Semisynthesis and Antifeedant Activity of New Derivatives of a Dihydro-β-Agarofuran from *Parnassia wightiana*

**Jiang-Jiang Tang** ¹,†, **Fei-Yu Zhang** ¹,†, **Dong-Mei Wang** ², **Jun-Mian Tian** ¹, **Shuai Dong** ¹ and **Jin-Ming Gao** ¹,*

¹ Shaanxi Engineering Center of Bioresource Chemistry & Sustainable Utilization, College of Science, Northwest A&F University, Yangling 712100, Shaanxi, China; E-Mails: tangjiang11@nwsuaf.edu.cn (J.-J.T.); feiyuzhang2013@outlook.com (F.-Y.Z.); tianjunmian@nwsuaf.edu.cn (J.-M.T.); phd@nwsuaf.edu.cn (S.D.)

² College of Forestry, Northwest A&F University, Yangling 712100, Shaanxi, China; E-Mail: dmwli@nwsuaf.edu.cn

† These authors contributed equally to this work.

* Author to whom correspondence should be addressed; E-Mail: jinminggao@nwsuaf.edu.cn; Tel.: +86-29-8709-2515; Fax: +86-29-8709-2226.

Received: 1 July 2013; in revised form: 22 August 2013 / Accepted: 10 September 2013 / Published: 26 September 2013

**Abstract:** Five new derivatives (2–6) were semi-synthesized using compound 1, a dihydro-β-agarofuran sesquiterpene with C-2 ketone obtained from *Parnassia wightiana*, as the starting material by acylation, oxidation, reduction, esterification, and amination, respectively. Structures of 2–6 were confirmed by 1D- and 2D-NMR and HR-ESI-MS spectra. In addition, antifeedant activities of these compounds (1–6) were tested against the 3rd-instar larvae of *Mythimna separata*. Antifeedant effects of compounds 2 and 4 were greater than the parent compound 1 whereas other compounds exhibited low to no feeding deterrent effects against third instar *M. separata* larvae in lab bioassays. Therefore, our results suggest that acylated and reduced derivatives at C-8 and C-2, respectively, of 1 may improve the antifeeding effect. This preliminary information will be useful in designing new insect control agents against *M. separata* and other important pests.

**Keywords:** sesquiterpene; agarofuran; derivatives; antifeedant activity
1. Introduction

The development of resistance to existing classes of pesticide and the increasing environmental pollution and toxicity generate a continuing need for the development of more active new classes of pest control agents against the target pests. During the last three decades, both synthetic and botanical pesticides from a variety of sources have been identified and developed, with most of the plant defensive chemicals discouraging insect herbivory, either by deterring feeding or by impairing larval growth, rather than by killing insects outright [1,2]. One of these pesticides structure is sesquiterpene polyesters with a dihydro-β-agarofuran skeleton such as Celangulin-V, which are the most widespread and characteristic group of secondary metabolites isolated mainly from Celastraceae family [3]. These compounds have attracted considerable attention from synthetic organic chemists and pharmacologists due to their complex structures and wide range of biological activities [4,5] including antifeedant activity, insecticidal activity, cytotoxicity [6,7], multidrug resistance (MDR) reversal activity [8–11], HIV inhibition [12] and antitumor activity [13]. Recently, Zhang and co-workers synthesized a series of ester [14], ether [15] and dimmer [16] derivatives using Celangulin-V as a lead compound, and found some potent insecticidal compounds.

Parnassia wightiana Wall. (Family: Saxifragaceae) is a widely occurring perennial rhizomatous herb in East Asia, North American and Europe [17]. This plant was characterized by flavonoids, alkaloids and sesquiterpenoids. In our previous report [18], a new dihydro-β-agarofuran sesquiterpene with C-2 ketone (compound 1, Figure 1) was isolated from this plant, showing significant cytotoxic activity against HepG2 (hepatocellular carcinoma) and MDA-10 (breast carcinoma) cells. To learn about the antifeedant activity of the natural sesquiterpene 1 and bioactivity changes resulting from altering the parent compound, 1 was used as the starting material and was transformed into derivatives 2–6 by acylation, oxidation, reduction, esterification and amination at C-2 or C-8 position of 1. Herein, we describe semisynthesis and antifeedant activity of these new derivatives.

Figure 1. Structure of the natural sesquiterpene 1 from P. wightiana.

2. Results and Discussion

The synthetic pathways of derivatives 2–6 are shown in Scheme 1. All the structures of these derivatives were established on the basis of 1D- and 2D-NMR and HR-ESI-MS spectroscopic analysis (Figures S1–S19). NMR spectra are shown in Tables 1 and 2.
Scheme 1. Semisynthesis of derivatives 2–6 from 1. 4-dimethylaminopyridine (DMAP), 1,1,1-tris(acetoxy)-1,1-dihydro-1,2-benziodoxol-3-(1H)-one (DMP), dichloromethane (DCM).

![Scheme 1](image)

Table 1. $^1$H NMR spectroscopic data of compound 2–6 (500 MHz, CDCl$_3$) $^a$.

| Position | 2  | 3  | 4  | 5  | 6  |
|----------|----|----|----|----|----|
| H-1      | 5.95 (s) | 5.96 (s) | 5.56 (d, 3.0) | 5.95 (s) | 5.95 (s) |
| H-2      | 2.09 (d, 3.0) | 3.40 (dd, 12.5, 7.5) | 3.73 (m) | 3.40 (dd, 12.8, 7.5) | 3.40 (dd, 12.5, 7.5) |
| H-3α     | 3.39 (dd, 12.5, 7.5) | 3.40 (dd, 13, 7.5) | 2.55 (m) | 2.32 (d, 13) | 2.32 (d, 13) |
| H-3β     | 2.30 (dd, 12.5, 1.5) | 2.37 (d, 13) | 2.67 (m) | 3.05 (m) | 3.03 (m) |
| H-4      | 3.02 (m) | 3.09 (m) | 2.67 (m) | 3.05 (m) | 3.03 (m) |
| H-6      | 6.04 (s) | 5.89 (1H, s) | 6.30 (s) | 5.99 (s) | 5.99 (s) |
| H-7      | 2.76 (d, 3.3) | 3.24 (1H, s) | 2.65 (d, 3.5) | 2.84 (d, 4.0) | 2.72 (d, 3.5) |
| H-8      | 5.39 (d, 3.3) | 4.99 (d, 3.3) | 5.17 (d, 3.3) | 5.08 (d, 3.3) | 5.08 (d, 3.3) |
| H-9      | 5.09 (s) | 5.49 (s) | 4.47 (s) | 5.52 (s) | 5.43 (s) |
| H-12     | 1.05 (d, 7.5) | 1.05 (d, 7.5) | 1.36 (d, 8.0) | 1.06 (d, 7.0) | 1.05 (d, 6.5) |
| H-13     | 1.45 (s) | 1.46 (3H, s) | 1.45 (s) | 1.45 (s) | 1.49 (s) |
| H-14     | 1.57 (s) | 1.54 (3H, s) | 1.56 (s) | 1.57 (s) | 1.56 (s) |
| H-15     | 2.29 (s) | 1.62 (3H, s) | 1.62 (s) | 1.61 (s) | 1.62 (s) |
| AcO-8 (CH$_3$) | 1.61 (s) |  |  |  |  |
| RCH$_2$COO-8 $^b$ |  |  | 4.35 (s) | 3.46 (s) |  |
| N(CH$_2$CH$_2$)$_2$O |  |  | 2.78 (m) |  |  |
| N(CH$_2$CH$_2$)$_2$O |  |  | 3.57 (m) |  |  |

$^a$ Data for additional ester groups are provided in the Experimental Section. $^b$ For compound 5, $R =$ Cl; for compound 6, $R = $ N(CH$_2$CH$_2$)$_2$O. Data are based on DEPT, HSQC, and HMBC experiments.

Compound 2, obtained through acylation [19] of 1 in the presence of triethylamine (Et$_3$N) and catalytic dimethylaminopyridine (DMAP) in dry dichloromethane (DCM), showed an accurate [M + Na]$^+$ ion at $m/z$ 615.2219 in the HR-ESI-MS, corresponding to the molecular formula C$_{33}$H$_{36}$O$_{10}$. In the $^1$H-NMR spectrum of 2, the signal of H-8 for 1 at 4.45 ppm had moved to 5.39 ppm. Further assignments were made based on DEPT, HSQC, and HMBC experiments spectra. Structure, numbering and key NOESY correlations of 2 are shown in Figure 2.
Table 2. $^{13}$C NMR Spectroscopic data of compound 2–6 (500 MHz, CDCl$_3$) $^a$.

| Position | 2   | 3   | 4   | 5   | 6   |
|----------|-----|-----|-----|-----|-----|
| C-1      | 77.4| 76.7| 74.9| 77.5| 77.4|
| C-2      | 204.3| 203.5| 70.8| 204.2| 204.3|
| C-3      | 43.9| 44.0| 34.1| 43.9| 43.9|
| C-4      | 38.7| 38.2| 34.0| 38.7| 38.7|
| C-5      | 89.5| 91.2| 90.8| 89.4| 89.5|
| C-6      | 76.1| 70.8| 76.2| 76.3| 76.0|
| C-7      | 53.1| 74.0| 55.7| 52.9| 53.2|
| C-8      | 75.9| 202.2| 74.6| 75.9| 77.2|
| C-9      | 77.2| 65.35| 80.4| 76.5| 76.3|
| C-10     | 55.0| 51.9| 55.6| 55.1| 59.4|
| C-11     | 82.9| 84.1| 81.7| 82.9| 82.9|
| C-12     | 18.1| 17.8| 18.7| 18.1| 14.2|
| C-13     | 20.2| 19.9| 19.4| 20.3| 20.3|
| C-14     | 31.0| 21.9| 25.7| 25.5| 25.5|
| C-15     | 25.6| 27.3| 29.7| 31.0| 31.0|
| AcO-8 $^b$ | 21.2, 169.3 | 41.0, 172.6 | 55.0, 165.3 | 53.3 | 67.0 |

RCH$_2$OCO-8 $^b$ 41.0, 172.6 55.0, 165.3
N(CH$_2$CH$_2$)$_2$O 53.3
N(CH$_2$CH$_2$)$_2$O 67.0

Data for additional ester groups are provided in the Experimental Section. $^b$ For compound 5, $R = \text{Cl}$; for compound 6, $R = \text{N(CH}_2\text{CH}_2\text{)}_2\text{O}$. Data are based on DEPT, HSQC, and HMBC experiments.

Figure 2. NOESY experiment of compound 2.

Compound 3 was obtained through mild Dess-Martin oxidation at the 8-OH using 1,1,1-tris(acetoxy)-1,1-dihydro-1,2-benziodoxol-3-(1H)-one (DMP). Signal of the C-8 for 3 appears at $\delta$ 202.2, stating existence of a carbonyl group. Reduction with NaBH$_4$ at the C-2 carbonyl of 1 at room temperature yielded mainly compound 4 with $\alpha$-OH configuration at position C-2. This result was supported by the presence of coupling constant ($J = 3.0$ Hz) of $H_{ax-1}$ through $H_{ex-2}$ being located at chemical shift $\delta$ 5.56 ppm. This is due to that coupling constant of $H_{ax}$ and $H_{ax}$ is 2~3 Hz, which is smaller than that of $H_{ax}$ and $H_{ax}$ ($J = 6$~8 Hz).

Compounds 5 and 6 were obtained using esterification of 1 followed by amination. It is well-known that morpholine is one of the commonly used chemical groups because it plays an important role in medicinal chemistry and agrochemicals, such as morpholine derivatives Amorolfine [20] used as
agricultural fungicides. To investigate whether hybrid of morpholine and compound 1 would display multiple activity or improved activity, we firstly prepared compound 5 through Yamaguchi esterification [21] at 8-OH of 1, which involves the reaction of a 2-chloroacetic acid with 2,4,6-trichlorobenzoyl chloride to form the mixed anhydride, and upon reaction with an 8-OH in the presence of DMAP to produce the 2-chloroacetic ester derivative 5 in 85% yield. Then, amination of compound 5 with morpholine afforded the compound 6.

Next, compound 1 and derivatives (2–6) was evaluated for their antifeedant activity against *M. separata* using a leaf disk choice bioassay [22]. The results are presented in Table 3. Some of the compounds caused more reduction in feeding of third instar larvae of *M. separata*, when given a choice between control and treated wheat leaf disks at 4 µg/cm² at 24 and 48 h. Compounds 2 and 4 were respectively 1.2 and 1.5 times more active as a feeding deterrent than the parent compound 1. Other compounds caused 0–15% feeding deterrent effect. The results showed that compounds 2 and 4 were more active than the parent compound 1 at 24 h and 48 h but less potent than the positive control probenazole (100% feeding deterrent effect), an agrochemical used for the protection of rice plants [23]. The other compounds (3, 5 and 6) showed weak or no feeding deterrent effect. Preliminary results suggested that acylated and reduced derivatives at C-8 and C-2, respectively, of 1 may improve the antifeedant effect, while oxidation, esterification and morpholine hybrids at C-8 of 1 may decrease the effect.

**Table 3.** Antifeedant effects of compounds (1–6) against 3rd-instar larvae of *M. separata*.

| Entry | 24 h-Antifeedant rate (%) | 48 h-Antifeedant rate (%) |
|-------|---------------------------|---------------------------|
| control | 0                         | 0                         |
| 1     | 55                        | 45                        |
| 2     | 65                        | 50                        |
| 3     | 15                        | 12.5                      |
| 4     | 80                        | 72.5                      |
| 5     | 0                         | 0                         |
| 6     | 0                         | 0                         |
| Probenazole | 100                       | 97.50                     |

*Antifeedant effects of compound 1, derivatives and probenazole (positive control) were measured at 4 µg/cm² in acetone. Three replications (N = 20 larvae/treatment, the weight of larva was 13–16 mg) were used in each experiment and experiment was repeated twice. Means were reproducible with deviation less than ± 15%.

3. Experimental Section

3.1. General

NMR spectra were recorded on a Bruker Advance III 500 instrument (Bruker Daltonics Inc., Bremen, Germany) in CDCl₃ with TMS as internal standard for protons and solvent signals as internal standard for carbon spectra. Chemical shift values are mentioned in δ (ppm) and coupling constants are given in Hz. Mass spectra (MS) were recorded on an ESI-esquire 3000 Bruker Daltonics instrument (Bruker Daltonics Inc., Bremen, Germany). HR-ESI-MS data were collected on Shimadzu liquid chromatography-mass spectrometry (LCMS)-iontrap (IT)-time of flight (TOF) (Shimadzu, Kyoto,
Further stirred for 4 h with monitoring by TLC analysis. The reaction mixture was poured into ice water (10 mL) and extracted with AcOEt (3 × 10 mL) and extracted with ethyl acetate (3 × 10 mL). The combined organic layer was washed with NaHCO₃ solution and dried over anhydrous Na₂SO₄. After evaporation in vacuum, the resulting solid was subjected to a silica-gel column chromatography with AcOEt/petroleum ether (1:6) as the eluent to give compound 2 in almost 65% yield. Yellow solid; AcO-1, BzO-6, BzO-9 was certified by ¹H NMR δ: 1.71 (s, 3H, AcO-1), 8.02 (d, 2H, J = 7.7 Hz, BzO-6), 7.50 (t, 2H, J = 7.7 Hz, BzO-6), 7.62 (t, 1H, J = 7.7 Hz, BzO-6), 8.04 (t, 2H, J = 7.7 Hz, BzO-9), 7.46 (t, 2H, J = 7.7 Hz, BzO-9), 7.58 (t, 1H, J = 7.7 Hz, BzO-6); ¹³C NMR δ: 169.8 (1C), 164.6 (1C), 20.0 (1C), 133.6 (1C), 129.9 (2C), 128.6 (2C), 128.8 (1C), 165.3 (1C), 133.7 (1C), 129.5 (3C), 128.5 (2C), other data see Tables 1 and 2; ESI-MS m/z: 593.96 [M + H]⁺, 615.94 [M + Na]⁺; HR-ESI-MS m/z: 615.2219 [M + Na⁺] (calcd. for C₃₃H₃₆O₁₀Na, 615.2201).

Compound 3 (1α-acetoxy-6β,9β-dibenzyloxy-2,8-dioxodihydro-β-agarofuran): To a stirred solution of compound 1 (11 mg, 0.02 mmol) in DCM (3 mL) was added DMP (0.024 mmol; J & K Scientific Ltd., Beijing, China) at room temperature, and then the mixture was further stirred for 3 h with monitoring by TLC analysis. The reaction mixture was poured into ice water (10 mL) and extracted with AcOEt (3 × 10 mL; Sinopharm Chemical Reagent Co. Ltd., Shanghai, China). The combined organic layer was washed with NaHCO₃ solution and dried over anhydrous Na₂SO₄. After evaporation in vacuum, the resulting solid was subjected to a silica-gel column chromatography with AcOEt/petroleum ether (1/6) as the eluent to give compound 3 in 90% yields. White solid; AcO-1, BzO-6, BzO-9 was certified by ¹H NMR δ: 1.67 (s, 3H, AcO-1), 8.06 (d, 2H, J = 7.7 Hz, BzO-6), 7.52 (t, 2H, J = 7.7 Hz, BzO-6), 7.65 (t, 1H, J = 7.7 Hz, BzO-6), 8.00 (t, 2H, J = 7.7 Hz, BzO-9), 7.43 (t, 2H, J = 7.7 Hz, BzO-9), 7.58 (t, 1H, J = 7.7 Hz, BzO-6); ¹³C NMR δ: 169.3 (1C), 20.2 (1C), 164.5 (1C), 133.9 (1C), 130.0 (2C), 128.6 (2C), 128.9 (1C), 128.3 (1C), 164.8 (1C), 134.1 (1C), 129.7 (1C), 128.8 (2C), 128.9 (1C), other data see Tables 1 and 2; ESI-MS m/z: 565.83 [M + NH₄]⁺, 571.22 [M + Na⁺]; HR-ESI-MS m/z: 571.1962 [M + Na⁺] (calcd. for C₃₃H₃₂O₉Na, 571.1939).

Compound 4 (1α-acetoxy-6β,9β-dibenzyloxy-2α,8α-dihydroxydihydro-β-agarofuran): To a stirred solution of compound 1 (22 mg, 0.04 mmol) in anhydrous methanol (10 mL) was added NaBH₄
(10 mg, 0.2 mmol) in 10 min at room temperature, and then the mixture was further stirred for 1 h with monitoring by TLC analysis. The reaction mixture was extracted with AcOEt (3 × 10 mL). The combined organic layer was washed with HCl (1 mol/L) solution to pH = 7 and dried over anhydrous Na2SO4. After evaporation in vacuum, the resulting solid was subjected to a silica-gel column chromatography with AcOEt/petroleum ether (1:3) as the eluent to give compound 4 in almost 85% yield. White solid; AcO-1, BzO-6, BzO-9 was certified by 1H NMR δ: 1.73 (s, 3H, AcO-1), 8.01 (d, 2H, J = 7.7 Hz, BzO-6), 7.53 (t, 2H, J = 7.7 Hz, BzO-6), 7.65 (t, 1H, J = 7.7 Hz, BzO-6), 8.10 (t, 2H, J = 7.7 Hz, BzO-9), 7.48 (t, 2H, J = 7.7 Hz, BzO-9), 7.60 (t, 1H, J = 7.7 Hz, BzO-6); 13C NMR δ: 170.0 (1C), 20.8 (1C), 165.5 (1C), 133.3 (1C), 129.7 (1C), 128.8 (1C), 130.2 (1C), 128.7 (1C), 129.6 (1C), 165.6 (1C), 133.3 (1C), 130.0 (1C), 128.6 (1C), 128.4 (1C), 128.6 (1C), 130.1 (1C), other data see Table 1 and 2; ESI-MS m/z: 552.77 [M + H]+, 575.12 [M + Na]+; HR-ESI-MS m/z: 575.2273 [M + Na]+ (calcd. for C31H36O8Na, 575.2252).

Compound 5 (1α-acetoxy-6β,9β-dibenzylxoy-8α-chloracetyl-2-oxodihydro-β-agarofuran): To a solution of chloroacetic acid (4 mg, 0.04 mmol) in toluene (0.2 mL) at room temperature were added Et3N (11 μL, 0.08 mmol) and 2,4,6-trichlorobenzoyl chloride (6.4 μL, 0.05 mmol). The resulting mixture was stirred for 1 h before a solution of compound 1 (66 mg, 0.12 mmol) and DMAP (6.3 mg, 0.05 mmol) in toluene (0.15 mL) added. The resulting mixture was heated to 80 °C and stirred for 24 h, then it was quenched with a NaHCO3 (1 mL). The layers were separated and the aqueous layer was extracted with EtOAc (3 × 10 mL). The combined organic layers were washed with brine (3 mL), dried (Na2SO4) and concentrated in vacuum. The resulting solid was subjected to a silica-gel column chromatography with AcOEt/petroleum ether (1:10) as the eluent to give compound 5 in almost 85% yield. White solid; AcO-1, BzO-6, BzO-9 was certified by 1H NMR δ: 1.72 (s, 3H, AcO-1), 8.02 (d, 2H, J = 7.7 Hz, BzO-6), 7.50 (t, 2H, J = 7.7 Hz, BzO-6), 7.63 (t, 1H, J = 7.7 Hz, BzO-6), 8.04 (t, 2H, J = 7.7 Hz, BzO-9), 7.47 (t, 2H, J = 7.7 Hz, BzO-9), 7.60 (t, 1H, J = 7.7 Hz, BzO-6); 13C NMR δ: 169.3 (1C), 20.0 (1C), 164.6 (1C), 133.8 (1C), 129.6 (1C), 128.7 (1C), 129.6 (1C), 166.0 (1C), 133.1 (1C), 128.8 (2C), 128.7 (2C), 128.2 (1C), other data see Tables 1 and 2; ESI-MS m/z: 626.71 [M + H]+, 649.01 [M + Na]+; HR-ESI-MS m/z: 649.1829 [M + Na]+ (calcd. for C33H35O11ClNa, 649.1811).

Compound 6 (1α-acetoxy-6β,9β-dibenzylxoy-8α-morpholinoacetoxy-2-oxodihydro-β-agarofuran): To a stirred solution of compound 5 (25 mg, 0.04 mmol) in DCM (10 mL) was added morpholine (0.05 mL) at room temperature, and then the mixture was further stirred for 20 h, with monitoring by TLC analysis. The reaction mixture was poured into ice water (10 mL) and extracted with ethyl acetate (3 × 10 mL). The combined organic layer was washed with NaHCO3 solution and dried over anhydrous Na2SO4. After evaporation in vacuum, the resulting solid was subjected to a silica-gel column chromatography with methanol/chloroform (1:50) as the eluent to give compound 6 in almost 50% yield. Yellow solid; AcO-1, BzO-6, BzO-9 was certified by 1H NMR δ: 1.71 (s, 3H, AcO-1), 8.02 (d, 2H, J = 7.7 Hz, BzO-6), 7.49 (t, 2H, J = 7.7 Hz, BzO-6), 7.62 (t, 1H, J = 7.7 Hz, BzO-6), 8.02 (t, 2H, J = 7.7 Hz, BzO-9), 7.47 (t, 2H, J = 7.7 Hz, BzO-9), 7.60 (t, 1H, J = 7.7 Hz, BzO-6); 13C NMR δ: 169.3 (1C), 20.0 (1C), 164.6 (1C), 133.9 (1C), 129.5 (2C), 128.8 (2C), 129.9 (1C), 168.9 (1C), 133.8 (1C), 129.4 (2C), 128.7 (2C), 128.4 (1C), other data see Tables 1 and 2; HR-ESI-MS m/z: 678.2915 [M + H]+ (calcd. for C37H44O11Na, 678.2909).
### 3.3. Antifeedant Test

Third instar larvae of *M. separata* Walker (armyworm), provided by the Institute of Pesticides, Northwest A & F University, were used as test insect for evaluation of antifeedant activity using a leaf disk choice bioassay [22]. *M. separata* was continuously maintained on a Petri dish (Ø 10 cm) in a growth chamber in our laboratory at a temperature of 27 °C and 60% relative humidity under a 16L:8D photoperiod. Leaf discs (0.5 × 0.5 cm) were cut from wheat leaves (*Triticum aestivum*; Poaceae). Control wheat leaf discs were painted on each side with 10 µL of the carrier solvent (acetone; Sinopharm Chemical Reagent Co. Ltd., Shanghai, China), and test wheat leaf discs with the same amount of the test solution at 100 µg/mL (4 µg/cm² of the final concentration). After the solvent had evaporated, it was treated and control discs were placed in each compartment of a plastic assay tray. Third instar larvae (weight 13–16 mg) starved for 5–7 h, was introduced gently into the center of each compartment. The number of larvae was 20 per treatment, and three replications were used in each experiment. Probenazole (J & K Scientific Ltd., Beijing, China) was used as a positive control. The amount of leaf area consumed at 24 and 48 h was estimated using squared graph paper. The antifeeding rate (%) was calculated using the formula \[\frac{(C - T)}{(C + T)} \times 100\], where C and T are areas consumed of the control and treated leaf disks, respectively. Experiment was repeated twice.

### 4. Conclusions

In conclusion, five new dihydro-β-agarofuran sesquiterpenes were semisynthesized using a dihydro-β-agarofuran with C-2 carbonyl (compound 1) from *P. wightiana* as a starting material. Some sesquiterpenes showed certain antifeedant activity, and structure-activity relationships suggested that the antifeedant effects of the derivatives depend on the position of hydroxyl and acetoxy substituents at C-8 and C-2 positions (compounds 2 and 4). This will be helpful in designing new dihydro-β-agarofuran insect control agents against *M. separata* and other important pests.

### Acknowledgments

This project was supported by the National Natural Science Foundation of China (31200254) and Chinese Universities Scientific Fund (QN2012047).

### Conflicts of Interest

The authors declare no conflict of interest.

### References

1. Agarwal, M.; Walia, S.; Dhingra, S.; Khambay, B.P.S. Insect growth inhibition, antifeedant and antifungal activity of compounds isolated/derived from *Zingiber officinale* Roscoe (ginger) rhizomes. *Pest. Manag. Sci.* 2001, 57, 289–300.
2. Akhtar, Y.; Isman, M.B.; Niehaus, L.A.; Lee, C.-H.; Lee, H.-S. Antifeedant and toxic effects of naturally occurring and synthetic quinones to the cabbage looper, *Trichoplusia ni*. *Crop Prot.* 2012, 31, 8–14.
3. Gao, J.M.; Wu, W.J.; Zhang, J.W.; Konishi, Y. The dihydro-β-agarofuran sesquiterpenoids. *Nat. Prod. Rep.* **2007**, *24*, 1153–1189.
4. Liu, G.Q.; Gao, J.M.; Wu, W.J. Adance in macrocyclic dihydroagarofuran pyriding alkaloids. *Chem. Res. Appl.* **2003**, *15*, 321–326.
5. Spivey, A.C.; Weston, M.; Woodhead, S. Celastraceae sesquiterpenoids: Biological activity and synthesis. *Chem. Soc. Rev.* **2002**, *31*, 43–59.
6. Kuo, Y.-H.; King, M.-L.; Chen, C.-F.; Chen, H.-Y.; Chen, C.-H.; Chen, K.; Lee, K.-H. Two new macrolide sesquiterpene pyride alkaloids from *Maytenus emarginata*: Emarginatine G and the cytotoxic emarginatine F. *J. Nat. Prod.* **1994**, *57*, 263–269.
7. Pérez-Victoria, J.M.; Tincusi, B.M.; Jiménez, I.A.; Bazzocchi, I.L.; Gupta, M.P.; Castanys, S.; Gamarrón, F.; Ravelo, A.G. New natural sesquiterpenes as modulators of daunomycin resistance in a multidrug-resistant *Leishmania tropica* Line. *J. Med. Chem.* **1999**, *42*, 4388–4393.
8. Duan, H.; Takaishi, Y.; Bando, M.; Kido, M.; Imakura, Y.; Lee, K. Novel sesquiterpene esters with alkaid and monoterpane and related compounds from *Tripterygium hypoglaucum*: A new class of potent anti-HIV agents. *Tetrahedron Lett.* **1999**, *40*, 2969–2972.
9. Munoz-Martinez, F.; Lu, P.; Cortés-Selva, F.; Perez-Victoria, J.M.; Jimenez, I.A.; Ravelo, A.G.; Shoram, F.J.; Gamarrón, F.; Castanys, S. Celastraceae sesquiterpenes as a new class of modulators that bind specifically to human P-glycoprotein and reverse cellular multidrug resistance. *Cancer Res.* **2004**, *64*, 7130–7138.
10. Cortés-Selva, F.; Jiménez, I.; Munoz-Martinez, F.; Campillo, M.; Bazzocchi, I.; Pardo, L.; Ravelo, A.; Castanys, S.; Gamarrón, F. Dihydro-β-agarofuran sesquiterpenes: A new class of reversal agents of the multidrug resistance phenotype mediated by P-glycoprotein in the protozoan parasite Leishmania. *Curr. Pharm. Des.* **2005**, *11*, 3125–3139.
11. Cortés-Selva, F.; Munoz-Martinez, F.; Ilias, A.; Jimenez, A.I.; Varadi, A.; Gamarrón, F.; Castanys, S. Functional expression of a multidrug P-glycoprotein transporter of Leishmania. *Biochem. Biophys. Res. Commun.* **2005**, *329*, 502–507.
12. Ujita, K.; Takaishi, Y.; Tokuda, H.; Nishino, H.; Iwashima, A.; Fujjita, A.T. Inhibitory effects of triptogelin A-1 on 12-O-tetradecanoylphorbol-13-acetate-induced skin tumor promotion. *Cancer Lett.* **1993**, *68*, 129–133.
13. Takaishi, Y.; Ujita, K.; Tokuda, H.; Nishino, H.; Iwashima, A.; Fujjita, T. Inhibitory effects of dihydroagarofuran sesquiterpenes on Epstein-Barr virus activation. *Cancer Lett.* **1992**, *65*, 19–26.
14. Zhang, J.; Hu, Z.; Li, S.; Wu, W. Synthesis and insecticidal activities of new ester-derivatives of Celangulin-V. *Int. J. Mol. Sci.* **2011**, *12*, 9596–9604.
15. Zhang, J.; Hu, Z.; Yang, H.; Wu, W. Synthesis and insecticidal activities of new ether-derivatives of Celangulin-V. *Nat. Prod. Commun.* **2010**, *5*, 845–848.
16. Zhang, J.; Li, S.; Ji, Z.; Hu, Z.; Wu, W. Synthesis and insecticidal activity of novel dimers of celangulin-V and podophyllootoxin. *Chem. Nat. Compd.* **2009**, *45*, 507–510.
17. Johnson, L.A.; Soltis, D.E. *matK* DNA sequences and phylogenetic reconstruction in Saxifragaceae. *Syst. Bot.* **1994**, *19*, 143–156.
18. Wang, D.-M.; Li, D.-W.; Pu, W.-J.; Liu, J.-J. Preparation and application of dihydro-β-agarofuran sesquiterpenes from *Parnassia wightiana* Wall. CN Patent 103073529, 2013.
19. Torres-Romero, D.; Jiménez, I.; Rojas, R.; Gilman, R.; Lópeze, M.; Bazzocchi, I. Dihydro-β-agarofuran sesquiterpenes isolated from Celastrus vulcanicola as potential anti-Mycobacterium tuberculosis multidrug-resistant agents. Bioorgan. Med. Chem. 2011, 19, 2182–2189.

20. Polak, A. Preclinical data and mode of action of amorolfine. Clin. Exp. Dermatol. 1992, 17, 8–12.

21. Dhimitruka, I.; SantaLucia, J., Jr. Investigation of the Yamaguchi esterification mechanism. Synthesis of a Lux-S enzyme inhibitor using an improved esterification method. Org. Lett. 2006, 8, 47–50.

22. Akhtar, Y.; Isman, M.B.; Paduraru, P.M.; Nagabandi, S.; Nair, R.; Plettner, E. Screening of dialkoxybenzenes and disubstituted cyclopentene derivatives against the cabbage looper, Trichoplusia ni, for the discovery of new feeding and oviposition deterrents. J. Agric. Food Chem. 2007, 55, 10323–10330.

23. Iwai, T.; Seo, S.; Mitsuhara, I.; Ohashi, Y. Probenazole-induced accumulation of salicylic acid confers resistance to Magnaporthe grisea in adult rice plants. Plant Cell Physiol. 2007, 48, 915–924.

© 2013 by the authors; licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution license (http://creativecommons.org/licenses/by/3.0/).