5'-OH groups are free and the 7-OH group is substituted by a methyl, and the only possible position of substitution of an acetyl group is position 8.

On the basis of the facts obtained, we correct the structure of glycoside isolated previously and propose for glycoside (I) the structure of 8-acetylmethylgossypetin 3-O- $\alpha$ -L-rhamnopyranoside, for its aglycone 8-acetyl-7-methylgossypetin, and for glycoside (II) 7-methylgossypetin 3-O- $\alpha$ -L-rhamnopyranoside. They are new natural compounds and we propose to call them pyrifolin, pyrifolidin, and pyrifolinin, respectively. It must be mentioned that the aglycone 7-methylgossypetin which we obtained by the acid hydrolysis of glycoside (II) has a different melting point from the 7-methylgossypetin isolated from the flowers of Lothus corniculatus [3, 4].

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## THE FLAVONOIDS OF Hieracium umbellatum

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We collected the plant <u>Hieracium umbellatum</u> L. (narrowleaf hawkweed) in the stage of full flowering in the environs of Vitebsk.

To isolate individual flavonoids, the epigeal part was extracted with ethanol in the boiling-water bath. The combined extracts were concentrated under vacuum to small volume, the residue was treated with carbon tetrachloride, and chromatography was carried out on a column of polyamide sorbent. Elution of the column with a mixture of ethanol and chloroform gave apigenin, luteolin, and a substance (III) with mp 266-268°C,  $[\alpha]_D$ -54.7° (c 0.6; formamide),  $\lambda_{max}$  255, 268, 350 nm. The product of the acid hydrolysis of (III) was an aglycone with mp 328-330°C giving an acetate with mp 224-226°C. A mixture of the aglycone of substance (III) with luteolin gave no depression of the melting point. The acid mother liquor was found by paper chromatography to contain glucose. On the basis of the results of UV, IR, and NMR spectroscopy it may be considered that the compound isolated was luteolin  $7-\beta$ -D-glucopyranoside.

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