

Copper (II) complexes supported by phenyl selanyl ligands: DNA binding and molecular docking studies

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Copper (II) complexes (**3a-d**) with (16E)-2,6-dimethyl-N-(2-phenylquinoline-4(1*H*)-ylidene)-5-(phenylselanyl)pyrimidine-4-amine ligands have been developed. They have been characterised by elemental analysis and several spectroscopic studies. Absorption spectra, fluorescence investigations, and viscosity tests reveal how the copper complexes interact with the calf thymus (CT-DNA). Furthermore, the ligand's ability to inhibit acetylcholinesterase (AChE) has been studied in order to establish its efficacy in the treatment of neurodegenerative diseases. Compared to normal Rivastigmine and Galantamine, the synthesised ligand **2c** shows selective inhibition (AChE and BuChE) with IC₅₀ values of 0.18 and 3.03 μ M. A molecular docking study has been carried out.

Keywords: Copper (II) complexes, Selenium, Acetylcholinesterase, Quinoline, Alzheimer's disease

Schiff bases bind redox active transition metal ions