Potential Switchable Circularly Polarized Luminescence from Chiral Cyclometalated Platinum(II) Complexes

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Supporting Information

ABSTRACT: A series of chiral cyclometalated platinum(II) complexes, [Pt(−-L1)(Dmpi)]Cl (−-1), [Pt(+-L1)(Dmpi)]Cl (+-1), [Pt(−-L2)(Dmpi)]Cl (−-2), [Pt(+-L2)(Dmpi)]Cl (+-2), [Pt3((+-)-L1)(Dmpi)]4[ClO4]4 ((+-)-3), and [Pt3((+-)-L2)(Dmpi)]4[ClO4]4 ((+-)-3)] [(−)-L2 = (−)-4,5-pine-6'-phenyl-2,2'-bipyrine, (+)-L1 = (+)-4,5-pine-6'-phenyl-2,2'-bipyrine, (+)-L2 = (+)-1,3-bis(2-(4,5-pine-6'-pyridyl)benzene, Dmpi = 2,6-dimethylphenyl isocyanide), have been designed and synthesized. In aqueous solutions, (−)-1 and (+)-1 aggregate into one-dimensional helical chain structures through Pt···Pt π–π and hydrophobic–hydrophobic interactions. (−)-3 and (−)-3 represent a novel helical structure with Pt···Pt bonds. The formation of helical structures results in enhanced and distinct chiroptical properties as evidenced by circular dichroism spectra. Circularly polarized luminescence (CPL) was observed from the aggregates of (−)-1 and (+)-1 in water, as well as (−)-3 and (−)-3 in dichloromethane. The CPL activity can be switched reversibly (for (−)-1 and (+)-1) or irreversibly (for (−)-3 and (−)-3) by varying the temperature.

INTRODUCTION

Circularly polarized luminescence (CPL) spectroscopy is the counterpart of circular dichroism (CD) in the characterization of emission. While CD spectroscopy has been widely used to investigate structural properties of the ground electronic states of a system, CPL has been proven to be powerful in studying the structural properties of the luminescent excited states. In some applications, CPL provides additional information concerning molecular dynamics and energetics by probing processes that occur between the excitation event and the emission. In spite of the fact that there are still many unresolved issues regarding the measurement and theoretical characterization, it is widely believed that CPL has great potential for the monitoring of chiral environments due to its high sensitivity and specificity. Moreover, CPL also has potential applications in photonic devices, such as light-emitting diodes, optical amplifiers, and optical information storage.

So far, the research field of CPL has been dominated by chiral lanthanide complexes. The intraconfigurational f ↔ f transitions from lanthanide metal ions, particularly those that obey magnetic dipole selection rules (J = 0 ± 1, except 0 ↔ 0), generally have large luminescence dissymmetry factors, δlum(λ), which facilitate the detection of the CPL signal. However, in recent years, there has been a growing interest in developing new CPL-active materials based on organic molecules or transition-metal complexes. Various strategies have been used to create CPL-emitting organic materials, including attaching chiral side chains to conjugated polymers, embedding organic chromophores in a chiral matrix, and developing inherently chiral conjugated molecules. The pinene group, a naturally occurring chiral element, is frequently introduced into pyridine ligands, leading to novel helical configurations and interesting circularly polarized luminescence.

CPL switches based on supramolecular systems are particularly interesting because CPL emission can be turned “on” and “off” by an external stimulus, including anion or ion pairing, temperature, solvent polarity, concentration, light irradiation, and mechanical stirring. Maeda et al. demonstrated that conformation changes by inversion (flipping) of two pyrrole rings as a result of anion binding can control the chiroptical properties of the anion receptors. Okano et al. showed that hydrogels with embedded Rhodamine B dye exhibited stir-induced circularly polarized luminescence, the...
Cyclometalated platinum(II) complexes have long been the subject of intense investigation as luminescent materials due to their intriguing stimulus-responsive structural and spectroscopic properties.\textsuperscript{12–15} Moreover, cyclometalated platinum(II) complexes have been proven to be biologically active as antitumor agents and DNA intercalators.\textsuperscript{16} Therefore, cyclometalated platinum(II) complexes with CPL emission would be very useful in organic light-emitting diodes (OLEDs) and chiral probes in biological systems. However, CPL emission from platinum(II) complexes has seldom been reported before.\textsuperscript{8b} Comparative studies of chiroptical memory, light-emitting devices, and multifunctional sensors.\textsuperscript{11}

In this work, we designed and synthesized a series of chiral cyclometalated platinum(II) complexes, namely, [Pt(−)-(−)-L\textsubscript{1})(Dmpi)]Cl ((−)-1), [Pt((+)-L\textsubscript{1})(Dmpi)]Cl ((+)-1), [Pt((−)-L\textsubscript{2})(Dmpi)]Cl ((−)-2), [Pt((+)-L\textsubscript{2})(Dmpi)]Cl ((+)-2), [Pt\textsubscript{3}((−)-L\textsubscript{1})(Dmpi)]\textsubscript{4}(ClO\textsubscript{4})\textsubscript{4} ((−)-3), and [Pt\textsubscript{3}((+)-L\textsubscript{1})(Dmpi)]\textsubscript{4}(ClO\textsubscript{4})\textsubscript{4} ((+)-3). [Pt\textsubscript{3}((−)-L\textsubscript{2})(Dmpi)]\textsubscript{4}(ClO\textsubscript{4})\textsubscript{4} ((−)-1) = (−)-4,5-pinepine-6′-phenyl-2,2′-bipyridine, [(+)-L\textsubscript{1}] = [(+)-4,5-pinepine-6′-phenyl-2,2′-bipyridine], (−)-L\textsubscript{2} = (−)-1,3-bis(2-(4,5-pinepine)pyridyl)benzene, [(+)-L\textsubscript{2}] = [(+)-1,3-bis(2-(4,5-pinepine)pyridyl)benzene, Dmpi = 2,6-dimethylphenyl isocyanide] (Chart 1). In aqueous solutions, (−)-1 and (−)-1 aggregate into one-dimensional helical chain structures through Pt···Pt π–π interactions. (−)-3 and (−)-3 represent a novel helical structure with Pt–Pt bonds. The formation of helical structures results in enhanced and distinct chiroptical properties. CPL was observed from the aggregates of (−)-1 and (+)-1 in water, as well as (−)-3 and (+)-3 in dichloromethane. The CPL activity can be switched reversibly for (−)-1 and (−)-1 or irreversibly for (−)-3 and (−)-3 by varying the temperature. These CPL-active materials based on our chiral Pt(II) complexes may have potential applications in photonic devices or biosensors.

![Chart 1. Molecular Structures of (−)-1, (+)-1, (−)-2, (+)-2, (−)-3, and (−)-3](image)

**EXPERIMENTAL SECTION**

**General Methods.** All reagents were purchased from commercial suppliers and used as received. Caution! Perchlorates are potentially explosive and must be handled with great care and in small amounts. Collision and friction must be avoided. 2,6-Dimethylphenyl isocyanide may cause irritation. All reactions should be performed in a fume hood. Compounds (−)-L\textsubscript{1}, (+)-L\textsubscript{1}, (−)-L\textsubscript{2}, (+)-L\textsubscript{2}, Pt((−)-L\textsubscript{1})Cl, Pt((+)-L\textsubscript{1})Cl, Pt((−)-L\textsubscript{2})Cl, and Pt((+)-L\textsubscript{2})Cl were prepared according to the methods reported previously.\textsuperscript{7,15} Mass spectra were acquired on an LCQ Fleet ESI mass spectrometer. The NMR spectra were obtained on a Bruker DRX-400, DRX-500, or DRX-600 spectrometer. Chemical shifts are reported relative to CH\textsubscript{3}OH/CD\textsubscript{3}OD (δ(H) = 3.31 ppm, δ(13C) = 49.0 ppm) or CH\textsubscript{2}Cl\textsubscript{2}/CD\textsubscript{2}Cl\textsubscript{2} (δ(H) = 5.32 ppm, δ(13C) = 53.8 ppm). Coupling constants are given in hertz. UV–vis absorption spectra were measured on a UV-3600 spectrophotometer (using a 10 mm quartz cell for a concentration of 5 \times 10^{-5} mol L^{-1}, using a 1 mm quartz cell for concentrations of 5 \times 10^{-4} and 5 \times 10^{-3} mol L^{-1}). The high-resolution transmission electron microscopy (HRTEM) images were obtained by employing a JEOL JEM-2100 transmission electron microscope with an acceleration voltage of 200 kV. The scanning electron microscopy (SEM) images were obtained on a Hitachi S-4800 scanning electron microscope with an acceleration voltage of 20 kV. The scanning electron microscopy (SEM) images were obtained on a Hitachi S-4800 scanning electron microscope at 20 kV. X-ray photoelectron spectroscopy (XPS) was carried out by a PHI 5000 VersaProbe (UHVAC-PHI), and binding energies were measured relative to the C 1s peak (284.8 eV) of internal hydrocarbon. The resonance light scattering (RLS) spectra were obtained by synchronously scanning the excitation and emission monochromators (namely, Δλ = 0.0 nm) of a Hitachi F-4600 fluorescence spectrophotometer in the wavelength region from 300 to 800 nm. CPL and total luminescence spectra were
recorded on an instrument described previously,\textsuperscript{17} operating in a differential photon-counting mode. The light source for excitation was a continuous wave 100 W xenon arc lamp from a Spex Fluorolog-2 spectrofluorometer, equipped with excitation and emission monochromators with a dispersion of 4 nm/mm (SPEX, 1681B). To prevent artifacts associated with the presence of linear polarization in the detection, a high-quality linear polarizer was placed in the sample compartment and aligned so that the excitation beam was linearly polarized in the direction of emission detection (z-axis). The key feature of this geometry is that it ensures that the molecules that have been excited and that are subsequently emitting are isotropically distributed in the plane (x, y) perpendicular to the direction of emission detection. The optical system detection consisted of a focusing lens, long-pass filter, and 0.22 μm monochromator. The emitted light was detected by a cooled EMI-9558B photomultiplier tube operating in photocounting mode. All measurements were performed with quartz cuvettes with a path length of 1.0 cm.

\[\text{[Pt(\text{--}L_2\text{Dmpzpy})(\text{ClO}_4)_2]} (\text{L} = \text{L}_1, \text{L}_2)\]

To a vigorously stirred solution of Pt(\text{--}L_2\text{Dmpzpy})(\text{ClO}_4) in 50 mL of dichloromethane was slowly added an equivalent amount of 2,6-dimethylphenyl isocyanide (1 mmol, 131 mg) in 50 mL of dichloromethane over 1 h. After reaction for 4 h at room temperature, the yellow phase was separated, the solvents were evaporated, and red powders were obtained (95%). The pure product was obtained after several recrystallizations in MeOH/H_{2}O solution. MS (ESI) (m/z): [M]^{+} calculated for C_{96}H_{98}N_{8}Pt_{3} (\text{ClO}_4), 692.0; found, 691.2. Anal. Calculated for C_{96}H_{98}N_{8}Pt_{3} (\text{ClO}_4): C, 49.61; H, 4.19; N, 4.74.\textsuperscript{13}C NMR (100 MHz, CD_{2}Cl_{2}, 273.15 K): δ 9.29 [s, 2H, H(17)], 7.79 [s, 2H, H(11)], 7.75 [d, J = 7.6 Hz, 2H, H(5)], 7.39–7.44 [m, 8H, H(8), H(4), H(38), H(14)], 7.25 [d, J = 7.6 Hz, 4H, H(37), H(39)], 7.21 [t, J = 7.6 Hz, 2H, H(48)], 7.04 [d, J = 7.6 Hz, 4H, H(47), H(49)], 6.62 [d, J = 7.2 Hz, 2H, H(5)], 3.36–3.41 [m, 2H, H(19b)], 3.20–3.25 [m, 2H, H(19a)], 2.71–2.75 [m, 6H, H(21b), H(29), H(26b)], 2.46 [s, 12H, H(41), H(42)], 2.24–2.44 [m, 2H, H(20)], 2.22–2.32 [m, 6H, H(22), H(26a), H(28b)], 2.17 [s, 12H, H(51), H(52)], 1.98–2.02 [m, 24H, H(27), H(28), H(30), H(31)], 1.13 [s, 6H, H(32)], 1.06 [s, 2H, H(21a)], 0.57 [s, 6H, H(25)], 0.33 [s, 6H, H(31)], −0.53 [m, 2H, H(28a)].\textsuperscript{15}C NMR (100 MHz, CD_{2}Cl_{2}, 273.15 K): δ 164.2 [C(7)], 160.0 [C(13)], 152.6 [C(9)], 149.5 [C(15)], 148.2 [C(43)], 148.0 [C(17)], 147.7 [C(33)], 147.2 [C(11)], 145.0 [C(8)], 144.5 [C(10)], 143.0 [C(22)], 142.8 [C(16)], 136.6 [C(36), C(40)], 136.0 [C(46), C(50)], 131.9 [C(4)], 131.5 [C(3)], 131.1 [C(48)], 129.1 [C(37), C(39)], 128.7 [C(47), C(49)], 127.9 [C(38)], 127.1 [C(5)], 125.2 [C(45)], 124.7 [C(35)], 123.4 [C(14)], 122.3 [C(11)], 119.4 [C(6)], 44.5 [C(22)], 43.9 [C(29)], 39.6 [C(20), 39.2 [C(30)], 39.0 [C(27)], 38.8 [C(25)], 34.0 [C(19)], 33.1 [C(26)], 32.0 [C(21)], 30.1 [C(28)], 25.6 [C(24)], 25.2 [C(32)], 22.1 [C(31)], 21.4 [C(25)], 19.2 [C(41), C(42)], 18.8 [C(51), C(52)]. XPS (eV): 73.3 [Pt 4f_{7/2}], 76.7 [Pt 4f_{5/2}].

**X-ray Structure Determination.** Yellow needles of complex (−)–1 were grown in acetonitrile/dichloromethane (1/1, v/v) solution at room temperature. Yellow rods of racemic 2-OTI were obtained by diffusion of diethyl ether into the acetonitrile solution of a mixture of (−)–2·OTF and (+)–2·OTF at room temperature, whereas green-yellow blocks of complexes (−)–3 and (+)–3 were isolated by recrystallization in dichloromethane solution at 273 K. The structure determination of (−)–1 is unsatisfactory, but it does provide accurate connectivity and packing information as well as valid bond distances and angles involving the heavy atoms in the structure. Single-crystal X-ray diffraction measurements were carried out on a Bruker SMART APEX charge-coupled device (CCD)-based diffractometer operating at room temperature. Intensities were collected with graphite-monochromatized Mo Kα radiation (\(λ = 0.71073\) Å) operating at 50 kV and 30 mA using the \(0/2θ\) scan mode. The data reduction was made with the Bruker SADABS program.\textsuperscript{19} Absorption corrections were performed using the SADABS program.\textsuperscript{20} The structures were solved by direct methods and refined on \(I^2\) by full-matrix least-squares using SHEXL-97 with anisotropic displacement parameters for all non-hydrogen atoms in all structures. Hydrogen atoms bonded to the carbon atoms were placed in calculated positions and refined in the riding mode, with C–H = 0.93 Å (methane) or 0.96 Å (methyl) and \(U_{eq}(H) = 1.2U_{eq}(C_{methyl})\) or \(U_{eq}(H) = 1.5U_{eq}(O)\). The water hydrogen atoms were located in the difference Fourier maps and refined with an O–H distance restraint of 0.85(1) Å and \(U_{eq}(H) = 1.5U_{eq}(O)\). All computations were carried out using the SHELXL-97 program package.\textsuperscript{21} CCDC 989466–989469 and 1006137 contain the supplementary crystallographic data.
graphic data for this paper. These data can be obtained free of charge from The Cambridge Crystallographic Data Centre via www.ccdc.ac.uk/data_request/cif.

■ RESULTS AND DISCUSSION

Synthesis and Characterization. The precursors Pt(−)-L1)Cl and Pt((−)-L2)Cl were prepared according to the methods reported previously.12c,d Complexes (−)-1 and (−)-2 can be obtained through ligand metathesis reaction of Pt((−)-L1)Cl and Pt((−)-L2)Cl with 2,6-dimethylphenyl isocyanide at room temperature.14b The unprecedented pair of trimeric helical complexes (−)-3 and (+)-3 were isolated at 273 K when excess 2,6-dimethylphenyl isocyanide was reacted with complexes (−)-2 and (+)-2, followed by addition of excess AgClO4 aqueous solution. Compounds (+)-1, (+)-2, and (+)-3 were obtained with the same procedures for (−)-1, (−)-2, and (−)-3, respectively. All the complexes ((−)-1, (−)-2, and (−)-3) have been fully characterized by elemental analysis, NMR, XPS, and ESI mass spectrometry (Figures S1–S22, Supporting Information).

Except (−)-1 and (+)-1, all these complexes are insoluble in water. (−)-1 and (+)-1 show aggregation behaviors in aqueous solutions. The 1H NMR spectra of (−)-1 show broadening and upfield shifts in the mixed solution of CD2OD and D2O (Figure 1), suggesting the presence of molecular association through

Pt···Pt, π−π, and hydrophobic–hydrophobic interactions when D2O was added. Similar upfield shifts have also been found in other related deaggregation–aggregation systems, in which broad and featureless NMR signals are exhibited in the more polar solvents.22 Upon increasing the concentration of water to ca. 10−2 mol L−1, (−)-1 and (+)-1 form hydrogels (Figure S23, Supporting Information). SEM and TEM also demonstrate fibrillar structures with an in-plane orientation, characteristic for aggregation of platinum(II) complexes (Figure S24, Supporting Information).22b No aggregation effect was observed for (−)-2, (+)-2, (−)-3, and (+)-3.

(−)-3 and (+)-3 are very stable in the solid state but partially decompose in dichloromethane solution at room temperature. As shown in Figure 2, complex (−)-3 starts to partially decompose at 293 K and turns into complex (−)-2, as evidenced by additional peaks at ca. 8.30, 1.48, and 0.76 ppm after decomposition. Therefore, all the spectroscopic data of (−)-3 and (+)-3 were measured at 273 K except where otherwise noted.

Crystal Structures. Complex (−)-1 crystallizes in the P21 space group of the monoclinic system (Table 1). Two isolated molecules are included in the asymmetrical unit of complex (−)-1 (Figure 3a). The Pt−C (1.86−2.02 Å) and Pt−N (1.91−2.12 Å) (Table S1, Supporting Information) bond lengths and relevant angles are in the typical ranges for Pt(C=N=N) isocyanide complexes (C=N=N denoted as (−)-L2).14 The rings of 2,6-dimethylphenyl isocyanide are almost coplanar with the Pt(C=N=N) unit, with dihedral angles between the two planes of 2.93° and 3.94°. The molecules are further slip-stacked in head-to-head mode along the a axis with effective π−π contacts (3.379 Å) (Figure 3b). The molecules are aligned in a zigzag style. The Pt−Pt distances between adjacent Pt atoms are 3.812 and 4.129 Å, respectively, indicating that no Pt−Pt interactions are involved.

Attempts to grow single crystals for (−)-2 and (+)-2 were not successful. However, we were able to obtain a racemic crystal of 2-OTf by substituting the Cl with OTf. As shown in Figure S25 (Supporting Information), the coordination environment of Pt(II) atoms in 2-OTf is very similar to that of (−)-1, showing a slightly distorted square-planar configuration. However, the torsion angle (17.33°) between the ring of 2,6-dimethylphenyl isocyanide and the Pt(N=C=N) plane in 2-OTf is more salient relative to that of (−)-1 (N=C=N denoted as (−)-L2). In the packing diagram of 2-OTf, two isolated molecules ((−)-2 and (+)-2 cations appear alternately) are slip-stacked in head-to-tail mode along the c axis (Figure S26, Supporting Information). Owing to the steric hindrance of pinene groups in 2-OTf, the neighbor molecules could not approach each other closely in a face-to-face manner and the nearest Pt···Pt distance is 6.01 Å, suggesting that neither distinct Pt···Pt nor π−π interactions are present.

The crystals of (−)-3 and (+)-3 recrystallized in CHCl3 at 273 K reside in the P21212 space group of the orthorhombic system (Table 1). In the trimeric structure of (−)-3, two Pt ions (top and bottom) coordinate with two carbon atoms of 2,6-dimethylphenyl isocyanide and one carbon atom and one nitrogen atom of the N=C=N ligands. These two Pt ions are

Figure 1. 1H NMR spectra of (−)-1 in a mixed solution of CD2OD and D2O in the ratios shown (Bruker DRX-500, rt).

Figure 2. 1H NMR spectra of (−)-3 in CD2Cl2 at different temperatures and (−)-2 in CD2Cl2 at room temperature (Bruker DRX-500).

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bridged by the central Pt ion, which is coordinated to nitrogen atoms of the N^C^N ligands (Figure 3c). The coordination number of the top and bottom Pt ions (Pt1 and Pt1A in Figure 3c) is five, and both of them exhibit the distorted tetragonal pyramid geometry, while the coordination environment of the central Pt ion (Pt2 in Figure 3c) with four-coordinate atoms is a distorted square-planar geometry. For the same N^C^N ligand, one side pyridine ring presents a twist of 44.63° relative to the central benzene plane, and the conformation of such complexes exhibit characteristic absorption bands for typical cyclometalated platinum(II) complexes, i.e., intraligand (IL) transitions (ε > 10^3 L mol^-1 cm^-1) in the region of 200–310 nm and metal-to-ligand charge transfer (MLCT) mixed with ligand-to-ligand charge transfer (LLCT) transitions (ε > 10^3 L mol^-1 cm^-1) in the region of 330–450 nm.13,14 The UV–vis absorption spectra of (-)-1, (+)-1, (-)-2, (+)-2, (-)-3, and (+)-3 are shown in Figure 4, and spectroscopic data are summarized in Table S2 (Supporting Information). All these complexes exhibit characteristic absorption bands for typical cyclometalated platinum(II) complexes, i.e., intraligand (IL) transitions (ε > 10^3 L mol^-1 cm^-1) in the region of 200–310 nm and metal-to-ligand charge transfer (MLCT) mixed with ligand-to-ligand charge transfer (LLCT) transitions (ε > 10^3 L mol^-1 cm^-1) in the region of 330–450 nm.13,14 The high-energy emissive state of (-)-3′ can be mainly assigned as the ligand-centered 3MLCT and 3LLCT excited states,14 while the highly structured emission spectra of (-)-3′, as shown in Figure S27 and Figure S28. Moreover, as compared to (-)-3, the molecules are more compact along the b axis in (-)-3′ (Figure S27 and Figure S28), which can also be reflected by the difference in their cell lengths.

**UV–Vis Absorption and Emission Spectra.** The UV–vis absorption spectra of (-)-1, (+)-1, (-)-2, (+)-2, (-)-3, and (+)-3 are shown in Figure 4, and spectroscopic data are summarized in Table S2 (Supporting Information). All these complexes exhibit characteristic absorption bands for typical cyclometalated platinum(II) complexes, i.e., intraligand (IL) transitions (ε > 10^3 L mol^-1 cm^-1) in the region of 200–310 nm and metal-to-ligand charge transfer (MLCT) mixed with ligand-to-ligand charge transfer (LLCT) transitions (ε > 10^3 L mol^-1 cm^-1) in the region of 330–450 nm.13,14 The high-energy emissive state of (-)-1 in methanol solution (λ_max = 321 nm) can be assigned to the combination of MLCT and LLCT excited states, while the highly structured emission spectra of (-)-2 and (-)-3 can be mainly assigned as the ligand-centered π→π* state.13,14,16 The profiles of emission spectra of complexes (-)-2 and (-)-3 are similar in dichloromethane solution (Figure S30), indicating that the presence of Pt→Pt bonds in (-)-3 and (+)-3 does not change the nature of the frontier orbitals.

When (-)-1 was dissolved in water, the absorption intensity in the range of 200–380 nm decreased while a new low-energy absorption in the range of 420–550 nm emerged with

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**Table 1. Crystallographic Data of (−)-1, 2-OTf, (−)-3, (+)-3, and (−)-3’**

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increasing H₂O content, resulting from different degrees of aggregation in mixed solvents with various ratios (Figure 5). One clear isosbestic point at 366 nm can be distinguished, indicating a clean conversion between the nonaggregate state and aggregate species. The color difference of the solution can intuitively reflect the aggregation process (Figure 6). As the content of H₂O increased, the color of the solution changed from pale yellow to orange. The orange solution can be attributed to formation of aggregate species though Pt···Pt, π−π, and hydrophobic−hydrophobic interactions. As a consequence, the absorption at around 500 nm is assigned to a metal−metal-to-ligand charge transfer (MMLCT) transition.¹⁴ When excited at 420 nm, a low-energy emission (λ<sub>em</sub> = 640 nm) was observed (Figure S29, Supporting Information) in aqueous solution which can be attributed to the 3MMLCT excited state.¹⁴b Similarly, the emission spectra of (−)-1 in water were concentration-independent, and the emission maximum remained at 640 nm.

This solvent-induced aggregation of (−)-1 was further evidenced by the enhanced signal of RLS spectra (Figure S33, Supporting Information). Upon increasing the H₂O content, the intensity of the RLS signal in the range from 350 to 600 nm was significantly strengthened, indicating the formation of aggregates in the solution and therefore the chromophores were strongly coupled.²⁴ As shown in Figure 7, the aggregation of (−)-1 is also sensitive to temperature. Upon increasing the temperature, the absorption peak at 460 nm associated with the MMLCT transition shows a slight drop in intensity; concomitantly the peaks at 350 and 250 nm related to the MLCT/LLCT transition become more intense. An isosbestic point can be found at 366 nm, resembling the solvent-induced absorption variance. These changes can be ascribed to the attenuation of Pt···Pt and π−π interactions at higher temperature, leading to well-defined aggregates have formed even in the diluted solution and therefore the aggregation is insensitive to concentration changes. Similar observations have been reported in the previous work that cyclometalated platinum(II) complexes can oligomerize in water even at a dilution of 2 × 10⁻⁵ mol·L⁻¹.¹⁴b Similarly, the emission spectra of (−)-1 in water were concentration-independent, and the emission maximum remained at 640 nm.
the deaggregation of the molecules. The emission intensity also presents a drastic drop, and the emission maximum has a slight blue shift upon increasing the temperature (Figure S34, Supporting Information), which further corroborates the deaggregation of molecules.

The UV–vis absorption spectra and emission spectra of (−)-3 measured at different temperatures above 273 K show a continuous variance due to decomposition (Figure S35 and S36, Supporting Information). As shown in Figure S35, when the temperature changes from 273 to 294 K, the absorbance at 394 nm decreases, while the absorptions of the peaks at 331 and 291 nm increase. Although the emission maximum of (−)-3 shows a small change upon increasing the temperature, the emission intensity of the shoulder peak at 525 nm exhibits a distinct drop (Figure S36).

**ECD Spectra.** Figure 4 shows the ECD spectra of (−)-1, (+)-1, (−)-2, (+)-2, (−)-3, and (+)-3. The ECD spectrum of (−)-1 shows moderate Cotton effects (positive, 240, 284, and 346 nm; negative, 303 nm) in methanol solution; however, a significantly different and enhanced ECD spectrum (positive, 250, 300, 370, and 480 nm; negative, 320 and 420 nm) was observed in water. The anisotropy $g_{abs}$ factor becomes higher with increasing H$_2$O content (Figure S5), and it reaches $+0.0022$ at 377 nm and $-0.0046$ at 420 nm in pure aqueous solution, which are 1 order of magnitude larger than those of the methanol solution at the corresponding wavelengths. Owing to the dissociation of molecules upon increasing the temperature, the chiral signals of (−)-1 in water solution decrease upon increasing the temperature of the sample from 298 to 353 K (Figure 7). When monitored at 370 nm, the chiral signals of (−)-1 can be reversibly tuned by changing the temperature, and no distinct degeneration can be found, indicating the persistence of temperature-induced ECD switching (Figure S36).

**Figure 4.** (a) UV–vis absorption and ECD spectra of (−)-1 and (+)-1 in MeOH and H$_2$O at 5 × 10$^{-5}$ mol·L$^{-1}$. (b) UV–vis absorption and ECD spectra of (−)-2 and (+)-2 in dichloromethane at 5 × 10$^{-5}$ mol·L$^{-1}$ (294 K) and (−)-3 and (+)-3 in dichloromethane at 2.5 × 10$^{-5}$ mol·L$^{-1}$ (273 K).

**Figure 5.** UV–vis absorption and ECD spectra and $g_{abs}$ factors of (−)-1 (5 × 10$^{-5}$ mol·L$^{-1}$) in a mixed solution of MeOH and H$_2$O in the ratios shown.

**Figure 6.** Photograph of (−)-1 (10$^{-4}$ mol·L$^{-1}$) in a mixed solution of MeOH and H$_2$O in the ratios shown (under ambient light).
bands at 350 and 395 nm in addition to the series of Cotton corresponding wavelengths. The intensity of the Cotton e of (−)-3 decreases upon increasing the temperature from 273 to 294 K owing to decomposition (Figure S35, Supporting Information).

The significant enhancement of chirality of (−)-1 and (+)-1 from MeOH to H₂O solutions, and from (−)-2 and (+)-2 to (−)-3 and (+)-3, could be elucidated from their structural changes. For (−)-1 and (+)-1 in MeOH solutions, the molecules are dissociated and the chirality of the slightly distorted square-planar complexes is rather weak. When (−)-1 and (+)-1 are dissolved in water, the molecules are inclined to aggregate into one-dimensional chain structures through Pt···Pt, π−π, and hydrophobic−hydrophobic interactions, as already reported in the literature and also evidenced by the appearance of an MMLCT in the absorption and emission spectra. Due to the steric hindrance of the bulky pinene groups in (−)-1 and (+)-1, the adjacent molecules have to be staggered from each other alongside the Pt···Pt chain, thus forming a helix and enhancing the chiral environment. Increasing the temperature disrupts the helical aggregation and subsequently results in a less pronounced chiral environment. As a consequence, the strong ECD signals of (−)-1 and (+)-1 in water should come from helical chirality of aggregated chromophores rather than intrinsic chirality of the separated molecules. Intermolecular Pt···Pt, π−π, and hydrophobic−hydrophobic interactions can be strengthened or weakened by variations of the solvent (MeOH/H₂O) ratios and temperature, producing aggregates of (−)-1 and (+)-1 with different helicity degrees and resulting in different ECD signals. For (−)-3 and (+)-3, the helical structures, which enhance the chiroptical activity, have been revealed by the single-crystal X-ray diffraction studies.

CPL. According to the temperature/solvent-induced variance of chirality in the ground state as evidenced in the ECD spectra, the chiral signals were enhanced through formation of an intramolecular helical structure or intermolecular helical aggregation. Therefore, we envision that interesting CPL activity in the corresponding excited state would be present. Therefore, the CPL spectra of (−)-1 and (+)-1 were measured in 1 mM MeOH and H₂O solutions, whereas the CPL spectra of (−)-2, (+)-2, (−)-3, and (+)-3 were measured in 1 mM dichloromethane solutions. Although (−)-1 and (+)-1 in MeOH solutions and (−)-2 and (+)-2 in dichloromethane solutions did not exhibit any CPL activity at 295 K, we were able to observe a measurable CPL activity for (−)-1 and (+)-1 in aqueous solutions at 295 K, as well as for (−)-3 and (+)-3 in dichloromethane solutions at 278 K (Figure 9). The values around the maximum emission wavelength are −0.0018/ +0.0012 for (−)-1/(+)−1 at 295 K and +0.0024/−0.0004 for (−)-3/(+)−3 at 278 K, respectively. Interestingly, the CPL activity was lost for aqueous solutions of (−)-1 and (+)-1 at 353 K (no conclusive CPL signal was detected; the data points are scattered and non-mirror-symmetric) (Figure S37, Supporting Information) but can be recovered upon cooling to 295 K. The CPL activity of the MeOH solution of (−)-3 and (+)-3 was weakened at 295 K (gₑm = +0.0016/−0.0016) as compared to that at 278 K (Figure S37), in agreement with the above-mentioned observation that part of the helical trinuclear complexes were decomposed into mononuclear complexes with weak chirality upon increasing the temperature. This process is irreversible.

It is noteworthy that, although CPL signals of platinum(II) complexes have been reported recently, the Pt atom is not the stereogenic center in these systems, and the CPL activity comes...
from the helical organic luminophore. However, in our study, the Pt atoms play a key role in the formation of helical aggregations or helical structures, which control the circularly polarized luminescent behavior of the chiral platinum(II) complexes. The asymmetric centers on the pinene groups are remote from the metal atom and have little direct effect on the CPL spectra, but they could have an influence on chiral coordination or the helical arrangement, causing measurable CPL signals. We have also observed the CPL-active $^3$MMLCT excited state, which is unprecedented in the literature to the best of our knowledge. Moreover, we demonstrated a very novel temperature-dependent tuning of the CPL activity.

**CONCLUSIONS**

In conclusion, two chiral cyclometalated platinum(II) complexes, [Pt((−)-L$_2$)(Dmp)]Cl ((−)-1) and [Pt((+)-L$_2$)(Dmp)]Cl ((+)-1), and [Pt((−)-L$_2$)(Dmp)]Cl ((−)-2) and [Pt((+)-L$_2$)(Dmp)]Cl ((+)-2), have been synthesized and characterized. (−)-1 and (+)-1 aggregated in water via Pt···Pt, π···π, and hydrophobic–hydrophobic interactions. (−)-2 and (+)-2 reacted with excess 2,6-dimethylphenyl isocyanide to give the trimetallic helical complexes [Pt$_3$((−)-L$_2$)$_2$(Dmp)]$_4$Cl$_4$ (CI0)$_4$ ((−)-3) and [Pt$_3$((+)-L$_2$)$_2$(Dmp)]$_4$Cl$_4$ (CI0)$_4$ ((+)-3). Significant enhancement and/or alteration of chirality was observed for (−)-1 and (+)-1 in aqueous solutions and (−)-3 and (−)-3, as revealed in the ECD spectroscopic study. The enhancement and/or alteration of chirality for (−)-1 and (−)-1 in aqueous solutions and (−)-3 and (−)-3 led to a detectable CPL activity for these platinum(II) complexes. More interestingly, the CPL activity can be switched reversibly (for (−)-1 and (+)-1) or irreversibly (for (−)-3 and (−)-3) due to deggregation or decomposition upon increasing the temperature. Extension of this work to other chiral luminescent Pt(II) complexes and studies aimed at utilizing these complexes in photonic devices or biosensors are under way and will be reported in due course.

**ASSOCIATED CONTENT**

Supporting Information

Synthesis and characterization of complexes (+)-1, (+)-2, and (+)-3; X-ray crystallographic data for complexes (−)-1, 2-OTf, (−)-3, (+)-3, and (−)-3$^3$ in CIF format, NMR, ESI mass, and XPS spectra for the complexes, photographs and SEM and TEM images of hydrogels of (−)-1, X-ray crystal structures and packing diagrams of 2-OTf, (−)-3, and (−)-3$^3$, emission spectra of (−)-1, (−)-2, and (−)-3, RLS spectra of (−)-1, UV–vis absorption and ECD spectra and g$_{av}$ factors of (−)-3 at various temperatures, CPL and total luminescence spectra of (−)-1 and (+)-1 in 1 mM aqueous solution at 353 K and (−)-3 and (+)-3 in 1 mM MeOH solution at 295 K, selected bond lengths and angles of complexes (−)-1, 2-OTf, (−)-3, (+)-3, and (−)-3$^3$, and absorption and emission data of complexes (−)-1, (−)-2, and (−)-3. This material is available free of charge via the Internet at http://pubs.acs.org.

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The authors declare no competing financial interest.

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**REFERENCES**


Figure 9. Circularly polarized luminescence (upper curves) and total luminescence (lower curves) spectra of (a) (−)-1 and (+)-1 (left panel) in 1 mM aqueous solutions at 295 K and (b) (−)-3 and (+)-3 (right panel) in 1 mM dichloromethane solutions at 278 K.
The degree of CPL is given by the luminescence dissymmetry ratio, \( g_{\Delta}(\lambda) = 2(1 - \epsilon_\lambda) / (\epsilon_\lambda + 1) \), where \( \epsilon_\lambda \) and \( \epsilon_\lambda' \) refer, respectively, to the intensities of the left and right circularly polarized emissions. The ECD anisotropy factor is defined as \( g_{\Delta} = \Delta \epsilon / \epsilon \), where \( \Delta \epsilon \) is the molecular extinction coefficient of the ECD spectrum and \( \epsilon \) is the molecular extinction coefficient of the UV–vis absorption spectrum.


