Chemical Constituents of the Leaves of *Campanula takesimana* (Korean Bellflower) and Their Inhibitory Effects on LPS-induced PGE$_2$ Production

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Abstract: *Campanula takesimana* Nakai (Campanulaceae; Korean bellflower) is one of the endemic herbs of Korea. The plant has been used as traditional medicines for treating asthma, tonsillitis, and sore throat in Korea. A hot water extract of the leaves of *C. takesimana* exhibited a significant inhibitory effect on lipopolysaccharide (LPS)-stimulated prostaglandin E$_2$ (PGE$_2$) production. Repetitive chromatographic separation of the hot water extract led to the isolation of three new neolignan glucosides, campanulalignans A–C ($^1$–$^3$), with 15 known compounds ($^4$–$^{18}$). The structures of new compounds ($^1$–$^3$) were elucidated by analyzing nuclear magnetic resonance (NMR) spectroscopic data, along with high resolution quadrupole time of flight mass spectrometric data. Among the isolates, simplidin ($^7$), 5-hydroxyconiferaldehyde ($^11$), icariside F$_2$ ($^{12}$), benzyl-α-L-arabinopyranosyl-(1″→6″)-β-D-glucopyranoside ($^{13}$), and kaempferol 3-O-β-D-apiosyl (1→2)-β-D-glucopyranoside ($^{15}$) were isolated from the Campanulaceae family for the first time. The isolates ($^1$, $^2$, and $^4$–$^{18}$) were assessed for their anti-inflammatory effects on LPS-stimulated PGE$_2$ production on RAW 264.7 cells. 7R,8S-Dihydrodehydrodiconiferyl alcohol ($^5$), 3′,4′-dimethylecedrusin 9-O-β-D-glucopyranoside ($^6$), pinoresinol di-O-β-D-glucoside ($^8$), ferulic acid ($^{10}$), 5-hydroxyconiferaldehyde ($^{11}$), and quercetin ($^{18}$) showed significant inhibitory effects on LPS-stimulated PGE$_2$ production.

Keywords: *Campanula takesimana*; Campanulaceae; campanulalignans; phenolic compounds; PGE$_2$ production

1. Introduction

*Campanula takesimana* Nakai (Campanulaceae), known as Korean bellflower, is a native herb of Korea, growing on Ulleng island. The leaves and stems of *C. takesimana* are shiny, and the flowers have spots on a pale purple background. It is clearly distinguished from *C. punctata*, commonly distributed in Korea, China, and Japan, by its smaller overall size and lighter purple color of flower [1]. Because it has long flowering period and shape of flower, this plant has been commonly cultivated for a decorative plant. Besides the value of horticulture, roots of *C. takesimana* have been used for herbal remedies for asthma, tonsillitis, and sore throat in traditional Korean medicine [2].
It was reported that the n-butanol and ethyl acetate fractions of *C. takesimana* exhibit free radical-scavenging, tyrosinase inhibition, and superoxide dismutase (SOD)-like activities [2]. Although extensive research has been done on the pharmacological activities and/or phytochemical constituents of the plants belonging to family Campanulaceae, yet only few studies have been conducted on the genus *Campanula*, despite their frequent traditional use [3]. Moreover, chemical constituents of *C. takesimana* have never been reported from previous studies.

As part of our continuing project to search for bioactive compounds with anti-inflammatory activities and identify chemical constituents of higher plants, we observed that a hot water extract from the leaves of *C. takesimana* showed a significant inhibitory effect on lipopolysaccharide (LPS)-stimulated prostaglandin E$_2$ (PGE$_2$) production in RAW 264.7 macrophages. LPS is one of the most potent activators of macrophages, and LPS-stimulated macrophages and monocytes are known to produce inflammatory mediators. In this regard, inflammatory mediators, such as PGE$_2$, nitric oxide (NO), and numerous pro-inflammatory cytokines released by the activated macrophages are important targets for the treatment of inflammation [4]. Thus, compounds that can regulate the production of pro-inflammatory mediators like PGE$_2$ may be promising therapeutic agents for the treatment of various inflammatory diseases.

In the present research, we concentrated on isolating secondary metabolites from the hot water extract and identifying the active constituents. Three new neolignan glucosides (1–3) and 15 known compounds (4–18) were identified by repetitive chromatographic separation of the hot water extract. The structure of the new compounds (1–3) was elucidated by analysis of one-dimensional (1D) and two-dimensional (2D) nuclear magnetic resonance (NMR) spectroscopic and high resolution quadrupole time of flight mass (HR-Q-TOF-MS) spectrometric data. The structures of the known compounds were identified in comparison with previously published values. Thereafter, the isolates were assessed for their inhibitory effects on the production of the pro-inflammatory mediator, PGE$_2$, in LPS-stimulated RAW264.7 macrophages. We describe in this paper the isolation of the secondary metabolites from the leaves of *P. japonicus*, structure elucidation of the three new neolignan glucosides (1–3), and inhibitory effects of the isolates on PGE$_2$ production.

2. Results

2.1. Structure Elucidation of New Compounds 1–3

By repetitive chromatographic separation of the hot water extract, three new neolignan glucosides (1–3) were isolated, along with 15 known compounds, including five lignans (4–8), five phenolic compounds (9–13), and five flavonoids (14–18), in the present research (Figure 1).

![Figure 1. Structures of compounds 1–18 isolated from Campanula takesimana.](image-url)
Compound 1 was isolated as a pale yellow powder. Based on its molecular ion peak [M–H]− at m/z 697.2360 (calcd for C32H41O17, 697.2344) in negative mode of HR-Q-TOF-MS (Figure S1), the molecular formula of 1 was established as C32H32O17. The infrared absorption (IR) spectrum displayed absorption bands at 3278, 2903, and 1580 cm−1, implying that compound 1 has phenolic group characteristics.

The 1H NMR spectroscopic data of 1 (Table 1, Figure S2) revealed one aromatic singlet signal (δH 6.71 (1H, H-6)) and 1,3,4,5-tetrasubstituted aromatic ring protons (δH 6.48 (1H, d, J = 2.0 Hz, H-2′) and 6.47 (1H, d, J = 2.0 Hz, H-6′)). The existence of two trans-olefinic groups was identified on the position of the proton signals and coupling constants (δH 6.86 (1H, d, J = 15.5 Hz, H-7), 6.24 (1H, dt, J = 15.0, 5.0 Hz, H-8) and 6.20 (1H, dt, J = 15.0, 5.0 Hz, H-8′), 6.14 (1H, d, J = 15.5 Hz, H-8′)). Additionally, two methylene (δH 4.24 (2H, dd, J = 5.5, 2.0 Hz, H-9) and 3.55 (2H, d, J = 5.5 Hz, H-9′)) and two methoxy proton signals (δH 3.86 (3H, s, OCH3-5), 3.81 (3H, s, OCH3-5′)) were also observed in the 1H NMR. Two anemic protons (δH 4.67 (1H, d, J = 8.0 Hz, Glc H-1), 4.63 (1H, d, J = 8.0 Hz, Glc H-1′)) and overlapped signals at δH 3.23–3.82 suggested that 1 has two sugars with β configuration [5]. The 13C NMR exhibited 32 signals, containing 12 signals for two glucopyranosyl moieties (Table 1, Figure S3). Enzyme hydrolysis of 1 and high performance liquid chromatography (HPLC) experiment led to the determination of the sugars of 1 as β-D-glucopyranose. In combination with 13C NMR and HMBC experimental results, compound 1 had two phenylpropanoid units that had π bond between α and β carbons (Figure S6). Each phenylpropyl moiety had one methoxy and one β-D-glucopyranosyl, approximate to demethyl syringin (9), supported by HMBC correlations (Figure 2). The differences were that one of the two phenylpropanoid units in 1 had 2-substituted aromatic ring and the other one had modified C-9′ into C-C bond instead of hydroxyl group. Furthermore, both of the substituted parts connected to each other were determined by HMBC correlation signal between H-9′ and C-2. Compound 1 was very similar to the known neolignan glucoside, tangshenoside III [6], except for missing one methoxy group in each of the two aromatic rings. Therefore, the structure of the new compound 1 was elucidated as 3,3′-demethyl tangshenoside III, and named as campanulalignan A.

![Figure 2. Key 1H-1H COSY (red) and HMBC (blue) correlations of compounds 1-3.](image-url)

Compound 2 was obtained as a pale yellow powder. The molecular formula of 2 was assigned as C32H32O17 by its HR-Q-TOF-MS ion peak at m/z 697.2354 [M–H]− (calcd for C32H41O17, 697.2344) in negative mode (Figure S7) and was the same as 1. The 1H and 13C NMR spectra of 2 strongly resembled those of 1. However, the carbon chemical shift of one methoxy group in aromatic ring moved to the downfield (δC 62.7) in 2 (Table 1, Figure S9). In the 1H NMR, the difference between compounds 1 and 2 was shown on the aromatic singlet signal of 2 shifted to the downfield (δH 6.87 (1H, s, H-6)), whereas two olefinic protons of 2 shifted to the upfield (δH 6.81 (1H, d, J = 15.5 Hz, H-7) and δH 6.17 (1H, dt, J = 15.5, 6.0 Hz, H-8)) (Table 1, Figure S8). By analyzing the HMBC spectrum, the carbon that linked to the methoxy proton showed correlation with methylene proton peaks at δH 3.55 (2H, m, H-9′) (Figure 2). For these reasons, the location of one methoxy group was suggested to be C-3 of aromatic ring, rather than C-5 of 1. Thus, the structure of the new compound 2 was elucidated as 5,3′-demethyl tangshenoside III, and named as campanulalignan B.
Table 1. $^1$H and $^{13}$C NMR spectroscopic data of compounds 1–3 (δ in ppm, CD$_3$OD, 500 and 125 MHz).

| Position $^a$ | 1      | 2      | 3      |
|---------------|--------|--------|--------|
|               | $^\delta$C | $^\delta$H $^b$ | $^\delta$C | $^\delta$H $^b$ | $^\delta$C | $^\delta$H $^b$ |
| 1             | 134.9  | 135.4  | 140.0  |
| 2             | 119.5  | 123.9  | 119.0  |
| 3             | 152.4  | 153.0  | 150.1  |
| 4             | 134.9  | 139.3  | 134.6  |
| 5             | 150.1  | 150.8  | 152.5  |
| 6             | 102.5  | 6.71 s  | 110.8  | 6.87 s  | 102.9  | 6.71 s  |
| 6′            | 129.3  | 6.86 d (15.5) | 128.9  | 6.81 d (15.5) | 72.1  | 5.40 d (5.5) |
| 7             | 132.0  | 6.24 dt (15.0, 5.0) | 130.0  | 6.17 dt (15.5, 6.0) | 141.7  | 6.01 ddd (16.5, 10.0, 5.0) |
| 9             | 62.2   | 4.24 dd (5.5, 2.0) | 63.9   | 4.21 dd (5.5, 2.0) | 115.2  | 5.26 d (17.0) |
| 1′            | 136.9  | 136.6  | 136.9  |
| 2′            | 108.2  | 6.48 d (2.0) | 108.2  | 6.49 d (2.0) | 103.1  | 6.47 d (1.5) |
| 3′            | 151.9  | 152.0  | 152.5  |
| 4′            | 134.5  | 134.6  | 134.6  |
| 5′            | 154.1  | 154.3  | 154.3  |
| 6′            | 103.0  | 6.47 d (2.0) | 103.1  | 6.48 d (2.0) | 108.4  | 6.48 d (1.5) |
| 7′            | 130.9  | 6.14 (15.5) | 130.4  | 6.13 d (15.5) | 131.0  | 6.18 d (16.0) |
| 8′            | 129.9  | 6.20 dt (15.0, 5.0) | 131.4  | 6.24 dt (15.5, 5.5) | 130.1  | 6.24 dt (15.5, 6.0) |
| 9′            | 29.9   | 3.55 d (5.5) | 30.2   | 3.55 d (5.5) | 29.5   | 3.54 d (6.0) |
| Glc-1         | 107.3  | 4.67 d (8.0) | 107.3  | 4.81 d (8.0) | 107.3  | 4.67 d (8.0) |
| Glc-2         | 71.0   | 3.45 overlap | 71.0   | 3.49 m  | 71.0   | 3.51 overlap |
| Glc-3         | 77.8   | 3.40 overlap | 78.0   | 3.43 overlap | 77.8   | 3.45 overlap |
| Glc-4         | 75.5   | 3.49 overlap | 75.5   | 3.43 overlap | 75.5   | 3.42 overlap |
| Glc-5         | 78.5   | 3.22 m  | 78.6   | 3.22 overlap | 78.7   | 3.23 overlap |
| Glc-6         | 62.0   | 3.68/3.82 overlap | 62.4   | 3.82 overlap | 62.3   | 3.86 m  |
| Glc-1′        | 107.0  | 4.63 d (8.0) | 107.0  | 4.64 d (8.0) | 107.1  | 4.63 d (7.5) |
| Glc-2′        | 71.0   | 3.47 overlap | 71.0   | 3.47 m  | 71.0   | 3.51 overlap |
| Glc-3′        | 77.8   | 3.43 overlap | 77.9   | 3.84 overlap | 77.8   | 3.45 overlap |
| Glc-4′        | 75.5   | 3.51 m  | 75.5   | 3.44 m  | 75.5   | 3.42 overlap |
| Glc-5′        | 78.5   | 3.30 m  | 78.5   | 3.29 m  | 78.7   | 3.23 overlap |
| Glc-6′        | 62.0   | 3.68/3.82 overlap | 62.2   | 3.22 overlap | 62.3   | 3.78 m  |
| OCH$_2$-5′    | 56.8   | 3.86 s  | 62.7   | 3.87s  | 56.8   | 3.84 s  |
| OCH$_3$-5′    | 56.8   | 3.81 s  | 56.8   | 3.81 s  | 56.7   | 3.81 s  |

$^a$ All assignments were based on $^1$H–$^1$H correlation spectroscopy (COSY), $^1$H–$^{13}$C heteronuclear single quantum coherence spectroscopy (HSQC), and $^1$H–$^{13}$C heteronuclear multiple bond correlation (HMBC) results. $^b$ δH (multi J in Hz).

Compound 3 had a molecular formula of C$_{32}$H$_{42}$O$_{17}$, as established by its molecular ion peak [M–H]$^-$ at m/z 697.2336 (calcd for C$_{32}$H$_{41}$O$_{17}$, 697.2344) in negative mode of HR-Q-TOF-MS (Figure S13) and was also the same as 1 and 2. The $^1$H and $^{13}$C NMR spectra of 3 were almost identical with those of 1 and 2, except for the existence of a vinyl alcohol group in 3 instead of an allyl alcohol group in 1 and 2. $^1$H NMR spectra of 3 exhibited a vinyl alcohol signals at δ$_{H}$ 5.40 (d, $J = 5.5$ Hz, H-7), 6.01 (dd, $J = 16.5$, 10.0, 5.0 Hz, H-8), 5.26 (d, $J = 17.0$ Hz, H-9a), 5.12 (d, $J = 10.5$ Hz, H-9b) (Table 1). The $^{13}$C NMR and HSQC of 3 indicated the presence of a vinyl alcohol group, which includes one oxygenated carbon signal (δ$_C$ 72.1 (C-7)), one olefinic carbon signal (δ$_C$ 141.7 (C-8)), and one exomethylene signal (δ$_C$ 115.2 (C-9)), (Table 1, Figure S16). The presence of the vinyl alcohol group in 3 was also supported by $^1$H–$^1$H COSY data, which showed the correlation at H-7/H-8 and H-8/H-9a, 9b (Figure 2, Figure S17). The location of the vinyl alcohol group was unambiguously determined by the HMBC correlation from H-6 to C-7 to be at C-1. Two β-d-glucopyranosyl groups and two methoxy groups were located at C-4, C-4′, C-5, and C-5′, respectively, as 1, through HMBC spectral interpretation (Figure 2). The chemical structure of 3 was established by the results noted above and by comparison with tangshenosides II and III [6,7], as shown in Figure 2, and named as campanulalignan C. However, the absolute configuration at C-7 of 3 was not determined due to the limited amount of sample obtained.

By comparison of their spectroscopic data with those published, the structures of the known compounds were identified as 3′,4-O-dimethylcedrusin (4) [8], 7R,8S-dihydrodehydrodiconiferyl
alcohol (5) [9], 3’,4-O-dimethylcedrusin 9-O-β-glucopyranoside (6) [10], simplidin (7) [11], pinoresinol di-O-β-d-glucoside (8) [12], demethyl syringin (9) [13], ferulic acid (10) [14], 5-hydroxyconiferaldehyde (11) [15], icariside F₂ (12) [16], benzyl-α-L-arabinopyranosyl-(1”→6’)-β-d-glucopyranoside (13) [17], astragalin (14) [18], kaempferol 3-O-β-d-apiosyl(1→2)-β-d-glucopyranoside (15) [19], nicotiflorin (16) [20], kaempferol 3,7-di-O-β-d-glucoside (17) [21], and quercetin (18) [22].

2.2. Inhibitory Effects of Compounds on LPS-Induced PGE₂ Production in RAW 264.7 Macrophages

The isolates 1, 2, and 4–18 were evaluated for their inhibitory effects of LPS-stimulated PGE₂ production in RAW 264.7 macrophages. Compound 3 was not subjected to this experiment due to the limited amount obtained. Except for 2, 3, and 4, all other compounds did not exhibit cytotoxicity in cell viability tests at concentrations up to 100 µM. The effects were assessed using the inhibitory rate on PGE₂, which are listed in Table 2. Quercetin (18) showed the most potent inhibitory effect on LPS-induced PGE₂ production in RAW 264.7 macrophages (97% inhibitory rate at 50 µM concentration), among isolated compounds from C. takesimana. 5-Hydroxyconiferaldehyde (11) also decreased remarkably the PGE₂ production with 85.17% inhibitory rate. Additionally, 7R,8S-dihydrodehydrodiconiferyl alcohol (5), simplidin (7), pinoresinol di-O-β-d-glucoside (8), and ferulic acid (10) showed inhibitory effects over 50% on the PGE₂ production at 50 µM concentration while the new compounds 1 and 2 did not exhibit a significant effect (<50% inhibitory rate). Among the active compounds, quercetin (18), 7R,8S-dihydrodehydrodiconiferyl alcohol (5), pinoresinol di-O-β-d-glucoside (8), and ferulic acid (10) have already been reported to have anti-inflammatory effects, whereas there is no report on biological activities related to anti-inflammatory effects for simplidin (7) and 5-hydroxyconiferaldehyde (11). Therefore, inhibitory effects of 7 and 11 on LPS-induced PGE₂ production at various concentrations were investigated.

| Compound | IC₅₀ (µM) Cytotoxicity | Inhibition Rate (%) a | Compound | IC₅₀ (µM) Cytotoxicity | Inhibition Rate (%) a |
|----------|------------------------|-----------------------|----------|------------------------|-----------------------|
| 1        | >50                    | 23.92                 | 10       | >50                    | 62.53                 |
| 2        | 33.66                  | 33.07                 | 11       | >50                    | 85.17                 |
| 3        | -                      | -                     | 12       | >50                    | 31.07                 |
| 4        | 30.69                  | 7.89                  | 13       | >50                    | 33.12                 |
| 5        | >50                    | 69.35                 | 14       | >50                    | 23.35                 |
| 6        | >50                    | 20.01                 | 15       | >50                    | 43.49                 |
| 7        | >50                    | 60.01                 | 16       | >50                    | 36.73                 |
| 8        | >50                    | 38.08                 | 17       | >50                    | 34.51                 |
| 9        | >50                    | 37.44                 | 18       | >50                    | 97.01                 |

a Cells were pretreated with 50 µM concentration of compounds for 1 h, then with LPS (100 ng/mL), and incubated for 24 h.

As shown in Figure 3A,B, simplidin (7) and 5-hydroxyconiferaldehyde (11) significantly and concentration-dependently suppressed LPS-stimulated PGE₂ production. Compounds 7 and 11 did not influence the RAW 264.7 cell viability in either the presence or absence of LPS at concentrations up to 50 µM for 24 h, suggesting that 7 and 11 may be major constituents of the leaves of C. takesimana showing anti-inflammatory activities. Accordingly, further pharmacological studies are demanded to investigate the mechanisms responsible for the inhibitory effects of 7 and 11 on inflammatory mediator in RAW 264.7 macrophages.

Table 2. The cytotoxicities and inhibitory effects of the isolates from C. takesimana on lipopolysaccharide (LPS)-induced prostaglandin E₂ (PGE₂) production in RAW 264.7 macrophages.
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Figure 3. Inhibitory effects of compounds 7 (A) and 11 (B) on LPS-stimulated PGE_2 production in RAW 264.7 macrophages. Cells were pretreated with different concentrations (20, 40, or 80 μM) of compounds 7 and 11 for 1 h, then with LPS (100 ng/mL), and incubated for 24 h. N-[2-(cyclohexyloxy)-4-nitrophenyl]-methanesulfonamide (NS-398, 10 μM) was used as a positive PGE_2 production inhibitor. Level of PGE_2 in culture media was quantified using enzyme-linked immunosorbent assay (ELISA) kits. (#: p < 0.05 as compared with the untreated group, ** p < 0.05, and ***p < 0.0002).

3. Discussion

The investigation on secondary metabolites in the leaves of C. takesimana was carried out for the first time. Consequently, three new neolignan glucosides, campanulalignans A–C (1–3), were isolated by repeated chromatography. Additionally, five lignans (4–8), five phenolic compounds (9–13), and five flavonoids (14–18) were obtained from the present study. To the best of our knowledge, 3’,4-O-dimethylcedrusin 9-O-β-glucopyranoside (6), simplidin (7), 5-hydroxyconiferaldehyde (11), icariside F2 (12), benzyl-α-L-arabinopyranosyl-(1′→6′)-β-d-glucopyranoside (13), and kaempferol 3-O-β-d-apiosyl-(1→2)-β-d-glucopyranoside (15) were isolated from the Campanulaceae family for the first time in this work.

All the isolates from the leaves of C. takesimana except for compound 3 were estimated for their inhibitory effects on the production of PGE_2 in the LPS-induced RAW 264.7 macrophages. Inflammatory mediators, such as NO and PGE_2, and numerous pro-inflammatory cytokines released by the activated macrophages are important targets for the treatment of various inflammatory diseases, including multiple sclerosis, rheumatoid arthritis, and obesity [23,24]. iNOS and COX-2 protein expression are remarkably elevated in the inflammatory processes, resulting in increased production of NO and PGE_2, respectively. These overexpressed pro-inflammatory mediators can lead to damage in living cells and tissues, resulting in various physiological disorders associated with inflammation [4,23]. Therefore, compounds that inhibit the production of pro-inflammatory mediators like NO and PGE_2 can be candidates for anti-inflammatory drugs [4].

Among the isolates, well known quercetin (18) showed the strongest inhibitory effect [25]. In addition, 7R,8S-dihydrodehydrodiconiferyl alcohol (5), pinoresinol di-O-β-p-glucoside (8), and ferulic acid (10) exhibited remarkable inhibitory activity on LPS-stimulated PGE_2 production in RAW264.7 cells and their inhibitory effects were well correlated with previous reports [26–28]. 7R,8S-Dihydrodehydrodiconiferyl alcohol (5), a neolignan, showed ABTS free radical scavenging and LPS-stimulated nitric oxide (NO) production inhibitory activities in a previous report [26]. Pinoresinol di-O-β-p-glucoside (8) possesses protective activity on human endothelial cells by inhibiting oxidized low density lipoprotein (oxLDL)-induced SOD activity, NO production, and endothelial nitric oxide synthase (eNOS) expression [27]. A prior study revealed that ferulic acid (10) significantly reduced LPS-induced PGE_2 levels [28].

Simplidin (7) and 5-hydroxyconiferaldehyde (11) also showed significant inhibitory effects on PGE_2 production in this study, but there was no investigation on biological activities
related to anti-inflammatory effects in previous studies. Simplidin (7), first isolated from Firmiana simplex (Sterculiaceae) [11], exhibited cytotoxicities against human cancer cell lines [29]. 5-Hydroxyconiferaldehyde (11) was discovered in the biosynthetic pathway for syringyl monolignol in angiosperms [30] and identified from the methanol extract of Carthamus tinctorius seed [15]. However, to the best of our knowledge, there have been no reports of any biological or pharmacological activities for 5-hydroxyconiferaldehyde (11). In our investigation and for the first time, simplidin (7) and 5-hydroxyconiferaldehyde (11) are suggested as active natural products with inhibitory effects on a pro-inflammatory mediator, PGE$_2$ production.

4. Materials and Methods

4.1. Plant Material

The leaves of Campanula takesimana Nakai (Campanulaceae; Korean bellflower) used in present study were provided by COSMAX BIO, Gyeonggi-do, Republic of Korea, in July 2019. The origin of the material was identified by D.S.J., one of authors, and a voucher specimen (CATA-2019) has been deposited in the Laboratory of Natural Product Medicine, College of Pharmacy, Kyung Hee University, Seoul, Republic of Korea.

4.2. General Experimental Procedures

General experimental procedures are in the Supplementary Materials.

4.3. Extraction and Isolation

Dried leaves of C. takesimana (2.5 kg) were extracted with distilled water (60 L) at 100 °C for 5 h, and the solvent was evaporated by rotary evaporator at 45 °C. The hot water extract (1.0 kg) was separated over column chromatography (CC) using Diaion HP-20 as the stationary phase with a gradient system of acetone-H$_2$O (0:100 to 100:0, v/v) to give 17 fractions (F1~F17). Fraction F10 was fractionated further using Sephadex LH-20 CC (20.2 cm, CH$_3$Cl-MeOH-H$_2$O = 80:18:2 to 70:27:3, v/v/v) to give 16 subfractions (F10-1~F10-16). Compounds 6 (175.3 mg, 0.00701%) was isolated from subfraction F10-2-6 by a flash chromatographic system with a Redi Sep-C18 cartridge (43 g, MeOH-H$_2$O, from 15:85 to 45:55, v/v). Subfraction F12-2-9 was also subjected to a flash chromatographic system by a Redi Sep-C18 cartridge (43 g, MeOH-H$_2$O, from 13:87 to 40:60, v/v) to afford 8 (14.3 mg, 0.000572%).

Subfraction F12-4-10 was fractioned using silica gel CC (230–400 mesh; 3.1 × 25.0 cm, CH$_2$Cl$_2$-MeOH-H$_2$O = 85:13.5:1.5 to 70:27:3, v/v/v) to produce 14 subfractions (F12-2-1~F12-2-14). Compound 6 (175.3 mg, 0.00701%) was isolated from subfraction F12-2-6 by a flash chromatographic system with a Redi Sep-C18 cartridge (43 g, MeOH-H$_2$O, from 15:85 to 45:55, v/v). Subfraction F12-2-9 was also subjected to a flash chromatographic system by a Redi Sep-C18 cartridge (43 g, MeOH-H$_2$O, from 13:87 to 40:60, v/v) to afford 8 (14.3 mg, 0.000572%).

Subfraction F12-4-10 was fractioned using silica gel CC (230–400 mesh; 3.1 × 25.0 cm, CH$_2$Cl$_2$-MeOH-H$_2$O = 85:13.5:1.5 to 70:27:3, v/v/v) to produce 14 subfractions (F12-2-1~F12-2-14). Compound 6 (175.3 mg, 0.00701%) was isolated from subfraction F12-2-6 by a flash chromatographic system with a Redi Sep-C18 cartridge (43 g, MeOH-H$_2$O, from 15:85 to 45:55, v/v). Subfraction F12-2-9 was also subjected to a flash chromatographic system by a Redi Sep-C18 cartridge (43 g, MeOH-H$_2$O, from 13:87 to 40:60, v/v) to afford 8 (14.3 mg, 0.000572%).

Subfraction F12-2-10 was fractioned by using silica gel CC (230–400 mesh; 3.1 × 25.0 cm, CH$_2$Cl$_2$-MeOH-H$_2$O = 85:13.5:1.5 to 70:27:3, v/v/v) to produce 14 subfractions (F12-2-1~F12-2-14). Compound 6 (175.3 mg, 0.00701%) was isolated from subfraction F12-2-6 by a flash chromatographic system with a Redi Sep-C18 cartridge (43 g, MeOH-H$_2$O, from 15:85 to 45:55, v/v). Subfraction F12-2-9 was also subjected to a flash chromatographic system by a Redi Sep-C18 cartridge (43 g, MeOH-H$_2$O, from 13:87 to 40:60, v/v) to afford 8 (14.3 mg, 0.000572%).
954:5:0.5 to 60:36:4 v/v/v). Fraction F15-5 was subjected to silica gel CC (230–400 mesh; 5.1 × 33.0 cm, CH₂Cl₂-MeOH-H₂O = 90:9:1 to 60:36:4, v/v/v) to isolate 10 (11.2 mg, 0.000448%), 14 (91.2 mg, 0.00365%), 15 (17.0 mg, 0.000704%), and 16 (32.4 mg, 0.0013%). Fraction F16 was separated into nine subfractions (F16-1~F16-9) using Sephadex LH-20 CC (4.5 × 45.9 cm) with 70% MeOH. Subfraction F16-1 was subjected to silica gel CC (230–400 mesh; 5.1 × 33.0 cm, CH₂Cl₂-MeOH-H₂O = 90:9:1 to 60:36:4, v/v/v) to afford 4 (293.0 mg, 0.0117%) and 5 (79.5 mg, 0.00318%). Compound 18 (10.1 mg, 0.000404%) was purified by recrystallizing subfraction F17 with MeOH.

4.3.1. Campanulalignan A (1)

Pale yellow powder; HR-Q-TOF-MS (negative mode) m/z = 697.2360 [M–H]⁻ (calcd for C₃₂H₄₁O₁₇, 693.2344); [α]D²⁰ = −12.7° (c 0.1, MeOH); UV (MeOH) λmax nm (log ε): 224 (3.31), 265 (3.08); IR (ATR) νmax 3278, 2903, 1580, 1448, 1348 cm⁻¹; ¹H and ¹³C NMR data, see Table 1.

4.3.2. Campanulalignan B (2)

Pale yellow powder; HR-Q-TOF-MS (negative mode) m/z = 697.2354 [M–H]⁻ (calcd for C₃₂H₄₁O₁₇, 693.2344); [α]D²⁰ = −14.5° (c 0.1, MeOH); UV (MeOH) λmax nm (log ε): 222 (3.24), 263 (3.00); IR (ATR) νmax 3302, 2920, 1580, 1421, 1347 cm⁻¹; ¹H and ¹³C NMR data, see Table 1.

4.3.3. Campanulalignan C (3)

Brown amorphous powder; HR-Q-TOF-MS (negative mode) m/z = 697.2336 [M–H]⁻ (calcd for C₃₂H₄₁O₁₇, 693.2344); [α]D²⁰ = −24.5° (c 0.1, MeOH); UV (MeOH) λmax nm (log ε): 230 (3.47), 267 (3.22); IR (ATR) νmax 3266, 2922, 1585, 1342, 1017 cm⁻¹; ¹H and ¹³C NMR data, see Table 1.

4.4. Enzymatic Hydrolysis of Compound 1

Compound 1 (1.0 mg) was incubated together with 2 drops of toluene, β-glucosidase (2.5 mg), and H₂O (1.5 mL) in a CO₂ incubator at 35 °C for 3 days. The reaction was stopped by addition EtOH to the reaction mixture and β-glucosidase was removed by filtration. The presence of glucose in 1 was confirmed by co-TLC (Rf 0.3, n-BuOH:acetic acid:H₂O = 2:1:1, v/v/v) with a standard compound.

4.5. Absolute Configurations of β-glucose in Compound 1

The absolute configuration of β-glucose in 1 was confirmed by the method from Tanaka et al. [31]. Hydrolysat was dissolved in pyridine (500 µL) and L-cysteine methyl ester hydrochloride (1.2 mg) was added and heated at 60 °C for 1 h. The mixture was heated again at 60 °C for 1 h after adding α-tolyl isothiocyanate (100 µL) and analyzed directly by HPLC under a gradient system (A: 0.1% (v/v) formic acid in water, B: 0.1% (v/v) formic acid in acetonitrile, 10 to 50% B, 45 min). The reaction mixture of 1 was detected at 27.0 min. The retention times of authentic L- and D-glucoses were 26.1 and 27.0 min, respectively, under the same HPLC conditions. Thus, the absolute configuration of β-glucose in 1 was confirmed as the D configuration.

4.6. Cell Culture

RAW 264.7 macrophages were purchased from the Korea Cell Line Bank (Seoul, South Korea). Cells were incubated in Dulbecco’s modified Eagle’s medium (DMEM) supplemented with penicillin-streptomycin sulfate (100 units/mL and 100 µg/mL) and 10% FBS at 37 °C humidified incubator containing of 5% CO₂.

4.7. Cell Viability Test

Cell viability test was assessed by the MTT (3-[4,5-dimethylthiazol-2-yl]-2,5-dipheyl tetrazoliumbromide; Sigma-Aldrich) assay. RAW 264.7 macrophages were grown with each isolated compound and LPS (100 ng/mL) for 24 h. After 24 h, cells were treated with MTT solution for 4 h.
MTT formazan was resolved by adding dimethyl sulfoxide (DMSO), and the absorbance of each well at 540 nm was read by a microplate reader.

4.8. Measurement of PGE$_2$ Production

RAW 264.7 cells were treated with compounds (50 µM) 1 h prior to LPS (100 ng/mL) stimulation for 24 h. Cells were pretreated with positive controls (NS398) for 1 h, and then stimulated with LPS (100 ng/mL) for 24 h. The supernatant was collected and PGE$_2$ production detected by using PGE$_2$ ELISA Kits (R&D Systems, MN, USA).

5. Conclusions

Three new neolignan glucosides (1–3) and 15 known compounds were isolated from the leaves of C. takesimana in the present study. We found that 7R,8S-dihydrodehydrodiconiferyl alcohol (5), simplidin (7), pinoresinol di-O-β-d-glucoside (8), ferulic acid (10), 5-hydroxyconiferaldehyde (11), and quercetin (18) inhibit production of an inflammatory mediator PGE$_2$. Among these, the inhibitory effect of simplidin (7) and 5-hydroxyconiferaldehyde (11) on PGE$_2$ production was reported for the first time in this work. Thus, simplidin (7) and 5-hydroxyconiferaldehyde (11) are worthy of further pharmacological evaluation for their potential as anti-inflammatory drugs.

Supplementary Materials: The following are available online at http://www.mdpi.com/2223-7747/9/9/1232/s1, Figure S1: HR-Q-TOF-MS spectrum of compound 1, Figure S2: The $^1$H-NMR (500 MHz, CD$_3$OD) spectrum of compound 1, Figure S3: The $^{13}$C-NMR (125 MHz, CD$_3$OD) spectrum of compound 1, Figure S4: The HSQC spectrum of compound 1 in CD$_3$OD, Figure S5: The COSY spectrum of compound 1 in CD$_3$OD, Figure S6: The HMBC spectrum of compound 1 in CD$_3$OD, Figure S7: HR-Q-TOF-MS spectrum of compound 2, Figure S8: The $^1$H-NMR (500 MHz, CD$_3$OD) spectrum of compound 2, Figure S9: The $^{13}$C-NMR (125 MHz, CD$_3$OD) spectrum of compound 2, Figure S10: The HSQC spectrum of compound 2 in CD$_3$OD, Figure S11: The COSY spectrum of compound 2 in CD$_3$OD, Figure S12: The HMBC spectrum of compound 2 in CD$_3$OD, Figure S13: HR-Q-TOF-MS spectrum of compound 3, Figure S14: The $^1$H-NMR (500 MHz, CD$_3$OD) spectrum of compound 3, Figure S15: The $^{13}$C-NMR (125 MHz, CD$_3$OD) spectrum of compound 3, Figure S16: The HSQC spectrum of compound 3 in CD$_3$OD, Figure S17: The COSY spectrum of compound 3 in CD$_3$OD, Figure S18: The HMBC spectrum of compound 3 in CD$_3$OD

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