Microbial Transformation of Two Prenylated Naringenins

Fubo Han and Ik-Soo Lee*

College of Pharmacy and Research Institute of Drug Development, Chonnam National University, Gwangju 61186, Republic of Korea

Abstract – Microbial transformation of (±)-6-(1,1-dimethylallyl)naringenin (6-DMAN, 1) and (±)-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 (±)-8-prenylnaringenin (3), (2S)-5,4’-dihydroxy-7,8-[R]-2-[1-hydroxy-1-methylethyl]-2,3-dihydrofurano[flavanone (4), (±)-5-(O-prenyl)naringenin-4’-acetate (5), (±)-naringenin-4’-acetate (6), and (±)-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

(±)-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 (±)-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. Biotransformation 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-d6 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 chromato-
graphy, the adsorbent used was silica gel 60 (70 - 230 mesh, Merck). HPLC was performed on a Waters 600E Multisolv 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. 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** 7613
- **Filobasidium neoformans** 7902
- **Fusarium merismoides** 6153
- **Gliocladium deliquescens** 6173
- **Glomerella cingulata** 6075
- **Hormoconis resinae** 6966
- **Kluyveromyces marxianus** 7155
- **Metschnikowia pulcherrima** 7605
- **Monascus ruber** 6122
- **Mortierella ramanniana** var. **angulispora** 6137
- **Mucor hiemalis** 26779
- **Penicillium chrysogenum** 6933
- **Pichia pastoris** 7190
- **Rhizopus oryzae** 6946
- **Saccharomyces ludwigii** 7126
- **Torulaspora 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. 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; 

$^1$H-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 = 8.5 Hz, H-2), 2.78 (1H, dd, J = 3.1, 17.0 Hz, H-3b), 1.60 (3H, s, H-4''), 1.57 (3H, s, H-5''); 13C NMR (CDCl3, 75 MHz) δ 196.6 (C-4), 164.5 (C-7), 163.9 (C-5), 160.4 (C-9), 156.0 (C-2'), 149.6 (C-2'), 130.7 (C-1'), 127.4 (C-2''/6''), 115.6 (C-3''/5''), 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-2''), 26.6 (C-5'').

(±)-5-(O-Prenyl)naringenin-4'-acetate (5) – A pale yellow amorphous powder; [α]D20: 0° (c 0.052, MeOH); UV (MeOH) λmax: 286, 343 nm; IR (KBr) νmax cm−1: 3345, 1688, 1620, 1596, 1443, 1372, 1208; 1H-NMR (DMSO-d6, 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''); 13C NMR (DMSO-d6, 75 MHz) δ 187.0 (C-4), 169.2 (C-1''), 166.5 (C-7), 164.0 (C-9), 161.3 (C-5), 150.3 (C-3''), 136.8 (C-3''), 136.4 (C-4'), 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 [M+H]+ (calcd for C23H23O6, 383.1495).

Naringenin-4'-acetate (6) – A pale yellow amorphous powder; [α]D20: 0° (c 0.024, MeOH); 1H-NMR (DMSO-d6, 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 amorphous solid; [α]D20: 0° (c 0.021, MeOH); 1H-NMR (DMSO-d6, 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, 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), 2.68 (1H, dd, J = 2.8, 17.0 Hz, H-3b).

Results and Discussion

6-DMAN (C20H20O5, MW 340) (1) and 5-O-PN (C20H23O3, 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 1H-NMR spectra clearly indicated the changes in the dimethyllallyl group of 1, thus the olefinic proton signals at C-2'' (δH 6.44) and C-3'' (δH 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, metabolite 3 was identified as (±)-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-methylethylidihydrofuranyl group. The absolute configuration of 4 at C-2 and C-2'' was considered to be 2S, 2'R based on its negative specific rotation ( [α]_D^22 = -120º). By comparison with the previously reported data, metabolite 4 was identified as (2S)-5,4''-dihydroxy-7,8-[((R)-2-(1-hydroxy-1-methylethyl)-2,3-dihydrofuranyl]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 C_{22}H_{29}O_{10}, 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 [α]_D^20 value of 5 was determined to be 0°, the same as rac-naringenin, indicating that 5 was also a racemate. On the basis of the spectral analyses and comparison with the spectral data of naringenin-4',7-diacetate, the chemical structure of 5 was assigned (±)-5-(O-prenyl)naringenin-4'-acetate.

Metabolite 6 was obtained as a pale yellow amorphous powder. The upfield signals at H-6 (δ_H 5.87) and H-8 (δ_H 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, metabolite 6 was identified as (±)-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, metabolite 7 was identified as (±)-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.

Acknowledgements

We thank Gwangju Branch of the Korea Basic Science Institute (KBSI) for running NMR and MS experiments. This study was financially supported by Chonnam National University, 2014.

References

(1) Sumathi, R.; Tamizharasi, S.; Sivakumar, T. Int. J. Curr. Adv. Res. 2015, 4, 234-236.
(2) Ruh, M. F.; Zacharewski, T.; Connor, K.; Howell, J.; Chen, I.; Safe, S. Biochem. Pharmacol. 1995, 50, 1485-1493.
(3) Zierau, O.; Gester, S.; Schwab, P.; Metz, P.; Kolba, S.; Wulf, M.; Vollmer, G. Planta Med. 2002, 68, 449-451.
(4) Meijanto, E.; Hermawan, A.; Anindiyaji. Asian Pac. J. Cancer Prev. 2012, 13, 427-436.
(5) Assini, J. M.; Mulvihill, E. E.; Huff, M. W. Curr. Opin. Lipidol. 2013, 24, 34-40.
(6) Zierau, O.; Hamann, J.; Tischer, S.; Schwab, P.; Metz, P.; Vollmer, G.; Gutzeit, H. O.; Scholz, S. Biochem. Biophys. Res. Commun. 2005, 326, 909-916.
(7) Seo, E.-K.; Silva, G. L.; Chai, H.-B.; Chagwedera, T. E.; Farnsworth, N. R.; Cordell, G. A.; Pezzuto, J. M.; Kinghorn, A. D. Phytochemistry 1997, 45, 509-515.
(8) Zierau, O.; Morrissey, C.; Watson, R. W. G.; Schwab, P.; Kolba, S.; Metz, P.; Vollmer, G. Planta Med. 2003, 69, 856-858.
(9) Tokalov, S. V.; Henker, Y.; Schwab, P.; Metz, P.; Gutzeit, H. O. Pharmacology 2004, 71, 46-56.
(10) Clark, A. M.; McClesney, J. D.; Hufford, C. D. Med. Res. Rev. 1985, 5, 231-253.
(11) Han, F.; Lee, I.-S. Phytochem Lett. 2016, 18, 136-139.
(12) Han, F.; Lee, I.-S. Nat. Prod. Res. 2017, 31, 883-889.
(13) Gester, S.; Metz, P.; Zierau, O.; Vollmer, G. Tetrahedron 2001, 57, 1015-1018.
(14) Kim, H. J.; Kim, S. H.; Kang, B. Y.; Lee, I.-S. Arch. Pharm. Res. 2008, 31, 1241-1246.
(15) Tabara, S.; Ingham, J. L.; Mizutani, J. Agric. Biol. Chem. 1987, 51, 211-216.
(16) Jiang, D. S.; Cuendet, M.; Hawthorne, M. E.; Kardono, L. B. S.; Kawaniishi, K.; Fong, H. H. S.; Mehta, R. G.; Pezzuto, J. M.; Kinghorn, A. D. Phytochemistry 2002, 61, 867-872.
(17) Kyriakou, E.; PrimiJryi, A.; Charisiadis, P.; Katsoura, M.; Gerothanassiss, I. P.; Stamatis, H.; Tsakos, A. G. Org. Biomol. Chem. 2012, 10, 1739-1742.
(18) Malteae, F.; Erkelens, C.; van der Kooy, F.; Choi, Y. H.; Verpoorte, R. Food Chem. 2009, 116, 575-579.