Antifungal and Insecticidal properties of the Phytoconstituents of *Drimys winteri* (Winteraceae) growing in Chiloé Island (Chile)

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Abstract

Seven drimane-type sesquiterpenes: polygodial, drimenol, drimenidol, drimenin, isodrimininol, isodrimenin and proximadiol (cryptomeridiol); three lignans: sesamin, cubebin and its epimer eudesmin, and one sterol: (β-sitosterol) were isolated from *Drimys winteri* (Winteraceae) from Chiloe Island, Chile. Eudesmin and its phytosterols were isolated from the plant for the first time, and sesamin was found to be the major product in the plant’s bark by a significant margin. Furthermore, qualitative and quantitative differences were detected between some products. Structures were established using 1D and 2D NMR (COSY, HSQC, and HMBC) spectra, HRESIMS and comparison with published data. Some isolated compounds were tested against the phytopathogenic fungi *Botrytis cinerea*, *Dothiorella sp.*, *Penicillium sp.* and *Cladosporium cladosporioides*; additionally, a study was carried out to determine the contact toxic action of leaf and stem bark extracts upon phloemocytic insect pests from the *Pseudococcus viburni* (Signoret) (Hemiptera: Pseudococcidae) mealy bug complex.

Keywords: *Drimys winteri*; Lignans; Sterols; Sesquiterpenoids; Antifungal activity; Insecticidal properties

Introduction

*Drimys winteri* J. R and G. Forster var. *chilensis* (DC.) A. Gray is a tree native to Central and Southern Chile, which grows in continental Chile from Coquimbo to Aysén, including Chiloé Island. It is abundant in wet swampy localities from sea level to an altitude of 1700 m [1]. In continental Chile, Winteraceae plants are represented by two species of Drimys: *D. winteri* J. R. Forster and G. Forster and *D. andina* Reiche [2]. *D. winteri* is a tree whereas *D. andina* is a shrub [3]. Marticorena and Quezada also distinguish the *chilensis* (DC.) A. Gray variety of *D. winteri* [4]. *Drimys winteri* is a sacred plant to the Mapuche indigenous people, who uses its aerial parts for the treatment of dermatitis, stomach pain, toothache, tumors, and other illnesses [5]. Extracts of the stem bark have long been used to treat human and bovine diseases [6]. In the past, the stem bark of *D. winteri* was exported to Europe as an antiscorbutic medicine [1,5]. The tree is also used for commercial purposes, in wood production, crafts, manufacturing and the pulp industry due to the high quality of its fibers. Its architectural characteristics and natural resistance to insects and microorganisms are advantageous for playgrounds in parks and gardens in a wide range of weather conditions. The wood of this species is highly valued and used to manufacture furniture and musical instruments as well as to protect crops [7]. Furthermore, extracts of aerial parts of *D. winteri* have industrial applications in products such as cosmetics, phytonutrients and pest repellant agents [7-9]. Previous reports on the chemical composition of continental *D. winteri* revealed the presence of tannins, flavonoids and essential oils containing sesquiterpene lactones [5]. The drimane-type sesquiterpenoids possess a wide variety of biological activities, including antimicrobial, pungency, antibacterial, antifungal, antifeedant, cytotoxic, molluscicidal, piscicidal, growth regulation and phytotoxic properties [9]. The pungency of several drimanes and their irritant properties on the skin (allergies) has also attracted attention [6-10]. To the best of our knowledge, as of the present, fourteen drimane-type sesquiterpenoids have been isolated from continental *D. winteri* and probably also from *D. andina* [5,6], as well as one aromadendrane derivative [11] and one drimane-type sesquiterpene trimer [12]. The sesquiterpenoids reported from the bark and leaves collected in Brazil [13] were not considered because *D. winteri* is not present in Brazil [3]. The main characteristics and properties of this tree have been attributed to the presence of certain sesquiterpenoids. The reactivity of the unsaturated dialdehyde functionality towards biological nucleophiles is considered responsible for the antifeedant activity of those compounds (polygodial and epipolygodial) [14,15]. The chemical composition of the essential oils obtained by hydrodistillation of the stem bark and leaves of continental Chile (Santiago) *D. winteri* was also studied using GC and GC/MS for which results have been published [16]. The essential oils from the stem bark and leaves of *D. winteri* from Chiloé Island and continental Chile (Santiago) were studied revealing certain differences in their chemical composition. Sesquiterpene hydrocarbons constituted the main chemical groups in the stem bark oils (in which α-santalene, trans-β-bergamotene curcumenes are the major components), whereas monoterpenes (in which (α) - pinene, β-pinene, and linalool are the main components) constituted the main chemical groups in the leaves of Island plants. Sesquiterpenes (germacrene) and phenylpropanoids are most abundant in the leaves of continental plants [16]. The aim of the present study was to compare the chemical profile of insular *D. winteri* from Chiloé Island (growing and flourishing in climate conditions of high humidity, abundant water and very scarce volcanic soil) that differ greatly from those where the continental population is found. We also separately investigated the components present in the bark and leaves and evaluated their

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insecticidal and antifungal properties for potential industrial use. The investigation also showed an unusual concentration of the sesamin lignan in bark from Chiloe canelo populations in comparison with continental populations, which would partially explain its extensive use by the local population as a treatment for certain conditions [17]; and the absence of bacteria and insect attacks on the species [18]. Until now, neither the existence of the lignan eudesmin nor its phytosterol content had been reported. This study also evaluates its possible use as a dietary supplement.

Materials and Methods

General

Column chromatography (GC) was carried out using silica gel 60G (Merck 7734, Germany). Thin layer chromatography (TLC) was performed on 2-mm thick silica gel F₂₅₄ plates (Merck 7731) and Chromatotron (Harrison-research 7924 T) or 2 mm thick discs, using silica gel 60 PF 254 (Merck 7749). Flash chromatography took place on silica gel 60 H (Merck 7739) with a hexane-acetate acetate gradient of 0-100% EtOAc. Mps are uncorrected. Optical rotations were measured with a Perkin Elmer 241 MC polarimeter. ¹HNMR spectra at 500MHz were recorded in CDCl₃, on a Bruker-DRX 500 MHz H Larmor frequency equipped with a 5-mm probe. ¹H, ¹³C and 2D (COSY, HMQC, HMBC) experiments were performed using standard Bruker pulse sequences. Capillary NMR spectra were recorded on a Varian Unity Inova 500 MHz spectrometer (Palo Alto, CA) equipped with a Protasit/MRM capNMR probe (Savoy, IL) in CDCl₃ and MD OD by direct injection of 7 µL samples. Proton assignment was determined using ¹H and COSY spectra while the carbon data was taken from HSQC and HMBC experiments.

Plant material

Drimys winteri var. chilensis was collected in February, June, September and December 2008 and 2009 on Chiloe Island, 30 km NW of Castro and 17 Kms SE of Chonchi Beach (Chile), within the same population. Voucher specimens are held at the Faculty of Sciences, Universidad de Talca (herbarium numbers 107-08). The plants were identified by Dr. José San Martín (Universidad de Talca, Instituto de Biología Vegetal y Biotecnología).

Extraction and isolation

Leaves were dried at room temperature for 3 weeks. Stem bark was dried in a forced-air oven at 40°C for 48 h.

1. Dried and powdered D. winteri stem bark (700 g) was extracted using n-hexane (3 x 2.0 L) at room temperature for 72 hours. After filtration, the residue was placed in a Soxhlet apparatus and extracted again using n-hexane for 24 hours. The solutions were combined and evaporated under reduced pressure (210 mbar) at 25-30°C to produce a yellowish oily residue (35.0 g). The crude extract was first subjected to flash CC (silica gel, 230–400 mesh, 650 g) and fractionated by gradient elution (100% n-hexane to 100% EtOAc), and individual fractions were then further purified by two CC on silica gel. Elution with n-hexane-ethyl acetate (98:2-90:10) yielded 2.84 g of a yellow mixture consisting of four compounds. Further separation using the Chromatotron (film thickness 2 mm) and elution with n-hexane-ethyl acetate (8:2) produced three main compounds which were further purified by CC (silica gel 230–400 mesh, 600 g) and fractionated by n-hexane-ethyl acetate (9:1 to 8:2) gradient elution yielding cubebin (0.12 g), its epimer (epi-cubebin (0.061 g) and eudesmin (0.041 g); Sesamin was also isolated.

Polygodial (Figure 1) (Scheme a): [α] D and spectroscopic data (NMR, MS) were identical to data previously reported on this compound [6].

Drimenol (Figure 2) (Scheme a): The MS, ¹HNMR and ¹³CNMR data on this compound correlated well with that reported elsewhere [12,19,20].

Drimedinol (Figure 3) (Scheme a): The ¹HNMR and ¹³CNMR spectral data on this compound correlated well with that reported elsewhere [6].

Drimenin: The MS, ¹HNMR and ¹³CNMR spectral data on this compound correlated well with reported data [20,21].

Isodrimenin: Already identified in D. winteri [6].

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Figure 1: Mortality with Treatment A (stem-bark) at different dilution rates.

Figure 2: Mortality with the Treatment B (leaves) at different dilution rates.
Antifungal activity

Antifungal activity against Botrytis cinerea Pers., Dothiorella sp., Penicillium sp. and Cladosporium cladosporoides was determined in triplicate experiments by microdilution [30-32] and the results are presented as the minimum inhibitory concentration (MIC). The spores were cultured on Sabouraud medium at 25°C for 7 days. The compounds were assessed in the 600-18.75 μg/mL dilution interval, and standard antifungal compounds were assessed in the of 250-3.9 μg/ml range. Spores were obtained from well-grown and sporulating fungal cultures maintained in potato-glucose agar medium by suspension in sterile distilled water, filtration on glass wool and centrifugation. The spores were counted in a Neubauer chamber and diluted with sterile distilled water to a final concentration of 10⁶ spores/mL. The assay was carried out in 96-well microtiter plates. 100 μl of the spore suspension was incubated with 100 μl of the compound sample suspended in Sabouraud medium. The final volume of the mixture was 200 μl. A spore germination control and a Sabouraud medium control were included in all of the experiments as well as the standard antifungal compounds. Mortality counts were assessed at days 5, 12 and 20 after spraying treatments. To determine full mortality, specimens were punctured slightly under a binocular microscope (Figure 1 and 2).

Analysis of phytosterols chemicals

n-Hexane and methanol (HPLC grade) from Th. Geyer (Renningen, Germany).

Technical grade ethanol (distilled prior to use) from BASF (Ludwigshafen, Germany). Potassium hydroxide (KOH, >85%) from Carl Roth (Karlsruhe/Germany). Pyridine (purity > 99.8%) from Sigma-Aldrich (Steinheim, Germany) and silylating agent consisting of 99% N, O-bis (trimethylsilyl) trifluoroacetamide (BSTFA) and 1 % trimethylsilyl chlorosilane (TMCS) from Supelco (Bellefonte, PA, USA).

Saponification and trimethylsilylation

For saponification, about 50 mg of plant material was heated in a test tube in 2 mL ethanolic KOH solution (prepared from 9 mL ethanol and 1 mL KOH 50% in water (w/w)) for 60 minutes at 80°C. After cooling to room temperature, 2 mL n-hexane and 2 mL water was added. The tube was closed and shaken vigorously. The organic phase was transferred into a vial. An aliquot (corresponding to ~0.1 mg of unsaponifiable matter) was transferred to another vial and the
solvent was stirred by means of a gentle stream of nitrogen. Pyridine (25 µL) and 50 µL of the silylation agent were added to the residue and the closed vial was heated for 30 min at 60°C. Afterwards, the solvent was stirred by means of a gentle stream of nitrogen and the residue was rediluted in 1 mL n-hexane. This solution was used for GC/MS analysis.

GC/MS conditions

Analyses were carried out using a 6890/5973N GC/MS system (Agilent, Waldbronn, Germany) equipped with an MPS2 autosampler system (Gerstel Mülheim/Germany). Injections (1 µL) were performed in splitless mode. The temperatures of the injector, transfer line, ion source and quadrupole were set to 250°C, 280°C, 230°C and 150°C, respectively. An HP-5MS U1 column (30 m, 0.25 mm i.d., 0.25 µm film thickness, Agilent, Waldbronn, Germany) was installed in the GC oven. Helium (purity 5.0) was used as carrier gas with a flow rate of 1.2 mL min⁻¹. The GC oven was programmed as follows: After 1 min at 55°C the temperature was increased by 10°C/min to 300°C. This temperature was held for 10 min. Data was recorded in full scan mode from m/z 50-550 after a solvent delay of 6 min.

Results and Discussion

In this paper, we have reported the compounds isolated from Chiloé Island *Drimys winteri*: six drimane-type sesquiterpenoids (a, 1-5); one sesquiterpene eudesmane (cryptomeridiol), the lignans sesamin (c), cubebin as a mixture of two epimers and the non-phenolic furofuran lignin (+) eudesmin. The complete 1H NMR and 13CNMR spectral data on this compound correlated well with that reported elsewhere [6,19,24,28,29,33,34] (Scheme a). All compounds, except eudesmin and β-sitosterol had been previously isolated from continental *D. winteri* from the continental population [26]. It is well-known insect antifeedant acting on several insects [8,14,15]. Poligodial was the most active compound against *Botrytis cinerea*; mixing poligodial with drimenin or with isodrimenin increased activity against *B. cinerea*, *Dothiorella* sp. and *Cladosporium cladosporoides*. No compound was active against *Penicillum* sp. The poligodial-drimenin mixture showed a major increase in the aforementioned activity. Table 1 shows the antifungal activity of the compounds and mixtures on phytopathogenic fungi, expressed in µg/mL. These results agree with a previous report [35], which showed poligodial to be the most potent antifungal compound, which became much more potent when combined with perillaldehyde. The synergy of poligodial and perillaldehyde against bacteria was the first example of the synergistic effect of poligodial against both Gram-positive and Gram-negative bacteria [36]. In insular *D. winteri* stem bark, drimenol (a, 2), was found at lower concentrations than poligodial. However, it was detected in continental *D. winteri* [6], in addition to drimenidol, a drimane sesquiterpenoid also previously isolated by Brown [11]. Drimenidol (a, 3) was also detected in the June and September collections. The sesamin lignin was also isolated from the stem bark. This compound is highly valued as an insect repellent for protecting buildings materials against undesired insect infestations. Because some lignans have potent antimicrobial, antifungal, antiviral, antioxidant, insecticidal and antifeedant properties, they probably play an important role in defending plants against various biological pathogens and pests [37]. At the ecological level, they are involved in plant interactions with other organisms, mainly due to their defensive role, and in protecting plants from physical damage. Methylendioxyphenyl (piperonyl) derivatives such as cubebin, sesamin have been described as feeding inhibitors, feeding deterrents, or larval growth inhibitors [38]. Substituents such as the methoxy or methylendioxy groups enhance this activity not only in lignans but also in simple phenylpropanoids [39]. This could explain the anti-insect activity of Chiloé *Drimys winteri*. The high quantities of this lignan found in stem bark from insular *D. winteri* in comparison with those found in continental species could also explain the insecticidal properties of the insular specimen (Table 1). Sesamin has a strong allelochemical biological-like activity, making it useful as a natural industrial insecticide and pest repellant. Consequently, the fungicidal and bactericidal actions of several purified extracts of insular *D. winteri* are under investigation [39] (Scheme c). A recent report on sesamin [17]...
describes this important lignan as chondroprotective and anti-inflammatory; therefore it is a new and powerful natural option for treating arthroses. Arthrosis, or osteoarthrosis, is a chronic and degenerative disease leading to deterioration and loss of hyaline articular cartilage, alterations in the subchondral bone and damage to soft tissues, including the synovial membrane. The high sesamin concentrations in Chilöe canelo bark offer a powerful natural alternative for the treatment of this condition, which is widespread in Chile. Due to their strong antifeedant, antimicrobial and fungicidal activity, we investigated naturally occurring sesquiterpenoid dialdehydes from the drimane series, such as polygodial and others [3,40]. In insular D. winteri, the main compounds found in stem bark and leaves are the polygodial, drimenol and drimendiol sesquiterpenoids and the sesamin, cubebin and eudesmin lignans respectively. However, the main fungistatic activity is in the leaves, mainly due to the presence of sesamin, cubebin, eudesmin and polygodial, although it is also found in the stem bark, due to the presence of polygodial. These results corroborate the variability of this plant’s chemical constituents. Variability in the chemical composition of continental Chile Drimys populations has already been studied, particularly for polygodial and drimenol. In dried leaves, the mean concentration is ca. 1% for polygodial and 0.011% for drimenol. This concentration depends on the population, regions, biological material studied and the time of harvest. In Chiloe Island, we found lower concentrations of drimenol in leaves, (ranging from undetectable quantities to almost 0.1% in the Southern Hemisphere winter, June to August); however bark drimenol concentrations are greater than in the leaves, whereas differences in polygodial content were not significant in comparison with continental populations. The presence of other sesquiterpenes such as drimenin (a, 4), isodrimeninol (a, 5), isodrimenin (b), also changes significantly in Drimys leaves and bark from Chiloe; the lowest mean value was observed in leaves; concentrations ranged from undetectable quantities to almost 0.2%. The main drimenidol concentration in summer (December - January) in Chiloe bark is 1.1%, while in winter or spring it is undetectable (Table 2). Proximadiol (cryptomeridiol) shows similar concentrations to those of continental plants, although it disappears in winter. The cubebin and eudesmin lignans were also detected and isolated in Drimys winteri leaves from Chiloé Island, but were not detected in bark. In leaves, the sesamin lignan was found in concentrations of almost 2.71%. Cubebin has also demonstrated biological activity, for which reason the use of Aristolochia esperanzae Kunze is indicated for treating rheumatoid arthritis. Phytochemical analysis of A. esperanzae stems reveals a mixture of cubebin [41]. Furthermore, Cubebin has significant anti-inflammatory and analgesic effects similar to those observed in non-steroidal drugs. It can be inferred that cubebin is one of the compounds responsible for the activity already known in popular medicine [42]. Finally, some of the isolated compounds were tested against the phytopathogenic fungi Botrytis cinerea, Dothiorella sp., Penicillium sp. and Cladosporium cladosporioides. Polygodial was the most active compound against Botrytis cinerea (Table 3). (Figures 1- 2) show the insecticidal properties of three treatments using different Drymis winteri stem and leaf extract dilution rates against fruit tree mealy bug Pseudococcus viburni reared on potato sprouts. Average mortality was rated per three replications (50 specimens) compared with a non-treated control colony. These results show that at the dilution rates used, the highest concentrations from both bark and leaf extracts achieved full mortality within one week of treatments. At 70% dilution rates, full mortality was achieved only 30 days after the evaluation period (data not provided). No significant mortality was found in either treatment at 30% dilution rates, indicating that the minimum effective dosage should be explored at levels not below 50%. In unsaponifiable matter (the sterol-containing fraction), we were able to detect various volatile compounds, which were most likely terpenoids [Figure 3]. In addition, phytol and n-alcohols from 18 to 28 were detected, of which 24:0 was the most abundant. The labeled compound and its structure achieved a good match with the NIST MS library; however this was unverifiable due to the lack of a standard (due to it being a sesamin-related compound).

GC/MS analysis of the unsaponifiable matter of sesame oil, where sesamine is a well known constituent, verified detection of sesamine in Drymis winteri. β-Sitosterol was the only sterol detected in unsaponifiable matter (Figure 3).

Conclusions

Seven drimane-type sesquiterpenes, namely polygodial, drimenol, drimendiol, drimenin, isodrimeninol, isodrimenin and proximadiol; the lignans sesamin, (the majority compound), cubebin, their epimers and eudesmin, and β-sitosterol were isolated from Drimys winteri (Winteraceae) collected from Chiloe Island, Chile. This is the first time that eudesmin has been isolated from this plant. Sesamin, β-sitosterol was the major products; and qualitative and quantitative differences were detected between products from Chiloé Island and Continental Drimys species. Some isolated compounds were tested against phytopathogenic fungi and assays were also carried out to determine the contact toxic action of leaf and stem bark extracts on phloematic pest insects.

Acknowledgments

The authors are indebted to Prof. Dr. Philippe Christen (Geneva University) for the acquisition of the NMR spectra.

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**Table 1:** Quantification of sesquiterpenes in Drimys winteri populations.

<table>
<thead>
<tr>
<th>Compound tested</th>
<th>Phytotoxic fungi</th>
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<tbody>
<tr>
<td></td>
<td>Botrytis cinerea</td>
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<tr>
<td>Polygodial</td>
<td>337.5</td>
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<tr>
<td>Sesamin</td>
<td>&gt;600</td>
</tr>
<tr>
<td>Drimenol</td>
<td>&gt;600</td>
</tr>
<tr>
<td>Eudesmin</td>
<td>585</td>
</tr>
<tr>
<td>Isodrimenin</td>
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</tr>
<tr>
<td>Polygodial + drimenin</td>
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</tr>
<tr>
<td>Polygodial + eudesmin</td>
<td>600</td>
</tr>
<tr>
<td>Iprodione1 (reference)</td>
<td>31.3</td>
</tr>
<tr>
<td>Myclobutanil2 (reference)</td>
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</tbody>
</table>

**Table 2:** Variability Sesquiterpenes in D. winteri from Continental and Island Populations.

### Drimys winterize

<table>
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<th>Place</th>
<th>Extract (gr)</th>
<th>Sesamin (mg)</th>
<th>Yield %</th>
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<td>50</td>
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<tr>
<td>Leaves</td>
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<td>0.84</td>
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<td>Leaves</td>
<td>35</td>
<td>950</td>
<td>2.71</td>
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</table>

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**Table 3:** Anti-fungal activity of some compounds isolated from Drymis winteri. Data reported as MIC in µg/mL.
References