

Antimicrobial Coumarins from the Oyster Culinary-Medicinal Mushroom, *Pleurotus ostreatus* (Agaricomycetes), from Kenya

Lilechi D. Baraza

Department of Pure and Applied Chemistry, Masinde Muliro University of Science and Technology, Kakamega, Kenya

Wekesa Nesor

Department of Pure and Applied Chemistry, Masinde Muliro University of Science and Technology, Kakamega, Kenya

Korir Cheruiyot Jackson

Department of Biological Sciences, Masinde Muliro University of Science and Technology, Kakamega, Kenya

Juma B. Fredrick

Department of Pure and Applied Chemistry, Masinde Muliro University of Science and Technology, Kakamega, Kenya

Ochieno Dennis

Department of Biological Sciences, Masinde Muliro University of Science and Technology, Kakamega, Kenya

Kamau R. Wairimu

Department of Pure and Applied Chemistry, Masinde Muliro University of Science and Technology, Kakamega, Kenya

Aggrey Osogo Keya

Department of Biological Sciences, Masinde Muliro University of Science and Technology, Kakamega, Kenya

Matthias Heydenreich

Universität Potsdam, Institut für Chemie, Potsdam, Germany

ABSTRACT

Pleurotus ostreatus has been widely used as food because of its nutritional and medicinal properties. These have been attributed to the presence of macronutrients, minerals, vitamins, and amino acids, among other secondary metabolites. There are, however, few reports on the antimicrobial activities of different classes of purified compounds from *P. ostreatus*. This led to the current study, the objective of which was to chemically characterize the antibiotic activities of *P. ostreatus* against selected human pathogenic bacteria and endophytic fungi. Chemical structures were determined using spectroscopic methods and by comparison with values of related structures reported in the literature. Pure compounds from *P. ostreatus* were tested *in vitro* against pathogenic bacteria (*Staphylococcus aureus* and *Escherichia coli*) and endophytic fungi (*Penicillium digitatum* and *Fusarium proliferatum*). A new compound, (*E*)-5,7-dimethoxy-6-(3-methylbuta-1,3-dienyl)-2*H*-chromen-2-one (5-methoxy-(*E*)-suberodiene) (compound 2), along with ergosterol (compound 1) and 5,7-dimethoxy-6-(3-methylbut-2-enyl)-2*H*-chromen-2-one (toddaculin; compound 3), were isolated from the fruiting bodies of *P. ostreatus*. The growth of *S. aureus*, *F. proliferatum*, and *P. digitatum* colonies was inhibited in media containing compound 2, with minimum inhibitory concentrations closely comparable to those of conventional antibiotics.