

**SYNTHESIS AND PHARMACOLOGICAL EVALUATION OF SOME
NOVEL 5-(PYRAZOL-3-YL)THIADIAZOLE AND OXADIAZOLE
DERIVATIVES AS POTENTIAL HYPOGLYCEMIC AGENTS**

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Abstract

Four series of 5-(pyrazol-3-yl) thiadiazole and oxadiazole derivatives (Va-c, VIa-c, VIIa-c, IXa-c) have been synthesized from 5-aryloxy-3-carbomethoxy-4-methoxy-1-phenylpyrazole (Ia-c) with a view to investigate their pharmacological activity. The structure of the synthesized products was inferred from elemental and spectral data. The hypoglycemic effect, antimicrobial activity and toxicity of these potential chemotherapeutic agents were evaluated. Nineteen of these products were effective, when administered at an oral dose of 100 mg/kg body weight in inducing a marked reduction in blood glucose level.

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SYNTHESIS

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Synthetic approaches to some new sulfur-containing heterocycles of anticipated immunosuppressive activity.

[Girges, M M](#); [Hanna, M A](#); [Ayyad, S N](#)

Abstract

Several new isolated or fused heterocyclic ring systems that accommodate the isothioureido functionality, often associated with immunosuppressive activity, were synthesized for possible use as immunosuppressive agents. Preparation of these anchored heterocycles was achieved via a multi-step synthesis starting with the key intermediate thiazolyl thiourea derivative (I). Structure of the newly synthesized products was confirmed using both of elemental and spectral analyses.

Address: Chemistry Department, Mansoura University, Egypt.

Research Areas: Pharmacology & Pharmacy; Microbiology; Allergy; Immunology (provided by Thomson Reuters)

Source: Bollettino chimico farmaceutico Volume: 134 Issue: 4 Pages: 204-8 Published: 1995-Apr

Address: Chemistry Department, Mansoura University, Egypt.

Synthetic approaches and biological evaluation of some new sulfonate ester-containing quinazoline derivatives as potentially active antimicrobial agents.

[Habib, O M](#); [Girges, M M](#); [Moawad, E B](#); [el-Shafei, A M](#)

Abstract

Several sulfonate ester-containing quinazolinone derivatives that are substituted with, isolated or fused, aryl, hetaryl or hetarylphenyl ring systems were synthesized with a view to evaluate their efficacy as new bactericidal and/or fungicidal agents. Synthesis of these products involved prior preparation of the 2-[4'-(benzenesulfonyloxy)phenyl]-3,1-4(H)-benzoxazin-4-one derivative (2) and subsequent reaction with several nitrogen nucleophiles. The structure of these products were established from elemental and spectral analyses. Most of the prepared compounds showed high antimicrobial activity, comparable to standard chemotherapeutic agents, when screened against different microorganisms.

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Research Areas: Pharmacology & Pharmacy; Microbiology; Mycology (provided by Thomson Reuters)

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