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Chromium(III) Complexes with Different Chromospheres Macrocyclic Ligands: Synthesis and Spectroscopic Studies

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Abstract: Symmetric tetradentate (L¹) and asymmetric pentadentate (L²) macrocyclic ligands that possess 2 dissimilar coordination sites were prepared. One has a 4-coordinate (N_4) and the other has a 5-coordinate (N_2O_3S) donor set. Trivalent Cr(III) complexes were synthesized with L¹ and L² and their structures were investigated using elemental analyses and magnetic moment, electronic, IR, ¹H NMR, ¹³C NMR and EPR spectra. All the Cr(III) complexes show magnetic moments corresponding to a high-spin configuration. Δ values indicate the energy difference between the principle bands, which are formed due to ligand field absorption. The spin-orbit coupling parameter, z, gives no significance because the splitting of doublet transition lines is too large to be explained by spin-orbit coupling. • values indicate that the complexes under study have a substantial covalent character. g-values were also calculated using the spin-orbital coupling constant, •.

Key Words: Macrocyclic ligands, ¹H NMR, ¹³C NMR, IR, EPR, Cr(III) complexes

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