

Área: INO

Antimicrobial activity analysis of coordination compounds containing Schiff Bases

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Highlights

Antibacterial evaluation of Pd/Ag-thiosemicarbazone coordination compounds. Pd-complex W2 showed dual antibacterial activity. Pd-complex W5 was active against *S. aureus* (MIC=16µg/mL). Metal complexes outperformed free ligands.

Abstract

The emergence of multidrug-resistant bacteria is a critical global health challenge, motivating the search for new antimicrobial agents [1]. Schiff base coordination compounds have garnered significant interest in this field due to their diverse biological activities [2]. This study evaluated a series of 12 coordination compounds (10 palladium and 2 silver complexes) and 9 thiosemicarbazone ligands (TSC), organized into five distinct groups based on their composition and molecular structure, with general formulae $[PdCl(N,S-TSC)PPh_3]Cl$, $[PdCl(N,N',C-TSC)]$, $[Pd(N,S,C-TSC)PPh_3]$, $[Pd(N,S,O-TSC)PPh_3]$, and $[AgCl(S-TSC)(PPh_3)_2]$, where PPh_3 = triphenylphosphine and N,S,O,C represent the donor atoms from the thiosemicarbazone ligand (TSC). The antibacterial potential was assessed against *Gram*-negative (*Escherichia coli*, *Pseudomonas aeruginosa*) and *Gram*-positive (*Staphylococcus aureus*) bacteria. The Minimum Inhibitory Concentration (MIC) was determined using the broth microdilution method, following CLSI guidelines [3]. While most compounds and ligands exhibited modest activity (MIC \geq 256 µg/mL), significant efficacy was observed in specific metal complexes. Pd-complex W2 exhibited promising dual activity against both *E. coli* and *S. aureus* (MIC = 32 µg/mL), while Pd-complex W5 displayed notable performance against *S. aureus* (MIC = 16 µg/mL). Crucially, the metal complexes significantly outperformed their corresponding free ligands (MIC > 256 µg/mL), underscoring the essential role of metal coordination in enhancing antibacterial properties. Subculture assays revealed a bactericidal mode of action for Pd-complex W2 and Ag-complex I3, while Pd-complexes A2 and W4 were bacteriostatic. These findings identify specific Pd(II)/Ag(I) thiosemicarbazone complexes as promising candidates for the development of new antibacterial therapies against relevant pathogens.

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