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## CONSTITUENTS OF ACANTHOPANAX TRIFOLIATUS

## FA-CHING CHEN, YUH-MEEI LIN and Po-LUNG YU

Chemistry Research Center, National Taiwan University, Taipei 107, Taiwan, Republic of China

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In continuation of our earlier work on the leaves of Acanthopanax trifoliatus, we have examined the stems, isolating KNO<sub>3</sub>, fatty acids, n-pentadecanoic, palmitic, margaric, stearic, and arachidic acids, a mixure of n-alkanes between C<sub>29</sub> and C<sub>31</sub> containing nonacosane as the major component, stigmasterol, sitosterol, and glucosides of the above two sterols, and three unidentified compounds.

## EXPERIMENTAL\*

The air dried and ground stems of Acanthopanax trifoliatus were extracted with benzene and then with MeOH. The MeOH extract was concentrated yielding  $KNO_3$ . The benzene extract was treated with n-hexane. The n-hexane soluble and insoluble fractions were chromatographed on  $SiO_2$  gave fractions I-III, and IV-V respectively.

Fraction I. Colourless plates, m.p.  $58-64^{\circ}$  (CHCl<sub>3</sub>), m/e 436 (M<sup>+</sup>),  $v_{\text{max}}$  (KBr) 2915, 2840, 1475, 1465, 728, 715 cm<sup>-1</sup>. GLC indicated n-C<sub>29</sub>H<sub>60</sub> 69%, n-C<sub>30</sub>H<sub>62</sub> 0·3%, and n-C<sub>31</sub>H<sub>64</sub> 30·7%.

Fraction II. Colourless plates, m.p. 162–164° (acetone), Liebermann–Buchard test positive,  $\nu_{\text{max}}$  (KBr), 3409 (OH), 2960, 2870, 1460, 1380, 1372 (–CH(CH<sub>3</sub>)<sub>2</sub>), 1630, 835, 800 cm<sup>-1</sup> (=C=C=H), 1660, 965 (H=C=C=H). Acetate, m.p. 138–140°. GLC showed a mixture of stigmasterol (97·2%) and sitosterol (2·8%).

Fraction III. Colourless plates, m.p. 54–59° (acetone),  $v_{\text{max}}$  (KBr), 3000–2700, 1700 (–COOH), seven bands in 1350–1180, 720 cm<sup>-1</sup> (–(CH<sub>2</sub>)<sub>n</sub>). Esterification with CH<sub>2</sub>N<sub>2</sub>, and analysis by GLC showed n-C<sub>14</sub>H<sub>29</sub>-COOH 0·5, n-C<sub>15</sub>H<sub>31</sub>COOH 71, n-C<sub>16</sub>H<sub>33</sub>COOH 1·5, n-C<sub>17</sub>H<sub>35</sub>COOH 15, and n-C<sub>19</sub>H<sub>39</sub>COOH 10%.

Fraction IV. Colourless plates, m.p. 151-154° (acetone), Liebermann-Buchard test positive. Analysed by GLC shown to be the mixture of sitosterol (52%) and stigmasterol (48%). Acetate, m.p. 125-127°.

Fraction V. Colourless powder, m.p. 278–280° (benzene–MeOH, 1:1),  $v_{\text{max}}$  (KBr), 3450 br., 2870, 1460, 1380, 1050 br. cm<sup>-1</sup>. Acetate, m.p. 271–273°. Hydrolysis of fraction V with 10% H<sub>2</sub>SO<sub>4</sub>–MeOH (1:1) gave glucose (identified by co-chromatography) and an aglycone mixture identical with fraction IV by IR and GLC.

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- \* GLC was carried out with Varian Aerograph Model 600D; Column, 2 m, 3 % SE-30 on Chromosorb W. Column temp.,  $240^{\circ}$  for alkane and methylester of fatty acid,  $250^{\circ}$  for sterols. Carrier gas:  $N_2$ , flow rate: 25 ml/min.
- <sup>1</sup> F. C. CHEN, Y. M. LIN and S. LIN, Phytochem. 11, 1486 (1972).