

## Rhenium(IV)-copper(II) heterobimetallic complexes: synthesis, crystal structure and magnetic properties.

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### Streszczenie

Two Re(IV)–Cu(II) heterometallic complexes  $\{(CuL_{\alpha})[ReCl_4(ox)]\}_n$  (where  $L_{\alpha} = N$ -meso-5,12-Me<sub>2</sub>-7,14-Et<sub>2</sub>-[14]-4,11-dieneN<sub>4</sub>), **1**, and  $(CuL_{\beta})[ReCl_4(ox)]$  ( $L_{\beta} = N$ -rac-5,12-Me<sub>2</sub>-7,14-Et<sub>2</sub>-[14]-4,11-dieneN<sub>4</sub>), **2**, were synthesized. The  $[CuL^{2+}]$  macrocyclic cation is coordinated from above and below by  $[ReCl_4(ox)]^{2-}$  units through the chloro-ligands and creates a chloro-bridged heterometallic Re<sup>IV</sup>–Cu<sup>II</sup> one-dimensional zig-zag chain. Compound **2** can be viewed as a heterobimetallic dinuclear unit, in which the Re(IV)-Cu(II) centers are linked by an oxalato bridge. The magnetic behavior of **1** and **2** has been investigated over the temperature range 1.8–300 K. Compound **1** behaves like a ferrimagnetic {Re(IV)–Cu(II)} bimetallic, one-dimensional chain with intrachain antiferromagnetic coupling. Compound **2** shows a weak antiferromagnetic interaction within the [Re(IV)–Cu(II)] unit along with a strong single-ion anisotropy,  $D(Re) = -63 \text{ cm}^{-1}$ .

### Słowa kluczowe

Chain structure, Chloro-bridged Re(IV) complexes, Macrocyclic complexes, Exchange interaction, Zero-field splitting

### Adres publiczny

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### Strona internetowa wydawcy

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