Bite-Angle-Regulated Coordination Geometries: Tetrahedral and Trigonal Bipyramidal in Ni(II) with Biphenyl-Appended (2-Pyridyl)alkylamine N,N′-Bidentate Ligands

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Supporting Information

ABSTRACT: Two simple biphenyl-appended (2-pyridyl)-alkylamine N-bidentate ligands, L⁶ and L⁸, having ethylene and methylene spacers between donor groups, with bite angles L⁶ ≈ 100° and L⁸ ≈ 80°, dictate pseudotetrahedral and trigonal-bipyramidal geometries in six high-spin Ni(II)-halide complexes, [Ni(L⁶)X₂] and [Ni(L⁸)X]ClO₄ (where X = Cl⁻, Br⁻, I⁻), respectively. The structures in the solid state, determined using X-ray crystallography, and in solution, determined using spectroscopic methods (UV–vis–NIR and paramagnetic ¹H NMR), which complement each other, are described.

INTRODUCTION

There is growing interest directed toward the modification of classical N-bidentate ligands, α-dimine (2,2′-bipyridine, 1,10-phenanthroline), β-diketiminate, diamine, and (2-pyrididy)-alkylamines to tune the steric and electronic properties to achieve novel and reactive transition metal complexes.¹ Low-coordinate (≤4) mononuclear nickel compounds with such N-bidentate ligands have attracted a great deal of recent interest in the field of bioinorganic chemistry, including activation of small molecules (O₂, N₂, N₂O, NO), oxygenation catalysis, olefin polymerization, electro- and photo-catalytic hydrogen production, and other areas of applications.²⁻⁴

Four-coordinate tetrahedral Ni(II)-halide complexes of sterically hindered pyridyl-imine ligands serve as alternative catalysts for traditional Ziegler–Natta polymerization of ethylene and α-olefins.⁵ Similarly, the tetrahedral Ni(II)-halide complex [Ni(bc)Cl₂] (bc = 2,9-dimethyl-4,7-diphenyl-1,10-phenanthroline) functions as a photocatalyst for production of hydrogen in aqueous medium.⁶ The classical five-coordinate Ni(II) complex, [Ni(Me₆tren)Cl]ClO₄, with magnetic anisotropy has drawn recent attention as a potential single-molecule magnet.⁷

Strangely, there are a limited number of four- and five-coordinate high-spin mononuclear Ni(II)-halide complexes with unhindered N-bidentate ligands, exhibiting tetrahedral (td) and trigonal-bipyramidal (tbp) geometries.⁸ This is due to the plasticity of Ni(II) coordination for a range of geometries: tetrahedral, square-planar, square-pyramidal, trigonal-bipyramidal, and octahedral, which has been elegantly demonstrated recently using a chelating hemilabile P,S-bidentate ligand.⁹ On the other hand, the use of soft donor bidentate diphosphine or hybrid P,N-ligands with methylene and ethylene spacers give low-spin square-planar Ni(II)-dichloride complexes.¹⁰ Even the N-bidentate dihydrobis(pyrazolyl)borates with bulky substituents on the ring give square-planar and occasionally octahedral geometries.¹¹ The most common N-bidentate ligands, like bpy, phen, and diamine (L), typically generate tris-chelated [NiL₃]⁺, hexacoordinate high-spin complexes,¹² and in solution, a dynamic equilibrium of species with different metal to ligand ratios exist.¹³ Even well-established versatile tridentate ligands, like triazacyclononane (tacn), hydrotris(pyrazolyl)borate anion (Tp⁻), and tetradeinate tris(2-pyridylmethyl)amine (tmpa), likewise give an octahedral geometry.¹⁴ A few tripodal tetraamine (tren) and tris(2-amino phenyl)amine derivatives with alkyl, aryl, amide, and urea substituents stabilize the Ni(II)-halides in the tbp geometry.¹⁵⁻¹⁶ On the other hand, bis(imino)-pyridine and terpyridine ligands mostly give square-pyramidal geometry.¹⁵ The modified bpy and phen ligands with bulky aryl/alkyl groups adjacent to N-donor atoms or alkylated diamines stabilize Ni(II)-halide in the tbp geometry,¹⁷⁻¹⁸ and less sterically hindered bidentate ligands often give halide-bridged binuclear Ni(II) complexes. However, the syntheses of such ligands with bulky substituents are sometimes challenging, either requiring expensive reagents or involving multiple steps. In addition, it has been shown that the bite angle of such rigid (bpy, phen) ligands remains unchanged (~81°) and the stabilization of tetrahedral geometry is due to steric protection.¹⁹

It is clear from the above survey that the structure and coordination geometries of Ni(II)-halide complexes are influenced by the nature of the ligand, that is, donor atom type, chelate ring size, and the steric bulk. However, control of their structures and coordination geometries by altering the bite
angles of bidentate ligands has not been recognized before. The vital roles of bite angles in regulating several physical and chemical properties of transition metal complexes have been documented. The influence of wide bite angle (120°) bidentate diphosphine ligands in hydroformylation of olefins using rhodium catalyst to give stereoselective linear aldehydes has been well established, wherein they stabilize reaction intermediates in trigonal bipyramidal geometry. Similarly, subtle changes in the bite angles of \( N \)-bidentate ligands influencing the spin-crossover (high-spin ↔ low-spin) transition of tris-chelated-iron complexes have been explained.

Interestingly, the use of simple \( N \)-substituted (2-pyridyl)-alkylamine ligands with different alkyl spacers to stabilize different coordinate geometries of nickel(II) complexes is scarce, even though they can offer a handle to alter the bite angles. We have employed two such simple biphenyl-appended (2-pyridyl)alkylamine \( N,N' \)-bidentate ligands having either ethyl or methyl alkyl spacers and shown how they regulate two different coordination geometries of Ni(II). The new ligands are easy to synthesize in one step and in high yields using readily available (2-pyridyl)alkylamines and 2,2'-bis-(bromomethyl)-1,1'-biphenyl. We have previously shown using one of the ligands (L\(^{m} \), Scheme 1) with copper compounds to give unusual structures and reactivity toward dioxygen and phosphodiester substrate.

### Scheme 1. Synthetic Route to Tetra- and Penta-Coordinate \( N \)(II)-Halide Complexes Dictated by the \( N \)-Bidentate Ligand Bite Angles

![Scheme 1. Synthetic Route to Tetra- and Penta-Coordinate \( N \)(II)-Halide Complexes Dictated by the \( N \)-Bidentate Ligand Bite Angles](image)

Herein, we describe the syntheses and characterization of six mononuclear high-spin Ni(II)-halide complexes, \([\text{Ni}(L)^{t}X_{2}]\) and \([\text{Ni}(L)^{m}X]^{+}\), in tetrahedral and trigonal-bipyraridal geometries, regulated by the bite angles of new \( N,N' \)-bidentate ligands, \( L' \) and \( L'' \), having ethylene (\( CH_{2}CH_{2} \)) or methylene (\( CH_{2} \)) spacers between the two donor groups (Scheme 1). The work presented here represents a rare example of controlling the coordination geometries of Ni(II) by merely altering the chelate bite angles of simple \( N \)-bidentate ligands. The structural characterizations of all of the high-spin Ni(II) complexes in the solid state by single crystal X-ray diffraction and in solution by spectroscopic (UV–vis–NIR, ESI-MS, and paramagnetic \( 'H \) NMR) methods are described.

### RESULTS AND DISCUSSION

Treatment of the \( N,N' \)-bidentate ligand, \( L' \), with anhydrous nickel(II)-halide salts, \( \text{NiX}_{2} \) (\( X = \text{Cl}^{+}, \text{Br}^{+}, \text{I}^{-} \)) in equimolar ratios in toluene at 50 °C for 2 h provided blush-violent to brownish-green crystalline solids, \([\text{Ni}(L)^{t}X_{2}]\) (where \( X = \text{Cl}^{-}, \text{Br}^{-}, \text{I}^{-} \), \( 1a-c \)) in good yield (70%). They were also prepared in tetrahydrofuran, \( CH_{3}CN \), or methanol using hydrated Ni-halide salts or Ni-perchlorate in the presence of a halide source, like \( Bu_{n}NX \) or \( NaX \) (\( X = \text{Cl}, \text{Br}, \text{I} \)). However, the yields were relatively lower because of their partial solubility in these solvents. These complexes were stable to air in the solid state and solution and soluble in chlorinated polar solvents. Efforts to prepare 1:2 Ni/L\(^{t} \) complexes were unsuccessful, in spite of using more than 2 equiv of ligand, clearly demonstrating the preference of ligand, L\(^{t} \), for four-coordinate complexes. Contrastingly, the reaction of L\(^{m} \) with hydrated Ni(II)-halide salts \( \text{NiX}_{2}\cdot xH_{2}O \) (\( X = \text{Cl}^{-}, \text{Br}^{-}, \text{I}^{-} \) in 1:2 molar ratio in methanol at room temperature, followed by the metathesis reaction with NaClO\(_{4} \) readily precipitated highly crystalline apple-green to yellow-green solids, \([\text{Ni}(L)^{m}X]^{+}\) (\( X = \text{Cl}^{-}, \text{Br}^{-}, \text{I}^{-} \), \( 2a-c \)) in good yield (70–75%). Like before, an attempt to prepare the 1:1 4-coordinate complex, \([\text{Ni}(L)^{m}Cl]^{+}\) was unsuccessful, which emphasizes the preference of L\(^{m} \) to stabilize five-coordinate complexes.

The molecular structures of all of the six Ni(II) complexes were determined by single crystal X-ray diffraction, and as a representative example, the results of \( 1a \) and \( 2a \) are shown in Figure 1. The comparable structures of \( 1b-c \) and \( 2b-c \) are shown in Figure S1, with selected bond lengths and angles listed in Table 1. The molecules \( 1a-c \) are neutral four-coordinate mononuclear Ni(II) with pseudo-tetrahedral geometry, and the metal is chelated by the \( N \)-bidentate ligand, \( L' \), and two terminal halide ions. The distortion in the geometry is based on the \( \tau_{4} \) criterion, 0.87 for \( 1a-b \) and 0.97 for \( 1c \).

Interestingly, the opening of the X1–Ni–X2 angles [130.74(2), 129.00(2), 115.53(2)°] compared to that in a regular tetrahedron is accompanied by closing of the ligand N–Ni–N bite angles [100.14(6), 100.73(8), 97.04(12)] in \( 1a-c \), respectively. This is not unusual as a similar pattern has been observed with related complexes containing bis-guanidine and other hindered bidentate ligands. It is important to point out that the ligand (\( L' \) \( N \)-Ni–\( N \) bite angle, \( \approx 100^\circ \) (av), is much wider than that observed (91°) for complexes containing propane-diamine, which also form a six-membered ring, chelate, like in \( L' \). The wide bite angle (>90°) bidentate diamines or (2-pyridyl)alkylamines are rare. The widening of the angle in the present case is due to the combined effect of both the ethyl spacer and sterically hindered biphenyl group. A similar example is apparent from careful analysis of a recent report on the Ni(II)-bromide complex of the \( N \)-substituted 2-
Table 1. Selected Bond Lengths (Å) and Bond Angles (deg) of [Ni(L\textsuperscript{a})Cl\textsubscript{2}], 1a; [Ni(L\textsuperscript{a})Br\textsubscript{2}], 1b; [Ni(L\textsuperscript{a})I\textsubscript{2}], 1c; [Ni(L\textsuperscript{m})\textsubscript{2}]Cl[(ClO\textsubscript{4})\textsubscript{2}]:CH\textsubscript{2}Cl\textsubscript{2}, 2a; [Ni(L\textsuperscript{m})\textsubscript{2}]Br[(ClO\textsubscript{4})\textsubscript{2}]:CH\textsubscript{2}Cl\textsubscript{2}, 2b; and [Ni(L\textsuperscript{m})\textsubscript{2}]I[(ClO\textsubscript{4})\textsubscript{2}], 2c Complexes

| X          | Cl (1a)   | Br (1b)   | I (1c)  |
|------------|-----------|-----------|---------|
| bond lengths |           |           |         |
| Ni1−N1     | 2.0463(14) | 2.049(2)  | 2.025(3) |
| Ni1−N2     | 1.9940(15) | 1.995(2)  | 2.001(3) |
| Ni1−N3     | 2.2261(6)  | 2.3582(5) | 2.5327(6) |
| Ni1−N4     | 2.503(6)   | 2.3791(5) | 2.5372(6) |
| bond angles |           |           |         |
| N1−N1−N2   | 100.14(6)  | 100.73(8) | 97.04(12) |
| N1−N1−X1   | 110.69(4)  | 111.17(6) | 115.06(9) |
| N1−N1−X2   | 107.00(4)  | 108.31(6) | 111.92(9) |
| N2−N1−X1   | 100.57(5)  | 100.35(6) | 110.94(9) |
| N2−N1−X2   | 102.80(5)  | 102.67(7) | 104.17(9) |
| X2−Ni−X1   | 130.74(2)  | 129.00(2) | 115.53(2) |
| bite angle  |           |           |         |
| N1−N1−N2   | 100.14(6)  | 100.73(8) | 97.04(12) |
| dihedral angle (ϕ) | |         |         |
| N1−N1−X2/N1−N1−X2 | 88.11      | 88.29    | 88.09   |
| N1−N1−X2/N1−N1−X1 | 87.77      | 86.17    | 86.63   |
| N1−N1−X1/N1−X2 | 86.15      | 87.67    | 84.73   |
| geometry index (r\textsubscript{g}) | 0.84 | 0.85 | 0.92 |

(2-pyridyl)ethylamine bidentate ligand having a hindered diethyl group, which offers a wide bite angle (97°) and stabilizes the four-coordinate tetrahedral geometry.\textsuperscript{25} The Ni−N\textsubscript{av} bond is longer than Ni−N\textsubscript{vis} as expected; this is due to the bulky biphenyl group on the amine nitrogen, which makes N\textsubscript{av} a weaker σ-donor. The coordination geometry and bond parameters of Ni(II) in 1a–c are comparable to those of related structures.\textsuperscript{6,16b,23} Furthermore, the molecules in the crystal lattice are stabilized by several strong noncovalent intermolecular interactions (Figure S2, Table S2), such as hydrogen bonding (C−H···Cl, 2.85 Å, 150.8°), π−π stacking (3.2 Å, py−py, face-to-face, interplanar; 3.17 Å, py−py, face-to-side, tilted 24°), and C−H−π (2.74 Å, CH\textsubscript{2}−π, 141°) giving them extra stability.

On the other hand, the five-coordinate Ni(II) complexes, 2a–c, adopt a distorted trigonal-bipyramidal geometry (Figure 1, 2a), ligated by two ligand molecules, L\textsubscript{m}, providing tetra-N coordination plus a chloride (halide) ion completes the pentacoordination. The equatorial plane is made of two tertiary amine-nitrogen atoms having a bulky biphenyl append and a chloride (halide) ion. The axial positions are occupied by the nitrogen atoms of two pyridyl rings, which are nearly orthogonal, with dihedral angles (ϕ) varying between 75.1 and 77.4° for 2a-c (Table 1). The metal atom sits almost in the trigonal plane of N\textsubscript{av}N\textsubscript{vis}Cl\textsubscript{av} atoms, and the sum of the trigonal angles is 360°. However, the geometry about the metal is distorted because of the acute ligand N−Ni−N bite angles (~80° av) and hence the axial and equatorial angles deviate from the ideal, 180 and 120°, by 9°. According to the r\textsubscript{g} (0.76) criterion,\textsuperscript{26} the pentagonal coordination for Ni(II) is identified as distorted tbp. The geometry and coordination bond parameters of 2a–c are comparable to those of other related Ni(II) complexes containing tripodal tetra-amine-derivatives.\textsuperscript{24} A similar example of 5-coordinate Ni(II) with tbp geometry has been recently reported with the N-substituted (2-pyridyl)methylamine ligand having a bulky (2,6-dimethyl)-phenyl group.\textsuperscript{24} It re-emphasizes the role of biphenyl append in the stabilization of such an uncommon tbp geometry with Ni(II). Furthermore, strong intermolecular hydrogen bonding and π−π stacking interactions, like those in 1a–c, are also observed (Figure S3).

The structures of 1a–c and 2a–c in solution, characterized by spectroscopic (UV–vis−NIR, 1H NMR, ESI-MS) methods and magnetic susceptibility measurements in solution (CD\textsubscript{2}Cl\textsubscript{2})
or CDCl₃) by Evans NMR method,²⁷ agree well with the structures observed in the solid state. The moments of 1a−c in solution with \( \mu_{\text{eff}} \approx 3.15 \) B.M. are consistent with those of Ni(II) \((S = 1)\) in tetrahedral geometry. Also, the moment of solid 1c (Figure S4), with a \( \chi_M T \) value of 1.1 emu mol\(^{-1}\) K \((\mu_{\text{eff}} = 3.0 \) B.M.) at 298 K, agrees with that measured in solution. The moment values, \( \mu_{\text{eff}} \approx 3.11, 3.47, \) and 3.28 B.M. for 2a−c, respectively, are slightly greater than the spin-only value due to the spin−orbital coupling contribution. The ESI-MS spectra of complexes 2a−c show molecular-ion peaks at \( m/z \), 667, 711, and 757, respectively, corresponding to \([\text{M}−\text{ClO}_4]^+\), and the observed and calculated isotopic patterns are consistent with the expected composition (Figure S5).

The electronic absorption spectra of Ni(II) complexes in solution are characteristic of their distorted-tetrahedral and tbp geometries. The spectra of 1a−c and 2a−c were measured in CH₂Cl₂, and as a representative example, the spectra of \([\text{Ni}(\text{Le})\text{Br}_2]\), 1b, and \([\text{Ni}(\text{Lm})_2\text{Br}](\text{ClO}_4)\), 2b are shown in Figure 2, and the comparable spectra of the remaining complexes are shown in Figure S6. The spectrum of 1b exhibits five bands in the visible−NIR region. Both the profile and features are typical of Ni(II) in a pseudotetrahedral geometry and comparable to those of \([\text{Ni}(\text{TpMe}_2)\text{(Br)}]\) (where TpMe² = hydrotris(3,5-dimethylpyrazole)borate) having a similar geometry.²⁸ The spectrum of 1c with two iodides is slightly different in the visible region due to a slight deviation in the coordination geometry. The spectra of complexes with tbp geometry 2b (Figure 2, 2b) and 2a,c (Figure S7) show three well-defined bands and a shoulder at 470 nm; besides, charge transfer bands are also observed for 2b−c in the UV-region. These features are characteristic of Ni(II)-halide complexes in the tbp geometry and compare well with other complexes containing tripod tetra-amine ligands.²⁹ Furthermore, the relative shift in the absorption wavelength position among 1a−c and 2a−c is in accordance with the spectrochemical series Cl < Br < I.

The geometries of Ni(II) complexes in solution were easily identified by proton NMR spectroscopy.³⁰ Their paramagnetic spectra differ significantly from the corresponding diamagnetic molecules. The spectra of six-coordinate Ni(II) compounds are normally broader, whereas those for tetrahedral and trigonal-bipyramidal complexes are sharper and shifted over a wide spectral window.¹³c The \(^1\)H NMR spectra of all six complexes, 1a−c and 2a−c, were either measured in CD₂Cl₂ or CDCl₃. As a representative example, the spectra of 1b and 2b are shown in Figure 3, and the rest are shown in Figures S8 and S9. The spectrum of 1b possesses seven hyperfine-shifted resonances due to the ligand (L¹) protons: one upfield shifted (−26.6 ppm), five downfield, and a broad signal in the diamagnetic region. We assume that the isotropic shifts in all of the resonances are dominated by Fermi-contact (\( \sigma \)-delocalization), and the pseudocontact contribution is negligible. Signals were assigned on the basis of chemical shifts, integral ratio, peak widths \((\nu_{1/2})\), proximity of protons to the metal center (Ni···H distances from the X-ray structure of 1b), and comparison with related examples from the literature.³² The resonance at −26.6 ppm is assigned to the methylene pair bound to the pyridine ring (H2), as it is common to observe this pair in the upfield region.³³ The other methylene pair (H1) appears at 194 ppm.
and the two methylene pairs attached to the biphenyl ring (H7) may also have overlapped with it. The remaining sharp signals at 228, 61, 53, and 31 ppm are typical of pyridine ring protons and assigned to H6, H5, H3, and H4, respectively. The biphenyl protons (H8) appear in the diamagnetic region, as they are too far away from the paramagnetic Ni(II) center. The spectra for the chloro and iodo derivatives, 1a and 1c, show a slight variation, but otherwise, the pattern remains the same. The spectral features of these complexes are comparable to those of other related tetrahedral Ni(II) complexes containing pyridine-type ligands.32 On the other hand, the spectrum of 1b showed signals in the upfield region. The few signals in the upfield region indicate a smaller magnetic anisotropy due to the small bite angle of two chelated ligand molecules (Lm).31b The two pyridine rings are equivalent, whereas the methylene groups are not.

**CONCLUSIONS**

In summary, we have reported the syntheses of six new high-spin Ni(II)-halide complexes in tetrahedral and trigonal-bipyramidal geometries, regulated by the bite angles of two simple biphenyl-appended (2-pyridyl)alkylamine N,N'-bidentate ligands having ethylene and methylene spacers. All of the complexes, 1a–c and 2a–c, were characterized by single-crystal X-ray diffraction. The electronic absorption and paramagnetic proton NMR spectra in solution complemented the solid-state structures. These findings demonstrate the potential of this and related systems to affect the metal coordination geometry and reactivity through simple ligand modifications. These complexes may have potential applications in the polymerization of olefins, similar to Brookhart catalysts, and could even stabilize Ni(II) with phosphines as coligands and provide a pathway for hydride complexes, which may function as organometallic catalysts for hydrogenation reactions.56

**EXPERIMENTAL SECTION**

All reactants were of reagent grade and used without further purification unless stated otherwise. Solvents were of HPLC quality and were freshly distilled under nitrogen before use. Dichloromethane was distilled by a standard procedure and, further, from calcium hydride. Toluene was distilled over sodium. Ethanol was purchased from Hayman and used as received. Electronic absorption spectra were recorded either on Shimadzu UV–vis 3100 or Jasco 650 spectrophotometers, and IR spectra were recorded on a Jasco 620 using the KBr pellet method. Proton nuclear magnetic resonance (1H NMR) spectra were recorded using Bruker Avance 400 and 500 spectrometers, and ESI-MS spectra were obtained on a Q-TOF-Mass microhybrid quadrupole time-of-flight mass spectrometer. The vibrating sample magnetometer measurements were carried out on a Lakeshore model 7407, with a maximum magnetic field value of 2.5 T.

**Synthesis of Metal Complexes.** Warning! Perchlorate salts are potentially explosive and should be handled only in small quantities and with care.37

The ligands, Lm and Lc, were prepared according to reported procedures.20

**Synthesis of [Ni(Lm)Cl]2.** A 2 mL MeOH solution of NiCl2·6H2O (42.7 mg, 0.179 mmol) was added to a 3 mL solution of the Lm ligand (103 mg, 0.359 mmol) in MeOH. The reaction mixture was stirred for about half an hour at room temperature to give a greenish-brown solution. An apple-green solid precipitated instantaneously when a 0.5 mL MeOH solution of NaClO4 (22.01 mg, 0.179 mmol) was added. The precipitate was filtered, vacuum dried, and weighed out to be 100 mg (73% yield). Deep-green block-shaped crystals were obtained by recrystal-
lization from CH$_2$Cl$_2$. FT-IR (KBr, cm$^{-1}$): 1609 (s), 1572 (w), 1482 (m), 1447 (s), 1300 (m), 1297 (w), 1286 (m), 1156 (m), 1109 (w), 1098 (w), 929 (m), 822 (m), 753 (vs), 622 (s), 504 (w), 426 (w). CHN analysis of $[\text{Ni(Lm)}_2]^+\text{Cl}^-$ (Evans, CDCl$_3$, RT): $\delta = -13.64$ (br), $\delta = -2.66$ (br), 25.46 (s), 54.44 (s), 65.40 (s), 96.01 (br), 113.31 (br), 117.35 (br), 193.96 (br), 330.56 (br). Magnetic moment (Evans, CDCl$_3$, RT): 3.11 $\mu$/Ni(II). ESI-MS (CH$_3$CN): $m/z$, 667 (85%), $[M - \text{Cl}]^+$; 471 (53%), $[\text{Ni(Lm)}^{+}]^+$; 287 (100%), $[\text{Ni(Lm)}^{+} - H]^+$.

**Synthesis of $[\text{Ni(Lm)}_2]\text{BrCl(O}_4\text{)}_2$.** In a reaction aimed at the preparation of 2b, a 3 mL MeOH solution of Ni(ClO$_4$)$_2$·6H$_2$O (128 mg, 0.350 mmol) was added to a solution of Lm (200 mg, 1.222 (14)). 1H NMR (CDCl$_3$, RT): δ $\delta = -14.10$ (s), $\delta = -4.63$ (s), $\delta = -3.22$ (s), $\delta = -2.66$ (br), 25.46 (s), 54.44 (s), 65.40 (s), 96.01 (br), 113.31 (br), 117.35 (br), 193.96 (br), 330.56 (br). Magnetic moment (Evans, CDCl$_3$, RT): 3.47 $\mu$/Ni(II). ESI-MS (CH$_3$CN): $m/z$, 711.34 (25%), $[M - \text{Cl}]^+$; 287 (100%), $[\text{Ni(Lm)}^{+} - H]^+$; 343 (20%), $[\text{Ni(Lm)}^{+} - 2\text{H}]^+$.

**Synthesis of $[\text{Ni(Lm)}_2]\text{Cl}_2$.** A 2 mL MeOH solution of Ni(ClO$_4$)$_2$·6H$_2$O (191.9 mg, 0.524 mmol) was added to 1 mL MeOH solution of Lm (300 mg, 1.05 mmol) under stirring at room temperature. The solution turned greenish-brown when stirred for about half an hour. Subsequently, the addition of NaI (78.6 mg, 0.524 mmol) resulted in a yellowish-green solid that weighed 215 mg (75% yield). The complex was found to be soluble in CH$_2$Cl$_2$, CH$_3$CN, and acetone and was sparingly soluble in MeOH and EtOH. The penta-green powder was recrystallized from CH$_2$Cl$_2$ to give dark green diamond-shaped crystals. FT-IR (KBr, cm$^{-1}$): 1609 (s), 1572 (w), 1483 (m), 1447 (m), 1400 (w), 1296 (w), 1199 (w), 1092 (vs), 928 (m), 865 (w), 821 (m), 755 (vs), 622 (s), 568 (w), 504 (w), 424 (w). CHN analysis of $[\text{Ni(Lm)}_2]^+\text{BrCl}^-$ (Evans, CDCl$_3$, RT): δ $\delta = -14.10$ (Br), $\delta = -4.63$ (Br), 24.30 (s), 52.84 (s), 65.13 (s), 94.41 (br), 116.54 (s), 125.01 (br), 187.28 (br), 341.62 (br). Magnetic moment (Evans, CDCl$_3$, RT): 3.47 $\mu$/Ni(II). ESI-MS (CH$_3$CN): $m/z$, 711.34 (25%), $[M - \text{Cl}]^+$; 287 (100%), $[\text{Ni(Lm)}^{+} - \text{H}]^+$; 343 (20%), $[\text{Ni(Lm)}^{+} - 2\text{H}]^+$.

**Single Crystal X-ray Diffraction.** Suitable single crystals of all six compounds were mounted on thin glass fibers with epoxy glue and optically aligned on a Bruker APEX II charge-coupled device X-ray diffractometer using a digital camera. Intensity data were measured at 25 °C using Mo Kα radiation ($\lambda = 0.7103$ A). APEX II software (Bruker AXS) was used for preliminary determination of the cell constants and data collection control. Determination of integral intensities and global refinement were performed using SAINT+ (Bruker AXS). The reflections were corrected for Lorentz and polarization effects, and empirical absorption corrections were applied using the SADABS program. The space groups were determined from systematic absences and confirmed by the results of refinement. The structures were solved by direct methods using the SHELXL software suite (SHELXS 86) and refined by the full-matrix least squares method (on F$_2$; SHELXL 97). All non-H (except some of the solvent molecules) were refined with anisotropic displacement parameters, and all H atoms of the organic ligands were placed at idealized positions and refined as riding atoms.
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