

2nd International Conference and Exhibition on Pharmacognosy, Phytochemistry & Natural Products

August 25-27, 2014 DoubleTree by Hilton Beijing, China

Anti-inflammatory effects of *M. pudica* (L.) constituents is mediated through inhibition of proinflammatory mediators in macrophages

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In the continuous search for newer anti-inflammatory leads from natural products, many plant extracts have been tested for their inhibitory effects towards pro-inflammatory cytokines. Among them, ethyl acetate fraction (MPE) of *M. pudica* whole plant have demonstrated the significant inhibition of NO, TNF- α and IL-1 β production in RAW 264.7 cells with an IC₅₀ of 34.4, 31.7 and 47.2 µg/mL respectively. Phytochemical investigation of the active ethyl acetate (MPE) fraction of *M. pudica*(L.) leaves yielded fourteen compounds. Among them, L-mimosine (IC₅₀=19.23 to 21.15 µM), crocetin (IC₅₀=23.45 to 25.57 µM) and its ester crocin (IC₅₀=27.16 to 31.53 µM) and jasmonic acid (IC₅₀=29.42 to 21.32 µM) were discovered as potent NO inhibitor when tested on the both cells. Towards TNF- α and IL-1 β inhibition, ethyl gallate, crocetin, crocin, gallic acid, L-mimosine, jasmonic acid and caffeic acid were found to be more active with half maximal concentration, 17.32 to 62.32 µM whereas the other compounds depicted moderate and mild effects (IC50= 59.32 to 95.01 µM). Results suggested that these compounds imparting greatly to anti-inflammatory effects of *M. pudica in vitro* through reduction of LPS-induced pro-inflammatory mediators which affirm the ethno-pharmacological use of this plant for prevention of inflammatory-related disorders.

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