ORIGINAL PAPER



α -Glucosidase and α -amylase inhibitory potential of main compounds and drug candidates from *Elaeagnus rhamnoides* (L.) A. Nelson

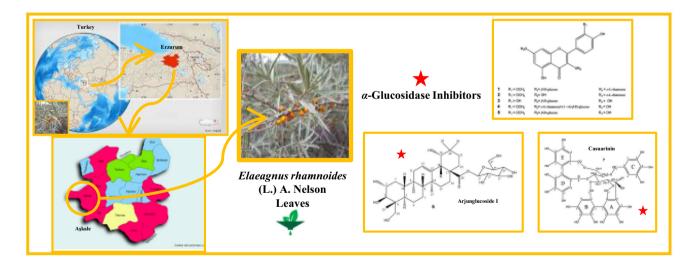
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Abstract

Elaeagnus rhamnoides (L.) A. Nelson (synonym: Hippophae rhamnoides) (Elaeagnaceae) is an important plant with multiple usages. The current study was laid on discovering the phytochemical profiling of *E. rhamnoides* leaves through antihyperglycemic and antioxidant effects. The ethyl acetate (IC₅₀=46.89 ± 2.18 μg/mL) and *n*-butanol extracts (IC₅₀=51.33 ± 2.53 μg/mL) possessed potent inhibitory activity against α-glucosidase enzyme as compared with standard compound, acarbose (IC₅₀=4212.62 ± 130.00 μg/mL). Seven compounds were isolated, and their structure was determined by 1D- and 2D-NMR. Isorhamnetin-3-*O*-β-D-glucopyranosyl-7-*O*-α-L-rhamnopyranoside (1), isorhamnetin-7-*O*-α-L-rhamnopyranoside (2), isoquercitrin (3), narcissin (4), isorhamnetin-3-*O*-β-D-glucopyranoside (5), arjunglucoside I (6), and casuarinin (7) were isolated from *n*-butanol extract. All isolated compounds, especially arjunglucoside I (IC₅₀=1074 ± 32 μM) and casuarinin (IC₅₀=21 ± 2 μM), showed higher α-glucosidase inhibitory activity than acarbose (IC₅₀=6561 ± 207 μM). Casuarinin displayed powerful scavenging activity against to both ABTS radical with 2±1 μM IC₅₀ value and DPPH radical with 14±1 μM IC₅₀ value while IC₅₀ values of trolox and α-tocopherol were 31 ± 1 and 50 ± 1 μM against ABTS radical, and 67 ± 2 and 95 ± 3 μM against DPPH radical, respectively. Arjunglucoside I was isolated for first time from this species and Elaeagnaceae family. Preparations prepared from *E. rhamnoides* leaf extracts standardized via casuarinin and arjunglucoside I could be potential phytotherapeutics for diabetes mellitus.

Graphic abstract



Keywords *Elaeagnus rhamnoides* · Elaeagnaceae · α -Glucosidase inhibition · Antioxidant activity · Arjunglucoside I · Casuarinin

Extended author information available on the last page of the article

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