

# Chemical and pharmacological evaluation of the non-flowering aerial parts of *Acacia modesta* Wall. cultivated in Egypt

*Mariam Abd Elhameid , Eman Mohamed Salah , Reham R. Ibrahim and Hesham S.M. Soliman*

## Abstract

**Background:** *Acacia modesta* Wall. (*A. modesta*), often recognized as Phulai, is belonging to family Fabaceae and sub-family Mimosaceae. *A. modesta* has many beneficial uses. Leaves, wood, flowers, and gum of *A. modesta* have been used frequently for multiple therapeutic purposes.

**Results:** The chemical investigation of butanol fraction of *A. modesta* non-flowering aerial parts yielded Vitexin-2''- $\beta$ D-glucopyranoside and Apigenin-6,8-di-C- $\beta$ -D-glucopyranoside in a flavone mixture as well as ( $\beta$ -D-glucopyranosyl (1-3)- $\beta$ -D-glucopyranosyl)-3- $\beta$ -hydroxy-11-oxo-olean-12-en-28-oic acid) an oleanane-type triterpenoidal saponin.

Metabolite profiling via ultra-performance liquid chromatography-electrospray ionization-mass spectrometry (UPLCESI-MS) of the ethyl acetate fraction resulted in recognizing of eighteen compounds tentatively compared with previously published data. Quantitative measurement of the overall value of flavonoids of *A. modesta* was found to be 2.824  $\mu$ g/100  $\mu$ g  $\pm$  0.01 calculated as quercetin. The acute toxicity study of the ethanol extract proved that the plant under investigation is safe and nontoxic to the male albino mice used. The anti-hyperglycemic activity of the ethanol extract performed on type 2 diabetic rats proved that the most potent dosage was 200 mg/kg b. wt. after 4 and 4 weeks of treatment respectively compared to metformin. Furthermore, evaluation of the hepato-protective activity of the ethanol extract of the plant under investigation showed that the most potent extract was with a dose level of 200 mg/kg b. wt. after 3 and 10 days of continuous treatment compared to silymarin.

**Conclusion:** It can be concluded that *A. modesta* Wall. cultivated in Egypt could be used as a promising antidiabetic agent and a hepato-protective agent against hepatocellular damage induced by hepatotoxins.

*Future Journal of Pharmaceutical Sciences 2020, December*