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Title: Synthesis and Characterization of Zwitterionic Zn(II) and Cu(II) Coordination Compounds with Ring-substituted 2,2-Biimidazole Derivatives

Year: 2016

Version:

Please cite the original version:

Tatikonda, R., Kalenius, E., & Haukka, M. (2016). Synthesis and Characterization of Zwitterionic Zn(II) and Cu(II) Coordination Compounds with Ring-substituted 2,2-Biimidazole Derivatives. *Inorganica Chimica Acta*, 453, 298-304.
<https://doi.org/10.1016/j.ica.2016.08.015>

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Accepted Manuscript

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PII: S0020-1693(16)30467-4
DOI: <http://dx.doi.org/10.1016/j.ica.2016.08.015>
Reference: ICA 17209

To appear in: *Inorganica Chimica Acta*

Received Date: 29 March 2016
Revised Date: 16 July 2016
Accepted Date: 8 August 2016

Please cite this article as: R. Tatikonda, E. Kalenius, M. Haukka, Synthesis and Characterization of Zwitterionic Zn(II) and Cu(II) Coordination Compounds with Ring-substituted 2,2'-Biimidazole Derivatives, *Inorganica Chimica Acta* (2016), doi: <http://dx.doi.org/10.1016/j.ica.2016.08.015>

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Synthesis and Characterization of Zwitterionic Zn(II) and Cu(II) Coordination Compounds with Ring-substituted 2,2'-Biimidazole Derivatives

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Abstract

Zwitterionic coordination compounds with strongly asymmetrical charge distribution were synthesized and characterized. Ring-substituted biimidazoles were used as the primary ligands for Zn and Cu compounds. Formation of Zwitterionic coordination compound was found to be strongly dependent on the pH of the reaction medium as well as on the ring and nitrogen substituents of the ligand. Reaction of the Df-R₂biim (Df-R₂biim = 2,2'-bi-1R-imidazole-5,5'-dicarboxaldehyde, R= Me, Et or Pr) with ZnCl₂ in neutral conditions led to binuclear compounds [Zn₂Cl₄(Df-R₂biim)₂] with two bridging ligands (**1a-c**). Reaction with CuCl₂·2H₂O gave neutral mononuclear compound [CuCl₂(Df-Me₂biim)] (**1d**) with chelating biimidazole ligand. In acidic conditions (pH = 3–4) the imidazole nitrogens were either fully or partially protonated, which prevented the bidentate coordination of the ligands. Furthermore, aldehyde substituents of Df-R₂biim ligands were involved in acetal formation with methanol solvent. Under acidic conditions the primary products were either ion pairs [MCl₄][C₁₄H₂₄N₄O₄] (M=Zn **3a**, M=Cu **3b**) with fully protonated H₂L²⁺ counter cation or desired Zwitterionic coordination compounds [ZnCl₃(C₁₆H₂₇N₄O₄)] (**2a**), [ZnCl₃(C₁₈H₃₁N₄O₄)] (**2b**), [CuCl₃(C₁₆H₂₇N₄O₄)] (**2c**), or [CuCl₃(C₁₈H₃₁N₄O₄)] (**2d**) with partially protonated monodentate HL⁺ ligand. Zwitterions were obtained only with ligands having both aldehyde groups as ring-substituents and longer alkyl chains (ethyl or propyl) as *N*-substituents. In other cases ion pairs were formed as final products. Zwitterions were found to decompose in alcoholic solutions. In methanol solution, Zwitterionic compounds released neutral ligand and HZnCl₃. All main products were characterized by NMR, elemental analysis and single-crystal X-ray diffraction.

Keywords.

Zinc • Copper • Biimidazole • Crystal structures • Zwitterionic and Non-coordinated compounds

1. Introduction.

Imidazole is an important aromatic N-heterocyclic compound because of its significant role in biosystems and its versatile coordination ability.^[1] The 2,2'-biimidazole molecule, H₂biim, is one of the most useful dimeric analogue of imidazole and well known ligand for transition metals. Compounds containing H₂biim moiety have been the focus of several investigations due to their biological and catalytic activity.^[2] Anti-protozoal,^[2d] anti-tuberculosis,^[2e] anti-cancer,^[2f] and cardiotoxic,^[2g] properties are examples of pharmacological functionalities of biimidazole based compounds. The conjugate polymers of imidazole and biimidazole containing metal compounds exhibit outstanding sensing properties to metal ions, anions, nitric oxide and amino acids.^[3] In H₂biim, two imidazole rings are capable to adopt coplanar orientation when the ligand reacts with a metal cation. This favors chelation as the primary coordination mode.^[4] However, few exceptions, where H₂biim behaves as a bridging ligand in binuclear and polynuclear compounds, are also known.^[5] If the N-H hydrogens are replaced with bulkier groups, chelation is less likely. Therefore, *N*-substituted R₂biim moieties are more commonly acting as bridging ligands in binuclear,^[6] oligonuclear or polynuclear^[7] compounds. So far, only few literature precedence exist of mononuclear compounds where 1,1'-dimethyl-2,2'-biimidazole, Me₂biim involved in chelation with the metal ion.^[8]

Coordination modes of unsubstituted biimidazole vary from monodentate to tetradentate depending on the reaction conditions and level of protonation (Fig. 1). In neutral medium the most common coordination mode of H₂biim is bidentate chelating bonding (Fig. 1a).^[9] In this mode the protonated nitrogens are available for H-bonding as H-bond donors.^[9] Removal of N-H protons produces typically tridentate (mono-deprotonated biimidazole, Hbiim⁻),^[10] or tetradentate (fully deprotonated biimidazole, biim²⁻)^[11] ligands (Fig. 1b,c). Basic medium usually favors these two coordination modes. In acidic medium biimidazole can be protonated to H₃biim⁺ or H₄biim²⁺. The cationic H₃biim⁺ (Fig. 1d) favors monodentate coordination. In H₄biim²⁺ (Fig. 1e) there are no available nitrogens for coordination and therefore H₄biim²⁺ is usually acting as counter cation in ionic systems.

Figure 1. Coordination modes of H₂biim (a) chelating (b) chelating-bridging (c) monodentate and ionic

The ring substituents (Fig. 2) have strong impact on physical and chemical properties of the biimidazole molecule. Ring substitution opens up a route for fine-tuning the properties of the ligand. However, only limited number of such biimidazole derivatives have been synthesized and structurally characterized and even fewer of those have been used as ligands in metal compounds.^[14,15]

R = Hydrogen or alkyl
R' = Nitro, Cyano, Aldehyde or Alkyl group

Figure 2. Ring -substituted (R') and *N*-substituted (R) 2,2'-Biimidazole.

This paper describes the synthesis and structural characterization of a series of zinc and copper coordination compounds with ring-substituted Df-R₂biim (2,2'-bi-1R-imidazole-5,5'-dicarboxaldehyde, R= Me, Et or Pr) derivatives. The goal was to use different ring substituents and the level of protonation as tools for preparation of polar and especially Zwitterionic coordination compounds. The products were characterized by NMR, IR, elemental analysis, molar conductivity and single crystal X-ray crystallography. The potential reaction route to Zwitterionic products is also briefly discussed.

2. Experimental Section

2.1 General Considerations. R₂biim and ring-substituted 2,2'-biimidazole derivatives, 2,2'-bi-1alkyl-imidazole-5,5'-dicarboxaldehyde (Df-R₂biim) were synthesized by following literature methods (see ESI).^[16,17] All other chemicals were obtained from commercial source. The NMR spectra (¹H and ¹³C) of ligands were recorded on Bruker Avance DRX 400 NMR spectrometer and chemical shifts were expressed in ppm. Elemental analyses were performed on VarioEL elemental analyzer. Molar conductivity experiments were performed by using handylab LF11 manufactured by Scott.

2.2. Synthesis

2.2.1. Synthesis of Binuclear Zn(II) and Mononuclear Cu(II) coordination compounds with Df-R₂biim (R= Me, Et or Pr) (1a-f)

A mixture of metal chloride (ZnCl₂ or CuCl₂·2H₂O) and Df-R₂biim (1:1) in methanol was stirred for 30 min to produce binuclear Zn(II) (white precipitate) and mononuclear Cu(II) compounds (blue precipitate). The compound was collected through filtration, washed with cold methanol and dried under vacuum. X-ray quality crystals were obtained with solvent diffusion process of methanolic metal solution to the ligand solution in chloroform. The yield was calculated based on metal salt.

[Zn₂Cl₄(C₁₀H₁₀N₄O₂)₂] **1a**. Yield: 82%. ¹H NMR (δ, 400 MHz, DMSO-D₆): 4.18 (s, 12H), 8.10 (s, 4H), 9.86 (s, 4H). ¹³C NMR (δ, 100 MHz, DMSO-D₆): 34.35, 132.63, 141.94, 142.02, 180.96. Anal. Calcd for C₂₀H₂₀N₈O₄Zn₂Cl₄·CH₃OH (741.038): C, 34.04; H, 3.26; N, 15.12. Found: C, 33.84; H, 3.34; N, 14.80. IR (ν, cm⁻¹): 1686 (CO).

[Zn₂Cl₄(C₁₂H₁₄N₄O₂)₂] 1b. Yield: 80%. ¹H NMR (δ, 400 MHz, DMSO-D₆): 1.33 (t, 12H), 4.69 (q, 8H), 8.13 (s, 4H), 9.84 (s, 4H). ¹³C NMR (δ, 100 MHz, DMSO-D₆): 16.12, 41.81, 131.99, 141.30, 143.08, 180.53. Anal. Calcd for C₂₄H₂₈N₈O₄Zn₂Cl₄ (765.102): C, 37.68; H, 3.69; N, 14.65. Found: C, 37.22; H, 3.76; N, 14.27. IR (ν, cm⁻¹): 1686 (CO).

[Zn₂Cl₄(C₁₄H₁₈N₄O₂)₂] 1c. Yield: 87%. ¹H NMR (δ, 400 MHz, DMSO-D₆): 0.79 (t, 12H), 1.66-1.75 (m, 8H), 4.69 (t, 8H), 8.14 (s, 4H), 9.84 (s, 4H). ¹³C NMR (δ, 100 MHz, DMSO-D₆): 10.58, 23.75, 47.46, 132.22, 141.56, 142.95, 180.64. Anal. Calcd for C₂₈H₃₆N₈O₄Zn₂Cl₄ (821.209): C, 40.95; H, 4.42; N, 13.64. Found: C, 40.68; H, 4.38; N, 13.34. IR (ν, cm⁻¹): 1682 (CO)

[CuCl₂(C₁₀H₁₀N₄O₂)] 1d. Yield: 74%. Anal. Calcd for C₁₀H₁₀N₄O₂CuCl₂ (352.664): C, 34.06; H, 2.86; N, 15.89. Found: C, 34.12; H, 2.92; N, 16.02. IR (ν, cm⁻¹): 1686 (CO).

[CuCl₂(C₁₂H₁₄N₄O₂)] 1e. Yield: 78%. Anal. Calcd for C₁₂H₁₄N₄O₂CuCl₂ (380.717): C, 37.86; H, 3.71; N, 14.72. Found: C, 37.56; H, 3.66; N, 14.44. IR (ν, cm⁻¹): 1687 (CO).

[CuCl₂(C₁₄H₁₈N₄O₂)] 1f. Yield: 77%. Anal. Calcd for C₁₄H₁₈N₄O₂CuCl₂ (408.770): C, 41.14; H, 4.44; N, 13.71. Found: C, 40.80; H, 4.42; N, 13.86. IR (ν, cm⁻¹): 1686 (CO).

2.2.2. Synthesis of Zwitterionic coordination compounds 2a-d

A binuclear (**1b** or **1c**, 0.1 mmol) or mononuclear (**1e** or **1f**, 0.1 mmol) coordination compounds were dissolved in 2 mL of methanol by adding 5-6 drops of Conc. HCl (pH = 3-4). The resulting solution was filtered off and the final crystalline product was obtained by slow evaporation of the solvent. After couple of days, colorless crystals for Zn²⁺ or blue colored crystals for Cu²⁺ were formed in filtrate at room temperature. During the reaction aldehyde groups from the ligand was involved in acetal formation in presence of methanolic HCl solutions. The yield was calculated based on metal salt.

[ZnCl₃(C₁₆H₂₇N₄O₄)] (2a). Yield: 62%. Anal. Calcd for C₁₆H₂₇N₄O₄ZnCl₃.CH₃OH (543.191): C, 37.59; H, 5.75; N, 10.31. Found: C, 37.36; H, 5.49; N, 10.45.

[ZnCl₃(C₁₈H₃₁N₄O₄)] (2b). Yield: 60%. Anal. Calcd for C₁₈H₃₁N₄O₄ZnCl₃.CH₃OH (571.244): C, 39.95; H, 6.18; N, 9.81. Found: C, 39.69; H, 5.96; N, 9.88.

[CuCl₃(C₁₆H₂₇N₄O₄)] (2c). Yield: 58%. Anal. Calcd for C₁₆H₂₇N₄O₄CuCl₃ (509.32): C, 37.73; H, 5.34; N, 11.00. Found: C, 37.62; H, 5.39; N, 11.02.

[CuCl₃(C₁₈H₃₁N₄O₄)] (2d). Yield: 57%. Anal. Calcd for C₁₈H₃₁N₄O₄CuCl₃ (537.37): C, 40.23; H, 5.81; N, 10.43. Found: C, 40.42; H, 5.92; N, 10.39.

2b1 is hydrolyzed product of **2b**, where acetal groups are hydrolyzed back to aldehyde. Compound **5** was synthesized following the same procedure but using HBr instead of HCl. Compound **2b1** and **5** was characterized only by single-crystal X-ray crystallography (For single crystal x-ray structure see ESI).

2.2.3. Synthesis of Cu(II) and Zn(II) ion pairs with Df-Me₂biim (3a)

Syntheses of ionic compounds were carried out by dissolving binuclear [Zn₂Cl₄(Df-Me₂Biim)₂] (**1a**, 0.1 mmol) or mononuclear [CuCl₂(Df-Me₂Biim)] (**1d**, 1 mmol) in 2 mL of methanol. The solution was made acidic by adding 5-6 drops of Conc. HCl. After couple of days, colorless or blue crystals were obtained in filtrate. The aldehyde group from the ligand was involved in acetal formation.

[ZnCl₄][C₁₄H₂₄N₄O₄] (**3a**). Yield: 64%. Anal. Calcd for C₁₄H₂₄N₄O₄ZnCl₄: C, 32.36; H, 4.66; N, 10.78. Found: C, 32.99; H, 4.72; N, 11.85.

[CuCl₄][C₁₄H₂₄N₄O₄] (**3b**). Yield: 54%. Anal. Calcd for C₁₄H₂₄N₄O₄CuCl₄: C, 32.48; H, 4.67; N, 10.82. Found: C, 33.10; H, 4.80; N, 10.98.

Ion pairs ([ZnCl₄]²⁻[C₁₀H₁₆N₄]²⁺: **4a** and [ZnCl₄]²⁻[C₁₂H₂₀N₄]²⁺: **4b**) were synthesized from the reaction of ZnCl₂ with R₂Biim (R = Et or Pr) in acidic medium (pH = 3–4). These compounds were characterized only by single crystal X-ray crystallography (see ESI).

2.3. X-ray Crystallography

X-ray Structure Determination. The crystals of **1-5** were immersed in cryo-oil, mounted in a MiTeGen loop and measured at 120-123 K. The X-ray diffraction data were collected on an Agilent Technologies Supernova or a Bruker AXS Kappa Apex diffractometer using Mo K α or Cu K α radiation. The *CrysAlisPro*^[18] or *Denzo-Scalepack*^[19] program packages were used for cell refinements and data reductions. Structures were solved by charge flipping method using *SUPERFLIP*^[20] program or by direct methods using *SHELXS-2014*^[21] program. An analytical, Gaussian, or multi-scan (*CrysAlisPro*^[18], *SADABS*^[22]) absorption correction was applied to all data. Structural refinements were carried out using *SHELXL-2014*^[21] or *SHELXL-97*^[21] software. In structure **2b1**, carbon atoms of some of the alkyl substituents (C9-C14) were slightly disordered but no disorder model was used in the final structure refinement. In **4a**, one of the ethyl substituents (C9C, C10C/C9D C10D) was disordered over two sites with occupancy ratio of 0.48/0.52. The carbon atoms of these disordered groups were restrained with effective standard deviation 0.01 so that their U_{ij} components approximate to isotropic behavior. In structure **5**, two of alkoxy groups (C13-C15B) were disordered over two sites with occupancy ratio of 0.57/0.43. Also, Br2 atom was slightly disordered but no disorder model was used for the final structure refinement. Hydrogen atoms were either placed in calculated positions or located from the difference Fourier map. In all cases hydrogen atoms were refined using riding model. The crystallographic details of **1b**, **1d**, **2a**, **2b**, **2c**, **3a**, and **3b** are summarized in Table S1. Crystallographic details of all other structures (**4** and **5**) are given in Table S3. Graph sets for intermolecular H-bonding were analyzed by using Mercury.^[23]

3. Results and discussion

A series of ring-substituted 2,2'-biimidazole derivatives were synthesized by adopting literature procedure.^[17] The ligand synthesis method was slightly modified by increasing the reaction time and molar amounts of reagents used to improve the yields and selectivity towards 2,2'-Bi-1R-imidazole -5,5'-dicarboxaldehydes (Df-R₂biim) (see ESI). The ligands (Df-R₂biim) synthesis involves direct lithiation followed by subsequent formylation with DMF in THF solution.

Reaction of Df-R₂biim with ZnCl₂ and CuCl₂·2H₂O in methanolic solution produced neutral binuclear Zn(II) (**1a-c**)^[23] and mononuclear Cu(II) (**1d**) compounds (Scheme 1). Reaction of Df-R₂biim with Zn²⁺ led to binuclear compounds with bridging Df-R₂biim ligands linking two tetrahedral Zn centers together. With Cu²⁺ Df-Me₂Biim formed mononuclear square planar structure, where the imidazole derivative was chelating (**1d**). Unlike in structures **1a-1c**, the imidazole rings in the chelating **1d** structure were nearly coplanar.

Reaction of Cu²⁺ with ethyl and propyl substituted ligands gave solid products but no X-ray quality crystals could be obtained. These products were characterized only by elemental analysis. Because of the lack of crystal structures, detailed geometries of products **1e** and **1f** could not be confirmed. Based on literature and obtained results, it can be assumed that the structure of **1e** and **1f** could resemble either mononuclear copper compound **1d** or a binuclear zinc compound **1a-c**.^[6, 24]

Scheme 1. Synthesis of Zn(II) and Cu(II) compounds in neutral and acidic medium.

When the compounds **1a-1f** were redissolved in methanol by adding few drops of concentrated HCl (pH = 3–4), either one or both metal-nitrogen bonds were broken and protonated. This led to two types of new products i.e. Zwitterionic (**2a-d**) coordination compounds (Scheme 1) or ion pairs of the type $[\text{MCl}_4]^{2-}$ $[\text{C}_{14}\text{H}_{24}\text{N}_4\text{O}_4]^{2+}$ (**3a,b**) or $[\text{MCl}_4]^{2-} [\text{H}_2\text{R}_2\text{biim}]^{2+}$ (**4a,b**) depending on the substituents of the biimidazole

ligand. Formation of Zwitterionic coordination compounds required low enough pH (3-4) to protonate one of the biimidazole nitrogens. In addition, the biimidazole ligand had to have aldehyde group as ring-substituents and longer alkyl chain (ethy or propyl) as *N*-substituents. The aldehyde groups were involved in acetal formation during the reaction and this was found to be crucial step to obtain Zwitterionic coordination compounds even if the detailed mechanism is not clear. On the other hand, aldehyde group alone was not enough to get Zwitterionic compounds (**3a,b**). The ligand also had to have longer chain alkyl groups as ring-substituents (Fig. 3).

Figure 3. Ligand required for the Zwitterion formation (R = Et, Pr).

If either aldehyde group or longer chain alkyl group was missing, only ion pairs were obtained (**3a,b** and **4a,b**). Interestingly, the acetal formation was found to be reversible. When the acetal containing Zwitterion compound **2b** was recrystallized from acetonitrile, the acetal groups were hydrolyzed back to aldehyde (**2b1**) due to residual water of the solvent.

The reaction of ZnCl₂ with Df-Et₂Biim was also performed in methanolic HBr solution. In this reaction, the formation of the Zwitterionic compound was accompanied by exchange of Cl ligands with Br (**5**).

Molar conductivity.

The stability of the Zwitterions in solution was studied by dissolving compounds **2a** and **2b** in methanol and measuring the conductivity of the solutions ($5 \cdot 10^{-4}$ M). The conductivities of the solutions of **2a** and **2b** were found to be 52.4 and 52.7 $\mu\text{S}/\text{cm}$ at 21°C respectively. Such values fall into the range of typical 1:1 electrolytes (the molar conductivity of $5 \cdot 10^{-4}$ M KCl solution in methanol was found as 52.7 $\mu\text{S}/\text{cm}$ at 21°C) indicating that the Zwitterions were decomposed. It is expected that the proton of the biimidazole is migrated to Zn, which results in release of neutral biimidazole compounds with acetal substituents and HZnCl₃ salt (Scheme 3).

Scheme 2. Decomposition of the Zwitterionic compound in methanolic solution.

The proposed decomposition route was further supported by $^1\text{H-NMR}$ and mass spectrometric analysis. In $^1\text{H-NMR}$ only one ring proton signal was observed at δ 7.65 ppm. Mass spectra confirmed the decomposition ($[\text{C}_{16}\text{H}_{27}\text{N}_4\text{O}_4]^+$ at m/z_{exp} 339.1892, m/z_{theor} 339.2027; $[\text{C}_{18}\text{H}_{31}\text{N}_4\text{O}_4]^+$ at m/z_{exp} 367.2305, m/z_{theor} 339.234 and $[\text{ZnCl}_3]^-$ at m/z_{exp} 168.7905, m/z_{theor} 168.8363).

3.1. Crystal Structures

The molecular structures with the numbering schemes are shown in Figures 4-7. Crystallographic data and selected bond distances, angles are summarized in the supplementary Table S1, S2 and S3.

$[\text{Zn}_2\text{Cl}_4(\text{Df-Et}_2\text{Biim})_2]$ (1b**).** As in the case of known $[\text{Zn}_2\text{Cl}_4(\text{Me}_2\text{Biim})_2]$ structure,^[24] the zinc atoms in **1b** are centrosymmetrically double-bridged by two biimidazolic nitrogens to form a binuclear compound. The planes of the imidazole rings of the ligands are nearly perpendicular with the angle of 85.67° . In **1b**, the typical tetrahedral coordination sphere of the Zn atoms are completed by two chlorido ligands. The Zn–N [$\text{Zn1-N2} = 2.064(5)$ Å, $\text{Zn1-N4} = 2.053(2)$ Å], Zn–Cl [$\text{Zn-Cl1} = 2.219(3)$ Å, $\text{Zn1-Cl2} = 2.207(5)$ Å] distances indicate slight asymmetry in the structure and Zn–N bond distances are almost identical to the $[\text{Zn}_2\text{Cl}_4(\text{Me}_2\text{Biim})_2]$ structure. The Zn1–N4 bond length ($2.053(2)$ Å) remains practically unchanged during conversion from the binuclear **1b** to Zwitterionic **2a** and Zn1–N2 bond ($2.064(5)$ Å) opens up allowing a third chloride ligand entering the coordination sphere of Zn.

$[\text{CuCl}_2(\text{Df-Me}_2\text{Biim})]$ (1d**).** The copper atom in this molecule is tetra-coordinated with square planar geometry and surrounded by two chlorido ligands and two nitrogen atoms from the chelating ligand. The bite angle [N2-Cu1-N2] of the chelating biimidazole ligand is 78.96° and the imidazole rings of the ligand are almost coplanar. The angle between the planes of the rings is only about 5.9° . The Cu–N distances ($1.988(2)$ and $2.2261(6)$ Å) are in the range of typical Cu–N_{im} ($1.94 - 2.03$ Å) distances.^[6, 26]

Figure 4. Molecular structure of **1b** (Left) and **1d** (right). Thermal ellipsoids are drawn at the 50% probability level.

[ZnCl₃(C₁₆H₂₇N₄O₄)] (2a). The coordination geometry around zinc atom is slightly distorted tetrahedron consisting of three chlorido ligands and one nitrogen atom from the singly protonated biimidazole ligand, [C₁₆H₂₇N₄O₄]⁺. The Zn–Cl bond lengths [Zn1–Cl1 = 2.242(3) Å, Zn1–Cl2 = 2.260(4) Å and Zn1–Cl3 = 2.270(6) Å] show again slight asymmetry of the structure. The angle between the planes of the two-imidazole rings is about 89.3°. In 2a, the intermolecular H-bonds are observed between two Zwitterionic molecules with the distance of Cl3...N4ⁱ = 3.144(2) Å (i = 1–X, 2–Y, 1–Z), and the angle of N–H...Cl = 154.5° in R²₂(14) synthon. The N...Cl distance falls in the range of typical N–H...Cl hydrogen bonds (2.91–3.62 Å).^[9d, 26a, 27]

[ZnCl₃(C₁₈H₃₁N₄O₄)] (2b). The coordination sphere of Zn consists of three chlorido ligands and one nitrogen atom from the biimidazole ligand [C₁₈H₃₁N₄O₄]⁺ as in 2a. The structure contains methanol of crystallization, which is hydrogen bonded to the protonated nitrogen of the imidazole ligand (N...O distance: 2.682 Å). The angle between the planes of imidazole rings is somewhat smaller (78.6°) than in 2a, but the imidazole rings are again clearly not co-planar. During crystallization process, methanol of crystallization was trapped into the crystal voids by H-bonds. The N–H and Cl groups of the neighboring compounds were involved in intermolecular H-bonding with methanol [N...O = 2.682(2) and O...Cl = 3.229(2) Å] and the angles of N–H...O and O–H...Cl are 164.9° and 157.9° respectively. Methanol solvent was acting as H-bonding acceptor in D¹₁ (2)a, donor in D¹₁ (2)b and as both acceptor and donor in R⁴₄(18) synthons.

Figure 5. Molecular structure and intermolecular H-bonding in structure **2a** (left) in $R^2_2(14)$ and **2b** (right) in $R^4_4(18)$ synthons. C-H hydrogens are omitted for clarity (50% probability ellipsoids).

[CuCl₃(C₁₆H₂₇N₄O₄)] (2c). The copper atom in this distorted tetrahedral molecule is surrounded by three chlorido ligands and one nitrogen from the imidazole ligand, [C₁₆H₂₇N₄O₄]⁺. The Cu–N2 distance (2.002(4) Å) is shorter than the corresponding Zn–N2 bond length, which is in the range of typical Cu–N distances (1.94–2.002 Å). The angle between the planes of the two imidazole rings is about 79.2°. Similarly to **2a**, intermolecular H-bonds in **2c**, are observed between two Zwitterionic molecules with the distance of Cl2...N4ⁱ = 3.141(6) Å (i = 2–X, 1–Y, –Z), and the angle of N–H...Cl = 158.9° in $R^2_2(14)$ synthon.

Figure 6. Molecular structure and intermolecular H-bonding in structure **2c** in $R^2_2(14)$ synthon. C-H hydrogens are omitted for clarity (50% probability ellipsoids).

No X-Ray quality crystals could be obtained from ([CuCl₃(C₁₈H₃₁N₄O₄))] **2d**.

[MCl₄][C₁₄H₂₄N₄O₄] (M=Zn, **3a**; M=Cu, **3b**). The ion pairs of **3a** and **3b** consist of anionic metal compounds ([CuCl₄]²⁻ and [ZnCl₄]²⁻) and fully protonated cationic [C₁₄H₂₄N₄O₄]²⁺. In **3a** and **3b**, intermolecular H-bonding was seen between cation and anionic part of the compound (R²₂(9) synthon). The distance between Cl2–N1ⁱ in **3a** is 3.172(2) Å (i= 1–X, +Y, 3/2–Z) and the corresponding distance Cl1–N2ⁱⁱ in **3b** 3.218(3) Å (ii= –X, +Y, 3/2–Z). In both structures, the imidazole rings of the cations are again not coplanar. The angles between the planes of imidazole rings are 69.3° and 56.9° for **3a** and **3b** respectively.

Figure 7. Molecular structure and H-bonding of **3a** (Zn, right) and **3b** (Cu, left). C-H hydrogens are omitted for clarity. Thermal ellipsoids are drawn at the 50% probability level. The distance of Cl2···N1ⁱ in **3a** is 3.172(2) Å (i= 1–X, +Y, 3/2–Z) and in **3b** is 3.218(3) Å (Cl1···N2ⁱⁱ; ii= –X, +Y, 3/2–Z).

Conclusions

A series of ring-substituted biimidazoles (Df-R₂biim; R=Me, Et or Pr) were used for the synthesis of binuclear Zn(II) and mononuclear Cu(II) compounds. Compound formation was found to depend strongly on the pH of the reaction medium. In neutral reaction conditions the biimidazole ligands were bidentate and acting as bridging ligands in binuclear Zn(II) compounds or chelating ligands in mononuclear Cu(II) compound. Lowering the pH to 3-4 led to partial or full protonation of the two N sites of the ligand. With fully protonated ligands only ion pairs (**3a,b**) were obtained while reactions with partially protonated biimidazoles gave Zwitterionic compounds (**2a-d**). In addition to pH, the formation of the Zwitterionic products was found to be dependent on the ring and N-substituents of the biimidazoles. Both of these substituents were needed for Zwitterion formation. In acidic media, the aldehyde substituent on the ring was involved in acetal formation and this process was a crucial step in obtaining Zwitterions. Longer alkyl N-substituents (ethyl or propyl) were also required for Zwitterion formation while methyl substituent favored ion pairs regardless of aldehyde substituent. In alcoholic solution the Zwitterionic compounds were decomposed to form free neutral ligand and HZnCl₃ salt.

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Electronic Supporting Information**for****Synthesis and Characterization of Zwitterionic Zn(II) and Cu(II) Coordination compounds with Ring-substituted 2,2'-Biimidazole Derivatives**

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*Department of Chemistry, University of Jyväskylä, P.O.Box 35, FI-40014 University of Jyväskylä, Finland*E-mail: matti.o.haukka@jyu.fi**1. Experimental Section**

The ligand synthesis method was slightly modified from the literature procedure^[17] by increasing the reaction time to ensure the completion of reaction. The molar amounts of reagents were optimized to improve the yield and selectivity towards 2,2'-Bi-1R-imidazole-5,5'-dicarboxaldehydes (Df-R₂biim). The reaction time was extended from 1h. to overnight and molar amounts of n-BuLi and DMF were taken as 5 times to the R₂biim molar amount.

General procedure for 2,2'-Bi-1R-imidazole-5,5'-dicarboxaldehyde (Df-R₂biim; R= Me, Et and Pr)

Under argon gas, a mixture of N,N'-disubstituted-2,2'-biimidazole (R₂biim, 2 mmol), 0.93 mL (6 mmol) of TMEDA and 20 mL of dried THF was cooled to -40°C. To the above cooled solution, 2.5 M n-BuLi in hexane (10 mmol) was added slowly with the help of syringe. After 15 min stirring at room temperature, a mixture of 0.76 mL (10 mmol) of DMF in 1 mL of THF was added drop wise over a period of 10 min. After complete addition of DMF, the reaction mixture was allowed for stirring overnight at room temperature. After the reaction time, the reaction mixture was diluted with 1N HCl, and extracted with ethyl acetate. The solvent was evaporated to get solid product. The product was recrystallized from ethanol.

2,2'-Bi-1methyl-imidazole-5,5'-dicarboxaldehyde (Df-Me2Biim)

1a was obtained by the general procedure as a pale yellow solid with 41% of yield. mp 159.2-162.4 °C. ¹H NMR (400 MHz CDCl₃): δ 4.34 (s, 6H), 7.89 (s, 2H), 9.86 (s, 2H). ¹³C NMR (100 MHz CDCl₃): δ 35.17, 133.00, 142.64, 142.87, 179.63. HRMS (*m/z*) Calcd for C₁₀H₁₁N₄O₂⁺ [M+H⁺]: 219.0882, found: 219.0873. IR (ν, cm⁻¹): 1667 (CHO).

2,2'-Bi-1ethyl-imidazole-5,5'-dicarboxaldehyde (Df-Et2Biim)

1b was obtained by the general procedure with 61% of yield. mp 146.5-149.8 °C. ¹H NMR (400 MHz CDCl₃): δ 1.41 (t, J = 7.0 Hz, 6H), 4.87 (q, J = 7.0 Hz, 4H), 7.88 (s, 2H), 9.82 (s, 2H). ¹³C NMR (100 MHz CDCl₃): δ 16.40, 42.63, 132.32, 142.22, 143.41, 179.15. HRMS (*m/z*) Calcd for C₁₂H₁₅N₄O₂⁺ [M+H⁺]: 247.1195, found: 247.1185. IR (ν, cm⁻¹): 1664 (CHO).

2,2'-Bi-1propyl-imidazole-5,5'-dicarboxaldehyde (Df-Pr2Biim)

1c was obtained by the general procedure with 72% of yield. mp 126.4-128.7 °C ¹H NMR (400 MHz CDCl₃): δ 1.41 (t, J = 7.4 Hz, 6H), 1.72-1.81 (m, 4H), 4.80 (t, J = 7.5 Hz, 4H), 7.87 (s, 2H), 9.82 (s, 2H). ¹³C NMR (100 MHz CDCl₃): δ 10.93, 24.45, 48.40, 132.52, 142.57, 143.31, 179.22. HRMS (*m/z*) Calcd for C₁₄H₁₉N₄O₂⁺ [M+H⁺]: 275.1508, found: 275.1506. IR (ν, cm⁻¹): 1662 (CHO).

2. X-ray Crystallography**Table S1.** Crystallographic Data for Compounds 1-3.

	1b	1d	2a	2b	2c	3a	3b
CCDC No.	1443455	1443456	1443457	1443458	1443461	1443462	1443459
Formula	C ₂₄ H ₂₈ N ₈ O ₄ Zn ₂ Cl ₄	C ₁₀ H ₁₀ N ₄ O ₂ CuCl ₂	C ₁₆ H ₂₇ N ₄ O ₄ ZnCl ₃	C ₁₈ H ₃₁ N ₄ O ₄ ZnCl ₃ , CH ₄ O	C ₁₆ H ₂₇ N ₄ O ₄ CuCl ₃	C ₁₄ H ₂₄ N ₄ O ₄ , ZnCl ₄	C ₁₄ H ₂₄ N ₄ O ₄ , CuCl ₄
Fw	765.08	352.66	511.13	571.23	509.30	519.54	517.71
Temp (K)	123	123	120	123	123	123	123
Cryst syst	Monoclinic	Monoclinic	Triclinic	Triclinic	Triclinic	Orthorhombic	Monoclinic
Space group	<i>P</i> 2 ₁ / <i>c</i>	<i>C</i> 2/ <i>c</i>	<i>P</i> $\bar{1}$	<i>P</i> $\bar{1}$	<i>P</i> $\bar{1}$	<i>P</i> bcn	<i>C</i> 2/ <i>c</i>
<i>a</i> (Å)	9.4812(2)	19.8441(1)	9.6893(2)	8.8362(4)	9.6212(18)	20.2149(6)	21.8044(18)
<i>b</i> (Å)	13.1493(3)	10.0122(4)	11.2715(4)	12.3593(9)	11.287(2)	8.9789(2)	7.8148(5)
<i>c</i> (Å)	13.5024(3)	6.9417(3)	11.6686(4)	13.1451(7)	11.8379(19)	11.5719(3)	12.4029(8)
α (°)	90	90.00	68.9230(1)	66.645(6)	67.948(16)	90.00	90.00
β (°)	108.213(2)	112.746(6)	69.619(2)	86.297(4)	68.641(16)	90.00	106.338(8)
γ (°)	90	90.00	81.483(2)	85.264(5)	77.051(15)	90.00	90.00
<i>V</i> (Å ³)	1599.02(6)	1271.94(11)	1114.28(6)	1312.66(13)	1104.0(4)	2100.41(1)	2028.1(3)
<i>Z</i>	2	4	2	2	2	4	4
<i>d</i> _{calc} (g/cm ³)	1.589	1.842	1.523	1.445	1.532	1.643	1.696
μ (mm ⁻¹)	5.293	6.339	1.490	1.276	4.992	6.557	6.629
Ref. collected	6291	2076	15345	10339	6759	5298	3613
Ind. reflections	3011	1219	5713	5435	4203	2148	1912
<i>F</i> (000)	776	708	528	596	526	1064	1060
GOF	1.079	1.059	1.107	1.033	0.960	1.062	1.060
<i>R</i> _{int}	0.0215	0.0377	0.0322	0.0149	0.0716	0.0215	0.0558
<i>R</i> ^a (<i>I</i> ≥ 2σ)	0.0320	0.0328	0.0366	0.0259	0.0578	0.0315	0.0467
w <i>R</i> ^b (<i>I</i> ≥ 2σ)	0.1170	0.0703	0.0695	0.0613	0.1187	0.0850	0.1210

$$^a R I = \sum |F_o| - |F_c| / \sum |F_o|. \quad ^b wR2 = [\sum (w(F_o^2 - F_c^2)^2) / \sum (w(F_o^2)^2)]^{1/2}.$$

Table S2. Selected bond lengths (Å) and angles (°) for Compounds **1-3**.

Compound 1b							
Zn1–Cl1	2.2193(6)	Zn1–Cl2	2.2072(5)	Zn1–N2	2.0637(2)	Zn1–N4	2.0535(2)
Cl1–Zn1–Cl2	123.87(3)	N4–Zn1–Cl1	104.22(5)	N4–Zn1–Cl2	111.10(5)	N2–Zn1–Cl1	102.39(5)
N2–Zn1–Cl2	110.83(5)	N4–Zn1–N2	102.04(7)				
Compound 1d							
Cu1–Cl1	2.2261(6)	Cu1–N2	1.988(2)				
Cl1 ¹ –Cu1–Cl1	96.45(3)	N2–Cu1–Cl1	169.78(6)	N2–Cu1–Cl1	92.52(6)	N2–Cu1–N2 ¹	79.01(12)
Compound 2a							
Zn1–Cl1	2.2422(6)	Zn1–Cl2	2.2603 (6)	Zn1–Cl3	2.2705(6)	Zn1–N2	2.0541(17)
Cl1–Zn1–Cl2	119.33(2)	Cl1–Zn1–Cl3	112.99(2)	Cl2–Zn1–Cl3	107.06(2)	N2–Zn1–Cl1	103.94(5)
N2–Zn1–Cl2	102.31(5)	N2–Zn1–Cl3	110.50(5)				
Compound 2b							
Zn1–Cl1	2.2494(5)	Zn1–Cl2	2.2664(5)	Zn1–Cl3	2.2299(5)	Zn1–N2	2.0519(14)
Cl1–Zn1–Cl2	106.641(2)	Cl1–Zn1–Cl3	119.22(2)	Cl2–Zn1–Cl3	110.153(2)	N2–Zn1–Cl1	108.46(4)
N2–Zn1–Cl2	108.12(4)	N2–Zn1–Cl3	103.82(4)				
Compound 2c							
Cu1–Cl1	2.2392(2)	Cu1–Cl2	2.2585(2)	Cu1–Cl3	2.2370(2)	Cu1–N2	2.002(4)
Cl1–Cu1–Cl2	100.56(6)	Cl1–Cu1–Cl3	133.58(7)	Cl2–Cu1–Cl3	102.89(6)	N2–Cu1–Cl1	97.36(13)
N2–Cu1–Cl2	130.63(14)	N2–Cu1–Cl3	96.89(13)				
Compound 3a							
Zn1–Cl1	2.2721(5)	Zn1–Cl2	2.2851(4)				
Cl1 ¹ –Zn1–Cl1	110.86(3)	Cl1 ¹ –Zn1–Cl2	109.817(17)	Cl1–Zn1–Cl2	107.900(16)	Cl1 ¹ –Zn1–Cl2 ¹	107.898(16)
Cl1–Zn1–Cl2 ¹	109.817(16)	Cl2–Zn1–Cl2 ¹	110.817(16)				
Compound 3b							
Cu1–Cl1	2.2562(8)	Cu1–Cl2	2.2447(8)				
Cl1 ¹ –Cu1–Cl1	129.59(5)	Cl1 ¹ –Cu1–Cl2	98.85(3)	Cl1–Cu1–Cl2	100.28(3)	Cl1 ¹ –Cu1–Cl2 ¹	100.28(3)
Cl1–Cu1–Cl2 ¹	98.85(3)	Cl2–Cu1–Cl2 ¹	134.08(5)

Figure S1. Molecular structures of **2b1**, **4ab** and **5**. C–H hydrogens are omitted for clarity. Thermal ellipsoids are drawn at the 50% probability level.

In crystal structure (**4a**) one of the ethyl substituent (C9C, C10D/C9D, C10D) was disordered over two sites with occupancy ratio of 0.48/0.52. The carbon atoms of the disordered groups were restrained with effective standard deviation 0.01 so that their U_{ij} components approximate to isotropic behavior. The highest peak is located 1.04 Å from atom C10 and the deepest hole is located 0.64 Å from atom Zn1B.

In crystal structure (**5**) some of the alkoxy groups (C13-C15B) were disordered over two sites with occupancy ratio of 0.57/0.43. The Br2 atom was also slightly disordered but no disorder model was used in the final structure refinement. The highest peak is located 0.74 Å from atom Br2 and the deepest hole is located 0.80 Å from atom Br2.

Table S3. Crystallographic data for Compounds **2b1**, **4ab** and **5**.

	2b1	4a	4b	5
CCDC No	1443460	1443464	1443463	1443465
Formula	C ₁₄ H ₂₁ Cl ₃ N ₄ O ₃ Zn	C ₁₀ H ₁₆ N ₄ ·ZnCl ₄	C ₁₂ H ₂₀ N ₄ ·ZnCl ₄	C ₁₆ H ₂₇ Br ₃ N ₄ O ₄ Zn
M.Wt	465.07	399.44	427.49	644.51
Temp (K)	123	123	120	123
Crystal size/mm ³	0.246 x 0.105 x 0.089	0.080 x 0.060 x 0.020	0.260 x 0.211 x 0.121	0.120 x 0.043 x 0.025
Crystal system	Monoclinic	Monoclinic	Monoclinic	Triclinic
Space group	<i>P</i> 2 ₁ /n	<i>P</i> 2 ₁ /c	<i>P</i> 2 ₁	<i>P</i> $\bar{1}$
a (Å)	14.32177(16)	15.2092(3)	7.89382(12)	9.7953(5)
b (Å)	16.70257(18)	25.9668(8)	7.77587(11)	11.4088(6)
c (Å)	17.48443(19)	12.0487(4)	14.6157(2)	11.9915(7)
α (°)	90	90	90	67.561(5)
β (°)	104.3992(12)	95.734(2)	96.2789(14)	68.652(5)
γ (°)	90	90	90	82.433(4)
Vol (Å ³)	4051.07(8)	4734.6(2)	891.75(2)	1153.60(12)
Z	8	12	2	2
d _{calc} (g/cm ³)	1.525	1.681	1.592	1.855
μ (mm ⁻¹)	1.628	8.326	1.974	7.811
F(000)	1904	2424	436	636
reflns collected	28961	20774	3102	8400
Independent reflections	8779	9732	1631	4682
GOF	1.053	1.082	1.045	1.042
R _{int}	0.0318	0.0242	0.0206	0.0386
R _{1a} ($I \geq 2\sigma$)	0.0389	0.0455	0.0214	0.0517
wR _{2b} ($I \geq 2\sigma$)	0.0958	0.1171	0.0488	0.1394

$$^a R I = \sum \|F_o\| - \|F_c\| / \sum \|F_o\|, \quad ^b wR2 = [\sum [w(F_o^2 - F_c^2)^2] / \sum [w(F_o^2)^2]]^{1/2}.$$

3. Solution studies of Zwitterionic coordination compounds (2a-b)

3.1 Molar conductivity.

Molar conductivity of Zwitterionic coordination compounds (**2a-b**) in methanol solution was performed by using handylab LF11 instrument manufactured by Scott.

Table S4. Molar conductivity of Zwitterionic coordination compounds (2a-d) in methanol solution

	Conc. (mmol)	K (μs/cm)	K (s/cm)	Molar conductivity Δm (Scm ² mol ⁻¹)
2a	1	93.3	0.0000933	0.0933
	0.5	52.4	0.0000524	0.0524
2b	1	92.4	0.0000924	0.0924
	0.5	52.7	0.000052.7	0.0527
KCl (Ref)	0.5	52.7	0.0000527	0.0527

3.2 Mass Spectrometry

Experimental

Mass spectrometry experiments were performed with ABSciex QSTAR Elite ESI-Q-TOF mass spectrometer equipped with an API 200 TurboIonSpray ESI source from AB Sciex (former MDS Sciex) in Concord, Ontario (Canada). The parameters were optimized to get maximum abundance of the ions under study. Both positive and negative polarization mass spectra were recorded. The measurements and data handling was accomplished with Analyst® QS 2.0 Software.

For ESI-MS measurements, solid **2a** and **2b** were dissolved in MeOH to obtain a stock solution of 1 mg mL⁻¹. The stock solution was diluted in MeOH to obtain final samples (20 µM). The samples were injected into the electrospray ionization (ESI) source with a flow rate of 5 µL min⁻¹. Mass spectra were externally calibrated by ESI Tuning mix (Agilent Technologies). The compositions of the ions were verified by comparing experimental and theoretical mass values as well as isotopic distributions.

Results

ESI-MS analyses for **2a** and **2b** were performed on positive and negative polarizations and they produced very similar spectra in methanol. In positive polarization spectra ions corresponding to [2-ZnCl₃]⁺ were observed as base peaks (see Fig. XX and Table X for interpretation and mass accuracies). In addition peaks for sodium adduct [2-ZnHCl₃+Na]⁺, methanol elimination product [2-ZnCl₃-MeOH]⁺ and dimeric ion [(2)₂+Zn]²⁺ were detected. On negative polarization from both compounds [ZnCl₃]⁻ at *m/z* 168 was observed as the most abundant peak.

Ion	Composition	<i>m/z</i> _{exp}	<i>m/z</i> _{theor}	Δ(<i>m/z</i>)
[2a-ZnCl ₃] ⁺	C ₁₆ H ₂₇ N ₄ O ₄	339.1892	339.2027	0.013
[2a-ZnCl ₃ -MeOH] ⁺	C ₁₅ H ₂₃ N ₄ O ₃	307.1743	307.1765	0.002
[2a-ZnHCl ₃ +Na] ⁺	C ₁₆ H ₂₆ N ₄ O ₄ Na	361.1805	361.1846	0.004
[(2a) ₂ +Zn] ²⁺	C ₃₂ H ₅₂ N ₈ O ₈ Zn	370.1675	370.1594	-0.008
[ZnCl ₃] ⁻	Cl ₃ Zn	168.7905	168.8363	0.046
[2b-ZnCl ₃] ⁺	C ₁₈ H ₃₁ N ₄ O ₄	367.2305	367.234	0.003
[2b-ZnCl ₃ -MeOH] ⁺	C ₁₇ H ₂₇ N ₄ O ₃	335.2088	335.2078	-0.001
[2b-ZnHCl ₃ +Na] ⁺	C ₁₈ H ₃₀ N ₄ O ₄ Na	389.2132	389.2159	0.003
[(2b) ₂ +Zn] ²⁺	C ₁₈ H ₃₁ N ₄ O ₄	398.1992	398.1907	-0.009

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Figure S3. ESI-MS spectra from **2b** 20 μ M in MeOH; a) positive and b) negative polarization.

Figure S4. ^1H NMR (top) and ^{13}C NMR (bottom) spectra of Df-Me₂biim in CDCl₃.

Figure S5. ^1H NMR (top) and ^{13}C NMR (bottom) spectra of Df-Et₂biim in CDCl₃.

Figure S6. ¹H NMR (top) and ¹³C NMR (bottom) spectra of Df-Pr₂biim in CDCl₃.

Figure S7. ¹H NMR (top) and ¹³C NMR (bottom) spectra of **1a** in DMSO-d₆.

Figure S8. ¹H NMR (top) and ¹³C NMR (bottom) spectra of **1b** in DMSO-d₆.

Figure S9. ¹H NMR (top) and ¹³C NMR (bottom) spectra of **1c** in DMSO-d₆.

Figure S10. Variable temperature ^1H NMR spectra of **2a** in MeOD-d_4 . (25°C (**a**), 0°C to -80°C (**b-i**) with 10°C decreasing steps).

Highlights

Highly polar Zwitterionic coordination compounds of Zn and Cu have been synthesized by using ring and *N*-substituted biimidazole ligands. The Zwitterion formation was found to be dependent on the acidity of the reaction media and the nature of the ligand substituents.