



University of Peradeniya
Postgraduate Institute of Science

PROCEEDINGS

**PGIS Research Congress
2022**



VOLUME 9

**POSTGRADUATE INSTITUTE OF SCIENCE
UNIVERSITY OF PERADENIYA
SRI LANKA**



**PGIS RESEARCH CONGRESS 2022
PROCEEDINGS
28th – 30th October 2022**

Copyright © 2022 by Postgraduate Institute of Science

All rights reserved. No part of this publication may be reproduced, distributed, stored in a retrieval system, and transmitted in any form or by any means, including photocopying, recording, or other electronic or mechanical methods, without the prior written permission of the publisher.

ISBN 978-955-8787-09-0

Published by

Postgraduate Institute of Science (PGIS)

University of Peradeniya

Peradeniya

20400

SRI LANKA

**ENZYME INHIBITORY ACTIVITY OF COMPOUNDS ISOLATED FROM
AN ENDOPHYTIC FUNGUS ASSOCIATED WITH *CURCUMA LONGA***

**J. Kalinga¹, K. Samarakoon¹, E.A.I.A. Perera¹, D. Yakandawala²,
N.S. Kumar¹, N.K.B. Adikaram¹, L. Jayasinghe^{1*}, H. Araya³ and Y. Fujimoto^{1,3}**

¹National Institute of Fundamental Studies, Kandy, Sri Lanka

²Department of Botany, Faculty of Science, University of Peradeniya, Peradeniya, Sri Lanka

³School of Agriculture, Meiji University, Kawasaki, Japan

*lalith.ja@nifs.ac.lk

Endophytic fungi are known as treasure houses of natural bioactive compounds. The objectives of this study are to isolate endophytic fungi from *Curcuma longa* L. (Zingiberaceae), known as turmeric, isolate their compounds, elucidate the structures and determine the bioactivities. After triple sterilisation, small rhizome pieces of *C. longa* were placed on Potato Dextrose Agar (PDA). The isolated fungus was tentatively identified as *Fusarium oxysporum* by analysis of ITS1 and ITS4 regions of the rDNA gene. Identification will be confirmed by using other specific gene regions. The fungus was grown on a large scale on Potato Dextrose Broth (PDB). After 4 weeks, the broth and the mycelium were extracted separately into ethyl acetate. After observing the similarities in their Thin Layer Chromatography (TLC), the two extracts were combined. The crude extract was subjected to chromatographic separation (silica gel column chromatography followed by Sephadex LH 20, HPLC and PTLC). This resulted in two pure compounds, and their structures were elucidated using spectral data as fusaric acid (1) and 9,10-dehydrofusaric acid (2). They were subjected to α -amylase, α -glucosidase, lipase and acetylcholinesterase enzyme inhibitory assays. The results revealed that Compound 2 has a strong potential to inhibit α -amylase enzyme activity (IC_{50} of 22.23 ± 4.63 mg L⁻¹) while the positive control acarbose showed IC_{50} of 1.3 ± 0.57 mg L⁻¹. Acetylcholinesterase enzyme activity was strongly inhibited by Compound 1 (IC_{50} of 23.6 ± 5.2 mg L⁻¹) and moderately inhibited by Compound 2 (IC_{50} of 85.8 ± 6.91 mg L⁻¹), whereas the positive control donepezil showed IC_{50} of 0.1 ± 0.01 mg L⁻¹. Both these compounds showed no inhibition of the activities of α -glucosidase and lipase enzymes. Based on these findings, both Compound 1 and Compound 2 have the potential to be used as natural enzyme inhibitors.

Financial assistance from National Science Foundation Research Grant RG/2017/BS/06 is gratefully acknowledged.

Keywords: Acetylcholinesterase, α -Amylase, *Curcuma longa*, Endophytic fungi