

## FLAVONOIDS OF AERIAL PARTS OF AN ENDEMIC SPECIES OF THE APIACEAE OF ALGERIA, *Ammoides atlantica*

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The genus *Ammoides* (Apiaceae) tribe of Ammineae includes two species in Algeria, one of which is endemic: *Ammoides atlantica* (Coss. et Dur.) Wolf; the other one, *Ammoides pusilla* (Brot.) Breistr, is widespread in the Mediterranean region [1].

The aerial parts of *Ammoides atlantica* were collected from Megress Mountain (Eastern Algeria) at 1500 m above sea level during June 2004 and identified by Dr. H. Laouer. A voucher specimen (B6308) has been deposited in the Museum d'Histoire Naturelle of Nice (France).

The air-dried powdered parts (700 g) of *A. atlantica* were macerated three times in boiling methanolic solution (70%). The MeOH extract was concentrated to dryness, the residue was dissolved in boiling water (600 mL) after filtration, and the residue was extracted successively three times with AcOEt and *n*-butanol (3×200 mL) to give 2.5 and 22.7 g of the respective residues. Solvents were evaporated and the residues of *n*-BuOH and AcOEt extracts were dissolved in small volumes of MeOH. Analysis by two-dimensional paper chromatography using 15% AcOH and BAW (*n*-BuOH–AcOH–H<sub>2</sub>O, 4:1:5, upper phase) as solvents indicated that extracts of *n*-BuOH and AcOEt contain identical separation of flavonoids. The AcOEt extract was applied to a column of silica gel and eluted with a gradient of *n*-hexane–AcOEt–MeOH with increasing polarity to obtain nine fractions (A–I). Concentration under reduced pressure of fractions B, D, and G gave three precipitates, which were filtered and washed with MeOH to obtain three compounds **1** (10 mg), **2** (32 mg), and **3** (40 mg). Fraction C was purified by preparative TLC on silica gel eluted with *n*-hexane–AcOEt (2:8) to afford compound **4** (10 mg). Final purification of each compound for spectral analysis was carried out using MeOH over Sephadex LH-20 (Pharmacia Fine Chemicals), prior to UV, MS, and <sup>1</sup>H spectral analysis [2, 3]. Hydrolysis of the glycosides **3** and **4** (HCl 0.1 N, 2 h) yielded the sugar residues and the aglycones, all of which were chromatographed with authentic samples. <sup>1</sup>H NMR spectra were recorded in CD<sub>3</sub>OD at 300 MHz. All the structures were in good agreement with literature data.

**Compound 1**, C<sub>15</sub>H<sub>10</sub>O<sub>5</sub>, mp 345°C, identified as apigenin [4].

**Compound 2**, C<sub>15</sub>H<sub>10</sub>O<sub>6</sub>, mp 330°C, identified as luteolin [5].

**Compound 3**, C<sub>21</sub>H<sub>20</sub>O<sub>10</sub>, mp 220–222°C identified as apigenin 7-*O*-glucoside [6].

**Compound 4**, C<sub>21</sub>H<sub>20</sub>O<sub>11</sub>, mp 239–242°C, identified as luteolin 7-*O*-glucoside [7].

Only one flavonoid, luteolin 7-*O*-glucoside [7], has recently been reported from an Egyptian *A. pusilla*. Compounds **1–4** were isolated for the first time from *A. atlantica*, and compounds **3, 4** are reported for a second time from a *Ammoides species* [6, 7]. Compounds **1** and **2** are new for the genus *Ammoides*.

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