Research paper

Can pendant pyridyl arm assist the proton delivery in electrocatalysis?

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Abstract

Two nickel(II) complexes [Ni(L1)]2Br2 and [Ni(L2)]2Br2 ([H2L1Cl = 1-benzyl-3-(pyridin-2-ylmethyl)benzimidazolium chloride; H2L2Cl = 1,3-bis(pyridin-2-ylmethyl)benzimidazolium chloride] were synthesized as non-noble metal catalysts for the electrocatalytic hydrogen evolution reaction. Single-crystal X-ray crystallography revealed that the nickel metal centers in both compounds are in a square-planar geometry with like donor atoms of the two bidentate ligands in cis positions. The redox properties of the two compounds were studied using cyclic voltammetry. Electrocatalytic proton reduction experiments using these complexes were performed in DMF with acetic acid as the proton source. Compound [Ni(L2)]2Br2 containing two free pyridyl groups, not only exhibits higher electrocatalytic activity, but also has a smaller overpotential for the reduction of protons. The comparison of these results provides convincing experimental evidence for the pyridyl group acting as proton relay during the proton reduction process, although unfortunately the performance of the compounds is rather low.

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1. Introduction

Ligands with pendant, non-coordinating Lewis bases have received ample attention from inorganic chemists, because the additional Lewis base might affect the performance of molecular catalysts by providing binding sites for protons [1–4]. Notably, the introduction of base-functionalized pendant arms in the ligands of molecular catalysts has been quite important in the electrochemical research area for assisting in proton-coupled electron-transfer processes [5–8]. For example, Ni-diphosphane complexes bearing pendant amines as proton-binding sites have been shown to exhibit extremely high activity (TOF > 100,000 s−1) in electrocatalytic H2 evolution [9]. The molecular design of catalysts using pendant Lewis bases is inspired by the structures of natural enzymes, in which nearby amino acids close to the active site play are involved in temporary proton storage [8–11]. With the aim to investigate the role of the pendant amine group in dihydrogen evolution, DuBois’ group compared the electrocatalytic activity of the compound [Co(PPh2N)2(CH3CN)2][BF4]2 ([PPh2N]2 = 1,3,5-tetraphenyln-1,5-diaza-3,7-diphosphacyclooctane) incorporating pendant amines, with that of the compound [Co(dppp)(CH3CN)2][BF4]2 ([dppp = 1,3-bis(diphenylphosphoryl)propane] without pendant amines [10]. Although the two cobalt compounds are equipped with similar diphosphane ligands, the investigations showed that only the former one possesses catalytic activity in proton reduction.

Not only secondary or tertiary amines but also the pyridyl group can realize the function as a proton relay [7,12–14]. Recent DFT calculations on a pyridyl-functionalized cobalt diimine-dioxalogen catalyst have indicated that the non-coordinated pyridyl pendant group may help to reduce the energetic barrier for hydride formation in electrolysis [7]. Indeed it has been suggested that the pyridyl group may facilitate the reaction of a metal-hydride intermediate with a nearby proton [14]. In the research described in this paper, we aimed to explore the role of uncoordinated pyridyl groups in the proton reduction activity of nickel compounds with carbene ligands.

In the design of metal-carbene complexes pyridine-functionalized (benz)imidazole-2-ylidene ligands have received a lot of attention. Coordination compounds of these functionalized carbenes have been reported for a large array of metal centers such as Ag, Hg, Pd, Au, Ni, Ru and Pt [15–20]. For one and the same ligand, the configuration of the complexes will largely depend on the properties of the metal center. Setting the ligand precursor 1,3-bis(pyridin-2-ylmethyl)benzimidazolium chloride H2L2Cl as an example, PdIV and PdII ions can coordinate with one ligand in a tridentate mode with both pyridines bound in square-planar geometries [15,20]; RuIII ions may bind two ligands in a tridentate mode in an octahedral geometry [20], whereas Ag+ and Au+ ions can bind either one [17] or two ligands [19] with both pyridine arms non-coordinated.
Herein, we report the synthesis and characterization of two new nickel complexes of pyridyl-functionalized carbene ligands based on benzimidazole and the study of their electrocatalytic activity in H₂ evolution. We sought to determine whether the presence of pendant pyridyl groups could enhance H₂ evolution by acting as an intramolecular proton shuttle in electrocatalysis by comparing these two related compounds. N-heterocyclic carbenes are widely recognized as innocent ligands in organometallic chemistry [21]. The influence of a pendant pyridine arm in electrocatalysis might be more readily understood in the absence of other redox-active functional groups.

2. Result and discussion

2.1. Synthesis of the complexes

The ligand precursors 1-benzyl-3-(pyridin-2-ylmethyl)benzimidazolium chloride (HL1Cl) [22] and 1,3-bis(pyridin-2-ylmethyl)benzimidazolium chloride (HL2Cl) [20] were synthesized following literature methods with small modifications, in good yields of 77% and 87%, respectively. The nickel(II) bromide compounds [Ni(L1)₂]Br₂ and [Ni(L2)₂]Br₂ were obtained as yellow-colored powders from a melt of nickel acetate with these ligand precursors in tetra- butylammonium bromide (TBAB) in vacuum (Scheme 1). The complexes were characterized by NMR spectroscopy and electrospray ionization mass spectrometry (ESI-MS). The successful creation of the carbene moieties and their coordination to the nickel(II) center is indicated by the absence in the ¹H NMR spectra of the characteristic downfield signals for the imidazolium NCHN protons (10–12 ppm) in both complexes. The sharp and clear ¹H NMR spectra of the imidazolium NCHN protons (10–12 ppm) could enhance H₂ evolution by acting as an intramolecular proton shuttle in electrocatalysis by comparing these two related compounds. The ESI-MS spectra exhibit base peaks corresponding to [Ni(L₂)₂]⁺, which must comprise only a very small part of the nickel compound [Ni(L₂)₂]Br₂, in solution (see Fig. S1). This broadening is likely is caused by a binding and dissociation equilibrium resulting in a mixture of the low-spin square-planar nickel complex with 5-coordinate or octahedral species containing high-spin nickel centers. As can be seen from the NMR spectrum in Fig. S1, this must comprise only a very small part of the nickel ions, as the NMR peaks are all in the diamagnetic region, but just are slightly broadened. The ESIMS spectra exhibit base peaks corresponding to the [M-2Br]²⁻ cationic complexes. Crystalline samples of the Ni compounds were obtained in vacuo before elemental analysis was performed; however, the analytical results indicate that the water molecules found in the crystal lattice were not totally removed. During synthesis of the complexes the chloride counter ions of the ligands were replaced by the bromide ion from TBAB.

2.2. Structural characterization of the complexes

Single crystals of [Ni(L1)₂]Br₂ and [Ni(L2)₂]Br₂ suitable for X-ray structure determination were obtained using the liquid-liquid diffusion method. The crystallographic and refinement data are collected in Table S1. The crystallographic data and refinement details are collected in Table S1. The projection of the structures of the cationic nickel centers in [Ni(L1)₂]Br₂ and [Ni(L2)₂]Br₂ are shown in Fig. 1. Selected bond distances and angles are provided in Table 1.

The compound [Ni(L1)₂]Br₂ crystallizes in the space group P-1, whereas the compound [Ni(L2)₂]Br₂ crystallizes in the monoclinic space group Cc. Both compounds crystallized with two independent cationic complexes, four bromide ions and some disordered solvent molecules in the asymmetric unit. The coordination environment of the nickel centers in the two independent molecules is very similar, and for both compounds, the relevant bond distances and angles from only one of the two molecules are provided. The nickel ion in [Ni(L1)₂]Br₂ is in a four-coordinate square-planar geometry in which two ligands are bound in a bidentate C, N fashion with the C and N donor atoms in mutual cis positions (Fig. 1a). Although the ligand L₂ potentially is a tridentate ligand, in [Ni(L₂)₂]Br₂ the nickel ion is also found in a square-planar geometry with donor atoms of the two bidentate ligands in cis positions (see Fig. 1b).

The Ni–C bond distances in the two compounds are around ca. 1.87 Å whereas the Ni–N bonds are ca. 1.94 Å. Both the Ni–C and Ni–N bond distances are shorter by 0.2 Å in comparison with the reported Ru [20], Pt [15,16], and Au compounds [19], which can be attributed to the smaller ionic radius of the nickel ion. In the compound [Ni(L₂)₂]Br₂ the nitrogen atom of one of the pendant pyridyl groups resides in an apical position of the nickel center at a distance of ca. 2.98 Å; if this interaction is taken into account the geometry of the nickel ion may be regarded as a distorted square pyramid (Fig. 1c). This additional weak interaction of the pyridyl group may be retained in solution, or even be stronger, thus causing the rather broad signals observed in the ¹H NMR spectra of this compound.

The square-planar geometries of the nickel ions in [Ni(L1)₂]Br₂ and [Ni(L2)₂]Br₂ were evaluated with an equation derived from a report by Yang [23,24]. The largest and second-largest L–Ni–L angles in [Ni(L₁)₂]Br₂ are 171.7(1)° and 174.0(1)°, resulting in a τ value of 0.100 indicative of only a very slight distortion of the square-planar geometry. For compound [Ni(L₂)₂]Br₂ a τ value of 0.023 is calculated, indicating that the geometry of the central ion (not taking into account the axial coordination of pyridine) is close to the ideal square-planar geometry.

Complexes of other metal ions bearing the ligand L₂ have been reported by Adhikary et al. [20], Pt(II) was found to coordinate with one ligand L₂ to form the complex [Pt(L₂)Cl]PF₆ with the Pt(II) ion in a square-planar geometry, whereas the Ru(II) ion binds two ligands L₂ to form the compound [Ru(L₂)₂](PF₆)₂ with the ruthenium ion in an octahedral geometry. A structure has been reported of a nickel compound containing two 1,3-bis(pyridin-2-ylmethyl)imidazolium ligands, in which similar to [Ni(L₂)₂]Br₂, the ligands are arranged with like donor atoms of the two bidentate ligands in cis positions [16]. The nickel ion in this compound is also in a square-planar geometry and one of the pendant pyridyl groups resides in an apical position of the nickel center at a distance of 3.345 Å, significantly larger than the one in [Ni(L₂)₂]Br₂.

2.3. Electrochemical properties of the complexes

The redox potentials of the nickel compounds were evaluated using cyclic voltammetry. The electrochemical potentials all are reported relative to the Ag/AgCl reference electrode. The cyclic voltammogram (CV) of [Ni(L1)₂]Br₂ shows two reductive waves, the first of which is reversible at a scan rate of 0.1 V/s (Fig. 2). The reversible redox process located at E₁/₂ = -1.00 V vs. Ag/AgCl is diffusion controlled, as shown by the linear dependence of the peak current on the square root of the scan rate (Fig. S2). This redox wave is characterized by a peak-to-peak separation ΔEₚ of 80 mV.
(in our conditions \( \Delta E_p \) for the Fe\(^{0/+}\) couple was found to be 95 mV), indicating a one-electron transfer. Thus, we assign this first wave to a Ni(II)/Ni(I) redox process. The quasi-reversible reduction event at \(-1.29\) V is tentatively ascribed to a Ni(I)/Ni(0) redox process. The assignment of these two redox waves to nickel-centered reductions is supported by the fact that the ligand precursor HL\(_1\)Cl is electrochemically inert in the range of 0 to \(-1.5\) V. The second reductive wave at \(-1.29\) V gradually becomes reversible with increasing scan rate, whereas the wave becomes less reversible with decreasing scan rates (see Fig. S3). This behavior indicates the occurrence of a dissociative electron transfer (DET) process \([25]\), meaning that after the reduction of [Ni\(^{II}(L_1)\)\(^2+\)] to [Ni\(^{II}(L_1)\)\(^+\)]

a stepwise chemical reaction is coupled to the second electron transfer, generating a new product.

Similarly, the CV of compound [Ni\(^{II}(L_2)\)\(^2+\)]Br\(_2\), containing two uncoordinated pyridyl groups, shows one reversible redox process at \(E_{1/2} = -1.05\) V followed by an irreversible reductive process at \(-1.27\) V (Fig. 2). The first reversible redox process of [Ni\(^{II}(L_2)\)\(^2+\)]Br\(_2\)

**Table 1**

|                  | [Ni\(^{II}(L_1)\)\(^2+\)]Br\(_2\) | [Ni\(^{II}(L_2)\)\(^2+\)]Br\(_2\) |
|------------------|----------------------------------|----------------------------------|
| Ni1–C12A         | 1.880(3)                         | 1.850(4)                         |
| Ni1–C42A         | 1.872(2)                         | 1.863(4)                         |
| Ni1–N22A         | 1.947(2)                         | 1.938(4)                         |
| Ni1–N52A         | 1.933(2)                         | 1.950(4)                         |
| Ni1–N62A         | -‡                               | 2.984(4)                         |
| N52A–Ni1–N22A    | 95.10(9)                         | 91.25(15)                        |
| N52A–Ni1–C42A    | 86.47(10)                        | 87.32(17)                        |
| C42A–Ni1–C12A    | 93.12(11)                        | 94.33(18)                        |
| C12A–Ni1–N22A    | 86.15(10)                        | 87.09(17)                        |
| N52A–Ni1–C12A    | 174.00(10)                       | 178.08(18)                       |
| C42A–Ni1–N22A    | 171.71(11)                       | 178.56(17)                       |

Fig. 2. CVs of the compounds [Ni\(^{II}(L_1)\)\(^2+\)]Br\(_2\) (solid line) and [Ni\(^{II}(L_2)\)\(^2+\)]Br\(_2\) (dashed line). Experiments were performed at a complex concentration of 1 mM in DMF with 100 mM TBAPF\(_6\) as the supporting electrolyte at scan rate 0.1 V/s, using a glassy carbon working electrode.

Fig. 1. Displacement ellipsoid plots of the cationic complexes (a) [Ni\(^{II}(L_1)\)\(^2+\)]Br\(_2\), (b) and (c) [Ni\(^{II}(L_2)\)\(^2+\)]Br\(_2\) are drawn at 50% probability level with selected atom numbering. For clarity the hydrogen atoms, bromide ions, solvent molecules and the lattice water molecules are omitted and parts of the ligands are displayed in wireframe.
is at a slightly more negative potential than that of \([\text{Ni(L1)}_2\text{Br}_2]\), indicating a slightly higher electron density at the nickel center, likely because of the interaction with one or both of the pendant pyridyl groups in solution. The reversibility of the second reductive wave at \(E_{pc2} = -1.29\ \text{V}\) of this tetrapyridyl-functionalized compound \([\text{Ni(L2)}_2\text{Br}_2]\) appeared not to depend on the scan rate. Even when the scan rate was increased to 500 mV/s the second reductive process still was quasi-reversible (see Fig. S4). The electrochemical data are summarized in Table 2.

### 2.4. Electrocatalytic activity for proton reduction in DMF

The two compounds were examined for their activity in the electrocatalytic proton reduction in DMF, using acetic acid (HOAc) as a weak proton source. Before each scan the working electrode was freshly polished.

Compared to the CV of acetic acid (10 mM) in DMF, after addition of either the complex \([\text{Ni(L1)}_2\text{Br}_2]\) or \([\text{Ni(L2)}_2\text{Br}_2]\) a relatively large current indicating the reduction of protons was observed at a potential around \(-2\ \text{V}\) vs Ag/AgCl (Fig. 3). Two small irreversible peaks appear before the electrocatalytic wave, approximately at \(-1.04\ \text{V}\) and \(-1.17\ \text{V}\) for \([\text{Ni(L1)}_2\text{Br}_2]\) and at \(-1.09\ \text{V}\) and \(-1.20\ \text{V}\) for \([\text{Ni(L2)}_2\text{Br}_2]\). These peaks were broad and close to each other making it difficult to identify them separately. Only a broad plateau can be recognized when showing these waves together with the following catalytic wave. A zoom of CVs (ranging from \(-0.5\ \text{V}\) to \(-1.5\ \text{V}\)) is provided in Fig. S5.

For both compounds the first small reductive wave occurs at the exact same potential where the metal-centered Ni\(^{II}/\text{Ni}\) reduction process was observed (see previous section and Fig. S5). The second small reductive potential occurs at a more positive potential compared to the ones in the absence of acid, indicating that there is a chemical step before the second electron transfer takes place. The electrocatalysis therefore most likely proceeds via an ECEC mechanism [26]. The catalytic onset potential for proton reduction in presence of \([\text{Ni(L2)}_2\text{Br}_2]\) at \(-1.6\ \text{V}\) occurs at a lower potential than in the presence of \([\text{Ni(L1)}_2\text{Br}_2]\) at \(-1.8\ \text{V}\). Furthermore, in the same conditions the compound \([\text{Ni(L2)}_2\text{Br}_2]\) shows the larger catalytic current, indicating a higher electrocatalytic activity than that of \([\text{Ni(L1)}_2\text{Br}_2]\) (Fig. 3).

With the aim to compare the electrocatalytic activity of the two compounds in a more quantitative way, controlled-potential coulometry experiments (CPC) were carried out. The charge consumptions over time of solutions containing just the pure acid or one of the nickel catalysts. The quantity of dihydrogen gas generated during the CPC experiments can be estimated from the charge consumption of solutions containing both the nickel compound and the acid, after subtraction of the charge consumption of the blanks. The two complexes \([\text{Ni(L1)}_2\text{Br}_2]\) or \([\text{Ni(L2)}_2\text{Br}_2]\) are estimated to produce \(3 \times 10^{-4}\ \text{mmol}\) and \(5 \times 10^{-4}\ \text{mmol}\ \text{H}_2\) respectively in 10 min in presence of \(6 \times 10^{-3}\ \text{mmol}\) catalyst at a potential of \(-1.80\ \text{V}\) (assuming that all electrons are used in proton reduction). Almost a doubled charge consumption was found for the complex \([\text{Ni(L2)}_2\text{Br}_2]\) equipped with two additional pendant pyridine groups, indicating a higher catalytic activity. However, despite this improvement, still the electrocatalytic activity of \([\text{Ni(L2)}_2\text{Br}_2]\) is disappointingly low.

Generally, the mechanism can be divided in three steps for a hydrogen-evolution process based on a metal-containing homogenous catalyst. The first step is the reduction of the metal center; if the metal center is equipped with an electron-withdrawing ligand, the metal center is more readily reduced. The second step is the formation of the metal hydride intermediate, which is facilitated by the presence of an electron-donating ligand. The third step is the generation of dihydrogen: the metal-hydride intermediate...
reacts with another proton and electron to release dihydrogen [27]. Based on this mechanism, generally a molecular catalyst will work at a small overpotential but with relatively low efficiency for electron-withdrawing ligands, or with high efficiency but at relatively larger overpotentials for electron-donating ligands.

As discussed in the previous section, the ligand \( L_2 \) is the more electron donating of the two in absence of acid, so that the nickel center in \([\text{Ni}(\text{L}_2)_2]\text{Br}_2\) is reduced at more negative potentials. Unexpectedly, the compound \([\text{Ni}(\text{L}_2)_2]\text{Br}_2\), not only shows higher electrocatalytic activity but also a smaller overpotential (Fig. 3). Thus, the presence of the free pyridyl pendant arms in the compound \([\text{Ni}(\text{L}_2)_2]\text{Br}_2\) apparently reduces the overpotential for the electrocatalytic proton reduction. The additional pyridine groups act as an intramolecular proton acceptor and proton shuttle assisting in the generation of dihydrogen [7]. Based on the assistance of the pyridyl groups for complex \([\text{Ni}(\text{L}_2)_2]\text{Br}_2\), the catalytic mechanism tentatively can be described as shown in Scheme 2. First the non-coordinated pyridyl groups are protonated in the presence of acid. Then the metal center is reduced to Ni\(^{1+}\), followed by the formation of a metal-hydride. Finally the hydride reacts with the proton on the pyridyl arm with the release of dihydrogen.

### 3. Conclusion

Two nickel(II) complexes \([\text{Ni}(\text{L}_1)_2]\text{Br}_2\) and \([\text{Ni}(\text{L}_2)_2]\text{Br}_2\) with different ligands were synthesized as non-noble metal molecular catalysts for the hydrogen evolution reaction. The nickel center in both compounds is in a square-planar geometry with two bidentate ligands in cis arrangements. The binding of two potentially tridentate ligands in this configuration in the structure of \([\text{Ni}(\text{L}_2)_2]\text{Br}_2\) results in the presence of two free pyridine groups. The redox properties of the two complexes were determined using cyclic voltammetry and the electrocatalytic dihydrogen generation with these complexes in DMF was studied using acetic acid as the proton source. The complex \([\text{Ni}(\text{L}_2)_2]\text{Br}_2\) containing two free pyridyl groups exhibited not only the higher activity in electrocatalysis, but also a smaller overpotential for proton reduction. This result indicates that the presence of the free pyridyl groups help to break the anti-correlation between the energy of the reduction and the catalytic activity. The comparison of the catalytic properties of these two related structures (both with redox-innocent carbene backbones) provide convincing experimental evidence of the pyridyl group acting as proton relay during the proton reduction process. Unfortunately, it appears that this type of carbene-based ligands in combination with nickel do not yield very active electrocatalysts for proton reduction.

### 4. Experimental

#### 4.1. Materials

Commercial chemicals were used without further purification. Acetonitrile and diethyl ether were obtained from a PureSolv MD5 solvent dispenser. Dry methanol and dimethylformamide were prepared by adding molecular sieves into commercial anhydrous solvent. All air-sensitive reactions were performed under argon or dinitrogen gas using standard Schlenk techniques unless mentioned otherwise.

#### 4.2. Analytical methods

\(^1\)H and \(^{13}\)C spectra were recorded on a Bruker 300 DPX/Bruker 400 AV spectrometer. Mass spectra were obtained using a Finnigan Aqua Mass Spectrometer (MS) with electrospray ionization (ESI). Elemental analyses were performed by the Mikroanalytisches Laboratorium Kolbe, Germany.

Cyclic voltammetry was recorded with an Autolab PGstat10 potentiostat controlled by GPE4 software under argon. A 3 mm diameter glassy carbon electrode was used as working electrode and a Pt wire as the counter electrode. The experimental reference was an Ag/AgCl electrode in the electrolyte solution. Cyclic voltammetry was performed in dry DMF with 0.1 M tetrabutylammonium hexafluoridophosphate (TBAPF\(_6\)) as the supporting electrolyte under a stream of argon at room temperature. All potentials are given versus Ag/AgCl. Ferrocene was added after each experiment as an internal standard. Under these conditions the Fe\(^{2+/3+}\) couple was located at \(E_{1/2} = 0.510\) V vs. Ag/AgCl with a \(\Delta E_p\) of 95 mV. The working electrode surface was polished, ultrasonically cleaned and rinsed before each single CV measurement.

Controlled-potential coulometry (CPC) experiments were carried out with an Autolab PGstat10 potentiostat controlled by GPE4 software under argon. A 3 mm diameter glassy carbon electrode was used as working electrode and a platinum wire as the counter electrode. 1 mM complex and 50 mM acetic acid were added in 6 mL dry degassed DMF with 0.1 M TBAPF\(_6\) as the supporting electrolyte under a stream of argon at room temperature. A CPC experiment was run at \(-1.8\) V for 600 s, while the solution was stirred continuously. The blank and reference CPC experiments (only acid or only catalyst added) were performed using the same conditions.

#### 4.3. Single crystal X-ray crystallography

All reflection intensities were measured at 110(2) K using a SuperNova diffractometer (equipped with Atlas detector) with Mo K\(\alpha\) radiation (\(\lambda = 0.71073\) Å) under the program CrysalisPro (Version 1.171.36.32 Agilent Technologies, 2013). The same program was used to refine the cell dimensions and for data reduction. The structure was solved with the program SHELXS-2014/7 [28,29], and was refined on \(F^2\) with SHELXL-2014 [28,29]. Numerical absorption correction based on gaussian integration or analytical numeric absorption correction over a multifaceted crystal model was applied using CrysalisPro. The temperature of the data collection was controlled using the system Cryojet (manufactured by Oxford Instruments). The H atoms were placed at calculated positions (unless otherwise specified) using the instructions AFIX 23 or AFIX 43 with isotropic displacement parameters having val-
ues 1.2 or 1.5 \text{eq} of the attached C atoms. For [Ni(L1)2]Br2, the H atoms attached to O1W (ordered lattice water molecule) and O2W (only for the major component of the disorder) could be retrieved from difference Fourier maps. Their coordinates were refined freely but the O–H and H...H distances were restrained to be 0.84(1) and 1.33(1) Å, respectively. For [Ni(L2)2]Br2, the O–H and H...H distances were restrained to be 0.84(3) and 1.33(3) using the DFIX instruction. The atoms attached to O4W, O5W, O6W, O7W, O8W and O9W (partially occupied or disordered lattice water molecules) could not be retrieved.

The structure of [Ni(L1)2]Br2 is partly disordered. One phenyl group (C61A-C66A) and one Br– counter ion are found disordered over two orientations; the occupancy factors of the major components of the disorder refine to 0.59(4) and 0.923(5), respectively. One of the three lattice water molecule (O2W) is disordered over three orientations, and the occupancy factors refine to 0.808(3), 0.109(3) and 0.083(3). The occupancy of the lattice water molecule O3W was refined freely, and its value refines to 0.639(8). The H atoms attached to O3W could not be retrieved. ADDSYM detects a pseudo translation of \(\frac{1}{2}a + \frac{1}{2}c\). However, the conformations of the two crystallographically independent molecules are different as the orientations of the phenyl groups are significantly different.

The structure of [Ni(L2)2]Br2 is partly disordered. One of the pyridine rings (from molecule A) is found to be disordered over two orientations, and the occupancy factor of the major component of the disorder refines to 0.52(2). In the asymmetric unit, there are five Br– sites with all occupancies lower than 1. Two sites (Br1 and Br2) are ordered, and their occupancy factors refine to 0.940(2) and 0.942(2), respectively. The remaining counter ions (Br3/Br3’, Br4/Br4’ and Br5/Br5’) are disordered over two orientations. The occupancy factors for Br3, Br4, Br4’, Br5 and Br5’ refine to 0.853(3), 0.086(3), 0.624(2), 0.0494(15), 0.391(2) and 0.1149(16), respectively. The sum of all occupancies factors for the Br– counter ion was restrained to be 4 using the SHELX instruction. The occupancy factors for O4W/O4W’ (disordered), O5W/O5W’ (disordered), O6W, O7W, O8W and O9W refine to 0.434(1)/0.566(1), 0.715(8)/0.285(8), 0.674(15), 0.671(16), 0.429(18) and 0.41(2), respectively.

CCDC-1511049 and 1,511,050 for [Ni(L1)2]Br2 and [Ni(L2)2]Br2 contain the supplementary crystallographic data for this paper. These data can be obtained free of charge from The Cambridge Crystallographic Data Center via www.ccdc.cam.ac.uk/data_request/cif.

4.4. Ligand and complex synthesis

N-Pyridylmethyl-benzimidazole, [30] ligand precursors HL1Cl [22], HL2Cl [20], and the complexes [Ni(L1)2]Br2 and [Ni(L2)2]Br2 [31–33] were synthesized following literature methods with small modifications.

4.5. [Ni(L1)2]Br2

HL1Cl (0.67 g, 2 mmol), anhydrous Ni(OAc)2 (0.17 g, 1 mmol) and tetrabutylammonium bromide (1.5 g) were weighed into a 10 mL round-bottomed flask and dried for 4 h at 80 °C in a Schlenk setup under vacuum. The reaction mixture was stirred for 48 h at 130 °C under vacuum. The mixture was then heated to room temperature and triturated with H2O. The water layer was washed with DCM (3 × 20 mL), and a brown crystalline solid was obtained from the water solution in 5% yield (40 mg). \(^1\)H NMR (300 MHz, DMSO-d6) \(\delta = 8.53, 8.32, 8.07, 7.57, 7.45, 7.41, 7.31, 7.08, 6.99, 6.43, 5.75, 4.96\). MS m/z found (calc): [M–2Br]\(^{2+}\) 329.1 (329.1). C\(_8\)H\(_{12}\)N\(_2\)NiBr\(_2\)-2.5H\(_2\)O. Calc. H 4.31, C 52.81, N 12.97; Found H 4.80, C 53.08, N 12.99.

Conflicts of interest

There are no conflicts to declare.

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Appendix A. Supplementary data

Supplementary data associated with this article can be found in the online version, at https://doi.org/10.1016/j.jica.2018.02.030.

References

[1] V.G. Organo, A.S. Filatov, J.S. Quartararo, Z.M. Friedman, E.V. Rybak-Akimova, Inorg. Chem. 48 (2009) 8456–8468.
[2] P. Zhang, M. Wang, Y. Yang, D. Zheng, K. Han, L. Sun, Chem. Commun. 50 (2014) 14153–14156.
[3] E.I. Musina, V.V. Khizasnovofa, I.D. Stelnik, M.I. Valtov, Y.S. Spirdonova, D.B. Krivolapov, I.A. Litvinov, M.K. Kadirov, P. Lonnecke, E. Hey-Hawkins, Y.H. Budnikova, A.A. Karasik, O.G. Sinyashin, Chem. Eur. J. 20 (2014) 3169–3182.
[4] J. Berding, T.F. van Dijkman, M. Lutz, A.L. Spek, E. Bouwman, Dalton Trans. (2009) 6964–6955.
[5] R. Tatevatsu, T. Ionomata, T. Ozawa, H. Masuda, Angew. Chem. 56 (2015) 5247–5250.
[6] A. Kocher, M. O’Hagan, E.S. Wiedner, M. van Gastel, Chem. Eur. J. 21 (2015) 10338–10347.
[7] P.F. Huo, C. Uyeda, J.D. Goodpaster, J.C. Peters, T.F. Miller, ACS Catal. 6 (2016) 6114–6123.
[8] D.L. DuBois, R.M. Bullock, Eur. J. Inorg. Chem. (2011) 5290–5293.
[9] M.L. Helm, M.P. Stewart, R.M. Bullock, M.R. DuBois, D.L. DuBois, Energy Environ Sci. 1 (2008) 167–174.
[10] P.D. Tran, J. Barber, Phys. Chem. Chem. Phys. 14 (2012) 13772–13784.
[11] D.J. Martin, B.D. McCarthy, C.L. Donley, J.L. Dempsey, Chem. Commun. 51 (2015) 5290–5293.
[12] N. Queyriaux, R. Tj, J. Jane, M. Massin, V. Artero, M. Chavarot-Kerdoud, Coord. Chem. Rev. 304–305 (2015) 3–19.
[13] J. Wang, C. Li, Q. Zhou, W. Wang, Y. Hou, B. Zhang, X. Wang, Catal. Sci. Techn. 6 (2016) 8482–8489.
[14] Q.-X. Liu, H.-L. Li, X.-J. Zhao, S.-S. Ge, M.-C. Shi, G. Shen, Y. Zang, X.-G. Wang, Inorg. Chem. Acta 376 (2011) 437–445.
[15] C. Chen, H. Qiu, W. Chen, J. Organomet. Chem. 696 (2012) 4166–4172.
[16] X. Zhang, S. Gu, Q. Xia, W. Chen, J. Organomet. Chem. 694 (2009) 2359–2367.
S. Luo et al. / Inorganica Chimica Acta 477 (2018) 24–30

[18] L.B. Munro, V.J. Catalano, Eur. J. Inorg. Chem. (2014) 4994–5007.
[19] K. Chen, M.M. Nenzel, T.M. Brown, V.J. Catalano, Inorg. Chem. 54 (2015) 6900–6905.
[20] S. Das Adhikary, T. Samanta, G. Roymahapatra, F. Loiseau, D. Jouvenot, S. Giri, P.K. Chattaraj, J. Dinda, New J. Chem. 34 (2010) 1974–1980.
[21] J.-N. Luy, S.A. Hauser, A.B. Chaplin, R. Tonner, Organometallics 34 (2015) 5099–5112.
[22] J. Liu, M. Liu, Y. Yue, M. Yao, K. Zhuo, Chin. J. Chem. 30 (2012) 644–650.
[23] M.H. Reineke, M.D. Sampson, A.L. Rheingold, C.P. Kubiak, Inorg. Chem. 54 (2015) 3211–3217.
[24] L. Yang, D.R. Powell, R.P. Houser, Dalton Trans. (2007) 955–964.
[25] G. Armendariz-Vidales, C. Frontana, Phys. Chem. Chem. Phys. 17 (2015) 29299–29304.
[26] C. Costentin, J.-M. Saveant, ChemElectroChem 1 (2014) 1226–1236.
[27] M.T.M. Koper, E. Bouwman, Angew. Chem. Int. Ed. 49 (2010) 3723–3725.
[28] G.M. Sheldrick, Acta Crystallogr. Sect. A: Found. Crystallogr. 64 (2008) 112–122.
[29] G.M. Sheldrick, Acta Crystallogr. C Struct. Chem. 71 (2015) 3–8.
[30] J. Dinda, S.D. Adhikary, S.K. Seth, A. Mahapatra, New J. Chem. 37 (2013) 431–438.
[31] J. Berding, M. Lutz, A.L. Spek, E. Bouwman, Organometallics 28 (2009) 1845–1854.
[32] S. Luo, M.A. Siegler, E. Bouwman, Eur. J. Inorg. Chem. (2016) 4693–4700.
[33] H.V. Huynh, C. Holtgrewe, T. Pape, L.L. Koh, E. Hahn, Organometallics 25 (2006) 245–249.