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## Abstract

A new lupene triterpenetriol was isolated from the acetone extract of the aerial parts of *Salvia sclareoides* and characterised as (1 $\beta$ ,3 $\beta$ )-lup-20(29)-ene-1,3,30-triol (**1**). In addition, nepetidin (**2**), nepeticin (**3**), lupendiol (**4**), (1 $\beta$ ,11 $\alpha$ )-dihydroxy-lup-20(29)-en-3-one (**5**), ursolic acid (**6**), sumaresinolic acid (**7**) and hederagenin (**8**), were identified in this *Salvia* sp. To the best of our knowledge, the compounds **2** and **7** are new constituents in *Salvia* spp. The acetone, ethanol, butanol and water extracts of the plant were screened for the in vitro inhibitory activity of acetylcholinesterase (AChE) and butyrylcholinesterase (BChE), enzymes which play a role in the Alzheimer disease. All extracts inhibited acetylcholinesterase activity at 10  $\mu$ g/ml, a remarkable activity since the standard drug rivastigmine does not inhibit acetylcholinesterase at the same concentration. Regarding the butyrylcholinesterase, the acetone extract at 1000  $\mu$ g/ml was able to inhibit completely the enzyme activity and the butanol and ethanol extracts, at this concentration, produced a potent inhibition of BChE.

**Keywords:** *Salvia sclareoides*; Medicinal-plants; essential oil; Triterpenoid; (1 $\beta$ ,3 $\beta$ )-Lup-20(29)-ene-1,3,30-triol; Acetylcholinesterase; Butyrylcholinesterase