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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 ($\epsilon = 1.51 \times 10^4 \text{ dm}^3 \text{ mol}^{-1} \text{ cm}^{-1}$) at $\lambda_{\text{max}} = 442 \text{ nm}$ and a strong phosphorescence emission ($\Phi = 0.045$) at $\lambda_{\text{max}} = 596 \text{ nm}$ in an air-saturated CH_2Cl_2 solution at room temperature, resulting from the $d\pi(\text{Pt}) \rightarrow \pi^*(\text{C}^{\wedge}\text{N}^{\wedge}\text{N})$ metal-to-ligand charge transfer (MLCT) mixing with the $\pi(\text{C}\equiv\text{C}-\text{Ar}) \rightarrow \pi^*(\text{C}^{\wedge}\text{N}^{\wedge}\text{N})$ 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 heterotrinnuclear ($\text{Pt}-\text{M}-\text{Pt}$) 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 $[\text{M}(\text{TPY})_2]^{2+}$ core and the terminal phosphorescent Pt(II) complex units.

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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-ethynylbenzo-15-crown-5,⁶ 5,17-diethynyl-25,27-dimethoxycalix[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 $[(\text{N}^{\wedge}\text{C}^{\wedge}\text{N}) \text{ motif}]^{10-16}$ and bipyridine $[(\text{C}^{\wedge}\text{N}^{\wedge}\text{N}) \text{ motif}]^{17-22}$ as well as the related diphenyl-substituted pyridine $[(\text{C}^{\wedge}\text{N}^{\wedge}\text{C}) \text{ motif}]^{23,24}$ 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 $[(\text{C}^{\wedge}\text{N}^{\wedge}\text{N})\text{Pt}(\text{II})]$ σ -alkynyl complex with a benzo-15-azacrown-5 receptor²⁷ and a bis(2-picoly)aniline (DPA) receptor²⁸ have been applied in ion-binding studies. Our group reported a cyclometalated $[(\text{C}^{\wedge}\text{N}^{\wedge}\text{N})\text{Pt}(\text{II})]$ 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 $[(\text{C}^{\wedge}\text{N}^{\wedge}\text{N})\text{Pt}(\text{II})]$ 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

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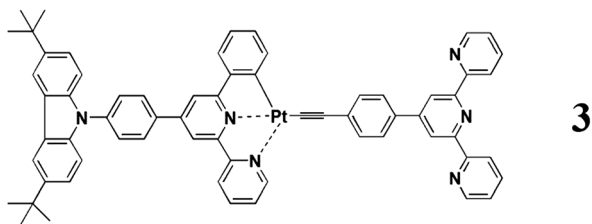


Chart 1 Molecular structure of complex 3.

and higher energy-donating component. Its synthesis, crystal structure, photophysical properties, and transition metal ion-binding luminescence 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 (Et_3N), phenylacetylene ($\text{C}_6\text{H}_5\text{C}\equiv\text{CH}$), anhydrous K_2CO_3 , CuI, and K_2PtCl_4 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. Diisopropylamine was treated with 5 Å molecular sieves 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 $\text{Zn}(\text{ClO}_4)_2$, $\text{Cd}(\text{ClO}_4)_2$, $\text{Fe}(\text{ClO}_4)_2$, NaClO_4 , KClO_4 , $\text{Ca}(\text{CH}_3\text{COO})_2$, $\text{Mg}(\text{CH}_3\text{COO})_2$, $\text{Mn}(\text{CH}_3\text{COO})_2$, $\text{Co}(\text{CH}_3\text{COO})_2$, $\text{Ni}(\text{CH}_3\text{COO})_2$, $\text{Cu}(\text{CH}_3\text{COO})_2$, and $\text{Pb}(\text{CH}_3\text{COO})_2$, were of analytical reagent grade and used as supplied to prepare the titrant CH_3OH solutions containing $3.5 \times 10^{-3} \text{ mol dm}^{-3}$ metal ions. Tetrabutylammonium perchlorate (Bu_4NClO_4), 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-ethynylphenyl)-2,2':6',2''-terpyridine have been described previously.²⁹

Synthesis of 4-[*p*-(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 K_2CO_3 (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, CH_2Cl_2 and H_2O were added. The aqueous phase was discarded, and the organic phase was washed with distilled

H_2O and then dried over anhydrous Na_2SO_4 . The evaporated residue was applied to a chromatography column (neutral Al_2O_3) and eluted with CH_2Cl_2 -petroleum ether (2 : 1). The pure product was obtained by recrystallization from its MeOH/ CH_2Cl_2 solution as a white crystal in a yield of 78%. ^1H NMR (400 MHz, CDCl_3): 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, $J = 8.4$ 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, $-\text{CH}_3$). ^{13}C { ^1H } NMR (100 MHz, CDCl_3): 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 ($-\text{C}(\text{CH}_3)_3$), 32.0 ($-\text{CH}_3$); elemental analysis calcd (%) for $\text{C}_{31}\text{H}_{28}\text{N}_4$: C, 81.55; H, 6.18; N, 12.27; found: C, 81.79; H, 6.20; N, 11.98.

Synthesis of 1

A mixture of HL (0.29 g, 0.5 mmol), K_2PtCl_4 (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/ CH_2Cl_2 solution to form the desired product as an orange crystal in a yield of 92%. ^1H NMR (300 MHz, $\text{DMSO}-d_6$): 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, $-\text{CH}_3$). Elemental analysis calcd (%) for $\text{C}_{42}\text{H}_{38}\text{ClN}_3\text{Pt}$: C, 61.87; H, 4.70; N, 5.15; found: C, 62.03; H, 4.83; N, 4.98.

Synthesis of 2

A mixture of 1 (0.24 g, 0.3 mmol), $\text{C}_6\text{H}_5\text{C}\equiv\text{CH}$ (0.10 g, 1.0 mmol), Et_3N (5 mL) and CuI (5.0 mg, 0.03 mmol) in degassed CH_2Cl_2 (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 Al_2O_3 , CH_2Cl_2 as eluent) and recrystallized from CH_2Cl_2 /MeOH solution to form the desired product as a red crystal in a yield of 84%. ^1H NMR (400 MHz, $\text{DMSO}-d_6$): 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, $-\text{CH}_3$). Elemental analysis calcd (%) for $\text{C}_{50}\text{H}_{43}\text{N}_3\text{Pt}$: C, 68.17; H, 4.92; N, 4.77; found: C, 68.45; H, 5.10; N, 4.62.

Synthesis of 3

To a stirred solution of 1 (0.14 g, 0.17 mmol) and 4'-(4-ethynylphenyl)-2,2':6',2''-terpyridine (0.07 g, 0.18 mmol) in degassed CH_2Cl_2 (20 mL), CuI (5.4 mg, 0.03 mmol) and Et_3N (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



Al₂O₃, CH₂Cl₂ as eluent) and recrystallized from CH₂Cl₂/MeOH solution to form the desired product as a red crystal in a yield of 89%. ¹H NMR (400 MHz, DMSO-*d*₆): δ 9.10 (d, *J* = 4.8 Hz, 1H), 8.80 (d, *J* = 4.4 Hz, 3H), 8.75 (s, 2H), 8.70 (d, *J* = 8.0 Hz, 3H), 8.46–8.38 (m, 4H), 8.35 (s, 2H), 8.06 (t, *J* = 8.4 Hz, 2H), 7.93–7.85 (m, 6H), 7.82 (d, *J* = 7.6 Hz, 1H), 7.60–7.52 (m, 6H), 7.45 (d, *J* = 8.8 Hz, 2H), 7.19 (t, *J* = 6.8 Hz, 1H), 7.13 (t, *J* = 7.4 Hz, 1H, *Ar*), 1.43 (s, 18H, –CH₃). Elemental analysis calcd (%) for C₆₅H₅₂N₆Pt: C, 70.19; H, 4.71; N, 7.56; found: C, 70.44; H, 4.86; N, 7.38. Matrix-assisted laser desorption/ionization time-of-flight mass spectrometry (MALDI-TOF MS): *m/z* = 1112.4 ([M], C₆₅H₅₂N₆Pt requires 1112.39).

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 experiment under the same instrument condition, in which 0.5 equiv. of the metal ions was added in each titrand solution containing 1.5×10^{-5} mol dm^{−3} **3**.

Physical measurements and instrumentation

¹H and ¹³C{¹H} NMR spectra were recorded on Bruker AV 300 and 400 spectrometers and referenced with respect to TMS internal standard. Elemental analyses were carried out using a Bio-Rad Co's elemental analytical instrument. MALDI-TOF MS analyses were performed on a Bruker Autoflex III with a negative reflector mode. UV-vis absorption spectra were obtained using the Perkin-Elmer Lambda 650s ultraviolet-visible spectrophotometer. Steady-state PL spectra were performed on the Perkin-Elmer LS50B spectrometer. Emission lifetimes and PL quantum yields were determined on the FLS-980 fluorescence spectrometer (Edinburgh Instruments Ltd.) with a microflash lamp as the excitation source. CV spectra were conducted on the CHI660A electrochemical workstation in CH₂Cl₂ solutions with a three-electrode electrochemical cell using Pt electrode, SCE and Pt wire as the working, reference and counter electrodes, respectively. ⁿBu₄NClO₄ (0.1 mol L^{−1}) was used as the supporting electrolyte and the scan rate was 50 mV s^{−1}.

X-ray crystallography

Diffraction data were collected on the Bruker SMART APEX II CCD diffractometer equipped with graphite-monochromated Mo-K_α radiation (λ = 0.71073 Å) at 296(2) K using the Φ - ω scan mode. The intensity data were corrected by *L*_p factors and empirical absorption. The structure was solved by direct methods and subsequent successive difference Fourier maps, and refined by full-matrix least-squares techniques on *F*². All of the non-hydrogen atoms were refined anisotropically. The organic hydrogen atoms were generated geometrically. All calculations were carried out with SHELXTL-97 program.³⁰

Determination of the association constant (*K*_a)

The binding constants of **3** toward Fe²⁺ and Zn²⁺ from the absorption titration data were calculated using the linear Benesi–Hildebrand equation.^{31–33} Moreover, the binding constants of **3** toward Fe²⁺ and Zn²⁺ were evaluated from their fluorescence titration data using the Stern–Volmer expression^{34,35} and the linear Benesi–Hildebrand equation, respectively.

Determination of the limit of detection (LOD)

The limits of detection of **3** for Fe²⁺ and Zn²⁺ ions were calculated using the following equation:³²

$$\text{LOD} = 3\sigma/S$$

where σ is the standard deviation of the observed absorption and emission data, and *S* is the slope of the calibration curve.

Results and discussion

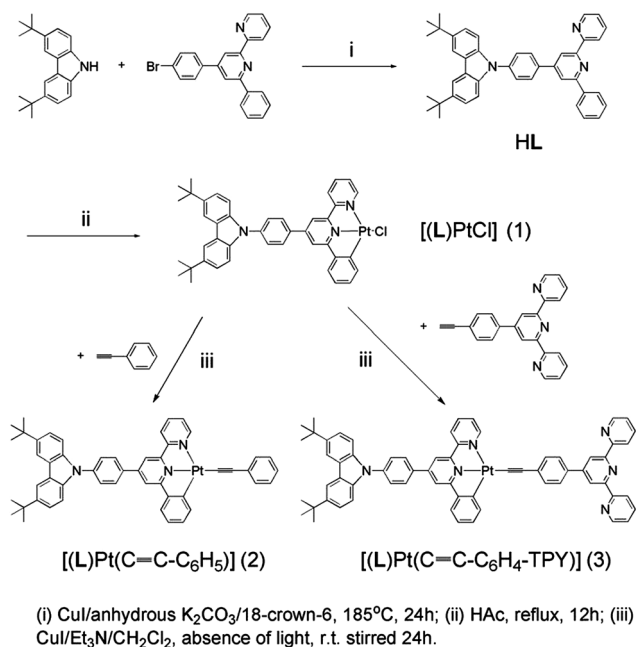
Synthesis and characterization

The synthesis of ligand HL and complexes **1–3** explored in this work is shown in Scheme 1. Furthermore, 4-(*p*-bromophenyl)-6-phenyl-2,2'-bipyridine was obtained using Kröhnke's method, and 4-{*p*-[9-(3,6-di-*tert*-butyl)carbazolyl]}-phenyl-6-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 **1** 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 perspective views are shown in Fig. 1 and 2. The coordination 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**, respectively, which are comparable to those of previously reported analogs.^{36–39} 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^NN) ligand is nearly planar. The dihedral angles between the phenyl ring at 4-position of the (C^NN) ligand and the plane of the [(C^NN)Pt] moiety are 32.00° for **2** and 34.11° for **3**. In complex





Scheme 1 Synthetic route of HL, 1, 2, and 3.

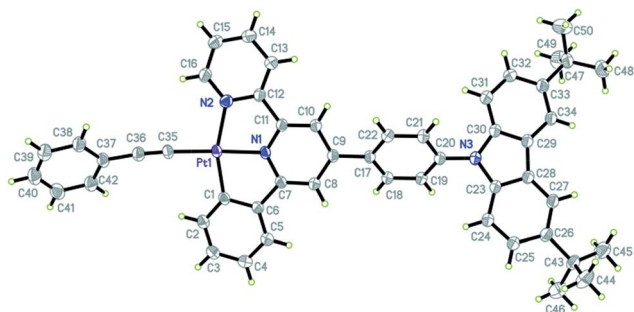


Fig. 1 Perspective view of the single crystal structure of 2 with 30% thermal ellipsoids (hydrogen atoms have been omitted for clarity).

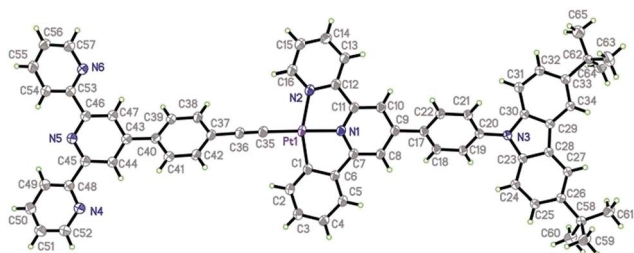


Fig. 2 Perspective view of the single crystal structure of 3 with 30% thermal ellipsoids (hydrogen atoms have been omitted for clarity).

3, as expected, three pyridine rings of TPY domain exhibit the transoid configuration about interannular C–C bonds, and the interplanar angles between two terminal pyridine rings and central pyridine ring are 13.79 and 18.05°, respectively. The terminal TPY and CBZ planes are basically coplanar to the central $[(C^{\wedge}N^{\wedge}N)Pt]$ unit, as evidenced by the dihedral angles of

10.27° and 9.88° between them, respectively, indicating its efficient π -conjugation molecule structure.

Instead of forming a continuous chain linked by the Pt–Pt interaction as in $[(C^{\wedge}N^{\wedge}N)PtCl]$,^{37,38} 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 $[(C^{\wedge}N^{\wedge}N)Pt]$ planes indicates that they are likely to be held together by the weak π – π interaction.

Photophysical properties

The UV-vis absorption spectra of complex 1–3 in CH_2Cl_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 ($C^{\wedge}N^{\wedge}N$) and alkynyl ($C\equiv C-Ar$) ligands, while the low-energy bands are attributed to the well-known spin-allowed singlet $d\pi(Pt) \rightarrow \pi^*(C^{\wedge}N^{\wedge}N)$ metal-to-ligand charge transfer (MLCT) mixing with the $\pi(Cl \text{ or } C\equiv C-Ar) \rightarrow \pi^*(C^{\wedge}N^{\wedge}N)$ 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.

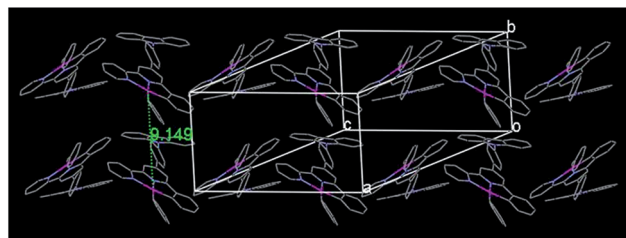
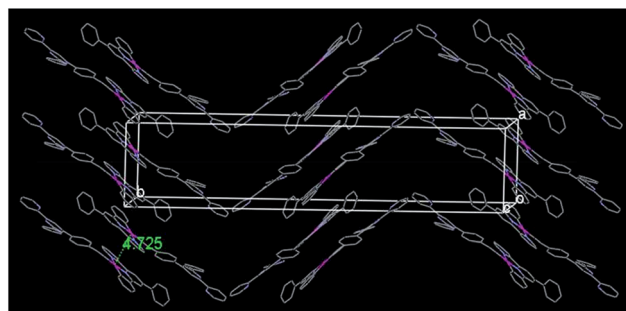
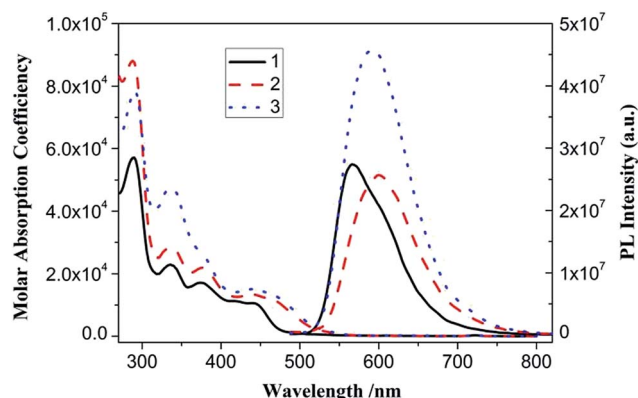
Fig. 3 Crystal packing diagram of 2 (hydrogen atoms and *tert*-butyl groups have been omitted for clarity).Fig. 4 Crystal packing diagram of 3 (hydrogen atoms and 3,6-di-*tert*-butylcarbazole units have been omitted for clarity).

Table 1 Photophysical properties of compounds 1–3^a

Compound	$\lambda_{\text{max,abs}}$ (nm) ($\epsilon \times 10^4 \text{ dm}^3 \text{ mol}^{-1} \text{ cm}^{-1}$)	$\lambda_{\text{max,em}}$ ^b (nm)	Φ_{em} ^{b,c}	τ_{em} ^{b,c} (μs)	k_{nr} ^d ($\times 10^5 \text{ s}^{-1}$)	k_{r} ^e ($\times 10^3 \text{ s}^{-1}$)
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) ^f	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) ^f	448 (3.20), 382 (sh, 2.53), 332 (4.16), 290 (8.42)	582	0.050	5.42	1.75	9.23

^a Measured in an air-saturated CH₂Cl₂ solution at room temperature. ^b Excited wavelength is 450 nm. ^c PL quantum yields and lifetime. ^d $k_{\text{nr}} = (1 - \Phi)/\tau$. ^e $k_{\text{r}} = \Phi/\tau$. ^f Measured after addition of 0.5 equiv. metal ion into an air-saturated 3 CH₂Cl₂ solution at room temperature.

Fig. 5 UV-vis absorption and PL emission spectra of 1, 2 and 3 in CH₂Cl₂ solution.

With an excitation at 450 nm, which is into their MLCT absorption bands, 1, 2 and 3 in CH₂Cl₂ solution emit reddish-orange luminescence (Table 1 and Fig. 5). For each of them, the time-resolved PL intensity decay is complex and is most adequately modeled using a sum of two single exponential functions (Fig. S1†). The average lifetimes [$\tau = (\alpha_1\tau_1^2 + \alpha_2\tau_2^2)/(\alpha_1\tau_1 + \alpha_2\tau_2)$], radiation (k_{r}) and non-radiation (k_{nr}) rate constants are calculated as listed in Table 1. The comparably long emission lifetimes (τ in μs scale), large Stoke's shifts, and k_{r} and k_{nr} values suggest that their emissions originate from the spin-forbidden triplet excited states. Complex 2 has longer emission wavelength and lifetime as well as larger quantum yield than 1, resulting from incorporation of the strong-field

acetylide ligand to extend its π -conjugated system and destabilize the non-radiative ligand-field transitions.^{39,40} With further introduction of the electron-withdrawing TPY unit, complex 3 shows a slightly blue-shifted ($\Delta\lambda_{\text{max,em}} = 4 \text{ nm}$) and apparently enhanced emission compared with 2, reflecting its lower HOMO level and stronger molecular rigidity. Despite the presence of the dynamical quenching effect of dissolved oxygen, complex 3 still possesses strong emission with comparative quantum yield in air-saturated CH₂Cl₂ solution at room temperature, which facilitates the consecutive titration and metal ion sensing studies mentioned later in this study.

In addition, selective optical-stimulation experiment was conducted to study the interaction between components of complex 3. When excited at 338 nm, which is into the CBZ absorption band in the UV region, the ³MLCT emission clearly increased compared with that of 3 excited at 450 nm (Fig. S2†). The enhanced luminescence efficiency ($[(I_2/I_1) - 1] \times 100(\%)$) is calculated to be ca. 175%, in which I_2 and I_1 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 CH₂Cl₂ 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 ($\Delta E = 133 \text{ mV}$, $i_{\text{p,a}}/i_{\text{p,c}} = 1.0$) at $E_{1/2} = +0.75 \text{ V}$. As the oxidation potential of the (ph-C⁺N⁺N) unit is beyond the potential window of the solvent, this wave is assigned to the mono radical cation formation of the CBZ group. For each of complexes 1–3, the cyclic voltammogram exhibits one reversible reduction couple at similar negative potentials, which presumably

Table 2 Electrochemistry data^a of the ligand and Pt(II) complexes

Compound	Oxidation ^b		Reduction $E_{1/2}$ (V)	HOMO ^c (eV)	LUMO ^c (eV)	E_{g} ^c (eV)
	Ligand-based, $E_{1/2}$ (V)	Metal-based, $E_{\text{p,a}}$ (V)				
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

^a Determined in CH₂Cl₂ at room temperature with 0.1 mol dm^{−3} ⁿBu₄NClO₄ as a supporting electrolyte, scanning rate: 50 mV s^{−1}. ^b Values are reported versus Fc⁺/Fc ($E_{1/2} = +0.46 \text{ V}$ vs. saturated calomel electrode (SCE)). ^c 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 $E = -(E_{\text{onset}} + 4.34) \text{ eV}$, respectively, while the energy gaps (E_{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(II)-based oxidation wave is shown as a shoulder peak at lower anodic potential than that of the CBZ moiety.⁴¹ The HOMO and LUMO levels of 1–3 were also estimated from their electrochemical data,⁴² which reveal that the HOMOs are basically Pt(II) acetylide based and the LUMOs predominantly correspond to the π^* orbitals of cyclometalated ligands.^{29,40,43–45} When the electron-withdrawing TPY group is attached to the phenylacetylide ligand of 2 at *para*-position, complex 3 possesses a clearly anodic-shifted Pt(II) oxidation peak ($\Delta E_{p,a} = 0.10$ V) and a slightly positive-shifted reduction potential of the (C[∧]N[∧]N) ligand ($\Delta E_{1/2} = 0.03$ V), suggesting that introduction of TPY mainly stabilizes its HOMO energy level ($\Delta E_g = 0.1$ eV). These results are consistent with the above-mentioned spectroscopic studies and theoretical calculations on the frontier molecular orbitals of phosphorescent cyclometalated Pt(II) acetylide.

Titration of 3 with Fe²⁺

As shown in Fig. 6a, besides the presence of the $d\pi(\text{Pt}) \rightarrow \pi^*(\text{C}^{\wedge}\text{N}^{\wedge}\text{N})$ ¹MLCT absorption band ($\lambda_{\text{max}} = 442$ nm) of 3, a new spin-allowed singlet $d\pi(\text{Fe}) \rightarrow \pi^*(\text{TPY})$ metal-to-ligand charge transfer (¹MLCT) transition^{46,47} appeared at $\lambda_{\text{max}} = 580$ nm upon addition of Fe²⁺ (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 Fe²⁺ (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 heterotrimeric complex (Pt–Fe–Pt) containing the [Fe(TPY)₂]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 [Fe²⁺] (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(II) complex-based chemosensor⁴⁸ and much lower than that of benzimidazole-based chemosensor.⁴⁹ By plotting the measured $[1/(A - A_0)]$ at 580 nm as a function of $1/[Fe^{2+}]$ 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 Fe²⁺, $K_a = 5.06 \times 10^4$ mol⁻¹ dm³, 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]²⁺ core in the Pt–Fe–Pt heterotrimeric complex,⁹ the photoinduced energy/electron transfer (PET) process from the peripheral cyclometalated Pt(II) 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 [Fe²⁺] (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.⁵⁰ In addition, the apparent association constant of 3 with Fe²⁺, $K_a = 3.14 \times 10^5$ mol⁻¹ dm³ ($R = 0.9934$), is determined from the Stern–Volmer plot (Fig. S7†). The inflection point of the curve measured the PL intensity at 596 nm as

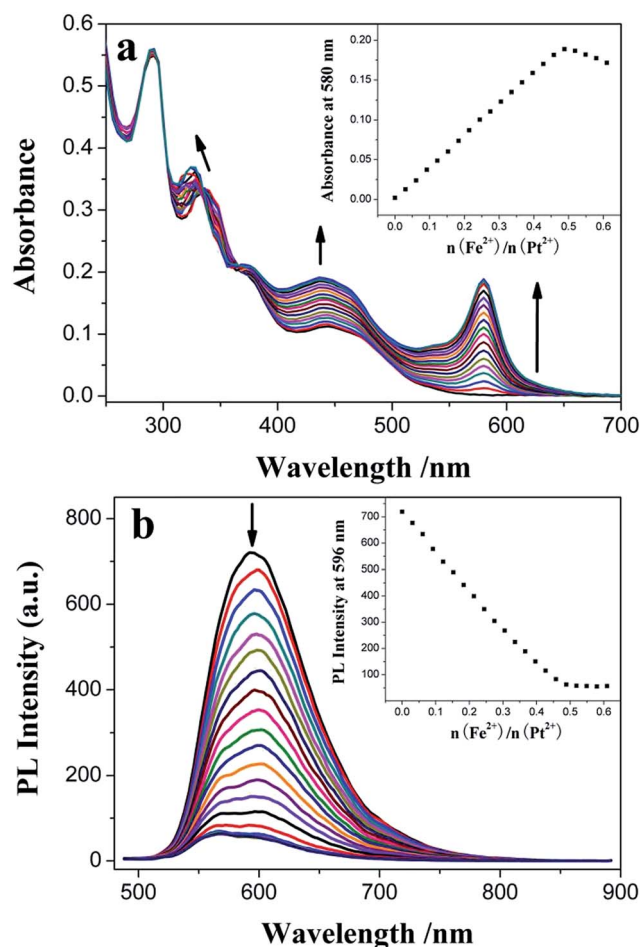


Fig. 6 Influence of consecutive Fe²⁺ titration on the UV-vis absorption (a) and PL emission (b) spectra of 1.50×10^{-5} mol dm⁻³ 3 in CH₂Cl₂ solution. The insets show plots of absorbance at 580 nm and integrated area of the emission wave in the range of 500–800 nm versus the molar ratio of added Fe²⁺ and the central Pt²⁺ ion of 3, respectively.

a function of the molar ratio of added Fe²⁺ and the central Pt²⁺ ion of 3 is also *ca.* 0.5 (Fig. 6b, inset), further confirming the 1 : 2 stoichiometry composition of the Pt–Fe–Pt heterotrimeric complex. Though the quenching effect is observed during the titration process, the MLCT emission of 3 is not fully quenched upon excessive addition of Fe²⁺ and the maximum PET efficiency ($[(I_0 - I_{\text{min}})/I_0] \times 100(\%)$) is estimated to be 92.4% (Table 1 and Fig. S8†).

Titration of 3 with Zn²⁺

A similar titration experiment was conducted in CH₂Cl₂ solution containing 1.50×10^{-5} mol dm⁻³ 3 using Zn²⁺ instead of Fe²⁺, but different absorption and emission changes were observed: (1) the $d\pi(\text{Pt}) \rightarrow \pi^*(\text{C}^{\wedge}\text{N}^{\wedge}\text{N})$ ¹MLCT absorption band of 3 is gradually increased with red-shift ($\Delta\lambda_{\text{max,abs}} = 6$ nm) of its wavelength maximum (Fig. 7a). In the range of $0\text{--}4.00 \times 10^{-6}$ mol dm⁻³, its intensity is linearly dependent on [Zn²⁺] ($R = 0.99833$) (Fig. S9†) and the colorimetric LOD is 2.42×10^{-7} mol dm⁻³. The



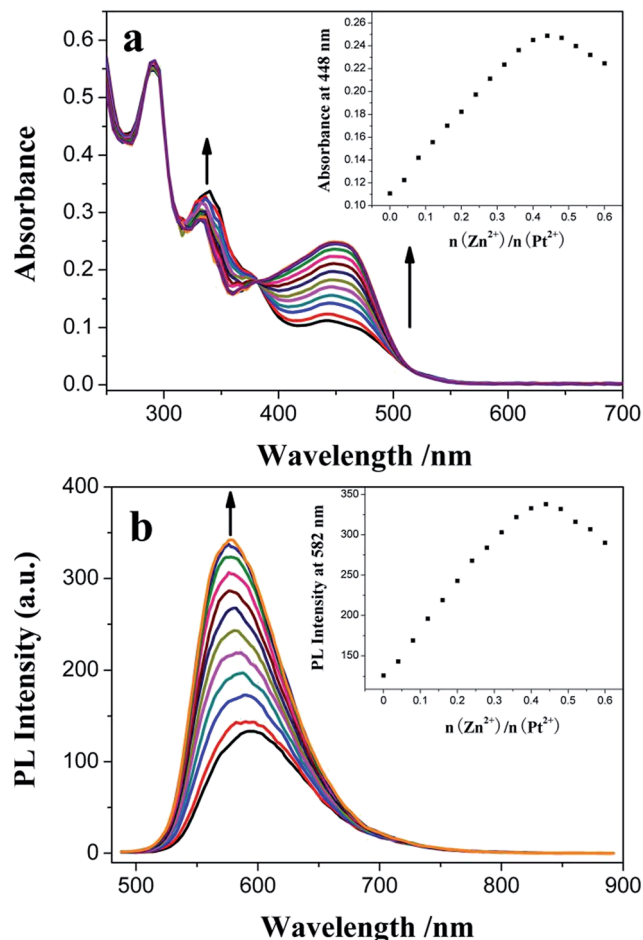


Fig. 7 Influence of consecutive Zn^{2+} titration on the UV-vis absorption (a) and PL emission (b) spectra of $1.50 \times 10^{-5} \text{ mol dm}^{-3}$ **3** in CH_2Cl_2 solution. The insets show plots of absorbance at 448 nm and integrated area of the emission wave in the range of 500–800 nm versus the molar ratio of added Zn^{2+} and the central Pt^{2+} ion of **3**, respectively.

association constant of **3** with Zn^{2+} is calculated to be $K_a = 5.16 \times 10^4 \text{ mol}^{-1} \text{ dm}^3$ ($R = 0.99967$) from its Benesi–Hildebrand plot (Fig. S10†). (2) And the MLCT emission band of **3** is also gradually increased, but with a clear blue-shift ($\Delta\lambda_{\text{max,em}} = 14 \text{ nm}$) of its wavelength maximum (Fig. 7b). The linear relationship ($R = 0.99632$, from 0 to $4.00 \times 10^{-6} \text{ mol dm}^{-3}$), the LOD value of $3.61 \times 10^{-7} \text{ mol dm}^{-3}$ and the association constant of **3** with Zn^{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^{51–54} and transition metal complex fluorophores.^{29,55,56} When the molar ratio of $\text{Zn}^{2+}/\text{Pt}^{2+}$ 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 heterotrimeric complex with 1 : 2 stoichiometry composition. The MLCT absorption sensitivity and emission enhancement effects²⁹ (Table 1, Fig. S3 and S8†) strongly suggest that the PET process from the central high energy $[\text{Zn}(\text{TPY})_2]^{2+}$ core to cyclometalated Pt(II) 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%.

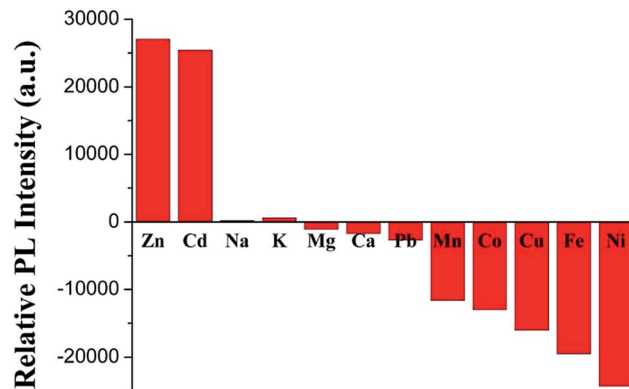


Fig. 8 PL responses of **3** toward various metal ions.

Moreover, introduction of Zn^{2+} , a strong Lewis acid, further stabilizes the Pt(II) acetylide-based HOMO level relative to the “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^+ and K^+), alkali-earth (Mg^{2+} and Ca^{2+}) and Pb^{2+} ions elicit no significant luminescence quenching or enhancing response. Similar to Fe^{2+} , the other common transition metal ions, such as Mn^{2+} , Co^{2+} , Cu^{2+} , and Ni^{2+} , also exhibit dramatic luminescence quenching effects. With the same filled d^{10} electronic configuration, Cd^{2+} displays a similar luminescence enhancement factor to Zn^{2+} , which is accordant with those of other TPY- and its analog-based chemosensors.^{28,40,57}

Conclusions

In this study, we synthesized a novel cyclometalated Pt(II) acetylide (**3**) derivative bearing a CBZ donor and a TPY receptor, whose molecular structure is clearly demonstrated by X-ray crystallography. This efficient π -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_2Cl_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^{2+} -quenched and Zn^{2+} -enhanced luminescence behaviors. The 3-Fe^{2+} and 3-Zn^{2+} binding constants are correspondingly calculated from their absorption ($K_a = 5.06 \times 10^4 \text{ mol}^{-1} \text{ dm}^3$ for 3-Fe^{2+} and $5.16 \times 10^4 \text{ mol}^{-1} \text{ dm}^3$ for 3-Zn^{2+}) and emission ($3.14 \times 10^5 \text{ mol}^{-1} \text{ dm}^3$ for 3-Fe^{2+} and $4.50 \times 10^4 \text{ mol}^{-1} \text{ dm}^3$ for 3-Zn^{2+}) titration data. The formation of the heterotrimeric (Pt–M–Pt) complex with 1 : 2 stoichiometry is proven by the consecutive titrations, and the intra-molecular energy transfers between the $[\text{M}(\text{TPY})_2]^{2+}$ core



and the terminal emission Pt(II) complex units are also quantitatively verified with the maximum PET efficiencies of 92.4% for 3-Fe²⁺ system and 62.7% for 3-Zn²⁺ system.

Conflicts of interest

There are no conflicts to declare.

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