ABSTRACT

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Summary extract obtained from dried aerial parts of *Vinca minor* L. was separated by column chromatography with petrol, chloroform and ethanol to 531 fractions. By further separation of fractions, following preparative thin layer chromatography and crystallization 2 alkaloidal compounds marked LB-2 and LB-3 were isolated. These compounds were identified by GC/MS, 1 H- and 13 C-NMR spectroscopy and by use of physical-chemical methods. The structure of compounds were elucidated as indole alkaloids (+)-vincaminoreine (LB-2) and (+)-vincamine (LB-3). Both substances were tested for their inhibition activity against human cholinesterases. (+)-Vincamine didn't exhibited in comparison with standard drugs (galanthamine IC₅₀ AChE: 1,710 \pm 0,065 μ M, IC₅₀ BChE: 42,30 \pm 1,30 μ M; huperzine A IC₅₀ AChE: 0,033 \pm 0,001 μ M) any inhibition activity. On the other hand (+)-vincaminoreine exhibited fairly strong selective inhibition of BChE (IC₅₀ = 8,71 \pm 0,49 μ M) with no inhibition of AChE.

Key words: *Vinca minor* L., indole alkaloids, vincamine, vincaminoreine, Alzheimer disease, acetylcholinesterase, butyrylcholinesterase