BMC Pharmacology



Poster presentation

Open Access

Ultraviolet B light initiating activity of two-stage mouse skin carcinogenesis and its regulation of cGMP

Harukuni Tokuda*1, Junko Takayasu1 and Akira Iida2

Address: ¹Dept. Mol. Biochem., Kyoto Pref. Univ. Med., Kyoto, Japan and ²Takasaki Univ. Health and Welfare, Takasaki, Japan Email: Harukuni Tokuda* - htokuda@koto.kpu-m.ac.jp

* Corresponding author

from 3^{rd} International Conference on cGMP Generators, Effectors and Therapeutic Implications Dresden, Germany. 15–17 June 2007

Published: 25 July 2007

BMC Pharmacology 2007, 7(Suppl 1):P61 doi:10.1186/1471-2210-7-S1-P61

This abstract is available from: http://www.biomedcentral.com/1471-2210/7/S1/P61

© 2007 Tokuda et al; licensee BioMed Central Ltd.

Ultraviolet (UV) light is the most common cause of skin cancer in humans and it is very cute problem in our life style. Several effects of UVB (290-320 nm) are thought to contribute to skin carcinogenesis. The generation of free radicals and related oxidants produced by UVB exposure. The UVB induced release of biologically active NO from these skin located stroes of NO-donors is biologically relevant and modulates several cellular process, and it increases cGMP formation that converts GTP in a NOdpendent manner. Tabebuia avellanedae (Bignoniaceae) (TA), which is native in South America from Brazil to northern Argentina, is well known in traditional folk medicine used for the treatment of various disease during five hundred years. The inner bark of this plant produced in Brazil is distributed in Asia as a herb tea and healthy purpose. Previously, we reported that extract essense of TA (TA ess.) and including naphthoquinones type compounds, NFD, inhibited TPA-induced in vitro assay. The present study purposed to evaluate for in vivo activity, using natural sourced materials. We have now extended these investigations to a new tumorigenesis model in which we initiated the tumors with UVB irradiation and promoted with 1.7 nmol of TPA in SENCAR mice. Oral feeding of 00025% of TA ess. two weeks before and after tumor initiation resulted in a highly significant reduction in tumor incidence (40%) accompanied by an extension (>20%) of the tumor latency TA ess. treatment also decreased the papilloma incidence and multiplicity when compared with the control and treatment during 20 weeks of the promotion. These results provide a basis for further development of TA ess. for human chemoprevention. To investigate the inhibitory effects of Tabebuia avellanedae and its effect of cGMP levels in mouse surface skin protein. cGMP contents was measured and protein expression was visualized by Western blotting analysis of MAP pathway. We posturate that these data suggest possible role of cancer main pathway and cGMP as regulatory mechanism of potent activity in UVB induced carcinogenesis