Effects of Microbial Transformation on the Biological Activities of Prenylated Chalcones from *Angelica keiskei*

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Abstract: Microbial transformation is an alternative method for structural modification. The current study aimed at application of microbial transformation for discovering new derivatives and investigating the structure-activity relationship of isobavachalcone (1), 4-hydroxyderricin (2), and xanthoangelol (3) isolated from the herb *Angelica keiskei*. In the initial screening process, 1–3 were incubated with microbes using a two-stage fermentation method and analyzed through TLC monitoring. The screening results showed that *Rhizopus oryzae* and *Mucor hiemalis* were able to transform 1 and 2, respectively. Additionally, *M. hiemalis* and *Mortierella ramanniana* var. *angulispora* were able to transform 3. Following scale-up fermentation, four new (4, 5, 7, and 10) and five known (6, 8, 9, 11, and 12) metabolites were produced. Cytotoxicity of all the compounds (1–12) was investigated using three human cancer cell lines including A375P, HT-29, and MCF-7 by MTT method. Meanwhile, the tyrosinase inhibitory activity of 1–12 was evaluated using l-tyrosine as a substrate. Overall, 1 and 3 displayed the highest cytotoxicity, and 5 and 7 exhibited the most potent tyrosinase inhibitory activity with relatively low cytotoxicity. This allowed us to postulate that the introduction of 4'-O-glcucopyranosyl group led to the reduction in cytotoxicity and improvement in tyrosinase inhibitory activity.

Keywords: *Angelica keiskei*; microbial transformation; isobavachalcone; 4-hydroxyderricin; xanthoangelol; cytotoxicity; tyrosinase inhibition

1. Introduction

*Angelica keiskei* (Umbelliferae) is a perennial leafy herb, mainly distributed in Asian countries, including Korea and Japan. The herb is named as ‘Myeong-il yeob’ in Korean and ‘Ashitaba’ in Japanese, both literally meaning tomorrow’s leaves. Another prevalent name in Korean for *A. keiskei* is ‘Sinsuncho’ meaning the herb of eternal youth [1]. In daily life, it is drunk as a tea and cooked as a vegetable. Traditionally, it is consumed as medicinal herb with tonic, mild cathartic, diuretic, and galactagogue effects [2]. Preceding phytochemical investigations on *A. keiskei* revealed the presence of abundant prenylated chalcones in its leaves, stems, and roots [3]. These chalcones are proposed as effective agents for diverse health-beneficial properties such as anti-tumor [4], anti-inflammatory [5], anti-bacterial [6], anti-diabetic [7], anti-melanogenic [8], and anti-obesity [9] effects. Particularly, 4-hydroxyderricin (HD) and xanthoangelol (XT) are the predominant prenlychalcones in this herb, with relative abundances of 1.97% and 5.05%, respectively [10]. HD and XT have been proposed as potent cytotoxic agents that promote apoptosis and suppress tumor-induced angiogenesis in cancer cells [11]. After surgically removing implanted tumors, the administration of HD or XT inhibited metastasis and increased overall survival in mice. They were considered as promising therapeutic agents for the treatment of melanoma [12,13]. Isobavachalcone (IBC) is another bioactive prenylchalcone isolated from *A. keiskei* [14]. Accumulative studies have demonstrated that IBC suppressed the proliferation and induced cell apoptosis in diverse cancer cell lines including colorectal cancer, liver cancer, breast cancer, leukemia, tongue squamous cell carcinoma [15]. Meanwhile,
IBC demonstrated reduced cytotoxicity toward normal cells in comparison with cancer cells. After oral administration in mice, IBC inhibited the growth of subcutaneous HL-60 xenograft tumor without obvious toxicity [16].

Microbial transformation has been used as an effective means for structural modification of natural products at non-activated positions. It has advantages over conventional chemical synthesis largely due to its simpler operation, higher selectivity, and milder conditions [17]. In some cases, biological activities may have been enhanced for the transformed metabolites [18,19]. Previous studies on the microbial transformation of natural bioactive chalcones identified diverse metabolites with promising activities [20–22]. The aim of the study was to apply microbial transformation of three major bioactive prenylchalcones (1–3) in order to identify new derivatives and evaluate their biological activities to understand their structure-activity relationship.

2. Materials and Methods

2.1. Materials and Reagents

The herbal material Angelica keiskei was obtained in September 2020 and identified by the herb company Damaon (Yeongcheon, Gyeongsangbuk-do, Korea). Voucher specimen (AK2009) has been deposited at the Herbarium of the College of Pharmacy, Chonnam National University (Gwangju, Korea).

The microbial media including malt extract, peptone, D-glucose, and potato dextrose were obtained from Becton, Dickinson and Co. (Sparks, MD, USA). Sucrose was obtained from Sigma-Aldrich (St. Louis, MO, USA). Dulbecco’s modified Eagle medium (DMEM), penicillin, and streptomycin were purchased from Gibco (Invitrogen, Carlsbad, CA, USA). Fetal bovine serum (FBS) was purchased from Welgene Inc. (Gyeongsan, Gyeongsangbuk-do, Korea). Phosphate-buffered saline (PBS) tablets were obtained from Takara Korea Biomedical Inc. (Seoul, Korea). 3-(4,5-Dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) was obtained from Thermo Fisher Scientific (Waltham, MA, USA). Demethylzeylasteral (DZ) used as a reference standard in the bioassay was purchased from Biopurify Phytochemicals, Ltd. (Chengdu, China). Tyrosinase from mushroom and L-tyrosine were purchased from Sigma-Aldrich (St. Louis, MO, USA) and Daejung Chemicals and Metals Co., Ltd. (Siheung, Gyeonggi-do, Korea), respectively. Kojic acid used as a reference standard in the bioassay was purchased from Tokyo Chemical Industry Co., Ltd. (Tokyo, Japan).

The microbial strains including Cunninghamella elegans var. elegans KCTC 6992, Mortierella ramanniana var. angulispora KCTC 6137, Mucor hiemalis KCCM 26779, Mucor plumbeus KCCM 60265, Rhizopus oryzae KCCM 60556 were purchased from the Korean Collection for Type Cultures (KCTC, Daejeon, Korea) or Korean Culture Center of Microorganisms (KCCM, Seoul, Korea). The human cancer cell lines A375P, HT-29, and MCF-7 were obtained from the Korean Cell Line Bank (Seoul, Korea).

2.2. Isolation and Identification of Chalcones 1–3 from the Aerial Parts of A. keiskei

The air-dried herbal material (2.0 kg) was powdered and extracted by soaking in 95% ethanol (10 L × 3). After concentration of the filtrate, the residue (432 g) was suspended in water and sequentially partitioned with dichloromethane (CH₂Cl₂) and ethyl acetate (EtOAc). The crude CH₂Cl₂ residue (59.0 g) was subsequently separated by silica gel column chromatography using a gradient of n-hexane/EtOAc (30:1–1:1) and CH₂Cl₂/methanol (10:1–0:1) to yield eleven fractions (Fr. A–K). Fr. D (3.76 g) was further subjected to a Sephadex LH-20 column chromatography using methanol to yield five subfractions D1–D5. Subfr. D5 was separated using high performance liquid chromatography (HPLC) to provide substrates 1 (50 mg), 2 (95 mg), and 3 (110 mg). The structures of 1–3 were established as isobavachalcone (1) [23], 4-hydroxyderricin (2) [24] and xanthoangelol (3) [25] by 1H- and 13C-nuclear magnetic resonance (NMR) spectral analysis (see Supplementary Materials Figures S2–S7).
2.3. Preparation and Analysis of Screening Samples for Microbial Transformation

Fermentation experiments were carried out in two kinds of media. The malt medium consisting of peptone (1 g), D-glucose (20 g), and malt extract (20 g) was prepared in 1 L of distilled water for the incubation of *M. hiemalis*. The potato sucrose medium consisting of sucrose (20 g) and potato dextrose (24 g) was prepared in 1 L of distilled water for the culture of *C. elegans* var. *elegans*, *M. ramanniana* var. *angulispora*, *M. plumbeus*, or *R. oryzae*.

Initial fermentations were performed in 50 mL media. A two-stage fermentation method was employed in all experiments [26]. After inoculation and continued incubation for 24 h, the solution of each substrate (1–3) in ethanol (20 mg/mL) was distributed to each flask. The concentration of each substrate in the media was 0.02 mg/mL. Incubation was further continued for 4–6 days. The two parallel controls were conducted under the same conditions, i.e., substrate controls (substrate in microorganism-free culture media) and culture controls (microorganisms in substrate-free culture media). Substrate-containing microbial media were sampled by withdrawing 5 mL of entire media at every two days after addition of substrate. They were extracted with equal volumes of EtOAc. The organic layers were concentrated and spotted on thin layer chromatography (TLC) plates, and developed with chloroform/methanol (6:1). After spraying anisaldehyde-sulfuric acid reagent, the TLC plates were heated over 100 °C for detection of potential metabolites (Scheme 1).

![Scheme 1. Two-stage fermentation method for screening.](image)

On the basis of TLC analyses, the microbes *M. plumbeus* KCCM 60556 and *M. hiemalis* KCTC 26779 were selected for scale-up fermentations of substrates 1 and 2, respectively, as they showed higher transformational efficiency, while the microbes *M. hiemalis* KCTC 26779 and *Mortierella ramanniana* var. *angulispora* KCTC 6137 were selected for scale-up fermentations of 3, as they showed potential capability of transforming 3 (See Supplementary Materials Figure S1).

2.4. Preparation and Isolation of Scale-Up Samples to Obtain Metabolites 4–11

Scale-up fermentation of each substrate (1–3) was conducted following the aforementioned procedures in 150 mL of pre-cultured media. The scale-up fermentation culture of each substrate (1–3) was extracted thrice with equal volume of EtOAc. The combined EtOAc extract was evaporated in vacuo to give a residue.

The organic residue of 1 cultured with *R. oryzae* was separated on HPLC eluted with 53% to 84% methanol to yield metabolites 4 (t<sub>R</sub> = 30.4 min, 2.8 mg) and 6 (t<sub>R</sub> = 34.9 min, 6.5 mg) together with fractions RO1 (t<sub>R</sub> = 34.1 min) and RO2 (t<sub>R</sub> = 36.8 min) at 2.0 mL/min. Metabolite 5 (t<sub>R</sub> = 32.0 min, 3.7 mg) was obtained by a further purification of fr. RO1 on
HPLC with 55% to 72% methanol. Metabolite 7 (t_{R} = 36.8 min, 3.2 mg) was obtained by a further purification of fr. RO2 on HPLC with a gradient of 53% to 85% methanol.

The organic residue of 2 cultured with *M. hiemalis* was separated on HPLC with 58% to 85% methanol to furnish 8 (t_{R} = 16.6 min, 1.8 mg) and 9 (t_{R} = 26.4 min, 2.2 mg) at 2.0 mL/min.

The organic residue of 3 cultured with *M. hiemalis* was separated on HPLC with 58% to 97% methanol to yield metabolite 10 (3.5 mg, t_{R} = 32.8 min) at 2.0 mL/min.

The organic residue of 3 cultured with *Mortierella ramanniana* var. *angulispora* was separated on HPLC with 58% to 78% methanol to provide fraction MR1 (t_{R} = 32.8 min). Metabolites 11 (2.4 mg, t_{R} = 14.8 min) and 12 (2.2 mg, t_{R} = 17.8 min) were yielded by a further purification of fr. MR1 on HPLC with n-hexane: anhydrous EtOH (85: 15, v/v) at 1.0 mL/min.

### 2.5. Structural Characterization of New Metabolites 4, 5, 7, and 10

#### 2.5.1. 3′-(3-Hydroxy-3-methylbutyl)-4′-O-β-d-glucopyranosyl-4,2′-dihydroxyxalchone (4)

Yellow solid; [α]_{D}^{20} = 25.2 (c 0.10, MeOH); UV (MeOH) λ_{max}: 371 nm; IR ν_{max}: 3333, 2855, 1632, 1233, 990 cm\(^{-1}\); HRESIMS m/z: 527.1892 [M+Na]\(^{+}\) (calcd. for C_{26}H_{32}O_{10}Na, 527.1893); \(^{1}\)H- and \(^{13}\)C-NMR spectral data: see Table 1.

#### Table 1. \(^{1}\)H- and \(^{13}\)C-NMR data for 4, 5, 7, and 10.

| C/H     | δ_{H} (J/Hz) | δ_{C} | δ_{H} (J/Hz) | δ_{C} | δ_{H} (J/Hz) | δ_{C} | δ_{H} (J/Hz) | δ_{C} |
|---------|--------------|-------|--------------|-------|--------------|-------|--------------|-------|
| α       | 7.66 d (15.5)| 118.4 | 7.67 d (15.2)| 118.4 | 7.65 d (15.3)| 118.4 | 7.80 d (15.3)| 117.1 |
| β       | 7.82 d (15.5)| 144.6 | 7.84 d (15.2)| 144.6 | 7.83 d (15.3)| 146.4 | 7.84 d (15.3)| 145.0 |
| C = O   |              |       |              |       |              |       |              |       |
| 1       | 194.5        | 127.9 | 194.5        | 127.9 | 194.5        | 127.9 | 195.4        | 125.6 |
| 2,6     | 7.64 d (8.6) | 132.1 | 7.66 d (8.6) | 132.1 | 7.64 d (8.6) | 132.1 | 7.79 d (8.5) | 131.5 |
| 3,5     | 6.85 d (8.6) | 117.1 | 6.87 d (8.6) | 117.1 | 6.85 d (8.6) | 117.1 | 6.85 d (8.6) | 115.9 |
| 4       | 161.9        | 161.9 | 161.9        | 161.9 | 161.9        | 161.9 | 161.9        | 160.6 |
| 1′      |              | 116.6 | 116.6        | 116.6 | 116.6        | 116.6 | 116.6        | 114.5 |
| 2′      |              | 164.1 | 164.2        | 164.2 | 164.2        | 164.2 | 164.2        | 162.4 |
| 3′      |              | 120.8 | 120.5        | 120.5 | 120.5        | 120.5 | 120.5        | 117.1 |
| 4′      |              | 162.6 | 162.6        | 162.6 | 162.6        | 162.6 | 162.6        | 161.0 |
| 5′      | 6.80 d (9.0) | 107.2 | 6.82 d (9.1) | 107.1 | 6.80 d (9.0) | 107.1 | 6.74 d (9.2) | 105.9 |
| 6′      | 7.98 d (9.0) | 130.6 | 8.00 d (9.1) | 130.6 | 7.98 d (9.0) | 130.6 | 8.21 d (9.2) | 130.6 |
| 1″      | 2.81 m       | 18.6  | 2.78 m       | 18.2  | 2.77 m       | 18.3  | 3.44 m, 3.24 | 21.4  |
| 2″      |              | 43.2  | 1.7 m        | 39.2  | 1.69 m       | 39.7  | 5.22 m       | 121.9 |
| 3″      |              | 72.0  | 76.9         | 76.7  | 1.90 m       | 39.3  |              |       |
| 4″      | 1.28 s       | 28.7  | 1.27 s       | 25.1  | 1.26 s       | 25.7  | 1.90 m       | 26.2  |
| 5″      | 1.26 s       | 29.8  | 1.25 s       | 25.8  | 1.24 s       | 26.6  | 1.99 m       | 26.2  |
| 6″      |              | 3.31 s | 49.8         | 3.55 q (7.0) | 58.0  | 5.04 t (7.0) | 124.2 |
| 7″      |              |       | 1.21 t (7.0) | 16.5  |              |       |              |       |
| 8″      |              |       |              | 1.59 s | 25.5  |              |       |              |       |
| 9″      |              |       |              | 1.53 s | 17.5  |              |       |              |       |
| 10″     |              |       |              | 1.75 s | 16.0  |              |       |              |       |
| 1‴      | 5.01 d (7.5) | 102.0 | 5.06 d (7.3) | 101.9 | 5.04 d (7.3) | 102.0 | 5.00 d (7.3) | 100.1 |
| 2‴      | 3.53 m       | 75.0  | 3.52 m       | 75.1  | 3.51 m       | 75.1  | 3.32 m       | 73.4  |
| 3‴      | 3.49 m       | 78.5  | 3.51 m       | 78.5  | 3.49 m       | 78.5  | 3.39 m       | 77.3  |
| 4‴      | 3.41 m       | 71.4  | 3.42 m       | 78.4  | 3.41 m       | 71.4  | 3.18 m       | 69.8  |
| 5‴      | 3.49 m       | 78.2  | 3.50 m       | 71.4  | 3.49 m       | 78.3  | 3.32 m       | 76.8  |
| 6‴      | 3.92 dd (12.1, 2.0) | 3.93 dd (12.1, 2.1) | 3.92 dd (12.3, 2.1) | 3.92 dd (12.3, 5.3) | 3.71 m | 3.46 m | 60.7 |
| 1‴      | 3.72 dd (12.1, 5.0) | 3.74 dd (12.3, 5.3) | 3.72 dd (12.3, 5.3) | 3.72 dd (12.3, 5.3) | 3.71 m | 3.46 m | 60.7 |

Assignments were achieved on the basis of 1D and 2D NMR data. The values of J are presented in parentheses.

*a* Data were acquired in methanol-d₄. *b* Data were acquired in DMSO-d₆.
2.5.2. 3′-(3-O-Methyl-3-methylbutyl)-4′-O-β-D-glucopyranosyl-4,2′-dihydroxychalcone (5)

Yellow solid; [α]_{D}^{20} = -26.9 (c 0.10, MeOH); UV (MeOH) \( \lambda_{\text{max}} \): 365 nm; IR \( \nu_{\text{max}} \): 3330, 2927, 1780, 1284, 1105, 1074 cm\(^{-1}\); HRESIMS \( m/z \): 541.2049 [M+Na]\(^+\) (calcd. for C\(_{27}\)H\(_{34}\)O\(_{10}\)Na, 541.2050); \(^{1}\)H- and \(^{13}\)C-NMR spectral data: see Table 1.

2.5.3. 3′-(3-O-Ethyl-3-methylbutyl)-4′-O-β-D-glucopyranosyl-4,2′-dihydroxychalcone (7)

Yellow solid; [α]_{D}^{20} = -46.5 (c 0.10, MeOH); UV (MeOH) \( \lambda_{\text{max}} \): 363 nm; IR \( \nu_{\text{max}} \): 3319, 2919, 1604, 1285, 1105, 835 cm\(^{-1}\); HRESIMS \( m/z \): 555.2208 [M+Na]\(^+\) (calcd. for C\(_{28}\)H\(_{36}\)O\(_{10}\)Na, 555.2206); \(^{1}\)H- and \(^{13}\)C-NMR spectral data: see Table 1.

2.5.4. 4′-O-β-D-Glucopyranosyl xanthoangelol (10)

Yellow solid; [α]_{D}^{20} = -28.0 (c 0.10, MeOH); UV (MeOH) \( \lambda_{\text{max}} \): 365 nm; IR \( \nu_{\text{max}} \): 3342, 2926, 1713, 1631, 1233, 1076 cm\(^{-1}\); HRESIMS \( m/z \): 577.2414 [M+Na]\(^+\) (calcd. for C\(_{31}\)H\(_{38}\)O\(_{9}\)Na, 577.2414); \(^{1}\)H- and \(^{13}\)C-NMR spectral data: see Table 1.

2.6. Acid Hydrolysis of 4, 5, 7, and 10

Each solution of compounds 4, 5, 7, and 10 (each 0.8 mg) in 2 mL 2N HCl was heated at 85 °C for 3 h. After cooling and neutralization, the reaction mixture was extracted with EtOAc. The water layer was concentrated and confirmed by TLC in comparison with authentic D-glucose as a standard. The organic layer containing the aglycone of 10 was determined as the corresponding substrate 3 by comparing their retention time on HPLC.

2.7. Cytotoxic Activity Evaluation

The cytotoxicity assays were conducted according to the MTT method as previously described [27] using human melanoma A375P, human colorectal HT-29, and human breast cancer MCF-7 cell lines. Briefly, cells were incubated in 96-well plates at an approximate density of 5.0 × 10\(^3\) cells/well with 100 µL DMEM medium supplemented with 5% FBS, penicillin (100 U/mL) and streptomycin (100 µg/mL) at 37 °C with 5% CO\(_2\) for 24 h. Then, cells were fed with different concentrations of the test compounds (1–12) for another 48 h. After gently removing the culture media, 100 µL of MTT solution (0.5 mg/mL) was applied for staining the cells. Finally, absorbance of each plate was observed at 490 nm by the microplate reader after removal of MTT solution and addition of 100 µL dimethyl sulfoxide.

2.8. Tyrosinase Inhibitory Activity

L-Tyrosine was used as a substrate to evaluate the tyrosinase inhibition of 1–12 following the previously developed method [28,29] with some modifications. The tyrosinase inhibition assay was performed using 0.1 M phosphate buffer (pH 6.5), 2 mM L-tyrosine, and test compound solution with different concentrations ranging from 10 µM to 100 µM. The mixture was prepared 5 min before treatment with the enzyme solution (150 U/mL). In the experiments, the reaction mixture without tyrosinase was used as a blank, and the reaction mixture without sample solution was used as a negative control. Kojic acid was used as a positive control. After incubation for 30 min at 37 °C, absorbance of the reaction mixture was recorded at 490 nm to evaluate the amount of dopachrome produced in the process. The extent of tyrosinase inhibition by compounds 1–12 is displayed as the half maximal inhibitory concentration (IC\(_{50}\)) in Table 2.
Table 2. Cytotoxic and tyrosinase inhibitory activities of 1–12 a.

| Compound | IC<sub>50</sub> ± SD (µM) | A375P | HT-29 | MCF-7 | Tyrosinase |
|----------|--------------------------|-------|-------|-------|-----------|
| 1        | 9.78 ± 1.06              | 26.33 ± 2.42 | 24.02 ± 0.36 | >100 |
| 2        | 25.34 ± 0.66             | 37.30 ± 4.86 | 29.71 ± 2.78 | >100 |
| 3        | 13.00 ± 1.28             | 24.35 ± 1.27 | 21.92 ± 2.30 | >100 |
| 4        | >80                      | >80     | >80   | 78.47 ± 3.13 |
| 5        | 77.96 ± 3.44             | 54.36 ± 1.03 | 32.44 ± 0.55 | >100 |
| 6        | >80                      | >80     | >80   | 33.32 ± 1.16 |
| 7        | >80                      | >80     | >80   | 43.12 ± 6.74 |
| 8        | 29.65 ± 2.09             | 54.36 ± 4.19 | 28.18 ± 1.34 | >100 |
| 9        | 21.18 ± 0.20             | >80     | >80   | 52.65 ± 2.85 |
| 10       | 29.03 ± 2.77             | >80     | >80   | 71.71 ± 9.22 |
| 11       | 69.77 ± 3.88             | >80     | >80   | >100 |
| 12       | 20.56 ± 2.17             | >80     | >80   | >100 |
| DZ       | 2.52 ± 0.23              | 8.68 ± 0.90 | 2.87 ± 0.38 | 77.74 ± 2.55 |
| Kojic acid|                         |         |       |       |

a Results are expressed as the mean values of three experiments ± SD.

3. Results

Microbial transformation of three bioactive prenylchalcones 1–3 is reported herein. Fermentation of 1 with *R. oryzae* KCCM 60,556 furnished three new (4, 5, and 7) and one known (6) metabolites. Fermentation of 2 with *M. hiemalis* KCTC 26,779 furnished two known (8 and 9) metabolites. Fermentation of 3 with *M. hiemalis* KCTC 26,779 furnished one new (10) metabolite, while that with *M. ramanniana* var. *angulispora* KCTC 6137 resulted in the production of two known (11 and 12) metabolites (Figure 1) (Figures S8–S41).

![Figure 1. Compounds 4–12 obtained by microbial transformation of 1–3.](image-url)
were quite similar to those of ortho-9.0) and 6.80 (1H, d, J = 8.6 Hz) and 6.85 (2H, d, J = 8.6 Hz). Meanwhile, an olefin moiety was identified by the observation of signals at δ_H 7.82 (1H, d, J = 15.5) and 7.66 (1H, d, J = 15.5). These typical resonances suggested the existence of a 4,2′,3′,4′-tetrasubstituted chalcone. The two multiplets at δ_H 2.81 (2H) and 1.66 (2H) and two singlets at δ_H 1.28 (3H) and 1.26 (3H) were attributed to the 3-hydroxy-3-methylbutyl group, whereas the anemic proton at δ_H 5.01 (1H, d, J = 7.5 Hz), together with the characteristic carbon signals at δ_C 102.0, 78.5, 78.2, 75.0, 71.4, and 62.6 demonstrated that the sugar moiety was a β-D-glucopyranosyl-(3-β-D-methyl-3-methylbutyl)-4′-O-β-D-glucopyranosyl-4,2′-dihydroxycalcone.

Compound 5 was assigned the composition C_{27}H_{34}O_{10} from its HRESIMS at m/z 541.2049 [M+Na]^+ (calcd. for C_{27}H_{34}O_{10}Na, 541.2050). The 1H- and 13C-NMR spectra of 5 were quite similar to those of 4, except for the resonances of a methoxy group (δ_C 33.1/δ_C 49.8) in 5. The attachment of the methoxy group was determined to be at C-3′′ according to the HMBC correlations from the methoxy protons H-6′′′/1.66 (H-2′′′) and δ_C 76.9 and C-2′′′ (δ_C 39.2). Thus, the structure of 5 was determined as 3′-(3-O-methyl-3-methylbutyl)-4′-O-β-D-glucopyranosyl-4,2′-dihydroxycalcone.

The molecular formula of compound 7 was determined as C_{28}H_{36}O_{10} using HRESIMS on the basis of a molecular ion at m/z 555.2208 [M+Na]^+ (calcd. for C_{28}H_{36}O_{10}Na,555.2206), which corresponded to 11 degrees of unsaturation. The NMR data of 7 were almost identical to those observed for 4 except for the appearance of one O-methylene (δ_C 3.55, 2H, q, J = 7.0 Hz/δ_C 58.0) and one methyl (δ_H 1.21, 3H, t, J = 7.0 Hz/δ_C 16.5) signals corresponding to H-6′ and H-7′′, readily attributable to an ethyloxy group. Further evidence was obtained from the obvious correlation between proton signals H-6′′ and H-7′′ (Figure 2) in the 1H-1H homonuclear correlation spectroscopy (COSY) experiment. HMBC correlation between H-6′′ and C-3′′ confirmed the location of the ethyloxy group at C-3′′. Accordingly, the structure of 7 was assigned 3′-(3-O-ethyl-3-methylbutyl)-4′-O-β-D-glucopyranosyl-4,2′-dihydroxycalcone.

Figure 2. Key COSY (1H-1H) and HMBC (1H→13C) correlations of compounds 7 and 10.

Compound 10 was confirmed to have C_{31}H_{38}O_{9} as the elemental composition according to the analysis of its HRESIMS data. The 1H-NMR spectrum of compound 10 displayed two sets of aromatic proton resonances at δ_H 8.21 (1H, d, J = 9.2) and 6.74 (1H, d, J = 9.2), 7.79 (2H, d, J = 8.5), and 6.85 (2H, d, J = 8.5) together with two olefinic proton resonances at δ_H 7.84 (1H, d, J = 15.3) and 7.80 (1H, d, J = 15.3) assignable to the 4,2′,3′,4′-tetrasubstituted chalcone. In addition, proton resonance signals for a geranyl group and
a sugar moiety were identified in the up-field region. Analysis of the COSY spectrum together with the heteronuclear single quantum coherence (HSQC) and HMBC spectra allowed the establishment of the structure as 4,2′,4′-trihydroxychalcone substituted with a geranyl group and a sugar moiety in 10 (Figure 2). The $^{13}$C-NMR data showed six signals at δC 100.1, 77.3, 76.8, 73.4, 69.8, and 60.7, which are characteristic of a glucose moiety [31]. The anomeric proton signal at δH 5.00 (1H, d, $J = 7.3$ Hz) indicated a β-configuration of the glucosidic bond. The connectivity of the glucose moiety in 10 was confirmed to be at C-4′ through an ether linkage by the HMBC correlation from the anomeric proton signal H-1′′ to C-4′. Presence of the glucose moiety and its aglycone after acid hydrolysis was further confirmed by comparison of their TLC and HPLC identification pattern with those of the authentic samples. Linkage of the geranyl group was deduced to be at C-3′ position by the observation of HMBC correlations from H-1′′ at 5.00 (1H, d, $J = 7.3$ Hz) to C-4′. The information could be beneficial for understanding the metabolic pathways of these compounds in this plant.

In this study, cytotoxicity of compounds 1–12 was determined by MTT assays against A375P, HT-29, and MCF-7 human cancer cells. As shown in Table 2, compounds 1 and 3 showed most potent cytotoxicity against A375P, HT-29, and MCF-7. Compound 2 exhibited moderate cytotoxic activity toward human cancer cell lines tested. However, all the metabolites 4–12 exhibited lower cytotoxicity compared with their corresponding substrates 1, 2, and 3. It appears that the C-prenyl or C-geranyl side chain is essential for cytotoxic activity. Meanwhile, glucosylation or methylation at 4′-OH decreased the cytotoxicity.

The capability of 1–12 to inhibit tyrosinase was determined using L-tyrosine as a substrate. The most potent compounds were 5 and 7, which displayed respectively 2.5- and 2-fold stronger inhibition compared with the positive control. Compound 4 showed moderate activity comparable to kojic acid. It was suggested that methyl or ethyl moiety introduced at the 3-O position of 3-hydroxy-3-methylbutyl group could improve the inhibitory activity against mushroom tyrosinase. The anti-tyrosinase potencies were enhanced in chalcone 4′-O-glucosides (6 and 10) compared with their corresponding aglycones (1 and 3). This allowed us to postulate that the introduction of 4′-O-glucopyranosyl group led to the improvement in the tyrosinase inhibitory capacity.

4. Discussion

In this work, three bioactive prenylchalcones, isobavachalcone (IBC, 1), 4-hydroxyderricin (HD, 2), and xanthoangelol (XT, 3), were isolated from A. keiskei and subjected to microbial transformation. The reaction types consisted of hydroxylation, methylation, ethylation, rearrangement, and glucosylation. These highly specific reactions are difficult to achieve by synthetic methods, especially under mild conditions. Therefore, microbial transformation is a useful way in structural diversification of chalcones to discover valuable derivatives. In addition, microbial transformation permits the possibility of understanding the metabolic pathways [37]. It is particularly noteworthy that microbial transformation of HD (2) with the microbe M. hiemalis produced two metabolites xanthoangelol H (8) and D (9), while microbial transformation of XT (3) with the microbe Mortierella ramanniana var. angulispora resulted in the production of two metabolites xanthoangelin A (11) and xanthoangelol B (12). All the four known metabolites (8, 9, 11, and 12) were previously identified from the herb A. keiskei. The information could be beneficial for understanding the metabolic pathways of these compounds in this plant.

Tyrosinase is a metalloenzyme containing two copper atoms and acts as a rate-limiting oxidase involved in the production of melanin [36]. During melanogenesis, tyrosinase is able to act as a hydroxylase to catalyze the conversion of L-tyrosine to L-3,4-dihydroxyphenylalanine (L-DOPA) and subsequently is able to act as an oxidase to catalyze the conversion of L-DOPA to L-dopaquinone, which results in the accumulation
of melanin and hyperpigmentation [39]. In recent years, it has been demonstrated that various dermatological disorders, such as age spots, melasma freckle, and sites of actinic damage, derive from the accumulation of an exaggerated level of epidermal pigmentation. Tyrosinase has been recognized as a significant target for the treatment of skin disorders related to irregular pigmentation [40]. Thus, the identification of new and potent leads with tyrosinase inhibitory activity has attracted considerable interest in medication and cosmetics. Chalcones represent a diverse class of compounds generally presented in higher plants. Structurally, compounds with the scaffold of 4-hydroxychalcone are considered as potent tyrosinase inhibitors owing to the structural similarity between 4-hydroxychalcone and tyrosine (Figure 3) [41]. In this study, the structure-activity relationship of twelve compounds (1–12) with the scaffold of 4-hydroxychalcone was investigated for its tyrosinase inhibitory activity. The results revealed that the glucosylation of the OH group at C-4′ position is important in the enhancement of the inhibition. Furthermore, the insertion of a methyl (compound 5) or an ethyl group (compound 7) into the 3-hydroxy-3-methylbutyl group at the 3-O position dramatically improved the inhibition. Considering that 5 and 7 showed the most potent tyrosinase inhibition with relatively low cytotoxicity, as shown in Table 2, they can act as potential leads for the development of effective and safe anti-browning and skin-whitening agents.

![Figure 3. The structural similarity between 4-hydroxychalcone and tyrosine.](image)

5. Conclusions

Microbial transformation of three bioactive prenylchalcones 1–3 isolated from A. keiskei afforded four new (4, 5, 7, and 10) and five known (6, 8, 9, 11, and 12) metabolites. The five known metabolites were all previously reported from natural sources. In the evaluation of all compounds for their cytotoxic and tyrosinase inhibitory activities, metabolites 5 and 7 exhibited the most potent tyrosinase inhibition with relatively low cytotoxicity. The two metabolites 5 and 7 can be considered as new leads for the further design of tyrosinase inhibitors. The present results indicate that microbial transformation can be an efficient and useful tool for the identification of natural product-like compounds as bioactive leads.

**Supplementary Materials:** The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/foods11040543/s1, Figure S1: TLC analyses for microbial transformation of 1–3 by selected microbes, Figures S2–S41: 1D, 2D NMR, and HRESIMS spectra of 1–12.

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References

1. Kil, Y.-S.; Pham, S.T.; Seo, E.K.; Jafari, M. Angelica keiskei, an emerging medicinal herb with various bioactive constituents and biological activities. Arch. Pharm. Res. 2017, 40, 655–675. [CrossRef] [PubMed]

2. Zhang, L.; Jiang, Y.; Pang, X.; Hua, P.; Gao, X.; Li, Q.; Li, Z. Simultaneous optimization of ultrasound-assisted extraction for flavonoids and antioxidant activity of Angelica keiskei using response surface methodology (RSM). Molecules 2019, 24, 3461. [CrossRef] [PubMed]

3. Caesar, L.K.; Cech, N.B. A review of the medicinal uses and pharmacology of ashitaba. Planta Med. 2016, 82, 1236–1245. [CrossRef] [PubMed]

4. Akihisa, T.; Tokuda, H.; Hasegawa, D.; Ukiya, M.; Kimura, Y.; Enjo, F.; Suzuki, T.; Nishino, H. Chalcones and other compounds from the exudates of Angelica keiskei and their cancer chemopreventive effects. J. Nat. Prod. 2006, 69, 38–42. [CrossRef] [PubMed]

5. Aoki, N.; Muko, M.; Ohta, E.; Ohta, S. C-geranylated chalcones from the stems of Angelica keiskei with superoxide-scavenging activity. J. Nat. Prod. 2008, 71, 1308–1310. [CrossRef] [PubMed]

6. Inamori, Y.; Baba, K.; Tsujiro, H.; Taniguchi, M.; Nakata, K.; KOZAWA, M. Antibacterial activity of two chalcones, xanthoangelol and 4-hydroxyderricin, isolated from the root of Angelica keiskei KOIDZUMI. Chem. Pharm. Bull. 1991, 39, 1604–1605. [CrossRef]

7. Enoki, T.; Ohnogi, H.; Nagamine, K.; Kudo, Y.; Sugiyama, K.; Tanabe, M.; Kobayashi, E.; Sagawa, H.; Kato, I. Antidiabetic activities of chalcones isolated from a Japanese herb, Angelica keiskei. J. Agric. Food Chem. 2007, 55, 6013–6017. [CrossRef]

8. Arung, E.T.; Furuta, S.; Sugamoto, K.; Shimizu, K.; Ishikawa, H.; Matsushita, Y.-I.; Kondo, R. The inhibitory effects of representative chalcones contained in Angelica keiskei on melanin biosynthesis in B16 melanoma cells. Nat. Prod. Commun. 2012, 7, 1007–1010. [CrossRef]

9. Zhang, C.; Wu, W.; Li, X.; Xin, X.; Liu, D. Daily supplementation with fresh Angelica keiskei juice alleviates high-fat diet-induced obesity in mice by modulating gut microbiota composition. Mol. Nutr. Food Res. 2019, 63, 1900248. [CrossRef]

10. Kim, D.W.; Curtis-Long, M.J.; Yuk, H.J.; Wang, Y.; Song, Y.H.; Jeong, S.H.; Park, K.H. Quantitative analysis of phenolic metabolites from different parts of Angelica keiskei by HPLC–ESI MS/MS and their xanthine oxidase inhibition. Food Chem. 2014, 153, 20–27. [CrossRef]

11. Sumiyoshi, M.; Taniguchi, M.; Baba, K.; Kimura, Y. Antitumor and antimetastatic actions of xanthoangelol and 4-hydroxyderricin isolated from Angelica keiskei roots through the inhibited activation and differentiation of M2 macrophages. Phytomedicine 2015, 22, 759–767. [CrossRef]

12. Kimura, Y.; Taniguchi, M.; Baba, K. Antitumor and antimetastatic activities of 4-hydroxyderricin isolated from Angelica keiskei roots. Planta Med. 2004, 70, 211–219.

13. Zhang, T.; Wang, Q.; Fredimoses, M.; Gao, G.; Wang, K.; Chen, H.; Wang, T.; Oi, N.; Zykova, T.A.; Reddy, K. The ashitaba (Angelica keiskei) chalcones 4-hydroxyderricin and xanthoangelol suppress melanomagenesis by targeting BRAF and PI3K. Cancer Prog. Res. 2018, 11, 607–620. [CrossRef]

14. Lee, H.; Li, H.; Kweon, M.; Choi, Y.; Kim, M.J.; Ryu, J.-H. Isobavachalcone from Angelica keiskei inhibits adipogenesis and prevents lipid accumulation. Int. J. Mol. Sci. 2018, 19, 1693. [CrossRef]

15. Wang, M.; Lin, L.; Lu, J.-J.; Chen, X. Pharmacological review of isobavachalcone, a naturally occurring chalcone. Pharmacol. Res. Commun. 2021, 165, 105483. [CrossRef]

16. Wu, D.; Wang, W.; Chen, W.; Lian, F.; Lang, L.; Huang, Y.; Xu, Y.; Zhang, N.; Chen, Y.; Liu, M. Pharmacological inhibition of dihydroorotate dehydrogenase induces apoptosis and differentiation in acute myeloid leukemia cells. Haematologica 2018, 103, 1472. [CrossRef]

17. Cao, H.; Chen, X.; Jassbi, A.R.; Xiao, J. Microbial biotransformation of active flavonoids. Biotechnol. Adv. 2015, 33, 214–223. [CrossRef]

18. Chen, G.; Yang, M.; Nong, S.; Yang, X.; Ling, Y.; Wang, D.; Wang, X.; Zhang, W. Microbial transformation of 20 (S)-protopanaxadiol by Absidia corymbifera. Cytotoxic activity of the metabolites against human prostate cancer cells. Fitoterapia 2013, 84, 6–10. [CrossRef]

19. Farooq, R.; Hussain, N.; Yousuf, S.; Ahmad, M.S.; Choudhary, M.I. Microbial transformation of mestanalone by Macrophomina phaseolina and Cunninghamella blakesleae and anticancer activities of the transformed products. RSC Adv. 2018, 8, 21985–21992. [CrossRef]

20. Akihisa, T.; Motoi, T.; Seki, A.; Kikuchi, T.; Fukatsu, M.; Tokuda, H.; Suzuki, N.; Kimura, Y. Cytotoxic activities and anti-tumor-promoting effects of microbial transformation products of prenylated chalcones from Angelica keiskei. Chem. Biodivers. 2012, 9, 318–330. [CrossRef]

21. Kozłowska, J.; Potaniec, B.; Żarowska, B.; Aniol, M. Microbial transformations of 4′-methylchalcones as an efficient method of obtaining novel alcohol and dihydrochalcone derivatives with antimicrobial activity. RSC Adv. 2018, 8, 30379–30386. [CrossRef]
22. Kim, H.J.; Yim, S.-H.; Han, F.; Kang, B.Y.; Choi, H.J.; Jung, D.-W.; Williams, D.R.; Gustafson, K.R.; Kennelly, E.J.; Lee, I.-S. Biotransformed metabolites of the hop prenylflavanone isoxanthohumol. *Molecules* 2019, 24, 394. [CrossRef] [PubMed]

23. Rong, Y.; Wu, J.; Liu, X.; Zhao, B.; Wang, Z. Study on structural and spectral properties of isobavachalcone and 4-hydroxyderricin by computational method. *Spectrochim. Acta A Mol. Biomol. Spectrosc.* 2014, 126, 254–259. [CrossRef] [PubMed]

24. Kim, J.H.; Son, Y.K.; Kim, G.H.; Hwang, K.H. Xanthoangelol and 4-hydroxyderricin are the major active principles of the inhibitory activities against monoamine oxidases on *Angelica keiskei* K. *Biomol. Ther.* 2013, 21, 234. [CrossRef] [PubMed]

25. Shin, J.E.; Choi, E.J.; Jin, Q.; Jin, H.-G.; Woo, E.-R. Chalcones isolated from *Angelica keiskei* and their inhibition of IL-6 production in TNF-α-stimulated MG-63 cell. *Arch. Pharm. Res.* 2011, 34, 437–442. [CrossRef] [PubMed]

26. Clark, A.M.; McChesney, J.D.; Hufford, C.D. The use of microorganisms for the study of drug metabolism. *Med. Res. Rev.* 1985, 5, 231–253. [CrossRef] [PubMed]

27. Mosmann, T. Rapid colorimetric assay for cellular growth and survival: Application to proliferation and cytotoxicity assays. *J. Immunol. Methods* 1983, 65, 55–63. [CrossRef]

28. Vincent, J.; Hearing, J. Mammalian monophenol monooxygenase (tyrosinase): Purification, properties, and reactions catalyzed. *Methods Enzymol.* 1987, 142, 154–165. [CrossRef]

29. Deering, R.W.; Chen, J.; Sun, J.; Ma, H.; Dubert, J.; Barja, J.L.; Seeram, N.P.; Wang, H.; Rowley, D.C. N-acyl dehydrotyrosines, tyrosinase inhibitors from the marine bacterium *Thalassotalea* sp. PP2-459. *J. Nat. Prod.* 2016, 79, 447–450. [CrossRef]

30. Ninomiya, M.; Efdi, M.; Inuzuka, T.; Koketsu, M. Chalcone glycosides from aerial parts of *Brassica rapa* L. ‘hidabeni’, turnip. *Phytochem. Lett.* 2010, 3, 96–99. [CrossRef]

31. Cioffi, G.; Escobar, L.M.; Braca, A.; De Tommasi, N. Antioxidant chalcone glycosides and flavanones from *Maclura* (Chlorophora) tinctoria. *J. Nat. Prod.* 2003, 66, 1061–1064. [CrossRef]

32. Kumazawa, S.; Suzuki, S.; Ahn, M.-R.; Kamihira, M.; Udagawa, Y.; Bang, K.-S.; Nakayama, T. A new chalcone from propolis collected on Jeju island, Korea. *Food Sci. Technol.* 2006, 12, 67–69. [CrossRef]

33. Sugamoto, K.; Matsusita, Y.-i.; Matsui, K.; Kurogi, C.; Matsui, T. Synthesis and antibacterial activity of chalcones bearing prenyl or geranyl groups from *Angelica keiskei*. *Phytochemistry* 2011, 67, 5346–5359. [CrossRef]

34. Baba, K.; Nakata, K.; Taniguchi, M.; Kido, T.; Kozawa, M. Chalcones from *Angelica keiskei*. *Phytochemistry* 1990, 29, 3907–3910. [CrossRef]

35. Shanu-Wilson, J.; Evans, L.; Wrigley, S.; Steele, J.; Atherton, J.; Boer, J. Biotransformation: Impact and application of metabolism in drug discovery. *ACS Med. Chem. Lett.* 2020, 11, 2087–2107. [CrossRef]

36. Deckert, H.; Schweikardt, T.; Tuczek, F. The first crystal structure of tyrosinase: All questions answered? *Angew. Chem. Int. Ed.* 2006, 45, 4546–4550. [CrossRef]

37. Chang, T-S. An updated review of tyrosinase inhibitors. *Int. J. Mol. Sci.* 2009, 10, 2440–2475. [CrossRef]

38. Nerya, O.; Musa, R.; Khatib, S.; Tamir, S.; Vaya, J. Chalcones as potent tyrosinase inhibitors: The effect of hydroxyl positions and numbers. *Phytochemistry* 2004, 65, 1389–1395. [CrossRef] [PubMed]