New Antifeedant Grayanane Diterpenoids from the Flowers of *Pieris formosa*

Chun-Huan Li 1,†, Shi-Hong Luo 2,†, Sheng-Hong Li 2,*,† and Jin-Ming Gao 1,*,†

1 Shaanxi Key Laboratory of Natural Products & Chemical Biology, College of Chemistry & Pharmacy, Northwest A & F University, Yangling 712100, China; chunhuanli@nwsuaf.edu.cn
2 State Key Laboratory of Phytochemistry and Plant Resources in West China, Kunming Institute of Botany, Chinese Academy of Sciences, Lanhei Road 132, Kunming 650201, China; lsh246967@126.com
* Correspondence: shli@mail.kib.ac.cn (S.-H.L.); jinminggao@nwsuaf.edu.cn (J.-M.G.);
Tel.: +86-871-652-23035 (S.-H.L.); +86-029-8709-2515 (J.-M.G.); Fax: +86-871-652-23035 (S.-H.L.)
† These authors contributed equally.

Received: 8 August 2017; Accepted: 24 August 2017; Published: 31 August 2017

Abstract: Three new grayanane diterpenoids, pierisoids C-E (1–3), as well as 10 known ones (4–13), were evaluated from the flowers of *Pieris formosa*, which is used as an insecticide in rural areas of China. Their structures were elucidated on the basis of extensive 1D and 2D NMR spectroscopic data analyses. Significant antifeedant activity of 1, 3 and 10 against the beet armyworm (*Spodoptera exigua*) was found, indicating that these diterpenoids might also be involved in the plant defense against insect herbivores.

Keywords: *Pieris formosa*; Ericaceae; grayanane diterpenoids; antifeedant activity

1. Introduction

Grayanoids represent a special type of diterpenoids, which have been limited to the plants of Ericaceae, such as the genera *Pieris*, *Rhododendron*, *Kalmia*, *Crabodendron* and *Leucothoe* [1,2]. The grayanoid diterpenoids, as the characteristic secondary metabolites of the plants of Ericaceae, have attracted much attention from chemists and biologists not only for their intriguing structure but also for their diverse bioactivity, especially for their toxicity, as well as their analgesic, significant antifeedant, and insecticidal activity [3–5]. To date, 15 types of diterpenoid skeleton have been reported, including grayanane (A-nor-B-homo *ent*-kaurane) [5], 1,5-secograyanane [6], 3,4-secograyanane [7], 9,10-secograyanane [8], 1,10:2,3-disecograyanane [9], leucothane (A-homo-B-nor grayanane) [10], kalmane (B-homo-C-nor grayanane) [11], 1,5-seckalmane [12], micranthane (C-homo grayanane) [13], mollane (C-nor-D-homo grayanane) [14], rhodomollane (D-homo grayanane) [15], *ent*-kaurane, 4,5-*sec*-ent-kauran [16], pierisketane (A-homo-B-nor-*ent*-kaurane) [17], and rhodomollane [18]. All of these diterpenoid types are assumed to be derived biogenetically from the *ent*-kanrane skeleton. Notably, eight types of skeletons have been produced by the narrow genus *Pieris*, whose representative plant is *Pieris formosa* D. Don. (Figure 1).

*P. formosa*, an evergreen shrub, is distributed mainly in the hilly regions and valleys of southern and southwestern China. The juice of both its fresh leaves and flowers are used as an insecticide and lotion for the treatment of ring worm and scabies, in folk medicine [19–21]. More than 60 grayanane diterpenoids with grayanane, 1,5-secograyanane, 3,4-*sec*-grayanane, 9,10-*sec*-grayanane, leucothane, 4,5-*sec*-ent-kauran, and pierisketane carbon skeletons have been isolated from the leaves, flowers, fruits, stems and roots of *P. formosa* [3,17,22–25]. In our previous work, two new highly esterified 3,4-*sec*-grayanane diterpenoids, pierisoids A and B, were reported from the flowers of *P. formosa* [20]. In our continuing endeavor to identify structurally unique and biologically diverse grayanoids from famous poisonous plants, three additional new grayanoids, pierisoids C-E (1–3) (Figure 2), together
with ten known ones (4-13) (Figure 2) were isolated from the flowers of *P. formosa*. Herein, the isolation and structural elucidation of 1-3 and their antifeedant activity against the generalist plant-feeding cotton bollworm are described.

**Figure 1.** Biogenetic relationships of grayanane carbon skeletons from the genus *Pieris*.

**Figure 2.** Structures of grayanane diterpenoids 1–13 isolated from *P. formosa*.
2. Results

2.1. Structural Elucidation of Compounds 1–3

Compound 1, [α]D24.8 = +14.3 (c = 0.2, MeOH), was isolated as colorless crystals. Its molecular formula was determined to be C25H40O3 according to its high-resolution (HR) ESI-MS (found: m/z 507.2567 [M + Na]+, calcd.: 507.2570) and 1H- and 13C-NMR spectra. IR absorptions at 3550, 3484, 1752 and 1729 cm⁻¹ were indicative of hydroxyl and ester carbonyl functional groups. In the 1H-NMR spectrum (Table 1), four tertiary methyls at δH 0.91, 1.11, 1.27 and 1.41 (each 3H of singlet), one acetyl methyl at δH 2.10 (s, 3H), and one primary methyl at δH 1.11 (3H, t, J = 7.6 Hz) were clearly observed. Additionally, four singlets (δH 3.06, 3.39, 4.93 and 5.59), two triplets (δH 3.60 and 3.62, J = 5.0 Hz), and one doublet (δH 4.20, d, J = 5.4 Hz) were ascribable to either oxygenated methine or free hydroxy groups. Other signals which were mostly overlapped centered between 1.58 and 2.85 ppm, resonating from either methine or methylene signals. The 13C-NMR spectrum revealed 25 carbon resonances, which were further classified by DEPT-90 and DEPT-135 spectra as six methyls, five methylenes, seven methines including four oxygenated ones (δC 72.4, 82.1, 82.5, and 85.0), seven quaternary carbons including three oxygenated ones (δC 78.4, 79.6, and 84.1), and two carbonyl carbons (δC 170.3 and 173.4). By analysis of the HSQC (H-NMR) and the HMBC spectra, all proton signals, except for the two singlets at δH 3.06 and 3.39 and the doublet at δH 4.20, could be assigned unambiguously to their respective carbons, suggesting that the signals at δH 3.06, 3.39 and 4.20 were assignable to free hydroxy groups. Moreover, the existence of a propionyloxy-group was determined from analysis of the 1H-13C COSY and HMBC spectra. The above spectroscopic evidence suggested a highly oxygenated grayanane diterpenoid for 1.

Detailed analysis of the 1D (1H and 13C) and 2D (1H-1H COSY, HSQC, and HMBC) NMR spectra (Figures S1–S6 in supplementary data) of 1 revealed that its structure closely resembled that of asebotoxin VIII [19,26], a known grayanane diterpenoid previously isolated from both P. japonica and P. formosa. The obvious difference between the two compounds was that the acetoxy group located at C-6 in asebotoxin VIII migrated to C-15 in 1, as indicated by the HMBC correlations from H-15 (δH 4.93, s) to the acetoxy carbonyl group at δC 170.3 and from H2-6 (δH 2.05, m) to C-5 (δC 84.1) and C-7 (δC 72.4) (Figure 3a). In the ROESY spectrum, the correlations of H-15 with Me-18 as well as Me-20 suggested that H-15, Me-18, and Me-20 were all in the same β-orientation (Figure 3d). In addition, the correlations of 3-OH with Me-18, and of 15-H with 7-H and Me-17 suggested that 3-OH, H-15, and Me-17 were in the same β-orientation (Figure 3d). Further analysis of the ROESY spectrum indicated the configurations of the remaining functional groups in 1 were the same as those in asebotoxin VIII, namely 3β, 5β, 7α, 10α, 16β-pentahydroxy, and 14β-propionyloxy. Accordingly, the structure of 1 was deduced as shown in Figure 2, and was named pierisoid C.

Compound 2 was obtained as colorless oil with a molecular formula of C23H36O8, as determined by a combination of HR-ESI-MS and NMR spectra (including 1H, 13C, and DEPT) (Figures S7–S2 in supporting information). The resemblance of the NMR spectra of 2 (Table 1) with those of 1 disclosed that 2 was another grayanane diterpenoid structurally similar to 1. The major difference was the replacement of a methylene carbon in 1 by an oxygen-occurring methine in 2 (δC 78.2), suggesting that either C-6, or C-11, or C-12 of 2 was oxygenated. In the HMBC spectrum of 2, the HMBC correlations from 5-OH to the methine carbon at δC 78.2 indicated that this methine was ascribable to C-6 (Figure 3b). Carefully comparison of 13C-NMR spectral data of 2 with those of 1 (Table 1) obviously found that the upfield-shift of C-15 (δC 68.6) and C-16 (δC 61.3) in 2, indicated an oxygen bridge, was formed between C-15 and C-16; this was supported by the HR-ESI-MS spectrum. In the ROESY spectrum of 2, the correlations of Me-17 with H-15; of 3-OH and 5-OH with Me-18; and of 5-OH with 6-OH and H-7 indicated that 3-OH, 5-OH, 6-OH, H-7, H-15, and Me-17 were in the same β-orientation (Figure 3e). Consequently, the structure of 2 was determined as shown in Figure 2 and was named pierisoid D.
Table 1. $^1$H- (400 MHz) and $^{13}$C- (100 MHz) NMR spectroscopic data of compounds 1–3 in acetone-$d_6$ ($\delta$ (ppm), $J$ (Hz)).

| No. | $^1$H | $^{13}$C | $^1$H | $^{13}$C | $^1$H | $^{13}$C |
|-----|-------|---------|-------|---------|-------|---------|
|     | $\delta$ (ppm) | $\delta$ (ppm) | $\delta$ (ppm) | $\delta$ (ppm) | $\delta$ (ppm) | $\delta$ (ppm) |
| 1   | 2.74 dd (11.6, 5.2) | 50.9 d | 2.77 m | 50.9 d | 3.08 m | 52.6 d |
| 2a  | 2.06 m | 35.4 t | 2.05 m | 35.9 t | 2.47 m | 32.8 t |
| 2b  | 2.27 m | 2.22 m | 2.73 dd (18.9, 11.8) | 35.4 t | 2.05 m | 2.22 m | 35.9 t | 2.47 m | 32.8 t |
| 3   | 3.60 t (5.0) | 82.5 d | 3.53 dd (1.5, 5.1) | 83.0 d | 83.0 d | 174.0 s |
| 4   | 51.4 s | 52.3 s | 62.5 s |
| 5   | 84.1 s | 83.1 s | 87.5 s |
| 6   | 2.05 m (2H) | 34.0 t | 3.91 overlap | 78.2 d | 6.04 d (9.6) | 69.8 d |
| 7   | 3.62 t (5.0) | 72.4 d | 3.41 overlap | 78.3 d | 5.23 d (9.7) | 68.7 d |
| 8   | 54.5 s | 53.8 s | 56.1 s |
| 9   | 2.08 m | 54.6 d | 2.08 m | 49.9 d | 2.32 m | 47.0 d |
| 10  | 78.4 s | 77.7 s | 77.4 s |
| 11a | 1.58 m | 21.8 t | 1.71 m | 21.6 t | 1.98 m | 2.03 m |
| 11b | 1.90 m | 1.81 m | 21.1 t | 21.1 t |
| 12a | 1.58 m | 27.0 t | 1.68 m | 28.1 t | 1.74 m | 2.08 m |
| 12b | 2.23 m | 2.35 m | 1.58 t | 25.8 t |
| 13  | 2.05 overlap | 52.9 d | 2.24 m | 47.6 d | 3.10 overlap | 46.4 d |
| 14  | 5.59 s | 82.1 d | 5.52 s | 76.4 d | 6.43 s | 79.3 d |
| 15  | 4.93 s | 85.0 d | 3.26 s | 68.6 d | 5.27 s | 86.2 d |
| 16  | 79.6 s | 61.3 s | 88.4 s |
| 17  | 1.27 s (3H) | 22.8 q | 1.45 s (3H) | 14.6 q | 1.55 s (3H) | 19.4 q |
| 18  | 1.11 s (3H) | 18.5 q | 1.23 s (3H) | 19.3 q | 2.49 d (5.0) | 52.8 t |
| 19  | 0.91 s (3H) | 23.0 q | 0.97 s (3H) | 23.2 q | 1.34 s (3H) | 17.4 q |
| 20  | 1.41 s (3H) | 27.8 q | 1.41 s (3H) | 28.2 q | 1.51 s (3H) | 33.6 q |
| 6-OAc | 2.02 s (3H) | 20.6 q | 0.20 s (3H) | 169.4 s |
| 7-OAc | 2.10 s (3H) | 21.6 q | 169.8 s |
| 14-OPr | 1.11 t (3H, 7.6) | 28.4 t | 2.08 t (3H, 7.6) | 28.5 t | 2.36 m (2H) | 9.1 q |
| 15-OAc | 1.98 s (3H) | 20.8 q | 171.6 s |
| 16-OAc | 1.98 s (3H) | 22.7 q | 169.8 s |

* Hydroxyl groups of 1: $\delta$H 4.20 (d, $J$ = 5.4 Hz, 3-OH), 3.39 (s, 10-OH), 3.06 (s, 16-OH).<br>B Hydroxyl groups of 2: $\delta$H 3.91 (overlap, 3-OH), 3.64 (s, 5-OH), 3.51 (s, 10-OH), 3.41 (overlap, 6-OH), 3.34 (d, $J$ = 7.8 Hz, 7-OH).

Compound 3, colorless crystals, has a molecular formula of C$_{31}$H$_{42}$O$_{14}$, as determined by a combination of HR-EI-MS and NMR spectra (including $^1$H-, $^{13}$C-, and DEPT) (Figures S13–S18 in supplementary data). Its spectroscopic data were very similar to those of secorhodomollolide B, a 3,4-secograyanane diterpenoid also isolated from *P. formosa* [27]. The only difference between them was that the terminal double bond between C-4 ($\delta$C 146.0) and C-18 ($\delta$C 116.7) in secorhodomollolide B was replaced by a 4,18-oxirane group ($\delta$C 62.5 and 52.8) in 3, which was confirmed by the HMBC correlations from Me-19 ($\delta$H 1.34, s) to C-4, C-5 ($\delta$C 87.5) and C-18 (Figure 3c). Such an oxirane moiety has also been found in pierisoid A, another 3,4-secograyanane diterpenoid we reported from the flowers of *P. formosa* [20]. In the ROESY spectrum of 3, the correlations of Me-19 with H-1, and of
Me-20 with H-7 indicated that Me-19 and H-1 were in α-orientation and Me-20 coupled with H-7 were in β-orientation (Figure 3f). Therefore, compound 3 was identified as shown in Figure 2 and was named pierisoid E.

Ten known diterpenoids (Figure 2), namely, pierisformotoxin C (4) [24], secorhodomollolides C (5), D (6), and F (7) [28], asebotoxins I (8), II (12), IV (10), and VIII (11) [26,29], pieristoxin I (9) [29], and pierisformosin (13) [30] were also isolated from P. formosa and were identified by comparison of their spectroscopic data with those reported in the literature.

Table 2. Antifeedant activity of compounds 1, 3, 4, and 10 against Spodoptera exigua.

| Compound | Molecular Formula | m/z | EC50 (µg/cm²) |
|----------|------------------|-----|--------------|
| 1        | C25H40O9         | 484 | 10.91        |
| 3        | C31H42O14        | 638 | 33.89        |
| 4        | C33H38O8         | 682 | NA           |
| 10       | C23H38O8         | 442 | 6.58         |
| Neem oil | -                | -   | 3.71         |

NA = Not active; “-” = No (Because neem oil is a mixture).

3. Discussion

Compounds 1–3 are highly oxygenated grayanane diterpenoids, which occur extensively in the plants of Ericaceae and exhibit remarkable biological activities, such as antifeedant and insecticidal...
activities. Compound 2 is a grayanane-type diterpenoid with an unprecedented 15,16-epoxy group in the grayanoids family. Compound 3, a 3,4-seco-grayanane diterpenoid, possesses a 4,18-epoxy substituent, which is also unusual in nature. Compared to compounds 1 and 3, 10 displayed more significant antifeedant activity against S. exigua, which was possibly attributed to the integrity of the A ring.

4. Materials and Methods

4.1. General Experimental Procedures

Melting points were recorded on an Aisey YLD-6000 instrument and are uncorrected. Column chromatography was performed on 200–300 mesh silica gel (Qingdao Marine Chemical Factory, Qingdao, China). Optical rotations were measured on a Horiba-SEAP-300 spectropolarimeter (Horiba, Tokyo, Japan). UV spectral data were obtained on a Shimadzu-210A double-beam spectrophotometer (Shimadzu, Tokyo, Japan). IR spectra were recorded on a Bruker-Tensor-27 spectrometer with KBr pellets (Bruker Optics, Ettlingen, Germany). NMR experiments were carried out on either a Bruker AV-400 or a DRX-500 spectrometer with tetramethyl silane (TMS) as an internal standard (Bruker, Karlsruhe, Germany). MS were recorded on a VG-Auto-Spec-3000 spectrometer (Waters Corp., Milford, MA, USA). TLC (Thin Layer Chromatography) spots were visualized under UV light, by dipping into 10% H$_2$SO$_4$ in EtOH followed by heating. All solvents including petroleum ether (60–90 °C) were distilled before use.

4.2. Plant Materials

The flowers of P. formosa were collected at Qu Jing, Yunnan province, China, in March 2008. The plant material was identified by Dr. Sheng-Hong Li.

4.3. Extraction and Isolation

Air-dried P. formosa flowers (3.5 kg) were powdered and extracted with MeOH (3 × 5 L) at room temperature. The extract was concentrated under reduced pressure and then partitioned between H$_2$O and EtOAc (1:1). The organic layer (EtOAc part) was concentrated and the residue (594 g) was purified by silica gel column chromatography with solvent mixtures of CHCl$_3$−Me$_2$CO (1:0, 9:1, 8:2, 7:3, 1:1 and 0:1) to afford six fractions. Fraction 2 (45.0 g, CHCl$_3$−Me$_2$CO, 9:1) was repeatedly chromatographed on silica gel (CHCl$_3$−Me$_2$CO, 10:1; petroleum ether−Me$_2$CO, 4:1), yielding subfractions A1−A3. Subfraction A1 (7.9 g) was further purified by silica gel column chromatography using petroleum ether-isopropanol (20:1), followed by purification on Sephadex LH-20 columns (CHCl$_3$−MeOH, 1:1) to obtain compounds 4 (6 mg), 5 (5 mg), and 7 (81 mg). Subfraction A2 (6.65 g) was repeatedly separated by silica gel column chromatography eluting with petroleum ether-acetone (6:1) and finally was purified by Sephadex LH-20 columns (CHCl$_3$−MeOH, 1:1; Me$_2$CO) to afford compounds 1 (27 mg), 3 (3 mg), and 6 (6 mg). Subfraction A3 (10.2 g) was separated by silica gel column chromatography eluting with petroleum ether−ethyl acetate (5:1) and petroleum ether−isopropanol (12:1), respectively, and then similarly purified by Sephadex LH-20 columns (CHCl$_3$−MeOH, 1:1; Me$_2$CO) to give 8 (78 mg) and 9 (7 mg). Fraction 3 (4.2 g, CHCl$_3$−Me$_2$CO, 8:2) was separated on a MCI (Middle Chromatogram Isolated) gel column employing solvent mixtures of MeOH−water (6:4, 7:3, 8:2, 9:1, and 1:0), and the resulting subfractions 2 and 3 (7:3 and 8:2 MeOH−water, respectively) were repeatedly chromatographed on silica gel (CHCl$_3$−EtOAc, 4:1; petroleum ether−isopropanol, 10:1) and Sephadex LH-20 (MeOH; Me$_2$CO) columns to yield compounds 2 (18 mg), 10 (45 mg), 11 (8 mg), 12 (9 mg) and 13 (6 mg).

Pierisoid C (1): Colorless crystals; mp 220–222 °C; [α]$_\text{D}$ = +14.3 (c = 0.2, MeOH); UV (MeOH) λ$_\text{max}$ (log ε): 209 (1.98), 193 (1.60) nm; IR (KBr) ν$_\text{max}$: 3550, 3484, 2976, 2942, 1752, 1729, 1408, 1373, 1261, 1153, 1047 cm$^{-1}$; ESI-MS: m/z 507 (50) [M + Na]$^+$; HR-ESI-MS: m/z 507.2567 [M + Na]$^+$ (calcd. for C$_{25}$H$_{40}$O$_9$Na, 507.2570); $^1$H- and $^{13}$C-NMR data: see Table 1.
**Pierisoid D** (2): Colorless oils; \([\alpha]_{D}^{23.8} = +16.7 (c = 0.1, \text{MeOH})\); UV (MeOH) \(\lambda_{\text{max}}\) (log \(\varepsilon\)): 202 (2.48) nm; IR (KBr) \(\nu_{\text{max}}\): 3441, 3432, 2971, 2939, 1735, 1730, 1630, 1382, 1179, 1025 cm\(^{-1}\); ESI-MS: \(m/z\) 463 (88) [M + Na]\(^{+}\); HR-EI-MS: \(m/z\) 440.2420 (calcd. for C\(_{23}\)H\(_{36}\)O\(_8\), 430.2410); \(^{1}\)H- and \(^{13}\)C-NMR data: see Table 1.

**Pierisoid E** (3): Colorless crystals; mp 271–272 °C; \([\alpha]_{D}^{23.6} = +12.6 (c = 0.1, \text{MeOH})\); UV (MeOH) \(\lambda_{\text{max}}\) (log \(\varepsilon\)): 202 (2.29) nm; IR (KBr) \(\nu_{\text{max}}\): 3441, 2942, 1787, 1734, 1631, 1372, 1264, 1237, 1053 cm\(^{-1}\); ESI-MS: \(m/z\) 661 (68) [M + Na]\(^{+}\); HR-EI-MS: \(m/z\) 638.2560 (calcd. for C\(_{31}\)H\(_{42}\)O\(_{14}\), 638.2575); \(^{1}\)H- and \(^{13}\)C-NMR data: see Table 1.

### 4.4. Antifeedant Activity

Beet armyworms (*Spodoptera exigua*) were purchased from the Pilot-Scale Base of Bio-Pesticides, Institute of Zoology, Chinese Academy of Sciences. A modified dual-choice bioassay was performed for an antifeedant test as previously described [20,31–33]. The larvae were reared on an artificial diet under a controlled photoperiod (light:dark, 12:8 h) and temperature (25 ± 2 ºC). The larvae were starved for 3–4 h before each bioassay. Fresh leaf disks were cut from *Brassica chinensis*, using a cork borer (1.1 cm in diameter). The treated leaf disks were painted with 10 µL of the test compound in acetone, and control leaf disks were treated with the same amount of acetone. After air drying, the tested and control leaf disks were set in alternating position in the same Petri dish (90 mm in diameter), with moistened filter paper at the bottom. Two-thirds of the instars were placed at the center of the Petri dish. Five replicates were run for each treatment. After feeding for 24 h, the areas of leaf disks consumed were measured. The antifeedant index (AFI) was calculated according to the following formula: AFI = \([\frac{C - T}{C + T}] \times 100\), where \(C\) and \(T\) represent the control and treated leaf areas consumed by the insect. The insect antifeedant potency of the test compound was evaluated in terms of the EC\(_{50}\) value, which was determined by probit analysis for each insect species.

### 5. Conclusions

Terpenoids play an important role in natural product chemistry and biology, such as antifungal and insecticidal activities [34]. Secondary metabolites, such as the grayanane diterpenoids that occur extensively in the plants of Ericaceae, are fascinating for their remarkable toxicity, as well as their significant antifeedant and insecticidal activity. In the current study, three new grayanane diterpenoids, pierisoids C–E (1-3), as well as 10 known ones (4-13), were identified from the flowers of *P. formosa* via their extensive 1D and 2D NMR spectroscopic data analyses. Notably, compounds 1, 3 and 10, especially 10, exhibited obvious antifeedant activity against the beet armyworm (*S. exigua*), suggesting that these diterpenoids were important defensive substances in *P. formosa* against natural enemies.

### Supplementary Materials

Supplementary Materials are available online. 1D (\(^{1}\)H- and \(^{13}\)C-) and 2D NMR spectra of compounds 1–3 are available online.

### Acknowledgments

This research was supported financially by the National Natural Science Foundation of China (Grant No. 31500275) and the Scientific Research Foundation of Northwest A & F Foundation (2452017175).

### Author Contributions

Designed the research: Sheng-Hong Li. Performed the experiments: Chun-Huan Li. Wrote the manuscript: Chun-Huan Li. Analyzed the data: Chun-Huan Li and Shi-Hong Luo. Evaluated the data and edited the manuscript: Sheng-Hong Li and Jin-Ming Gao. Contributed to discussion: Chun-Huan Li and Jin-Ming Gao.

### Conflicts of Interest

The authors declare no conflict of interest.

### References

1. Cao, L.; Li, Y.; Li, H.; Liu, D.; Li, R. New 3,4-seco-Grayanane Diterpenoids from the Flowers of *Pieris japonica*. *Chem. Pharm. Bull.* 2016, 64, 1222–1225. [CrossRef] [PubMed]

2. Li, Y.; Liu, Y.B.; Yu, S.S. Grayanoids from the Ericaceae family: Structures, biological activities and mechanism of action. *Phytochem. Rev.* 2013, 12, 305–325. [CrossRef]
3. Niu, C.-S.; Li, Y.; Liu, Y.-B.; Ma, S.-G.; Li, L.; Qu, J.; Yu, S.-S. Analgesic diterpenoids from the twigs of *Pieris formosa*. Tetrahedron 2016, 72, 44–49. [CrossRef]  
4. Chen, X.; Gao, L.; Li, Y.; Li, H.; Liu, D.; Liao, X.; Li, R. Highly Oxygenated Grayanane Diterpenoids from Flowers of *Pieris japonica* and Their Structure-Activity Relationship of Antifeedant Activity Against *Pieris brassicae*. *J. Agric. Food Chem.* 2017, 65, 4456–4463. [CrossRef] [PubMed]  
5. Im, N.-K.; Zhou, W.; Na, M.; Jeong, G.-S. Pierisformoside B exhibits neuroprotective and anti-inflammatory effects in murine hippocampal and microglial cells via the HO-1/Nrf2-mediated pathway. *Int. Immunopharmacol.* 2015, 24, 353–360. [CrossRef] [PubMed]  
6. Wang, W.G.; Li, H.M.; Li, H.Z.; Wu, Z.Y.; Li, R.T. New grayanol diterpenoid and new phenolic glucoside from the flowers of *Pieris formosa*. *J. Asian Nat. Prod. Res.* 2010, 12, 70–75. [CrossRef] [PubMed]  
7. Wang, S.J.; Lin, S.; Zhu, C.G.; Yang, Y.C.; Li, S.A.; Zhang, J.J.; Chen, X. G.; Shi, J. G. Highly Acylated Diterpenoids with a New 3,4-Secograyanane Skeleton from the Flower Buds of Rhododendron molle. *Org. Lett.* 2010, 12, 1560–1563. [CrossRef] [PubMed]  
8. Wu, Z.Y.; Li, H.Z.; Wang, W.G.; Li, H.M.; Chen, R.; Li, R.T.; Luo, H.R. Lyonin A, a new 9,10-Secograyanotoxin from *Lyonia ovalifolia*. *Chem. Biodivers.* 2011, 8, 1182–1187. [CrossRef] [PubMed]  
9. Li, Y.; Liu, Y.-B.; Zhang, J.-J.; Li, Y.-H.; Jiang, J.-D.; Yu, S.-S.; Ma, S.-G.; Qu, J.; Lv, H.-N. Mollolide A, a Diterpenoid with a new 1, 10,2,3-diecograyanaketone skeleton from the roots of *Rhododendron molle*. *Org. Lett.* 2013, 15, 3074–3077. [CrossRef] [PubMed]  
10. Wang, L.Q.; Chen, S.-N.; Cheng, K.-F.; Li, C.-J.; Qin, G.-W. Diterpene glucosides from *Pieris formosa*. *Phytochemistry* 2000, 54, 847–852. [CrossRef]  
11. Burke, J.W.; Doskotch, R.W.; Ni, C.Z.; Clardy, J. Kalmanol, a pharmacologically active diterpenoid with a new ring skeleton from *Kalma angustifolia*. *J. Am. Chem. Soc.* 1989, 111, 5831–5833. [CrossRef]  
12. Zhou, S.-Z.; Yao, S.; Tang, C.; Ke, C.; Li, L.; Lin, G.; Ye, Y. Diterpenoids from the flowers of *Rhododendron molle*. *J. Nat. Prod.* 2014, 77, 1185–1192. [CrossRef] [PubMed]  
13. Zhang, M.; Zhu, Y.; Zhan, G.; Shu, P.; Sa, R.; Lei, L.; Xiang, M.; Xue, Y.; Luo, Z.; Wan, Q. Micranthanone A, a new diterpenoid with an unprecedented carbon skeleton from *Rhododendron molle*. *Org. Lett.* 2013, 15, 3094–3097. [CrossRef] [PubMed]  
14. Li, Y.; Liu, Y.-B.; Liu, Y.-L.; Wang, C.; Wu, L.-Q.; Li, L.; Ma, S.-G.; Qu, J.; Yu, S.-S. Mollanol A, a diterpenoid with a new C-nor-D-homograyanane skeleton from the roots of *Rhododendron molle*. *J. Nat. Prod.* 2013, 76, 4320–4323. [CrossRef] [PubMed]  
15. Li, Y.; Liu, Y.-B.; Yan, H.-M.; Liu, Y.-L.; Li, Y.-H.; Lv, H.-N.; Ma, S.-G.; Qu, J.; Yu, S.-S. Rhodomollins A and B, two Diterpenoids with an Unprecedented Backbone from the Fruits of *Rhododendron molle*. *Sci. Rep.* 2016, 6, 1–6. [CrossRef] [PubMed]  
16. Wang, L.-Q.; Qin, G.-W.; Chen, S.-N.; Li, C.-J. Three diterpene glucosides and a diphenylamine derivative from *Pieris formosa*. *Fitoterapia* 2001, 72, 779–787. [CrossRef]  
17. Niu, C.-S.; Li, Y.; Liu, Y.-B.; Ma, S.-G.; Liu, F.; Li, L.; Xu, S.; Wang, X.-J.; Wang, R.-B.; Qu, J. Pierisketolide A and Pierisketones B and C, Three Diterpenes with an Unusual Carbon Skeleton from the Roots of *Pieris formosa*. *Org. Lett.* 2017, 19, 906–909. [CrossRef] [PubMed]  
18. Zhou, J.; Zhan, G.; Zhang, H.; Zhang, Q.; Li, Y.; Xue, Y.; Yao, G. Rhodomollanol A, a Highly Oxygenated Diterpenoid with a 5/7/5/5 Tetracyclic Carbon Skeleton from the Leaves of *Rhododendron molle*. *Org. Lett.* 2017, 19, 3935–3938. [CrossRef]  
19. Wang, L.Q.; Chen, S.N.; Qin, G.W.; Cheng, K.F. Grayanane diterpenoids from *Pieris formosa*. *J. Nat. Prod.* 1998, 61, 1473–1475. [CrossRef] [PubMed]  
20. Li, C.-H.; Niu, X.-M.; Luo, Q.; Xie, M.-J.; Luo, S.-H.; Zhou, Y.-Y.; Li, S.-H. Novel polyesterified 3,4-seco-grayanane diterpenoids as antifeedants from *Pieris formosa*. *Org. Lett.* 2010, 12, 2426–2429. [CrossRef] [PubMed]  
21. Wang, L.Q.; Ding, B.Y.; Zhao, W.M.; Qin, G.W. Grayanane diterpenoids from *Pieris formosa*. *Chin. Chem. Lett.* 1998, 9, 465–467. [CrossRef]  
22. Wang, L.-Q.; Ding, B.-Y.; Qin, G.-W.; Lin, G.; Cheng, K.-F. Grayanoids from *Pieris formosa*. *Phytochemistry* 1998, 49, 2045–2048. [CrossRef]  
23. Wu, Z.Y.; Li, H.Z.; Wang, W.G.; Li, X.L.; Li, R.T. Secopieristoxins A and B, two unusual diterpenoids with a new 9,10-secograyanane skeleton from the fruits of *Pieris formosa*. *Phytochem. Lett.* 2012, 5, 87–90. [CrossRef]
24. Wang, W.G.; Wu, Z.Y.; Chen, R.; Li, H.Z.; Li, H.M.; Li, Y.D.; Li, R.T.; Luo, H.R. Pierisformotoxins A–D, Polyesterified Grayanane Diterpenoids from Pieris formosa and Their cAMP-Decreasing Activities. Chem. Biodivers. 2013, 10, 1061–1071. [CrossRef] [PubMed]

25. Wu, Z.Y.; Li, Y.D.; Wu, G.S.; Luo, H.R.; Li, H.M.; Li, R.T. Three New Highly Acylated 3,4-seco-Grayanane Diterpenoids from the Fruits of Pieris formosa. Chem. Pharm. Bull. 2011, 59, 492–495. [CrossRef] [PubMed]

26. Hikino, H.; Ogura, M.; Fushiya, S.; Konno, C.; Takemoto, T. Stereostructure of asebotoxin VI, VIII, and IX, toxins of Pieris japonica. Chem. Pharm. Bull. 1977, 25, 523–524. [CrossRef]

27. Wang, S.; Lin, S.; Zhu, C.; Yang, Y.; Li, S.; Zhang, J.; Chen, X.; Shi, J. Highly acylated diterpenoids with a new 3,4-secograyanane skeleton from the flower buds of Rhododendron molle. Org. Lett. 2010, 12, 1560–1563. [CrossRef] [PubMed]

28. Wu, Z.Y.; Li, H.M.; Li, Y.D.; Li, H.Z.; Li, R.T. Highly Acylated 3,4-Secograyanane Diterpenoids from the Fruits of Pieris formosa. Helv. Chim. Acta 2011, 94, 1283–1289. [CrossRef]

29. Katai, M.; Fujiwara, M.; Terai, T.; Meguri, H. Studies on the Constituents of the Leaves of Pieris japonica D. DON. Chem. Pharm. Bull. 1980, 28, 3124–3126. [CrossRef]

30. Zhou, C.; Li, X.; Li, H.; Li, R. Chemical constituents from the leaves of Craibiodendron yunnanens. Biochem. Syst. Ecol. 2012, 45, 179–182. [CrossRef]

31. Li, C.H.; Jing, S.X.; Luo, S.H.; Shi, W.; Hua, J.; Liu, Y.; Li, X.N.; Schneider, B.; Gershenzon, J.; Li, S.H. Peltate Glandular Trichomes of Colquhounia coccinea var. mollis Harbor a New Class of Defensive Sesterterpenoids. Org. Lett. 2013, 15, 1694–1697. [CrossRef] [PubMed]

32. Li, C.H.; Liu, Y.; Hua, J.; Luo, S.H.; Li, S.H. Peltate glandular trichomes of Colquhounia seguini harbor new defensive clerodane diterpenoids. J. Integr. Plant Biol. 2014, 56, 928–940. [CrossRef] [PubMed]

33. Luo, S.H.; Luo, Q.A.; Niu, X.M.; Xie, M.J.; Zhao, X.; Schneider, B.; Gershenzon, J.; Li, S.H. Glandular Trichomes of Leucoceptrum canum Harbor Defensive Sesterterpenoids. Angew. Chem. Int. Edit. 2010, 49, 4471–4475. [CrossRef] [PubMed]

34. Wang, D.-M.; Zhang, C.-C.; Zhang, Q.; Shafiq, N.; Pescitelli, G.; Li, D.-W.; Gao, J.-M. Wightianines A-E, Dihydro-beta-agarofuran Sesquiterpenes from Parnassia wightiana, and Their Antifungal and Insecticidal Activities. J. Agric. Food Chem. 2014, 62, 6669–6676. [CrossRef] [PubMed]

Sample Availability: Samples of the compounds 1–13 are available from the authors.