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New copper(II) complex with bioactive 2–acetylpyridine-4N-p-chlorophenylthiosemicarbazone ligand:

Synthesis, X-ray structure, and evaluation of antioxidant and antibacterial activity

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Abstract

2-acetylpyridine-4N-*p*-chlorophenylthiosemicarbazone (HL) ligand derived the

condensation of 2-acetylpyridine and p-chlorophenylthiosemicarbazide (pClT), and the

corresponding Cu(II) complex, [Cu(L)Cl] were synthesized and characterized by different

spectroscopic methods. Single crystal X-ray study of [Cu(L)Cl] complex displayed a square-

planar coordination geometry around Cu(II) ion. The in vitro antibacterial activities of pClT, HL

and [Cu(L)Cl] compounds were studied against a series of gram positive and gram negative

bacteria, using the zone inhibition methods. Experimental results revealed that the involvement

of pyridine cycle in the ligand backbone enhanced the antibacterial activity of the

thiosemicarbazone ligand in comparison with its thiosemicarbazide precursor (pCIT). In

addition, the [CuLCl] complex had higher antibacterial activity than the free ligand. Further, the

investigation of antioxidation properties showed that the thiosemicarbazone ligand (HL) and

pClT have considerable in vitro radical scavenging activity over the Cu (II) complex, [CuLCl].

Keywords: Thiosemicarbazones; Copper(II) complex; Biological evaluation; Antibacterial;

Antioxidant.

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