

3rd International Conference on **Alzheimer's Disease & Dementia**

August 31 - September 02, 2015 Toronto, Canada

Amla enhances autophagy and modulates beta amyloid metabolism in an *in vitro* model of Alzheimer's disease

Elham Teimouri¹, Stephanie R Rainey-Smith^{1,2}, Prashant Bharadwaj^{1,3}, Paul Fraser⁴, Giuseppe Verdile^{1,2,3} and Ralph N Martins^{1,2,5}

¹Edith Cowan University, Australia

²Sir James McCusker Alzheimer's Disease Research Unit (Hollywood Private Hospital), Australia

³Curtin University, Australia

⁴University of Toronto, Canada

⁵University of Western Australia, Australia

Alzheimer's disease (AD) is a progressive, fatal neurodegenerative disease characterized by extensive neuronal loss associated with increased accumulation of the beta amyloid (A β) protein. Reducing production, preventing aggregation and improving clearance of A β are areas of active research in the development of therapeutic agents to ameliorate neurodegeneration in AD. The Indian plant amla (*Emblica officinalis*), commonly known as Indian gooseberry, has widely been utilized in traditional Ayurvedic medicine preparations in the treatment of a variety of disease conditions including cardiovascular disease and diabetes: accumulating evidence also suggests that amla may be beneficial in AD. Amla exhibits antioxidant, anti-inflammatory, and anti-apoptotic mechanisms and more recently has been shown to modulate autophagy; a vital protein degradation pathway involved in the clearance of damaged organelles and aggregate proteins in cells. Our own recent *in vitro* work shows that amla extract enhances autophagy and modulates accumulation of proteolytic products of Amyloid precursor protein (APP) such as APP-C terminal fragments (C99, C83). Amla treatment (50-300 μ g/ml) induced a dose-dependent increase in autophagic flux, as measured by Western blotting utilizing an LC3 directed antibody as an autophagosome marker. At similar concentrations, amla treatment also reduced accumulation of APP C-terminal fragment levels by 33 to 77%. However, no significant changes were observed in APP levels, indicating that amla did not alter APP production. Overall, our findings suggest that amla may confer beneficial effects through modulating autophagy and A β metabolism, and warrants further investigation as a potential therapeutic agent in AD.

Biography

Elham Teimouri is a PhD candidate based at Edith Cowan University in Western Australia, working on a project investigating the utility of nutraceutical compounds as potential therapeutic agents in Alzheimer's disease. Elham is supervised by Professor Ralph Martins, Dr Stephanie Rainey-Smith, Prashant Bharadwaj and Associate Professor Giuseppe Verdile. At the time of writing, Elham has published one peer-reviewed scientific publication with an additional three in preparation.

eteimoori@gmail.com

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