<u>ISSN (ONLINE) : 2395-695X</u> ISSN (PRINT) : 2395-695X Available online at <u>www.ijarbest.com</u>



International Journal of Advanced Research in Biology, Engineering, Science and Technology (JARBEST) Vol. 2, Special Issue 8, February 2016 in association with KAMARAJ COLLEGE OF ENGINEERING AND TECHNOLOGY, VIRUDHUNAGAR DEPARTMENT OF BIOTECHNOLOGY ORGANIZES

DBT, NEW DELHI SPONSORED NATIONAL LEVEL CONFERENCE ON CONTEMPORARY TRENDS IN BIOENERGY AND GREEN TECHNOLOGY: CHALLENGES AND OPPORTUNITIES [ORA-2016] (25-26TH FEBRUARY 2016)

INHIBITION OF AGES FORMATION AND CYCLOOXYGENASE -2 BY SODIUM 9-AMINO-4-METHOXYUNDECANOATE FROM LYNGBYA SP

Sameer Kumar Rai^a, Ganeshan Shakambari^a, M. J. AngelaaLincy^a, Prabu Manoharan^b, Balasubramaniem Ashokkumar^c, Perumal Varalakshmi^a* ^aDepartment of Molecular Microbiology, School of Biotechnology, Madurai Kamaraj University, Madurai, Tamil Nadu, India ^bCentre for Excellence in Bioinformatics, School of Biotechnology, Madurai Kamaraj University, Madurai, Tamil Nadu, India ^cDepartment of Genetic Engineering, School of Biotechnology, Madurai Kamaraj University, Madurai, Tamil Nadu, India ^cCorresponding author: pvlakshmi.biotech@mkuniversity.org

Abstract

Microbial natural products are important secondary metabolites which have potentials anti-cancer, antidiabetetic and anti-inflammatory activity. In this study, Lyngbya sp., a marine cyanobacterium, collected from Gulf of Mannar, Rameswaram was prospected for novel bioactive compounds with anti-diabetic and antiinflammatory potentials. The compound was purified from the extracts of Lyngbya sp. by column chromatography and its structure was elucidated by FT-IR, HPLC, ESI-MS and NMR to be Sodium 9-amino-4*methoxyundecanoate* (VS). The anti-oxidant activity potential of the compound was lower IC₅₀ (25.89 \pm 0.21 µg) comparison to standard ascorbic acid (46±0.8 µg) by DPPH method. This compound showed significant antidiabetic and anti-inflammatory activity by in vivo and in vitro. The anti-diabetic activity of the compound was examined in vitro by BSA-glycation inhibition assay where IC_{50} value of VS (16.42±0.28 µg/mL) was lesser then pholoroglucinol used as control. The compound showed anti-diabetic activity in C. elegans used as animal model, *in vivo* conditions, where hyperglycemia was induced in the nematodes by cultivating under high glucose conditions and then treated by the compound VS. The compound VS showed protective effect against AGE formation in hyperglycemia induced *C. elegans*. The effect of purified compound on expression of COX-2 in LPS induced RAW 264.7 macrophage cell lines was further tested using fluorescence activated cell sorting (FACS) analysis, and a significant reduction in COX-2 expression in the induced RAW cells were observed. This study implicates that the VS may be prospected as a potential therapeutic drug against inflammatory diseases. These results showed that VS has significant inhibition against the formation of AGE's in hyperglycemic C. elegans and it also showed potent anti-inflammatory activity by inhibiting COX-2 expression in LPS stimulated RAW 264.7 macrophage cell-lines.

Conclusively, the new bioactive compound VS from *Lyngbya sp.* was explored for protective action against diabetic complication as well as anti-inflammatory responses and hence it is of high interest as a promising natural therapeutic molecule for treating diabetes mellitus and inflammatory diseases.

Keywords: anti-diabetic; COX-2; anti-inflammation.