



Microbial Transformation of Two Prenylated Naringenins

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Abstract – Microbial transformation of (\pm)-6-(1,1-dimethylallyl)naringenin (6-DMAN, **1**) and (\pm)-5-(*O*-prenyl)naringenin-4',7-diacetate (5-*O*-PN, **2**) was performed by using fungi. Scale-up fermentation studies with *Mucor hiemalis*, *Cunninghamella elegans* var. *elegans*, and *Penicillium chrysogenum* led to the isolation of five microbial metabolites. Chemical structures of the metabolites were determined by spectral analyses as (\pm)-8-prenylnaringenin (**3**), (2*S*)-5,4'-dihydroxy-7,8-[(*R*)-2-(1-hydroxy-1-methylethyl)-2,3-dihydrofuran]flavanone (**4**), (\pm)-5-(*O*-prenyl)naringenin-4'-acetate (**5**), (\pm)-naringenin-4'-acetate (**6**), and (\pm)-naringenin (**7**), of which **5** was identified as a new compound.

Keywords – Microbial transformation, 6-(1,1-dimethylallyl)naringenin, 5-(*O*-prenyl)naringenin-4',7-diacetate, *Mucor hiemalis*, *Cunninghamella elegans* var. *elegans*, *Penicillium chrysogenum*

Introduction

Naringenin, a colorless flavanone present in a variety of herbs and fruits including grapefruit and tomatoes, has been found to be highly beneficial to human health due to its strong antioxidant potential.¹ It was reported that naringenin has anti-inflammatory, antiproliferative, anticancer, chemopreventive and estrogenic properties,²⁻⁴ and has beneficial influence on lipid metabolism and insulin sensitivity.⁵ The pharmacological application of naringenin-type flavonoids has become more attractive due to the possibility of the chemical synthesis of its prenylated derivatives.⁶

(\pm)-6-(1,1-Dimethylallyl)naringenin (6-DMAN, **1**), a natural naringenin-type flavonoid isolated originally from the leaves of the African tree *Monotes engleri* (family Dipterocarpaceae),⁷ has been found to exhibit estrogenic and antiandrogenic properties,^{3,8} and also found to display cytotoxic activity against several human cancer cell lines.⁷ (\pm)-5-(*O*-Prenyl)naringenin-4',7-diacetate (5-*O*-PN, **2**), a synthetic derivative of 6-DMAN, showed high cytotoxicity in HL-60 and MCF-7 cell lines.⁹

No metabolism study has yet been carried out to identify the metabolic fate of 6-DMAN and 5-*O*-PN. Hence, metabolic processes of **1** and **2** were investigated

by using microorganisms in the present study. Bio-transformation studies using microorganisms are well known as an important tool for the conversion of natural compounds. It has been used successfully as in vitro models to mimic and predict the metabolic fate of pharmaceutical agents in mammalian systems.¹⁰⁻¹² Scale-up studies with three microbial cultures, *Mucor hiemalis*, *Cunninghamella elegans* var. *elegans*, and *Penicillium chrysogenum*, have resulted in the production of five microbial metabolites (Fig. 1).

Experimental

General experimental procedures – The ¹H and ¹³C NMR spectra were obtained in CDCl₃ and DMSO-*d*₆ on a Varian Unity Inova 300 spectrometer at 300 and 75 MHz, respectively. The chemical shift values (δ) are reported in ppm units, and the coupling constants (*J*) are in Hz. UV spectra were recorded on a JASCO V-530 spectrophotometer, and IR spectra were obtained on a JASCO FT/IR-300E spectrometer in KBr discs. Optical rotations were measured with a Perkin Elmer 343 Plus polarimeter. ESIMS and HRESIMS analyses were performed on a Micromass QTQF2 mass spectrometer. TLC analyses were carried out on precoated silica gel 60 F₂₅₄ plates (Merck, Darmstadt, Germany). The developing system used was chloroform:methanol (9:1, v/v) solution, and visualization of the TLC plates was performed using anisaldehyde-H₂SO₄ spray reagent. For column chromatography

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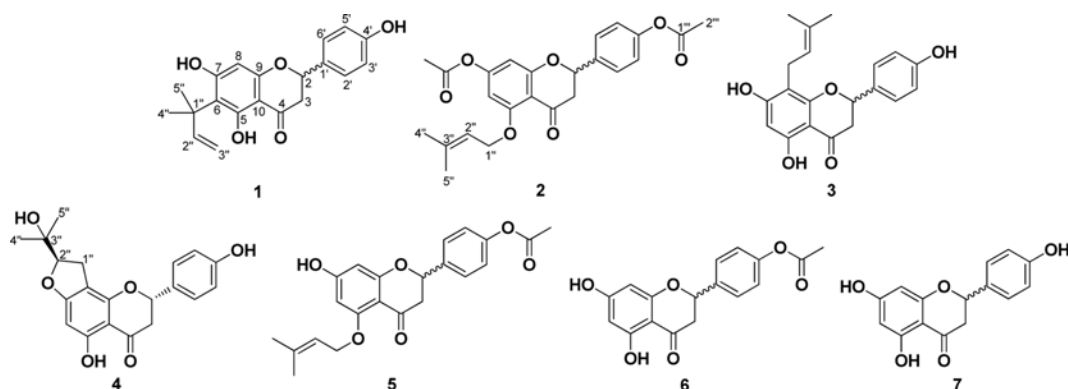


Fig. 1. Chemical structures of the compounds 1 - 7.

graphy, the adsorbent used was silica gel 60 (70 - 230 mesh, Merck). HPLC was performed on a Waters 600E Multisolvant Delivery System (Waters Corp., Milford, MA, USA) connected to a Waters 486 detector using Phenomenex C18 column (10 × 250 mm, 5 μm) with MeOH:H₂O at a flow rate of 2.0 mL/min.

Materials and microorganisms – 6-DMAN (**1**) and 5-*O*-PN (**2**) were semi-synthesized from *rac*-naringenin.¹³ All of the ingredients for microbial media, including dextrose, peptone, malt extract, yeast extract, and potato dextrose broth were purchased from Becton, Dickinson and Company (Sparks, MD, USA).

All the microorganisms were obtained from the Korean Collection for Type Cultures (KCTC). Twenty-four cultures were used for the preliminary screening process and are listed below: *Absidia coerulea* 6936, *Alternaria alternata* 6005, *Aspergillus fumigatus* 6145, *Candida famata* 7000, *Cunninghamella elegans* var. *elegans* 6992, *Debaryomyces hansenii* var. *hansenii* 7645, *Debaryomyces occidentalis* var. *occidentalis* 7194, *Filobasidium neoformans* 7902, *Fusarium merismoides* 6153, *Gliocladium deliquescens* 6173, *Glomerella cingulata* 6075, *Hormoconis resinae* 6966, *Kluyveromyces marxianus* 7155, *Metschnikowia pulcherrima* 7605, *Monascus rubber* 6122, *Mortierella ramanniana* var. *angulispora* 6137, *Mucor hiemalis* 26779, *Penicillium chrysogenum* 6933, *Pichia pastoris* 7190, *Rhizopus oryzae* 6946, *Saccharomycodes ludwigii* 7126, *Torulasporea delbrueckii* 7116, *Trichoderma koningii* 6042, *Trigonopsis variabilis* 7263.

Three types of media were used in the fermentation experiments and are listed below: *A. coerulea* and *M. hiemalis* were cultured on malt medium (malt extract 20 g/L, dextrose 20 g/L, peptone 1 g/L); *C. elegans* var. *elegans* was cultured on potato dextrose medium (24 g/L); other microorganisms were cultured on yeast-malt medium (dextrose 10 g/L, peptone 5 g/L, malt extract 3 g/L,

and yeast extract 3 g/L).

Screening procedures – Microbial metabolism studies were carried out according to the standard two-stage procedure.¹⁰ Briefly, the actively growing microbial cultures were inoculated in 100 mL Erlenmeyer flasks containing 25 mL of a suitable medium, and incubated with gentle agitation (200 rpm) at 25 °C in a temperature-controlled shaking incubator. The DMSO solutions (10 mg/mL, 100 μL) of **1** and **2** were added to each flask 24 h after inoculation, and further incubated at the same conditions for another 8 days. Sampling and TLC monitoring were performed at an interval of 24 h. Culture controls consisted of fermentation cultures in which the microorganisms were grown without addition of substrates.

Scale-up fermentations of 6-DMAN and 5-*O*-PN – Scale-up fermentations were carried out under the same temperature-controlled shaking conditions with three or eight 500 mL Erlenmeyer flasks each containing 125 mL of a suitable medium, and 45 mg of **1** and 100 mg of **2** dissolved in DMSO were distributed evenly among flasks, respectively. After incubation for 7 days, the microbial culture broth was extracted with EtOAc (400 mL × 3), and the organic layers were combined and concentrated *in vacuo*. The EtOAc extract (80 mg) of **1** from *M. hiemalis* culture broth was chromatographed by semi-preparative reversed-phase HPLC with 80% MeOH as mobile phase to give metabolite **3** (3.0 mg). The EtOAc extract (70 mg) of **1** from *C. elegans* var. *elegans* culture broth was chromatographed by semi-preparative reversed-phase HPLC with 80% MeOH to give metabolite **4** (4.4 mg). The EtOAc extract (200 mg) of **2** from *P. chrysogenum* culture broth was chromatographed by semi-preparative reversed-phase HPLC with 65% MeOH to afford metabolites **5** (4.5 mg), **6** (1.5 mg), and **7** (3.0 mg).

(±)-6-(1,1-Dimethylallyl)naringenin (**1**) – A white solid; ¹H-NMR (CDCl₃, 300 MHz) δ 13.11 (1H, s, 5-OH), 7.48

(1H, s, 7-OH), 7.31 (2H, d, $J = 8.9$ Hz, H-2'/6'), 6.86 (2H, d, $J = 8.9$ Hz, H-3'/5'), 6.44 (1H, dd, $J = 10.6, 17.9$ Hz, H-2''), 5.93 (1H, s, H-8), 5.43 (1H, dd, $J = 0.8, 17.9$ Hz, H-3'a), 5.36 (1H, dd, $J = 0.8, 10.6$ Hz, H-3'b), 5.31 (1H, dd, $J = 3.1, 12.7$ Hz, H-2), 5.10 (1H, s, 4'-OH), 3.07 (1H, dd, $J = 12.2, 17.0$ Hz, H-3a), 2.78 (1H, dd, $J = 3.1, 17.0$ Hz, H-3b), 1.60 (3H, s, H-4''), 1.57 (3H, s, H-5''); ^{13}C NMR (CDCl₃, 75 MHz) δ 196.2 (C-4), 164.5 (C-7), 163.9 (C-5), 160.4 (C-9), 156.0 (C-4'), 149.6 (C-2''), 130.7 (C-1'), 127.4 (C-2'/6'), 115.6 (C-3'/5'), 113.5 (C-3''), 111.5 (C-6), 103.0 (C-10), 96.7 (C-8), 78.4 (C-2), 43.3 (C-3), 40.7 (C-1''), 27.2 (C-4''), 26.6 (C-5'').

(±)-5-(*O*-Prenyl)naringenin-4',7-diacetate (**2**) – A white solid; $^1\text{H-NMR}$ (CDCl₃, 300 MHz) δ 7.46 (2H, d, $J = 8.4$ Hz, H-2'/6'), 7.14 (2H, d, $J = 8.4$ Hz, H-3'/5'), 6.42 (1H, d, $J = 2.0$ Hz, H-6), 6.32 (1H, d, $J = 2.0$ Hz, H-8), 5.53 (1H, t, $J = 6.5$ Hz, H-2''), 5.43 (1H, dd, $J = 2.7, 12.5$ Hz, H-2), 4.62 (2H, d, $J = 6.5$ Hz, H-1''), 3.01 (1H, dd, $J = 12.5, 16.9$ Hz, H-3a), 2.82 (1H, dd, $J = 2.7, 16.9$ Hz, H-3b), 2.31 (3H, s, 7-OAc), 2.30 (3H, s, 4'-OAc), 1.79 (3H, s, H-4''), 1.74 (3H, s, H-5'').

(±)-8-Prenylnaringenin (**3**) – A white solid; $[\alpha]_{\text{D}}^{20} : 0^\circ$ (c 0.028, MeOH); $^1\text{H-NMR}$ (DMSO- d_6 , 300 MHz) δ 12.18 (1H, s, 5-OH), 10.78 (1H, s, 7-OH), 9.69 (1H, s, 4'-OH), 7.30 (2H, d, $J = 8.5$ Hz, H-2'/6'), 6.76 (2H, d, $J = 8.5$ Hz, H-3'/5'), 5.96 (1H, s, H-6), 5.42 (1H, dd, $J = 2.8, 12.5$ Hz, H-2), 5.08 (1H, t, $J = 6.2$ Hz, H-2''), 3.20 (1H, dd, $J = 12.5, 16.3$ Hz, H-3a), 3.08 (2H, d, $J = 6.2$ Hz, H-1''), 2.71 (1H, dd, $J = 2.8, 16.3$ Hz, H-3b), 1.58 (3H, s, H-4''), 1.53 (3H, s, H-5''); ^{13}C NMR (DMSO- d_6 , 75 MHz) δ 196.7 (C-4), 164.3 (C-9), 161.1 (C-5), 159.7 (C-7), 157.5 (C-4'), 130.1 (C-3''), 129.2 (C-1'), 128.0 (C-2'/6'), 122.6 (C-2''), 115.1 (C-3'/5'), 106.9 (C-8), 101.7 (C-10), 95.2 (C-6), 78.2 (C-2), 41.9 (C-3), 25.5 (C-5''), 21.2 (C-1''), 17.6 (C-4'').

(2*S*)-5,4'-Dihydroxy-7,8-[(*R*)-2-(1-hydroxy-1-methylethyl)-2,3-dihydrofuran]flavanone (**4**) – A pale yellow amorphous powder; $[\alpha]_{\text{D}}^{20} : -120^\circ$ (c 0.056, MeOH); $^1\text{H-NMR}$ (DMSO- d_6 , 300 MHz) δ 12.43 (1H, s, 5-OH), 9.69 (1H, s, 4'-OH), 7.32 (2H, d, $J = 8.5$ Hz, H-2'/6'), 6.80 (2H, d, $J = 8.5$ Hz, H-3'/5'), 5.96 (1H, s, H-6), 5.49 (1H, dd, $J = 3.0, 12.2$ Hz, H-2), 4.74 (1H, t, $J = 8.5$ Hz, H-2''), 3.26 (1H, dd, $J = 12.2, 17.3$ Hz, H-3a), 2.90 (2H, dd, $J = 3.5, 8.5$ Hz, H-1''), 2.72 (1H, dd, $J = 3.0, 17.3$ Hz, H-3b), 1.11 (3H, s, H-4''), 1.08 (3H, s, H-5''); ^{13}C NMR (DMSO- d_6 , 75 MHz) δ 196.6 (C-4), 168.9 (C-7), 164.5 (C-5), 158.2 (C-4'), 157.4 (C-9), 129.2 (C-1'), 128.7 (C-2'/6'), 115.6 (C-3'/5'), 105.2 (C-8), 102.7 (C-10), 92.0 (C-2''), 90.9 (C-6), 78.9 (C-2), 70.4 (C-3''), 42.4 (C-3), 26.5 (C-1''), 26.1 (C-5''), 25.5 (C-4'').

(±)-5-(*O*-Prenyl)naringenin-4'-acetate (**5**) – A pale yellow amorphous powder; $[\alpha]_{\text{D}}^{20} : 0^\circ$ (c 0.052, MeOH); UV (MeOH) λ_{max} : 286, 343 nm; IR (KBr) ν_{max} cm⁻¹: 3345, 1688, 1620, 1596, 1443, 1372, 1208; $^1\text{H-NMR}$ (DMSO- d_6 , 300 MHz) δ 7.53 (2H, d, $J = 8.5$ Hz, H-2'/6'), 7.16 (2H, d, $J = 8.5$ Hz, H-3'/5'), 6.04 (1H, d, $J = 1.6$ Hz, H-6), 5.95 (1H, d, $J = 1.6$ Hz, H-8), 5.48 (1H, dd, $J = 2.8, 12.5$ Hz, H-2), 5.39 (1H, m, H-2''), 4.50 (2H, d, $J = 6.2$ Hz, H-1''), 2.97 (1H, dd, $J = 12.5, 16.3$ Hz, H-3a), 2.60 (1H, dd, $J = 2.8, 16.3$ Hz, H-3b), 2.27 (3H, s, -OAc), 1.74 (3H, s, H-4''), 1.69 (3H, s, H-5''); ^{13}C NMR (DMSO- d_6 , 75 MHz) δ 187.0 (C-4), 169.2 (C-1''), 165.6 (C-7), 164.0 (C-9), 161.3 (C-5), 150.3 (C-4'), 136.8 (C-3''), 136.4 (C-1'), 127.7 (C-2'/6'), 121.9 (C-3'/5'), 119.9 (C-2''), 104.2 (C-10), 95.8 (C-6), 94.9 (C-8), 77.4 (C-2), 65.1 (C-1''), 44.9 (C-3), 25.5 (C-4''), 20.9 (C-2''), 18.1 (C-5''); HR-ESI-MS m/z 383.1494 $[\text{M}+\text{H}]^+$ (calcd for C₂₂H₂₃O₆, 383.1495).

(±)-Naringenin-4'-acetate (**6**) – A pale yellow amorphous powder; $[\alpha]_{\text{D}}^{20} : 0^\circ$ (c 0.024, MeOH); $^1\text{H-NMR}$ (DMSO- d_6 , 300 MHz) δ 12.16 (1H, s, 5-OH), 7.55 (2H, d, $J = 8.6$ Hz, H-2'/6'), 7.19 (2H, d, $J = 8.6$ Hz, H-3'/5'), 5.87 (1H, d, $J = 1.2$ Hz, H-6), 5.84 (2H, d, $J = 1.2$ Hz, H-8), 5.56 (1H, dd, $J = 2.7, 13.0$ Hz, H-2), 3.26 (1H, dd, $J = 13.0, 17.2$ Hz, H-3a), 2.74 (1H, dd, $J = 2.7, 17.2$ Hz, H-3b), 2.29 (3H, s, -OAc).

(±)-Naringenin (**7**) – A white solid; $[\alpha]_{\text{D}}^{20} : 0^\circ$ (c 0.021, MeOH); $^1\text{H-NMR}$ (DMSO- d_6 , 300 MHz) δ 12.17 (1H, s, 5-OH), 7.30 (2H, d, $J = 8.5$ Hz, H-2'/6'), 6.80 (2H, d, $J = 8.5$ Hz, H-3'/5'), 5.86 (2H, s, H-6/8), 5.41 (1H, dd, $J = 2.8, 13.0$ Hz, H-2), 3.25 (1H, dd, $J = 13.0, 17.0$ Hz, H-3a), 2.68 (1H, dd, $J = 2.8, 17.0$ Hz, H-3b).

Results and Discussion

6-DMAN (C₂₀H₂₀O₅, MW 340) (**1**) and 5-*O*-PN (C₂₄H₂₄O₇, MW 424) were obtained as white amorphous powders by semi-synthetic method from *rac*-naringenin.¹³

A total of 24 microorganisms were evaluated for their ability to metabolize 6-DMAN and 5-*O*-PN using the usual two-stage fermentation procedure.¹⁰ Thin layer chromatographic analyses of the culture extracts during the screening studies indicated that *M. hiemalis*, *C. elegans* var. *elegans* and *P. chrysogenum* were capable of metabolizing **1** and **2**. Preparative scale fermentations of **1** and **2** led to the isolation of one new and four known metabolites (**3-7**).

Metabolite **3** was obtained as a white solid. Its $^1\text{H-NMR}$ spectra clearly indicated the changes in the dimethylallyl group of **1**, thus the olefinic proton signals at C-2'' (δ_{H} 6.44) and C-3'' (δ_{H} 5.36 and 5.43) in **1**

disappeared, and two new proton signals at δ_{H} 5.08 (t, $J=6.2$ Hz) and 3.08 (d, $J=6.2$ Hz) were shown in **3**. It was suggested that the 1,1-dimethylallyl group of **1** was rearranged and migrated to form the prenyl group at C-8 position of **3**. By comparison with the previously reported data,¹⁴ metabolite **3** was identified as (\pm)-8-prenylnaringenin.

Metabolite **4** was obtained as a pale yellow amorphous powder. ¹H-NMR spectral analyses indicated that the rearrangement of 1,1-dimethylallyl group in **1** was also involved in the formation of the prenyl group at C-8 of **4**. In addition, the two methyl groups at C-4" and C-5" of **4** were observed in the upfield region at δ_{H} 1.11 and 1.08, indicating that the prenyl group was further oxidized and cyclized to form the 1-hydroxy-1-methylethyldihydrofuranyl group. The absolute configuration of **4** at C-2 and C-2" was considered to be 2*S*, 2"*R* based on its negative specific rotation ($[\alpha]_{\text{D}}^{20}$ -120°).¹⁵ By comparison with the previously reported data,¹⁶ metabolite **4** was identified as (2*S*)-5,4'-dihydroxy-7,8-[(*R*)-2-(1-hydroxy-1-methylethyl)-2,3-dihydrofuran]flavanone.

Metabolite **5** was obtained as a pale yellow amorphous powder. HRESIMS spectrum gave a quasi-molecular ion $[M+H]^+$ peak at m/z 383.1494 suggesting the molecular formula of **5** to be C₂₂H₂₂O₆, which is 42 mass units lower than that of **2**. The ¹H- and ¹³C-NMR spectra of **5** also indicated the absence of one acetyl group compared with that of **2**. Thus, the proton signals at C-6 (δ_{H} 6.42) and C-8 (δ_{H} 6.32) of **2** moved to the upfield region at δ_{H} 5.95 and 6.04 in **5**, clearly indicating that the acetyl group at C-7 position was hydrolyzed. The $[\alpha]_{\text{D}}^{20}$ value of **5** was determined to be 0°, the same as *rac*-naringenin, indicated that **5** was also a racemate. On the basis of the spectral analyses and comparison with the spectral data of naringenin-4',7-diacetate,¹³ the chemical structure of **5** was assigned (\pm)-5-(*O*-prenyl)naringenin-4'-acetate.

Metabolite **6** was obtained as a pale yellow amorphous powder. The upfield signals at H-6 (δ 5.87) and H-8 (δ 5.84) together with the existence of 5-OH signal (δ_{H} 12.16) in its ¹H-NMR spectra clearly indicated that the acetyl group was substituted at C-4' position. By comparison with the previously reported data,¹⁷ metabolite **6** was identified as (\pm)-naringenin-4'-acetate.

Metabolite **7** was obtained as a white solid. No methyl group signal was observed in its ¹H-NMR spectra, indicating that all the substituted groups in **2** were hydrolyzed, and by comparison with the reported data,¹⁸ metabolite **7** was identified as (\pm)-naringenin.

In summary, the microorganisms were capable of transforming 6-DMAN (**1**) and 5-*O*-PN (**2**) into related metabolites. **1** was rearranged to **3** by *M. hiemalis*, which

was further oxidized and cyclized to form **4** by *C. elegans* var. *elegans*. **2** was regioselectively hydrolyzed to **5**, **6** and **7** by *P. chrysogenum*. It is considered that this work might thus contribute to a better understanding of the metabolic process of 6-DMAN and 5-*O*-PN in biological systems.

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