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Please use this identifier to cite or link to this item: http://hdl.handle.net/123456789/206 Synthesis, spectroscopic and structural characterization of Co(II), Ni(II) and Cu(II) complexes of Title: substituted 2-pyridyl amine based [N,N] chelating ligand Authors: Prashanth, B. (/jspui/browse?type=author&value=Prashanth%2C+B.) Karanam, M. (/jspui/browse?type=author&value=Karanam%2C+M.) Choudhury, A.R. (/jspui/browse?type=author&value=Choudhury%2C+A.R.) Singh, Sanjay (/jspui/browse?type=author&value=Singh%2C+Sanjay) Keywords: Cobalt complex Copper complex Hydrogen bonding Issue 2012 Date: Publisher: Elsevier B.V. Citation: Polyhedron, 47 (1), pp. 112-117. Abstract: The reaction of N-benzylaminopyridine with the imidoylchloride of N-(2,6-iPr2C6H3)acetamide in the presence of Et3N affords a new neutral [N,N] chelating ligand, (PhCH2)N(2-pyridyl)C{(Me)(N-2,6-iPr2C6H3)} (L). The reaction of equimolar quantities of L with Cu(NO3)2, CuCl2 and NiBr2, respectively, in DCM, acetonitrile and DME yields the corresponding mononuclear complexes L·Cu(NO3)2 (1), L·CuCl2 (2) and L·NiBr2 (3). Whereas, the reaction of L with CoCl2·6H2O leads to the formation of [HL·CoCl3] (4) with pyridine nitrogen coordinated to cobalt. Solid state structure of L and compounds 1-4 have been investigated by single crystal X-ray structural analysis. The ligand L shows the E-anti arrangement in the solid state and its mononuclear complex 1 shows six coordinated Cu in a quasi square planar geometry with two long distanced donors; complexes 2 and 3 show distorted tetrahedral arrangement of the substituents around metal ions. Interestingly, the solid state structure of complex 4 reveals C-H····Cl intra-molecular hydrogen bonding and N-H··· Cl and C-H···Cl inter-molecular hydrogen bonds. These hydrogen bonding interactions in complex 4 facilitate the formation of an extended 2D network structure. URI: http://www.sciencedirect.com/science/article/pii/S0277538712005931 (http://www.sciencedirect.com/science/article/pii/S0277538712005931)  $http://dx.doi.org/10.1016/j.poly.2012.08.019 \ (http://dx.doi.org/10.1016/j.poly.2012.08.019) \ (http://dx.doi.org/10.1016/j.poly.201$ 

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