

New Flavonoids from the Farina of *Pityrogramma* Species

Christiane Hitz, Karin Mann,
and Eckhard Wollenweber

Institut für Botanik der TH Darmstadt, Schnittpahn-
straße 3, D-6100 Darmstadt

Z. Naturforsch. **37 c**, 337 – 339 (1982);
received November 16, 1981

Pityrogramma calomelanos, *P. triangularis*, Pteridophyta,
Frond Exudate, Chalcone, Dihydrochalcone, Galangin-5,7-
dimethyl Ether

2',6',4'-OH, 4'-OMe-chalcone (neosakuranetin) and
2',6',4'-OH, 4'-OMe-dihydrochalcone (asebogenin) were
found as major constituents of the frond exudate on one in-
dividual plant of *Pityrogramma calomelanos*. Galangin-5,7-
dimethyl ether was isolated from the farina on fronds of a
certain population of *Pityrogramma triangularis* as a novel
natural flavonol.

In a previous report on the composition of the
flavonoid exudate on ferns of the genus *Pity-
rogramma* the occurrence of a new chalcone was
mentioned [1]. This compound, 2',6',4'-OH, 4'-OMe
chalcone, had been detected only in the farina of two
vouchers of *Pityrogramma tartarea* var. *aurata*
(Moore) Tryon. These are J. A. Ewan 15672 from
Bolivia (at US) and C. Vargas 22355 from Peru (at
CR). Of course, only small fragments of these her-
barium specimens were available and hence the
chalcone could be identified only tentatively, by
direct comparison on TLC. Meanwhile we found the
same flavonoid in the farina of one plant of *Pity-
rogramma calomelanos* (L.) Link cultivated in a
greenhouse [2]. Of this flavonoid a minute amount
could now be isolated and identified unambigu-
ously. From the same plant we could also isolate the
corresponding dihydrochalcone, which had not been
seen previously in any other *Pityrogramma* farina
analysed. The third flavonoid we wish to report here
is a novel natural methyl derivative of galangin,
isolated from the farina of a distinct population of
Pityrogramma triangularis Max.

Material and Methods

Pityrogramma calomelanos is cultivated in a green-
house at the Botanical Garden of Darmstadt Uni-

Reprint requests to Prof. Dr. E. Wollenweber.

0341-0382/82/0300-0337 \$ 01.30/0

versity. *Pityrogramma triangularis* was collected by
E. W. at School Land Gulch at the Pardee Reser-
voir/California, in July, 1980. Vouchers are kept in
the private herbarium of E. W. in Darmstadt (*P.
calomelanos*, 'plant 11'; *P. triangularis*, EW-2). Fern
fronds were rinsed with acetone to dissolve the exu-
date material. This was worked up by column chro-
matography on silica and on polyamide SC-6 in the
usual manner. Preparative TLC and TLC compari-
sons were done on silica or on polyamide DC-11 (for
details cf. [3]). – A sample of neosakuranetin was
prepared by treatment of a solution of sakuranetin
(naringenin-7-methyl ether) with alkali. The re-
action product was, after slight acidification, rapidly
extracted with diethyl ether. – Demethylation of
comp. **III** was done with pyridine-HBr according to
[4].

Results and Discussion

From the few leaves of *Pityrogramma calomelanos*
'plant 11' available a small amount of compound **I**
could be isolated, but not crystallized, so we could
not determine the m.p. The material was sufficient,
however, to run a mass spectrum, which shows the
following fragmentation. MS m/z (rel. int.): 286 (M^+ ,
53), 285 (23), 180 (18), 167 (78), 166 (32), 138 (37),
120 (84). These data indicate a chalcone with 3 OH-
groups and 1 OMe-group 5 as well as a B-ring with 1
OH-group. The chromatographic identity of comp. **I**
with the reaction product of alkali treatment of
naringenin-7-methyl ether (sakuranetin), together
with the mass spectral result shows that this com-
pound is indeed 2',6',4'-OH,4'-OMe-chalcone. Fur-
thermore the UV spectra of both products are
identical (± 2 nm). UV $\lambda_{\max}^{\text{MeOH}}$ 365, 295 nm; + AlCl_3
405, 312 nm; + $\text{AlCl}_3 + \text{HCl}$ 395, 310 nm; no shifts
with NaOAc and H_3BO_3 .

This chalcone, as to our knowledge, has not been
found in nature previously [2]. However, its 6'-
O-glucoside was isolated as early as 1954 from the
ethanolic extract of *Prunus pudum* stem bark [6]. As
this glycoside is known under the name of neosaku-
ranin, we proposed the trivial name neosakuranetin
for the aglycone **I** [1, 2].

Compound **II** was isolated as light brownish crys-
tals, m.p. 144 °C. Its chromatographic properties in-
dicated a dihydrochalcone and this assumption is
supported by the UV spectrum. UV $\lambda_{\max}^{\text{MeOH}}$ 268,



Dieses Werk wurde im Jahr 2013 vom Verlag Zeitschrift für Naturforschung
in Zusammenarbeit mit der Max-Planck-Gesellschaft zur Förderung der
Wissenschaften e.V. digitalisiert und unter folgender Lizenz veröffentlicht:
Creative Commons Namensnennung-Keine Bearbeitung 3.0 Deutschland
Lizenz.

Zum 01.01.2015 ist eine Anpassung der Lizenzbedingungen (Entfall der
Creative Commons Lizenzbedingung „Keine Bearbeitung“) beabsichtigt,
um eine Nachnutzung auch im Rahmen zukünftiger wissenschaftlicher
Nutzungsformen zu ermöglichen.

This work has been digitalized and published in 2013 by Verlag Zeitschrift
für Naturforschung in cooperation with the Max Planck Society for the
Advancement of Science under a Creative Commons Attribution-NoDerivs
3.0 Germany License.

On 01.01.2015 it is planned to change the License Conditions (the removal
of the Creative Commons License condition "no derivative works"). This is
to allow reuse in the area of future scientific usage.

229 nm; + AlCl₃ 370, 312, 226 nm; + NaOH 364, 296, 242, 220 nm. MS *m/z* (rel. int.) 288 (52, M⁺), 182 (13), 167 (83, C₈H₇O₃⁺), 140 (19, C₇H₈O₃⁺), 120 (31, C₈H₈O⁺), 107 (29, C₇H₇O⁺). The data show that this dihydrochalcone has also 3-OH-groups and 1 OMe-group. One OH- group is obviously located at ring B, which points to 2',6',4-OH, 4'-OMe-dihydrochalcone. This structure is supported by the PMR spectrum. ¹H-NMR (90 MHz, DMSO-d₆; ppm/TMS) 12.3 (1 H, s; OH at C-2' or at C-6'; further OH-protons exchanged), 7.02 and 6.68 (2 H each, d; AA'BB' spin system, B-ring p-substituted), 6.0 (2 H, s; H-3'/H-5'), 2.66 (3 H, s; OCH₃), ~3.2–2.6 (4 H, m; α- and β-methylens). Hence to compound II the structure of 2',6',4-OH,4'-OMe- dihydrochalcone is assigned. UV, MS, and PMR data are in agreement with those reported in literature [7]. The m.p. is 18° lower, but this certainly is due to impurity.

This dihydrochalcone has been found in nature only once before as an aglycone. In leaves of three *Rhododendron* species it occurs as such, jointly with its 2'-O-glucoside [7]. As this glycoside is named asebotin, we proposed the trivial name asebogenin for the aglycone (cf. 2).

Further compounds found in the farina of our distinct plant of *Pityrogramma calomelanos*, in trace amounts, are 2',6'-OH, 4'-OMe-dihydrochalcone, 2',6'-OH,4',4'-OMe-dihydrochalcone 2',6'-OH,4',4'-OMe-chalcone, naringenin-7,4'-dimethyl ether and naringenin-7-methyl ether (sakuranetin), and the complex flavonoid "D-1" (compound A in [8]).

Of *Pityrogramma triangularis* (EW-2) 56 g of air-dried fronds were available. The yellow farina was dissolved yielding 3.82 g of exudate material. From this we isolated, by column chromatography and finally by preparative TLC, a small amount of compound III, distinguished as a bright yellow fluorescent spot on polyamide TLC. From boiling ethanol it forms light yellow crystals, m.p. 174°. The bright yellow fluorescence on polyamide layer points to a flavonoid with either an OMe-group or no substituent at C-5. MS *m/z* (rel. int.) 298 (21, M⁺), 297 (100, M-1), 279 (22), 267 (26), 252 (67), 181 (61, "Pic D" according to [9]), 105 (45, "Pic C"), 77 (29, "C-28"). M⁺ 298 indicates a flavone or a flavonol with 1 OH and 2 OMe, and fragments *m/z* 181, 105 and 77 indicate an unsubstituted B-ring. Compound III could hence be a dimethyl ether of galangin, as this flavonol itself as well as its 7-methyl ether (izalpinin) are present in this fern farina, too. According to the

fluorescence of the spot it should be the 5,7-dimethyl derivative. As a matter of fact, demethylation of compound III with pyridin-hydrobromide yields a mixture of galangin with gal-7-Me. This proves unambiguously that compound III is indeed galangin-5,7-dimethyl ether. Compound III exhibits the following UV spectrum. UV λ_{max}^{MeOH} (402), 352 (305), 265 nm; + AlCl₃ 406 (330), 273, 252 nm; + NaOH 395, 271, 250 nm. — This is the first report of this flavonol as a natural product (cf. [10]).

On *Pityrogramma calomelanos* the farina usually is either white, consisting mainly of 2',6'-OH,4'-OMe-dihydrochalcone and/or 2',6'-OH,4',4'-OMe-dihydrochalcone (*P. calomelanos* var. *calomelanos*), or it is yellow, then mainly consisting of 2',6'-OH,4'-OMe-chalcone and/or 2',6'-OH,4',4'-OMe-chalcone (*P. calomelanos* var. *aureoflava* (Hook.) Weath. = *P. austroamericana* Domin). Yellow farina can also consist of a mixture of the cited dihydrochalcone(s) and chalcone(s) [1]. The farina produced by 'plant 11' in our greenhouse is only slightly yellowish and it is produced in a rather low amount only. It is composed mainly of 2',6',4-OH,4'-OMe-dihydrochalcone (asebogenin) and the corresponding 2',6',4-OH,4'-OMe-chalcone (neosakuranetin), while the other chalcones and dihydrochalcones are only minor constituents. In the yellow farina of some other plants of *P. calomelanos* var. *aureoflava* cultivated in our greenhouse neosakuranetin can be detected as a trace constituent. All plants were originally grown from spores obtained from one plant cultivated at the Botanical Garden in Munich. They differ markedly in the shape of pinnae and pinnules as well as in the farina composition. This obviously is due to the well-known phenomenon of very easily occurring hybridization in *Pityrogramma* [11].

P. triangularis (EW-2), in its farina composition, resembles none of the previously described varieties or chemotypes of this species complex (cf. [12]). The novel flavonol, galangin-5,7-dimethyl ether, is present as a minor constituent only, but by its fluorescence it dominates the flavonoid pattern on TLC. A series of further flavonoids could be identified in this farina, which is still under investigation. The results will be reported and discussed elsewhere. The population from which this plant material has been collected represents either a new chemotype or a hybrid.

Acknowledgements

Thanks are due to A. Ritter, H. Groh, and F. Axel for cultivation of *Pityrogramma calomelanos* in the greenhouse and to James Lattie, Oakland/California, for rendering possible to E. W. to collect *Pityrogramma triangularis*. We wish to thank also to M.

Fischer, Darmstadt and to Dr. G. Schilling and Mrs. U. Blaser, both Heidelberg, for running the MS- and PMR-spectra. The investigation was financially supported by the Deutsche Forschungsgemeinschaft with a grant to E. W. which is gratefully acknowledged.

- [1] E. Wollenweber and V. H. Dietz, *Biochem. Syst. Ecol.* **8**, 21 (1980).
- [2] E. Wollenweber and V. H. Dietz, *Phytochemistry* **20**, 869 (1981).
- [3] E. Wollenweber, V. H. Dietz, D. Schillo, and G. Schilling, *Z. Naturforsch.* **35c**, 685 (1980).
- [4] G. Howard and T. J. Mabry, *Phytochemistry* **9**, 2413 (1970).
- [5] E. Wollenweber and V. H. Dietz, *Phytochem. Bull.* **12**, 48 (1979).
- [6] B. Puri and T. R. Seshadri, *J. Sci. Industr. Res.* **13B**, 698 (1954).
- [7] T. J. Mabry, M. Sakakibara, and B. King, *Phytochemistry* **14**, 1448 (1975).
- [8] H. Wagner, O. Seligmann, M. V. Chari, E. Wollenweber, V. H. Dietz, D. M. X. Donnelly, M. J. Meegan, and B. O'Donnell, *Tetrahedron Letters* **1979**, 4269.
- [9] H. Audier, *Bull. Soc. Chim. France* **1966**, 2892.
- [10] E. Wollenweber, in: *The Flavonoids – Advances in Research 1975–1980*. Chapman & Hall, London 1982.
- [11] K. Domin: *Rozpr. České Akad. Trida II*, **38**, 1 (1929).
- [12] D. M. Smith: *Bull. Torrey Bot. Club.* **107**, 134 (1980).