Studies on the Constituents of Artemisia annua
Part II

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Abstract

The present paper is a continuation of our study on the Chinese traditional herb Artemisia annua L. [1—5], describing several additional constituents: quinghaosu IV and V (V, VII), quinghao acid (VIII) [6], chrysosplenol (VIa) [7] and a paraffinic alcohol; V, VII and VIII are compounds with unreported structures.

Introduction

A number of our earlier papers have been devoted to studies of chemical constituents isolated from Artemisia annua L. (Compositae), the most notable constituent being the antimalerial quinghaosu (I) [1—4], a peroxidic lactone with unique structure. Other constituents include quinghaosu-I—III (II-IV), a flavonol (VI), scopoletin and a few terpenes from the essential oil [5]. This paper deals with the isolation and characterization of V, VIa, VII, VIII and a paraffinic alcohol.

It is pertinent here to point out the close stereochemical kinship among the previously established structures I—IV. They all belong to the amorphane series (IX) [9], which has a cis decalin skeleton with the isopropyl group trans to the hydrogen on the ring juncture. Compounds I and IV are further distinguished by the presence of a modified seven-membered A-ring as the result of insertion of an extra ether oxygen.
Results and Discussion

Qinghaosu-IV (V) is a colourless crystalline compound with m.p. 172–173°. MS molecular weight 282 (M⁺) agreed with C₁₅H₂₂O₅. Hydroxy and lactonic carboxyl groups can be inferred from its IR spectrum (3450, 1728 cm⁻¹). The ¹H NMR spectrum displayed the following features: δ 0.95 (d, J = 6 Hz, 10-CH₃), 1.20 (d, J = 6, 11-CH₃), 1.65 (s, 4-CH₃), 3.20 (m, H-C-11), 3.60 (br, H-C-3), 5.60 (s, H-C-5) and an exchangeable OH group at 1.88. Except for the two signals at δ 1.88 and 3.60, the NMR spectra of IV and V are almost superposable, thus leading to the conclusion that the extra oxygen of V is in the form of a hydroxyl group with only four possible places (C-2, 3, 8 and 9) for its accommodation. Eu(fod)₃ was used for its allocation. Addition of successive aliquots of the shift reagent gave rise to linear changes of chemical shifts. In the case of qinghaosu III (IV), we have for 10-CH₃, δ 0.95 → 0.96 → 1.16 → 1.16; for 11-CH₃, 1.20 → 1.42 → 2.20 → 2.40; for 4-CH₃, 1.54 → 1.55 → 1.74 → 1.75; and for H-C-5, 5.55 → 5.75 → 6.20 → 6.40. Qinghaosu IV (V) gave the corresponding shifts: 10-CH₃, δ 1.04 → 1.46 → 1.77 → 2.15; 11-CH₃, 1.27 → 1.99 → 2.60 → 3.42; 4-CH₃, 1.65 → 3.93 → 5.74 → 7.25; H-C-5, 5.70 → 6.91 → 8.06 → 9.18. The marked shifts for 4-CH₃ and H-C-5 in compound V is compatible only with an OH group at position-3, and the W₁/₂ 6 Hz) of the H-C-3 multiplet (hence equatorial) indicates an α-orientation for the OH group (axial). The structure of IV has been firmly established by its preparation from qinghaosu (I) by catalytic hydrogenation [5].

Chrysosplenol (6a) has very similar UV and ¹H NMR spectra to eupatin (3, 5, 3'-triOH, 6, 7, 4'-triOMe) [7]. However, large discrepancies in m.p. of the acetates (155–157°; 219–221° for eupatin acetate) indicated α, β-unsaturated lactone functionalities. The ¹H NMR spectrum showed two methyl groups at δ 0.85 (d) and 1.36 (s), and terminal methylene protons at 5.56 (s) and 16.16 (s). The carbonyl hydrogen at C-5 (δ 3.82) is a doublet eith J = 3, hence should be in cis relationship with the hydroxyl on the ring juncture. Further studies have been thwarted by scanty supply of material, and the structure as shown by VII is thus tentatively proposed, leaning heavily on biogenetic considerations.

We also isolated a straight chain fatty alcohol, m.p. 74–76°, characterized by its IR and ¹H NMR spectra. The MS peak at m/e 392 (M-18) [11] showed it to be octacosanol (C₂₈H₅₇OH), probably contaminated by some C₃₀ alcohol (ca. 5 %) as evidenced by a tiny peak at m/e 402. Further fragmentations of interest involved successive losses of 28 units from m/e 392, giving peaks at m/e 364 and 336. The last mentioned peak was stronger than usual [11], indicating the possible contamination by a C₂₀ alcohol. ULBELLEN et al. [12] reported the isolation of a C₂₀ alcohol from the same species, using elemental analysis as the main evidence. Since C₂₀ and C₂₅ alcohols cannot be adequately differentiated by elemental analysis, there is room for the possibility of their sample being also octacosanol.

Experimental

Melting points were not corrected. IR spectra were taken with KBr discs on an IR-S spectrometer. ¹H NMR spectra were taken with CDC₁₃ solutions on WH-90, with TMS as the internal standard. MS were recorded with MM70–70H spectrometer.

Plant Material

Artemisia annua L. is a regular commodity, available in practically all warehouses for Chinese herbs. However, there might well be variations in chemical constituents with different localities, which were therefore specified below.

Silica gel columns and plates were used and eluted with the mixed solvent of petroleum ether and ethyl acetate in individually specified proportions.
**Qinghaosu-IV**

Plant material from Sichuan Province was extracted with petroleum ether and the solvent removed. The crude extract was chromatographed. Qinghaosu-IV displayed only a single spot on TLC (1:1 mixed solvent, 2% phosphomolybdic acid spray). It was purified by recrystallization from ethanol. MS, m/e (%): 305 (M+2, 100), 283 (4), 261 (4), 240 (4), 218 (6), 196 (16), 174 (21), 152 (21), 130 (21), 118 (21), 96 (21), 74 (100).

**Qinghaosu-III**

The crude material was twice recrystallized from ethyl alcohol. MS, m/e (%): 234 (M, 100), 212 (34), 190 (26), 168 (15), 146 (15), 124 (15), 102 (15), 79 (26), 71 (100).

**Qinghaosu-I**

The crude acid displayed a single spot on TLC (9:1 mixed solvent, 2% vanillin-H2SO4 spray). It was purified by recrystallization from ethanol. MS, m/e (%): 382 (M, 5), 360 (75), 338 (31), 316 (16), 294 (20), 272 (11), 250 (11), 228 (10), 206 (11), 184 (11), 162 (11), 140 (11), 118 (11), 96 (11), 74 (100).

**Qinghaosu-IV**

The crude material was isolated from the plant material of the Beijing area. The ethereal extracts were shaken with 2% aqueous sodium hydroxide, which upon acidified by recrystallization from ethanol. MS, m/e (%): 282 (M, 5), 260 (3), 238 (3), 216 (3), 194 (4), 172 (4), 150 (6), 128 (6), 106 (6), 84 (6), 62 (6), 40 (6). The fatty alcohol obtained was an oil. HNMR: 0.85 (d, J = 6.5 Hz), 1.11 (s, 3H), 3.35 (br, 2H), 3.60 (br, 2H), 4.10 (br, 2H), 4.20 (br, 2H). Mass spectrometry: m/e (%) 282 (M, 5), 280 (15), 278 (15), 256 (15), 254 (15), 232 (15), 210 (15), 188 (15), 166 (15), 144 (15), 122 (15), 100 (15), 78 (15), 56 (15), 34 (100).

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**References**


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