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Towards Trigonal Prismatic Hexanuclear Copper Complexes For Catalytic Water Oxidation

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FLORIDA INTERNATIONAL UNIVERSITY

Miami, Florida

TOWARDS TRIGONAL PRISMATIC HEXANUCLEAR COPPER COMPLEXES
FOR CATALYTIC WATER OXIDATION

A dissertation submitted in partial fulfillment of the

requirements for the degree of

DOCTOR OF PHILOSOPHY

in

CHEMISTRY

by

David Ian James Kreiger

2019

To: Dean Michael R. Heithaus
College of Arts, Sciences and Education

This dissertation, written by David Ian James Kreiger, and entitled Towards Trigonal Prismatic Hexanuclear Copper Complexes for Catalytic Water Oxidation having been approved in respect to style and intellectual content, is referred to you for judgment.

We have read this dissertation and recommend that it be approved.

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ABSTRACT OF THE DISSERTATION
TOWARDS TRIGONAL PRISMATIC HEXANUCLEAR COPPER COMPLEXES
FOR CATALYTIC WATER OXIDATION

by

David Ian James Kreiger

Florida International University, 2019

Miami, Florida

Professor Raphael G. Raptis, Major Professor

Our planet is running out of energy resources and traditional renewables are not easily transported nor will be sufficient to cover the void left by fossil fuels. Therefore, a new energy storage system needs to be adopted. The medium with highest energy density that is readily available is hydrogen from water. However, to access hydrogen, water needs to be oxidized and water oxidation catalysts (WOCs) will be required for widespread adoption. In pursuit of WOCs, several hexanuclear copper complexes were synthesized and characterized, including the isolation and characterization of a mixed-valent Cu_6 (formally, $\text{Cu}^{\text{II}}_5\text{Cu}^{\text{III}}$) complex. Furthermore, early in the research, several novel coordination polymers (CPs) were serendipitously synthesized.

This dissertation is divided into three major parts:

(i) A series of trigonal prismatic Cu_6 -pyrazolato complexes (Cu_6) form a unique arrangement of two O^{2-} ions in a favorable position to form an O-O bond without much external influence. The Cu_6 complexes undergo two reversible one-electron oxidations and a structurally characterized one-electron oxidized mixed-valent

complex displays an O...O distance ~ 0.3 Å closer than the homovalent compound, which highlights the propensity of the system towards O-O bond formation. Spectroscopic findings and DFT calculations on the electronic structure of the mixed-valent Cu₆-complex are reported. The results support the hypothesis that a variation of a Cu₆ motif could function as a water oxidation catalyst.

(ii) Three new coordination polymers containing trinuclear Cu(II)-4-formyl pyrazolato units connected by formyl group coordination were prepared and the crystal structures are reported: a 1D polymer formed by linking secondary building units (SBUs) and two 3D polymers with novel topologies consist of 14-nodal $3^8 4^2, 5^4$ -c and 3-nodal 3,3,4-c nets, respectively.

(iii) Two new motifs of Cu₆-pyrazolato complexes were prepared. One motif has two pyrazolato bridges between the trinuclear copper subunits with a μ_4 -Cl situated inside the open cage, and the other has the subunits bridged by a single pyrazolato ligand. Variable low-temperature NMR and magnetic studies were conducted. And for the first time, far-infrared spectroscopy studies were used to determine the characteristic absorptions of various coordination bonds in copper pyrazolate complexes.

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ABBREVIATIONS AND ACRONYMS

AB.....	Acid-Base Mechanism
ATR-IR.....	Attenuated Total Reflectance Infrared
Bpe.....	bispyridylethylene
B/R.....	Below Range
CP.....	Coordination Polymer
DC.....	Direct Coupling
DCM.....	Dichloromethane
DFT.....	Density Functional Theory
EPR.....	Electron Paramagnetic Resonance
Eq.....	Equivalents
Exptl.....	Experimental
FTIR.....	Fourier Transform Infrared
GOF.....	Goodness of Fit
HIPPO.....	Hexanuclear Interacting Prismatic Pyrazolato Open-Cage Complex
HV.....	homovalent
IR.....	Infrared
IVCT.....	Intervalent Charge Transfer
L.....	Ligand
MeCN.....	Acetonitrile
MV.....	Mixed-Valent
N/A.....	Not Applicable
NMR.....	Nuclear Magnetic Resonance
PDB.....	Protein Data Bank
PPN.....	Bis(triphenylphosphine)iminium
PSII.....	Photosystem II
Py.....	Pyridine

Pz ⁻	Pyrazolate/pyrazolato
PzH.....	Pyrazole
PzH*.....	Functionalized Pyrazole
RT.....	Room Temperature
SBU.....	Secondary Building Unit
TGA.....	Thermogravimetric Analysis
THF.....	Tetrahydrofuran
TOF.....	Turn Over Frequency
TON.....	Turn Over Number
TPP.....	Tetraphenylphosphonium
OTf.....	Triflate (CF ₃ SO ₃) ⁻
WOC.....	Water Oxidation Catalyst
XAS.....	X-Ray Absorption Spectroscopy

Chapter 1 : General Introduction

1.1. Importance and Overview of Water Oxidation Catalysts

There are many challenges for humanity to overcome in the coming decades. The most important of these challenges is, arguably, energy security. The world's oil supply is dwindling and is projected to be depleted within 150 years at the current consumption rate. With the growing population of the planet and the increased economic development of India and China, it may be depleted much sooner.¹ Beyond the impending collapse of current living standards if an alternative energy source is not identified, there is the moral question of using our oil reserves for something as paltry and insignificant as moving metal boxes around and turning on light bulbs. The hydrocarbons in the oil reserves are of vital importance in the pharmaceutical and material science industries.

It has been suggested that a shift to other fossil fuels could compensate for this, however, while there is enough natural gas to last 200-600 years and enough coal to power the earth for another 1000-2000 years, the carbon emissions could lead to a runaway greenhouse effect.¹

The runaway greenhouse effect leaves us with renewable energy resources for our energy needs. These too, have their problems, although they can be overcome. The main issue with renewable resources is distribution. Winds and rivers are not uniformly distributed across the planet, and therefore a way to distribute the energy is needed. Solar energy, on the other hand, is available

almost everywhere, although typically for only half the day. Thus, a system for storage and transportation must be devised.

Currently, batteries are used for energy storage and transportation. But with an energy density of only 0.5 - 3.0 MJ·kg⁻¹, batteries cannot store enough power to make renewable resources viable.^{2,3} The energy storage medium should be a gas or a liquid as they have a high energy density and can easily be transported. Nuclear power would be ideal for stationary energy (or on massive vehicles like aircraft carriers), unfortunately the world is afraid of nuclear energy. Traditional liquid fuels have an energy density of ~50 MJ·kg⁻¹, which is the benchmark for useful energy storage. Of the many potential targets, hydrogen gas would be the most desirable as its only combustion product is water and it has an energy density of 140 MJ·kg at 700 atm.³ Molecular hydrogen effectively does not exist in nature and must be created from other hydrogen containing sources. The best option would be the electrolysis of cheap and plentiful water, according to equation 1.

This is a redox reaction with two half-reactions (Equations 2 and 3).

Scheme 1. Lattimer diagram for H₂O

While the energy requirement for water oxidation is acceptable for laboratory purposes, it is too large to be implemented on the scale needed to support the renewable energy transition. That is why it is imperative that a catalyst be developed to lower the activation energy, increasing the rate of formation. Ideally, the activation energy should be low enough to be in the range of photocatalysts.

1.1. Water Oxidation Catalysts

The first known water oxidation catalyst (WOC) is photosystem II (PSII), which reduces the oxidation potential by 300 mV.⁵ The active site of photosystem II was developed by Nature 2.5-3.5 billion years ago and most likely formed from the precipitation of manganese oxides that were common in the ocean at that time.⁶⁻⁸ The active site of PSII is a distorted chair-shape with the chemical formula $\text{Mn}_4\text{O}_5\text{Ca}$ (Figure 1).^{9,10}

Figure 1. The $\text{Mn}_4\text{O}_5\text{Ca}$ active center of PSII.¹⁰ Mn, purple; Ca, yellow; O, red.

Photosystem II is the active center in the chlorophyll of plants that generate most of the molecular oxygen on Earth and is essential to life as we know it. It is a very efficient catalyst with a turnover number (TON) in excess of 180,000 and a turnover frequency (TOF) of 400 s^{-1} .^{11,12} In efforts to match the efficiency of Nature's catalyst, many of the catalysts being developed are multinuclear like the PSII active center. The first synthetic WOC, the "blue dimer" (Figure 2) was synthesized in 1982 by Meyer *et al.* and had a TON of just 13.2, using cerium ammonium nitrate as a sacrificial oxidant.¹³ Although the blue dimer had a poor TON, it nevertheless caused an explosion of interest in synthetic WOCs. Since then, there has been rapid growth and development in the area of WOCs.

Figure 2. The “Blue Dimer”.¹³

In the 30 some years since the first WOC was reported many schemes and metals have been tested. There is little consensus on the best metal or the number of metal centers required for an efficient WOC and it is impossible to empirically compare them because of differences in the conditions for the catalytic testing, such as pH, oxidant, and stirring speed.^{13,15} Some Ti, Mn, Fe, Ni, Cu, Ru, Ir and Os complexes and their oxides can catalyze the water oxidation reaction;^{13,16–19} and a fully organic WOC has also been reported.²⁰ It appears that electron releasing ligands as well as their rigidity and accessibility of their reaction site will increase the reactivity of WOCs.^{13,16,21,22} Although the most efficient synthetic WOCs are mononuclear, effective WOCs tend to be complexes that have antiferromagnetically coupled high-spin metal centers.^{16,22–26} The most efficient catalyst under mild conditions to date is an iridium half-sandwich complex with a TON near 400,000 although it has a low TOF of less than 0.2 s^{-1} (Figure 3).²⁶ Ruthenium catalysts show a much higher TOF, up to an astounding TOF of $50,000 \text{ s}^{-1}$, which far surpasses PSII, but require a high pH and an over-potential of 0.7

V. (Figure 4).²⁷ However, both metals are rare and expensive and will not serve for the large scale water oxidation catalysts needed to support a global economy.

Figure 3. WOC with the highest TON. X = Cl, I, OTf.²⁶

Figure 4. WOC with the highest TOF.²⁷

As mentioned above, even the best WOCs fall far short of PSII or require harsh conditions. That is because the mechanisms for the oxidation of water by PSII and synthetic WOCs are different.^{17,22} Photosystem II has been extensively studied for decades and the reaction pathway is confirmed to proceed through the coupling of one oxygen radical to a bridging oxygen atom within the active center cluster.^{17,23,28} The pathway is significantly different from the reaction mechanism of synthetic

WOCs, which either proceed through a nucleophilic attack by water or hydroxide to a coordinated oxygen, called the acid-base mechanism (AB), or the coupling of two oxygen radicals on adjacent metal sites, known as direct coupling (DC) (Scheme 2).^{5,16,17,22} Because of the involvement of highly oxidized metal centers and oxygen radicals, synthetic WOCs suffer from ligand degradation and dissociation, and thus, catalyst deactivation.^{16,22} Catalyst deactivation is the limiting factor that keeps WOCs from being practical. Although PSII is also degraded through the generated radicals, Nature is constantly repairing the protein.^{17,22,23} A common feature of PSII and synthetic catalysts alike is that all involve two or more oxo ligands in proximity, allowing the formation of an O-O bond. Although, to our knowledge, there are no discrete molecular synthetic WOCs that are known to proceed through the reaction pathway of PSII.

Scheme 2. Direct coupling (top); Acid-base coupling (bottom).¹⁷

1.2. Polynuclear Copper Centers in Nature - Oxygen Reduction Enzymes in Comparison to Trigonal Copper Complexes

Multicopper oxidases, such as tree and fungal laccases, ascorbate oxidase, ceruloplasmin, Fet3p, CueO, and CotA will facilitate the $4e^-$ reduction of O_2 to

water; three examples are shown in Figure 5.^{29–39} The cycle for dioxygen reduction is shown in Figure 6, and according to the principle of microscopic reversibility, multicopper enzymes should also function as catalysts for the reverse reaction, water oxidation. Several of the enzymes have a triangular Cu arrangement in their active site, and in the native-intermediate form, possess a μ_3 -O bridging the three Cu atoms, which are positioned on average about 3.30 Å apart, according to calculations,^{29,32–41} while the crystal structure of CueO places the copper atoms 3.39 Å, 3.67 Å, and 3.82 Å apart,⁴² the distances are shown in Figure 7 along with a Cu-pyrazolate for comparison. Furthermore, it has been determined, using DFT and magnetic susceptibility, that the copper atoms in the enzymes and the Cu₃-pyrazolato complexes are antiferromagnetically coupled with an overall ground state spin of $S=1/2$.^{29,33,36,37,43–48} The copper oxidases cycle between [Cu^{II}₂Cu^{III}] and [Cu^{II}₂Cu^I] active centers, and involves two redox processes, which is required to facilitate the reduction of O₂ to water.^{32,34,36,41} The active center, having two redox processes is similar to the hexanuclear copper complexes from Raptis' group in that regard.^{44–55}

Figure 5. Multicopper enzymes with a trinuclear active site. A: McoC, PDB:3ZX1; B: CueO, PDB:1N68; C: CotA laccase, PDB:1W6L.³² Cu atoms are orange spheres. Note the trinuclear Cu active site in all three enzymes.

Figure 6. Catalytic cycle of a trinuclear copper enzyme.³⁸

Figure 7. Left: Crystallographically determined Cu distances in the natural intermediate of CueO (a multicopper oxidase);⁴² right: Crystallographically determined distances for Cu₃-pyrazolate.⁴⁶

1.3. Classification of Mixed-Valence Complexes

As mentioned, the multicopper enzymes require a mixed-valent system to facilitate the reduction of oxygen, therefore it is important to understand the various classes of mixed-valence and how that relates to our target compounds. Like the multicopper enzymes, the trinuclear copper-pyrazolato complexes, when oxidized, are formally Cu^{II}₂Cu^{III}, although the copper complexes are more accurately described in most cases as Cu^{2.33+3},^{47,49,50} due to charge delocalization at most time scales and temperatures. The delocalization helps stabilize the high-valent Cu center in the oxidized state. According to the Robin-Day three-class system for electron delocalization (Scheme 3),⁵⁶ these Cu-pyrazolate compounds fall into the Robin-Day class II, or partial delocalization, with class I being fully localized and class III being fully delocalized. The degree of electron delocalization is dependent

on various things, such as the nature of metals and bridging ligands and the topology of the assembly. The extent of delocalization can be determined by analysis of the inter-valence charge transfer (IVCT) band in the near infrared (NIR) region of their absorption spectra, generally between $5000 - 10000 \text{ cm}^{-1}$.^{57,58}

Scheme 3. The three classes of mixed-valence compounds according to Robin-Day classification system. A: class I, fully localized; B: class II, partially delocalized; C: class III, fully delocalized.⁵⁶

1.4. Cofacial Porphyrins and Pacman Complexes

There has been much interest in porphyrin dimer complexes (Figure 8a) and subsequently Schiff-base calixpyrrole complexes (Figure 8b), collectively called Pacman complexes, since the late 1970's when a cobalt porphyrin dimer was shown to catalyze oxygen reduction to water.^{59,60} Since then, various oxygen reduction and water oxidation catalysts were developed using Pacman-type motifs.⁶¹⁻⁶⁷ Pacman complexes show a penchant for catalysis because of their ability to allow substrates to bind to both metal centers inside the cleft, which helps facilitate multielectron processes while inhibiting side reactions.^{62,64,67,68} Various

alterations to complexes of cofacial porphyrins and calixpyrroles have emerged since that time, such as mixed metal complexes,^{69–75} two different porphyrins,^{71,76,77} multiple pillared bisporphyrins,^{78–81} and multinuclear variations.^{74,82,83} When the porphyrins are bridged by long flexible linkers, there is a great degree of conformational freedom. Another complex of note, a hexanuclear copper complex that is reminiscent of multipillared cofacial porphyrin complexes (Figure 9), was reported by the Raptis' group and consists of a two trinuclear Cu(I) pyrazolate subunits connected by organic chains in three positions.⁸⁴ However, bispyrazoles, like those used in the hexanuclear Cu(I) complex reported by Raptis, can easily lead to coordination polymers under various reaction conditions.

Figure 8. A, example of a cofacial porphyrin complex;⁸¹ B, example of a Schiff-base calixpyrrole complex.⁶⁶

Figure 9. Crystal structure of Cu_6 with trinuclear subunits connected by bispyrazoles.⁸⁴ Black, C; Blue, N; Cyan, Cu. Hydrogens omitted for clarity.

1.5. Coordination Polymer Overview

First developed in the late 1990's,⁸⁵ permanently porous coordination polymers (CPs) have attracted a lot of attention for their potential use in various applications. Coordination polymers generally involve repeating metal centers or clusters held together by organic bridging ligands, although there is no requirement for the compound to have metals or organic linkers.^{86–88} Often coordination polymers will have secondary building units (SBUs) that consist of metal clusters held together with charged bridging ligands to form a well-defined structure. The SBUs are then

connected by neutral or charged ligands forming a 1D, 2D, or 3D topology. Coordination polymers can theoretically maintain their porosity when guest molecules are removed, making them attractive for gas sorption applications. The stability makes CPs comparable to zeolites.⁸⁹

The SBUs made of metals with well understood chemistry are generally preferred as the pore size can be adjusted by using linkers that are topologically similar, but of varying lengths to form isorecticular CPs. However, the coordination and overall topology of a coordination polymer may be affected by the conditions under which the synthesis was performed, including temperature, solvent, and concentrations.⁸⁹

Specifically, trinuclear copper SBU based CPs have been shown to be effective catalysts (e.g., for waste water treatment),^{90–92} magnetic materials,^{91,92} semiconductors,^{90,92} cathode/anode materials for rechargeable batteries, and be useful for selective gas sorption.^{90,93}

1.6. Brief Overview of Pyrazolate Coordination Chemistry

Pyrazoles (pzH) and pyrazolates (pz⁻) have rich coordination chemistry. Pyrazoles are monodentate while pyrazolates have a wide variety of coordination modes available (Scheme 4),⁹⁴ although only a few are commonly found (e.g., κ^1 , κ^2 , $\kappa^1:\kappa^1-\mu$) with the other coordination modes are restricted to more exotic reaction conditions. In copper chemistry, the $\kappa^1:\kappa^1-\mu$ coordination mode features prominently in many compounds. The $\kappa^1:\kappa^1-\mu$ coordination mode allows for the

formation of triangular complexes as the bite angle between the two nitrogen atoms is 60° . The angle is similar to carboxylate ligands, but pyrazolates are less labile than carboxylates, allowing for reversible redox processes. There are also fewer atoms between the metal centers with pyrazolate ligands (N-N) compared with carboxylates (O-C-O), which allows for greater electronic and magnetic communication between the metal centers. Pyrazoles can also be functionalized at the 3, 4, and 5 positions, allowing control of electronic and steric properties, in turn enabling metal pyrazolate complexes to be modified in such a way as to preferentially favor particular geometries, nuclearities, and electrochemical properties.^{94–96}

Scheme 4. Different coordination modes of pyrazoles and pyrazolates.⁹⁰

1.7. Objectives and Hypotheses:

The conversion of water to dioxygen is a four-electron process that requires two oxygens being brought into close proximity. Hexanuclear copper pyrazolato complexes consist of two triangular subunits assembled from three copper atoms and three pyrazolates with a central $\mu_3\text{-O}^{2-}$ ligand (Figure 10). The subunits are held together by three bridging pyrazolates that place the $\mu_3\text{-O}$ atoms ~ 3.5 Å apart with little coercion. If these atoms were brought closer and the O-p_z orbitals start to overlap, upon oxidation, it may be possible to form a formal bond (Scheme 5). Once this two-electron process is complete, the $\mu_6\text{-dioxygen}$ ligand would have to be released, at which point the O_2^{2-} would easily lose two more electrons to the surrounding environment to yield an O_2 molecule, or disproportionate into O_2 and 2O^{2-} .

Scheme 5. Qualitative molecular orbital diagram if the O-p_z orbital overlaps and oxidation was centered on the $\mu_3\text{-O}$ ligands.

Figure 10 Archetypal hexanuclear copper pyrazolate.⁴⁹ Black, C; white, H; red, O; blue, N; cyan, Cu. Anions are omitted for clarity.

With the goal to enable Cu₆ motifs to function as catalysts, several aspects of these compounds must be explored or modified:

- 1) Obtain evidence that the oxidation processes of the Cu₆ motif could be μ_3 -O centered.

Currently, it is known that the Cu₆-pyrazolato complex has two redox processes, which have been shown to lead to an IVCT band, meaning that the electrons are delocalized over the Cu atoms. However, if these processes remain Cu-centered even if the μ_3 -O ligands were brought close enough together to facilitate the O-p_z orbital overlap (Scheme 5), no bond will form if the electrons are lost preferentially from the Cu centers. That is

why it is of utmost importance to isolate the oxidized Cu_6 and characterize it by single-crystal X-ray diffraction, magnetic susceptibility, K- and L-edge absorption, and NMR to determine if there is any indication that an O-O bond could form.

- 2) Bring the $\mu_3\text{-O}$ ligands closer together to facilitate O- p_z overlap which will allow for a formal O-O bond to form upon oxidation.

Overlap of the O- p_z orbitals can be accomplished by allowing the Cu_3 subunits to rotate into a staggered position. Currently, the Cu_6 motifs in the literature are held together by rigid linkers that keep the Cu atoms in an eclipsed position. As the Cu atoms are larger than the central $\mu_3\text{-O}$ ligand, they prevent sufficient overlap of the O- p_z orbitals to make a formal bond upon oxidation. Thus, it is necessary to employ flexible ligands (Scheme 6) to allow for the Cu atoms to rotate into a staggered position, thereby bringing the O atoms closer and facilitating the O- p_z overlap, which is similar to a compound Raptis has already reported.⁸⁴

Scheme 6. Qualitative example of the rotation flexible ligands would facilitate.

While this particular goal eluded this study, three new coordination polymers were synthesized, two with unique topologies, so it was decided to pursue studies on them.

3) Synthesize “open-cage” Cu₆ Motifs.

In the literature, all Cu₆ motifs are fully enclosed with three bridging ligands. If a formal O-O could be achieved, the O₂²⁻ ion would have to be released from the Cu₆ motif. To facilitate this, an open cage would present an easy route for the O₂²⁻ ion to be released (Scheme 7). Furthermore, this open cage motif would be structurally similar to the Pacman complexes which have been shown to catalytically facilitate O₂ reduction.

Scheme 7. Qualitative example of O₂²⁻ release from an open cage. Charges only account for Cu and O atoms

Chapter 2 : Mixed-Valent Cu₆-Pyrazolato Complex as a Candidate for Water Oxidation

2.1. Introduction

One of the major challenges facing humanity in recent years the identification of candidate compounds that can convert water to hydrogen and oxygen for use as an energy source. Existing technology is either energetically or economically unfeasible.¹ For example, fuel cells can execute water oxidation, but they require an exorbitant amount of energy to do so. There are plenty of water oxidation catalysts containing precious elements, such as ruthenium or iridium, but their practical applicability is limited by their cost. Scientists have been searching for an ideal compound for water oxidation, preferably using first row transition metals, such as cobalt, manganese or iron, and to lesser extent, copper, nickel, and titanium.^{11,19,97,98} Whereas, there are various examples of water oxidation catalysts in the literature, there is room for improvement: With the highest TON of 15,000, for Mn based catalyst,⁹⁹ and highest TOF of 1400 s⁻¹, for a cobalt complex,¹⁰⁰ both requiring an overpotential too great for mass commercial adoption or become easily poisoned.

We turned to copper as the metal of choice for water oxidation because of the following reasons: its versatile coordination geometry, tunable redox-potential defined by its coordination environment and above all, the ease with which one can obtain copper. Copper is virtually everywhere, and if we can repurpose copper from wires into a water oxidation catalyst, it is going to be a great achievement.

Copper(II) is also present in the active sites of many enzymes that catalyze energy intensive reactions, such as methane oxidation to methanol in multicopper oxidases, some examples of these enzymes are tyrosinase and ascorbate oxidase.^{29,30,33,35} A common feature in several of the Cu-containing enzymes is the presence of a multicopper center carrying out the electron transfer process in an efficient manner.

In addition to the literature examples of polynuclear Cu complexes with reversible redox processes,^{14,101–109} we have had a hand at a series of trinuclear Cu-pyrazolato complexes that undergo a reversible one-electron oxidation process.^{48,50} The redox processes are achieved at a relatively low potential and can be modified by the substitution of the pyrazolate ligands.⁵⁰ Additionally, we have reported the synthesis, electronic structure, magnetic properties, and redox processes of several hexanuclear trigonal prismatic Cu₆-pyrazolato complexes earlier.^{48,49} The Cu₆-pyrazolato complexes display two step-wise, one-electron redox processes.⁴⁸ It was also envisaged that, upon oxidation, depending on the substitution on the pyrazolato groups, it may be possible to transfer the oxidation from Cu-centers to the oxygen atoms to promote the formation of a favorable O...O interaction. In this paper, we present for the first time, structural and spectroscopic characterization of a mixed-valent (MV) hexanuclear complex where the oxidation has been predominantly centered on the Cu-centers. However, there is also some shortening of O...O distance compared to its homovalent (HV) analog and

increased spin density on the μ_3 -O atoms. Cu K-edge X-ray absorption spectroscopy provides support for Cu-based oxidation.

2.2. Experimental

2.2.1. Materials and Methods

All commercially available reagents were used as received. Solvents were purified by standard purification procedures. **PPN[1⁻]** was synthesized by following the previously reported procedure.⁴⁹

2.2.2. UV-vis-NIR

The UV-vis-NIR spectra were recorded on an Agilent 5000 spectrophotometer in a quartz cuvette and dichloromethane solution.

2.2.3. Infrared Spectroscopy

The IR spectra were recorded on Agilent Technologies Cary 600 Series FTIR spectrometer. The sample was mixed with CsI at an approximately 1:10 ratio ground into a powder and pressed into a pellet.

2.2.4. Single-Crystal X-ray Diffractometry

Single crystals of the compounds were mounted on a Mitegen® Cryoloop™ using Parabar™ oil and diffraction data were collected on a Bruker D8 Quest diffractometer equipped with a Photon 100 CMOS using Mo-K α radiation. Data were collected using APEX 3; structure solution was obtained using intrinsic phasing with SHELXT.¹¹⁰ Data were refined by least-square methods available within SHELXL¹¹¹ using the Olex2 interface.¹¹²

2.2.5. X-ray Absorption Spectroscopy

The Cu K-edge X-ray absorption spectroscopy experiments were performed on beamline 6-2 at and Cu L-edge absorption spectroscopy on beamline 8-2 at Stanford Synchrotron Radiation Lightsource in Stanford, CA. The beamline was equipped with Si(111) double crystal monochromator. The sample data were internally calibrated using a Cu foil edge with the first inflection point assigned to 8980.3 eV.

2.2.6. Synthesis of [1]

To synthesize [1], **PPN[1⁻]** (0.025 mmol, 25 mg) and benzoyl peroxide (0.0125 mmol, 7 mg) were mixed in 5 mL CH₂Cl₂ at room temperature for 3 hours. The reaction mixture was filtered through celite and condensed to dryness. The resulting solid was washed with cold methanol to remove excess benzoyl peroxide and the PPN-benzoate salt that was generated during the reaction. X-ray quality single crystals of [1] were obtained after one week by slow evaporation from an NMR tube. Crystalline yield 29.5%.

2.3. Results and Discussion

2.3.1. Synthesis

Mixed-valent compound [1], [$\text{Cu}_3(\mu_3\text{-O})(\mu\text{-pz})_3)_2(\mu\text{-3,5-Ph}_2\text{pz})_3$] was synthesized from reacting the previously reported⁴⁹ **PPN[1⁻]**, where PPN = bis(triphenylphosphoranilydinium), $[\text{Ph}_3\text{P}=\text{N}=\text{PPh}_3]^+$ and pz is pyrazolato anion ($\text{C}_3\text{H}_3\text{N}_2$)⁻. The oxidation was effectively performed by oxidative equivalents of $(\text{NH}_4)_2[\text{Ce}(\text{NO}_3)_6]$, H₂O₂, PhIO or benzoyl peroxide. X-ray quality single crystals

were obtained from the reaction of **PPN[1]** and 0.7 eq. benzoyl peroxide in CH₂Cl₂. **[1]** is relatively stable at room temperature but decomposes slowly to the parent **[1]** or to various Cu₃ compounds. When more than 2 eq. of oxidizing reagent is used, **[1]** decomposes.

2.3.2. UV-vis-NIR absorption Spectroscopy and UV-vis-NIR Spectroelectrochemistry

The UV-vis-NIR spectroelectrochemistry helped confirm that complexes are indeed mixed-valent as **[1]** and **[1⁺]** showed IVCT bands (Figure 11, 9700 and 10100 cm⁻¹ for **[1]** and **[1⁺]**, respectively) ~400 cm⁻¹ away from each other. The presence of IVCT bands indicates that a valence-trapped situation in **[1]** or **[1⁺]** does not exist in the electronic spectroscopy time-scale (10⁻¹⁵ s). A Γ value of ~0.45 for both **[1]** and **[1⁺]** (determined by equation 4) suggests that the system at hand is a weakly coupled delocalized system, pertaining to Robin-Day Class II (0 < Γ < 0.5).⁵⁶ H_{ab} value for **[1]** was calculated to be 4600 cm⁻¹ using (equation 5) with the r_{ab} of 3.315 Å, average intra-trimer Cu...Cu distance from the crystal structure of **[1]**.

$$\Gamma = 1 - \left(\frac{\Delta\nu}{\Delta\nu_{1/2}^0} \right)$$

Equation 4

Where $\Delta\nu$ is the energy of the band maximum and $\Delta\nu_{1/2}^0$ is the full width-at-half-height of the band.

$$H_{ab} = \frac{\mu_{12}\Delta\nu}{eR}$$

Equation 5

Where H_{ab} is the electronic coupling parameter, μ_{12} is the transition dipole moment of the IVCT band, $\Delta\nu$ is the energy of the band maximum, e is the electric charge, and R is the distance (Å) between the redox centers involved in electron transfer.

Figure 11. NIR region of the UV-vis-NIR spectra **[1]** (left) and **[1•⁺]** (right)

2.3.3. X-ray Crystallography

The electrochemical reversibility from their voltammetric study tells us that there is minimal structural alteration upon oxidation of the parent Cu^{II}_6 species. Therefore, we attempted to prepare the one- and two-electron oxidized analogues in bulk, chemically, and isolate them. Using 0.7 equivalents of benzoyl peroxide reagent, **[1]** was generated in solution, as indicated by UV-vis-NIR spectrum. After 3 hours, the reaction mixture was filtered through celite and condensed to dryness. The resulting solid was washed with cold methanol to remove excess benzoyl peroxide

and PPN-benzoate that was generated during the reaction. X-ray quality single crystals of **[1]** were obtained by slow evaporation in an NMR tube.

Scheme 8. Stepwise oxidation of **PPN[1⁻]**

Complex **[1]** crystallizes in the **hexagonal** space group $P6_3m$ and the crystallographic data are collected in Table 1 and selected angles and distances in Table 2. The standout difference between the two structures is the O...O distance: 3.467(4) Å in **[1⁻]** and 3.17(1) Å in **[1]**. The neutral compound has 3-fold symmetry and is reflected in the crystallographic hexagonal symmetry. One sixth of **[1]** is present in the asymmetric unit. The symmetry is preserved in solution, which is clear from the NMR spectrum (Figure 17). The two Cu₃-pyrazolate triangles are arranged in a trigonal prismatic manner by 3,5-diphenylpyrazolato clips. Inter-trimer Cu...Cu distances in **[1]** are 2.975(1) Å, similar to those in **[1⁻]**, which are 2.975(3) – 3.020(3) Å. Cu...Cu distances within trimers are slightly longer in **[1⁻]**.

Table 1. Crystallographic Data for **[1]**.

Identification code	[1]
Empirical formula	C ₁₂₆ H ₉₀ Cu ₁₂ N ₃₆ O ₈
Formula weight	2998.93
Temperature/K	273.15
Crystal size/mm ³	0.08 × 0.037 × 0.027
Crystal system	hexagonal
Space group	<i>P</i> 6 ₃ /m
a (Å)	13.617(2)
b (Å)	13.617(2)
c (Å)	21.124(4)
α (°)	90
β (°)	90
γ (°)	120
V (Å ³)	3392.1(1)
Z	1
ρ _{calc} (g/cm ³)	1.468
μ/mm ⁻¹	1.905
Total data	63945
Unique Data (R _{int})	2387 (R _{int} = 0.1340)
2θ range (°)	5.984 to 52.818
Goodness of fit (GOF)	1.051
R ₁ /wR ₂ (<i>I</i> > 2σ(<i>I</i>))	0.0538/0.1241
R ₁ /wR ₂ (all data)	0.0993/0.1390

Figure 12. Crystal structure of **[1]**. Black, C; white, H; blue, N; red, O; cyan, Cu.

Table 2. Structural parameters between **PPN[1⁻]** and mixed-valent **[1]**.

	[1⁻] ([Cu ₆ O ₂] ¹²⁺), Å	[1] ([Cu ₆ O ₂] ¹³⁺), Å
Cu—O	1.883(5), 1.894(5), 1.886(5), 1.890(5), 1.886(5), 1.883(5)	1.9001(2)
Cu—N _{pz}	1.957(7), 1.964(6), 1.934(6), 1.942(6), 1.946(7), 1.949(7), 1.940(6), 1.943(6), 1.943(6), 1.953(6), 1.946(6), 1.949(6)	1.9319(2)
Cu—N _{Ph2pz}	2.020(6), 2.043(7), 2.050(6), 2.051(6), 2.007(6), 2.026(6)	2.0072(2)
Cu···Cu (inter-trimer)	2.999(3), 2.975(3), 3.028(3)	2.9738(2)
Cu···Cu (intra-trimer)	3.206(3), 3.263(3), 3.253(3), 3.213(3), 3.279(3), 3.239(3)	3.2868(4)
O···O	3.461(9)	3.1681(6)
Cu-O-Cu (°)	118.5(3), 116.6(2), 120.3(3), 116.6(2), 119.6(2), 119.2(2)	119.745(9)

2.3.4. Cu K-edge X-ray absorption spectroscopy (XAS)

The K-edge XAS is instrumental in finding the oxidation state of a metal in a complex. Only rarely has the K-edge technique been used to describe high-valency in a multinuclear system.^{113,114} And when it comes to Cu(III) complexes, only a limited number of examples are available in the literature,^{113–116} where the K-edge spectrum of a mononuclear Cu(II) complex originally was compared with that of its Cu(III) congener to determine the characteristic pre-edge energies.^{113,114} The pre-edge energies for the Cu(II) is at 8979.2 eV whereas that for the Cu(III) complex is at 8980.2 eV. The intensity difference of the pre-edge features shows increased symmetry for Cu(III) – higher intensity in Cu(II) results from a greater distortion from the square planar geometry. The K-edge data generated from mixing two dinuclear Cu^{II}/Cu^{III} complexes reproduce that of a mixed-valent dinuclear Cu^{II}/Cu^{III} complex well. Isolated Cu^{II}/Cu^{III} complexes are distinguished easily by the energy shifts of the pre-edge peaks.¹¹³ However, the mixed-valent

complex and the stoichiometric mixture do not show two peaks at 8979 and 8981 eV. Rather, they show a slight shoulder in the pre-edge region.¹¹³ The second derivatives of the spectra show that the $1s \rightarrow 4p$ + shakedown transition is split indicating the Cu^{II} and Cu^{III} energies at 8985 and 8987 eV, respectively.

Unlike previous systems, the ones studied herein comprise the homovalent and the mixed-valent analogue of the $[\{\text{Cu}_3(\mu_3\text{-O})\}_2]$ -complex. Based on these findings, the K-edge spectra of **PPN[1⁻]** (all-Cu^{II}) and **[1]** (formally, Cu^{II}₅Cu^{III} or Cu₆¹³⁺) were analyzed. The spectra were recorded on powdered samples at 10 – 15 K mounted on an aluminum slit at the Stanford Synchrotron Radiation Laboratory (see Experimental Section). Maintaining the samples at low temperatures prevents photoreduction of the samples in high energy X-rays. The homovalent complex **[1⁻]** showed a typical K-edge spectrum with a pre-edge absorption at 8977.5 eV and a peak in the rising edge at 8982 eV (Figure 13). The pre-edge absorption shifts slightly in the mixed-valent complex to 8978.0 eV, while the rising edge shifts to 8982.5. The pre-edge peaks at 8977.5 eV (HV) and 8978.0 eV (MV) can be assigned to $1s \rightarrow 3d$ electronic transition in Cu.^{113,114,116} The shifts to higher energy for the pre-edge and rising edge absorptions are, presumably, consistent with the higher nuclear charge of the mixed-valent complex. The shifts are only fractional (0.5 eV, $\sim 4000\text{ cm}^{-1}$), as expected for a +0.165 overall charge difference per Cu-center between the two complexes, which is similar to what is found in the $[\text{Cu}_3(\text{O})]$ system (Figure 14),¹¹⁷ and indicates that the charge is localized on one of the two trimers. A trinuclear Cu^{III}Cu^{II}₂ system was analyzed using K-edge XAS and a shift

of 0.5 eV to higher energy indicates the presence of an additional +0.33 charge on each copper.¹¹⁴

Figure 13. (Top) Cu K-edge X-ray absorption spectrum of [1⁻] (black) and [1] (red) and the second derivate of the spectra (bottom).

Figure 14. L-edges of Cu_3^{2-} (orange) and Cu_3^- (blue).¹¹⁷

Figure 15. L-edges of **PPN[1⁻]** (orange) and **[1]** (blue).

Figure 16. Close-up of shift in L-edges of **PPN[1⁻]** (orange) and **[1]** (blue).

2.3.5. NMR

The NMR of **[1]** (Figure 17) is vastly different from **[1^{•-}]**. The paramagnetic shift of **[1^{•-}]** is almost nonexistent, with all of the peaks falling in or near the aromatic region. The peaks do not match the free ligands, indicating that the NMR is indeed of **[1]**. The NMR spectrum also contains peaks for benzoate/benzoic acid and PPN, as well as minor peaks from unoxidized **[1^{•-}]**. However, the NMR was sufficiently clean to identify the protons corresponding to **[1]**. The NMR and comparison of the proton signals from **[1^{•-}]** and **[1]** can be seen in Table 3 (labels seen in Scheme 9). The smaller paramagnetic shift indicates the spin density is farther away from the protons in **[1]**.

Figure 17. NMR of **[1]**

Scheme 9. Proton labels for **PPN[1⁻]** and **[1]**

Table 3. Proton shifts for **PPN[1⁻]** and **[1]**

	H ^a	H ^b	H ^c	H ^d	H ^e	H ^f
PPN[1⁻]	35.35	18.86	28.67	11.93		
[1]	8.07	7.91	5.81	7.37	7.64	7.11

2.3.6. Far-IR

The far-IR (Figure 18) does not show any shift in the Cu-O bond absorption. Although, this is not unreasonable. Since the copper atoms of one subunit is only reduced by 0.165 electrons the strength of the Cu-O bond will not be very different from **PPN[1⁻]**. What can be noted from the IR is the reduction in width of the peaks in the aromatic distortion frequencies, 675-775 cm⁻¹, and shows that the counterions of **PPN[1⁻]** are no longer present, indicating that the IR spectrum is, in fact, of **[1]**.

Figure 18. IR and Far IR of **PPN[1⁻]** (black) and **[1]** (red).

2.4. Conclusions

Here we present solid evidence that there is a possibility that Cu₆-pyrazolato motifs could function as water oxidation catalysts. The crystal structure of the MV-Cu₆ compound shows that the distance between the two μ_3 -O ligands is reduced by approximately 0.3 Å, to 3.17(1). The combined van der Waals radii of the two oxygen atoms is 3.04, very close to the crystallographic distance of **[1]**. The reduction of the O...O distance, suggests that upon further oxidation, and perhaps functionalization of the pyrazolates, the O atoms can be brought close enough for O-p_z overlap to occur. The Cu K-edge XAS indicates that the charge is delocalized among all the Cu atoms. The NMR shows a remarkably reduced paramagnetic shift.

Preliminary magnetism and DFT calculations performed by our collaborators point to an increased spin density on the oxygen ligands (Figure 19).

Figure 19. Qualitative map of spin density on **[1]**.

Chapter 3 : Coordination polymers based on pyrazole-4-carboxaldehyde-containing Cu₃N₆ metallacycles as building units.¹¹⁸

3.1. Introduction:

In pursuit of the aim of my research, a flexible motif needed to be synthesized. Therefore, pyrazoles with a formyl group at the 4' position were used to create the Cu₃ subunits with the idea of further modification using condensation reactions. The Cu₃ subunits with terminal Cl⁻ ligands and an overall 2⁻ charge were synthesized and isolated. When reacted with *m*-xylene diamine or 1,4-diaminobutane, an insoluble powder was obtained. Originally thought to be a polymer, the reaction was repeated by adding the reactants, on a 0.25 mmol scale, dropwise over the course of 4 hours into 500 mL of DCM to prevent polymerization. The same insoluble powder was obtained, which led to the conclusion that the product may be insoluble because of the overall charge of the resulting product. Thus, a positively charged variation with terminal pyridines and an overall charge of 1⁺ was seen as a possible solution to the insolubility problem (Scheme 10).

Scheme 10. Proposed trimer with terminal pyridine and an overall charge of 1⁺.

During the crystallization step of the Cu₃ subunits, it was discovered that the reaction had led to various coordination polymers. Studies of the polymers were pursued as they were possessed a rare Cu-aldehyde bond. And while the coordination chemistry of carbonyl compounds has been well explored in carboxylic acids, and less frequently in amides, aldehydes seldom feature in coordination complexes, except when completing a chelate bite. For example, pyridine-2-carboxaldehyde is a common chelate, whereas pyridine-bis-2,6-carboxaldehyde has been used as a pincer ligand for metals in labile oxidation states.^{119–123} Aromatic aldehydes, e.g., benzaldehyde, tend to form few first row, σ -bonded coordination complexes, except with Zn(II).^{120,124–133} In a few cases, aldehydes bind Zn(II) stronger than some anionic ligands.^{134,135} A search in the Cambridge Structural Database for coordinated aldehydes (with H-atoms on β -carbons) turns up approximately one hundred crystal structures, of which fewer than one half involve first row transition metal complexes with either a σ - or π , η^2 -bound aldehyde. Some of the frequently encountered Zn-aldehyde complexes have been studied as models of liver alcohol dehydrogenase.^{136,137} Magnetic exchange mediated by bridging, σ -bound $-\text{CH}(\text{O})$ groups has been described in a manganese 2D-coordination polymer [O-Mn: 2.197(7) Å].¹³³ To the best of our knowledge, only four coordination polymers containing CH(O)-bound Cu centers have been reported: Two containing only fairly long Cu-O interactions of 2.723(7) Å¹²⁰ and 2.877(3) Å,¹³⁸ while significantly shorter Cu-O bonds of 2.296(6) Å and 2.005(4) Å were measured in the other two.^{139,140} Polynuclear Cu(II)-pyrazolato cluster chemistry is rich with nuclearities ranging from two to 34.¹⁴¹ 4-Substituted

pyrazolates typically yield trinuclear metallacycles: We and others have published pyrazolato complexes of the general formula $[\text{Cu}_3(\mu_3\text{-E})(\mu\text{-4-R-pz})_3\text{X}_3]^z$, where E = O, OH, Cl₂, Br₂, OMe, OEt; R = H, Cl, Br, NO₂, Me, Ph, CH(O); X = Cl, Br, SCN, NO₂, CF₃CO₂, PhCOO, py and z = 1-, 2-, 1+ or 2+.^{46,142–145} The trinuclear subunits can also be strung together as structural building units (SBUs) with suitable linkers to form 1D-chains, 2D-sheets and 3D-lattices.²⁹ We have also reported the pyrazole-4-carboxaldehyde-containing, 0D-complex (PPN)₃[Cu₃(μ₃-O)(μ-4-CH(O)-pz)₃Cl₃]Cl.¹⁴² Here, we are now showing that, in the absence of other terminal or bridging ligands, 4-CH(O)-pz⁻ can be used as a μ₃,η³-linker by itself to form extended networks. Specifically, we are reporting three new coordination polymers (CPs), **[3](CF₃SO₃)₂**, **[4](CF₃SO₃)₂** and **[5](ClO₄)₂**, constructed by assemblies of [Cu^{II}₃(μ₃-OH)(μ-4-CH(O)-pz)₃]²⁺ cations clipped together by short intermolecular Cu-(O)HC bonds, along with the corresponding Cl-terminated SBU, **(Ph₄P)₂[1]**, and one bipyridine-linked CP, **[2](CF₃SO₃)₂**. Topological, geometrical and 3D-structural properties of four extended network compounds are also discussed.

3.2. Experimental:

3.2.1. Materials and Methods

The CuCl₂·2H₂O, Cu(ClO₄)₂·6H₂O, tetraphenylphosphonium chloride, 1,2-bispyridylethylene (1,2-bpe, Acros), Cu(CF₃SO₃)₂ (Alfa-Aesar), and 4-CH(O)-pzH (Combi Blocks, Inc.) were purchased from commercial sources and used without further purification. Solvents were purified using standard procedures.¹⁴⁶ All reactions were conducted open to the atmosphere.

3.2.2. X-ray Crystallography

Single crystals of the compounds were mounted on a MiteGen® CryoLoop™ or on a glass fiber and diffraction data were collected on a Bruker D8 Quest diffractometer equipped with a Photon 100 CMOS or a Bruker APEX II diffractometer with a SMART detector using Mo-K α radiation. Data were collected using APEX 3 or APEX 2 suite; structure solution was obtained using intrinsic phasing with SHELXT.¹¹⁰ Data were refined by least-square methods available within SHELXL¹¹¹ using the Olex2 interface.¹¹² Electron densities corresponding to disordered interstitial solvent molecules were accounted for by using the MASK routine available within Olex2.

3.2.3. X-ray Powder Diffraction

Powders of samples were loaded in a capillary and mounted on the Bruker D8 Quest diffractometer and 360° phi-scan images were collected. The scans were integrated using Fit2D to obtain powder patterns.^{33,34}

3.2.4. Topology

Topological analysis was performed with the ToposPro program package and the TTD collection of periodic network topologies.¹⁴⁷

3.2.5. Void Volume

Solvent accessible void volume for **[2](CF₃SO₃)₂**, **[4](CF₃SO₃)₂** and **[5](ClO₄)₂** was calculated using Mercury program package.¹⁴⁸

3.2.6. IR data

The FT-IR data were collected using an Agilent Technologies Cary 600 Series FT-IR spectrometer with 16 scans from 4000-500 cm^{-1} . Data were analyzed using Spectragryph 1.2.10 software.¹⁴⁹ Powdered samples were placed on the ATR stage and data were collected, then minimal acetonitrile was added to dissolve the compounds and data were collected from the resulting solutions.

3.2.7. Thermogravimetric Analysis

The TGA was performed with a TA instruments SDT Q600 from RT to 900 °C at a heating rate of 5 °C/min with He as the carrier gas.

3.3. Synthesis

3.3.1. $(\text{Ph}_4\text{P})_2[\text{Cu}_3(\mu_3\text{-O})(\mu\text{-4-CH(O)-pz})_3\text{Cl}_3]$, $(\text{Ph}_4\text{P})_2[\mathbf{1}]$

To Synthesize $(\text{Ph}_4\text{P})_2[\mathbf{1}]$, $\text{CuCl}_2 \cdot 2\text{H}_2\text{O}$ (1 mmol, 170.5 mg), 4-CH(O)-pzH (1 mmol, 96.1 mg) NaOH (2 mmol, 80 mg, and Ph_4PCI (0.66 mmol, 262.4 mg) were mixed in 25 mL CH_2Cl_2 overnight. The mixture was filtered to remove excess NaOH and NaCl. The filtrate was concentrated under reduced pressure and crushed out with diethyl ether. The resulting solid was filtered off and air-dried overnight. X-ray quality single crystals (0.21 mmol, 277 mg; yield 65%) were grown by slow diethyl ether vapor diffusion into a CH_2Cl_2 solution of $(\text{Ph}_4\text{P})_2[\mathbf{1}]$.

3.3.2. $[\text{Cu}_3(\mu_3\text{-OH})(\mu\text{-4-CH(O)-pz})_3(\mu\text{-1,2-bpe})_2(1,2\text{-bpe})_2(\text{H}_2\text{O})](\text{CF}_3\text{SO}_3)_2 \cdot 3\text{H}_2\text{O}$, $[\mathbf{2}](\text{CF}_3\text{SO}_3)_2$

To synthesize $[\mathbf{2}](\text{CF}_3\text{SO}_3)_2$, $\text{Cu}(\text{CF}_3\text{SO}_3)_2$ (0.60 mmol, 225 mg), 4-CH(O)-pzH (0.56 mmol, 53 mg) and NaOH (0.94 mmol, 38 mg) were mixed in 25 mL MeOH. 1,2-bpe (0.96 mmol, 187 mg) was dissolved in 25 mL MeOH and was added dropwise to the Cu salt solution over 2 h. The reaction mixture was stoppered and stirred overnight at room temperature. A small amount of brown precipitate was filtered off and the filtrate was allowed to evaporate slowly to yield blue-green crystals (0.11 mmol, 181 mg; yield 57%) of $[\mathbf{2}](\text{CF}_3\text{SO}_3)_2$ over two weeks. Analysis calculated for $\text{C}_{62}\text{H}_{58}\text{N}_{14}\text{Cu}_3\text{O}_{14}\text{F}_6\text{S}_2$: C, 46.78; H, 3.67; N, 12.32. Found: C, 46.63; H, 3.66; N, 12.34.

3.3.3. $\{[\text{Cu}_3(\mu_3\text{-OH})(\mu\text{-4-CH(O)-pz})_3(\text{H}_2\text{O})_3](\text{CF}_3\text{SO}_3)_2\}_n$, $[\mathbf{3}](\text{CF}_3\text{SO}_3)_2$

To synthesize $[\mathbf{3}](\text{CF}_3\text{SO}_3)_2$, $\text{Cu}(\text{CF}_3\text{SO}_3)_2$ (1 mmol, 361 mg), 4-CH(O)-pzH (1 mmol, 96 mg) and NaOH (2 mmol, 80 mg) were mixed in 25 mL of pyridine overnight at room temperature. The reaction mixture was filtered, and the solvent was removed under reduced pressure. The residue was suspended in 10 mL of THF and filtered (0.08 mmol, 66 mg; yield 24%). X-ray quality dark-blue single crystals of $[\mathbf{3}](\text{CF}_3\text{SO}_3)_2$ were obtained after one week.

3.3.4. $\{[\text{Cu}_3(\mu_3\text{-OH})(\mu\text{-4-CH(O)-pz})_3(\text{py})_3](\text{CF}_3\text{SO}_3)_2\}_n$, $[\mathbf{4}](\text{CF}_3\text{SO}_3)_2$

The coordination polymer $[\mathbf{4}](\text{CF}_3\text{SO}_3)_2$ was obtained as an impurity of the synthesis of $[\mathbf{3}](\text{CF}_3\text{SO}_3)_2$ and was isolated as a few X-ray quality crystals from the bulk powder of $[\mathbf{3}](\text{CF}_3\text{SO}_3)_2$.

3.3.5. $\{[\text{Cu}_3(\mu_3\text{-OH})(\mu\text{-4-CH(O)-pz})_3](\text{ClO}_4)_2\}_n$, $[\text{5}](\text{ClO}_4)_2$

The coordination polymer $[\text{5}](\text{ClO}_4)_2$ was prepared using the same procedure as $[\text{3}](\text{CF}_3\text{SO}_3)_2$, using $\text{Cu}(\text{ClO}_4)_2 \cdot 6\text{H}_2\text{O}$ (1 mmol, 370 mg) instead of $\text{Cu}(\text{CF}_3\text{SO}_3)_2$. X-ray quality crystals were obtained by slow diffusion of diethyl ether vapor into a THF solution of $[\text{5}](\text{ClO}_4)_2$ (0.23 mmol, 162 mg; yield 70%).

3.4. Crystallography

Crystallographic details on the compounds discussed here are summarized in Table 4 and selected bond lengths and angles are listed in Table 5. $[\text{2}](\text{CF}_3\text{SO}_3)_2$, $[\text{3}](\text{CF}_3\text{SO}_3)_2$, and $[\text{5}](\text{ClO}_4)_2$ contained interstitial solvent molecules that were removed using the SQUEEZE protocol available with PLATON.¹⁵⁰ The amount of solvent molecules was determined from a combination TGA and elemental analyses. In $[\text{2}](\text{CF}_3\text{SO}_3)_2$, the $(\mu_3\text{-OH})$ is located on a C_2 -axis and coplanar with the three Cu-atoms. However, its thermal ellipsoid is vertically elongated, indicating that the pyramidal OH-group is disordered among two sites, above and below the Cu_3 -plane.

Table 4. Crystallographic data parameters for compounds in Chapter 2

Compound	(Ph ₄ P) ₂ [1]	[2](CF ₃ SO ₃) ₂	[3](CF ₃ SO ₃) ₂	[4](CF ₃ SO ₃) ₂	[5](ClO ₄) ₂
Formula	C ₆₀ H ₄₉ Cl ₃ Cu 3N ₆ O ₄ P ₂	C ₆₂ H ₄₉ Cu ₃ F ₆ N ₁₄ O ₁₃ S ₂	C ₂₈ H ₂₀ Cu ₆ F ₁₂ N ₁₂ O ₂₉ S ₄	C ₃₈ H ₄₀ Cu ₆ F ₁₂ N ₁₄ O ₂₀ S ₄	C ₁₂ H ₉ Cl ₂ Cu ₃ N ₆ O ₁₂
<i>f_w</i> (amu)	1276.96	1566.89	1726.04	1740.24	689.78
Temperature (K)	293(2)	296(2)	296(2)	293(2)	296(2)
Size (mm ³)	0.31 x 0.12 x 0.05	0.27 x 0.26 x 0.23	0.10 x 0.17 x 0.17	0.54 x 0.46 x 0.24	0.28 x 0.28 x 0.18
Crystal System	Triclinic	Monoclinic	Monoclinic	Ortho- rhombic	Cubic
Space group	<i>P</i> $\bar{1}$	<i>C</i> 2/c	<i>P</i> 2 ₁ /n	<i>P</i> na2 ₁	<i>I</i> 43d
<i>a</i> (Å)	12.858(1)	11.903(1)	18.9259(8)	17.989(1)	23.8296(8)
<i>b</i> (Å)	13.770(1)	25.050(2)	15.1550(7)	17.989(1)	23.8296(8)
<i>c</i> (Å)	18.617(2)	23.801(2)	20.6283(9)	24.289(2)	23.8296(8)
α (°)	76.345(1)	90	90	90	90
β (°)	75.103(1)	97.605(1)	91.700(1)	90	90
γ (°)	66.284(1)	90	90	90	90
<i>V</i> (Å ³)	2883.3(4)	7034(1)	5914.0(5)	7860.3(8)	13531.6(1)
<i>Z</i>	2	4	4	4	16
ρ_{calc} (g cm ⁻³)	1.471	1.480	1.936	1.471	1.354
μ (mm ⁻¹)	1.343	1.044	2.390	1.792	2.074
Total data	33257	40230	1574738	146084	140252
Unique data (<i>R</i> _{int})	13010 (0.0285)	8364 (0.0252)	12220 (0.0441)	16148 (0.0351)	2327 (0.0488)
2 Θ range (°)	3.832 to 55.312	4.00 to 46.48	5.728 to 53.01	5.518 to 52.92	6.398 to 52.832
Goodness of fit (GOF)	1.025	1.034	1.040	1.038	1.068
<i>R</i> ₁ / <i>wR</i> ₂ (<i>I</i> > 2 σ (<i>I</i>))	0.0393/ 0.0970	0.0637/ 0.1799	0.0508/ 0.1384	0.0371/ 0.0935	0.0349/ 0.0930
<i>R</i> ₁ / <i>wR</i> ₂ (all data)	0.0679/ 0.1104	0.0817/ 0.1961	0.0673/ 0.1510	0.0416/ 0.0970	0.0383/ 0.0963

Table 5. Selected bond lengths and angles for complexes 1-5 in chapter 2

	[1] ²⁻	[2] ²⁺	[3] ²⁺	[4] ²⁺	[5] ²⁺
Cu-μ ₃ -O(H)	1.887(2)- 1.899(2)	1.978(2)- 1.986(4)	1.986(3)- 2.030(3)	1.981(4)- 2.009(4)	2.000(4)
Cu-N	1.948(2)- 1.963(2)	1.951(2)- 1.984(3)	1.936(4)- 1.960(4)	1.932(5)- 2.002(4)	1.93(1)
Cu-L	2.283(1)- 2.315(1)	2.099(4)	1.982(3), 2.023(3)	2.010(6)- 2.017(4)	N/A
Cu-bpe	N/A	2.045(3), 2.493(4)	N/A	N/A	N/A
O-Cu-Bpe	N/A	159.2(1)	N/A	N/A	N/A
O...Cu-plane	N/A	N/A	0.517, 0.534	0.508, 0.567	0.603
Cu-OCIO ₃ /OTf	N/A	N/A	2.427(5)- 2.565(5)	2.19(1)- 2.497(7)	2.579(3), 2.726(5)
Cu-O-Cu	118.5(1)- 120.7(1)	120.0(1)	110.0(1)- 115.5(2)	110.5(2)- 118.1(2)	111.1(2)
N-Cu-N	160.3(1)- 176.5(1)	169.3(1)- 177.3(2)	169.7(2)- 179.5(2)	167.8(2)- 177.9(2)	175.7(2)
O-Cu-O	N/A	N/A	167.5(1)- 175.5(1)	165.5(2)- 178.6(2)	171.1(2)
O-Cu-L	158.1(1)- 170.3(1)	180.0(1)	174.3(1)- 178.6(1)	162.1(2), 178.2(2)	N/A
C=O (Cu-bound)	N/A	N/A	1.239(6), 1.234(6)	1.177(8) – 1.256(8)	1.226(7)
C=O (free)	1.19(2) - 1.204(4)	1.156(6), 1.202(8)	1.183(8), 1.204(7), 1.202(8)	N/A	N/A
(C=O)O-Cu	N/A	N/A	2.020(3)	1.973(4)- 2.479(5)	1.982(4)

3.5. Results and Discussions

3.5.1. Synthesis

All five compounds form by self-assembly, upon mixing a Cu(II) salt, 4-CH(O)-pzH, NaOH, a counterion and pyridine or 1,2-bpe in an organic solvent. The only discrete trinuclear complex is **(Ph₄P)₂[1]** and has terminal chloride ligands instead of a bridging linker. Coordination polymer **[2](CF₃SO₃)₂** was obtained in a soluble form only from a dilute reaction mixture to which the linker was added slowly; Fast addition of the linker forces rapid precipitation of **[2](CF₃SO₃)₂** as an amorphous solid whose composition was determined by elemental analysis. Crystals of **[2](CF₃SO₃)₂** were formed in good yield by slow evaporation of the filtrate. In the structures of **(Ph₄P)₂[1]** and **[2](CF₃SO₃)₂**, the 4-CH(O) substituents are spectator groups.

In the absence of competing ligands (such as Cl in **(Ph₄P)₂[1]** and 1,2-bpe in **[2](CF₃SO₃)₂**), the formyl group participates in completing the coordination sphere of the Cu²⁺. The only 1D-chain in this series of compounds is **[3](CF₃SO₃)₂**, as two of the three Cu-centers of its trinuclear SBU are terminated by water ligands. In **[4](CF₃SO₃)₂** one of the Cu-centers is terminated by pyridine. Interestingly, **[3](CF₃SO₃)₂** and **[5](ClO₄)₂** formed from reaction mixtures that also contained pyridine, yet no pyridine molecule was coordinated to the Cu centers.

Figure 20. Ball-and-stick diagram of **12⁻**, counter cation omitted for clarity. Color coding: Blue, Cu; green, Cl; red, O; light blue, N, black, C and pink, H.

3.5.2. Crystal Structure Descriptions

3.5.2.1. (Ph₄P)₂[1]

Compound **(Ph₄P)₂[1]** crystallizes in the triclinic $P\bar{1}$ space group with the whole trinuclear complex in the asymmetric unit (Figure 20). Its bond lengths and angles are unexceptional (Table 5) and in agreement with several similar trinuclear Cu(II)-pyrazolates in the literature.⁴⁶ The carbonyl C-O bonds are 1.194(1) – 1.204(4) Å long.

3.5.2.2. [2](CF₃SO₃)₂

Compound **[2](CF₃SO₃)₂** crystallizes in the monoclinic $C2/c$ space group with one half molecule in the asymmetric unit. A C_2 axis passes through one of the Cu-atoms and the center of a pyrazole ring, bisecting the molecule and imposing a crystallographic disorder on one carbonyl group. The dangling carbonyl groups

in this compound are non-coordinating, with typical C-O double bond lengths of 1.156(6) and 1.202(8) Å. The Cu-centers are in 5-coordinate square pyramidal geometry. One of the bidentate bpe ligands links two Cu₃-SBUs and one is terminal (i.e., monodentate, non-catenating). One of the terminal positions is also occupied by a water molecule, which in turn is H-bonded to the non-catenating bpe linker (OH...N distance: 2.802(5) Å). A crystal packing diagram shows parallel pillars along the crystallographic *c*-axis with no π - π interactions between adjacent pyridyl rings of the 1,2-bpe linkers (ring centroid-to-centroid distances: 3.661 and 3.714 Å). In the literature, 1,2-bpe has been reported as a component of pillared structures in some soft-porous materials.¹⁵¹ In addition to the common 3D network structures in **dia** topology, **[2](CF₃SO₃)₂** is 3-fold interpenetrated (

Figure 21), as is quite common among lattices containing long bis-pyridyl linkers.^{152,153} Due to interpenetration, the void volume of this lattice is only 5.8%. Pairs of adjacent C=C bonds in **[2](CF₃SO₃)₂** are only 3.791(6) Å and 3.603(6) Å apart. Vittal et al. have reported Single-Crystal-to-Single-Crystal cycloaddition reactions under similar conditions (3.79, 3.77 and 3.99 Å).¹⁵⁴

Figure 21. Diagram of three-fold interpenetration of $[2]^{2+}$ viewed along the crystallographic c -axis. H-atoms and counter anions are not shown for clarity.

3.5.2.2.1. Topology of $[2](CF_3SO_3)_2$

The underlying topology of $[2](CF_3SO_3)_2$, is a 3-nodal 3,4,5- c net of 3,4,5T174 topological type (Figure 22). The connectivity of the copper clusters in $[2]^{2+}$ is described by the 4- c uninodal net of **dia** topological type (Figure 23). The coordination polymer structure comprises three interpenetrating networks, which are related by translations along the $[100]$ direction, thus being characterized by interpenetration Class Ia .¹⁵⁵

Figure 22. Standard representation of 2^{2+} : a) The net obtained after primary simplification; b) The 3,4,5-c underlying net of 3,4,5T174 topological type obtained after secondary simplification procedure.

Figure 23. Underlying net of 2^{2+} obtained by the cluster representation procedure at 6-rings: a) Dissection of the structure by means of the cluster simplification procedure as implemented in ToposPro; b) The 4-c underlying net of **dia** topological type obtained.

3.5.2.3. [3](CF₃SO₃)₂

Compound [3](CF₃SO₃)₂ crystallizes in the monoclinic *P*2₁/*n* space group, with two Cu₃ SBUs in the asymmetric unit, connected via the carbonyl oxygen of the 4-CH(O)-pz ligand forming an 1D chain. Two of its three Cu-centers are terminated by water molecules, preventing extension into 2D- or 3D-network (Figure 24). The overall backbone structure of the Cu₃-units is similar to that found in the literature with other Cu₃N₆ metallacycles and the two μ₃-O(H) are located 0.536(1) and 0.516(1) Å above their respective Cu₃-planes. The Cu-centers are all 5-coordinate in a square pyramidal N₂O₃ environment. Axial coordination is provided by a CF₃SO₃ counter-anion, while pyrazolato-N, μ₃-O and water or the carbonyl-O atoms are in the equatorial plane. As mentioned earlier, carbonyl groups of two (out of six) 4-CH(O)-pz are involved in catenation, causing elongation of the C-O bond distances of the coordinating carbonyl groups: 1.234(1) Å and 1.239(1) Å, while the C-O bonds of non-coordinating carbonyl groups are shorter than 1.210 Å and close to a typical C-O bond in an aldehyde and also those seen in 1²⁻. While 3D structures are fairly common with Cu₃-pyrazolato subunits, 1D chains and 2D sheets are rather uncommon, with just a few examples in the literature reported by us and others.^{93,156–167} The two Cu₃-SBUs differ from each other in that two loosely associated water molecules are coordinated perpendicular to the Cu₄-Cu₅-Cu₆ subunit.

Figure 24. Molecular structures of one of the chains of **3²⁺** (top); Packing diagram showing the wave-like arrangement parallel to the crystallographic a-axis (bottom). CF₃SO₃ anions are not shown. Color coding: Blue, Cu; red, O; light blue, N, black, C and pink, H.

3.5.2.4. [4](CF₃SO₃)₂

Compound **[4](CF₃SO₃)₂** has a 3D-network structure that crystallizes in the orthorhombic *Pna*2₁ space group with two Cu₃ SBUs in the asymmetric unit (Figure 25). The Cu₃N₆ metallacycles have Cu-centers with either a 5-coordinated square pyramidal or a pseudo-octahedral geometry. The axial positions are generally occupied by triflate ions with bond distances of 2.19(1)-2.497(7) Å. An interesting feature of **[4](CF₃SO₃)₂** is that all carbonyl groups are involved in catenation and all but two C-O distances are over 1.2 Å. The two shorter carbonyl C-O bonds of 1.177(8) Å and 1.179(9) are weakly coordinating at the axial position of Cu atoms at 2.315(5) Å and

2.479(5) Å, respectively. These two bonds are ~0.3-0.4 Å longer than the other CO-Cu bond distances. The shortest CO-Cu bond in **[4](CF₃SO₃)₂** is 1.973(4) Å, far shorter than analogous Cu-OC bonds hitherto reported.^{120,138–140} The solvent accessible void volume is 2982.98 Å³ (29%). The structure differs from that of **[5](ClO₄)₂** (*vide infra*) in the pyridine coordination. The Cu₃(μ₃-OH)-unit is also similar to those found in the literature¹⁴⁵ with the μ₃-O(H)-located 0.560(4) and 0.508(4) Å above the Cu₃-planes.

Figure 25. Molecular packing diagram for **4**²⁺ viewed along the crystallographic a-axis. Color coding: Blue, Cu; red, O; light blue, N and black, C. H-atoms and counter anions are not shown for clarity.

3.5.2.4.1. Topology of $[4](\text{CF}_3\text{SO}_3)_2$

In the standard simplification procedure,¹⁵⁵ the Cu atoms are shown, while ligands are represented by their centers of mass, keeping the connectivity with their neighbors (Figure 26a). Upon secondary simplification of the obtained net, by removing of the 0- and 1-coordinated ligands (nodes) and replacing the 2-coordinated ligands (nodes, or bridge structural groups) by net edges. The resulting topology is a 14-nodal $3^8,4^2,5^4$ -c net. The point symbol for this net is $\{3.4^2\}_2\{3.8.9\}_2\{3^2.4^2.5.6.8.9.10^2\}_2\{3^2.4^2.5.6.8.9^2.10\}\{3^2.4^2.5.6.8^2.9.10\}\{4.8.10\}\{4.8.9\}\{4.8^2\}_2\{4.2.5.8.10^2\}\{4^2.5.8^2.10\}$, and is not contained in the TOPOS Topological Database. (Figure 26b).

Figure 26. Standard representation of 4^{2+} : a) The 14-nodal $3^8,4^2,5^4$ -c net obtained after primary simplification. Red spheres correspond to copper atoms, purple spheres represent hydroxyl anions, grey spheres - 4-CH(O)-pz ligands and violet spheres – pyridine molecule; b) The underlying net of new topological type obtained after secondary simplification procedure.

Using a structure simplification procedure in ToposPro, the $C_{16}H_{12}N_6O_7F_6S_2Cu_3$ were identified as having the connectivity described by the 6-c uninodal net of the **sol** topological type (Figure 27).¹⁵⁵ There is only one other CP that shows this topology.¹⁶⁸

Figure 27. Underlying net of 4^{2+} obtained by the cluster representation procedure at 6-rings: a) Dissection of the structure to clusters by ToposPro; b) The 6-c underlying net of the **sol** topological type obtained after secondary simplification procedure.

3.5.2.5. $[5](ClO_4)_2$

Compound $[5](ClO_4)_2$, another 3D polymer, crystallizes in the highly symmetric cubic space group $\bar{4}3d$ with the asymmetric unit containing one-third of the Cu_3 SBU, i.e., one Cu^{2+} , one 4-CH(O)-pz⁻, one-third μ_3-OH^- and two-thirds ClO_4^- counter anion. In the extended structure, all carbonyl groups are involved in Cu-coordination (Figure 28) with CO-Cu bonds of 1.982(4) Å, the second shortest bond among the structures reported here, and a long C-

O bond of 1.226(7) Å. The structure is characterized by six contacts around the Cu₃ SBU, three dative bonds to neighboring SBUs, and three dative bonds from neighboring SBUs. The solvent accessible void volume is 4294.78 Å³ (31.7%).

Figure 28. Molecular packing diagram of **5²⁺** viewed along the crystallographic [111] plane. Counter anions and H-atoms are not shown for clarity. Color coding: Blue, Cu; red, O; light blue, N, and black, C.

3.5.2.5.1. Topology of $[5](\text{ClO}_4)_2$

After procedure for simplification of the network, described for $[4](\text{CF}_3\text{SO}_3)_2$, a 3-nodal 3,3,4-c net is determined; this connectivity is not encountered in the Topos Topologic Database (Figure 29). The point symbol for the net is $\{4.8^2\}_3\{4^2.6.8^2.10\}_3\{4^3\}$.

Figure 29. Standard representation of 5^{2+} : the 3,3,4-c underlying net of unknown topological type obtained after secondary simplification procedure.

Cluster simplification revealed that the net created by $\text{C}_9\text{H}_6\text{N}_6\text{OCu}_3$ clusters is described by the 6-c uninodal net of the **bcs** topological type (Figure 30).¹⁵⁵ There are two other CPs that display **bcs** topology.^{169,170}

Figure 30. Underlying net of 5^{2+} obtained by the cluster representation procedure at 5-rings: a) Dissection of the structure to clusters by ToposPro; b) The 6-c underlying net of the **bcs** topological type obtained after secondary simplification procedure.

3.5.3. Powder X-Ray Diffraction Patterns

The powder diffractions of bulk $[2](CF_3SO_3)_2$ and $[5](ClO_4)_2$ match well with the simulated patterns, Figure 31 and Figure 33, respectively. Single crystals of compounds $[3][CF_3SO_4]_2$ (major product) and $[4][CF_3SO_4]_2$ (minor product) were isolated from the same reaction mixture. The experimental diffraction pattern of bulk reaction product reflects this and is a good match to the overlapping simulated patterns of the two components (Figure 32).

Figure 31. Powder diffraction of $[2](CF_3SO_3)_2$. Red, experimental; black, simulated

Figure 32. Powder diffraction of $[3](CF_3SO_3)_2$. Experimental, purple; simulated pattern of $[3](CF_3SO_3)_2$, green; simulated pattern of $[4](CF_3SO_3)_2$, blue.

Figure 33. Powder pattern (red) of **[5](ClO₄)₂** compared to the simulate (green).

3.5.4. Thermogravimetric Analysis

Thermogravimetric analyses of **[2](CF₃SO₃)₂**, **[3](CF₃SO₃)₂** and **[5](ClO₄)₂** indicate that all three release solvent molecules before decomposition. Approximately 5 weight % of **[2](CF₃SO₃)₂** is lost at < 95 °C corresponding to two interstitial MeOH molecules, followed by decomposition at ~230 °C. Approximately 10 weight % of **[3](CF₃SO₃)₂** and **[5](ClO₄)₂** is lost at < 95 °C corresponding to five H₂O and one THF molecules, respectively. Upon further heating, **[3](CF₃SO₃)₂** undergoes an additional ~ 10 weight % loss, then decays sharply at 510 °C. Compound **[5](ClO₄)₂** behaves similarly to **[2](CF₃SO₃)₂** (Figure 34).

Figure 34. TGA of **[2](CF₃SO₃)₂** (blue), **[3](CF₃SO₃)₂** (black) and **[5](ClO₄)₂** (red).

3.6. Experimental Applications

As proof of concept that the synthesis of lattices **3²⁺** and **5²⁺** (and presumably also **[4](CF₃SO₃)₂**) can be reversed while maintaining intact the Cu₃-SBUs, their dissolution was studied by IR and the ¹H-NMR spectra were examined. Compound **[3](CF₃SO₃)₂** displays absorptions in the IR spectrum for both coordinated (1616 cm⁻¹) and uncoordinated (1663 cm⁻¹) carbonyl groups. Upon dissolution in acetonitrile, a slight, but visible, color change occurred and only one carbonyl absorption appeared at 1680 cm⁻¹ (Figure 35). The IR spectrum of **[5](ClO₄)₂** shows one carbonyl absorption at 1614 cm⁻¹,

associated with the coordinated aldehyde. Once **[5](ClO₄)₂** was dissolved in acetonitrile, this band shifted to 1675 cm⁻¹ (Figure 36), indicating that the aldehyde was no longer coordinated. Coordination polymer **[2](CF₃SO₃)₂** is not soluble in MeCN, its IR spectrum showing only the 1668 cm⁻¹ absorption corresponding to uncoordinated CH(O) groups (Figure 37). The NMR spectrum of **[5](ClO₄)₂** in CD₃CN has two paramagnetically shifted resonances at 37.69 ppm (1H, *w*_{1/2} = 1033 Hz, C(O)H) and 19.85 ppm (2H, *w*_{1/2} = 312 Hz, H^{3,5}) corresponding to a Cu^{II}₃-pyrazolato complex (Figure 38). The chemical shifts of these resonances are similar to those of analogous Cu₃ compounds we have previously reported,¹⁴² clearly indicating the presence of a trinuclear species in the solution of **[5](ClO₄)₂**.

Figure 35. IR of **[3](CF₃SO₃)₂** before (blue) and after (orange) adding acetonitrile.

Figure 36. IR of $[5](\text{ClO}_4)_2$ before (blue) and after (red) adding acetonitrile.

Figure 37. IR of $[2](\text{CF}_3\text{SO}_3)_2$ before (blue) and after (orange) adding acetonitrile.

Figure 38. ^1H NMR of $[5](\text{ClO}_4)_2$ in acetonitrile- d_3

3.7. Conclusions

By reporting the first three CPs based on Cu_3 -4-formyl pyrazolate metallacycles, we have shown that formyl groups can be exploited as ligands for $\text{Cu}(\text{II})$ centers (even in the absence of chelation, such as seen with pyridine-2-carboxaldehyde) and further exploited this property towards catenation of the trinuclear SBUs. The extended network structures exhibit uncommon topologies, such as **sol** and **bcs** (in 4^{2+} and 5^{2+} , respectively). When competing, stronger-coordinating ligands are present, the aldehyde groups remain free and either catenation is accomplished through other bridging ligands (e.g., 1,2-bpe in 2^{2+}), or 0-D Cu_3 complexes are formed

when terminal ligands (e.g., chlorides in **1**²⁺) are present. Finally, we have shown that the assembly of the Cu₃-SBUs into the formyl-linked CPs **3**²⁺ - **5**²⁺ is reversible (Scheme 11).

Scheme 11. Reaction pathway determined in the study.

Chapter 4 : Synthesis and Characterization of Hexanuclear Interacting Prismatic Pirazolato Open-Cage Complexes (HIPPOs); Structural, VT-NMR, Magnetic, and Far-IR Studies

4.1. Introduction

For years, various groups have been on a quest to characterize metal-metal interaction of copper complexes comprised of trigonal Cu_3 subunits. Specifically, how the various connections, such as covalent or supramolecular interactions, between the copper clusters effect the magnetics, EPR, NMR, and redox processes. This quest started in 2002 when Kamiyama et al. synthesized the first trigonal prismatic Cu_6 pyrazolato complex.¹⁷¹ What makes these complexes so interesting is the electronic tunability of ligands and the plethora of ways to connect the Cu_3 subunits together. The connectivity found in literature (Scheme 12) include supramolecular conjugation through weakly binding facially oriented counterions (a),^{172,173} a hydrogen being shared by two $\mu_3\text{-O}$ (b),¹⁷⁴ coordinated carboxylates orienting the subunits linearly (c),⁵¹ carboxylates orienting the subunits facially (d),⁵¹ a terminal anion weakly binding a neighboring Cu_3 (e),¹⁷⁵ through direct μ -pyrazolate bridging ligands (f),^{47,48,170} a single halide (g),¹⁷⁷ direct μ -pyrazole bridges and an encapsulated halogen (h),^{171,53,55} a coordinated copper atom (i),¹⁷⁸ or pyrazoles and another ligand (j).¹⁷⁶

Scheme 12. Representations of the various motifs of Cu_6 present in literature.

The magnetic interactions in Cu_3 -pyrazolato subunits are complicated by spin frustration, with the magnetic relationship being dictated by the μ_3 -capping ligand. The magnetic interaction switches from antiferromagnetic to ferromagnetic occurring as the μ_3 -capping ligand rises above the basal plane to make more acute

angles with the copper centers, with the transition occurring at $\phi = 92^\circ$.^{44,45} What is less understood however, is how the Cu₃ subunits interact when connected in various ways. What is known so far is that when the Cu₃ subunits are interacting through supramolecular interactions, there is a weak antiferromagnetic coupling between the Cu₃ subunits^{172,174,177} and direct pyrazolate bridging leads to strong antiferromagnetic coupling between the Cu₃ subunits.^{48,53,171} Furthermore, encapsulating an ion inside a Cu₆ cage leads to significantly weaker antiferromagnetic coupling within each subunit as it pushes the μ_3 -OH ligand farther out of the basal plane,⁵³ and unsymmetric ligands, specifically ones that lead to a non-parallel basal plane reduce the antiferromagnetic coupling between the Cu₃ subunits.¹⁷⁶ However, there are many more ways one can imagine connecting Cu₃ subunits. Another aspect of these compounds that has been grievously neglected is the Far-IR absorptions, which are diagnostic of Cu-halogen, Cu-oxygen or Cu-nitrogen bond strengths, with no far-IR data in the literature even in regards to simple Cu₃-pyrazolato complexes, of which, the first motif was reported in 1983.¹⁸⁰ Far-IR assignments of Cu-N, O, or X bonds are rare for all Cu-complexes, especially involving bridging ligands, with only a few studies being conducted.^{181–187} Here, we present the first open-cage Hexanuclear Interacting Prismatic Pyrazolato Open-cage complexes, “HIPPOs”, which have a structure reminiscent of “Pac-Man” complexes,^{188,189} with Cu₆ compound having the most similar structure being reported by Coronado.¹⁷⁶ In this paper the crystal structures of various HIPPOs are discussed as well as magnetic susceptibility, NMR and the Far-IR absorptions of HIPPOs and related complexes.

4.2. Experimental

4.2.1. Materials and Methods

All reagents were purchased from commercial sources and used without further purification. 4-Ph-pzH,¹⁹⁰ 4-Cl-pzH,¹⁹¹ and compounds **[1]** and **PPN[2]** were synthesized according to published procedures.^{49,53} Solvents were purified using standard procedures.¹⁴⁶ CuCl₂·2H₂¹⁸O was prepared by drying CuCl₂·2H₂O in an oven at 125°C overnight until the brown anhydrate was obtained, then H₂¹⁸O was added to the salt. Na¹⁸OH was prepared by adding H₂¹⁸O to Na metal under argon. All reactions were performed under N₂ unless mentioned otherwise. Elemental analysis was performed at Galbraith Laboratories, Inc.

4.2.2. Variable Low-Temperature NMR

¹H-NMR was recorded in acetone-d₆, THF-d₈, or CD₂Cl₂ in a Bruker Avance 400 NMR spectrometer. T₂ times were calculated using the equation $T_2^* = 1/(w_{\frac{1}{2}}\pi)$, assuming $T_2 \sim T_2^*$, where $w_{\frac{1}{2}}$ is width-at-half-height, at room temperature unless otherwise noted.

4.2.3. IR and Far-IR

Far-IR data were collected using an Agilent Technologies Cary 600 Series FTIR spectrometer with 4 scans from 3500-250 cm⁻¹. Data were analyzed using Spectragryph 1.2.10 software.¹⁴⁹ The samples were mixed with cesium iodide salt, at an approximate 1:10 ratio, and ground into a fine powder. The powder was pressed into a pellet and was used for IR collection.

4.2.4. Computational Methods for Calculated IR

The geometry optimizations and frequency calculations of all copper complexes have been carried out by Dr. Daniel Belisario Lara using DFT with the B3LYP functional included with the GAUSSIAN-09 package.¹⁹² Non-metal atoms were calculated using a triple- ζ valence polarized basis set. Copper atoms were treated using the LANL2DZ basis effective core potentials.¹⁹³ Comparisons were calculated between treating the copper atoms at the B3LYP/TVZP level and the B3LYP/LANL2DZ level, with closer agreement on both bond distances and frequencies found with the effective core potential. The Cu-Cl distances were found to be overestimated by 0.1-0.5 Å when compared to values obtained via X-ray diffraction. Bond distances for other components and ligands showed similar deviations in agreement with accepted values. Frequencies were typically in agreement to experimental data by 30 cm⁻¹, except for one frequency associated with Cu-Cl. The peak shapes of simulated spectra were reasonable approximations of the experimental data. Counterions were not included in the calculations in order to off-set some of the computational cost of these relatively large complexes. Phenyl rings attached to pyrazolate were substituted with hydrogen atoms to further offset computational time.

4.2.5. Magnetic Data Acquisition

Temperature-dependent ($T = 1.9 - 300$ K, $B = 1.0$ T) and field-dependent ($B = 0 - 9$ T, $T = 2, 5$ and 10 K) magnetization measurements were performed by Dr. R.

Herchel using a PPMS Dynacool VSM magnetometer (Quantum Design). The magnetic data were corrected for the diamagnetism of the constituent atoms.

4.2.6. Theoretical Calculations for Magnetic Data

The DFT theoretical calculations were carried out with the ORCA 3.0 computational package.¹⁹⁴ The B3LYP functional was used for calculations of the isotropic exchange constants J ,^{195–197} following the Ruiz approach,^{198,199} by comparing the energies of high-spin and broken-symmetry spin states. The relativistic effects were treated with the 0th order regular approximation (ZORA) using a relativistic version of the polarized triple- ζ quality basis set, ZORA-def2-TZVP(-f) for Cu, N, Cl and O atoms, and ZORA-def2-SVP basis set for C and H atoms.²⁰⁰ The calculations utilized the RI approximation with the decontracted auxiliary SARC/J Coulomb fitting basis set²⁰¹ and the chain-of-spheres (RIJCOSX) approximation to exact exchange as implemented in ORCA.^{202,203} Increased integration grids (Grid5 and Gridx5 in ORCA convention) and tight SCF convergence criteria were used in all calculations. The molecular fragment used in the calculations was extracted from the experimental X-ray structure. The calculated spin density was visualized with VESTA 3 program.²⁰⁴

4.2.7. X-ray Crystallography

Single crystals of the compounds were mounted on a MiteGen® CryoLoop™ or on a glass fiber and diffraction data were collected on a Bruker D8 Quest diffractometer equipped with a Photon 100 CMOS or a Bruker APEX II diffractometer with a SMART detector using Mo-K α radiation. Data were

collected using APEX 3 suite; structure solution was obtained using intrinsic phasing with SHELXT.¹¹⁰ Data were refined by least-square methods available within SHELXL¹¹¹ using the Olex2 interface.¹¹² Electron densities corresponding to disordered interstitial solvent molecules were accounted for by using the MASK routine available within Olex2.

4.3. Synthetic procedures

4.3.1. H-, Cl- and Br-HIPPO

The **H-**, **Cl-**, and **Br-HIPPO**, $\text{TPP}[\{\text{Cu}_3(\mu_3\text{-OH})(\mu\text{-4-R-pz})_3\text{Cl}\}_2(\mu\text{-4-R-pz})_2(\mu_4\text{-Cl})]$, (R= H for **H-HIPPO**, Cl for **Cl-HIPPO**, and Br for **Br-HIPPO**) were synthesized on a 1 mmol scale with 3:2:11:1 ratio of $\text{CuCl}_2 \cdot 2\text{H}_2\text{O}$, the appropriate 1H-pyrazole, NaOH, and tetraphenylphosphonium chloride, respectively. The reactants were stirred overnight in 25 mL THF under a blanket of N_2 . The reaction mixtures were then collected, and the filtrate was concentrated using a rotary evaporator. The product was crushed out by the addition of diethyl ether or hexane. The product was collected by suction filtration and allowed to air dry overnight. The powdered sample was collected and redissolved in 4 mL THF and then ~7.5 mL of toluene was added to the solution. The solutions were then layered with 15 mL of hexane and allowed to crystallize. The **Cl-HIPPO** was obtained in 71.7 % yield with respect to Cu. The **Br-HIPPO** and **H-HIPPO** were obtained in 32.4 and 34 % yields, respectively.

Elemental analysis for $\text{Ph}_4\text{P}[\{\text{Cu}_3(\mu_3\text{-OH})(\mu\text{-4-Cl-pz})_3\text{Cl}\}_2(\mu_4\text{-Cl})](\mu\text{-4-Cl-pz})_2\cdot\text{thf}$, Compound **2**, ($\text{C}_{52}\text{H}_{46}\text{Cl}_{11}\text{Cu}_6\text{N}_{16}\text{O}_3\text{P}$), calculated/found (%): C, 35.79/36.06; H, 2.66/2.85; N, 12.84/12.84.

4.3.2. Ph-HIPPO

The synthesis of $[\{\text{Cu}_3(\mu_3\text{-OH})(\mu\text{-4-Ph-pz})_3\text{Cl}\}_2(\mu\text{-4-Ph-pz})_2(\mu\text{-Cl})]$, **Ph-HIPPO**, was carried out on a 0.2 mmol scale in 15 mL of MeCN, using a 1:1:1:2 ratio of $\text{CuCl}_2\cdot 2\text{H}_2\text{O}$, 4-Ph-pzH, pyridine, and NaOH. It was stirred for 4 hours and subsequently filtered using suction. The filtrate was allowed to stand until all solvents evaporated and X-ray quality crystals of **Ph-HIPPO** were obtained after 2-3 weeks. The **Ph-Hippo** was obtained in 50.1% yield with respect to Cu. Elemental analysis for $[\{\text{Cu}_3(\mu_3\text{-OH})(\mu\text{-4-Ph-Pz})_3\text{pyCl}\}_2(\mu_2\text{-Cl})(\mu\text{-4-Ph-pz})]$, ($\text{C}_{73}\text{H}_{61}\text{Cl}_3\text{Cu}_6\text{N}_{16}\text{O}_2$) $\cdot 2\text{NaCl}\cdot\text{MeCN}$ calculated/found (%): C, 49.56/48.7; H, 3.77/3.87; N, 13.10/13.34.

4.3.3. ^{18}O -H-HIPPO

The synthesis of $\text{TPP}[\{\text{Cu}_3(\mu_3\text{-}^{18}\text{OH})(\mu\text{-4-R-pz})_3\text{Cl}\}_2(\mu\text{-pz})_2(\mu_4\text{-Cl})]$ used specially prepared reagents on a 1 mmol scale with 3:2:11:1 ratio of $\text{CuCl}_2\cdot 2\text{H}_2^{18}\text{O}$, 1H-pyrazole, Na^{18}OH , and tetraphenylphosphonium chloride, respectively. Before use, the pyrazole was placed in a desiccator for a week. The reagents were stirred overnight in 25 ml of dry, degassed THF under a blanket of argon with a 50 μl aliquot of H_2^{18}O . The resulting solution was filtered and set for diethyl ether vapor diffusion which yielded X-ray quality crystals of **TPP** $[\{\text{Cu}_3(\mu_3\text{-}^{18}\text{OH})(\mu\text{-4-R-pz})_3\text{Cl}\}_2(\mu\text{-pz})_2(\mu_4\text{-Cl})]$ after two days. The crystalline product yield was 22.8 %.

4.4. Results

4.4.1. Crystallography

4.4.1.1. H-, Cl-, and Br-HIPPO

Three HIPPOs, **H-**, **Cl-**, and **Br-HIPPO** consist of two triangular $\text{Cu}_3(\mu\text{-4-R-pz})$ -subunits held together by two bridging $\mu\text{-4-R-pz}$ ligands and a $\mu_4\text{-Cl}$. This structural feature is unique in this series of complexes. The **H-HIPPO** (Figure 39), **Cl-HIPPO** (Figure 40) and **Br-HIPPO** (Figure 41) crystalized in the triclinic $P\bar{1}$ space group and are isostructural. The asymmetric unit encompasses an entire molecule with the TPP counter ion tucked inside the cage opening (Figure 42). In **H-HIPPO**, a THF solvent molecule is hydrogen bonded to the $\mu_3\text{-hydroxo}$ proton on one of the subunits. Crystallographic details are listed in Table 6, with selected bond lengths listed in Table 7.

In all three of these structures a $\mu_4\text{-Cl}$ ligand resides within the “jaws” of the complex, with Cu-Cl bond lengths between 2.560(1) and 3.040(1). The $\mu_3\text{-OH}$ ligands sit 0.555 to 0.608 Å above the plane, which is in agreement with previous literature.^{53,179} The two panes formed by the Cu atoms of the trimer subunits form 73.9, 66.8, and 66.1° angles for **H-HIPPO**, **Cl-HIPPO**, and **Br-HIPPO**, respectively

Figure 39. **H-HIPPO** – front view. Color code: Black, C; Grey, H; Red, O; Blue, N; Cyan, Cu; Green, Cl. Counter anions omitted for clarity.

Figure 40. **Cl-HIPPO** – side view. Color code: Black, C; Grey, H; Red, O; Blue, N; Cyan, Cu; Green, Cl. Counter anions omitted for clarity.

Figure 41. **Br-HIPPO** – front view. Color code: Black, C; Grey, H; Red, O; Blue, N; Cyan, Cu; Green, Cl. Counter anions omitted for clarity.

Figure 42. Counter ion placement in **H-HIPPO**.

4.4.1.2. Ph-HIPPO

The **Ph-HIPPO** (Figure 43) crystallizes in space group P_{bcn} with one half of the molecule in the asymmetric unit. The copper trimers in **Ph-HIPPO** are bridged by a single 4-phenylpyrazolate ligand, the Cu-N bond length of 2.026(3) Å. The other two Cu atoms are terminated by a chloride and pyridine in each subunit. The **Ph-HIPPO** has a Cl⁻ in between the two trigonal trimers similar to the other HIPPOs, it is only bound to four of Cu atoms, two have a Cu-Cl bond length of 2.668(1) Å, the other two have a bond length of 2.941(2) Å. The non-bonded Cu-Cl distances are 3.535(1) Å. This Cl-atom bridges diagonally across the opening with an angle of 159.51(7)°. The μ_3 -OH ligands sit 0.589 Å above the plane, again similar to literature.¹⁷⁹ The Cu plane in the singly bridged **Ph-Hippo** is at a 31.2° angle, much closer to being cofacial than in the other HIPPOs.

Figure 43. **Ph-HIPPO** – front view. Color code: Black, C; Grey, H; Red, O; Blue, N; Cyan, Cu; Green, Cl.

Table 6. Crystal parameters for chapter 4.

Identification code	H-HIPPO	Cl-HIPPO	Br-HIPPO	Ph-HIPPO
Formula	C ₁₁₂ H ₁₂₄ Cl ₆ Cu ₁₂ N ₃₂ O ₈ P ₂	C ₄₈ H ₃₈ Cl ₁₁ Cu ₆ N ₁₆ O ₂ P	C ₉₆ H ₇₆ Br ₁₆ Cl ₆ Cu ₁₂ N ₃₂ O ₄ P ₂	C ₇₃ H ₆₁ Cl ₃ Cu ₆ N ₁₆ O ₂
fw (amu)	3083.54	1673.10	4057.56	1681.96
Size (mm ³)	0.30 × 0.21 × 0.19	0.20 × 0.13 × 0.12	0.84 × 0.39 × 0.16	0.26 × 0.21 × 0.10
Temperature (K)	293(2)	301.24	299.23	273.15
Crystal system	triclinic	triclinic	triclinic	orthorhombic
Space group	$P\bar{1}$	$P\bar{1}$	$P\bar{1}$	P_{bcn}
a (Å)	12.043(1)	13.5291(7)	13.737(1)	18.284
b (Å)	17.014(2)	16.0367(9)	16.191(1)	21.158(4)
c (Å)	17.162(2)	17.2368(9)	17.272(1)	18.284(3)
α (°)	101.787(3)	109.995(2)	110.347(2)	90
β (°)	100.776(3)	92.558(2)	93.031(2)	90
γ (°)	106.228(3)	103.732(2)	104.189(2)	90
V (Å ³)	3191.6(6)	3381.3(3)	3451.6(4)	7073.1(2)
Z	2	2	2	4
ρ _{calc} (g/cm ³)	1.604	1.643	1.952	1.579
M (mm ⁻¹)	2.171	2.360	6.633	1.944
Total data	45811	43290	41176	114916
unique data (R _{int})	15909 (0.0686)	13912 (0.0336)	14006 (0.0328)	7433 (0.0826)
2θ range (°)	5.628 to 56.788	5.586 to 52.98	5.998 to 52.79	5.89 to 53.428
Goodness of fit (GOF)	0.986	1.038	1.054	1.035
R ₁ /wR ₂ (I > 2σ(I))	0.0522/0.1087	0.0449/0.0887	0.0465/0.1094	0.0546/0.1140
R ₁ /wR ₂ (all data)	0.1186/0.1323	0.0763/0.0986	0.0745/0.1204	0.1005/0.1340

Table 7. Selected interatomic lengths (Å) and angles (°) for chapter 4.

Parameter	H-HIPPO	Cl-HIPPO	Br-HIPPO	Ph-HIPPO
Cu-- μ_3 -O(H))	1.938(3)-2.033(3)	1.963(2)- 2.050(3)	1.956(3)- 2.047(3)	1.977(3)- 2.022(3)
Cu-N (intra-trimer)	1.939(3)-1.970(3)	1.937(3)- 1.977(3)	1.929(4)- 1.979(4)	1.937(4)- 1.958(3)
Cu-N (inter-trimer)	1.962(3)-1.995(3)	1.979(3)-1.991(3)	1.976(4)- 1.987(4)	2.026(3)
Cu-Cl(terminal)	2.246(1),2.263(1)	2.246(1), 2.254(1)	2.244(1)- 2.247(1)	2.271(1)
Cu-- $\mu_{4/2}$ -Cl	2.584(1),2.869(1)	2.5598(9)- 3.038(1)	2.546(1)- 3.040(1)	2.6677(6), 2.941(2)
μ_3 -O(H)···(Cu3- plane)	0.590(3),0.607(4)	0.555(2), 0.608(3)	0.561(3), 0.605(3)	0.588(4)
Cu···Cu (inter-rear)	3.6044(8), 3.6536(8)	3.6288(6), 3.6321(6)	3.6301(7), 3.6352(7)	3.607(1)
Cu···Cu (inter-front)	7.203(1)	6.8924(6)	6.8681(8)	5.235(1)
Cu···Cu (intra-trimer)	3.1027(6)- 3.4066(8)	3.1718(7)- 3.4231(7)	3.1745(9)- 3.4125(9)	3.2235(9)- 3.4078(9)
O···O	5.906(5)	5.785(3)	5.783(4)	5.819(8)
Cu-Cl-Cu (intra-trimer)	69.23(3),71.73(3)	68.83(1)- 71.69(3)	68.78(3)- 71.87(3)	69.97(3)
Cu-Cl-Cu (inter-trimer)	81.79(3),86.16(3)	77.45(0), 88.13(3)	77.89(3)- 88.01(4)	71.65(2), 75.65(5)
Cu-Cl-Cu (trans)	122.68(4), 125.40(4)	121.95(4), 122.33(8)	122.37(4), 122.49(5)	159.51(5)
Cu-O-Cu	100.6(1)-118.1(1)	103.9(1)- 119.4(1)	103.9(1)- 119.6(2)	105.8(2)- 116.9(2)
N-Cu-N	153.8(1)-170.6(1)	147.8(1)- 169.6(1)	147.0(2)- 170.3(2)	163.9(2)- 172.4(2)
O-Cu-L	146.1(1)-176.0(1)	162.63(8)- 178.2(1)	159.5(1)- 178.6(2)	159.4(1)- 176.3(2)
Cu plane angle	73.9	66.8	66.1	31.2

4.4.2. NMR

4.4.2.1. NMR of H-HIPPO

The NMR spectrum of **H-HIPPO** (Figure 44) has peak integration 3:2:2:2:2:2:1:1:1 with the two additional peaks corresponding to the THF that is hydrogen-bonded to **H-Hippo**. The peak for H^c (36.87 ppm) envelopes the OH proton peak throughout the temperature range examined which leads to an integration of three protons. H^a and H^b (36.48, slopes are -0.023 and -0.087 respectively) are overlapped at RT and separate by -5°C. There are three peaks between 39 and 41 ppm at room temperature, which correspond to the THF (H^h and Hⁱ) and H^d, however when the temperature is decreased they become overlapped and appear as two peaks. H^f and H^g are assigned to 56.64 and 53.81 ppm, respectively. The last peak, H^e, is assigned to 52.91 ppm. The peaks were assigned based on distance from the bridging pyrazolates, as the spin density is through the two pyrazolates as opposed to three in previous literature,⁴⁹ which is assumed to push the relative shift farther upfield. The NMR spectra can be seen in Figure 44, with the proton names in Scheme 13. The shifts, slopes, and T₂ times are listed in Table 8.

Figure 44. Variable Low-Temperature NMR plot of **H-HIPPO**. Red, RT; green, -5 °C; blue, -15 °C, purple -25 °C.

Table 8. NMR shifts (ppm) of **H-HIPPO**, with T_2 times, shifts, and slope.

Proton	T_2 at RT (seconds)	Temperature				Slope (ppm/°C)
		20 °C	-5 °C	-15 °C	-25 °C	
H ^a	0.0849*	36.48	37.27	37.47	37.57	-0.023
H ^b	0.0849*	36.48	38.78	39.77	40.80	-0.087
H ^c	0.0353	36.87	38.50	39.19	39.87	-0.061
H ^d	0.0571	39.69	42.07	43.04	44.15	-0.090
H ^e	0.1208	52.91	56.61	58.22	59.95	-0.142
H ^f	0.0739	56.64	60.73	62.53	64.47	-0.157
H ^g	0.1083	53.81	57.61	59.27	61.03	-0.145
H ^h	0.0924	40.30	42.40	43.26	44.15	-0.078
H ⁱ	0.1653	40.64	42.34	42.97	43.57	-0.060

Scheme 13. Proton labels for **H-HIPPO**.

4.4.2.2. NMR of **Cl-HIPPO**

The ^1H NMR spectrum of **Cl-HIPPO** is similar to the one of **H-HIPPO**, however it has fewer peaks, due to the 4' position Cl and the lack of a THF molecule. The **Cl-HIPPO** shows a 2:2:2:2:1 ratio for the 18 protons a, c, d, f and the OH proton peak. The NMR spectrum of **Cl-HIPPO** shows five peaks at RT. There are two well-defined peaks at 58.72 ppm and 41.30 ppm, which have a 2:2 integration. There are also three overlapping peaks, around 39.5 ppm that integrates to 5 protons. These peaks resolve throughout the temperature range investigated. The presence of Cl in the 4' position allowed for easy assignment of the $\mu_3\text{-OH}$ group, as it only integrates to half of the other peaks, and its relative shift and slope confirm the assignment for **H-HIPPO**. The other protons are assigned by the distance from the bridging pyrazolates, in the same manner as **H-HIPPO**. This leads to H^a being the least shifted. The three signals at approximately 39 - 40.5

ppm are overlapping at RT but are resolved at -5°C. The NMR spectra can be seen in Figure 45. Proton labels are listed in Scheme 14. The shifts, slopes, and T₂ times are listed in Table 9.

Scheme 14. Proton labels for **CI-HIPPO**.

Table 9. NMR shifts (ppm) of **CI-HIPPO**, with T₂ times, shifts, and slope.

Proton	T2 at RT (seconds)	Temperature				Slope (ppm/°C)
		20 °C	-5 °C	-15 °C	-23 °C	
Ha	0.0628	39.71	40.77	41.08	41.23	-0.034
Hc	0.0365	39.90	42.47	43.49	44.48	-0.075
Hd	0.0483	41.41	43.49	44.27	44.97	-0.096
Hf	0.0628	58.68	62.80	64.52	66.12	-0.156
OH	0.0430	39.30	40.99	41.61	42.13	-0.060

Figure 45. Variable Low-Temperature NMR plot of **CI-HIPPO**. Red, RT; green, -5 °C; blue, -15 °C, purple -23 °C.

4.4.2.3. NMR of Ph-HIPPO

The NMR spectrum of **Ph-HIPPO** has many overlapping peaks in the aromatic region. It appears that many of the protons are not heavily affected by paramagnetic shift, and the resulting spectra are overlapping peaks and multiplets (Figure 46), with one peak around 38 ppm (Figure 47) that integrates to 5 protons. However, peak assignments and T_2 calculations were not possible.

Figure 46. Variable Low-Temperature NMR plot (downfield) of **Ph-HIPPO**. Red, RT; green, -5 °C; blue, -15 °C, purple -25 °C.

Figure 47. Variable Low-Temperature NMR plot (upfield) of **Ph-HIPPO**. Red, RT; green, -5 °C; blue, -15 °C, purple -25 °C.

4.4.2.4. NMR of [1]

The NMR spectrum of compound **[1]** shows most proton signals to be significantly shifted downfield compared to the parent Cu₆ **PPN[2]**. This shift could be attributed to the encapsulated F⁻ ion as **H-HIPPO** and **CI-HIPPO** are also shifted in the same direction, albeit to a smaller extent, which could be due to the partial encapsulation, the larger Cu-O-Cu angles, or the difference in electronegativity of the encapsulated ion. Compound **[1]** shows the OH proton peak at a very high shift, 70.69, in CH₂Cl₂ which with **H-HIPPO** and **CI-HIPPO** is the first time a OH proton signals has been observed in hexanuclear Cu complexes. Perhaps the most interesting information from the spectra is the slopes of H^c and H^d are slightly positive. The peaks were assigned using the RT spectrum and previous literature.⁵³ The differences in relative shift, slope, and T₂ are listed in Table 10 with the NMR spectra at the various temperatures shown in Figure 48 and Figure 49. The proton labels are listed in Scheme 15.

Table 10. NMR of **[1]**, with T₂ times, shifts, and slope.

Proton	T ₂ at RT (seconds)	Temperature				Slope (ppm/°C)
		20 °C	-5 °C	-15 °C	-25 °C	
H ^a	0.1745	47.03	50.22	51.38	52.61	-0.113
H ^b	0.1083	33.81	35.55	36.23	36.94	-0.063
H ^c	0.1795	7.67	7.64	7.64	7.63	0.001
H ^d	0.1745	8.07	8.06	8.04	8.03	0.001
H ^e	0.0751	17.18	17.97	18.29	18.65	-0.030
H ^f	0.1964	9.00	9.09	9.13	9.17	-0.003
OH	0.1102	70.69	74.53	76.01	77.56	-0.139

Figure 49. Variable Low-Temperature NMR plot (downfield) of compound **5**. Red, RT; green, -5 °C; blue, -15 °C, purple -25 °C.

Scheme 15. Proton labels for compound **1**.

4.4.2.5. NMR of PPN[2]

PPN[2] has shifts and slopes that differ significantly from the derivative compounds. Most likely due to the lack of encapsulated anion. The decrease in temperature shifts all the peaks downfield, although H^d and H^f are only very slightly shifted over the range. The shifts mirror **[1]**, which is explained by their location on the phenyl rings which are far from the paramagnetic centers and are unaffected by paramagnetism. However, H^c has the second largest slope, -0.107, which is in contrast to the most comparable compound, **[1]**, where the shift is 0.001. Peaks were assigned using RT spectra and literature assignments.⁴⁹ The proton labels are shown in Scheme 16, differences in relative shift, slope, and T_2 are listed in Table 11 with the NMR spectra at the various temperatures shown in Figure 50 and Figure 51.

Scheme 16. Proton labels for **PPN[2]**.

Table 11. NMR of **PPN[2]**, with T₂ times, shifts, and slope.

Proton	T ₂ at RT (seconds)	Temperature				Slope (ppm/°C)
		20 °C	-5 °C	-15 °C	-25 °C	
H ^a	0.2094	35.50	38.96	40.52	42.23	-0.135
H ^b	0.1337	18.17	19.70	20.42	21.24	-0.061
H ^c	0.2856	27.96	30.46	31.80	33.36	-0.107
H ^d	0.1904*	7.70	7.74	7.75	7.77	-0.001
H ^e	0.0582	11.82	12.18	12.35	12.53	-0.014
H ^f	0.1571*	8.01	8.12	8.16	8.18	-0.004

Figure 50. Variable Low-Temperature NMR plot (upfield) of **PPN[2]**. Red, RT; green, -5 °C; blue, -15 °C, purple -25 °C.

Figure 51. Variable Low-Temperature NMR plot (downfield) of **PPN[2]**. Red, RT; green, -5 °C; blue, -15 °C, purple -25 °C.

4.4.3. IR Analysis

4.4.3.1. IR of H-HIPPO

There are three absorptions of importance in the spectrum of **H-HIPPO** (Figure 52). Two absorptions are medium-strong absorptions at 282 cm^{-1} and 352 cm^{-1} which correspond to the terminal chloride and Cu-N stretch respectively. There is another medium strength absorption at 500 cm^{-1} that corresponds to another Cu-N stretch. The peaks correspond well to the range described in previous literature. Another two peaks at 542 cm^{-1} and 528 cm^{-1} correspond to Cu-O stretch and

vibrations. This is confirmed by ^{18}O substitution (selected portion of IR shown in Figure 53) and DFT calculations.

Figure 52. Far-IR of **H-HIPPO**.

Figure 53. IR Cu-O stretching bands for ^{16}O - and ^{18}O -**H-HIPPO**.

4.4.3.2. Far-IR of **CI-HIPPO**

The IR spectra of **CI-HIPPO** (Figure 54) shows an absorption at 295 cm^{-1} , which corresponds to Cu-N stretch and is confirmed by the calculations (simulated spectra shown in Figure 55) which places two absorptions at 301.71 cm^{-1} and 306.46 cm^{-1} , the latter is coupled with pyrazolate distortions. Calculations place

two more Cu-N absorptions at 504.16 cm^{-1} and 506.22 cm^{-1} which have merged into one observed peak that appears at 529 cm^{-1} . The peak at 319 cm^{-1} , which is a Cu-O stretch coupled with pyrazole distortions, DFT places at 309.86 cm^{-1} . Additionally, calculations show that there are Cu-O absorptions at 346.63 , 452.77 and 455.12 . These peaks are seen in the experimental spectrum as a weak peak at 337.3 and a broad peak with a shoulder at 483 . At 263 cm^{-1} a Cu-Cl absorption coupled with Cu-N is observed, DFT places this at 239.00 . Additionally, there is a shoulder at 280 cm^{-1} which corresponds to two calculated stretches, 256.57 and 256.92 cm^{-1} , associated with Cu-Cl, Cu-N coupled modes. At 250 cm^{-1} , at the edge of the capabilities of our instrument, the start of an absorption is noticeable, DFT indicates that the peak corresponds to Cu-Cl and calculates two frequencies at 208.89 and 212.47 cm^{-1} .

Figure 54. Far-IR of **CI-HIPPO**.

Figure 55. Simulated spectra of **CI-HIPPO** determined by DFT.

4.4.3.3. Far-IR of Ph-HIPPO

In the spectrum of **Ph-HIPPO** (Figure 56), calculations place two peaks at 277.58 cm^{-1} and 277.91 cm^{-1} observed as a single peak at 280 cm^{-1} , corresponding to a Cu-N stretch. Calculated frequencies at 337.73 cm^{-1} , 339.5 cm^{-1} , 344.86 cm^{-1} , and 346.22 cm^{-1} , can be seen as a peak at 341.5 cm^{-1} with a shoulder at 353 cm^{-1} . Absorptions associated with Cu-O bonds are seen 374.7 cm^{-1} and strong absorption at 430 cm^{-1} with a slight inflection at 442.3 cm^{-1} , the calculated absorptions are at 440.58, 445.16, which have pyrazolate distortions, and 447.03 cm^{-1} . The Cu-Cl absorption that appears at 251.4 cm^{-1} corresponds to a calculated frequency coupled with pyrazolate distortions at 242.89 cm^{-1} . There is also a small, rounded absorption at 305 cm^{-1} , which corresponds to Cu-Cl, Cu-O coupled modes calculated to be at 311.87 cm^{-1} and 313.77 cm^{-1} .

Figure 56. Far-IR of **Ph-HIPPO**.

4.4.3.4. Far-IR of [1]

The spectra of **[1]** (Figure 57), lacks an absorption in the terminal Cl range, which is to be expected. The Cu-N stretch appears at 364 cm^{-1} . A much weaker absorption at 534 cm^{-1} corresponding the Cu-O bonds.

Figure 57. Far-IR of **[1]**.

4.4.3.5. Far-IR of PPN[2]

The spectrum of **PPN[2]** (Figure 58) has peaks associated with Cu-N stretches at 282 (a small peak), 314.5 (a shoulder), and 353 cm^{-1} with an inflection at 341 cm^{-1} . The calculated peaks for Cu-N stretch are shown at 294.84 cm^{-1} , 313.83 cm^{-1} ,

341.85 cm^{-1} , 345.7 cm^{-1} , 347.59 cm^{-1} , and 353.83 cm^{-1} . The peaks at 345.7 cm^{-1} , 347.59 cm^{-1} , and 353.83 cm^{-1} are seen as one large broad peak in the experimental spectrum. There are four Cu- μ_3 -O absorptions seen in the experimental spectrum. One medium absorption at 433 cm^{-1} , and three strong absorptions at 501 cm^{-1} , 534.9 cm^{-1} , and 546.9 cm^{-1} . The calculated absorptions for the Cu- μ_3 -O stretches are at 427.05 cm^{-1} , 487.28 cm^{-1} , 506.2 cm^{-1} , and 520.2 cm^{-1} , respectively.

Figure 58. Far-IR of **PPN[2]**.

4.4.3.6. IR Compared to Literature Values

Overall, the IR absorption frequencies are in agreement with previous literature, which place the vibration frequency for Cu-Cl bonds at 218-345 cm^{-1} ,^{181,182,185,187,191,205,206} Cu-N at 183-421 cm^{-1} ,^{181,206,207} and Cu-O at 529-594 cm^{-1} .^{182,183,187} Observed and calculated data for **CI-HIPPO**, **Ph-HIPPO**, and **PPN[2]** are listed on Table 12.

Table 12. Comparison of calculated absorptions and observed absorptions.
* indicates coupling with pz⁻ deformation.

Bond	CI-HIPPO		Ph-HIPPO		PPN[2]	
	Calculated	Exptl	Calculated	Exptl	Calculated	Exptl
Cu-N	227.81	B/R	277.58 277.91	280	233.22 (bridging) 237.16	B/R
Cu-N	301.71 306.46*	295	337.73 339.5 344.86 346.22	341.5 353	294.84	282
Cu-N	504.16 506.22	529.3	N/A	N/A	313.83 (bridging)	314.5
Cu-N	N/A	N/A	N/A	N/A	341.58 345.7 347.59 353.83	341 353
Cu-O	N/A	N/A	375.9: 379.71	374.7	427.05	433
Cu-O	309.86*	319	440.58* 445.16* 447.03	430 442.3	487.28	501
Cu-O	346.63	337.3	N/A	N/A	506.2	534.9
Cu-O	452.77 455.12	457:483	N/A	N/A	520.2	546.9
Cu-Cl	208.89 212.47	250	264.9*	251.4	N/A	N/A
Cu-O Cu-N	223.75	N/A	N/A	N/A	N/A	N/A
Cu-Cl Cu-N	208.89	B/R	234.4 237.3 242.89	B/R	N/A	N/A
Cu-Cl Cu-N	239.0	262.8	N/A	N/A	N/A	N/A
Cu-Cl Cu-N	256.57 256.92	280	N/A	N/A	N/A	N/A
Cu-Cl Cu-O	N/A	N/A	311.87 313.77	305	N/A	N/A

4.4.4. Magnetism

The magnetic properties were deeply studied for selected compounds, **CI-HIPPO** and **Ph-HIPPO**, by acquiring temperature and field dependent magnetization data as showed in Figure 59. The room temperature values of the effective magnetic

moment (μ_{eff}) are $4.28 \mu_B$ for **CI-HIPPO** and $3.85 \mu_B$ for **Ph-HIPPO** and these values are close to the theoretical values of $4.24 \mu_B$ for six non-interacting Cu(II) ions with $S_i = 1/2$ and $g_i = 2.0$. The μ_{eff} is continuously decreasing on lowering the temperature down to ca 50 K and then significant drop of μ_{eff} is observed down to values of $0.47 \mu_B$ for **CI-HIPPO** and $1.67 \mu_B$ for **Ph-HIPPO** at $T = 1.9$ K. This magnetic behaviour reflects strong antiferromagnetic exchange interactions within each $\text{Cu}_3(\mu_3\text{-OH})$ triangles resulting in $S_{\text{triangle}} = 1/2$ spin state and further weak antiferromagnetic exchange between these two triangles leading to $S = 0$ molecular ground state. Here we can expect that difference on low temperature magnetism is related to strength of the antiferromagnetic exchange between each $\text{Cu}_3(\mu_3\text{-OH})$ triangles in **CI-HIPPO** and **Ph-HIPPO** reflecting their structural differences. Moreover, the triangular systems of Cu(II),^{48,53,208} Ni(II),²⁰⁹ Fe(III)²¹⁰ complexes possessing large antiferromagnetic exchange are well known to exhibit also the magnetic anisotropy of $S = 1/2$ ground state due to the antisymmetric exchange interaction also known as Dzyaloshinsky–Moriya interaction.²¹¹ Therefore, the following spin Hamiltonian was postulated to interpret magnetism of **CI-HIPPO**

$$\hat{H} = -J_1(\mathbf{S}_1 \cdot \mathbf{S}_2 + \mathbf{S}_1 \cdot \mathbf{S}_3 + \mathbf{S}_4 \cdot \mathbf{S}_5 + \mathbf{S}_4 \cdot \mathbf{S}_6) - J_{1b}(\mathbf{S}_2 \cdot \mathbf{S}_3 + \mathbf{S}_5 \cdot \mathbf{S}_6) - J_2(\mathbf{S}_2 \cdot \mathbf{S}_5 + \mathbf{S}_3 \cdot \mathbf{S}_6) + \mathbf{d} \cdot (\mathbf{S}_1 \times \mathbf{S}_2 + \mathbf{S}_2 \times \mathbf{S}_3 + \mathbf{S}_3 \times \mathbf{S}_1 + \mathbf{S}_4 \times \mathbf{S}_5 + \mathbf{S}_5 \times \mathbf{S}_6 + \mathbf{S}_6 \times \mathbf{S}_4) + \mu_B \mathbf{B} \cdot \mathbf{g} \cdot \sum_{i=1}^6 \mathbf{S}_i$$

Equation 6

and **Ph-HIPPO**

$$\hat{H} = -J_1(\mathbf{S}_1 \cdot \mathbf{S}_2 + \mathbf{S}_2 \cdot \mathbf{S}_3 + \mathbf{S}_{1'} \cdot \mathbf{S}_{2'} + \mathbf{S}_{2'} \cdot \mathbf{S}_{3'}) - J_{1b}(\mathbf{S}_1 \cdot \mathbf{S}_3 + \mathbf{S}_{1'} \cdot \mathbf{S}_{3'}) - J_2(\mathbf{S}_1 \cdot \mathbf{S}_{1'}) + \mathbf{d} \cdot (\mathbf{S}_1 \times \mathbf{S}_2 + \mathbf{S}_2 \times \mathbf{S}_3 + \mathbf{S}_3 \times \mathbf{S}_1 + \mathbf{S}_{1'} \times \mathbf{S}_{2'} + \mathbf{S}_{2'} \times \mathbf{S}_{3'} + \mathbf{S}_{3'} \times \mathbf{S}_{1'}) + \mu_B \mathbf{B} \cdot \mathbf{g} \cdot \sum_{i=1}^6 \mathbf{S}_i \quad \text{Equation 7}$$

where the isotropic exchange parameters J_1 and J_{1b} are characterizing magnetic coupling with $\text{Cu}_3(\mu_3\text{-OH})$ triangles and J_2 between two triangles, the antisymmetric exchange is characterized by vector parameter $\mathbf{d} = (d_x, d_y, d_z)$, which was simplified thanks to Moriya symmetry rules²¹² to $\mathbf{d} = (0, 0, d_z)$. Next, the molar magnetization was calculated from the partition function Z as

$$M_{\text{mol}}(\theta, \varphi) = N_A kT \frac{\partial \ln Z}{\partial B_{\theta, \varphi}} \quad \text{Equation 8}$$

and finally, the averaged molar magnetization was calculated as an integral average to properly simulated powder sample magnetic measurements

$$M_{\text{mol}} = \frac{1}{4\pi} \int_{\varphi=0}^{2\pi} \int_{\theta=0}^{\pi} M_{\text{mol}}(\theta, \varphi) \sin \theta \, d\theta d\varphi \quad \text{Equation 9}$$

where the magnetic field vector is defined with the help of the spherical coordinates.

The both temperature and field dependent magnetization data were fitted simultaneously and the best-fitted parameters for **CI-HIPPO** are $J_1 = -240 \text{ cm}^{-1}$, $J_{1b} = -225 \text{ cm}^{-1}$, $J_2 = -26 \text{ cm}^{-1}$, $|d_z| = 17.0 \text{ cm}^{-1}$ with an isotropic g -factor $g = 2.65$, and for **Ph-HIPPO** are $J_1 = -208 \text{ cm}^{-1}$, $J_{1b} = -145 \text{ cm}^{-1}$, $J_2 = -17 \text{ cm}^{-1}$, $|d_z| = 34.3 \text{ cm}^{-1}$ with an isotropic g -factor $g = 2.22$ (Figure 59). Also, the temperature-

independent magnetism was treated with fixed value of $\chi_{\text{TIP}} = 0.88 \times 10^{-9} \text{ m}^3 \text{mol}^{-1}$ per copper atom.

Figure 59. Magnetic properties of compounds **2** and **4**. Each plot shows the temperature dependence of the effective magnetic moment and in the inset, the isothermal magnetizations measured at $T = 2, 5$ and 10 K are given. Experimental data – empty symbols, full red lines – calculated data with spin Hamiltonian in eq.1 and parameters in the text.

The analysis of the experimental data confirmed strong antiferromagnetic coupling within $\text{Cu}_3(\mu_3\text{-OH})$ triangles mediated by pyrazolato and hydroxo bridging ligands. Interestingly, there is sizeable difference in J_2 between **Cl-HIPPO** and **Ph-HIPPO**, which most probably can be assigned to difference in the electronic properties of pyrazolato bridging ligands with chloro or phenyl substituent.

4.4.5. Theoretical calculations

To support our analysis of the magnetic data of **Cl-HIPPO** and **Ph-HIPPO**, the Density Functional Theory (DFT) theoretical calculations were performed aiming to estimate the isotropic exchange parameters J . Based on our previous

experience with polynuclear systems, B3LYP functional was chosen and the Ruiz's approach for calculation of J 's were followed employing the broken-symmetry DFT calculations with computational package ORCA 3.0.

In case of compound **2**, the spin Hamiltonian was formulated as

$$\hat{H} = -J_{12}(\mathbf{S}_1 \cdot \mathbf{S}_2) - J_{13}(\mathbf{S}_1 \cdot \mathbf{S}_3) - J_{23}(\mathbf{S}_2 \cdot \mathbf{S}_3) - J_{45}(\mathbf{S}_4 \cdot \mathbf{S}_5) - J_{46}(\mathbf{S}_4 \cdot \mathbf{S}_6) - J_{56}(\mathbf{S}_5 \cdot \mathbf{S}_6) - J_{25}(\mathbf{S}_2 \cdot \mathbf{S}_5) - J_{36}(\mathbf{S}_3 \cdot \mathbf{S}_6)$$

Equation 10

and several broken-symmetry spin states were calculated and compared to the energy of high-spin state, which enabled us to derive these equations for the values of the isotropic exchange parameters in equation 11 as

$$\begin{aligned} J_{12} &= (\Delta_1 - \Delta_3 - \Delta_5 + \Delta_{25} + \Delta_{123})/2 \\ J_{23} &= (-\Delta_1 + \Delta_2 + \Delta_3 - \Delta_{123})/2 \\ J_{13} &= (\Delta_1 + \Delta_3 + \Delta_5 - \Delta_{25} - \Delta_{123})/2 \\ J_{45} &= (-\Delta_2 + \Delta_4 - \Delta_6 + \Delta_{25} + \Delta_{123})/2 \\ J_{56} &= (-\Delta_4 + \Delta_5 + \Delta_6 - \Delta_{123})/2 \\ J_{46} &= (-\Delta_2 - \Delta_4 - \Delta_6 + \Delta_{25} + \Delta_{123})/2 \\ J_{25} &= (\Delta_2 + \Delta_5 - \Delta_{25})/2 \\ J_{36} &= (-\Delta_2 - \Delta_5 + \Delta_{25} + 2\Delta_{123})/2 \end{aligned}$$

Equation 11

As a result, we obtained the isotropic exchange parameters of **CI-HIPPO** as $J_{12} = -100 \text{ cm}^{-1}$, $J_{23} = -86 \text{ cm}^{-1}$, $J_{13} = -240 \text{ cm}^{-1}$, $J_{45} = -171 \text{ cm}^{-1}$, $J_{56} = -111 \text{ cm}^{-1}$, $J_{46} = -168 \text{ cm}^{-1}$, $J_{25} = -23.4 \text{ cm}^{-1}$, $J_{36} = -38.9 \text{ cm}^{-1}$.

The similar procedure was applied to compound **Ph-HIPPO** with spin Hamiltonian defined as

$$\hat{H} = -J_{12}(\mathbf{S}_1 \cdot \mathbf{S}_2 + \mathbf{S}_{1'} \cdot \mathbf{S}_{2'}) - J_{13}(\mathbf{S}_1 \cdot \mathbf{S}_3 + \mathbf{S}_{1'} \cdot \mathbf{S}_{3'}) - J_{23}(\mathbf{S}_2 \cdot \mathbf{S}_3 + \mathbf{S}_{2'} \cdot \mathbf{S}_{3'}) - J_{11'}(\mathbf{S}_1 \cdot \mathbf{S}_{1'})$$

Equation 12

and the relationships for J 's were derived as

$$\begin{aligned} J_{12} &= (\Delta_{11'} + \Delta_{22'} - \Delta_{33'})/4 \\ J_{23} &= (-\Delta_{11'} + \Delta_{22'} + \Delta_{33'})/4 \\ J_{13} &= (\Delta_{11'} - \Delta_{22'} + \Delta_{33'})/4 \\ J_{11'} &= \Delta_{123} \end{aligned}$$

Consequently, the isotropic exchange parameters were derived as $J_{12} = -174 \text{ cm}^{-1}$, $J_{23} = -124 \text{ cm}^{-1}$, $J_{13} = -141 \text{ cm}^{-1}$, $J_{11'} = -221 \text{ cm}^{-1}$.

Generally, the DFT calculations showed large antiferromagnetic exchange within each $\text{Cu}_3(\mu_3\text{-OH})$ triangles both in **CI-HIPPO** and **Ph-HIPPO**. Moreover, there is obvious magneto-structural correlation between Cu-O-Cu angle and the value of the respective J -parameter as showed in Figure 60 and it holds that the larger angle which Cu-O-Cu, the stronger antiferromagnetic exchange was calculated. Next, the values of inter-triangular exchange (J_{25} and J_{36}) are quite reasonable for **CI-HIPPO** and follows the analysis of the experimental magnetic data, however, the situation is very different for **Ph-HIPPO**, where $J_{11'}$ adopted very large value incompatible with the experiment.

Figure 60. The magneto-structural correlation between the angle Cu-O-Cu in $\text{Cu}_3(\mu_3\text{-OH})$ triangles of **Cl-HIPPO** and **Ph-HIPPO** extracted from X-ray structures and magnetic coupling parameters J calculated by DFT using B3LYP functional.

4.5. Conclusions

Here the first four open cage Cu_6 motifs are reported. This shows that Cu-pyrazolate motifs can be exploited in new ways. Furthermore, we characterized for the first time Cu-O, Cu-N, and Cu-Cl bonds in Cu-pyrazolate compounds. Magnetic susceptibility was measured for the new motifs and will be compared to other Cu_6 compounds and supramolecular Cu_3 compounds to further understand how the magnetism is affected by the distance, connectivity, and angle of the Cu_3 subunits. Variable low-temperature NMR studies were also conducted.

5. Summary and Future Work

The purpose of this project was to determine if a Cu₆ motif could function as a water oxidation catalyst. Toward this end, Cu₆ has been oxidized and characterized in a mixed-valent state. The data support the conclusion that a Cu₆ motif could be used as a WOC, due to the evidence that significant spin density is located on the μ_3 -oxo ligands. Furthermore, the crystal structures of the parent Cu₆ and the MV-Cu₆ show that upon oxidation, the central μ_3 -O ligands move ~ 0.3 Å closer.

The attempted synthesis of an elusive positively charged Cu₃ compound with 4-formyl-pzH and terminal pyridine led to novel Cu₃ coordination polymers with rare Cu-aldehyde coordination and new topology.

Finally, two new open-cage Cu₆ motifs have been synthesized and characterized. The open-cage motifs were characterized using variable low-temperature NMR, magnetic susceptibility, and IR spectroscopy. The IR-spectra have been used to characterize the various copper bonds that have never been investigated in copper-pyrazolate motifs.

In future studies, it will be vital to synthesize a flexible rigid Cu₆ motif. While such complexes eluded this study, there are many synthetic techniques that could still be employed, such as using the Cu-CPs presented here as a starting material for the flexible Cu₆ motifs. It was demonstrated that with coordinating solvents, the coordination polymers can be broken apart into the Cu₃ subunits.

After the synthesis for flexible a Cu₆ has been perfected, the synthesis should be modified to allow for a flexible two pillar an open-cage variation that will facilitate the release of the generated O₂²⁻ ion.

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Kreiger, D. I., Mathivathanan, L., Herchel, R., and Raptis, R. G., (2012) *Magnetic and Structural Properties of Open-Caged Hexanuclear Copper Pyrazolato Complexes*. Poster presented at FAME.

Kreiger, D. I., Mathivathanan, L., Herchel, R., and Raptis, R. G., (2019) *Synthesis, structure and magnetic studies of novel open-cage hexanuclear copper pyrazolato complexes*. Paper presented at National ACS, Orlando.

Kreiger, D. I., Mathivathanan, L., and Raptis, R. G., Coordination polymers based on pyrazole-4-carboxaldehyde-containing Cu₃N₆ metallacycles as building units. *CrystEngComm* 2019, 21, 3047-3055. DOI: 10.1039/c9ce00421a.