TRITERPENE ACIDS OF SOME REPRESENTATIVES
OF THE GENUS Thymus
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We have previously reported the isolation of ursolic acid from Thymus dimorphus Klok [1].
A chromatographic analysis of chloroform extracts from the herbs Thymus dimorphus Klok., Th. nummularius M. B., Th. Kotschyanus Boiss et Hohen, and Th. transcaucasicus Ronn. on paper [solvent systems: 1) petroleum ether-benzene $-\mathrm{CH}_{3} \mathrm{COOH}(5: 20: 2)$; 2) petroleum ether-methanol $-\mathrm{H}_{2} \mathrm{O}$ (10:5:5)] showed that all the species studied contain two substances giving spots with $\mathrm{R}_{f} 0.71$ and 0.81 (system 1) and 0.10 and 0.30 (system 2), respectively. The spots were revealed with a saturated solution of antimony trichloride in chloroform and with a $25 \%$ ethanolic solution of phosphotungstic acid.

A chloroform extract from Th. dimorphus Klok., after the removal of the ursolic acid [1], was evaporated to a dry residue, and this was treated repeatedly with hot methanol. The methanolic extracts were combined and evaporated, and after repeated recrystallization a white crystalline substance with the composition $\mathrm{C}_{30} \mathrm{H}_{48} \mathrm{O}_{3}, \mathrm{mp} 301-304^{\circ} \mathrm{C}$ (from methanol) (A) was obtained.

A chloroformic extract of Th. transcaucasicus Ronn. gave a white crystalline substance with the composition $\mathrm{C}_{30} \mathrm{H}_{48} \mathrm{O}_{3}$, mp 279-282 ${ }^{\circ} \mathrm{C}$ (from ethanol) (B).

Substances $A$ and $B$ gave a positive reaction with chlorosulfonic acid and positive LiebermannBurchard and Sal'kovskii reactions: " The $\mathrm{R}_{f}$ values of substance A coincide with those of an authentic sample of oleanolic acid, and the $\mathrm{R}_{f}$ values of substance $B$ coincide with those for ursolic acid.

For substance $A,[\alpha]_{D}^{20}+76^{\circ}$ (c 0.82 , chloroform), and for substance $B,[\alpha]_{D}^{20}+68^{\circ}$ (c 0.90 , chloroform).
In the UV spectra of sulfuric acid solutions of the substances isolated (sulfuric acid with sp. gr. 1.835; c 0.4) [3], $\lambda_{\text {max }}$ is 310 nm for both substances, but the $\log \varepsilon$ values differ. For $A, \log \varepsilon=4.08$, and for $B, \log \varepsilon=4.11$.

In the IR spectra of both substances, absorption bands appear at $1714 \mathrm{~cm}^{-1}$ ( $>\mathrm{C}=\mathrm{O}$ ) and also bands characteristic for ursolic acid at 1392,1383 , and $1250 \mathrm{~cm}^{-1}$ (geminal $-\mathrm{CH}_{3}$ groups) (substance B) and bands characteristic for oleanolic acid at $1700,1620,1600,1390,1320,1290,1260$, and $1230 \mathrm{~cm}^{-1}$ (substance A) [4].

The product of the acetylation of substance $B$ (in a mixture of pyridine, benzene, and acetic anhydride) has mp $284-286^{\circ} \mathrm{C}$ (from ethanol), $[\alpha]_{\mathrm{D}}^{20}+69^{\circ}$ (c 0.72 , chloroform).

The acetyl derivative of substance A was obtained, with mp $258-260^{\circ} \mathrm{C}$ (from methanol), $[\alpha]_{\mathrm{D}}^{20}+79^{\circ}$ (c 0.72 , chloroform).

All the species of Thymus L. that we studied contained ursolic acid (substance B) and oleanolic acid (substance A).

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