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# Low spin Fe(II) complexes formed of monosubstitued 2,6-bis(2-benzimidazolyl)pyridine ligands

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#### Abstract

Five ligands as monosubstituted derivatives of the basic skeleton of 2,6-bis(2benzimidazyl)pyridine were synthesized, and characterized by NMR and IR spectra along with the X-ray structure analysis. Their complexation gave a set of hexacoordinate Fe(II) complexes showing predominantly the diamagnetism until ambient temperature. Some temperature-independent paramagnetism along with an onset of the spin transition is also detected.

#### Key words

Crystal structure, derivates of 2,6-bis(2-benzimidazolyl)pyridine, Fe(II) complex, magnetic data.

#### 1. Introduction

The condensation product of the *o*-phenylenediamine with pyridine-2,6-dicarboxylic acid results in the formation of 2,6-bis(2-benzimidazolyl)pyridine, abbr. *bzimpy*. The *bzimpy* ligand became one of the most useful starting materials for more complicated organic compounds e.g. anchoring and ancillary ligands used to coordination chemistry [1-3]. The benzimidazole group is also known as a good provider of greater coordination bonding due to strong  $\sigma$  donor effect [4]. This ligand has been widely combined with metal ions yielding a number of transition metal complexes that were studied from various aspects in detail [5]. The high and low spin Fe(II) complexes with six-memberd ring ligands were studied largely [6-13]. One target of investigation was oriented to magnetic studies as Fe(II) complexes of the formula [Fe(*bzimpy*)<sub>2</sub>]X<sub>2</sub>·*sol* are susceptible for spin crossover from the low-spin state S = 0 to the high-spin state S = 2 [14-16]; X – monoanion, *sol* – crystal solvent. Also deprotonation of the basic ligand *bzimpy* to *bzimpy*-<sub>1H</sub> gave neutral Fe(II) complex with interesting magnetic behaviour [17].

Herein we are reporting about synthesis and characterization of new ligands  $L^2$  through  $L^6$  as derivatives of the basic ligand  $L^1 = bzimpy$  (Scheme 1). Their complexation with Fe(II) salts produced a set of hexacoordinate complexes [Fe( $L^n$ )<sub>2</sub>](ClO<sub>4</sub>)<sub>2</sub> and [Fe( $L^n$ -<sub>1H</sub>)<sub>2</sub>] whose magnetic properties, after structure determination, were studied in detail (Scheme 2).

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