moiety are 32.00° for 2 and 34.11° for 3. In complex
As expected, three pyridine rings of TPY domain exhibit the transoid conformation about interannular C–C bonds, and the interplanar angles between two terminal pyridine rings and central pyridine ring are 13.79\(^\circ\) and 18.05\(^\circ\), respectively. The terminal TPY and CBZ planes are basically coplanar to the central [(C^N^N)Pt] unit, as evidenced by the dihedral angles of 10.27\(^\circ\) and 9.88\(^\circ\) between them, respectively, indicating its efficient \(\pi\)-conjugation molecule structure.

Instead of forming a continuous chain linked by the Pt–Pt interaction as in [(C^N^N)PtCl]\(^3^-\) the crystal lattice of 2 is packed by one type of alternating arranged dimeric units along the c axis (Fig. 3), in which two molecules are stacked in parallel with a Pt–Pt distance of 9.149 Å. However, complex 3 shows another type of continuously stacked dimers in a head-to-tail fashion along the a axis (Fig. 4). The Pt–Pt distance of 4.725 Å also suggests no metal–metal interaction, but the vertical distance of 3.489 Å between the neighboring [(C^N^N)Pt] planes indicates that they are likely to be held together by the weak \(\pi\)-\(\pi\) interaction.

Photophysical properties

The UV-vis absorption spectra of complex 1–3 in CH\(_2\)Cl\(_2\) solution exhibit intense bands at 250–400 nm and less intense bands at 400–550 nm (Table 1 and Fig. 5). With reference to previous spectroscopic work on phenylbipyridyl platinum(II) complexes,\(^39\) the high-energy intense absorption bands are clearly assigned to the singlet \(\pi\rightarrow\pi^*\) intraligand (IL) transitions of phenylbipyridine (C^N^N) and alkynyl (C≡C–Ar) ligands, while the low-energy bands are attributed to the well-known spin-allowed singlet \(d\pi(Pt)\rightarrow\pi^*(C^N^N)\) metal-to-ligand charge transfer (MLCT) mixing with the \(\pi(Cl\text{ or }C≡C–Ar)\rightarrow\pi^*(C^N^N)\) ligand-to-ligand charge transfer (LLCT) transitions. The low-energy absorption bands obey Beer’s law in the range of 10\(^{-6}\) to 10\(^{-4}\) mol dm\(^{-3}\), suggesting no dimerization or oligomerization of these complexes within this concentration range.
The highest occupied molecular orbital (HOMO) and the lowest unoccupied molecular orbital (LUMO) levels are calculated from the onset value of their absorption bands, \( \lambda_{\text{onset}} + 4.34 \) eV, respectively, while the energy gaps (\( \epsilon_{\text{g}} \)) are the diagonalization between the HOMO and LUMO levels.

The enhanced luminescence efficiency ([\( I_L/\bar{I} \) − 1] \( \times 100\% \)) is calculated to be ca. 175%, in which \( I_L \) and \( \bar{I} \) stand for the integrated area from 480 to 750 nm in the emission spectra of 3 excited at 338 and 450 nm, respectively.

The electrochemical behaviors of HL and 1–3 in CH$_2$Cl$_2$ solution at room temperature were also investigated by cyclic voltammetry (CV), and the data are listed in Table 2. The cyclic voltammogram of HL shows a quasi-reversible redox wave (\( \Delta \epsilon = 133 \) mV, \( 1/2 \ell_{\text{pa}}/\ell_{\text{pc}} = 1.0 \)) at \( E_{1/2} = +0.75 \) V. As the oxidation potential of the (ph-C$^N$N)$_2$ unit is beyond the potential window of the solvent, this wave is assigned to the mono radical couple at similar negative potentials, which presumably

Table 1 Photophysical properties of compounds 1–3

| Compound | \( \lambda_{\text{max,abs}} \) (nm) \( (\epsilon \times 10^4 \text{ dm}^{-1} \text{ mol}^{-1} \text{ cm}^{-1}) \) | \( \lambda_{\text{max,em}} \) (nm) | \( \phi_{\text{em}} \) | \( \tau_{\text{em}} \) (ns) | \( k_{\text{flu}} \) \( (\times 10^3) \) (s$^{-1}$) | \( k_{\text{r}} \) \( (\times 10^3) \) (s$^{-1}$) |
|----------|-------------------------------------------------|-----------------|---------------|-----------------|-----------------|-----------------|
| 1        | 432 (1.06), 414 (sh, 1.12), 376 (1.71), 336 (2.29), 290 (5.71) | 566, 598 (sh) | 0.019 | 5.10 | 1.92 | 3.73 |
| 2        | 438 (1.34), 376 (2.20), 338 (2.84), 288 (8.80) | 600 | 0.026 | 5.24 | 1.86 | 4.96 |
| 3        | 442 (1.51), 370 (sh, 2.79), 338 (4.73), 292 (7.82) | 596 | 0.045 | 5.89 | 1.62 | 7.64 |
| 3 + Fe$^{2+}$ (2 : 1) | 580 (1.98), 440 (2.58), 366 (sh, 2.97), 326 (5.06), 290 (8.03) | 568 | 0.0065 | 5.32 | 1.87 | 1.22 |
| 3 + Zn$^{2+}$ (2 : 1) | 448 (3.20), 382 (sh, 2.53), 332 (4.16), 290 (8.42) | 582 | 0.050 | 5.42 | 1.75 | 9.23 |

* Measured in an air-saturated CH$_2$Cl$_2$ solution at room temperature. * Excited wavelength is 450 nm. * PL quantum yields and lifetime. * \( k_{\text{flu}} = (1 - \phi/\tau) \). * \( k_{\text{r}} = \phi/\tau. * Measured after addition of 0.5 equiv. metal ion into an air-saturated 3 CH$_2$Cl$_2$ solution at room temperature.

Table 2 Electrochemistry data of the ligand and Pt(s) complexes

| Compound | Ligand-based, \( E_{1/2} \) (V) | Metal-based, \( E_{\text{p,a}} \) (V) | Reduction \( E_{1/2} \) (V) | HOMO$^\text{L}$ (eV) | LUMO$^\text{L}$ (eV) | \( E_{\text{g}} \) (eV) |
|----------|-----------------|-----------------|-----------------|---------------|---------------|-----------------|
| HL       | +0.75 | — | — | −5.0 | −2.4 | 2.6 |
| 1        | +0.76 | +0.40 (sh) | −1.76 | −4.7 | −2.7 | 2.0 |
| 2        | +0.77 | +0.50 (sh) | −1.77 | −4.7 | −2.7 | 2.0 |
| 3        | +0.78 | +0.60 (sh) | −1.74 | −4.8 | −2.7 | 2.1 |

* Determined in CH$_2$Cl$_2$ at room temperature with 0.1 mol dm$^{-3}$ Bu$_4$NClO$_4$ as a supporting electrolyte, scanning rate: 50 mV s$^{-1}$. * Values are reported versus Fe$^{3+}$/Fe (\( E_{1/2} = +0.46 \) V vs. saturated calomel electrode (SCE)). * The highest occupied molecular orbital (HOMO) and the lowest unoccupied molecular orbital (LUMO) levels are calculated from the onset values of their first anodic and cathodic waves according to the equation \( E = (E_{\text{onset}} + 4.34) \) eV, respectively, while the energy gaps (\( E_{\text{g}} \)) are the differences in their HOMO and LUMO levels.
corresponds to the metal-perturbed one-electron reduction of the (C’N’N) ligand. In addition, the irreversible Pt(n)-based oxidation wave is shown as a shoulder peak at lower anodic potential than that of the CBZ moiety. The HOMO and LUMO levels of 1–3 were also estimated from their electrochemical data, which reveal that the HOMOs are basically Pt(n) acetylidy based and the LUMOs predominantly correspond to the π* orbitals of cyclometalated ligands. When the electron withdrawing TPY group is attached to the phenylacetylide position, complex 3 possesses a clearly anodic-shifted Pt(n) oxidation peak (ΔE_{pa} = 0.10 V) and a slightly positive-shifted reduction potential of the (C’N’N) ligand (ΔE_{1/2} = 0.03 V), suggesting that introduction of TPY mainly stabilizes its HOMO energy level (ΔE_g = 0.1 eV). These results are consistent with the above-mentioned spectroscopic studies and theoretical calculations on the frontier molecular orbitals of phosphorescent cyclometalated Pt(n) acetylidy.

**Titration of 3 with Fe^{2+}**

As shown in Fig. 6a, besides the presence of the φ (Pt) → π*(C’N’N) \(^1\)MLCT absorption band (λ_{max} = 442 nm) of 3, a new spin-allowed singlet φ (Fe) → π*(TPY) metal-to-ligand charge transfer (MLCT) transition \(\pi^*(\text{MLCT})\) appeared at \(λ_{max} = 580 \text{ nm}\) upon addition of Fe\(^{2+}\) (Table 1 and Fig. S3†). Both of them are gradually increased, and they finally reach their maxima when titrated with 0.5 molar amount of Fe\(^{2+}\) (Fig. 6a, inset). These results suggest that the added iron ions bond to the free TPY receptor of 3 with 1:2 stoichiometry to form the heterotrinuclear complex (Pt–Fe–Pt) containing the [Fe(TPY)]\(^{2+}\) core. A perfect linear relationship \((R = 0.99983)\) is established from the absorption titration data for the plot, and absorbance was measured at 580 nm as a function of \(1/[\text{Fe}^{2+}]\) (from 0 to 4.50 × 10\(^{-6}\) mol dm\(^{-3}\)) (Fig. S4†). The colorimetric LOD is 8.21 × 10\(^{-8}\) mol dm\(^{-3}\), which is comparable to that of the ruthenium(n) complex-based chemosensor and much lower than that of benzimidazole-based chemosensor. By plotting the measured \([1/A - A_0]/C_0\) at 580 nm as a function of \([1/[\text{Fe}^{2+}]\) based on the well-known Benesi–Hildebrand expression (Fig. S5†), a good linear relationship \((R = 0.99958)\) is obtained, and the association constant of 3 with Fe\(^{2+}\;, K_a = 5.06 \times 10^4 \text{ mol}^{-1}\text{ dm}^3\), is calculated from the slope and intercept of this linear plot. Owing to its lower energy excited state of the as-formed [Fe(TPY)]\(^{2+}\) core in the Pt–Fe–Pt heterotrinuclear complex, the photoinduced energy/electron transfer (PET) process from the peripheral cyclometalated Pt(n) complex units to the central Fe–TPY complex quenches the MLCT emission of 3 with a much smaller quantum yield of 0.0065 (Fig. 6b). A good linear relationship \((R = -0.99861)\) between the PL intensity and [Fe\(^{2+}\)] (from 0 to 4.50 × 10\(^{-6}\) mol dm\(^{-3}\)) is also obtained from the quenching titration data (Fig. S6†), and the LOD is 2.38 × 10\(^{-7}\) mol dm\(^{-3}\), which is an order of magnitude lower than that of the benzoimidazoquinazoline-based fluorescent chemosensor. In addition, the apparent association constant of 3 with Fe\(^{2+}\); \(K_a = 3.14 \times 10^5 \text{ mol}^{-1}\text{ dm}^3\) (\(R = 0.9934\)) is determined from the Stern–Volmer plot (Fig. S7†). The inflection point of the curve measured the PL intensity at 596 nm as a function of the molar ratio of added Fe\(^{2+}\) and the central Pt\(^{2+}\) ion of 3, respectively.

**Titration of 3 with Zn\(^{2+}\)**

A similar titration experiment was conducted in CH\(_2\)Cl\(_2\) solution containing 1.50 × 10\(^{-5}\) mol dm\(^{-3}\) 3 using Zn\(^{2+}\) instead of Fe\(^{2+}\), but different absorption and emission changes were observed: (1) the φ (Pt) → π*(C’N’N) \(^1\)MLCT absorption band of 3 is gradually increased with red-shift (Δ\(\lambda_{max,abs} = 6 \text{ nm}\)) of its wavelength maximum (Fig. 7a). In the range of 0–4.00 × 10\(^{-6}\) mol dm\(^{-3}\), its intensity is linearly dependent on [Zn\(^{2+}\)] (\(R = 0.99833\)) (Fig. S9†) and the colorimetric LOD is 2.42 × 10\(^{-7}\) mol dm\(^{-3}\). The...
increased, but with a clear blue-shift (Δλ\text{max,em} = 14 nm) of its wavelength maximum (Fig. 7b). The linear relationship \( R = 0.99632 \), from 0 to 4.00 \( \times 10^{-5} \text{ mol dm}^{-3} \), the LOD value of 3.61 \( \times 10^{-7} \text{ mol dm}^{-3} \) and the association constant of 3 with Zn\text{2+} \( (K_a = 4.50 \times 10^{4} \text{ mol}^{-1} \text{ dm}^{3}, R = 0.99789) \) are also obtained from its PL titration data (Fig. S11 and S12). The LOD is lower than or comparable to those of organic\text{21-24} and transition metal complex fluorophores.\text{29,35,36} When the molar ratio of Zn\text{2+}/Pt\text{2+} is near to 0.5 (Fig. 7, inset), the MLCT absorption band at \( \lambda_{\text{max}} = 448 \text{ nm} \) and the corresponding emission band at \( \lambda_{\text{max}} = 582 \text{ nm} \) reach their maxima, indicating the formation of the Pt–Zn–Pt heterotrinuclear complex with 1:2 stoichiometry composition. The MLCT absorption sensitivity and emission enhancement effects\text{29} (Table 1, Fig. S3 and S8) strongly suggest that the PET process from the central high energy [Zn(TPY)]\text{2+} core to cyclometalated Pt\text{n+} complex units is present in this multi-component complex with a maximum efficiency \((I_{\text{max}} - I_0)/I_0 \times 100(\%)) of 62.7%.

Moreover, introduction of Zn\text{2+}, a strong Lewis acid, further stabilizes the Pt\text{n+} acetylide-based HOMO level relative to the \text{"free"} TPY complex 3, resulting in the blue-shift of its emission band, which is accordant with its spectroscopic and electrochemical studies.

**Selectivity**

The binding behaviors of 3 toward different metal cations (0.5 equiv. of 3) were monitored by PL spectroscopy under the same condition. As expected, in Fig. 8, alkali (Na\text{+} and K\text{+}), alkali-earth (Mg\text{2+} and Ca\text{2+}) and Pb\text{2+} ions exhibit no significant luminescence quenching or enhancing response. Similar to Fe\text{2+}, the other common transition metal ions, such as Mn\text{2+}, Co\text{2+}, Cu\text{2+}, and Ni\text{2+}, also exhibit dramatic luminescence quenching effects. With the same filled d\text{10} electronic configuration, Cd\text{2+} displays a similar luminescence enhancement factor to Zn\text{2+}, which is accordant with those of other TPY- and its analog-based chemosensors.\text{28,40,57}

**Conclusions**

In this study, we synthesized a novel cyclometalated Pt\text{n+} acetylide (3) derivative bearing a CBZ donor and a TPY receptor, whose molecular structure is clearly demonstrated by X-ray crystallography. This efficient \( \pi \)-conjugation system displays an intense \( ^1 \text{MLCT} \) absorption band (\( \epsilon = 1.51 \times 10^{4} \text{ dm}^{3} \text{ mol}^{-1} \text{ cm}^{-1} \)) at \( \lambda_{\text{max}} = 442 \text{ nm} \) and a strong \( ^3 \text{MLCT} \) phosphorescence emission (\( \Phi = 0.045 \)) at \( \lambda_{\text{max}} = 596 \text{ nm} \) in air-saturated CH\text{2}Cl\text{2} solution at room temperature. Owing to the presence of the free TPY receptor, complex 3 possesses the expected luminescence responses toward common transition metal cations, e.g. the Fe\text{2+}-quenched and Zn\text{2+}-enhanced luminescence behaviors. The 3–Fe\text{2+} and 3–Zn\text{2+} binding constants are correspondingly calculated from their absorption \((K_a = 5.06 \times 10^{4} \text{ mol}^{-1} \text{ dm}^{3} \text{ for } 3–\text{Fe}^2+ \text{ and } 5.16 \times 10^{4} \text{ mol}^{-1} \text{ dm}^{3} \text{ for } 3–\text{Zn}^2+)\) and emission \((3.14 \times 10^{5} \text{ mol}^{-1} \text{ dm}^{3} \text{ for } 3–\text{Fe}^2+ \) and \( 4.50 \times 10^{5} \text{ mol}^{-1} \text{ dm}^{3} \text{ for } 3–\text{Zn}^2+) \) titration data. The formation of the heterotrinuclear (Pt–M–Pt) complex with 1:2 stoichiometry is proven by the consecutive titrations, and the intra-molecular energy transfers between the [M(TPY)]\text{2+} core
and the terminal emission Pt(n) complex units are also quan-
titatively verified with the maximum PET efficiencies of 92.4% for 3–Fe$^{2+}$ system and 62.7% for 3–Zn$^{2+}$ system.

**Conflicts of interest**

There are no conflicts to declare.

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