

Coordination properties of diethyl (pyridin-2-ylmethyl)phosphate (2-pmOpe) ligand with perchlorate transition metal salts: crystal structure of $[\text{Co}(\text{2-pmOpe})_2(\text{H}_2\text{O})_2](\text{ClO}_4)_2$.

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A new series of the perchlorate transition-metal complexes containing the diethyl (pyridin-2-ylmethyl)phosphate (2-pmOpe) ligand of general formula $[\text{M}(\text{2-pmOpe})_2(\text{H}_2\text{O})_2](\text{ClO}_4)_2$ (M = Cu, Ni, Co) were prepared. The complexes were identified and characterized by elemental analysis, spectroscopic and magnetic studies. The ligand containing two donor atoms, heterocyclic pyridyl nitrogen and phosphoryl oxygen atoms, binds in a bidentate chelate manner in all complexes. The crystal structure for $[\text{Co}(\text{2-pmOpe})_2(\text{H}_2\text{O})_2](\text{ClO}_4)_2$ was determined by the X-ray diffraction method. The geometry of CoN_2O_4 chromophore shows an elongated octahedron, resulting from the bidentate N,O-bonded two chelate ligands and two coordinated water molecules. Every mononuclear unit is linked to other ones via two extended $\text{CoOH}\cdots\text{OClO}\cdots\text{HOCO}$ bridges, forming one-dimensional (1D) hydrogen bond system. Additionally, weak $\text{CH}\cdots\text{O}$ and $\pi\text{-}\pi$ interactions link the chains resulting in a three-dimensional (3D) (supramolecular architecture) polymeric network arrangement. Spectroscopic and magnetic results are presented in the light of the crystal structure. Magnetic data of mononuclear Co(II) compound indicate an influence of a large zero-field splitting (ZFS) and very weak intermolecular exchange interaction ($zJ' = -0.48 \text{ cm}^{-1}$) on magnetic behavior. The spectroscopic and magnetic properties of Cu(II) and Ni(II) complexes indicate octahedral and mononuclear structure of both compound

Słowa kluczowe

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