

## Supramolecules Self-assembled by Nickel(II) Hexaazamacrocycles Bearing Imidazole Pendants

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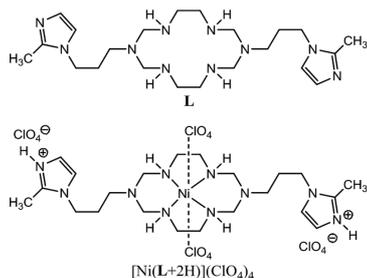
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Metallo-supramolecules self-assembled by a discrete number of molecules attract attention due to their fascinating structural features as well as potential applications in catalysis, gas storage and separation, molecular magnets, and molecular devices.<sup>1–6</sup> Non-covalent intermolecular interactions such as hydrogen bonds,  $\pi$ - $\pi$  interactions and C-H... $\pi$  interactions in addition to metal-ligand covalent bonds are frequently observed in the process of self-assembling discrete molecules to form large supramolecular structures.<sup>7–11</sup> Among various metallomacrocycles, the complexes of a macrocycle **L** (**L** = 3,10-bis(3-(2-methyl-1-imidazolyl)propyl)-1,3,5,8,10,12-hexaazacyclotetradecane) bearing two imidazole pendants on the macrocycle are of special interest owing to their favorable structural features for the formation of metallo-supramolecules. The imidazole pendants on **L** are expected to participate in intermolecular interactions as hydrogen bond donors and/or acceptors. The involvement of C-H... $\pi$  interactions and  $\pi$ - $\pi$  interactions between the imidazole pendants are another type of intermolecular forces to be anticipated in the self-assembly process of metallomacrocycles with **L**. With the structural advantages of **L** described above in mind for the assembling metallo-supramolecules, we attempted and successfully self-assembled the nickel(II) complexes of macrocycle **L**.

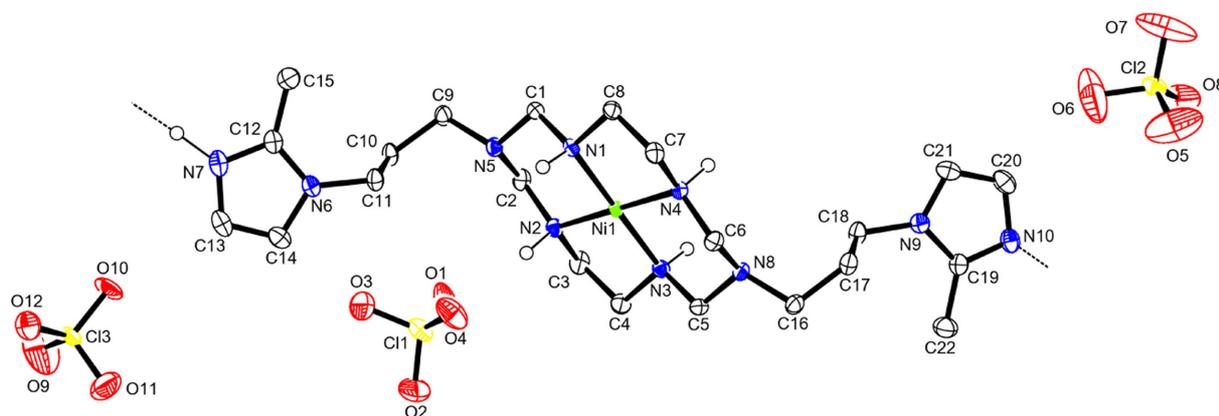
Herein, we report two macrocyclic nickel(II) supramolecules  $[\text{Ni}(\text{L}+\text{H})](\text{ClO}_4)_3$  (**1**) and  $[\text{Ni}(\text{L})](\text{ClO}_4)_2 \cdot 2\text{H}_2\text{O}$  (**2**) with their structural details.



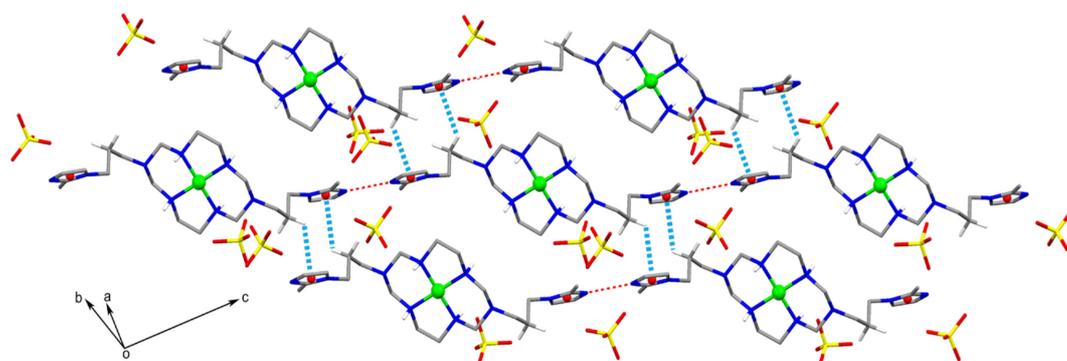
The nickel(II) precursor complex  $[\text{Ni}(\text{L}+2\text{H})](\text{ClO}_4)_4$  was synthesized and isolated by a well-known one-pot template condensation reaction of  $\text{Ni}(\text{OAc})_2 \cdot 4\text{H}_2\text{O}$ , ethylenediamine, paraformaldehyde, and 1-(3-aminopropyl)-2-methyl-1*H*-imidazole, followed by acidification of imidazole pendants with  $\text{HClO}_4$ .<sup>7–9</sup> The title complexes  $[\text{Ni}(\text{L}+\text{H})](\text{ClO}_4)_3$  (**1**) and  $[\text{Ni}(\text{L})](\text{ClO}_4)_2 \cdot 2\text{H}_2\text{O}$  (**2**) were obtained by the deprotonation of  $[\text{Ni}(\text{L}+2\text{H})](\text{ClO}_4)_4$  with triethylamine. The partly deprotonated complex **1** (orange needles) and the fully deprotonated complex **2** (yellow plates) were formed simultaneously in the deprotonation and crystallization processes even under the excess use of triethylamine. Crystals of **1** and **2** were able to be separated out manually with a needle under the microscope as their colors and shapes are quite different from each other.

The crystal structure of **1** with selected interatomic distances and angles is shown in Fig. 1. The structure of **1** consists of the macrocyclic nickel(II) cation  $[\text{Ni}(\text{L}+\text{H})]^{3+}$  and three perchlorate anions. In **1**, the coordination geometry about the nickel(II) ion exhibits a square-plane by coordinating to the four secondary nitrogen atoms of the macrocycle. The N-H interatomic distances vary from 1.9297(17) to 1.9374(17) Å. They are slightly longer than those expected in square-planar nickel(II) macrocycles (Ni-N = 1.88–1.91 Å for square planar species, Ni-N = 2.07–2.10 Å for octahedral species).<sup>12–14</sup> As is common in the nickel(II) macrocycles, the skeleton of the macrocycle adopts the most stable *trans-III* (*R,R,S,S*) configuration.<sup>15,16</sup>

The pertinent feature of the structure of **1** is that one of the imidazole pendants remains protonated. Thus, the protonated nitrogen atom of the imidazole pendant acts as hydrogen bond donor to the deprotonated nitrogen atom of the imidazole pendant belonging to the neighboring nickel(II) macrocycle. The hydrogen bonding interaction between the protonated nitrogen atoms and deprotonated nitrogen atoms results in the formation of a 1D macro-



**Figure 1.** Molecular structure of  $[\text{Ni}(\text{L}+\text{H})](\text{ClO}_4)_3$  (**1**) with atom-labeling scheme. Hydrogen atoms are omitted for clarity other than those on secondary nitrogen atoms and one of the protonated nitrogen atom of imidazole rings. Selected interatomic distances ( $\text{\AA}$ ) and angles ( $^\circ$ ). Ni1-N1, 1.9353(17); Ni1-N2, 1.9325(17); Ni1-N3, 1.9374(17); Ni1-N4, 1.9297(17); N1-Ni1-N2, 93.21(7); N1-Ni1-N4, 86.73(7); N2-Ni1-N3, 86.82(7); N3-Ni1-N4, 93.24(7).



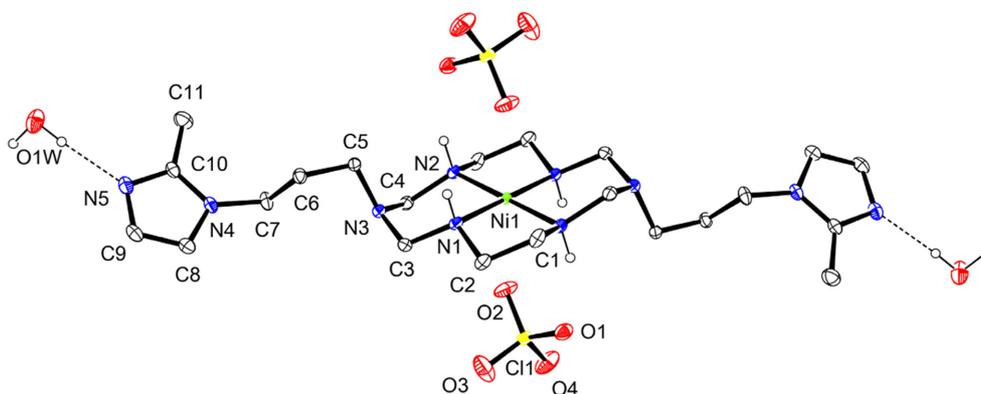
**Figure 2.** View of 2D supramolecular structure of **1** by hydrogen bonds (···) and C-H··· $\pi$  interactions (···). Hydrogen atoms are omitted for clarity other than those participating in hydrogen bonds and C-H··· $\pi$  interactions.

cyclic nickel(II) supramolecule (Fig. 2,  $\{\text{D-H}\cdots\text{A} = \text{N7-H7N}\cdots\text{N10}$ ;  $d(\text{H}\cdots\text{A}) = 1.526 \text{ \AA}$ ,  $d(\text{D}\cdots\text{A}) = 2.691 \text{ \AA}$ ,  $\angle(\text{DHA}) = 177.69^\circ$ ).

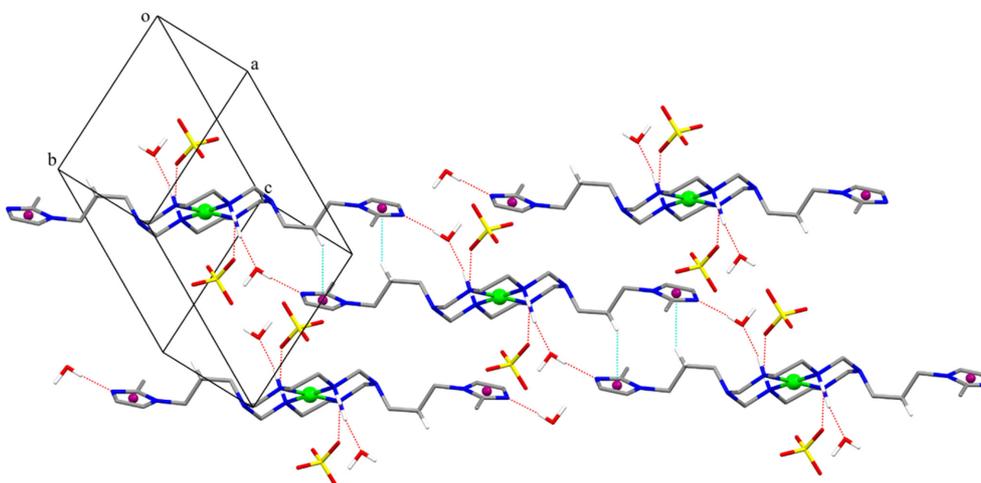
The 1D chains formed by N-H···N hydrogen bonds are further extended by weak C-H··· $\pi$  interactions between the methylene groups of aminopropyl-methyl-imidazole pendants and adjacent imidazole rings, forming a 2D supramolecule (Fig. 2,  $\{\text{C-H}\cdots\pi = \text{C10-H10A}\cdots\text{Im}_{\text{centroid}}$ ;  $d(\text{H}\cdots\pi) = 3.054 \text{ \AA}$ ,  $d(\text{C}\cdots\pi) = 3.468 \text{ \AA}$ ,  $\angle(\text{CH}\pi) = 106.52^\circ$ , Im = imidazole}). The C-H··· $\pi$  interaction observed in **1** is close to type III ( $d(\text{H}\cdots\pi) \leq 3.05 \text{ \AA}$ ,  $\alpha < 150^\circ$ ) according to the classification of Malone *et al.*<sup>17</sup> It is generally understood that even weaker intermolecular interactions could affect profoundly on the molecular-packing patterns of molecules.<sup>18</sup>

The molecular structure of **2** is given in Fig. 3. As shown in Fig. 3, the structure is composed of the macrocyclic nickel(II) cation, two perchlorate counter anions and two lattice water molecules. Both of the nitrogen

atoms of the imidazole pendants are deprotonated. The coordination environment around the nickel(II) ion is square-planar. The Ni-N interatomic distances of 1.9341(14)  $\text{\AA}$  and 1.9340(14)  $\text{\AA}$  are normal and comparable to those found in **1** and in related square-planar nickel(II) macrocycles.<sup>13</sup> Lattice water molecules and perchlorate anions participate in multiple types of hydrogen bonds with nitrogen atoms of the macrocycle as well as the nitrogen atoms of secondary amines of the macrocycle. In **2**, the important intermolecular force for the formation of 1D supramolecule is C-H··· $\pi$  interactions. Similar to that observed in **1**, the presence of C-H··· $\pi$  interaction between the methylene group of the aminopropyl-methyl-imidazole and imidazole ring links macrocyclic nickel(II) units together to form a 1D supramolecule {Fig. 4, C-H··· $\pi = \text{C6-H6B}\cdots\text{Im}_{\text{centroid}}$ ;  $d(\text{H}\cdots\pi) = 2.905 \text{ \AA}$ ,  $d(\text{C}\cdots\pi) = 3.650 \text{ \AA}$ ,  $\angle(\text{CH}\pi) = 132.77^\circ$ , Im = imidazole}. It looks that the C-H··· $\pi$  interaction observed in **2** is more effective than that in **1** from the point of view of interaction distances and angles.



**Figure 3.** Molecular structure of  $[\text{Ni}(\text{L})](\text{ClO}_4)_2 \cdot 2\text{H}_2\text{O}$  (**2**) with atom-labeling scheme. Hydrogen atoms are omitted for clarity other than those on secondary nitrogen atoms and lattice water molecules. Selected interatomic distances (Å) and angles ( $^\circ$ ). Ni1-N1, 1.9341(14); Ni1-N1#1, 1.9340(14); Ni1-N2, 1.9293(14); Ni1-N2#1, 1.9293(14); N1-Ni1-N2, 93.58(6); N1-Ni1-N2#1, 86.42(6). Symmetry transformations used to generate equivalent atoms: #1  $-x+1, -y+1, -z+1$ .



**Figure 4.** View of a 1D supramolecular structure of **2** by hydrogen bonds ( $\cdots$ ) and C-H $\cdots\pi$  interactions ( $\cdots$ ). Hydrogen atoms are omitted for clarity other than those participating in hydrogen bonds and C-H $\cdots\pi$  interactions.

Again, the pattern of the C-H $\cdots\pi$  interaction belongs to type III.<sup>17</sup> The type and parameters of the C-H $\cdots\pi$  interaction in **2** are comparable to those of our early reports  $\{[\text{Ni}(\text{L1+H})(\text{CH}_3\text{CN})_2](\text{ClO}_4)_3$  (**L1** = 3,10-bis{3-(1-imidazolyl)propyl}-1,3,5,8,10,12-hexaazacyclotetradecane):<sup>9</sup> C-H $\cdots\pi$ :  $d(\text{H}\cdots\pi) = 2.952 \text{ \AA}$ ,  $d(\text{C}\cdots\pi) = 3.607 \text{ \AA}$ ,  $\angle(\text{CH}\pi) = 125.21^\circ$ ;  $\{[\text{Cu}(\text{L1+2H})\}_2(\mu\text{-tp})](\text{ClO}_4)_6 \cdot 2\text{H}_2\text{O}$  (tp = terephthalate):<sup>11</sup> C-H $\cdots\pi$ :  $d(\text{H}\cdots\pi) = 3.128 \text{ \AA}$ ,  $d(\text{C}\cdots\pi) = 3.739 \text{ \AA}$ ,  $\angle(\text{CH}\pi) = 121.24^\circ$ , C-H $\cdots\pi$ :  $d(\text{H}\cdots\pi) = 3.345 \text{ \AA}$ ,  $d(\text{C}\cdots\pi) = 3.961 \text{ \AA}$ ,  $\angle(\text{CH}\pi) = 122.05^\circ$ ;  $[\text{Cu}(\text{L1})(\text{CH}_3\text{CN})_2](\text{ClO}_4)_2$ :<sup>11</sup> C-H $\cdots\pi$ :  $d(\text{H}\cdots\pi) = 3.568 \text{ \AA}$ ,  $d(\text{C}\cdots\pi) = 4.272 \text{ \AA}$ ,  $\angle(\text{CH}\pi) = 130.81^\circ$ .

Consequently, it is believed that the introduction of imidazole pendants on the macrocycle by using imidazole containing primary amines as a padlock is a good strat-

egy for the formation of self-assembled macrocyclic nickel(II) supramolecules. In addition, the alkyl chain of a primary padlock amine plays a significant role in intermolecular C-H $\cdots\pi$  interactions together with imidazole rings.

Elemental analyses for complexes **1** and **2** agree well with the structures determined by X-ray diffraction methods, respectively. A weak band at  $3196 \text{ cm}^{-1}$  for **1** and  $3187, 3116 \text{ cm}^{-1}$  for **2** in the IR spectra were assigned to N-H stretchings of the macrocycle, respectively. The strong bands due to perchlorate ions were appeared at  $1090 \text{ cm}^{-1}$  ( $\nu_{\text{as}}\text{Cl-O}$ ) and  $625 \text{ cm}^{-1}$  ( $\delta\text{O-Cl-O}$ ) for **1** and  $1096 \text{ cm}^{-1}$  ( $\nu_{\text{as}}\text{Cl-O}$ ) and  $621 \text{ cm}^{-1}$  ( $\delta\text{O-Cl-O}$ ) for **2**. The shapes of the bands indicate that the perchlorate ions are not coordinated to nickel(II) ions in **1** and **2**.<sup>19</sup> Electronic spectra of **1**

and **2** in DMF solutions show band maxima at 450 nm which are typical for square-planar nickel(II) macrocycles.<sup>20,21</sup>

In summary, we self-assembled and structurally characterized two macrocyclic nickel(II) supramolecules **1** and **2**. In **1**, the macrocyclic nickel(II) unit extends its structure by N-H...N hydrogen bonds and C-H... $\pi$  intermolecular interactions to form a 2D supramolecule, whereas the C-H... $\pi$  interactions act as intermolecular forces to interconnect macrocyclic nickel(II) units, resulting in the formation of a 1D supramolecule in **2**.

## EXPERIMENTAL

### Materials and Measurements

All chemicals were commercially available from Aldrich and were used without further purification. Water was distilled before use for all procedures. IR spectra were recorded on a JASCO FT-IR-4000 spectrophotometer with Nujol mull (KBr discs) in the 4000-400  $\text{cm}^{-1}$ . Elemental analyses were performed on a VarioMICRO analyzer.

Caution! The perchlorate salts are potentially explosive and should be handled in small quantities.

### Synthesis of precursor complex [Ni(L+2H)](ClO<sub>4</sub>)<sub>4</sub>

The slightly modified literature method was adopted to prepare the precursor complex [Ni(L+2H)](ClO<sub>4</sub>)<sub>4</sub> by

using 1-(3-aminopropyl)-2-methyl-1*H*-imidazole instead of 4-(aminomethyl)pyridine or 1-(3-aminopropyl)imidazole.<sup>7-9</sup> Typical synthetic procedures are as follows. To a stirred methanol (40 mL) solution of Ni(OAc)<sub>2</sub>·4H<sub>2</sub>O (3.1 g, 12.5 mmole) were dropwise added ethylenediamine (1.7 mL, 25 mmole), paraformaldehyde (1.5 g, 50 mmole), and 1-(3-aminopropyl)-2-methyl-1*H*-imidazole (3.63 mL, 25 mmole). The mixture was refluxed for 24 hr, cooled and added 7 mL of 60% HClO<sub>4</sub> slowly, and stored in the refrigerator until yellow solid formed. The solid filtered, washed with methanol and dried in vacuo. Yield: ~10%. Anal. Calc. for [Ni(L+2H)](ClO<sub>4</sub>)<sub>4</sub> C<sub>22</sub>H<sub>44</sub>N<sub>10</sub>O<sub>16</sub>Cl<sub>4</sub>Ni: C, 29.20; H, 4.86; N, 15.47%. Found C, 29.53; H, 4.92; N, 15.38%. IR [Nujol,  $\text{cm}^{-1}$ ]: 3195 ( $\nu_{\text{NH}}$ ), 1105, 1068 ( $\nu_{\text{asCl-O}}$ ), 625 ( $\delta_{\text{O-Cl-O}}$ ).

### Syntheses and obtaining crystals of [Ni(L+H)](ClO<sub>4</sub>)<sub>3</sub> (**1**) and [Ni(L)](ClO<sub>4</sub>)<sub>2</sub>·2H<sub>2</sub>O (**2**)

To an CH<sub>3</sub>CN/H<sub>2</sub>O (~20 mL) suspension of the precursor complex [Ni(L+2H)](ClO<sub>4</sub>)<sub>4</sub> (1 g) was added an excess amount of triethylamine (~1 mL). The suspension turned clear immediately by the addition of triethylamine. The mixture was stored in the refrigerator until crystals formed. Suitable crystals for **1** (orange needles) and **2** (yellow plates) were manually harvested under a microscope for X-ray diffraction studies and subsequent spectroscopic measurements, respectively. Anal. Calc. for [Ni(L+H)](ClO<sub>4</sub>)<sub>3</sub> (**1**) C<sub>22</sub>H<sub>43</sub>Cl<sub>3</sub>N<sub>10</sub>NiO<sub>12</sub>: C, 32.83; H, 5.34; N, 17.40%. Found

**Table 1.** Crystal data and structure refinement for **1** and **2**

	1	2
Empirical formula	C <sub>22</sub> H <sub>43</sub> Cl <sub>3</sub> NiN <sub>10</sub> O <sub>12</sub>	C <sub>22</sub> H <sub>46</sub> Cl <sub>2</sub> NiN <sub>10</sub> O <sub>10</sub>
Formula weight	804.72	740.30
Temperature(K)	147(2)	147(2)
Wavelength(Å)	0.71073	0.71073
Crystal system	Triclinic	Triclinic
Space group	P-1	P-1
<i>a</i> (Å)	9.3358(13)	8.4728(10)
<i>b</i> (Å)	11.4175(15)	9.6582(11)
<i>c</i> (Å)	15.782(2)	11.0023(13)
$\alpha$ (°)	96.898(3)	72.716(3)
$\beta$ (°)	98.548(3)	74.918(2)
$\gamma$ (°)	101.836(3)	74.368(2)
<i>V</i> (Å <sup>3</sup> )	1608.8(4)	811.84(16)
<i>Z</i>	2	1
<i>D</i> <sub>calc</sub> (Mg/m <sup>3</sup> )	1.661	1.514
Absorption coefficient (mm <sup>-1</sup> )	0.929	0.829
Independent reflections	7286 [R(int) = 0.0252]	3712 [R(int) = 0.0320]
Goodness-of-fit on <i>F</i> <sup>2</sup>	1.042	1.074
Final R indices [ <i>I</i> > 2 $\sigma$ ( <i>I</i> )]	R <sub>1</sub> =0.0399, wR <sub>2</sub> =0.1043	R <sub>1</sub> =0.0335, wR <sub>2</sub> =0.0963
R indices (all data)	R <sub>1</sub> =0.0442, wR <sub>2</sub> =0.1080	R <sub>1</sub> =0.0374, wR <sub>2</sub> =0.0989

C, 32.90; H, 5.32; N, 17.43%. IR [Nujol,  $\text{cm}^{-1}$ ]: 3196 ( $\nu\text{NH}$ ), 1090 ( $\nu_{\text{asCl-O}}$ ), 625 ( $\delta\text{O-Cl-O}$ ). UV/vis [DMF;  $\lambda_{\text{max}}/\text{nm}$ ]: 550. Anal. Calc. for  $[\text{Ni}(\text{L})](\text{ClO}_4)_2 \cdot 2\text{H}_2\text{O}$  (**2**)  $\text{C}_{22}\text{H}_{46}\text{Cl}_2\text{N}_{10}\text{NiO}_{10}$ : C, 35.69; H, 6.21; N, 18.91%. Found C, 35.59; H, 6.29; N, 18.88%. IR [Nujol,  $\text{cm}^{-1}$ ]: 3565, 3328 ( $\nu\text{OH}$ ), 3187, 3116 ( $\nu\text{NH}$ ), 1096 ( $\nu_{\text{asCl-O}}$ ), 621 ( $\delta\text{O-Cl-O}$ ). UV/vis [DMF;  $\lambda_{\text{max}}/\text{nm}$ ]: 550.

### X-ray crystallography

Crystallographic data for **1** and **2** are summarized in Table 1. Bruker Kappa APEX-DUO CCD X-ray diffractometer with  $\text{Mo-K}\alpha$  radiation ( $\lambda = 0.71073 \text{ \AA}$ ) was used for data collection. To collect sufficient data, a combination of  $\phi$  and  $\omega$  scans with  $\kappa$  offsets were used. The data frames were integrated and scaled using the Denzo-SMN package.<sup>22</sup> The structure was solved and refined, using the SHELXTLPC V6.1 package.<sup>23</sup> Refinement was performed by full-matrix least squares on  $F^2$ , using all data (negative intensities included). Hydrogen atoms were included in calculated positions.

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**Supplementary Materials.** CCDC Nos. 2015621 (**1**) and 2015622 (**2**) contain the supplementary crystallographic data. These data can be obtained free of charge via [http://www.ccdc.cam.ac.uk/data\\_request/cif](http://www.ccdc.cam.ac.uk/data_request/cif).

### REFERENCES

- Piguet, C.; Bernardinelli, G.; Hopfgartner, G. *Chem. Rev.* **1997**, *97*, 2005.
- Leong, W. L.; Vittal, J. J.; *Chem. Rev.* **2011**, *111*, 688.
- Eddaoudi, M.; Moler, D. B.; Li, H.; Chen, B.; Reineke, T. M.; O'Keefe, M.; Yaghi, O. M. *Acc. Chem. Res.* **2001**, *34*, 319.
- Suh, M. P.; Cheon, Y. E.; Lee, E. Y. *Coord. Chem. Rev.* **2008**, *252*, 1007.
- Moulton, B.; Zaworotko, M. J. *Chem. Rev.* **2001**, *101*, 1629.
- Eryazici, I.; Moorefield, C. N.; Newkome, G. R. *Chem. Rev.* **2008**, *108*, 1834.
- Min, K. S.; Suh, M. P. *Chem. Eur. J.* **2001**, *7*, 303.
- Min, K. S.; Suh, M. P. *Eur. J. Inorg. Chem.* **2001**, 449.
- Han, S.; Kim, T.; Lough, A. J.; Kim, J. C. *Inorg. Chim. Acta* **2011**, *370*, 170.
- Han, S.; Lough, A. J.; Kim, J. C. *J. Coord. Chem.* **2010**, *63*, 2069.
- Kim, T.; Lough, A. J.; Kim, J. C. *Bull. Korean Chem. Soc.* **2013**, *34*, 1913.
- Misra, T. K.; Chung, C. -S.; Cheng, J.; Lu, T. -H. *Polyhedron*, **2001**, *20*, 3149.
- Kim, J. C.; Lough, A. J.; Kim, H. *Inorg. Chem. Commun.* **2002**, *5*, 771.
- Kim, J. C.; Fettingner, J. C.; Kim, Y. I. *Inorg. Chim. Acta* **1999**, *286*, 67.
- Hunter, T. M.; McNae, I. W.; Liang X.; Bella, J.; Parsons, S.; Walkinshaw, M. D.; Sadler, P. J. *Proc. Natl. Acad. Sci. USA* **2005**, *102*, 2288.
- Jang, C.; Kim, J. C. *J. Korean Chem. Soc.* **2014**, *58*, 234.
- Malone, J. F.; Murray, C. M.; Charlton, M. H.; Docherty, R.; Lavery, A. J. *J. Chem. Soc., Faraday Trans.* **1997**, *93*, 3429.
- Ferguson, G.; Gallagher, J. F.; Li, Y.; McKervey, M. A.; Madigan, E. M.; Malone, J. F.; Moran, M. B.; Walker, A. *Supramol. Chem.* **1996**, *7*, 223.
- Tasker, P. A.; Sklar, L. *J. Cryst. Mol. Struct.* **1975**, *5*, 329.
- Szalda D. J.; Fujita E.; Sanzenbacher, R; Paulus, H.; Elias, H. *Inorg. Chem.* **1994**, *33*, 5855.
- Kang, S.-G.; Kweon, J. K.; Jung, S.-K. *Bull. Korean Chem. Soc.* **1991**, *12*, 483.
- Otwinowski, Z.; Minor, W. *In Methods in Enzymology, Macromolecular Crystallography, Part A; Carter, C. W., Sweet, R. M. Eds, Academic Press: London* **1997**, 276, 307.
- Sheldrick, G. M. SHELXTL\PC V6.1, Bruker Analytical X-ray Systems, Madison, WI (**2001**).