

(40 l.) and the extract concentrated *in vacuo* below 50°. The syrupy material obtained was extracted successively with C_6H_6 (6 l.), EtOAc (6 l.) and *n*-BuOH (4 l.). C_6H_6 and EtOAc fraction gave compounds reported earlier^{1,2} and also tetramethylellagic acid (10%, EtOAc in C_6H_6) and epioleonic acid (20%, EtOAc in C_6H_6). *n*-BuOH extract was freed of the solvent and the residue treated with Me_2CO . Me_2CO soluble fraction gave **1** while Me_2CO insoluble part afforded ellagic acid.

Genkwanin 4'-galactoside (**1**). Me_2CO soluble part of the *n*-BuOH extract, deposited a solid which was recrystallized from EtOH (Found: C, 58.70; H, 5.20. $C_{22}H_{22}O_{10}$ required: C, 59.10; H, 4.91%). Acid hydrolysis and extraction with Et_2O afforded genkwanin, m.p. 267° (3 crystallization gave m.p. 280°) (Found: C, 67.10; H, 4.30. $C_{16}H_{12}O_5$ required: C, 67.60; H, 4.20%).

Diacetate. M.p. 204° (Found: C, 64.10; H, 4.10. $C_{20}H_{16}O_7$ required: C, 64.60; H, 4.30%). The aq. hydrolysis soln was neutralized ($BaCO_3$) and PC indicated that the only sugar present was galactose.

Ellagic acid. The Me_2CO insoluble fraction from the *n*-BuOH extract was extracted (Soxhlet) with hot Me_2CO . Concentration and standing in the cold furnished a yellow solid. This on recrystallization gave yellow needles, m.p. > 360° (Found: C, 56.10; H, 2.10. $C_{14}H_6O_8$ required: C, 56.60; H, 1.98%).

Tetramethyl ellagic acid. Obtained as needles when crystallized from MeOH and dioxane m.p. 340° (Found: C, 59.80; H 4.20. $C_{18}H_{14}O_8$ required: C, 60.33; H, 3.91%).

Epioleonic acid. Crystallized from MeOH, m.p. 302° (Found: C, 78.50; H, 9.86. $C_{30}H_{48}O_3$ required: C, 78.94; H, 9.86%).

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ACORENONE-B IN *ANGELICA LUCIDA* OIL

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Plant. *Angelica lucida* L. (\equiv *Coelopleurum actaeifolium* (Michx.) Coult. & Rose, *Archangelica gmelini* (specimen retained in Herbarium at University of Waterloo, NA 5422). *Source*. Blue Rocks, Lunenburg, Nova Scotia, Canada. *Uses*. Tender parts of plant used as food and tonic¹ by Eskimos.

Present work. The dried seeds were water distilled to yield 0.15% essential oil. The oil, which was analysed by a combination of techniques described previously² was found to contain acorenone-B (43.1%), β -phellandrene (16.1%), myrcene (8.0%), caryophyllene (7.5%), α -pinene (2.0%), phenyl ethyl isovalerate (2.0%), bornyl acetate (1.6%), benzyl isovalerate (1.1%), camphene (0.3%), β -pinene (0.2%) and sabinene (0.2%).

Comment. Acorenone B, which was recently isolated from *Bothriochloa intermedia* (R.Br.) A. Camus (Gramineae) and synthesized,³ was found to be identical to acorenone previously isolated from *Acorus calamus* L. (Araceae).⁴

¹ FRENCH, D. H. (1971) *The Biology & Chemistry of the Umbelliferae* (HEYWOOD, V. H., ed.), pp. 385–412, Publ. for Linn. Soc., Academic Press, London.

² LAWRENCE, B. M. (1971) *Can. Inst. Food Technol. J.* **4**, A44.

³ McCCLURE, R. J. (1969) Ph.D. Thesis, Georgia Inst. Technol.

⁴ VRKOC, J., HEROUT, V. and SORM, F. (1961) *Coll. Czech. Chem. Commun.* **26**, 3183.