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## Investigating the isolation and interconversion of two diastereoisomers in an octahedral $\text{Ni}^{\text{II}}\text{A}_2(\widehat{\text{BC}})_2$ system†

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Two diastereoisomers of bis(2-benzoylpyridine-*O,N*)dichloridonickel(II) have been isolated from the reactions of nickel(II) chloride hexahydrate and the organic ligand depending on the reaction solvent (ethanol, acetonitrile) used. DFT studies indicate that one of the isomers is more stable in both solvents, but nevertheless different diastereoisomers are crystallized from the two solvents. The two complexes represent the first pair of well characterized diastereoisomers in the coordination chemistry of 2-pyridyl ketones, providing a new example in the chemistry of octahedral  $\text{MA}_2(\widehat{\text{BC}})_2$  complexes (A = a monodentate ligand,  $\widehat{\text{BC}}$  = an unsymmetrical bidentate chelate).

Diastereoisomeric and enantiomeric metal complexes (especially those of the d-block elements) have played a central role in the development of modern inorganic chemistry.<sup>1–5</sup> An isomer is one of several chemical species that have the same atom composition (molecular formula), but have different constitutional formulae (atom connectivities) or different stereochemical formulae.<sup>1</sup> Isomers exhibit different physical and/or chemical properties. If two complex molecules or ions have the same molecular formula and the same atom connectivity, but differ in the spatial arrangement of different atoms or groups about a central metal ion, then the coordination compounds are stereoisomers.<sup>1</sup> Diastereoisomers (which are the focus of the present work) are stereoisomers that are not mirror-images of one another, whereas enantiomers are stereoisomers that are mirror-images of one another. A famous example is the diastereoisomers in the pair of the square planar

complexes  $[\text{PtCl}_2(\text{NH}_3)_2]$ ; the *cis* isomer is a well-known anti-tumor drug, but the *trans* isomer is pharmacologically inactive.<sup>6</sup> Diastereoisomerism continues to attract the intense interest of synthetic and physical inorganic chemists. For instance, recent examples include *cis-trans* interconversion in octahedral bis(2,2'-bipyridine) Ru(II) complexes;<sup>7</sup> this results in specific synthetic applications of the *trans* species, potentially useful as a platform for designing highly efficient visible light-activated caged compounds. Single-crystal X-ray crystallography is the state-of-the-art method for identifying diastereoisomers in the solid state, while advanced theoretical methods can explain the preference of a given system for one diastereoisomer and predict/explain interconversions.

The introduction of chelating ligands in six-coordinate octahedral complexes usually acts to limit the number of diastereoisomers (or geometrical isomers), or at least not to extend the number.<sup>2</sup> For example, the coordination of three symmetrical bidentate chelates ( $\widehat{\text{BB}}$ ) can take place in only one way, in the same sense that six identical monodentate ligands can coordinate in only one way. Likewise,  $\text{MA}_2(\widehat{\text{BB}})_2$  compounds (where A refers to a monodentate group) have precisely the same number and type of diastereoisomers as the all-monodentate analogue  $\text{MA}_2\text{X}_4$ . The same holds true for the  $\text{MA}_2(\widehat{\text{BC}})_2$  and the all-monodentate  $\text{MA}_2\text{X}_2\text{Y}_2$  complexes, where  $\widehat{\text{BC}}$  refers to an unsymmetrical bidentate chelate (X and Y are monodentate ligands bearing the same donor atoms that are present in the B and C parts, respectively, of  $\widehat{\text{BC}}$ ); for these compounds there are now five diastereoisomers (Scheme 1). Contrary to  $\text{MA}_2(\widehat{\text{BB}})_2$  diastereoisomers,<sup>1–5</sup> studies in  $\text{MA}_2(\widehat{\text{BC}})_2$  diastereoisomers are generally lacking. As a continuation of our interest in the chemistry of diastereoisomers possessing asymmetrical bidentate chelating ligands,<sup>8</sup> we report here experimental (synthetic, structural, spectroscopic) and theoretical studies on two nickel(II) diastereoisomers that belong to the  $\text{MA}_2(\widehat{\text{BC}})_2$  class. We selected 2-benzoylpyridine (or phenyl 2-pyridyl ketone), (py)(ph)CO, for

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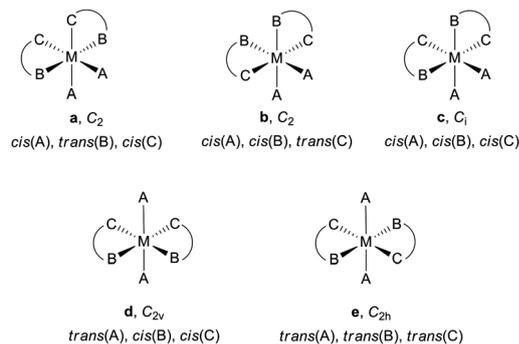
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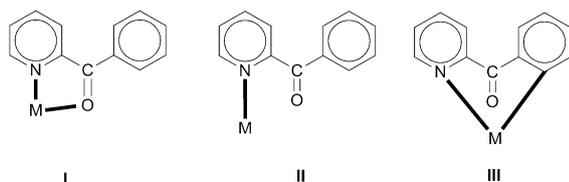
E-mail: attsipis@uoi.gr; Tel: +30 26510 08333

† Electronic supplementary information (ESI) available: Full synthetic and characterization (spectroscopic, theoretical and structural) details for the two complexes (text, figures, tables). Cifs and checkcifs for the two compounds. CCDC 1949392 and 1949393. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c9nj04876f





**Scheme 1** Diastereoisomers (with indication of point symmetry) possible for octahedral complexes  $MA_2(\overline{BC})_2$  (in the present work  $M = Ni$ ,  $A = Cl$  and  $(\overline{BC}) = 2$ -benzoylpyridine). B (O in the present case) and C (N in the present case) denote the donor atoms of the unsymmetrical (heterotopic) bidentate chelating ligand 2-benzoylpyridine; **b** corresponds to complex **2** of this work and **e** corresponds to **1**·0.4H<sub>2</sub>O·0.6EtOH.

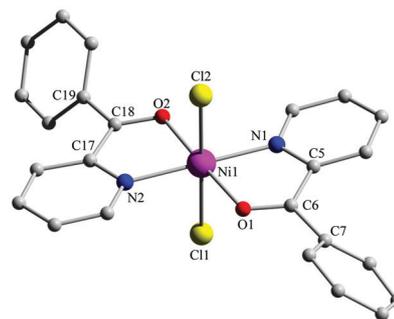


**Scheme 2** The to-date crystallographically established coordination modes of (py)(ph)CO. The ligand is neutral in I (the most common ligation mode) and II, and anionic organometallic in III.

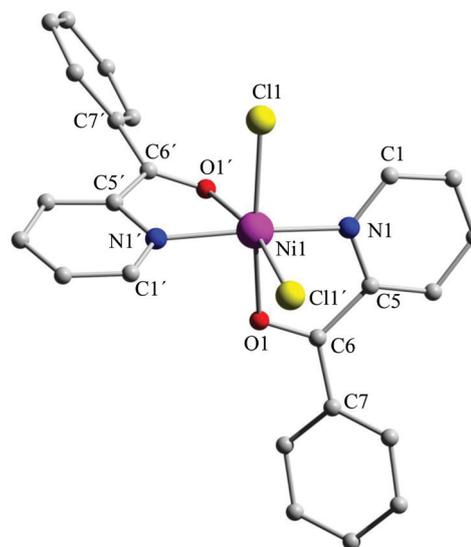
two reasons. First, this molecule normally behaves as a neutral, *O,N*-bidentate chelating ligand,<sup>9,10</sup> (exceptions are known,<sup>11–14</sup> Scheme 2) and, second, it can undergo impressive metal ion-assisted transformations,<sup>15,16</sup> a topic that is currently one of the central themes in our research efforts.<sup>17–19</sup>

Treatment of NiCl<sub>2</sub>·6H<sub>2</sub>O with (py)(ph)CO (1:2) in EtOH gave an olive green solution from which were subsequently isolated dark orange-red crystals of *trans,trans,trans*-[NiCl<sub>2</sub>{(py)(ph)CO}<sub>2</sub>]<sub>2</sub>·0.4H<sub>2</sub>O·0.6EtOH (1·0.4H<sub>2</sub>O·0.6EtOH) in moderate yields (50–55%) upon vapour diffusion with Et<sub>2</sub>O. The 1:2 reaction between NiCl<sub>2</sub>·6H<sub>2</sub>O and (py)(ph)CO (1:2) in MeCN gave a green solution from which were subsequently isolated green crystals of *cis,cis,trans*-[NiCl<sub>2</sub>{(py)(ph)CO}<sub>2</sub>]<sub>2</sub> (**2**) in rather high yields (~70%) upon liquid diffusion with Et<sub>2</sub>O. The first prefix in the formulae refers to the chlorido groups, the second to the O and the third refers to the N donor atoms (*vide infra*). Complex **1** is readily transformed into **2** upon dissolution in MeCN and either slow solvent evaporation or precipitation with Et<sub>2</sub>O. Solid **2** reverts to **1** in EtOH under heating and precipitation with Et<sub>2</sub>O. The IR  $\nu(C=O)$  mode appears at ~1625 cm<sup>-1</sup> in the spectra of the complexes (which are similar but not identical) and is shifted to lower wavenumbers by ~45 cm<sup>-1</sup> (compared with the free ligand) due to coordination.<sup>20</sup>

The mononuclear octahedral complexes **1** and **2** (Fig. 1, 2 and Fig. S4, S8a, ESI<sup>†</sup>) are diastereoisomers. In the former, all the chemically similar donor atoms are in *trans* positions. The latter possesses a crystallographic 2-fold axis which is parallel



**Fig. 1** The structure of the molecule *trans,trans,trans*-[NiCl<sub>2</sub>{(py)(ph)CO}<sub>2</sub>]<sub>2</sub> that is present in the crystal of **1**·0.4H<sub>2</sub>O·0.6EtOH. Bond lengths (Å) in the coordination sphere of Ni<sup>II</sup>: Ni1–Cl1 2.417(1), Ni1–Cl2 2.397(1), Ni1–O1 2.095(2), Ni1–O2 2.088(2), Ni1–N1 2.046(2), Ni–N2 2.049(3).



**Fig. 2** The structure of the molecule *cis,cis,trans*-[NiCl<sub>2</sub>{(py)(ph)CO}<sub>2</sub>]<sub>2</sub> that is present in the crystal of **2**. Bond lengths (Å) in the coordination sphere of Ni<sup>II</sup>: Ni1–Cl1/Cl1' 2.376(1), Ni1–O1/O1' 2.160(1), Ni1–N1/N1' 2.054(2). The N1–Ni1–N1' angle is 169.9(1)°. Symmetry operation: (')  $x, 1 - y, 3/2 - z$ .

to the *a* axis and passes through Ni1 bisecting the Cl1–Ni1–Cl1' angle; the molecule is the *cis,cis,trans* isomer considering the positions of the chlorido ligands, the carbonyl oxygen atoms and the 2-pyridyl nitrogen atoms. In both complexes, the (py)(ph)CO molecules behave as *O,N*-bidentate chelating ligands (I in Scheme 2). The distortion from the regular octahedral geometry in the two complexes is primarily attributed to the somewhat small bite angles of the two 5-membered chelating rings (the N–Ni–O angles for a given ligand are in the range 77.2(1)–78.9(1)°). There is one crystallographically independent (and its symmetry equivalent) intramolecular H bond in **2** with the aromatic carbon atom C1 as donor and the chlorido group as acceptor (Fig. S8a and Table S5, ESI<sup>†</sup>). Intermolecular H-bonding interactions (1·0.4H<sub>2</sub>O·0.6EtOH), and both intermolecular H bonds and  $\pi$ - $\pi$  stacking interactions (**2**) are responsible for the formation of interesting 3D architectures in the two structures (see the ESI<sup>†</sup> for full details).





solution turned to olive green. The reaction solution was stirred for a further 15 min and filtered. Vapour diffusion of Et<sub>2</sub>O into the reaction solution gave X-ray quality, dark orange-red crystals of the product after 21 d. The crystals were collected by filtration, washed with ice-cold EtOH (2 × 0.5 mL) and Et<sub>2</sub>O (3 × 2 mL), and dried *in vacuo* over P<sub>4</sub>O<sub>10</sub> overnight. Typical yields were in the range 50–55%. The product was analyzed satisfactorily as lattice solvent-free, *i.e.* **1**. Anal. calcd for C<sub>24</sub>H<sub>18</sub>N<sub>2</sub>O<sub>2</sub>Cl<sub>2</sub>Ni: C, 58.11; H, 3.67; N, 5.65%. Found: C, 57.92; H, 3.73; N, 5.33%. Diagnostic IR (KBr, cm<sup>-1</sup>) band: 1624 (s). Solid-state electronic spectrum (λ/nm): 405, 696, 810 (sh), 1080.

### Preparation of *cis,cis,trans*-[NiCl<sub>2</sub>{(py)(ph)CO}<sub>2</sub>] (**2**)

To a pale green solution of NiCl<sub>2</sub>·6H<sub>2</sub>O (0.059 g, 0.25 mmol) in MeCN (15 mL) was added solid (py)(ph)CO (0.092 g, 0.50 mmol). The solid was soon dissolved, and the resultant green solution was stirred for a further 15 min. The reaction solution was layered with Et<sub>2</sub>O (20 mL). Slow mixing gave X-ray quality, green crystals of the product over a period of 10 d. The crystals were collected by filtration, washed with Et<sub>2</sub>O (4 × 2 mL), and dried in air. The yield was 67%. Anal. calcd for C<sub>24</sub>H<sub>18</sub>N<sub>2</sub>O<sub>2</sub>Cl<sub>2</sub>Ni: C, 58.11, H, 3.67, N, 5.65%. Found: C, 58.27; H, 3.80; N, 5.47%. Diagnostic IR (KBr, cm<sup>-1</sup>) band: 1622 (s). Solid-state electronic spectrum (λ/nm): 397, 425, 710 (sh), 1010.

XRD data for 1·0.4H<sub>2</sub>O·0.6EtOH: monoclinic, *P*<sub>2</sub><sub>1</sub>/*n*, *a* = 10.5379(2) Å, *b* = 9.1919(2) Å, *c* = 25.3211(5) Å, β = 95.362(1)°, *V* = 2441.95(9) Å<sup>3</sup>, *Z* = 4, *R*<sub>1</sub>[*I* > 2σ(*I*)] = 0.0449, *wR*<sub>2</sub> (all data) = 0.1441. XRD data for **2**: orthorhombic, *P**ca**n*, *a* = 7.9540(1) Å, *b* = 13.9375(3) Å, *c* = 16.6479(4) Å, *V* = 2178.14(7) Å<sup>3</sup>, *Z* = 4, *R*<sub>1</sub>[*I* > 2σ(*I*)] = 0.0306, *wR*<sub>2</sub> (all data) = 0.0819.

### Conversion of **1** into **2**

A green solution of **1** (0.099 g, 0.20 mmol) in MeCN (20 mL) was stirred overnight. The solution was layered with Et<sub>2</sub>O (25 mL). Slow mixing gave X-ray quality, green crystals in a period of one week; the unit cell of a crystal was almost identical with that of the authentic **2** prepared as described above. The crystals were collected by filtration, washed with Et<sub>2</sub>O (4 × 2 mL), and dried in air. The yield was ~55%. Analytical data calcd for C<sub>24</sub>H<sub>18</sub>NiN<sub>2</sub>O<sub>2</sub>Cl<sub>2</sub> (found values are in parentheses): C 58.11 (58.43), H 3.67 (3.51), N 5.65 (5.70)%. The IR and diffuse reflectance spectra of this sample are superimposable with the corresponding spectra of the authentic complex **2**. Effective magnetic moment (μ<sub>eff</sub>) at 22 °C: 3.36 BM. The complex can be isolated (albeit in a rather poor crystal quality and a lower yield of ~20%) upon slow solvent evaporation of the green solution at room temperature over a period of 1 week.

All DFT calculations were performed using the Gaussian program suite employing the 1999 hybrid functional of Perdew, Burke and Ernzerhof (denoted as PBE0). For Ni, the relativistic effective core potentials (RECP) were used in conjunction with the SDD basis set to describe the metal valence electrons, and two sets of *f* polarization functions were added; the 6-31G(d,p) basis set was used for all other main group elements, E. The method used is abbreviated as PBE0/SDD(Ni) ∪ 6-31G(d,p)(E).

## Conflicts of interest

There are no conflicts to declare.

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