SULFORAPHANE ATTENUATES NEUROINFLAMMATION, OXIDATIVE STRESS AND COGNITIVE IMPAIRMENT ON ALUMINUM-INDUCED ALZHEIMER’S DISEASE IN WISTAR RATS

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Abstract

Alzheimer’s disease is the most common form of dementia related to neuropathological and neurobehavioral changes. This study is aimed at evaluating the neuroprotective activities of sulforaphane on aluminum chloride induced Alzheimer’s disease in wistar rats. Twenty-eight male wistar rats were divided into four groups (n=7). Group A (control) received normal saline solution orally; Group B received 200 mg/kgbw of Aluminum chloride orally; Group C received 200 mg/kgbw of sulforaphane and 200 mg/kgbw of Aluminum chloride orally, while Group D received 200 mg/kgbw of sulforaphane alone orally. The experiment lasted for 45 days after which behavioural test (Morris water maze, Y maze, open field test) were conducted, serum was obtained for electrolyte concentration analysis (Na, K, Ca, Zn and Cu) and brain tissues were processed for histology, antioxidant parameters (CAT, GSH, SOD), oxidative stress parameters (MDA, H2O2 and Nitrite Levels), brain monoamine neurotransmitters (Dopamine, Serotonin and Norepinephrine), enzymes activity (AChE and Na-K ATPase).

Results showed that Sulforaphane improved learning and memory in rats treated with Sulforaphane and Aluminum chloride compared with Aluminum chloride only group. Also, the Aluminum chloride treated rats showed decreased SOD, CAT and GSH activities as opposed to the sulforaphane treated group with improved antioxidant enzymes level. Furthermore, the levels of MDA, H2O2 and nitrite in Aluminum chloride group showed significant increase as compared to sulforaphane treated group. In addition, administration of sulforaphane with Aluminum chloride improved the monoamine neurotransmitters, brain enzymes activities, serum electrolyte concentration and protects hippocampal histomorphology when compared with Aluminum chloride only group. In conclusion, Sulforaphane ameliorates the neurotoxic effects of aluminum induced rat model of Alzheimer’s disease by improving memory and learning skills, alleviating the neurological disorders thereby acting as powerful antioxidant.

Keywords: Alzheimer’s disease, Antioxidant, Behavioural stress test, Oxidative stress.

Biography

Dr. Babatunde OGUNLADE, completed his PhD at the age of 33 years in June, 2018 from University of Lagos, Nigeria on the thesis titled “Histomorphological and Immunohistochemical Responses to 1-Isothiocyanato-4-(Methylsulfonyl) Butane and D-Ribose-L-Cysteine on Puncture Induced intervertebral Disc Degeneration in Rabbit Model”. He is currently seeking postdoctoral fellowship in research institution outside my country.

He is a lecturer and independent neurobiology researcher at the Federal University of Technology, Akure, Nigeria, in the Department of Human Anatomy, School of Health and Health Technology. He possessed good theoretical and technical laboratory skills such as Electrophysiology, Immuno-cytchemistry, Immuno-histochemistry, Neurobiochemistry, Cell culture, and Computed reconstruction and quantitation (stereology) among others. He have published more than 37 papers in reputed journals. He is positive about academic excellence, motivation for research, professionalism, personal experience, passionate dedication or interpersonal skills.