

Novel metal complexes derived from N₂S₂ donor sets; synthesis, structural characterisation and biological activities

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Abstract

The synthesis of ligand systems with N₂S₂ donor sets that include imine, an amide, thioether and thiolate moieties and their metal complexes are achieved. The new Schiff-base ligands; N-(2-((2,4-diphenyl-3-azabicyclo[3.3.1]nonan-9-ylidene)amino)ethyl)-2-((2-mercaptoethyl)thio)acetamide (H₂L¹) and N-(2-((2,4-di-p-tolyl-3-azabicyclo[3.3.1]nonan-9-ylidene)amino)ethyl)-2-((2-mercaptoethyl)thio)acetamide (H₂L²) were synthesised from the reaction of amine precursors with 1,4-dithian-2-one in the presence of triethylamine as a base in the CHCl₃ medium. Complexes of the general formula K₂[M(Lⁿ)Cl₂], (where: M = Mn (II), Co(II) and Ni(II)) and [M(Lⁿ)], (where: M = Cu(II), Zn(II) and Cd(II); n =1-2, expect [Cu(HL²)Cl]) were reported. The entity of ligands and complexes including their purity were confirmed using elemental microanalysis (C.H.N.S), atomic absorption (A.A), chloride content, molar conductance, melting point and thermal analysis technique. while the molecular structures were elucidated with FT-IR, UV-Vis, magnetic susceptibility, ¹H, ¹³C and DEPT ¹³CNMR and mass spectroscopy. The prepared ligands and their complexes were tested against a range of bacterial strains (G⁺ and G⁻) and fungi species. The tested compounds indicated that; the ligands have not shown any antimicrobial activity against *Escherichia coli*. The Cd(II) complex for ligands H₂L¹ and H₂L² display the higher antimicrobial activity, compared with the other complexes. The H₂L¹ and H₂L² ligands have not shown any activity against *Candida albicans*. All complexes for ligands (H₂L¹ and H₂L²) exhibited less activity against *Candida albicans*, compared with other types of fungi. These results are quite comparable with some commercial drugs that used for bacterial and fungi species.