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<td>TERAZAWA, Minoru</td>
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PHENOLIC COMPOUNDS IN LIVING TISSUES OF WOODS. VII.*

(+)-Pinoresinol monoglucoside in Fraxinus mandshurica Rupr. var. Japonica Maxim. (Oleaceae)

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1. Introduction

In the previous papers, the isolation of some phenolic compounds such as (±)-syringaresinol, fraxino, and a lignan from the sapwood of yachidamo Fraxinus mandshurica Rupr. var. japonica Maxim. and coniferin and syringin, coumarin glucosides, phenylethanol derivatives (secoiridoid glucosides) from the inner bark of the same wood were reported (Fig. 1). During in this course of the investigations of phenolic compounds in living tissues of woods, two additional phenolic compounds FGa and FG were isolated from the inner bark of yachidamo.

This paper deals with the isolation and structural elucidation of the two phenolic compounds FGa and FG and the seasonal variations of the compound FG in the young shoot of the wood.

2. Results

Two phenolic compounds FGa and FG were isolated from the ethanol extractives of the inner bark of yachidamo (F. mandshurica) by silica gel column chromatography using ethyl acetate saturated with water as a developing solvent. The compound FGa came out in an aglycon fraction and purified by repeated silica gel column chromatography using n-hexane and ethyl acetate (3:1—1:1, v/v, gradient). It was obtained as colorless crystals, melting at 122°C with $[\alpha]_D^0 +91.6^\circ$ (c=1.0, in MeOH). The compound FG
was obtained from a glycoside fraction as colorless crysyls, melting at 107-109°C with $[\alpha]_D +16^\circ$ (c=0.5, in EtOH). The compound FG yielded an aglycon and glucose by hydrolysis with $\beta$-glucosidase. The Rf value and color reaction of the aglycon on thin layer chromatography (TLC) were identical with those of the compound FGa isolated from the aglycon fraction. The results indicated that the compound FG was a glucoside of FGa,
The proton nuclear magnetic resonance (\(^1\text{H-NMR}\)) spectrum of the compound \(\text{FGa}\) indicated that it is a lignan with two guaiacyl nuclei and a tetrahydrofurofuran ring: six protons of two aromatic methoxyl groups at \(\delta 3.80\), two protons of two phenolic hydroxyl groups at \(\delta 7.37\), six aromatic protons of two guaiacyl nuclei between \(\delta 6.98-6.65\); two protons of two \(\text{C}_a\) methine groups at \(\delta 4.70\), two protons of two \(\text{C}_a\) methine groups at \(\delta 3.05\), and four protons of two \(\text{C}_a\) methylene groups at \(\delta 4.1\) and 4.05. The data are identical with those of pinoresinol published.\(^{12}\) It was identified to be \((+)-\text{pinoresinol}\) by the comparison of the chromatographic behaviors of an authentic specimen on TLC. Melting point and specific rotation were identical with those of \((+)-\text{pinoresinol}\) published.\(^{10}\)

The \(^1\text{H-NMR}\) spectrum of \(\text{FG acetate}\) shows three protons of one aromatic acetoxyl group at \(\delta 2.30\), three protons of aromatic methoxyl group at \(\delta 3.84\) and 3.83 and the multiplet between \(\delta 6.90-7.11\) corresponding to the six aromatic protons of the two guaiacyl nuclei. Two protons of two \(\text{C}_a\) methine groups are observed at \(\delta 4.80\) and 4.76 and two protons of two \(\text{C}_a\) methine groups are observed between \(\delta 3.10-3.07\). Four protons of two \(\text{C}_a\) methylene groups are observed at \(\delta 3.94\) and 3.93 (\(\text{C}_a\), double doublets, \(J=4.5\) Hz) and \(\delta 4.25\) and 4.30 (\(\text{C}_a\), multiplets), respectively. In addition to these signals derived from pinoresinol moiety, signals derived from glucosyl residue are observed: twelve protons of four methyl groups of alcoholic acetoxyl groups at \(\delta 2.03\) and 2.01, seven protons of five methine groups and one methylene group between \(\delta 3.74-5.30\). The data indicate that the compound \(\text{FG}\) is a pinoresinol monoglucoside.

The highest mass ion at \(m/z 730\) in the mass spectrum (MS) of \(\text{FG acetate}\) corresponds to the molecular ion (\(M^+\)) of pinoresinol monoglucoside pentaacetate \((\text{C}_{95}\text{H}_{120}\text{O}_{16}, \text{MW}=730)\). The fragment ions at \(m/z 400\) and 331 are corresponding to those of pinoresinol monoacetate and the glucosyl residue, respectively. The fragment ion peak at \(m/z 358\) is corresponding to pinoresinol itself. The ionization voltage 20 eV was used for detection of these high mass fragment ions including the molecular ion. However, the intensities of these high mass ions were very weak. In contrast, field desorption (FD) mass spectrum of \(\text{FG acetate}\) shows the molecular ion clearly at \(m/z 730\) and a fragment ion corresponding to \([\text{M}-42]^+\) at \(m/z 688\). FD-MS spectrum of the compound \(\text{FG}\) shows the molecular ion (\(M^+\)) at \(m/z 520\) as pinoresinol monoglucoside \((\text{C}_{95}\text{H}_{120}\text{O}_{11}, \text{MW}=520)\) and peak at \(m/z 358\) corresponding to \([\text{M}+\text{H}-163]^+\). The peak at \(m/z 163\) is corresponding to glucosyl residue.\(^9\) These mass spectral data indicate that one of the two phenolic hydroxyl groups of \((+)-\text{pinoresinol}\) is blocked by a glucosyl residue.
The $^{13}$C-NMR spectral data ($^1$H decoupled (COM)) of the compound FG well support the conclusion derived from the data of $^1$H-NMR and MS spectra. The peaks at $\delta$ 84.91 and 85.20 are those of $C_\alpha$ and $C_\alpha$, and peaks at $\delta$ 70.97 and 71.06 are corresponding to $C_\beta$ and $C_\beta$, respectively. The peaks at $\delta$ 53.60 and 53.75 are corresponding to $C_\gamma$ and $C_\gamma$. The peaks at $\delta$ 55.77 and 55.67 are two carbons of aromatic methoxyl groups. The peaks of aromatic carbons are observed as follows: 147.57, 148.99 ($C_4$, $C_4'$), 145.90, 145.95 ($C_3$, $C_3'$), 132.26, 135.30 ($C_\delta$, $C_\delta'$), 118.18, 118.68 ($C_6$, $C_6'$), 115.20, 115.33 ($C_5$, $C_5'$) and 110.46, 110.65 ($C_\alpha$, $C_\alpha'$). In addition to these peaks derived from pinoresinol moiety, the peaks corresponding to glucose carbons are observed. The peak of C-1 is observed in the lowest field among the carbons of glucosyl residue at $\delta$ 100.24 and the peak of C-6 in the highest field at $\delta$ 60.75. The other peaks at $\delta$ 76.88, 76.86, 73.26 and 69.75 are corresponding to C-3, C-5,
C-2 and C-4, respectively. The data of the off resonance $^1$H decoupling experiment (OFR) of the compound FG show C-H coupling. The information is useful for the assignment of each peak in the COM spectrum.

The compound FG was methylated with diazomethane (CH$_2$N$_2$) and the methyl ether derivative was hydrolyzed with 3% sulfuric acid. The aglycon obtained was trimethylsilylated and analyzed by gas-liquid chromatography mass spectroscopy (GC/MS). The highest mass ion at $m/z$ 444 is corresponding to pinoresinol monomethyl ether mono-TMS ether (C$_{29}$H$_{50}$O$_6$T$_1$, MW=444, Fig. 2). Fragment ions at $m/z$ 277 and 219 are attributed to the fragment [A] and [B], respectively. And the other main fragment ions are corresponding to those containing 3, 4-dimethoxyphenyl nucleus and those containing 3-methoxyl-4-OTMS-phenyl nucleus: [C]: 177 and 235, [D]: 166 and 224, [E]: 165 and 223, and [F]: 151 and 209.

The glucose as the glucosyl residue of the compound FG, was confirmed by the comparison of the data of gas-liquid chromatography (GLC), and paper partition chromatography (PPC) with those an authentic glucose in the same way as described in the previous papers.$^{25,26}$

The part of the GLC on the periodical analysis of the ethanol extractives of the inner bark and the xylem of young shoots of yachidamo was shown in the previous paper.$^{25}$ Among the many glycosides, (+)-pinoresinol glucoside marked as FG was observed at Rt 87.5 min. The seasonal variations of (+)-pinoresinol monoglucoside in the young shoots of yachidamo are shown in Fig. 3. The amounts of (+)-pinoresinol monoglucoside in the inner bark were between 20-55 mg/g and...
changed little during the growing season. They reduced after August and increased again toward winter after showing a minimum at the end of September. On the other hand, the variation of (+)-pinoresinol monoglu-
coside in the xylem was more drastic than that in the inner bark, even its
amounts in the xylem were 1/10 of those in the inner bark: (+)-pinoresinol
monogluicoside showed a maximum at the end of August and a minimum
at the end of September and increased against toward winter. Judging
from its amount in June, the amount of (+)-pinoresinol monogluicoside seems
to reduce toward coming spring.

3. Discussion

3.1 Distribution of glycosides of pinoresinols

Phillyrin\(^{10,20}\) and symplocosin\(^{13}\) were isolated from Forsythia species and
from kuroki (Sympliocos lucia), respectively as the glucoside of pinoresinol
derivatives. The former is monogluicoside of (+)-epipinoresinol monomethyl
ether and the latter is monogluicoside of (-)-epipinoresinol monomethyl ehter.
These glucosides are not optical antipodes because the glucosyl residues are
attached to the different aromatic rings each other. Monogluicoside of (+)-
epipinoresinol was also isolated from Forsythia species.\(^{20}\) These are all
glucosides of epi-type pinoresinols. Dimethyl ehter of (+)-pinoresinol was

\[\text{Table 1. Glycosides of pinoresinol derivatives in plants}\]

<table>
<thead>
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<th>Pinoresinols</th>
<th>Plants</th>
<th>Organs</th>
<th>Reference</th>
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<tr>
<td>(-)-Epipinoresinol monomethyl ether monogluicoside = Symplocos (I)</td>
<td>Sympliocos</td>
<td>Bark</td>
<td>[13]</td>
</tr>
<tr>
<td>(+)-Epipinoresinol monogluicoside (II)</td>
<td>Forsythia</td>
<td>Bark</td>
<td>[29]</td>
</tr>
<tr>
<td>(-)-Pinoresinol monogluicoside (IV)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>dimethyl ether = Eudesmin (V)*</td>
<td>Eucalyptus</td>
<td>Bark</td>
<td>[5]</td>
</tr>
<tr>
<td>(+)-Pinoresinol monogluicoside (VI)</td>
<td>Fraxinus</td>
<td>Bark</td>
<td>[20, 21]</td>
</tr>
<tr>
<td>dimethyl ether (VII)</td>
<td>Liriodendron</td>
<td>Bark</td>
<td>[6]</td>
</tr>
<tr>
<td>digluicoside (VIII)</td>
<td>Forsythia</td>
<td>Bark</td>
<td>[2]</td>
</tr>
<tr>
<td>(±)-Pinoresinol digluicoside (IX)</td>
<td>Ligustrum</td>
<td>Bark</td>
<td>[9]</td>
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<tr>
<td></td>
<td>Magnolia</td>
<td>Bark</td>
<td>[11]</td>
</tr>
<tr>
<td></td>
<td>Liriodendron</td>
<td>Bark</td>
<td>[6]</td>
</tr>
<tr>
<td></td>
<td>Inomenium</td>
<td>Bark</td>
<td>[15]</td>
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* (+)-Diaedesmin was isolated from Piper as a diastereomer of V and VII [1].
(+)-PINORESINOL MONOGLUCOSIDE IN FRAXINUS MANDSHURICA

Fig. 4. Pinoresinols in plants.

Legends: I.: R = Me, R' = Glu: (-)-epipinoresinol monomethyl ether monoglucoside (symplocosin); II.: R = H, R' = Glu: (+)-epipinoresinol monoglucoside; III.: R = Glu, R' = Me: (+)-epipinoresinol monomethyl ether monoglucoside (phillyrin); IV.: R = H, R' = Glu: (-)-pinoresinol monoglucoside; V.: R = R' = Me: (-)-pinoresinol dimethyl ether (eudesmin); VI: R = H, R' = Glu: (+)-pinoresinol monoglucoside; VII: R = R' = Me: (+)-pinoresinol dimethyl ether; VIII: R = R' = Glu: (+)-pinoresinol diglucoside. IX.: R = R' = Glu: (±)-pinoresinol diglucoside.

found as one of lignans in Magnolia (1972). Glycosides of (+)-pinoresinol had not been isolated until Terazawa et al. isolated (+)-pinoresinol monoglucoside from the inner bark of yachidamo (Fraxinus mandshurica Rupr. var. japonica Maxim.) and named as pinoresinoside tentatively (in 1972 and 1973). Fujimoto et al.6 showed later the evidence of the existence of the mono- and diglucosides of (+)-pinoresinol accompanied by the mono- and diglucosides of (+)-syringaresinol and (+)-medioresinol in the bark of Liliodendron tulipefera, although they could not isolate them (1975-1977). Kudo et al.9 isolated a lignan glucoside, which was identified to be (+)-pinoresinol monoglucoside, in addition to many lignan glucosides from the bark of Ligustrum japonicum (1976-1980). Chiba et al. also isolated it as one of lignan glucosides from the fruit buds of Forsythia suspensa (1978) and reported its 13C-NMR data (1980). However, no glycosides of (-)-pinoresinol have not been reported so far, although dimethyl ether of (-)-pinoresinol (eudesmin), as a derivative of (-)-pinoresinol, was reported (Table 1). Sih et al. isolated diglucoside of (±)-pinoresinol from Eucommia ulmodes (1976) (Fig. 4).
3.2 Biogenesis of lignans

Terazawa and Sasaya isolated (±)-syringaresinol from the ethanol extractives of the sapwood of yachidamo (1972). The occurrence of the mixture of (+)- and (−)-syringaresinol seemed rather common. However, no reports on the isolation of (±)-pinoresinol had been published at that time. Based on these facts, they discussed the possibility that (±)-syringaresinol in the xylem might be one of the dimers remained not to be incorporated into lignin macromolecules during lignification in the xylem.

They pointed out the considerable reason why (±)-pinoresinol was difficult to find: coniferyl alcohol and pinoresinol have active sites at 5 and 5' of the guaiacyl nucleus(ei) to form C–C bonding when they are dehydrogenated by peroxidase in the process of lignification. This nature results in the formation of dehydrogenative polymerization products of higher molecular weight than those of the dimers. Thus there are less chance to isolate (±)-pinoresinol itself in contrast to the frequent encounter to (±)-syringaresinol which dose not have active site at 5, or 5'. They mentioned the result that (±)-syringaresinol was the predominant dimerization products derived from sinapyl alcohol by the catalytic reaction of laccase or peroxidase in vitro.

In this respect, it is interesting that SiH et al. isolated a diglucoside of (±)-pinoresinol from Tu-Chung (Eucommia ulmoides) (1976). Namely, the existence of (±)-pinoresinol in the plant suggests the possibility of the existence of the specific enzyme systems to produce them predominantly instead of dehydrogenative polymerization products of coniferyl alcohol produced by catalytic reaction of peroxidase which is involving in lignification process. In the other words, the enzyme systems must have the specific ability to locate the radical at C₅ of coniferyl alcohol to lead C₅–C₅ coupling as an initial reaction to form pinoresinols. The finding of the potential existence of the specific enzyme systems is very important because the C₅–C₅ coupling is the key point as the initial reaction of the formation of ordinary lignans, even though the enzymes in Eucommia lack ability to control the reactions after C₅–C₅ coupling to make optically active pinoresinol.

SiH et al. used a microorganism Caldariomlces fumago, which produce chloroperoxidases, for preparing (±)-pinoresinol and (±)-cis-dehydrodiconiferyl alcohol from coniferyl alcohol. The enzyme systems in the microorganism can turn coniferyl alcohol into (±)-pinoresinol and (±)-cis-dehydrodiconiferyl alcohol in approximately 1:1 ratio, respectively. These enzyme systems also seem to have the specific ability to locate predominantly the radical at C₅ of coniferyl alcohol together with the ability to locate the
radical at 5 position of aromatic ring.

The findings of naturally occurring (±)-pinoresinol as its diglucoside in *Eucommia* and of the enzyme systems to form (±)-pinoresinol in *Caldariomlces* suggest that (±)-syringaresinol in hardwoods7,8,14,19 might be formed by the similar enzyme systems in *Eucommia* rather than be formed by per-oxidase which is involving in lignin formation.

Stockigt et al. studied the biosynthesis of optically active lignans16 and found that glucoferulic acid, glucoferulic aldehyde and coniferin were good precursors for phillyrin ((+)-epipinoresinol monomethyl ether monoglucoside) but 3, 4-dimethoxyxynamic acid could not be the intermediate. They postulated a biosynthesis way that the first dimerization occurs at C9 carbon atoms of the two coniferyl alcohol units, which are formed by hydrolysis of the glucosidic intermediate coniferin with β-glucosidase. All the reactions including C9-C9 coupling and ring formation to form (+)-epipinoresinol as the intermediate of phillyrin should be performed by enzyme systems stereospecifically but not through random free radical reaction as seen in polymerization of 4-hydroxycinnamyl alcohols during the formation of optically inactive lignins. Methylation is performed in the final step.

We are now discussing on the enzyme systems which can convert coniferyl alcohol to specific lignan structures. In the next step, we need to find the enzyme systems which can produce optically active lignans in vitro. If we could find a pair of enzyme systems which can produce (+)- and (−)-type of a lignan independently, for example (+)-pinoresinol and (−)-pinoresinol, the problem on the natural occurrence of the racemates would be answered somehow.

3.3 Seasonal variation of (+)-pinoresinol monoglucoside (FG) in yachidamo

The seasonal variations of secoiridoid glucosides such as ligustoroside and oleuropein in yachidamo were reported in the previous paper20. The increasement of the glucosides toward winter was demonstrated and their decreasement toward spring after showing a maximum in winter was estimated judging from their amounts in June. The patterns of seasonal variations of (+)-pinoresinol monoglucoside (FG) in the inner bark and the xylem demonstrated in this work were similar to those of secoiridoid glucosides reported.20 On the contrary, the patterns were different from those of hirsutoside in hannoki (*Alnus hirsuta* Turcz)23, and platyphylloside in shira-kamba (*Betula platyphylla* Sukachev var. *japonica* Hara24, which decreased toward winter after showing maxima during the growing season.

The physiological significance of these difference in the seasonal variations
of the phenolic glycosides in the living tissues of woods remains uncertain.

4. Experimental

4.1 Isolation of (+)-pinoresinol (FGa) and (+)-pinoresinol monoglucoside (FG)

Two hundred microgram of the compound FGa ((+)-pinoresinol) and 2.5 g of the compound FG ((+)-pinoresinol monoglucoside) were isolated from 50 g of the ethanol extractives of the inner bark of yachidamo by silica gel column chromatography using ethyl acetate saturated with water in the same way as described in the previous papers.17-20

4.2 Physico-chemical properties of (+)-pinoresinol (FGa) and (+)-pinoresinol monoglucoside (FG)

The conditions for UV, IR, MS and NMR spectroscopies are same as described in the previous papers.24,26,28 The part of NMR (1H-NMR, 13C-NMR) and FD-MS spectroscopies were conducted by a JNM FX-200 FT NMR Spectrometer and by a JOEL JMS-01SG-2 Mass Spectrometer, respectively. \( [\alpha]_b \) was measured by a JASCO DIP-360.

4.2.1 (+)-Pinoresinol (FGa): Mp 122°C. \( [\alpha]_b +96.8^\circ \) (c = 1.0, in MeOH). TLC (SG-III): Rf 0.6, pinkish orange with diazotized sulfanilic acid (DSA). Anal. Calcd. for C₂₀H₂₂O₆: C, 67.02; H, 6.19. Found: C, 67.11; H, 6.06. 1H-NMR (10% in d₆-acetone) \( \delta \) (ppm): 7.37 (2 H, broad s, 2 Ar-OH), 6.98-6.65 (6 H m, two aromatic nuclei), 4.70 (2 H, d, J=4.5 Hz, two C₆ methine protons), 4.07-4.12 (2 H, m, C₇ methylene), 4.05 (2 H, m, C₇ methylene), 3.80 (6 H, s, two Ar-OMe), and 3.05 (2 H, m, two C methine protons).

4.2.2 (+)-Pinoresinol di-TMS ether: (+)-Pinoresinol (FGa) was treated with TMS reagent and the reactant was subjected to GC-MS. MS (20 eV) m/z: 502 (M⁺), 487 (M-15), 471 (M-31), 430, 306, 294, 277, 235, 223, 209, and 131.

4.2.3 (+)-Pinoresinol dimethyl ether: (+)-Pinoresinol (FGa) was treated with CH₃N₂ and the reactant was subjected to GC-MS. MS (20 eV) m/z: 386 (M⁺), 371 (M-15), 355 (M-31), 379, 378, 365, 353, 341, 331, 320, 308, 296, 284, 272, and 151.

4.2.4 (+)-Pinoresinol monomethyl ether mono-TMS ether: (+)-Pinoresinol monoglucoside (FG) was dissolved in small amount of ethanol and added the excess amount of CH₃N₂ ether solution and the reaction mixture was evaporated to dryness. The methyl ether was hydrolyzed by a 3% H₂SO₄ solution with refluxing for 30 min., and the cooled hydrolyzate was extracted with ether. The ether soluble part was concentrated to dryness and treated with TMS reagent, then the reactant was subjected to GC-MS. MS (20 eV) m/z: 444 (M⁺), 429 (M-15), 423 (M-31), 430, 306, 294, 277[A],
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4.2.5 (+)-Pinoresinol monoglucoside (FG): Mp 106-107°C. [α]_D^20 + 16° (c = 0.5, in EtOH). UV λmax nm (log e): 282 (3.89). TLC (AEAW): Rf 0.4, pinkish orange with DSA. Anal. Calcd. for C_{36}H_{46}O_{11}: C, 59.99; H, 6.20. Found C, 59.90; H, 6.22. FD-MS (JOEL JMS-10SG-2, EC 15-16 MA) m/z (relative intensities %): 522 (10.2, M+2), 521 (32.1, M+1), 520 (100.0, M+), 358 (7.87, M+H-163(glu»), 248 (7.7), 179 (7.3). "C-NMR (2% d_6-DMSO, JNM FX-200 FT NMR) δ (ppm): COM: 53.60, 53.75 (C_r, C_r'), 84.91 (C_a), 85.20 (C_a'), 70.97, 71.06 (C_p, C_p'), 55.77, 55.67 (Ar-OCH_aX2), 132.26 (C_1), 135.30 (C_r), 110.46, 110.65 (C_2, C_2'), 145.90, 145.95 (C_3, C_3'), 147.57, 148.99 (C_4, C_4'), 115.20, 115.33 (C_5, C_5'), 118.18, 118.68 (C_6, C_6'), 55.67, 55.77 (Ar-OCH_a×2), 100.24 (G-C_1), 73.26 (G-C_2), 76.88 (G-C_3), 69.75 (G-C_4), 76.86 (G-C_5), 60.75 (G-C_6).

4.2.6 (+)-Pinoresinol monoglucoside pentaacetate: (+)-pinoresinol monoglucoside (FG) was treated with pyridine and acetic anhydride at 55°C for 8 hours and colorless crystals, melting at 109-110°C, were obtained from aqueous ethanol. Anal. Calcd. for C_{86}H_{42}O_{16}: C, 59.14; H, 5.75. Found: C, 58.56, H, 5.75. IH-NMR (2% in CDCl_3, JNM FX-200 FT NMR) δ (ppm): 3.843, 3.833 (6 H, two s., Ar-OCH_a), 2.308 (3 H, s., Ar-OCOCH_a), 6.897-7.108 (6 H, m., Ar-H, two quaiacyl nuclei), 4.801, 4.756 (2 H, two d., J=5 Hz, C_r, C_r' methines), 3.104-3.065 (2 H, m., C_p, C_p' methines), 3.942, 3.930 (2 H, two d., J=4.5 Hz, C_r methylene), 4.247-4.312 (2 H, m., C_r' methylene), 2.034, 2.012 (12 H, two s., four alc-OCOCH_a), 3.739-4.312 (3 H, m., glu resid.), 4.928-5.295 (4 H, m., glu resid.). MS (20 eV) m/z: 730 (M^+, 688 (M-42), 672, 658, 642, 629, 611, 689, 569, 537, 521, 509, 495, 491, 476, 443, 425, 400, 369, 358, (pinoresinol) 341, 331 (glu resid.), 289, 262, 261, 229, 221, 205, 197, 189, 187, 175, 169, 161, 150, 141. FD-MS (JOEL JMS-01SG-2) m/z (relative intensities %): 732 (16.60, M+2), 731 (45.52, M+1), 730 (100.0, M^+), 688 (11.0, M-42) 399 (7.2), 331 (10.2, glu resid.).

4.2.7 Glucose as glucosyl residue of the compound FG: The hydroylizates obtained by hydrolysis of the compound FG with β-glucosidase and by hydrolysis of FG monomethyl ether with 3% sulfuric acid solution were neutralized with BaCO_3 and filtrated through Toyo Roshi No. 5C and then evaporated the filtrates to dryness. The parts of them were treated with TMS reagent and subjected to GLC (OV-1). The parts of them were also subjected to PPC (BAW). The Rt and Rf obtained were the same as those of the authentic glucose.²⁰

4.2.8 Conditions of chromatographies: GLC, TLC and PPC were
the same as described in the previous papers\textsuperscript{22,24,26}. GLC conditions for the periodical analysis of FG were the same as described in the previous paper.\textsuperscript{25}

\textbf{Acknowledgement}

The author express his appreciation to Professor A. SAKAKIBARA of Hokkaido University for the gift of the authentic (+)-pinoresinol and his helpful advice during this work, to Professor M. MIYAKE of Obihiro University of Agriculture and Veterinary Medicine for his interest and advice during this work and to Mr. M. SASAKI for his experimental assistance. The author express his gratitude to Professor T. KAYAMA of Hokkaido University for his critical reading of this manuscript.

\textbf{Summary}

The compounds FG\textsubscript{a} and FG, which were isolated from the ethanol extractives of the inner bark of yachidamo \textit{Fraxinus mandshurica} RUPR. var. \textit{japonica} MAXIM., were identified to be (+)-pinoresinol and (+)-pinoresinol monoglucoside, respectively.

Biogenesis of lignins with and without optical activity was discussed. The seasonal variation of the compound FG in the xylem and the inner bark of the same wood showed two maxima during a year: the growing season and winter. The patterns were similar to those of secoiridoid glucosides in the same wood but different from those of diaryleptanoid glucosides in the woods of Betulaceae.

\textbf{Literature Cited}

(+)-PINORESINOL MONOGLUCOSIDE IN FRAXINUS MANDSHURICA


