Scientific paper

Cu(I) Arylsulfonate π-Complexes with 3-Allyl-2-thiohydantoin: The Role of the Weak Interactions in Structural Organization

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Abstract

The present work is directed toward preparation and structural characterization of two novel Cu(I) arylsulfonate π-complexes with 3-allyl-2-thiohydantoin, namely $[Cu_2(Hath)_4](C_6H_5SO_3)_2$ (**1**) and $[Cu_2(Hath)_4](p\text{-}CH_3C_6H_4SO_3)_2\cdot 2H_2O$ (**2**) (*Hath* = 3-allyl-2-thiohydantoin), obtained by the means of alternating current electrochemical synthesis and studied with X-ray diffraction method. In both structures, the inner coordination sphere is represented by the cationic dimer $[Cu_2(Hath)_4]^{2+}$ with one crystallographically independent copper(I) atom which has a trigonal pyramidal coordination environment formed by three *Hath* thiogroup S atoms and double C=C bond of its allyl group. $[Cu_2(Hath)_4]^{2+}$ fragments in both coordination compounds are very similar, despite some divergences such as a big difference in Cu−S distance to the apical S atom (3.0374(8) Å in **1** and 2.7205(9) Å in **2**). This difference was explained by the impact of the system of weak interactions, which are quite different.

Keywords: copper(I) arylsulfonate, π-complex, thiohydantoin, weak interaction, crystal structure.

1. Introduction

In the last years both 2-thiohydantoin (2-thioxoimidazolidin-4-ones) and hydantoin core fragments were studied as useful scaffolds in medicinal chemistry due to their synthetic feasibility and

versatility of substituents. [1-5] Moreover, their derivatives are already used in commercially available drugs such as Phenytoin, Dantrolene and Allantoin. The presence of several active chemical groups, as well as simple synthetic route advantages, could make them interesting not only as biologically active compounds but also as compounds of great interest in the chemistry of coordination compounds too. Although, due to the presence of capable for complex compound formation functional groups, 2-thiohydantoin based compounds were found to have an application in the analytical chemistry as reagents for the determination of several *d*-metals (including Cu and Ag). [6, 7] Despite the small number of such representatives (according to the Cambridge Crystallographic Database only 12 copper coordination compounds with 2-thiohydantoin still known), [8] it was already investigated that they could have a potential interest as optical materials due to their fluorescence sensing properties and luminescence towards Cu^{+} and Cu^{2+} as well as the possibility of successful usage in crystal engineering of organometallics. [9, 10] Moreover, previously, we have shown, that Cu(I) π-complexes with allyl derivatives of heterocycles can possess noticeable non-linear optical properties. [11-15]

This work is the continuation of our previous studies, in which we have studied the coordination behavior of 3-allyl-2-thiohydantoin (*Hath*) regarding Ag(I), where obtained coordination compounds have already shown interesting structural peculiarities, confirming the status of 2-thiohydantoin-based molecules as the potential ligands in the structural engineering. [16, 17]

2. Experimental

2.1 General consideration

Unless otherwise mentioned, all chemicals were obtained from a commercial source (Sigma Aldrich) and used without further purification. The NMR experiments: ^{1}H NMR (500 MHz), ^{13}C{^{1}H} NMR (125 MHz) spectra for *Hath* were recorded on a Bruker Avance 500 MHz NMR spectrometer. The chemical shifts are reported in ppm relative to the residual peak of the deuterated CD_3OD for the ^{1}H and ^{13}C{^{1}H} NMR spectra. The infrared (IR) spectrum for *Hath* was recorded on the Bruker IFS-88 spectrometer as nujol mulls. Diffraction data for **1** and **2** were collected on a Gemini+ diffractometer with Mo K_a radiation (λ = 0.71073 Å) and Atlas CCD detector.

2. 2. Preparation of 3-allyl-2-thioxoimidazolidin-4-one (Hath)

Ligand *Hath* was synthesized from allylisothiocyanate and glycine at the presence of triethylamine and pyridine, in accordance with the reported method. [16]

^{1}H NMR (500 MHz, CD_3OD) *δ*, 5.84 p.p.m. (ddt, *J* = 17.1, 10.4, 5.6 1H, =CH), 5.18 p.p.m. (ddd, *J* = 17.1, 2.9, 1.5 Hz, 1H, CH_2=), 5.14 p.p.m. (ddd, *J* = 10.5, 2.5, 1.0 Hz, 1H, -CH_2=), 4.38 p.p.m. (dt, *J* = 5.6 Hz, 1.5 2H, CH_2), 4.14 (s, 2H, CH_2).

^{13}C{^{1}H} NMR (125 MHz, CD_3OD) *δ*, p.p.m. 185.74 p.p.m. (-C=S), 174.17 p.p.m. (-C=O), 132.62 p.p.m. (=CH), 117.94 p.p.m. (CH_2=), 49.48 p.p.m. (CH_2), 43.80 p.p.m. (CH_2).

IR (nujol, cm^{-1}): 3488 (w), 3225 (s), 3091 (w), 3011 (vw), 1864 (vw), 1751 (vs), 1650 (m), 1524 (vs), 1431 (vs), 1367 (w), 1344 (vs), 1306 (s), 1289 (w), 1260 (s), 1176 (vs), 1106 (m), 1048 (m), 1029 (m), 994 (m), 977 (w), 930 (s), 893 (m), 755 (vw), 719 (w), 700 (vs), 610 (m), 581 (m), 563 (m), 541 (m), 515 (m), 474 (m), 440 (vw).

2. 3. Preparation of complexes

*2. 3. 1. Preparation of $[Cu_2(Hath)_4](C_6H_5SO_3)_2$ (**1**)*

To the solution of $Cu(C_6H_5SO_3)_2 \cdot 6H_2O$ (0.157 g, 0.4 mmol) in 4.5 ml of *n*-propanol 0.156 g (1 mmol) of 3-allyl-2-thiohydanthoine (*Hath*) was added and obtained mixture was stirred. The resulting orange-brown solution was placed into a 5 mL test tube and then copper-wire electrodes in cork were inserted. The inner mixture in the obtained cell was subjected to alternating-current electrochemical recovery (0.6 V, 50 Hz) for 5 days. [18] Crystals **1**, suitable for X-ray diffraction studies, were formed on copper wires while maintaining this reactor for 2 weeks at -3°C.

*2. 3. 2. Preparation of $[Cu_2(Hath)_4](p\text{-}CH_3C_6H_4SO_3)_2 \cdot 2H_2O$ (**2**)*

To 5 ml of a solution of 0.198 g (0.4 mmol) of $Cu(p\text{-}CH_3C_6H_4SO_3)_2 \cdot 6H_2O$ in *n*-propanol was added 0.156 g (1 mmol) of 3-allyl-2-thiohydanthine (*Hath*) and stirred. The resulting orange-brown solution was placed into a 5 mL test tube and then copper-wire electrodes in cork were inserted. The inner mixture in the obtained cell was subjected to alternating-current electrochemical recovery (0.6 V, 50 Hz) during 2 days. Crystals **2**, suitable for X-ray diffraction studies, were formed on copper wires.

2. 4. X–Ray crystal structure determination

The collected data for **1** & **2** were processed with CrysAlis Pro program. [19] The structures were solved by dual-space algorithm using SHELXT and refined by least squares method on F^2 by SHELXL-2014 with the following graphical user interfaces of OLEX^2. [20-22] Atomic displacements for non-hydrogen atoms were refined using an anisotropic model. Hydrogen atoms were placed in ideal positions and refined as riding atoms with relative isotropic displacement parameters. The figures were prepared using DIAMOND 3.1 software. Crystal parameters, data collection and the refinement parameters are summarized in Table 1.

Table 1. Selected crystal data and structure refinement parameters of **1** and **2**.

	$[Cu_2(Hath)_4](C_6H_5SO_3)_2$ (**1**)	$[Cu_2(Hath)_4](CH_3C_6H_4SO_3)_2\times$ $\times 2H_2O$ (**2**)
Formula weight ($g\cdot mol^{-1}$)	533.11	565.15
Crystal system and space group	Triclinic, $P\ \bar{1}$	Triclinic, $P\ \bar{1}$

a(Å)	9.1664(6)	9.4494(3)
b(Å)	10.7387(6)	10.3035(4)
c(Å)	12.4008(6)	14.4419(6)
α(º)	93.096(4)	94.228(3)
β(º)	95.581(5)	94.821(3)
γ(º)	114.254(6)	116.983(4)
V(Å^3)	1101.71(12)	1238.64(9)
Z	2	2
D (g/cm^3)	1.607	1.515
μ (mm^{-1})	1.31	1.18
F(000)	548	584
Crystal size (mm)	0.53×0.42×0.34	0.60×0.36×0.25
θ range for data collection (º)	3.7 – 28.9	3.5–28.8
Index ranges	$-12 \le h \le 9$ $-14 \le k \le 13$ $-16 \le l \le 16$	$-11 \le h \le 11$ $-13 \le k \le 13$ $-19 \le l \le 18$
Measured reflections	7786	8562
Independent reflections	4534	5143
Reflections with $I > 2\sigma(I)$	3954	4274
Refined parameters	288	302
R_{int}	0.022	0.024
$R[F^2 > 2\sigma(F^2)]$	0.038	0.047
$wR(F^2)$	0.091	0.132
$\Delta\rho_{max}/\Delta\rho_{min}$(e/Å^3)	1.11, −0.65	1.04, −0.78

3. Results and Discussion

π-Complex $[Cu_2(Hath)_4](C_6H_5SO_3)_2$ (**1**) forms triclinic crystals in the centrosymmetric space group $P\ \bar{1}$. This compound is composed of the centrosymmetric cationic $[Cu_2(Hath)_4]^{2+}$ dimers (Fig. 1) with one crystallographically independent Cu atom and outer coordination sphere represented by benzenesulfonate anions. Cu(I) atom in **1** has a trigonal pyramidal coordination environment (geometric index τ_4 = 0.80). [23] Sulfur atom and C5A=C6A olefine group from one *Hath* moiety and S centre from another ligand unit form basal plane of the metal ion coordination sphere, and sulfur atom from one more *Hath* molecule is located at the apical position. The distance to the apical sulfur atom Cu1–S1^i is equal to 3.0374(8) Å and is significantly greater than Cu–S distances to the equatorial S1 and S2 atoms (Table 2).

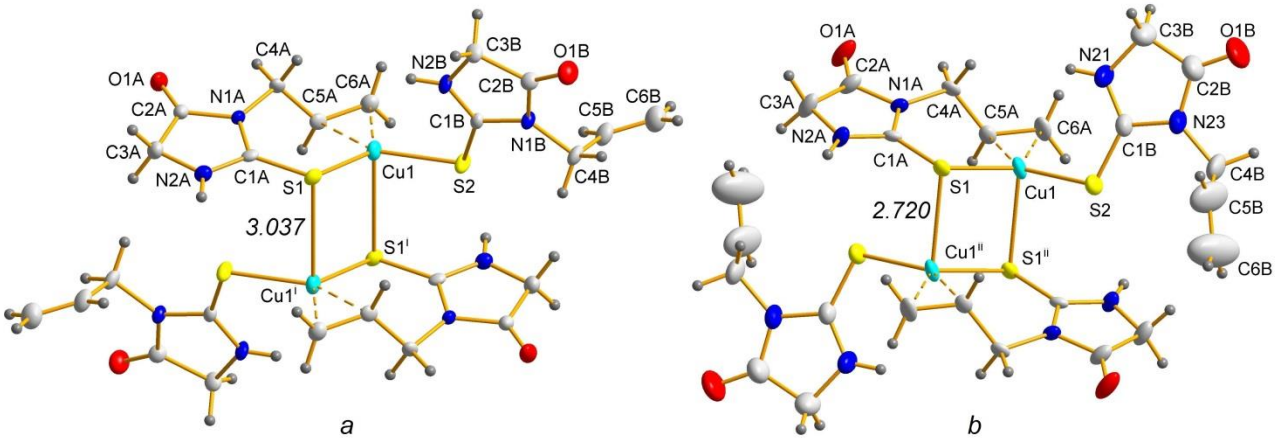

Figure. 1. Cationic fragment $[Cu_2(hath)_4]^{2+}$ in the structures **1** (*a*) and **2** (*b*).
Symmetry codes: (*i*) 1−*x*, 2−*y*, 1−*z*; (*ii*) 2−*x*, 1−*y*, 1−*z*.

In the mentioned cationic fragment $[Cu_2(Hath)_4]^{2+}$ there are two crystallographically independent molecules of organic ligand *Hath*. One of them is coordinated to the Cu atom by the thiogroup S1 atom and double C=C bond of the allyl group. Thus, the first ligand molecule possesses a bidentate chelate function, forming a seven-membered {C_4NSCu} cycle. The aforementioned S1 atom of the first ligand is also coordinated to the $Cu1^i$ atom, and, due to the cetrosymmetricity of the structure, a flat four-membered {Cu_2S_2} cycle with a S1–Cu1–$S1^i$ angle of 97.04(3)° is formed. Second ligand molecule is coordinated to the Cu1 atom only through its S2 atom, complementing copper's coordination number to four. Accordingly – allyl group of the second ligand molecule doesn't participate in the metal bonding and is freely located in the crystal structure with anticlinal conformation (119.8(3)º) relative to the C4B–C5B bond.

The structure of the complex $[Cu_2(Hath)_4](p\text{-}CH_3C_6H_4SO_3)_2\cdot 2H_2O$ (**2**) is quite similar to the structure **1**. It has similar cationic $[Cu_2(Hath)_4]^{2+}$ fragments, but the outer coordination sphere is filled with *p*-toluenesulfonate anions and water molecules (Fig. 3). Regarding the cationic fragment, the main differences are in the noticeable shortening of the Cu−S distance to the apical atom (2.7205(9) Å in **2** compared to 3.0374(8) Å in **1**) (Table 2) and a slightly different conformation of the uncoordinated ligand molecule, in particular, the torsion angle N23−C4B−C5B−C6B is equal to 142.6(7)º (in contrast to 119.8(3)º in **1**). These differences can be explained, taking into account some features of the differences in the outer coordination sphere (Fig 2).

Table 2. Selected bond lengths (Å) and angle values (º) in **1** and **2**.

	1	**2**
Bond	*d*, Å	
Cu1–S1	2.2573(8)	2.2785(8)
Cu1–S2	2.2556(7)	2.2644(9)
Cu1–$S1^X$	3.0374(8)	2.7205(9)

Cu1–m^*	1.978(3)	1.994(3)
C5A–C6A	1.361(4)	1.363(5)
Angles	ω, °	
S2–Cu1–S1	112.97(3)	112.06(3)
S1–Cu1–$S1^{X**}$	97.04(3)	97.21(3)
m–Cu1–S1	117.03(8)	116.37(3)
m–Cu1–S2	129.53(8)	128.32(3)

$^*m$ – middle point of C5A–C6A bond; $^{**}S1^X$ – $S1^i$ atom for **1** and $S1^{ii}$ atom for **2**

Symmetry codes: (*i*) 1−*x*, 2−*y*, 1−*z*; (*ii*) 2−*x*, 1−*y*, 1−*z*.

Benzenesulfonate anions are located in the outer coordination sphere and take part in the formation of weak bonding in the structure **1**. Two out of three oxygen atoms of the same anion (O13 and O33) participate in N–H···O bonding (Table 3, Fig. 2) with N–H groups of different cationic fragments, connecting them into infinite H-bonded chain (Fig 3). Also, one of the oxygen atoms of $C_6H_5SO_3^-$ anion (O13) forms a weak Cu–O contact with a bond length of 3.409(5) Å. Nevertheless, this distance is noticeably longer than the sum of van der Waals radii of Cu and O by Bondi (2.92 Å), [24, 25] it is still less than the corresponding value according to both Batsanov and Alvarez studies - namely 3.55 Å and 3.88 Å respectively. [26, 27]

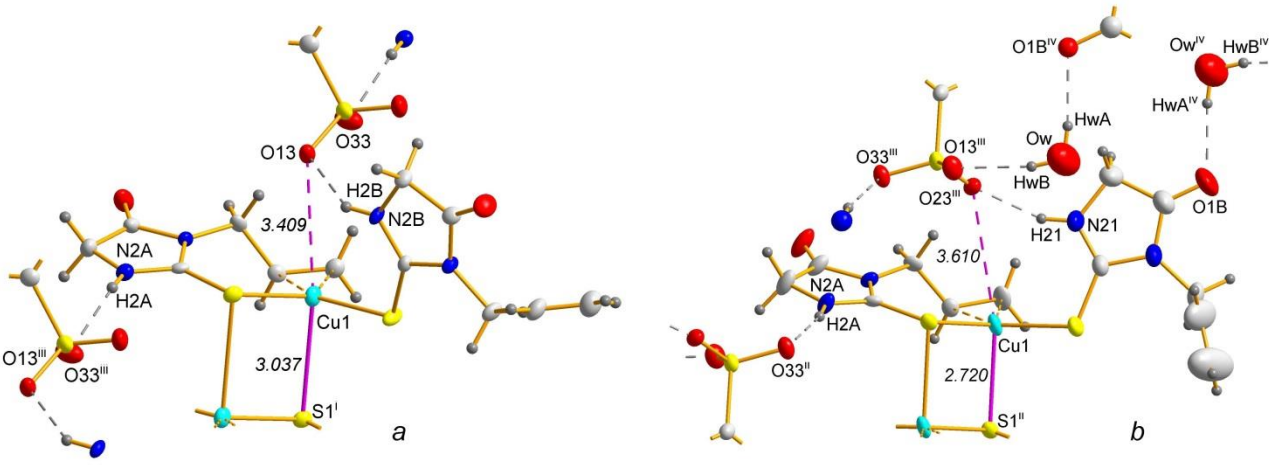

Figure. 2. Systems of weak bonding in the structure **1** (*a*) and **2** (*b*).

Symmetry codes: (*i*) 1−*x*, 2−*y*, 1−*z*; (*ii*) 2−*x*, 1−*y*, 1−*z*; (*iii*) *x*−1, *y*, *z*; (*iv*) 1− *x*, −*y*, −*z*.

Table 3. Geometry of selected hydrogen bonds in **1**.

Atoms involved	Distances, Å			Angle, deg
D–H···A	D···H	H···A	D···A	D–H···A
N2A–H2A···$O33^{iii}$	0.86	1.99	2.743(3)	146
N2B–H2B···O13	0.86	2.01	2.774(3)	148

Symmetry code: (*iii*) *x*−1, *y*, *z*.

The system of hydrogen bonding in **2** is much more complicated than in **1** due to the presence of the water molecule (Table 4, Fig 3). As one of the results of its presentence in **2** all three anion's O atoms participate in a formation of hydrogen bonding – two of them (O23 and O33) form N–H···O bonding, like in **1**, and the third one (O13) is connected with water molecule H*w*B atom. Second water hydrogen H*w*A atom is involved in O*w*–H*wA*···O1*B*^{iv} H-bonding with carbonyl C=O *Hath* group. As a result of all bonding, 2*D* H-bonded net is formed (Fig 3). Also, like in **1**, there is a weak Cu–O bonding but in structure **2**, the corresponding distance is noticeably greater and is equal to 3.610(3) Å, which is still less than the sum of van der Waals radii by Alvarez (3.88 Å).

Table 4. Geometry of selected hydrogen bonds in **2**.

Atoms involved	Distances, Å			Angle, deg
D–H···A	D···H	H···A	D···A	D–H···A
N21–H21···O23^{iii}	0.88	1.93	2.761(4)	156
N2A–H2A···O33^{ii}	0.88	1.91	2.764(3)	164
O*w*–H*wA*···O1*B*^{iv}	0.85	2.00	2.841(6)	173
O*w*–H*wB*···O13^{iii}	0.85	2.08	2.914(5)	168

Symmetry codes: (*ii*) 2−*x*, 1−*y*, 1−*z*; (*iii*) *x*−1, *y*, *z*; (*iv*) 1−*x*, −*y*, −*z*.

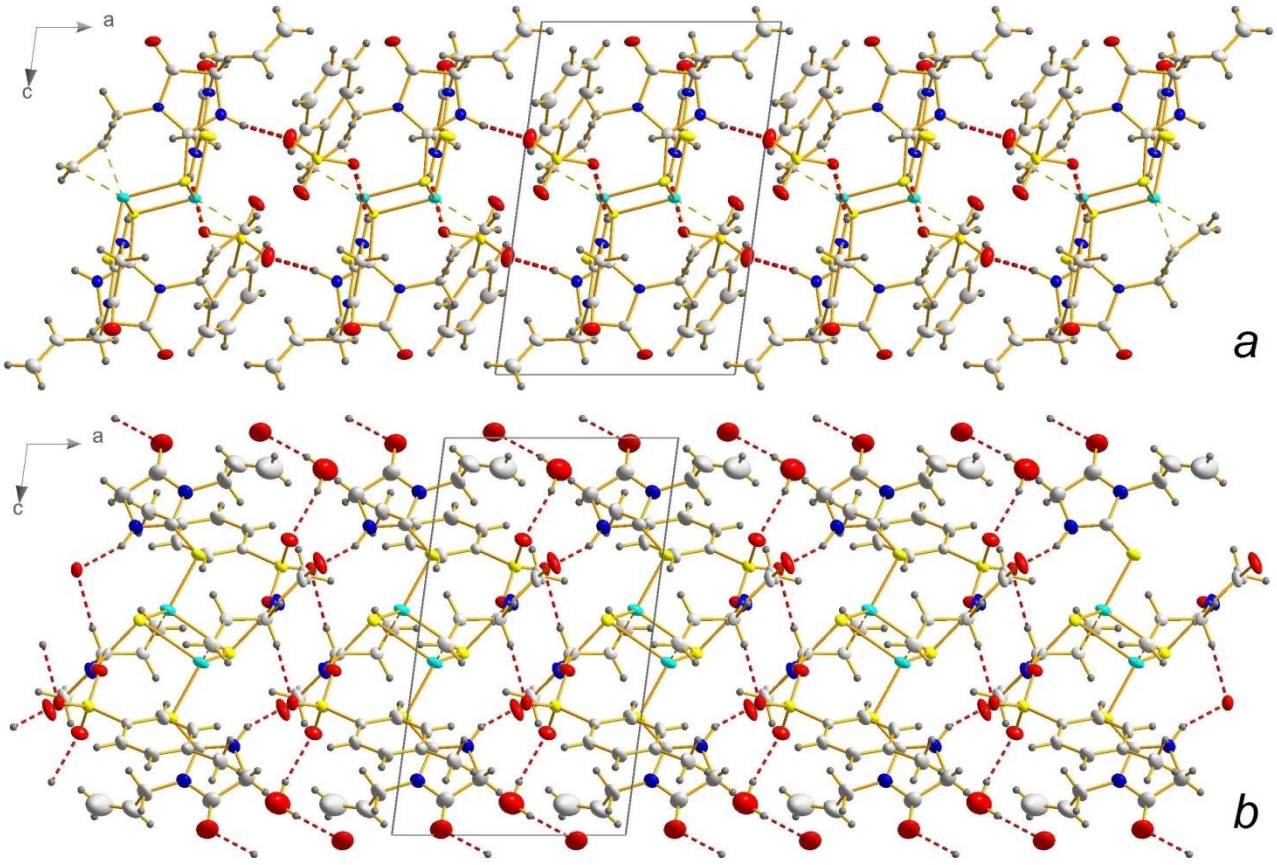

Figure. 3. 1*D* H-bonded chain in **1** (*a*) and 2*D* H-bonded net in **2** (*b*).

As it was shown before, Cu1–S1^{X} distance to the apical S1^{X} atom (X = *i* in **1** and *ii* in **2**) is noticeably different in these two structures. Moreover, since all other distances and angles within the coordination environment of Cu atom are almost the same (Table 2) as well as a composition of the cationic fragment, it can be concluded that the cause of these differences should be found in the outer coordination sphere. As a possible reason, we would like to introduce the confrontation of Cu–S bonding and Cu–O weak interaction. One can notice that in **1**, where Cu–S interaction is comparably weaker (3.0374(8) Å) Cu–O interaction is stronger (3.409(3) Å) and wise versa – in **2** Cu–S is stronger (2.7205(9) Å), but Cu–O interaction is weaker (3.610(3) Å) (Fig 2). In our previous works we have shown an impact of a weak interactions, including hydrogen bonding, on the structural organization in π-complex compounds. [28-30] The weakening of the Cu–O interaction in **2** (and the strengthening of Cu–S as a result) can be explained by the fact, that the anion in **2** is involved in a bigger number of the H-bonding, due to the presence of the water molecule, all of which competes with the Cu–O interaction. Finally, since the water was presented in both reaction mixtures, but co-crystallized only in **2**, its presentence in its crystal structure should be explained by the different character of benzene- and *p*-toluenesulfonate anions.

4. Conclusions

Two novel copper(I) π-complexes $[Cu_2(Hath)_4](C_6H_5SO_3)_2$ (**1**) and $[Cu_2(Hath)_4]\cdot 2CH_3C_6H_4SO_3\cdot 2H_2O$ (**2**) (*Hath* = 3-allyl-2-thiohydantoine) were synthesized and characterized by the X-ray diffraction method. Both structures contain centrosymmetric cationic dimmer $[Cu_2(Hath)_4]^{2+}$ in which Cu(I) atom has a trigonal pyramidal coordination environment. Nevertheless, both $[Cu_2(Hath)_4]^{2+}$ are very similar, there are some differences, including the difference in the Cu1−S1^{X} distance (3.0374(8) Å in **1** and 2.7205(9) Å in **2**). As a possible explanation, the confrontation of Cu–S and Cu–O weak interaction was proposed. Due to this model this difference is explained by the fact, that the anion in **2** is involved in a bigger number of the H-bonding, due to the presence of the water molecule in it, and act on Cu(I) atom weaker than in **1**.

5. Supplementary material

CCDC numbers 1997084 (**1**) and 1997085 (**2**) contains the supplementary crystallographic data for this paper. Copies of the data can be obtained free of charge on applications to the Director, CCDC, 12 Union Road, Cambridge CB2 1EZ, UK (Fax: int. code +(1223)336–033; e-mail for inquiry: fileserv@ccdc.cam.ac.uk).

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