

New xanthone and bioactive constituents from *Garcinia subelliptica*

Jih-Jung Chen (陳日榮)^{1*}, Hsiang-Ruei Liao (廖庠睿)², Zou-Chu Chen (陳柔竹)¹,
Hui-Hsuan Chand (張德玟)¹, Yi-Ting Yang (楊依庭)¹, Min-Hsiu Kuo (郭旻修)³,
Yu-Yan Kuo (郭毓晏)⁴, Jhy-Yih Chen (陳志益)¹, and Dau-Chang Wei (魏道昌)¹

¹Department of Pharmacy, Tajen University, Pingtung 907, Taiwan

²Graduate Institute of Natural Products, Chang Gung University, Taoyuan 333, Taiwan

³National Chiayi Senior High School, Chiayi 600, Taiwan

⁴National Chiayi Girls' Senior High School, Chiayi 600, Taiwan

Garcinia subelliptica Merr. (Guttiferae) is a small-to-medium-sized dioecious tree, distributed throughout Philippines, the Ryukyus, and Taiwan. Benzopyrans, biphenyls, benzophenones, xanthenes, benzoylphloroglucinols, phloroglucinols, terpenoids, and their derivatives are widely distributed in plants of the genus *Garcinia*. Many of these compounds exhibit diverse biological activities, including anti-tubercular, anti-inflammatory, and cytotoxic activities. In a preliminary screening, the methanolic extract of the root of this species showed anti-inflammatory activities *in vitro*.

Phytochemical investigation of the stem of this plant has led to the isolation of a new xanthone derivative, garcisubellone (**1**), and four known compounds (**2–5**). The structure of new compound **1** was determined through spectroscopic and MS analyses. The anti-inflammatory effects of the isolated compounds from the roots of *G. subelliptica* were evaluated by suppressing fMet-Leu-Phe (fMLP)-induced O₂⁻ generation by human neutrophils. Among the isolated compounds, 1,4,5-trihydroxyxanthone (**4**) exhibited potent inhibition, with IC₅₀ value of 15.6 ± 2.8 μM, against formyl-L-methionyl-L-leucyl-L-phenylalanine (fMLP)-induced super-oxide anion (O₂⁻) generation.