THE ASSESSMENT OF THE THERAPEUTIC AND TOXICOLOGICAL PROPERTIES OF CARPOBROTUS ACINACIFORMIS AND SCHKUHRIA PINNATA USED IN TRADITIONAL MEDICINE IN SOUTH AFRICA



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Declaration

I, Pasquale Jada Yengkopiong, hereby declare that the work on which this dissertation is based, is original (except where acknowledgements indicate otherwise); and that neither the whole nor any part of it has been, is being, or shall be submitted for another degree at this or any other university, institution for tertiary education or examining body.

PJ Yengkopiong

Dedication

To Alisandro Loku, my father.

To Bernadetha Juka, my mother.

To Mathias Wani, my brother.

Acknowledgements

Although my name appears on the cover of this dissertation as the author, it is the result of the dedicated effort of many individuals, some of whom deserve special mention. My supervisor, Prof. CP Kahler for her patience and commitment to guiding me to reach this point of no return; to her, I owe my future. I would like to acknowledge my former lecturers for having guided me to this point. Today I see my future take a sharp turn to Pharmacology and Therapeutics.

Special thanks and appreciation go to the administration of the University of Limpopo, Faculty of Medicine, Department of Pharmacology and Therapeutics for having given me this opportunity to invest in my future in this institution. You are my pillars on which I put the roof.

I want to mention here earlier that whether this dissertation will be accepted or not, I am confident I shall be acquitted of having acted recklessly. In all I have done I have had some conviction for the faith that is in me, and I am driven by the reasons that guide all of us in research: – first to investigate and second to progress and advance science.

ON mend of count

PJ Yengkopiong.

List of abbreviations

AIDS Acquired immunodeficiency syndrome.

ATP Adenosine triphosphate.

α1PI Alpha-1-proteinase inhibitor.

ACN Acetonitrile.

CAL Carpobrotus acinaciformis leaf.

Ca Calcium.

Cl⁻ Chloride.

Cu Copper.

CO₂ Carbon dioxide.

CNTL Control.

CNTN Contraction.

DNA Deoxyribonucleic acid.

g Gram.

GC Gas Chromatography.

GI tract Gastrointestinal tract.

HIV Human immunodeficiency virus.

HPLC High pressure liquid chromatography.

H₂ Hydrogen.

HOCl Hypochlorous acid.

Kg Kilogram.

K Potassium.

KHCO₃ Potassium bicarbonate.

LDL Low density lipoprotein.

Mg Magnesium.

NADPH Nicotinimide adenine dinucleotide phosphate reduced.

NH₄Cl Ammonium chloride.

Na⁺ Sodium ion.

O₂ Oxygen.

pH Negative logarithm to the base 10 of the hydrogen ion concentration.

PMA Phorbol myristate acetate.

ROS Reactive oxygen species.

rpm Revolutions per minute.

SPL Schkuhria pinnata leaf.

Tris-EDTA Tri-ethylenediaminetetraacetic acid.

TLC Thin layer chromatography.

Zn Zinc.

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Summary

The study assessed the effect of water and ethyl alcohol extracts from different parts of the plants Carpobrotus acinaciformis and Schkuhria pinnata on the contraction of the guinea pig trachea, superoxide production and ATP extraction from human neutrophils and guinea pig whole blood, antibiotic activity against some strains of E. coli and fungi to determine their therapeutic and toxicological effects of when used by humans. The studies with the guinea pigs fed on the water extracts of these plants and those not fed did not show any significant differences in tracheal contraction with either methacholine or histamine, p > 0.05, five degrees of freedom. However, there was a significant decrease in the time of contraction in the tracheas treated with the extracts than those not treated. Superoxide production was decreased in those guinea pigs fed on the plant extracts than those not, p < 0.05, five degrees of freedom. There was also a significant difference in superoxide production by human neutrophils. ATP extraction from the neutrophils was significantly decreased. The water and ethyl alcohol extracts of C. acinaciformis inhibited the growth of all E. coli strains for at least 24 hours; the water extracts of S. pinnata inhibited for only 24 hours and growth resumed thereafter. Ethyl alcohol extracts of S. pinnata inhibited the growth for more than 72 hours. The water extracts of both plants also prevented the growth of fungi in the laboratory settings for at least three months while the ethyl alcohol extracts inhibited the growth for more than a year. The minimum volume of ethyl alcohol required to prevent fungal growth was 10 per cent. Chromatographic techniques showed at least one band or peak and at most five. Thus, the pre-treatment of animals with the extracts of these plants increased the force of tracheal contraction. Treatment of the neutrophils with the extracts reduced superoxide production and ATP extraction. The extracts inhibited the growth of some strains of E. coli and fungi.

Key words

Therapeutic; Toxicological; *C. acinaciformis*; *S. pinnata*; Diseases; Scavengers; Reactive oxygen species; Antioxidants.