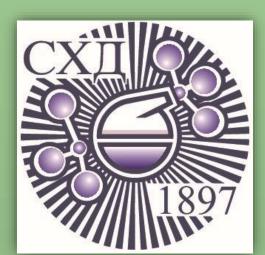
Coordination preferences of NNO and NNS Schiff base ligands with Co(III) complexes

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Two Co(III) complexes $[Co(\mathbf{L}^1)_2]BF_4 \cdot H_2O(\mathbf{1})$, and $[Co(\mathbf{L}^2)(N_3)_3](\mathbf{2})$ with condensation product of thiosemicarbazide and 2-acetylthiazole (\mathbf{HL}^1) and the condensation product of 2-acetylpyridine and Girard's P reagent (\mathbf{HL}^2Cl) and $Co(BF_4)_2 \cdot 6H_2O$ have been synthesized Scheme 1 and 2. Complexes were characterized by elemental analysis, IR and NMR spectroscopy and X-ray crystallographic analysis.

$$\begin{array}{c|c} & & & & \\ & &$$

Scheme 1. Synthesis of complex $[Co(L^1)_2]BF_4 \cdot H_2O$. (1)

octahedral complex in which two deprotonated ligand molecules coordinate in a *mer* arrangement through two NNS sets of donor atoms, while with **HL**²Cl, the ligand is coordinated to Co(III) ion in tridentate fashion through NNO set of donor atoms, and the other three coordination sites of a monokis octahedron are occupied by meridionally coordinated azide anions (2).

Cobalt(III) complex (1) with HL¹ ligand is bis

Scheme 2. Synthesis of complex $[Co(L^2)(N_3)_3]$. (2)

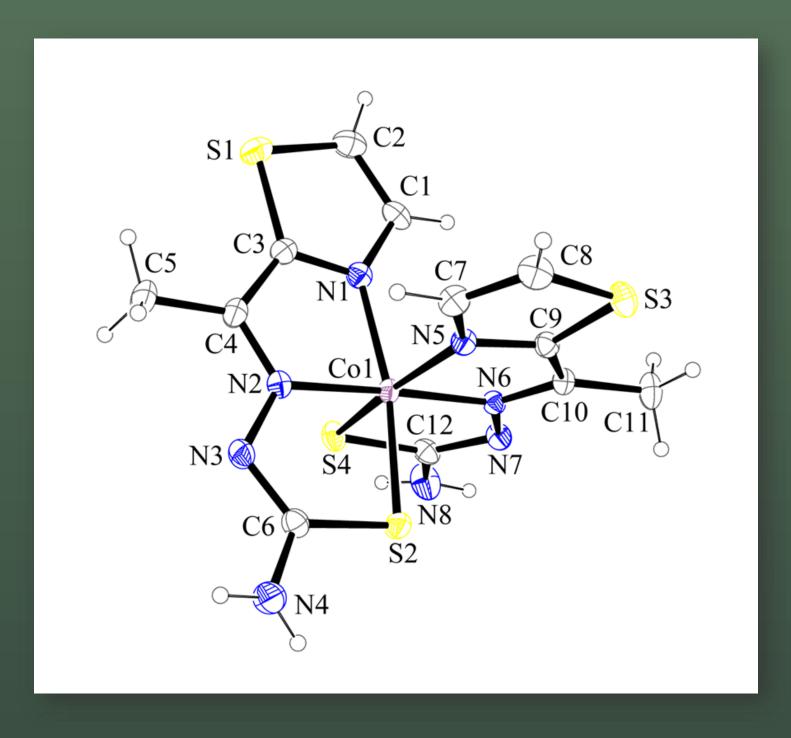


Fig 1. The ORTEP drawing of $[CoL_2^1]^+$ complex cation in 1. Thermal ellipsoids are drawn at the 30% probability level. (1)

Fig 1. The complex 1 crystallizes in the orthorhombic space group Pbca.

The asymmetric unit of 1 consists of $[Co(\mathbf{L}^1)_2]^+$ complex cation, BF_4^- counter anion, and one solvent water molecule.

Fig 2. The complex 2 crystallizes in the triclinic space group P-1.

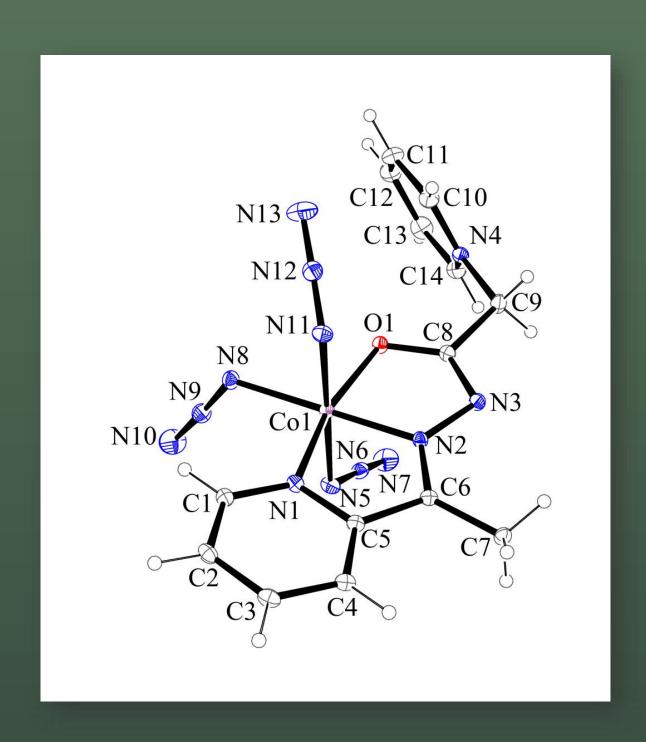


Fig 2. The ORTEP drawing of the $[Co(L^2)(N_3)_3]$ (2) complex. Thermal ellipsoids are drawn at the 30% probability level. (2)