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Revised Manuscript

**Doubly Phenoxide-Bridged Binuclear Copper(II) Complexes with ONO
Tridentate Schiff Base Ligand: Synthesis, Structural, Magnetic and
Theoretical studies**

**Néstor Novoa^{a,b}, Frédéric Justaud^b, Paul Hamon^b, Thierry Roisnel^c, Olivier Cador^b,
Boris Le Guennic^b, Carolina Manzur^a, David Carrillo^{a,*}, Jean-René Hamon^{b,*}**

^a *Laboratorio de Química Inorgánica, Instituto de Química, Pontificia Universidad
Católica de Valparaíso, Campus Curauma, Avenida Universidad 330, Valparaíso, Chile*

^b *UMR 6226 «Institut des Sciences Chimiques de Rennes», CNRS-Université de Rennes 1,
Campus de Beaulieu, 35042 Rennes Cedex, France*

^c *UMR 6226 «Institut des Sciences Chimiques de Rennes», CNRS-Université de Rennes 1,
Centre de Diffractométrie X, Campus de Beaulieu, 35042 Rennes Cedex, France*

Dedicated to Dr. Claude Lapinte for his outstanding contribution to
organometallic molecular wires

- Corresponding authors. *Phone:* 56 32 227 49 14, *E-mail address:* david.carrillo@ucv.cl (D. Carrillo); *Phone:* 33 223 23 59 58, *E-mail address:* jean-rene.hamon@univ-rennes1.fr (J.-R. Hamon).

Abstract

The tridentate ONO-donor Schiff base ligand H₂L, derived from the condensation of 1-anisyl-1,3-butanedione and 2-aminophenol, was generated *in situ* and reacted with Cu(NO₃)₂•3H₂O to yield two doubly phenoxo bridged di-copper(II) complexes depending on the nitrogenous base used. [Cu₂L₂] (**1**) is obtained in 85% and 75% yield in the presence of pyridine or 4-picoline, respectively, and [(py-*t*Bu)₂Cu₂L₂] (**2**) is isolated in 75% yield in the presence of 4-*tert*-butylpyridine. Compounds **1** and **2** were characterized in the solid-state by elemental analysis and FT-IR spectroscopy. Single crystal X-ray diffraction study reveals that in **1** the two four-coordinated copper atoms adopt a square planar geometry, whereas in **2** each Cu(II) metal ion shows a five coordinate square pyramidal (ONO,N + O) geometry. In each dimer, two μ-phenolic oxygen atoms bridge the two half-units forming a planar Cu₂O₂ core. EPR studies in fluid solutions indicate that the dimeric structure of **1** and **2** is destroyed upon dissolution. In the solid-state, **1** is EPR silent, whereas **2** presents an unresolved broad resonance (ΔH peak-to-peak = 71.5 G) with g = 2.071 at 298 K, along with the triplet state (S = 1) signature at g = 4.181. Variable temperature (2-300 K) magnetic susceptibility measurements exhibit strong antiferromagnetic interactions between the Cu(II) centers with a J value of -397 cm⁻¹ for **1**, while no interaction operates between the two spins localized on Cu(II) metal ions in **2**. Ab initio calculations were also performed to supplement the experimental results.

Keywords: Copper(II); ONO Tridentate Schiff base ligand; magnetic properties; structure determination; EPR study; Ab initio calculation.

1. Introduction

In recent years, there has been continuing interest in using bridging ligands in the synthesis of polynuclear complexes of paramagnetic transition metal ions. In particular, ligands which contain potentially bridging phenoxo, alkoxo or hydroxo oxygen and nitrogen donor atoms have been widely employed to build up multinuclear copper(II) complexes [1-5]. Such symmetrically or asymmetrically dibridged complexes with μ -phenoxo, μ -alkoxo or μ -hydroxo di-copper(II) core have been the subject of a considerable amount of work in terms of correlating structure and magnetic properties [1,2,7-11]. Moreover, doubly phenoxo bridged binuclear Cu(II) complexes have also considerable interest as they provide the simplest case of magnetic interaction including only two unpaired electrons [2,6,11,12]. And it is now established that the major factor controlling the singlet-triplet energy gap, J , between the metal centers is the Cu-O-Cu angle [8,13-15]. Di-copper(II) derivatives are also of importance as precursors in the chemistry of supramolecular [16] and discrete molecular high-nuclearity copper(II) complexes [17-21].

Diprotic Schiff bases are easily prepared by 1:1 condensation of appropriate salicylaldehyde or β -diketone reagents with appropriate amino alcohol substrates under mild conditions. Upon double deprotonation, they form a group of di-negative tridentate O,N,O-donor ligands that react readily with cupric salts to produce variable-nuclearity compounds depending on the coligand [19,22]. Thus, a N-donor monodentate coligand results in the formation of neutral mononuclear square planar ternary copper(II) complexes while square pyramidal Cu(II) compounds are obtained in the presence of a N-N chelate coligand [19,23-27]. Moreover, the very popular bridging ligand 4,4'-bipyridine has been used also to construct homobinuclear copper(II) complexes [28], acting as a linear spacer between the mononuclear units, thus allowing the investigation of the intramolecular magnetic interaction between the magnetic centers [29].

We were interested in preparing neutral ternary mononuclear metal complexes of the type [LCu(II)(py)] with differently substituted di-anionic tridentate ONO Schiff base ligands (L^{2-}) and pyridine as ancillary ligand, as building blocks for the construction of novel self-assembled push-pull systems. However, it turned out that the synthesis using three closely related nitrogenous bases as coligand, leads to the formation of two different dimeric Schiff base complexes than those expected. Herein, we report on the synthesis, spectral and structural characterization, magnetic properties and ab initio study of both compounds formulated as [Cu₂L₂] (**1**) and [(py-*t*Bu)₂Cu₂L₂] (**2**), where H₂L is the mono-condensation

product of 1-anisyl-1,3-butanedione and 2-aminophenol (see Scheme 1). An EPR study of both complexes was also carried out.

2. Experimental

2.1. Materials

Manipulations of air-sensitive compounds were performed under Argon atmosphere using standard Schlenk techniques. Solvents were dried and distilled according to standard procedures [30]. 4-methoxyacetophenone, potassium *tert*-butoxide, 2-aminophenol, pyridine, 4-*tert*-butylpyridine, 4-picoline, Ethyl acetate (99%) and copper(II) nitrate trihydrate were purchased from commercial sources and used without further purification. Compound 1-anisyl-1,3-butanedione was synthesized according to the literature procedure [31].

2.2. Characterization and instrumentation.

Infrared spectra were recorded on a Perkin-Elmer, Model Spectrum One, FT-IR spectrophotometer as KBr disks in the 4000 to 400 cm^{-1} range. Elemental analyses were conducted on a Thermo-FINNIGAN Flash EA 1112 CHNS/O analyzer by the Microanalytical Service of the CRMPO at the University of Rennes 1, France. Electron paramagnetic resonance (EPR) spectra were recorded at 298 K with a Bruker EMX-8/2.7 (X-band) spectrometer. The temperature dependences of the magnetizations for powdered samples have been measured with a SQUID magnetometer (Quantum design MPMS-XL5) operating between 2 and 300 K at a constant field of 2 kOe. The experimental data have been corrected from the sample holder diamagnetism and the intrinsic diamagnetism estimated from the Pascal's tables [32]. Melting points were determined in evacuated capillaries on a Kofler Bristoline melting point apparatus and were not corrected.

2.3. Preparations

2.3.1. Preparation of $[\{4\text{-MeO-C}_6\text{H}_4\text{-C(O)CH=C(CH}_3\text{)N-}o\text{-C}_6\text{H}_4\text{-O}\}\text{Cu}]_2$ (**1**)

A Schlenk tube was charged with a magnetic stir bar, 200 mg (1.04 mmol) of 1-anisyl-1,3-butanedione, 114 mg (1.04 mmol) of 2-aminophenol and 10 mL of toluene, and the mixture was refluxed for 2 hours. After cooling, a yellow solid deposited. The precipitate was filtered off and washed with a petroleum ether: diethyl ether (1:1 v:v) mixture. Then, 292 mg (2.6 mmol) of potassium *tert*-butoxide and 2 mL of THF were added and the reaction mixture was stirred at r.t. for 20 min, upon which time it turned dark-red. Pyridine (160 μL , 2.08 mmol) was then added, before dropwise addition of a solution of $\text{Cu}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O}$ (252 mg, 1.04

mmol) in 5 mL of THF. The stirring was continued overnight. The reaction was quenched with 15 mL of EtOH, releasing a dark-green microcrystalline precipitate that was filtered off and washed with cold EtOH (3 x 10 mL) and diethyl ether (4 x 10 mL). The dark-green solid residue was dried under vacuum for 2 hours and dissolved in dichloromethane. The solution was subjected to slow evaporation for 5 days, affording 300 mg (0.44 mmol, 85% Yield) of black microcrystals. A crystal from this crop was used for X-ray structure determination. M.p. 215-217 °C. Anal. Calcd for C₃₄H₃₀Cu₂N₂O₆ (681.71 g mol⁻¹): C, 59.21; H, 4.38; N, 4.06. Found: C, 59.15; H, 4.33; N, 4.16%. Under the same conditions, complex **1** was isolated in 75% yield when its preparation was carried out with 4-picoline (185 µL, 2.08 mmol) instead of pyridine.

2.3.2. Preparation of [{4-MeO-C₆H₄-C(O)CH=C(CH₃)N-*o*-C₆H₄-O}Cu(NC₅H₄-4-CMe₃)]₂ (**2**)
This complex was synthesized following a similar procedure to that described above for the preparation of **1**, using in this case, 200 mg (1.04 mmol) of 1-anisyl-1,3-butanedione, 114 mg (1.04 mmol) of 2-aminophenol, 292 mg (2.6 mmol) of potassium *tert*-butoxide, 0.44 mL (3.0 mmol) of 4-*tert*-butylpyridine and 300 mg (1.24 mmol) of Cu(NO₃)₂ • 3H₂O. Recrystallization by slow evaporation of a saturated CH₂Cl₂ solution yielded 506 mg (75%) of a dark-green crystalline solid. A crystal from this crop was used for X-ray structure determination. M.p. 171-173 °C. Anal. Calcd for C₅₂H₅₆Cu₂N₄O₆ (960.12 g mol⁻¹): C, 65.05; H, 5.88; N, 5.84. Found: C, 64.74; H, 5.44; N, 5.58%.

2.4. X-ray crystal structure determination

Well-shaped single crystals of **1** and **2** of suitable dimensions were coated in Paratone-N oil, mounted on a Kaptan loop and transferred to the cold gas stream of the cooling device. Intensity data were collected at T = 150(2) K on a APEXII, Bruker-AXS diffractometer, Mo-Kα radiation (λ = 0.71073 Å), equipped with a bidimensional CCD detector and were corrected for absorption effects using multiscanned reflections. The two structures were solved by direct methods using the *SIR97* program [33], and then refined with full-matrix least-square methods based on *F*² (*SHELXL-97*) [34] with the aid of *WINGX* program [35]. All non-hydrogen atoms were refined with anisotropic atomic displacement parameters. All the hydrogen atoms were placed in their geometrically idealized positions and constrained to ride on their parent atoms. A summary of the details about crystal data, collection parameters and refinement are documented in Table 1, and additional crystallographic details are in the CIF files. ORTEP views are drawn using *Olex2* software [36].

Table 1 Crystallographic data, details of data collection and structure refinement parameters for compounds **1** and **2**

	1	2
Empirical formula	C ₃₄ H ₃₀ Cu ₂ N ₂ O ₆	C ₅₂ H ₅₆ Cu ₂ N ₄ O ₆
Formula mass, g mol ⁻¹	689.68	960.09
Collection T, K	150(2)	150(2)
Crystal system	Monoclinic	Triclinic
space group	<i>P</i> 2 ₁ / <i>c</i>	<i>P</i> $\bar{1}$
a (Å)	9.0297(2)	9.4434(3)
b (Å)	14.3356(4)	11.3431(4)
c (Å)	11.4572(3)	11.4661(4)
α (°)	90	101.758(2)
β (°)	109.8300(10)	109.5860(10)
γ (°)	90	92.739(2)
V (Å ³)	1395.15(6)	1123.86(7)
Z	2	1
D _{calcd} (g cm ⁻³)	1.642	1.419
Crystal size (mm)	0.15 x 0.07 x 0.04	0.41 x 0.11 x 0.06
Crystal color	black	black
F(000)	708	502
abs coeff (mm ⁻¹)	1.577	1.002
θ range (°)	3.41 to 27.48	2.96 to 27.48
range h,k,l	-11/11, -17/18, -12/14	-12/12, -13/14, -14/14
No. total refl	12339	18161
No. unique refl	3183	5123
Comp. to θ_{\max} (%)	99.6	99.4
Max/min transmission	0.939/0.856	0.942/0.807
Data/Restraints/Parameters	3183/0/201	5123/0/294
Final R	R ₁ = 0.0285	R ₁ = 0.0286
[I > 2 σ (I)]	wR ₂ = 0.0691	wR ₂ = 0.0712
R indices (all data)	R ₁ = 0.0378 wR ₂ = 0.0736	R ₁ = 0.0331 wR ₂ = 0.0737
Gof / F ²	1.016	1.055
Largest diff. Peak/hole (eÅ ⁻³)	0.430/-0.324	0.365/-0.406

2.5. Computational details

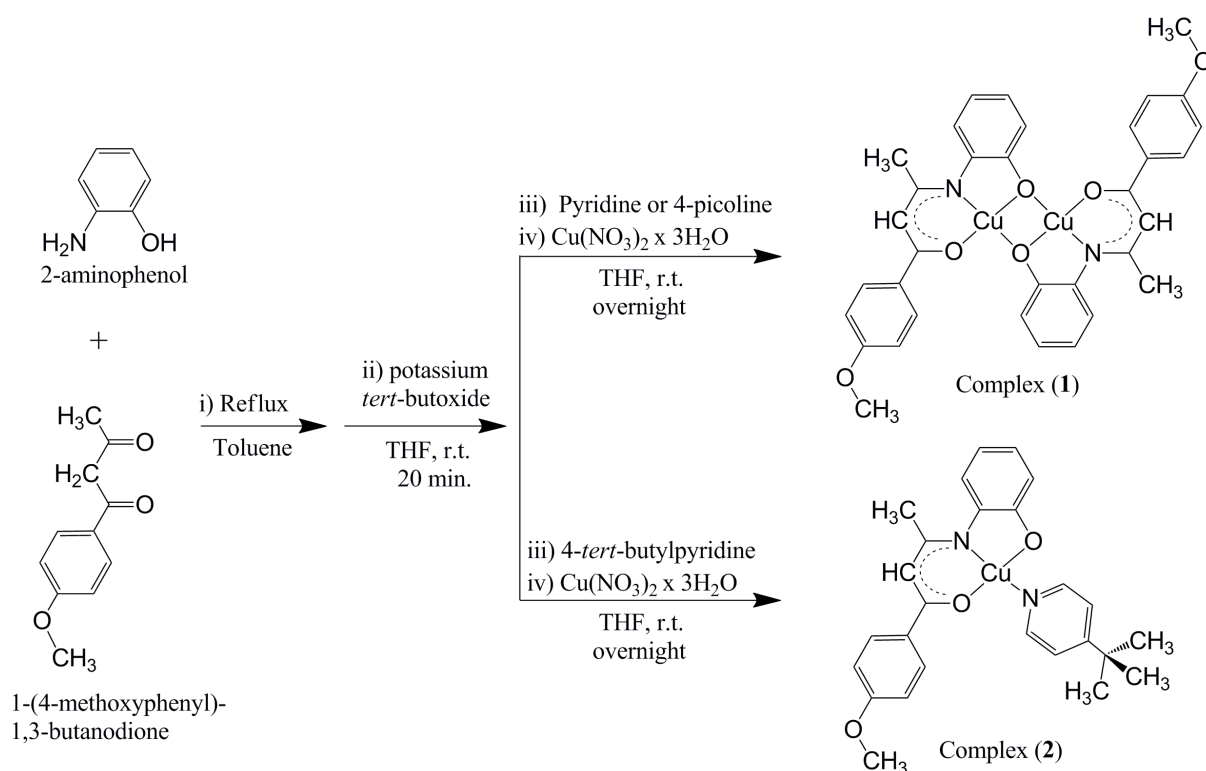
DFT calculations [37] were carried out with the Amsterdam Density Functional (ADF) program [38]. The Becke approximation for exchange [39] and the Perdew expression for correlation [40] (BP) has been chosen with the TZ2P basis set [41]. The optimized geometries of both compounds **1** and **2** were characterized as true minima on the potential energy surfaces using vibrational frequency calculations.

To compute exchange couplings in **1** and **2**, Complete active space self-consistent field (CASSCF) calculations [42], including two electrons in two molecular orbitals (MOs), have been performed using the MOLCAS 7.6 package [43] to generate a reference space (CAS[2,2]). The dynamical correlation effects were then incorporated by using the dedicated difference configuration interaction (DDCI) method [44] implemented in the CASDI code [45]. With this approach, one concentrates on the differential effects rather than on the evaluation of the absolute energies. DDCI1 (i.e. CAS+S) involves one hole and one particle (1h, 1p, 1h1p) single excitations on the full active space. DDCI2 also accounts for the two holes or two particles diexcitations (2h, 2p). Finally, the two holes/one particle (2h1p) and one hole/two particles (1h2p) excitations are taken into account in DDCI3 (i.e. CAS+DDCI). Since the DDCI philosophy relies on the simultaneous characterization of different spin states, which share similar spatial descriptions, one has to initially determine a set of common MOs to build up the CI space. Computed J values strongly depend on the quality of the magnetic orbitals [46]. In the present work, one uses orbitals obtained for the lowest triplet state. Calculations have been done with C_i symmetry on model dimers ensuing from the crystallographic data without any geometry optimization. All methyl groups have been replaced by hydrogen atoms. All atoms have been depicted with ANO-RCC type basis sets. The Cu, N, C and H atoms have been described with (21s15p10d6f4g2h)/[5s4p2d1f], (14s9p4d3f2g)/[3s2p1d], (14s9p4d3f2g)/[3s2p] and (8s4p3d1f)/[2s] contractions, respectively [47]. A (14s9p4d3f2g)/[3s2p1d] contraction has been used for the O atoms directly coordinated to copper whereas other O atoms have been described with a (14s9p4d3f2g)/[3s2p] contraction.

3. Results and Discussion

3.1. Synthesis and characterization

The dideprotonated ONO tridentate Schiff base ligand, $[4\text{-MeO-C}_6\text{H}_4\text{-C(O)CH=C(CH}_3\text{)N-}o\text{-C}_6\text{H}_4\text{-O}]^{2-}$ (L^{2-}), employed in this work was generated in a two-step one-pot reaction by facile condensation of 1-anisyl-1,3-butanedione with 2-aminophenol, in 1:1 molar ratio, in refluxing toluene for 2 hours, followed by the double deprotonation of the yellow diprotic precipitate formed on cooling, with 2.5 equiv of $t\text{BuO}^-\text{K}^+$ in THF. Pyridine based nitrogenous base (4- $\text{NC}_5\text{H}_4\text{R}$, $\text{R} = \text{H, CH}_3, \text{C(CH}_3\text{)}_3$) was then added to the dark red reaction mixture before dropwise addition of a THF solution of the copper(II) nitrate salt. The solutions were stirred overnight to separate microcrystalline compounds. Subsequent recrystallization of the product from dichloromethane solution gave dark green crystals of the binuclear ONO Cu(II) Schiff base complexes **1** and **2**, respectively (Scheme 1).



Scheme 1 Preparation of complexes **1** and **2**

Whatever the amount of pyridine used (1 to 4 equivalents), the doubly phenoxo bridged binuclear complex **1** was always obtained and isolated in excellent yields of 80-85%. The same binuclear derivative **1** was also isolated, albeit in a lower yield of 75%, when 4-picoline (4-C₅H₄CH₃), a more donating ligand, was used under the same conditions in place of pyridine. It is worth noting that among the nitrogenous coligands used, only the sterically more demanding 4-*tert*-butylpyridine does coordinate to the metal center to form the binuclear Cu(II) Schiff base complex **2** that is isolated in 75% yield (Scheme 1). Addition of an excess of 4-*tert*-butylpyridine to a THF solution of the binuclear compound **1** did not afford its mononuclear counterpart **2**. The extra stability of compound **1** presumably arises from its symmetrical nature which in turn might inhibit its dissociation to generate the monomeric species [Cu(II)L].

Both compounds **1** and **2** are thermally stable, air and moisture insensitive on storage under ordinary conditions, exhibiting good solubility in common polar organic solvents but are not soluble in ethanol, diethyl ether and hydrocarbon solvents. The molecular identity and geometry of both complexes were elucidated by X-ray crystal structure determination (see below). Analysis by ESI-MS did not give the parent ions or indeed any useful information, except a prominent fragment ion observed at $M/z = 345.0429$ in the mass spectrum of **1** that is assigned to the cation $[L^{63}\text{Cu-H}]^+$ ($M/z = 345.04262$). However, satisfactory analytical data on crystalline material demonstrates purity of both compounds (see Section 2.3).

3.2. Infrared spectral study

The solid-state FT-IR spectra of **1** and **2** were assigned on the basis of frequency calculations on DFT optimized geometries of **1** and **2** [37-41]. Table 2 gathers both computed and experimental major frequencies and their respective attributions, showing a good agreement between the calculated and observed spectral data. The most salient feature found in the spectra of **1** and **2** is the characteristic strong intensity bands in the ranges 1587-1466 and 1604-1584 cm⁻¹, respectively, assigned to the $\nu(\text{C}^{\equiv}\text{C})$, $\nu(\text{C}^{\equiv}\text{N})$, $\nu(\text{C}^{\equiv}\text{O})$ stretching vibrations of the Schiff base skeleton, suggesting that the imine nitrogen and carbonyl oxygen atoms coordinate to the Cu(II) metal ion. The weak bands observed at 3049 and 3040 cm⁻¹ for **1** and **2**, respectively, are attributed to the aromatic $\nu(\text{C-H})$ vibrations, while the bands found at 2932 and 2830 cm⁻¹ in the spectrum of **1** and at 2926 and 2864 cm⁻¹ in that of **2**, are characteristic of aliphatic $\nu(\text{C-H})$ vibrations. The observed medium intensity band at 1244 and 1242 cm⁻¹ for **1** and **2**, respectively, are due to the ν_{asym} vibration mode of the CH₃-O-aryl

group. The deformation modes of the C-H bonds in **1** and **2** show up respectively at 772 and 734 cm⁻¹. Thus, the infrared spectral data are consistent with the structural features of both compounds (see below Section 3.3).

Table 2 Computed and experimental infrared frequencies^a for compounds **1** and **2**

	Computed	1 ^b	2 ^b
$\nu(\text{C-H aryl})$	3089(w)	3049(w)	3040(w)
$\nu(\text{C-H, -OCH}_3)$	2992(w)	2932(w)	2926(w)
$\nu_{\text{sym}}(\text{CH}_3)$	2949(w)	2830(w)	2864(w)
$\nu(\text{C}\equiv\text{O}), \nu(\text{C}\equiv\text{N}), \nu(\text{C}\equiv\text{C})$	1587(s)-1466(s)	1581(s)-1485(s)	1604(s)-1584(s)
$\nu_{\text{asym}}(\text{C-H, C}_6\text{H}_4\text{-O-CH}_3)$	1240(m)	1244(m)	1242(m)
$\delta(\text{C-H})$	719(m)	772(m)	734(m)

^a in cm⁻¹. ^b recorded as KBr disk.

3.3. Description of the molecular and crystal structures

Perspective views of binuclear compounds **1** and **2**, including the atom labeling scheme, are shown in Figs. 1 and 2, while selected bond lengths and angles are listed in Table 3. Compound **1** crystallizes in the monoclinic system with space group P2₁/c, while complex **2** crystallizes in the triclinic crystal system P $\bar{1}$ space group. In both cases, the asymmetric unit consists of a neutral centrosymmetric bis(μ -phenoxo)di-copper(II) complex with the two halves of the dimeric unit related by a crystallographic inversion center in the middle of the four membered Cu₂O₂ core.

In each doubly phenoxo bridged dimer, each μ -phenolic oxygen atom bridges in an antisymmetric fashion two Cu(II) metal ions leading to a Cu₂O₂ rhombic core with two distinct bridging Cu(1)-O(1) and Cu(1)-O(1') bond lengths of 1.9357(14) and 1.9579(13) Å in **1**, and 1.9247(11) and 2.445(3) Å in **2**, respectively, while Cu-O-Cu bridge angles are of 102.95(6)° in **1** and of 95.506(3)° in **2**. In both compounds, the Cu(1)-O(1)-Cu(1')-O(1') torsion angle is 0.0°, indicating that the Cu₂O₂ core is strictly planar. Furthermore, the intramolecular non-bonding Cu(1)...Cu(1') distance is 3.0462(4) in **1** and 3.253(3) Å in **2**. All these bond distances and angles are in good agreement with that recently reported for analogous doubly phenoxo bridged square planar and square pyramidal Cu(II) species [19,20,24,48,49].

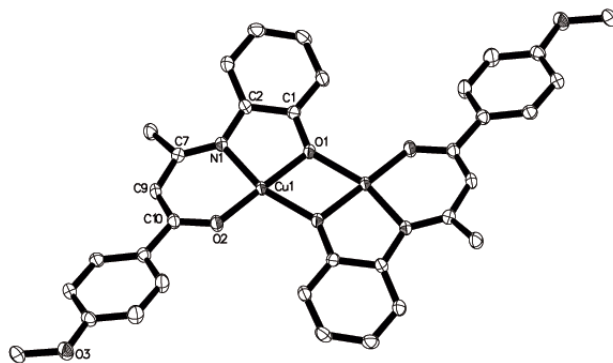


Fig. 1 Molecular structure of complex **1** showing partial atom numbering scheme. Hydrogen atoms are omitted for clarity. Thermal ellipsoids are drawn at 60% probability.

In compound **1**, the dinuclear unit presents two tetracoordinated cupric ions in a slightly distorted square-planar environment (Fig. 1), with the coordination sphere formed by the deprotonated amide nitrogen atom and the carbonyl and phenoxo oxygen atoms of the acyclic Schiff base ligand, and the bridging oxygen atom of the second half-unit. The four bond lengths span the range 1.8749(14)-1.9579(13) Å (Table 3). The sum of the angles around the Cu(II) center is 359.97°, indicating a slightly distorted square-planar geometry around the metal atom. The diagonal angles are, indeed, found to be 176.42(6)° [O(1)-Cu(1)-O(2)] and 161.35(6)° [N(1)-Cu(1)-O(1')]. Those values deviate somewhat from idealized transoid angles of 180° expected for a perfectly square-planar compound. Deviations of the coordinating O(1), O(1'), O(2) and N(1) atoms around Cu(II) center from the least-square mean planes through them are 0.0224(11), -0.0198(9), 0.0166(8) and -0.0192(9) Å, respectively, and that of the Copper atom from the same plane is 0.0290(6) Å.

Table 3 Selected bond distances (Å) and angles (°) for compounds **1** and **2**

	1	2
Bond distances		
Cu(1)-O(1)	1.9579(13)	1.9247(11)
Cu(1)-N(1)	1.9183(16)	1.9729(13)
Cu(1)-O(2)	1.8749(14)	1.9155(11)
Cu(1)-X ^a	1.9579(13)	2.0467(13)
Angles		
O(1)-Cu(1)-O(2)	176.42(6)	174.93(5)
O(1)-Cu(1)-N(1)	84.30(6)	84.60(5)
O(2)-Cu(1)-N(1)	97.76(6)	95.20(5)
N(1)-Cu(1)-X ^a	161.35(6)	162.66(5)
O(1)-Cu(1)-X ^a	77.05(6)	90.86(5)
O(2)-Cu(1)-X ^a	100.86(6)	90.78(5)

^aX = O(1)^{#1} for **1**, X = n(2) for **2**; #1 Symmetry transformations used to generate equivalent atoms: -x+1,-y,-z+1.

By contrast, the binuclear unit of **2** exhibits two pentacoordinated Cu(II) centers in a square pyramidal (4+1) coordination geometry (Fig. 2). The basal plane consists of the same tridentate ONO- donor set of atoms as in **1** and the nitrogen atom of the monodentate 4-*tert*-butylpyridine, while the apical site is occupied by the phenoxido oxygen atom of another ONO-tridentate dianion that is basal to the second copper(II) metal ion of the dimer. The copper atom is deviated by 0.1022 Å from the basal plane due to the formation of the axial bond. The bond distances of the donor atoms of the tridentate Schiff base ligand to the central metal atom are in the range 1.9155(11) to 1.9729(13) Å, and are typical for Cu(II) complexes [19,20,23-27,50,51]. The Cu(II)-N_{py} bond distance of 2.0467(13) Å is slightly longer than those observed in related ternary square-planar Cu(II) complexes (1.90-1.96 Å) [23-27]. Such a lengthening may explain an easy dissociation of less donating and less sterically demanding monodentate N-donor ligand leading to the formation of square planar dimeric complex **1**, as observed here with pyridine and 4-picoline. Obviously, the Cu(1)-O(1') axial bond length (2.445(3) Å) is much longer than the equatorial distances which may be ascribed to Jahn-Teller distortion. The Addison parameter (τ) [52] which is an index of distortion from the square-pyramidal to the trigonal-bipyramidal geometry is calculated to be 0.072, indicating that the pentacoordinated geometry is very little distorted from a perfectly square pyramid ($\tau = 0$). The dihedral angle between the bridging Cu₂O₂ plane and the mean basal plane of the copper square pyramid is 80.713(4)°.

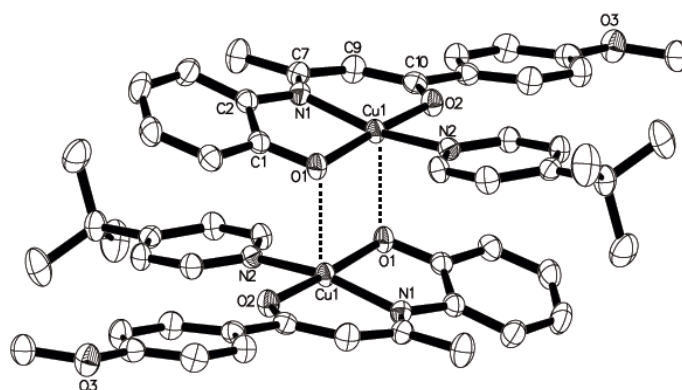


Fig. 2 Molecular structure of complex **2** showing partial atom numbering scheme and apical short contact Cu-O interactions in dashed-line. Hydrogen atoms are omitted for clarity. Thermal ellipsoids are drawn at 60% probability.

In both complexes, the fused five- and six-membered heterometallacycles formed upon the Schiff base condensation of anisoylacetone with aminophenol and subsequent chelation of the Cu(II) ion, are essentially co-planar. Moreover, the six-membered metallacycle is also co-planar with the anisyl substituent with dihedral angles of 3.202(3) and 0.868(2)° in **1** and **2**, respectively, and of 25.60(3)° with the 4-*tert*-butylpyridine ligand in **2**. In addition, interatomic distances and angles (Tables 3 and 4) are indicative of substantial π delocalization of the electron density through the chelate rings.

3.4. EPR spectroscopy

The X-band EPR spectra of compounds **1** and **2** have been recorded in fluid solutions and as powdered samples at room temperature (298 K). The spectra of compound **1**, recorded on a polycrystalline powder, do not show any EPR signal. In contrast, EPR spectrum of **2**, obtained in the polycrystalline state at room temperature (Fig. 3), exhibits unresolved broad-band resonances, due to exchange coupling between the copper(II) ions, at $g = 2.071$ with a ΔH peak-to-peak values of 71.5 G. Moreover, the spectrum showed a weak $\Delta M_S = \pm 2$ half field transition (Fig. 3, insert) at $g = 4.181$, thus giving unequivocal proof of the existence of the triplet state ($S = 1$) in the solid.

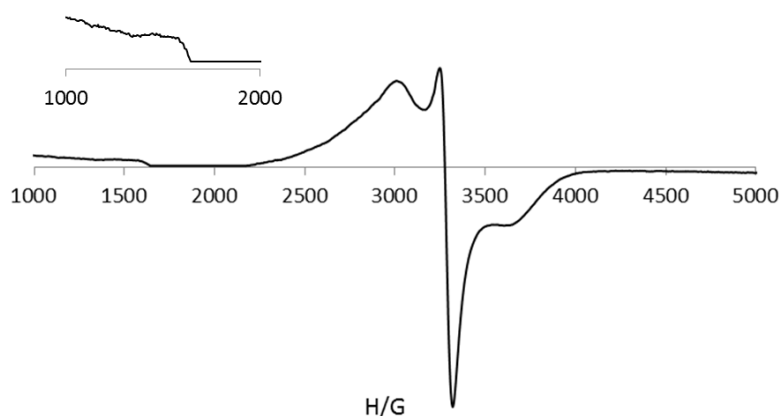


Fig. 3 Experimental EPR spectrum of complex **2** recorded in the solid state at 298 K with an expansion of the half-field region.

The fluid spectrum of compound **1** dissolved in pure pyridine, is typical of mononuclear Cu(II) centers and shows a well-resolved hyperfine structure with isotropic g

value of 2.115 with hyperfine coupling constants A_{Cu} of 76 G. This observation suggests that, after dissolution, the dimeric structure of **1** is destroyed, as noticed for other polynuclear Cu(II) compounds [5b,54].

On the other hand, at 298 K in CH_2Cl_2 :1,2- $\text{C}_2\text{H}_4\text{Cl}_2$ (1:1 v:v) solution (Fig. 4), the spectrum of compound **2** exhibits partially resolved superhyperfine structure (5 lines, $A_{\text{N}} = 11.5$ G) superimposed on the quartet hyperfine structure centered at $g_{\text{iso}} = 2.101$ with hyperfine coupling A_{Cu} value of 90 G. Those data are consistent with a mononuclear copper(II) species ($S = 1/2$), with the Cu(II) ion in a slightly distorted square planar or square pyramidal coordination environment [26,50,53,55].

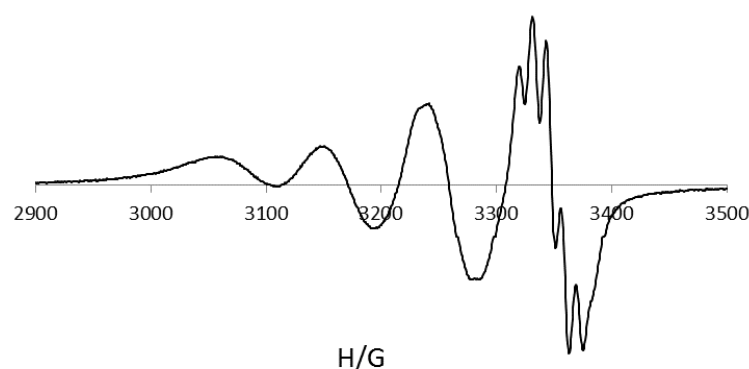


Fig. 4 X-band EPR spectrum of **2** recorded in CH_2Cl_2 : 1,2- $\text{C}_2\text{H}_4\text{Cl}_2$ (1:1 v:v) mixture at 298 K.

3.5. Magnetic properties

The magnetic properties of compounds **1** and **2** have been measured from room temperature to 2 K using a constant applied field of 2 kOe. On cooling a powdered sample of **1** $\chi_{\text{M}}T$ (χ_{M} the magnetic molar susceptibility and T the temperature in Kelvin) decreases continuously from $0.375 \text{ cm}^3 \text{ K mol}^{-1}$ down to almost zero ($0.02 \text{ cm}^3 \text{ K mol}^{-1}$). This is a clear indication that very strong antiferromagnetic interaction operates between the two spins. The experimental temperature dependence of $\chi_{\text{M}}T$ of two spins interacting through the Heisenberg-Dirac-van Vleck Hamiltonian ($H = -J \mathbf{S}_A \cdot \mathbf{S}_B$) is reproduced by a slightly modified Bleaney-Bowers equation [11,56]:

$$\chi_M T = (1-x) \frac{2Ng^2\beta^2}{k} \frac{1}{3 + \exp\left(-\frac{J}{kT}\right)} + x \frac{2Ng^2\beta^2}{3k} S(S+1)$$

N , β , g , and k represent the Avogadro number, the Bohr magneton, the Boltzmann constant and the Zeeman factor respectively, while x represents the fraction of paramagnetic impurities in the sample. The best agreement between experiment and theory (Fig. 5) is obtained with $g = 2.15$ and $J = -397 \text{ cm}^{-1}$. Such a strong antiferromagnetic spin-coupling involving two dx^2-y^2 orbitals of the metal centers could be expected owing to magnetic interactions mediated by equatorial-equatorial bridging ligands with a Cu-O-Cu angle of $102.95(6)^\circ$ [8,13-15], agreeing well with those reported for other analogous compounds [19,20].

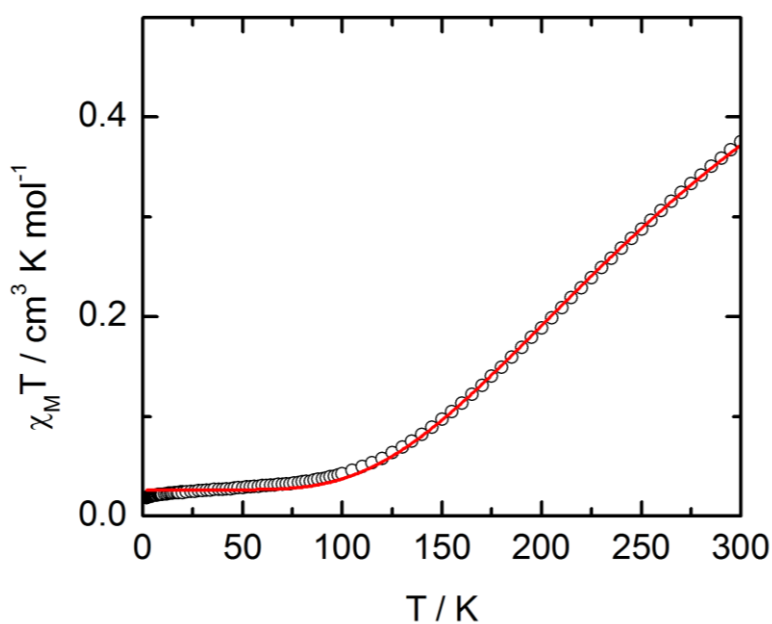


Fig. 5 Temperature dependence of $\chi_M T$ for compound **1** (empty circles) with the best-fitted curve (full red line).

Compound **2** behaves differently from **1**. Indeed, $\chi_M T$ (χ_M the magnetic molar susceptibility and T the temperature in Kelvin) for compound **2** is constant and equal to $0.87 \text{ cm}^3 \text{ K mol}^{-1}$ (Fig. 6) in the whole temperature range which witnesses that no interaction operates between the two spins localized on Cu(II) metal ions. With this Curie constant we can determine the average Zeeman value $g = 2.152$ per Cu(II) center. The simulated Brillouin curve for two uncoupled Cu(II) perfectly reproduces the field variation of the magnetization at 2 K (inset of

Fig. 5). Compound **2** shows a slightly distorted square-pyramidal geometry, thus the magnetic interactions that are mediated by apical-equatorial bridges are expected to be negligible as the dx^2-y^2 and dz^2 orbitals of the Cu(II) centers are orthogonal to each other (see below Section 3.6), and the axial Cu-O bond is significantly longer (0.5203 Å) than the equatorial ones.

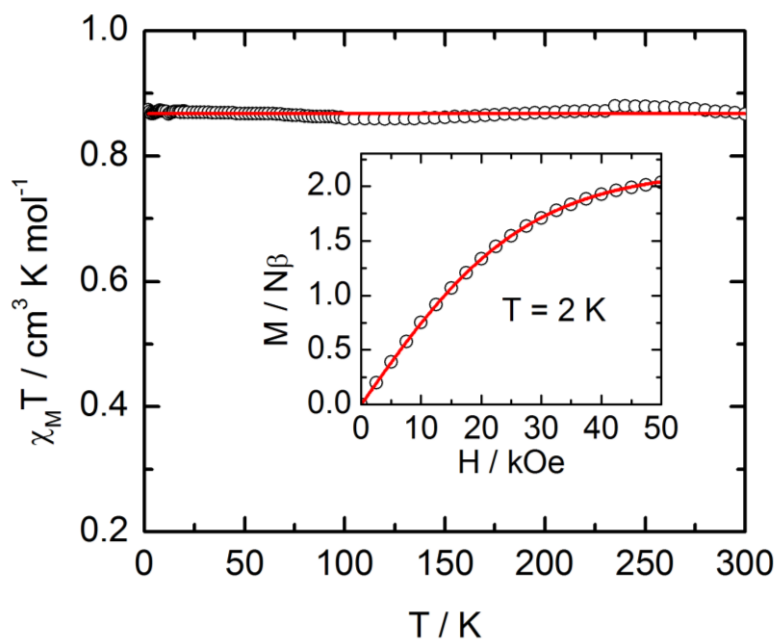


Fig. 6 Temperature dependence of $\chi_M T$ for compound **2** with the best fitted curve with a Curie law (full line). Inset: field variation of the magnetization for compound **2** at 2 K with the best fitted curve with a Brillouin law (full line).

3.6. Theoretical investigations

To complete magnetic measurements, *ab initio* calculations of the magnetic interactions in **1** and **2** have been performed at the CAS[2,2]SCF/DDCI level (see computational details). Such a computational approach usually gives excellent agreement with respect to experimental findings [57], even for through-space interactions such as in **2** [58]. Of course, DFT (within the broken symmetry (BS) approximation) could have been used instead of CASSCF/DDCI approach [59]. However, major drawbacks, among them the spin contamination of the BS wavefunction and the choice of the exchange-correlation functional, explain the present choice of wavefunction-based multireference approaches. Results are gathered in Table 4 and the magnetic orbitals of **1** are given in Fig. 7. At the best level of calculation, i.e. CAS+DDCI, the strong antiferromagnetic behaviour of **1** is recovered whereas no interaction is calculated in the case of **2** due to a poor through-space overlap

between the magnetic orbitals. Both results confirm the experimental data although in the case of **1**, slight discrepancy remains between the measured and calculated exchange couplings.

Table 4 Calculated exchange coupling constant J (cm^{-1}) in model complexes of **1** and **2**

	1	2
CAS[2,2]SCF	-40.0	0.5
CAS	-27.5	0.5
CAS+S	-115.3	-0.6
CAS+DDCI	-256.0	3.1

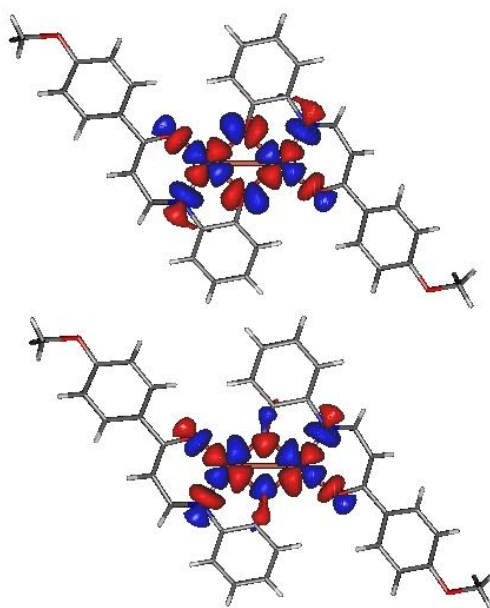


Fig. 7 Magnetic MOs extracted from a CAS[2,2]SCF calculation over the triplet state.

4. Conclusion

In this report, we have shown that treatment of the dianionic ONO-tridentate ligand derived from the Schiff base condensation of 1-anisyl-1,3-butanedione and 2-aminophenol, with Cu(II) ions leads to the dinuclear complex **1** when the reaction is carried out in the presence of pyridine or 4-picoline, while the same reaction carried out in the presence of 4-*tert*-butylpyridine results in the formation of the binuclear species **2**. Both compounds have been authenticated by single crystal X-ray diffraction analysis that showed that dimeric complex **1** presents two tetracoordinated Cu(II) centers in a square planar environment and that dimer **2** exhibits two pentacoordinated Cu(II) centers with a square pyramidal coordination sphere. The two halves of the dimeric unit are related through a planar Cu_2O_2

rhombic core in both cases. EPR study in fluid solutions indicates that the dimeric structure of **1** and **2** is destroyed upon dissolution. In the solid-state, **1** is EPR silent whereas both $\Delta MS = \pm 1$ and $\Delta MS = \pm 2$ transitions are observed for **2**. The variable temperature magnetic measurements reveal that dimer **1** exhibits a strong antiferromagnetic interaction due to the coplanar disposition of the Cu(II) metal ions and acute Cu-O-Cu angle (102°), whereas dimer **2** shows negligible antiferromagnetic coupling caused by the basal-apical phenoxido oxygen bridging mode. The magnetic findings have been corroborated by theoretical Ab initio calculations.

Appendix A. Supplementary data

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References

- [1] D. Venegas-Yazigi, D. Aravena, E. Spodine, E. Ruiz, S. Alvarez, *Coord. Chem. Rev.* 254 (2010) 2086.
- [2] E. Ruiz, S. Alvarez, A. Rodríguez-Fortea, P. Alemany, Y. Pouillon, C. Massobrio, in: J.S. Miller, M. Drillon (Eds.), *Electronic Structure and Magnetic Behavior in Polynuclear Transition-Metal Compounds*, vol. 2, Wiley-VCH, Weinheim, 2001, p. 227.
- [3] S. Anbu, M. Kandaswamy, *Polyhedron* 30 (2011) 123.

- [4] R. Vafazadeh, B. Khaledi, A.C. Willis, M. Namazian, *Polyhedron* 30 (2011) 1815.
- [5] (a) A. Jana, S. Mohanta, *Inorg. Chim. Acta*, 405, (2013) 265;
(b) S. Saha, A. Sasmal, C. R. Choudhury, C. J. Gómez-García, E. Garribba, S. Mitra, *Polyhedron* 69 (2014) 262.
- [6] D. Gatteschi, O. Kahn, R.D. Willet, *Magnetostructural Correlations in Exchange Coupled Systems*, Reidel, Dordrecht, 1984.
- [7] V.H. Crawford, H.W. Richardson, J.R. Wasson, D.J. Hodgson, W.E. Hatfield, *Inorg. Chem.* 15 (1976) 2107.
- [8] L. Merz, W. Haase, *J. Chem. Soc., Dalton Trans.* (1980) 1594.
- [9] T.N. Doman, D.E. Williams, J.F. Banks, R.M. Buchanan, H.-R. Chang, R.J. Webb, D.N. Hendrickson, *Inorg. Chem.* 29 (1990) 1058.
- [10] G. Dutta, R. K. Debnath, A. Kalita, P. Kumar, M. Sarma, R. B. Shankar, B. Mondal, *Polyhedron* 30 (2011) 293.
- [11] O. Kahn, *Molecular Magnetism*, VCH Publishers, New York, 1993.
- [12] J.-P. Launay, M. Verdaguer, *Electrons in Molecules: From Basic Principles to Molecular Electronics*, 1st Edn, OXFORD University Press, Oxford, UK, 2014, Chapter 2, p. 145.
- [13] L.K. Thompson, S.K. Mandal, S.S. Tandon, J.N. Bridson, M.K. Park, *Inorg. Chem.* 35 (1996) 3117.
- [14] E. Ruiz, P. Alemany, S. Alvarez, J. Cano, *Inorg. Chem.* 36 (1997) 3683.
- [15] E. Safaei, A. Wojtczak, E. Bill, H. Hamidi, *Polyhedron* 29 (2010) 2769.
- [16] S. Gupta, A. Mukherjee, M. Nethaji, A. R. Chakravarty, *Polyhedron* 24 (2005) 1922.

- [17] A. Mukherjee, R. Raghunathan, M.K. Saha, M. Nethaji, S. Ramasesha, A.R. Chakravarty, *Chem.-Eur. J.* 11 (2005) 3087.
- [18] S. G. Naik, A. Mukherjee, R. Raghunathan, M. Nethaji, S. Ramasesha, A. R. Chakravarty, *Polyhedron* 25 (2006) 2135.
- [19] P. Bhowmik, N. Aliaga-Alcalde, V. Gómez, M. Corbella, S. Chattopadhyay, *Polyhedron* 49 (2013) 269.
- [20] N.E. Borisova, A. Kostin, T.V. Magdesieva, M.D. Reshetova, O. Nikitin, V. Paredes-García, M.T. Garland, P. Hermosilla-Ibáñez, W. Cañon-Mancisidor, A. Rodionov, D. Venegas-Yazigi, E. Spodine, *New J. Chem.* 38 (2014) 709.
- [21] S. Khanra, T. Weyhermüller, P. Chaudhuri, *Dalton Trans.* (2009) 3847.
- [22] C.R. Bhattacharjee, P. Goswami, H.a.R. Pramanik, P.C. Paul, P. Mondal, *Spectrochim. Acta A* 78 (2011) 1408.
- [23] R.N. Patel, V.L.N. Gundla, D.K. Patel, *Polyhedron* 27 (2008) 1054.
- [24] R. Vafazadeh, R. Esteghamat-Panah, A. C. Willis, A. F. Hill, *Polyhedron* 48 (2012) 51.
- [25] P.A.N. Reddy, M. Nethaji, A.R. Chakravarty, *Inorg. Chim. Acta* 337 (2002) 450.
- [26] A. Garcí'a-Raso, J. J. Fiol, B. Adrover, P. Tauler, A. Pons, I. Mata, E. Espinosa, E. Molins, *Polyhedron* 22 (2003) 3255.
- [27] R.N. Patel, N. Singh, V.L.N. Gundla, *Polyhedron* 25 (2006) 3312.
- [28] C. Biswas, S. Chattopadhyay, M.G.B. Drew, A. Ghosh, *Polyhedron* 26 (2007) 4411.
- [29] V. Tudor, V. Kravtsov, M. Julve, F. Lloret, Y.A. Simonov, J. Lipkowski, V. Buculei, M. Andruh, *Polyhedron* 20 (2001) 3033.

- [30] W.L.F. Armarego, C.L.L. Chai, Purification of Laboratory Chemicals, Fifth ed., Butterworth-Heinemann, 2003.
- [31] V.V. Popic, S.M. Korneev, V.A. Nikolaev, I.K. Korobitsyna, Synthesis (1991) 195.
- [32] P. Pascal, Ann. Chim. Phys. 19 (1910) 5.
- [33] A. Altomare, M.C. Burla, M. Camalli, G. Cascarano, C. Giacovazzo, A. Guagliardi, A.G.G. Moliterni, G. Polidori, R. Spagna, J. Appl. Crystallogr. 32 (1999) 115.
- [34] G.M. Sheldrick, Acta Crystallogr. A64 (2008) 112.
- [35] L.J. Farrugia, J. Appl. Crystallogr. 45 (2012) 839.
- [36] O.V. Dolomanov, L.J. Bourhis, R.J. Gildea, J.A.K. Howard, H. Puschmann, J. Appl. Crystallogr. 42 (2009) 339.
- [37] E.J. Baerends, D.E. Ellis, P. Ros, Chem. Phys. 2 (1973) 41.
- [38] (a) G.T. Velde, F.M. Bickelhaupt, E.J. Baerends, C.F. Guerra, S.J.A. van Gisbergen, J.G. Snijders, T. Ziegler, J. Comput. Chem. 22 (2001) 931; (b) C.F.; Guerra, J.G. Snijders, E.J. Baerends, Theor. Chem. Acc. 99 (1998) 391; (c) ADF2012.01, SCM, Theoretical Chemistry, Vrije Universiteit, Amsterdam, The Netherlands, <http://www.scm.com>.
- [39] A. Becke, Phys. Rev. A 38 (1998) 3098.
- [40] J.P. Perdew, Phys. Rev. B 33 (1986) 8822.
- [41] E. Van Lenthe, E.J. Baerends, J. Comput. Chem. 24 (2003) 1142.
- [42] B.O. Roos, Adv. Chem. Phys. 69 (1987) 399.

- [43] F. Aquilante, L. DeVico, N. Ferre, G. Ghigo, P.A. Malmqvist, P. Neogrady, T. Bondo Pedersen, M. Pitonak, M. Reiher, B.O. Roos, L. Serrano-Andres, M. Urban, V. Veryazov, R. Lindh, *J. Comput. Chem.* 31 (2010) 224.
- [44] (a) J. Miralles, J.P. Daudey, R. Caballol, *Chem. Phys. Lett.* 198 (1992) 555; (b) J. Miralles, O. Castell, R. Caballol, J.P. Malrieu, *Chem. Phys.* 172 (1993) 33.
- [45] N. Ben Amor, D. Maynaud, *Chem. Phys. Lett.* 286 (1998) 211.
- [46] (a) C. Angeli, C.J. Calzado, *J. Chem. Phys.* 137 (2012) 034104; (b) N. Suaud, R. Ruamps, N. Guihery, J.P. Malrieu, *J. Chem. Theory Comput.* 8 (2012) 4127.
- [47] (a) B.O. Roos, R. Lindh, P.A. Malmqvist, V. Veryazov, P.O. Widmark, *J. Phys. Chem. A* 108 (2004) 2851; (b) B.O. Roos, R. Lindh, P.A. Malmqvist, V. Veryazov, P.O. Widmark, *J. Phys. Chem. A* 109 (2005) 6575.
- [48] A. Mukherjee, M.K. Saha, M. Nethaji, A.R. Chakravarty, *Polyhedron* 23 (2004) 2177.
- [49] A. Jakob, C.C. Joubert, T. Ruffer, J.C. Swarts, H. Lang, *Inorg. Chim. Acta* 411 (2014) 48.
- [50] R.N. Patel, S.P. Rawat, M. Choudhary, V.P. Sondhiya, D.K. Patel, K.K. Shukla, D.K. Patel, Y. Singh, R. Pandey, *Inorg. Chim. Acta* 392 (2012) 283.
- [51] Y. Yahsi, H. Kara, *Inorg. Chim. Acta* 397 (2013) 110.
- [52] A.W. Addison, T.N. Rao, J. Reedijk, J. van Rijn, G.C. Verschoor, *J. Chem. Soc., Dalton Trans.* (1984) 1349.
- [53] B. Sarkar, M.G.B. Drew, M. Estrader, C. Diaz, A. Ghosh, *Polyhedron* 27 (2008,) 2625.
- [54] (a) I.A. Koval, M. Sgobba, M. Huisman, M. Lüken, E. Saint-Aman, P. Gamez, B. Krebs, J. Reedijk, *Inorg. Chim. Acta* 359 (2006) 4071; (b) C.P. Pradeep, S.K. Das,

- Polyhedron 28 (2009) 630; (c) S. Thakurta, P. Roy, G. Rosair, C.J. Gómez-García, E. Garribba, S. Mitra, Polyhedron 28 (2009) 695.
- [55] A. Trujillo, F. Justaud, L. Toupet, O. Cadot, D. Carrillo, C. Manzur, J.-R. Hamon, New J. Chem. 35 (2011) 2027.
- [56] B. Bleaney, K. D. Bowers, Proc. Roy. Soc. London, A 214 (1952) 451.
- [57] J.-P. Malrieu, R. Caballol, C.J. Calzado, C. de Graaf, N. Guihéry, Chem. Rev. 114 (2014) 429.
- [58] (a) B. Le Guennic, N. Ben Amor, D. Maynau, V. Robert, J. Chem. Theory Comput. 5 (2009) 1506; (b) J. S. Costa, N.A.G. Bandeira, B. Le Guennic, V. Robert, P. Gamez, G. Chastanet, L. Ortiz-Frade, L. Gasque, Inorg. Chem. 50 (2011) 5696; (c) B. Kozlevcar, N. Kitanovski, Z. Jaglicic, N.A.G. Bandeira, V. Robert, B. Le Guennic, P. Gamez, Inorg. Chem. 51 (2012) 3094; (d) N.A.G. Bandeira, D. Maynau, V. Robert, B. Le Guennic, Inorg. Chem. 52 (2013) 7980.
- [59] (a) I. de P.R. Moreira, F. Illas, Phys. Chem. Chem. Phys. 8 (2006) 1645; (b) A. Bencini, Inorg. Chim. Acta 361 (2008) 3820; (c) A. Bencini, F. Totti, J. Chem. Theory Comput. 5 (2009) 144; (d) F. Neese, Coord. Chem. Rev. 253 (2009) 526; (e) C.J. Cramer, D.G. Truhlar, Phys. Chem. Chem. Phys. 11 (2009) 10757; (f) A.J. Cohen, P. Mori-Sánchez, W. Yang, Chem. Rev. 112 (2012) 289.