Two Types of Pyridine Ligands in Mononuclear and Dinuclear Copper(II) Carboxylates

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Copper(II) acetate [Cu(OOCCH₃)₂(L₁)₂] (L₁ = 2,6-diaminopyridine) (1), [Cu(OOCCH₃)₂(L₂)₂] (L₂ = 2-amino-6-methylpyridine) (2) and benzoate compounds [Cu₂(OOCC₆H₅)₄(L₁)₂]·2CH₃CN (3), [Cu₂(OOCC₆H₅)₄(L₁)₂] (4), [Cu₂(OOCC₆H₅)₄(L₂)₂] (5), [Cu(OOCC₆H₅)₂(L₂)₂] (6), were synthesized and characterized. X-ray structure analysis revealed monomeric structure in 1 and 6. In 1, cis arrangement of the ligands was found, and trans in 6, in an elongated octahedral CuO₄N₂ chromophore. The basal plane in 1 and 6 is formed by one co-ordinated oxygen atom from both carboxylates (Cu–O 1.9560(12)–2.007(3) Å) and pyridine nitrogen atom from two pyridine ligands (Cu–N 2.013(3)–2.282(14) Å), forming a CuO₂N₂ plane, while the second carboxylate oxygen atoms are more distant (Cu–O 2.488(3)–2.7648(16) Å). The dinuclear paddle-wheel central core was found in 5 (Cu–O 1.942(6)–1.992(6) Å), with pyridine nitrogen atoms in the axial positions (Cu–N 2.283(7), 2.284(7) Å). All compounds were characterized by magnetic measurements, electronic and vibrational spectroscopy and tested for fungal growth retardation activity.

Key words
copper
π-stacking
structure
spectroscopy
carboxylates
pyridine ligands

INTRODUCTION

Two main types of copper(II) carboxylates, mononuclear (distorted octahedral) and dinuclear (paddle-wheel), respectively, have been observed.¹² The diversity of these coordination compounds largely depends on the carboxylate radical tail and even more on the additional ligand, both directing the path of the synthesis via their size, shape, substituents, etc. The stabilization of these compounds often takes place via non-covalent intra or inter-molecular forces as hydrogen-bonding or π-stacking interactions.³ Very common additional ligands are based on the pyridine molecule, due to a strong nitrogen coordination site in the ring. The groups on positions 2 and 6 on pyridine ring are playing especially important role in this type of complexes. Alkyl groups may sterically hinder also pyridine nitrogen coordination site, while amino groups may stabilize the compound with hydrogen bonds in the same coordination sphere or inter-molecular. Additionally, alkyl and amino groups are electron donating and as such disfavor π-stacking interactions.³ The aim of this work is to correlate different carboxylates and pyridine-based ligands in copper(II) complexes with the selected properties and structural parameters of the investigated compounds.

In this paper, synthesis, characterization and fungicidal activity of copper(II) acetates and benzoates with 2,6-diaminopyridine and 2-amino-6-methylpyridine are presented.

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EXPERIMENTAL

Materials

Grey-black 2,6-diaminopyridine L1 (2.00 g) was recrystallized from 50.0 mL of chloroform. The mixture was heated up to the boiling temperature, filtered, and left at 253 K for three hours. Precipitated colourless crystals were filtered off and dried in a desiccator over KOH for a day (yield 40%).

[Cu₂(OOCC₆H₅)₄(CH₃COOH)₂] was prepared from hot aqueous solution of copper(II) sulphate, benzoic acid and sodium benzoate by precipitation method. The other starting substances and solvents were purchased from commercial sources and used without previous purification.

Synthesis

[Cu(OOCC₆H₅)₂(L1)₂] (L1 = 2,6-diaminopyridine) (1)

Grey-black 2,6-diaminopyridine L1 (2.00 g) was recrystallized from 50.0 mL of chloroform. The mixture was heated up to the boiling temperature, filtered, and left at 253 K for three hours. Precipitated colourless crystals were filtered off and dried in a desiccator over KOH for a day (yield 40%). 

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The procedure was similar as already described.4 [Cu₂(OOCC₆H₅)₄(H₂O)₂] was dissolved in 15.0 mL of acetonitrile, while heated and than added to the L1 solution. The obtained blue-green solution was left at 279 K for 24 hours. Dark green crystals were filtered off and then dried on glass fryth and for 24 hours in a desiccator (yield 44 %). λ_max / nm 340, 580; μ eff / BM 1.94.

Anal. Calcd for C₁₄H₂₀CuN₆O₄ (Mᵣ = 399.90): C 42.1, H 5.04, N 10.7, Cu 15.4 %; found: C 42.2, H 4.92, N 20.8, Cu 15.9 %.

[Cu₂(OOCC₆H₅)₄(L1)₂] (L1 = 2,6-diaminopyridine) (2)

The procedure was similar as already described.4 [Cu₂(OOCC₆H₅)₄(L1)₂] (0.40 g) was dissolved in 22.0 mL of acetonitrile. The solution was added to the acetonitrile (4.0 mL) solution of L1 (0.65 g). After 24 hours at 279 K, violet crystals were filtered off and dried for 24 hours in a desiccator (yield 91 %). λ_max / nm 315, 400sh (sh = shoulder), 545, 690; μ eff / BM 1.94.


[Cu₂(OOCC₆H₅)₄(L1)₂]·2CH₃CN (3)

The procedure was similar as already described.5 0.43 g of [Cu₂(OOCC₆H₅)₄(CH₃COOH)₂] was dissolved in acetonitrile (10.0 mL) and added to acetonitrile solution (10.0 mL) of 2,6-diaminopyridine L1 (0.22 g). After 24 hours at 279 K, green needle-like crystals were filtered off and left for a day in a desiccator (yield 85 %). λ_max / nm 335, 400sh, 735; μ eff / BM 1.43.


[Cu₂(OOCC₆H₅)₄(L1)₂] (4)

0.22 g of 2,6-diaminopyridine L1 was dissolved in 2.0 mL of acetone. In the other flask, 0.43 g of [Cu₂(OOCC₆H₅)₂]·(C₆H₅COOH)₂ was dissolved in 9.0 mL of acetone while heated, and than added to the L1 solution. Green solution was left at 279 K for 2 hours. The green needle-like crystals were filtered off (they dim on air – unstable) and left for a day in a desiccator (yield 55 %). λ_max / nm 330, 400sh, 730; μ eff / BM 1.45.

Anal. Calcd for C₃₀H₃₂CuN₆O₈ (Mᵣ = 829.81): C 55.0, H 4.13, N 10.1, Cu 15.3 %; found: C 54.8, H 4.08, N 9.94, Cu 15.3 %.

[Cu₂(OOCC₆H₅)₄(L2)₂] (5)

In 7.5 mL of acetonitrile, 0.34 g of [Cu₂(OOCC₆H₅)₄(CH₃COOH)₂] was dissolved, and added to the solution of 0.11 g of 2-amino-6-methylpyridine L2 in 0.5 mL of acetonitrile. The solution was left at 279 K for 24 hours. Dark green crystals were filtered off and than dried on fryth and for 24 hours in a desiccator (yield 44 %). λ_max / nm 315, 390sh, 735; μ eff / BM 1.39.

Anal. Calcd for C₁₄H₁₈CuN₄O₇ (Mᵣ = 587.84): C 58.0, H 4.38, N 6.77, Cu 15.4 %; found: C 57.7, H 4.18, N 7.11, Cu 15.3 %.

[Cu(OOCC₆H₅)₂(L2)₂] (6)

0.34 g of [Cu₂(OOCC₆H₅)₄(CH₃COOH)₂] was dissolved in 15 mL of acetonitrile, and added to the solution of 0.34 g of 2-amino-6-methylpyridine L2, dissolved in 3 mL of acetonitrile. Obtained blue-green solution was left at 279 K for 24 hours. In the first step, monomeric 6 and dimeric 5 benzoate compounds precipitated, however during next 24 hours, green crystals of 5 were dissolved and only dark violet crystals of 6 were filtered off. They were dried on glass fryth and 24 hours in a desiccator (yield 79 %). λ_max / nm 320, 405, 550, 705; μ eff / BM 1.95.

Anal. Calcd for C₁₈H₂₀CuN₄O₈ (Mᵣ = 522.06): C 59.8, H 5.02, N 10.7, Cu 12.2 %; found: C 59.7, H 4.86, N 10.9, Cu 12.5 %.

d-spacings (Å) and their relative intensities, from X-ray powder diffraction data for 1, 2, 4, 5 and 6, are in agreement with the calculated values8,9 from crystal structure analysis. X-ray powder diffraction pattern for 4 is completely reproducible for different samples (structure not solved).

X-ray Diffraction Work

Single crystal diffraction measurements were carried out on an Enraf-Nonius CAD4 diffractometer with graphite-monochromated Mo-Kα radiation. Unit cell dimensions were obtained from 25 reflections in the θ range 8–16°. Common data corrections for variations in reference reflections and Lorentz polarization effects were applied.8 Absorption correction was performed by Gaussian integration method for 5 and 6. No absorption correction was performed for compound 1. Structures were solved by direct methods implemented in SHELXS-97 and refined by full matrix least squares on F² by SHELXL-97.9 Non-hydrogen atoms were refined anisotropically, hydrogen atoms were generated geometrically, assigned appropriate isotropic thermal displacement parameters and allowed to ride on their parent atoms.
Physical Measurements

Metal analysis was carried out electrogravimetrically with Pt electrodes. C,H,N analysis was performed with a Perkin Elmer, Elemental Analyzer 2400 CHN. Interplanar spacings were obtained by the Guinier camera (Huber and Enraf-Nonius) with Cu-Kα radiation. The magnetic susceptibility of the substances was determined at room temperature by powdered samples with a Sherwood Scientific MSB-1 balance, using Hg[Co(NCS)₄] as a calibrant. Diamagnetic corrections were estimated from Pascal’s constants. Infrared spectra were measured on mineral mulls between CsI plates, using a Perkin-Elmer FT-IR 1720X spectrometer. Electronic spectra were recorded as nujol mulls with a Perkin-Elmer UV/VIS/NIR spectrometer Lambda 19. Compounds were tested for fungicidal activity for wood decay fungi *Trametes versicolor* (L. ex Fr.) Pilat and *Antrodia vaillantii* (DC. ex Fr.) Ryv. as described previously.

RESULTS AND DISCUSSION

Description of the Crystal Structures

The coordination sphere in *Cu(OOCCCH₃)₂(L₁)₂* (*L₁* = 2,6-diaminopyridine) (1) is composed of two didentate acetate anions and two monodentate *L₁* molecules in *cis* conformation around the central copper(II) ion, forming a distorted octahedron (Figure 1). The crystallographic details are summarized in Table I. In the basal plane of the octahedron, one oxygen atom from both acetate groups is present (*Cu-O₁1* 2.007(3) Å, *Cu-O₂₁* 1.971(3) Å) and the pyridine nitrogen atom from both pyridine-based ligands.

![Figure 1. *cis* arranged ligands in a distorted octahedral environment around copper ion in *Cu(OOCCCH₃)₂(L₁)₂* (1), with numerous H-bonds in 3D network.](image)

<table>
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<th>Table I. Crystal data and data collection summary</th>
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ands (Cu–N2 2.013(3) Å, Cu–N1 2.025(3) Å). Significant difference is noticed in the distances of the remaining acetate oxygen atoms, that are semi-coordinated (Cu–O12 2.488(3) Å, Cu–O22 2.657(3) Å) on axial positions. The structure is stabilized by several hydrogen bonds among acetate oxygen atoms and amino groups from the pyridine ligands. The closer axial oxygen atom O12 participates in three H-bonds, intra-molecular to two pyridine ligands (N11–H···O12 3.119(5); N21–H···O12 2.893(5) Å) and one inter-molecular (N12–H···O12 2.900(4) Å). The semi-coordinated oxygen atom from the other acetate ion is forming one intra-molecular H-bond (N12–H···O12 2.910(5) Å) and one inter-molecular H-bond (N11–H···O22 3.035(4) Å). Only one oxygen atom in the first coordination sphere (Cu–O ≈ 2.0 Å), participate in intra-molecular H-bond (N21–H···O21 3.040(4) Å). Interestingly three amino groups are included (Figure 1), each in two H-bonds, while for the fourth (N22) no such interactions were noticed. Extensive hydrogen-bonding network is obviously playing a key role in the asymmetric cis arrangement of the ligands in 1.

The crystal structure analysis of [Cu(OOCC₆H₅)₂(L₂)] (6) revealed trans arrangement of benzoate anions and 2-amino-6-methylpyridine molecules L₂ around the central copper(II) ion, lying on the inversion center. Like in 1, one oxygen atom from both carboxylate anions (Cu–O 1.9580(12) Å) and pyridine nitrogen atom from 2-amino-6-methylpyridine molecules (Cu–N 2.028(14) Å) are composing the basal plane, while the remaining benzoate oxygen atoms from the two benzoates (Cu–O12 2.7648 (16) Å) are occupying the axial positions. Similar geometry was observed in 2.⁴ The amino group is forming an intra-molecular H-bond to the coordinated oxygen atom (N–H···O11 2.926(3) Å), and an inter-molecular H-bond to the semi-coordinated oxygen atom (N–H···O12 2.947(2) Å). Additional stabilization is found trough π-stacking interactions between the aromatic rings of 2-amino-6-methylpyridine molecules (Figure 2) and also between benzoates; pyridine rings: d(c₇–c₇) 3.9270(12) Å, α 0 °, β 26.53 °, benzene rings: d(c₇–c₇) 3.9142(22) Å, α 0 °, β 24.98 °, c₇ – centroid of the ring, α – angle between the planes of the rings, β – angle between c₉–c₈ vector and normal to plane of one ring.

The copper(II) ions in [Cu₂(OOCC₆H₅)₄(L₂)₂] (5), are bridged by four benzoate anions via carboxylate groups. The Cu–Cu distance of 2.7233(15) Å, is typical for the dinuclear paddle wheel type of copper coordination compounds.¹² The axial positions in the isolated dinuclear units are occupied by pyridine nitrogen atoms of 2-amino-6-methylpyridine molecules (Cu–N1 2.283(7), Cu–N2 2.284(8) Å). Each pyridine based ligand is forming an intra-molecular H-bond to one carboxylate oxygen atom (N11–H···O11 3.019(13) Å, N22–H···O32 3.133(12) Å). Both H-bonds in a dinuclear unit are trans oriented with respect to the center of the tetracarboxylate central core (Figure 3). Pyridine rings in the same dinuclear unit are almost co-planar (11.1 °). The pyridine rings are involved also in inter-dinuclear π-stacking interactions (d(c₉–c₇) 3.7109(80) Å, α 0.03 °, β 19.33 °), which influence the crystal packing in the structure.

**Magnetic Measurements and Spectroscopy**

The results of magnetic measurements are in agreement with mononuclear (μₑffective > 1.73 BM, for 1, 2,⁴ 6) and dinuclear (μₑffective < 1.73 BM, for 3,⁵ 4, 5) nature of the compounds.

The electronic spectra of the dinuclear complexes 3–5 are similar, representing the bands of the d-d transition at 730 nm and LMCT at 400 nm and 320 nm, as observed in the spectrum of copper acetate hydrate.¹³ Nev-
ertheless, the UV region band partially originates also in π–π transitions, since it is observable in the spectra of pure ligands.

In the spectra of mononuclear compounds 2 and 6 (trans oriented ligands), two d-d transition bands were observed (∼550 nm d_{xz}, d_{yz} → d_{x^2–y^2}, ∼700 nm d_{xy} → d_{x^2–y^2})^{13,14} and two LMCT transitions in the region towards higher energy (320, 400 nm).^{14} Such observations are in agreement with small rhombic deviations from axially elongated octahedral environment (Cu–O1 < Cu–N < Cu–O2), found in 2 (Ref. 4) and 6. On the other hand, only two bands (d-d at 580 nm, LMCT at 340 nm) were noticed in the spectrum of 1, where cis arrangement of the ligands was found and less rhombic distortion of elongated axial octahedron (Cu–O1 ~ Cu–N < Cu–O2). The presence of two signals instead of four is probably a consequence of smaller energy difference between orbitals from the electron absorption to d_{x^2–y^2} take place (d_{xy}, d_{xz}, d_{yx}, d_{x^2} for d-d and ligand orbitals for LMCT transitions). In the other monomeric cis complex [Cu(OOCCCH3)2(2-aminopyridine)2]^{15} also very small rhombic distortion was found, however the distance to axial semi-coordinated oxygen atoms do not differ so much as in 1. In the spectrum of 2-aminopyridine complex, only one band was found in the 450–800 nm region as for 1, while two bands in the lower energy region (310, 410 nm) as in 2 and 6.

NH2 stretching vibrations in the IR spectra of dinuclear complexes 3–5 are noticed at 3500 cm⁻¹ (asymmetric ν(NH2)as) and at 3380 cm⁻¹ (symmetric ν(NH2)s), as found for their analogues with 2-aminopyridine.^{16} Both signals (ν(NH2)as, ν(NH2)s) split to three signals in the spectrum of solvated complex 3. An analogy to 2-aminopyridine complexes was observed also for mononuclear compounds, where ν(NH2)as and ν(NH2)s are found at lower energies as for the dinuclear compounds. Additional band at 3436 cm⁻¹ for 6 and two in the same region for 1 are probably due to H-bonding network. In the region between 1800 and 1300 cm⁻¹ the bands for δ(NH2), ν(C–C)ar (ar – aromatic) ν(CO2)as, ν(CH3), and ν(CO2)s, are found (Table II).^{17} Splitting of the carboxylic group stretching bands for 1 is probably related to the nonequivalency of both carboxylate groups around the central copper ion. Similar splitting was noticed for not equivalent acetate groups in cis mononuclear 2-aminopyridine complex.^{16} Lower energy of ν(CO2)s in benzoates than in related acetates is also an analogy to observed behavior in 2-aminopyridine complexes.^{16}

### The Influence of the Compounds on Fungal Growth

The tests on fungal growth for the species *Trametes versicolor* showed stronger activity for benzoate compounds than for acetates at concentration 1.0×10⁻³ mol L⁻¹, as already observed.^{11,18} Stronger retardation for benzoates was noticed also for *Antrodia vaillantii*, already at lower concentration 5.0×10⁻⁴ mol L⁻¹. No significant difference among mononuclear 6 and dinuclear 3–5 benzoates was noticed, therefore the results are not in agreement with predicted stronger fungicidal activity of the dinuclear complexes over mononuclear compounds.^{19} Nevertheless, we should not neglect the possibility, that the structures of solid compounds may change by dissolving in DMSO for the activity tests.

### CONCLUSIONS

Monomeric complexes 1 and 2 (Ref. 4) differ in cis and trans arrangement of the ligands. In the structural database CSD, another two monomeric acetate complexes [Cu(OOCCCH3)2(L)2] with similar ligands 2-aminopyridine L3 (Ref. 15) and 2,6-dimethylpyridine L4 (Ref. 20) were found. The L3 complex is cis type as in 1, while L4 complex is trans type, as found in 2. In the cases of 2,6-dimethylpyridine and 2-methyl-6-aminopyridine, trans orientation is more stable, possibly due to less steric hindrance of methyl groups in trans than in cis conformation. On the other hand, in 2-aminopyridine and 2,6-diaminopyridine, amino groups enable strong hydrogen-bonding network that favorize cis orientation with respect to trans. Interestingly, in the acetate family with these four ligands [Cu(OOCCCH3)2(L)2] (L = L1, L2, L3, L4), only one dinuclear species [Cu2(OOCCCH3)4(L)2] with L = L3, 2-aminopyridine, was found.^{16}

For all four acetate mononuclear complexes (with L1, L2, L3 and L4) no π-stacking interactions were found, however two types (pyridine-pyridine and benzene-benzene) were noticed in the monomeric trans benzoate complex [Cu(OOCCCH3)2(L)2] (6). All the other obtained benzoate complexes with 2-amino-6-methylpyridine and 2,6-diaminopyridine 3–5, show the dinuclear struc-

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**TABLE II. IR (mineral oil mull) wave numbers/cm⁻¹**

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ture, a significant difference from acetates with the same pyridine-based ligands (1, 2).

The comparison of the investigated complexes 1–6, show differentiation of acetates and benzoates in appearance of the mononuclear or the dinuclear complex structures. The reason for one mononuclear benzoate 6 (the others (3–5) are dinuclear) might be in π-stacking interactions, observed in 6 but not in the acetate mononuclear compounds 1 and 2. The leading role of the methyl and amino groups in pyridine-based ligands, was observed in monomeric complexes 1, 2 and 6, where their presence or absence seems to direct the path of the ligand arrangement around the central copper(II) ion in the octahedral coordination sphere.

**Supplementary Materials.** – CCDC– 222673 (1), 222674 (5), and 222672 (6) contain the supplementary crystallographic data for this paper. These data can be obtained free of charge at www.ccdc.cam.ac.uk/conts/retrieving.html on quoting the deposition numbers [or from the Cambridge Crystallographic Data Centre, 12, Union Road, Cambridge CB2 1EZ, UK; fax: (internat.) +44-1223/336-033; E-mail: deposit@ccdc.cam.ac.uk].

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**REFERENCES**


**SAŽETAK**

**Dva tipa piridinskih liganada u mononuklearnim i dinuklearnim kompleksima Cu(II) karboksilata**

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Pripravljeni su i karakterizirani bakar(II) acetati [Cu(OOCCH3)2(L1)2] (L1 = 2,6-diaminopiridin) (1), [Cu2(OOCCH3)4(L2)2] (L2 = 2-amino-6-metilpiridin) (2) te benzoati [Cu2(OOCC6H5)4(L1)2]2CH3CN (3), [Cu2(OOCC6H5)4(L2)2] (4), [Cu2(OOCCH3)2(L2)2] (5), i [Cu(OOCCH3)2(L2)2] (6). Rentgenska je strukturalna analiza pokazala monomere u 1 i 6. Izduženi oktaedarski CuO4N2 kromofor u 1 ima cis–, a u 6 trans-smještene ligande. Glavnu ravinu u 1 i 6 čine po jedan kisikov atom iz obje karboksilne skupine [Cu–O 1,9560(12)–2,007(3) Å] i dušikovi atomi dvaju piridinskih liganada [Cu–N 2,013(3)–2,282(14) Å] tvoreći Cu2O2N2 ravinu, dok drugi kisikov atom karboksilne skupine čini dulju vezu [Cu–O 2,4883(8)–2,7648(16) Å]. Dinuklearnar centralna jedinica konfiguracije mlinskoga kola nađena je u 5 [Cu–O 1,942(6)–1,992(6) Å] s dušikovim atomima piridina u aksijalnim položajima [Cu–N 2,283(7), 2,284(7) Å]. Svi spojevi su karakterizirani magnetskim mjerjenjima, elektronskom i vibracijskom spektroskopijom i provjereni na antifungalno djelovanje.