transition metal elements TM = E8 - E10. In the system Ba - Ni - Ge, two homogeneity ranges with different Ni content are structurally characterized, which are both related to the clathrate-I structure. Compounds with low Ni content, $\text{Ba}_8\text{Ni}_x\text{Ge}_{42\pm1}$ ($0 < x \le 0.6$), can be considered as solution of Ni in the binary $\text{Ba}_8\text{Ge}_{43}\square_3$ (space group Ia-3d, a = 21.307(2) Å). The subcell with space group Pm-3n is observed for the compositions $\text{Ba}_8\text{Ni}_x\text{Ge}_{42\pm1}$ ($0.6 < x \le 4.2$). The change in Ni content is associated with the respective change in lattice parameters (a = 10.657(1) - 10.681(1) Å), but also confirmed by microprobe analyses.

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Keywords: clathrate-I; defects; supercell

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Valence Tautomerism of a Novel Polymeric Mn(II) Species. <u>Francesco Caruso</u>^a, Elizabeth Chacon-Villalba^b, Miriam Rossi^c. ^aIstituto di Chimica Biomolecolare, CNR, Piazzale Aldo Moro 5, 00185, Rome, Italy. ^bCEQUINOR, U.N.L.P. La Plata 1900, Argentina. ^cVassar College, Department of Chemistry, Poughkeepsie, NY, 12604-0484, USA. E-mail: <u>caruso@vassar.edu</u>

 $Mn(Lap)_2$, HLap = lapachol, is a polymeric compound showing extended conjugated double bonds, where the 3 O atoms of lapachol act in a novel way, two chelating one Mn and the 3rd generating a coordinative bond towards another metal unit. X-ray structure determinations at 125K, 197K and 300K describe the progressive trend of a Mn contribution to Mn-O1 bond length as a function of T. The Mn-O1 bond distance increases with temperature and may be therefore associated with a lapachol semiquinonate action at low T by the carbonyl O1 donor (and corresponding to Mn(III)). It transforms to a more classical coordinative bond at room T and stabilizes a Mn(II) species; this is a reversible phenomenon involving Mn(II)-Mn(III) oxidation states. A related UV-visible band shows in the near IR spectrum. The extended network of mesomerism allows for potential transfer of charge through the whole crystal. There are few precedents of this phenomenon in the literature [1].

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Keywords: Mn(II); valence tautomerism; electron transfer

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Structural Analysis, Magnetic Properties of the Bi₂Sr₂Ca(Cu_{2-x}Co_x)O_{8+d} System. N. Boussouf^a, M. F. Mosbah^a, A. Amira^b, T. Guerfi^c. ^aUniversité Mentouri de Constantine, Laboratoire de Couches Minces et Interfaces, Campus de Chaabet-Erssas-25000 Constantine- Algérie. ^bLaboratoire des Essais Non Destructifs (LEND), Université de Jijel. B.P. 98, 18000 Jijel, Algeria. ^cUniversité de Boumerdes,

Faculté des Sciences, département de Physique. E-mail: <u>boussoufnora@yahoo.fr</u>

The effects of Co substitution on the structural and superconducting properties of the Bi₂Sr₂Ca(Cu_{2-x}Co_x)O_{8+d} compound have been studied. as a function of doping concentrations. The $Bi_2Sr_2Ca(Cu_{2-x}Co_x)O_{8+d}$ samples were prepared by conventional solid state reaction method and characterized by X-ray diffraction (XRD), DC resistivity, AC susceptibility, XRD investigations showed multiphase in the Co-substituted samples with slight reflection peaks of impurity and Bi2201 low-Tc (LTc) phase, these configurations suggest the insufficient ionic diffusion to form the superconducting crystals [1]. The refinement result for XRD data shows that the a-axis expands, while the c-axis contract slightly with increasing of Co. The value Tc increased significantly with increasing the Co concentration in the system. The relation between Tc and the c-axis lattice parameter suggests that the change of Tc is correlated with a redistribution of holes between the Bi-O layers and the Cu-O planes [2].



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Keywords: superconductors; Bi2212 doped; magnetic susceptibility