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Isolation and phototransformation of enantiomerically pure iridium(III) bis[(4,6-difluorophenyl)pyridinato-N,C²]picolinate[†]

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Here we report the resolution of phosphorescent light-emitting iridium(III) bis[(4,6-difluorophenyl)pyridinato-N,C²]-picolinate into its respective enantiomers by using chiral HPLC and the photo-induced transformation of the isolated enantiomers.

Phosphorescent light-emitting diodes (LEDs) are drawing more attention than fluorescent LEDs^{1–6} because the fluorescent LEDs have shown maximum internal quantum yields of *ca.* 25 while the phosphorescent LEDs can in theory reach internal quantum yields of nearly 100%.¹ Therefore, various iridium complexes have been studied; with the blue light-emitting iridium bis[(4,6-difluorophenyl)pyridinato-N,C²]picolinate (FIRpic) being a representative complex.² FIRpic is a heteroleptic, octahedral complex, for which four geometric isomers are possible, and the complex from commercial sources is known to be the *mer* isomer with *cis* C–C and *trans* N–N geometry, for which Λ - and Δ -enantiomers exist (Fig. 1, left pair).

Herein we present the isolation of the Λ and Δ -enantiomers of FIRpic by preparative HPLC and their photo-induced transformation to the enantiomers of the *mer-cis* isomer (Fig. 1, right pair). Several examples of enantiomers of octahedral Ir complexes have been reported.^{3–10} Recently, the isolation and characterization of chiroptical properties of FIRpic enantiomers were reported by Cannazza and co-workers¹⁰ while the separation of these enantiomers in the present work was conducted independently. The phototransformation of an enantiomerically pure FIRpic complex has not been previously

accomplished, to the best of our knowledge; the ability to carry out this transformation is important for developing applications for circularly polarized light (CPL) emitter designs where excited-state stability of light-emitting species would play a crucial role. In addition to LED applications, Ir complexes have been used as valuable catalysts in excited states¹¹ where phototransformation of an Ir complex would have significant influences on the catalyzed reactions.

Full details of the current experiments are given in the ESI.[†] Preparative chromatographic resolution experiments were conducted using a Daicel ChiralPak IC column (25 cm \times 2 cm (i.d.)), and analytical resolution using analytical-scale ChiralPak IC, IA, and OD columns (25 cm \times 0.46 cm (i.d.)). These columns use polysaccharide derivatives as chiral selectors and have been applied for the resolution of a wide variety of racemic compounds.¹² When carrying out analytical-scale resolution on the IA column, a mixture of hexane, 2-propanol, and dichloromethane as eluent led to a clearly better separation than did a mixture of hexane and 2-propanol (Fig. S1 in ESI[†]), and slightly better resolution than did a mixture of hexane, 2-propanol, and chloroform (Fig. S2 in ESI[†]). In the analytical resolution on the analytical IC column using a mixture of hexane, 2-propanol, and dichloromethane, base-line separation was achieved where the ratio of the three components affected the resolution profiles (Fig. S3 in ESI[†]). On the basis of these observations, preparative resolution was performed using a 40/

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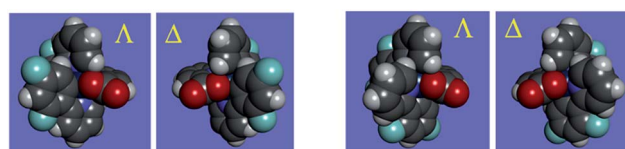


Fig. 1 Structures of FIRpic: Λ - and Δ -enantiomers of *mer-cis*-C–C–*trans*-N–N isomer (left pair) and those of the *mer-cis* isomer (right pair) optimized by DFT calculations. Detailed conditions of calculations are found in ESI.[†]



40/20 (v/v/v) mixture of hexane, 2-propanol, and dichloromethane as the eluent.

The Chiralcel OD column was also effective in resolving Firpic into its enantiomers; the circular dichroism (CD) signs at 250 nm of the first- and second-eluting enantiomers were opposite those reported in ref. 10 under the same separation condition (Fig. S4 in ESI†).

Fig. 2A(1–4) show the analytical high-performance liquid chromatography (HPLC) profiles obtained with the IA column, and CD, UV spectra, and CPL emission spectra¹³ of the isolated enantiomers, and theoretical CD spectra of Λ - and Δ -enantiomers obtained from density functional theory (DFT) calculations. The HPLC chromatograms (Fig. 2A) indicated that the enantiomers obtained by preparative HPLC resolution of Firpic were almost enantiomerically pure; the separation factor in Fig. 1A-2 (α) was 1.29. The isolated enantiomers exhibited mirror-image CD spectra in methanol over the visible and UV ranges (Fig. 2B).

Furthermore, the isolated enantiomers showed clear, mirror-image CPL emission spectra¹³ in the range 450–550 nm on excitation at 286 nm (Fig. 2D) where maximum anisotropy (g_{lum}) values were 2.0×10^{-3} at 489 nm (Λ -enantiomer) and 2.1×10^{-3} at 490 nm (Δ -enantiomer); anisotropy in emission (g_{lum}) is defined as $2(I_L - I_R)/(I_L + I_R)$ where I_L and I_R are L- and R-CPL emission intensities, respectively. These results showed the isolated enantiomers to have enantiomeric structures both in the ground state and excited states. The clear CPL emission spectra of the Firpic enantiomers indicated they are potentially applicable for CPL-emitting LEDs.

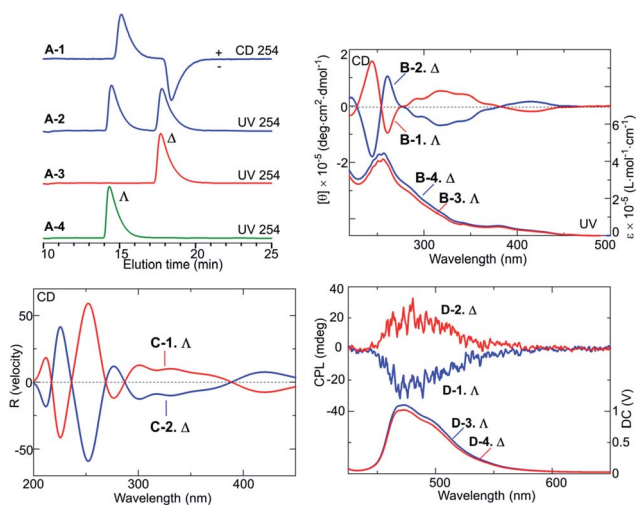


Fig. 2 Analytical results on the racemic *mer-cis-C-C-trans-N-N* Firpic isomer: HPLC profiles of the racemic mixture ((A-1) CD detection at 254 nm; (A-2) UV detection at 254 nm, Δ -enantiomer ((A-3) UV detection at 254 nm), and Λ -enantiomer ((A-4) UV detection at 254 nm) and experimental CD ((B-1) Λ ; (B-2) Δ) and UV ((B-3) Λ ; (B-4) Δ) spectra, theoretical CD spectra ((C-1) Λ ; (C-2) Δ), and CPL emission spectra ((D-1) Λ ; (D-2) Δ) with DC spectra (total emission spectra) ((D-3) Λ ; (D-4) Δ) of the Λ - and Δ -enantiomers. Conditions: eluent = hexane/2-PrOH/ CH_2Cl_2 mixture (65/15/20 (v/v/v)), flow rate = 0.5 mL min^{-1} (A); solvent = MeOH, conc. = 1.0×10^{-4} M, cell path = 1 mm (B); solvent = THF, conc. 2.0×10^{-5} M (D).

In addition, by comparing the experimental and theoretical CD spectra, the absolute configurations of the resolved enantiomers were determined. The theoretical spectra in Fig. 2C were obtained by carrying out TDDFT calculations with the B3LYP functional^{14,15} and the SDD basis set¹⁶ for Ir using the Gaussian 09 program package.¹⁷ The theoretical CD spectrum showed a rather sharp and intense negative Cotton band and a broad, weaker negative Cotton band in the range of ca. 300–380 nm, which roughly corresponded to the spectrum of the first enantiomer eluted in the chiral HPLC resolution of Firpic (Fig. 2B-1). On the basis of these observations, the first-eluting enantiomer from the HPLC column was assigned to be the Λ -enantiomer and the second-eluting enantiomer was assigned to be the Δ -enantiomer. Thus, the Λ - and Δ -enantiomers of the *mer-cis-C-C-trans-N-N* isomer were successfully isolated and their absolute configurations were assigned.

Furthermore, the isolated Δ - and Λ -enantiomers were found to undergo photo-isomerization and degradation. Since optically active Ir complexes may be used as CPL LED materials, the stability levels of such complexes in their excited states would be an important criterion in molecular design. In order to shed light on this aspect, the Δ - and Λ -enantiomers were independently irradiated in a MeOH solution (1.0×10^{-4} M) using non-polarized light (NPL) (ca. 200 mW cm^{-2}) from an Hg-Xe lamp in a MeOH solution under an N_2 atmosphere.

Fig. 3A and B show the changes in the CD and UV spectra of the Λ - and Δ -enantiomers, respectively, upon irradiation under an N_2 atmosphere. CD spectral intensity remarkably decreased upon irradiation, and the shapes of the UV spectra also changed. The changes in UV spectra suggested that the CD

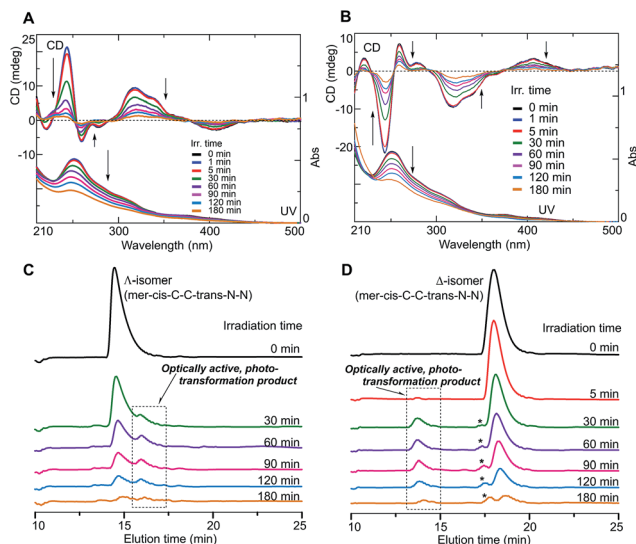


Fig. 3 CD and UV spectral changes of the Λ -enantiomer (A) and Δ -enantiomer (B) (*mer-cis-C-C-trans-N-N*) in MeOH on NPL irradiation, and HPLC profile changes detected at UV 254 nm for the Λ -enantiomer (C) and Δ -enantiomer (D). Conditions: conc. = 1.0×10^{-4} M, cell path = 1 mm (A and B); eluent = hexane/ CH_2Cl_2 /2-PrOH mixture (v/v/v), flow rate = 0.5 mL min^{-1} , sample conc. before irradiation = 1.0×10^{-4} M, injected sample volume = 10 μL (C and D). * denotes optically inactive impurity signals.



spectral changes were not due to simple racemization but some other structural transformation. This assumption was supported by the HPLC profiles of the irradiated samples (Fig. 3C and D); upon irradiation, the intensity of the Λ - and Δ -enantiomer signals decreased and, at the same time, new signals emerged at retention times of 16.5 min and 13.5 min, respectively, for the systems with the Λ - and Δ -enantiomers: the new species isolated using HPLC exhibited clear mirror image CD spectra (Fig. 4A-1 and A-2).

Note the slightly different intensities of the CD spectra in Fig. 2B-1 and B-2; the Λ -isomer yielded a slightly lower intensity than did the Δ -isomer. A small amount of the Λ -isomer might have undergone the phototransformation before its CD spectrum was acquired.

In addition, a new species was obtained from the Δ -enantiomer as indicated from mass spectrometry results, which showed a main signal at a mass number (m/z) of 718, corresponding to an ion of Flrpic in the form of an $[M + Na]^+$ sodium adduct, along with other weaker signals (Fig. S6 in ESI[†]). These results suggested that the Λ - and Δ -enantiomers of the *mer-cis-C-C-trans-N-N* isomer underwent, under the present experimental conditions, stereomutation to enantiomers of one of the three other geometric isomers of Flrpic, *i.e.*, *mer-cis*, *fac-cis*, and *mer-trans-C-C-cis-N-N*, rather than simple racemization.

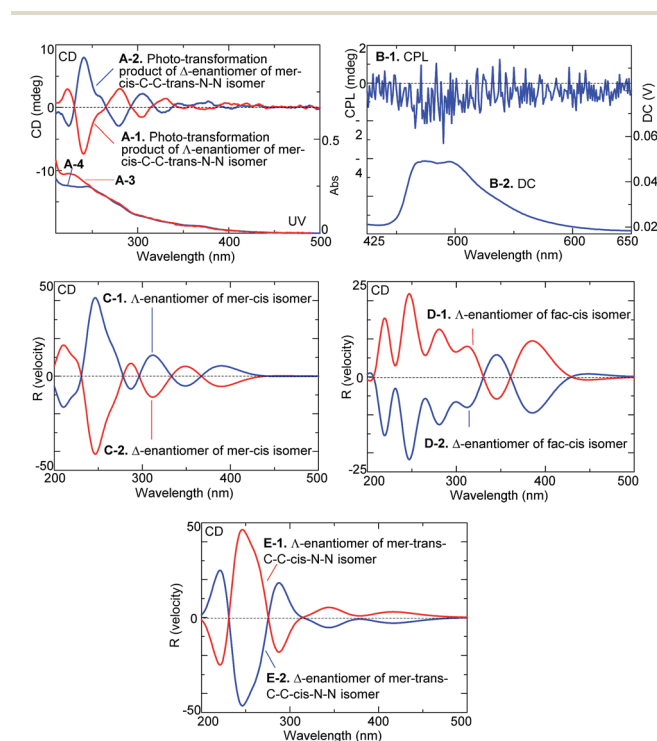


Fig. 4 CD/UV spectra of the enantiomeric compounds obtained by chiral HPLC separation of the photo-irradiation products of the Δ -enantiomer ((A-1) CD; (A-3) UV) and Δ -enantiomer ((A-2) CD; (A-4) UV) of *mer-cis-C-C-trans-N-N* isomer, CPL/DC spectra of the photo-irradiation product of Δ -enantiomer of *mer-cis-C-C-trans-N-N* isomer ((B-1) CPL; (B-2) DC), and theoretical CD spectra of enantiomers of the *mer-cis* isomer ((C-1) Λ ; (C-2) Δ), of the *fac-cis* isomer ((D-1) Λ ; (D-2) Δ) and of the *mer-trans-C-C-cis-N-N* isomer ((E-1) Λ ; (E-2) Δ).

Moreover, the phototransformation product of the Δ -enantiomers of the *mer-cis-C-C-trans-N-N* isomer yielded CPL emission spectra with a maximum g_{lum} value appearing to be at about 2×10^{-3} (Fig. 4B-1 and B-2).

Since attempts at crystallizing the isolated, transformed enantiomers for X-ray analysis failed, their structures were identified by acquiring and comparing their experimental and theoretical CD spectra (Fig. 4C-1 and E-2). The theoretical CD spectra of the enantiomers of the three geometric isomers differed from one another. The shapes of the experimental spectra in Fig. 4A-1 and A-2 appeared most similar to those of the theoretical spectra of the enantiomers of the *mer-cis* isomer in Fig. 4C-1 and C-2: both the theoretical spectrum of the Λ -enantiomer (in Fig. 4C-1) and the experimental spectrum (in Fig. 4A-2) showed three positive and three negative Cotton bands in the range 200–400 nm and such a feature was not seen for the enantiomers of *fac-cis* isomer and *mer-trans-C-C-cis-N-N* isomer (Fig. 4D-1 and E-1). We hence propose that the Δ -enantiomer of the *mer-cis-C-C-trans-N-N* isomer mutated into the Λ -enantiomer of the *mer-cis* isomer.

Note that the intensities of all of the signals in the HPLC profiles in Fig. 3C and D decreased as the irradiation time was increased, and, after 180 min of irradiation, the overall intensities became much lower than those in the initial state (Fig. S7 in ESI[†]). The decreases in the CD intensities in Fig. 3A and B were thus not due only to the generation of the optically active transformed products but also to disappearance or decomposition of the enantiomers and their corresponding transformed derivatives.

In addition, changes in the CD/UV spectra and in the HPLC profiles of the Δ -enantiomer occurred more rapidly upon irradiation using NPL with a relatively high irradiation power (of *ca.* 200 mW cm⁻²) than when using LPL (*ca.* 40 mW cm⁻²) under an N₂ atmosphere. Moreover, upon irradiation of the Δ -enantiomer using NPL, the changes in its CD and HPLC results were faster in the presence of air than under N₂ (Fig. S7 in ESI[†]). These results suggested that the stereomutation and decomposition of the Λ - and Δ -enantiomers occurred in excited states, and were accelerated by O₂ contained in air, possibly through interactions between unpaired electrons of O₂ with electrons of Flrpic in its higher-energy orbitals. The photoreactions of Flrpic are assumed to have occurred through its triplet excited states. A photo-induced transformation of the Ir complex has been reported for iridium(III) tris(2-phenylpyridinato-N,C^{2'}) (Ir(ppy)₃)¹⁸ but has never been reported for Flrpic. The current work is the first to report on the photo-isomerization of enantiomers of Flrpic.

In conclusion, commercial Flrpic, *i.e.*, the *mer*-isomer of Flrpic with *cis* C–C and *trans* N–N arrangements, was resolved into optically pure Λ - and Δ -enantiomers by using chiral HPLC, and upon photo-irradiation, the enantiomers underwent stereomutation and decomposition: we propose that the Λ - and Δ -enantiomers of the *mer-cis-C-C-trans-N-N* isomer mutated into the Δ - and Λ -enantiomers of the *mer-cis* geometrical isomer, respectively. The mutation and decomposition occurred not only in the presence of air but also under an N₂ atmosphere. These results revealed an important cautionary note, that



optically active FIRpic may be transformed in excited states even in a sealed device, which would need to be taken into consideration when designing LEDs using optically active FIRpic enantiomers. In addition, transformation of FIRpic may also play a role in catalyses using Ir complexes.¹¹

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