

ISOLATION AND SYNTHESIS OF PINOCEMBRIN AND PINOSTROBIN FROM *Artocarpus odoratissimus*

(Pemencilan dan Sintesis Pinoembrin dan Pinostrobin dari *Artocarpus odoratissimus*)

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Abstract

Artocarpus odoratissimus is native to Borneo and Mindanao Island. In Sarawak, it is locally known as Terap. Previous studies showed *Artocarpus* species are rich in phenolic compounds, including flavonoids, stilbenoids and arylbenzofurans. The chemical profile of the indigenous plants from Sarawak, such as *A. odoratissimus* has not been studied intensively and the analyses of medicinal properties have not been explored. Thus, a phytochemical study is carried out on the root extracts of *A. odoratissimus* using various chromatographic methods and it has led to the isolation of two known flavonoids, namely pinocembrin (1) and pinostrobin (2). The structures are identified by comparison of their ¹H and ¹³C NMR data with those reported in the literature. Flavonoid molecule incorporated as multifunctional in the pharmaceutical industry. It has a vast range of pharmacological activities, such as antimicrobial, anti-inflammatory, antioxidant, and anticancer activities. Pinocembrin and pinostrobin which are successfully synthesized with 2-hydroxy,4,6-methoxyacetophenone and benzaldehyde as starting materials were discussed.

Keywords: *Artocarpus odoratissimus*, terap, pinocembrin, pinostrobin

Abstrak

Artocarpus odoratissimus berasal dari Borneo dan Pulau Mindanao. Di Sarawak ia dikenali sebagai Terap. Kajian terdahulu menunjukkan yang spesis *Artocarpus* kaya dengan sebatian fenolik, termasuk flavonoid, stilbenoid dan arylbenzofuron. Profil kimia tentang tumbuhan yang berasal dari Sarawak, seperti *A. odoratissimus* masih belum dikaji secara intensif dan analisis tentang khasiat perubatan masih belum lagi diterokai. Oleh itu, kajian fitokimia yang telah dilakukan ke atas krud ekstrak akar *A. odoratissimus* dengan menggunakan pelbagai kaedah kromatografi telah berjaya memencilkan dua sebatian flavonoid, iaitu pinosembrin (1) dan pinostrobin (2). Bentuk struktur telah dikenal pasti melalui perbandingan data ¹H dan ¹³C NMR dengan kajian literatur. Molekul flavonoid mempunyai pelbagai fungsi di dalam industri farmaseutikal. Ia mempunyai aktiviti farmakologi yang sangat meluas seperti antimikrob, anti-radang, antioksidan, dan antikanser. Pinosembrin dan pinostrobin yang telah berjaya disintesis dengan menggunakan 2-hidroksi,4,6-metoksiasetofenon dan benzaldehid sebagai bahan pemula telah dibincangkan.

Kata kunci: *Artocarpus odoratissimus*, terap, pinosembrin, pinostrobin

Introduction

Artocarpus odoratissimus is native to Borneo. It is cultivated in the Philippines and is also found in the wild abundantly in Brunei Darussalam, Sabah and Sarawak [1]. *Artocarpus* species are used as traditional medicines such as treatment against inflammation, malarial fever, diarrhea, diabetes and tapeworm infection. According to Jagtap and Bapat [2] compounds from *Artocarpus* species exhibited diverse biological activities including antibacterial, antitubercular, antiviral, antifungal, antiplatelet, antiarthritic, tyrosinase inhibitory and cytotoxicity. Some *Artocarpus* species are used as traditional medicines. For example *Artocarpus odoratissimus* (terap) is used by local communities in Sarawak as an antidote against centipede and scorpion stings by applying the ash from the leaves on the wounds, while for treatment of scabies the ash is added with a little amount of coconut oil [3]. Ee et al. [4] reported that the isolation of a new prenylated pyranoflavone derivatives, artosimmin and traxateryl acetate. Artosimmin exhibited significant cytotoxicity against cancer cell lines (HL-60 and MCF-7) and showed antioxidant properties.

Artocarpus species also rich in phenolic compounds, including flavonoids, stilbenoids and arylbenzofurans. Most of the pharmacological effects can be explained by the presence of phenolic compounds [5]. However, there are very few studies on the *Artocarpus* species from Sarawak, therefore it would be interesting to develop a chemical profile of such Sarawakian species. Thus, a phytochemical study is carried out on the root extracts of *Artocarpus odoratissimus* using various chromatographic methods and it has led to the isolation of two known flavonoids, namely pinocembrin and pinostrobin.

Pinocembrin is a flavanone, a type of flavonoid. It is an antioxidant found in damiana, honey, and propolis. Pinocembrin is a major flavonoid molecule incorporated as multifunctional in the pharmaceutical industry. Its vast range of pharmacological activities has been well studied including antimicrobial, anti-inflammatory, antioxidant, and anticancer activities. Pinocembrin has potential as a drug to treat cerebral ischemia, neurodegenerative diseases, cardiovascular diseases and atherosclerosis as well as other clinical conditions [6]. Pinocembrin also exhibits various interesting biological activities such as antibacterial [7], antifungal [8] and anti-inflammatory [9], [10]. Furthermore, pinocembrin plays a pivotal role in prevention and cure of various neurodegenerative diseases, such as Alzheimer's disease (AD), Parkinson's disease and other neuronal dysfunctions [11].

Pinostrobin is a potent flavonoid inducer of mammalian phase 2 chemoprotective and antioxidant enzymes [12]. Pinostrobin isolated from *Polygonum lapathifolium* nodosum could quickly penetrate through cytoplasm to the cellular nucleus of the cultured cells and gives intensive apoptotic response in stimulating leukemic cells *in vitro*. As such, it may be considered as a good candidate for a leukemia chemopreventive agent [13]. Pinostrobin from Temu Kunci (*Kaempferia pandurata* Roxb) has anticancer activity againsts benzo(a)pyrene-induced fibrosarcoma cancer on mice *in vivo* [14]. With increasing concentration, pinostrobin caused a gradual leakage, also contributing to breakage of the envelope and virus inactivation. Treatment effect of oral pinostrobin *in vivo* showed that pinostrobin possesses definite therapeutically effect in the development of lesion score [15].

The interest in the biological properties of pinocembrin and pinostrobin attracted the intense synthetic efforts towards the synthesis of various derivatives compounds. By synthesis we can also save the usage of natural resources of raw material.

Materials and Methods

Plant materials

The root of *Artocarpus odoratissimus* was collected from Kampung Krokong Pisa, Bau Sarawak. Voucher specimen of the plant is kept in the School of Chemistry and Environmental Studies, Faculty of Applied Sciences at the University Teknologi MARA (UiTM) Kota Samarahan, Sarawak, Malaysia.

Extraction and isolation

The air-dried root (722.14 g) of *Artocarpus odoratissimus* was extracted by a cold extraction method for 24 hours and repeated thrice using n-hexane, ethyl acetate and methanol as solvents. The removal of solvents under reduced pressure gave hexane (AORH), ethyl acetate (AOEA), and methanol (AORM) crude extracts respectively. The crude extracts were fractionated using silica gel vacuum liquid chromatography (VLC) with a stepwise gradient

solvents system such as hexane, hexane-ethyl acetate, ethyl acetate-methanol and methanol, followed by column chromatography to give major fractions. Fractions with spots of the same R_f values were combined and rechromatographed in appropriate solvent systems until pure isolates were obtained. The crude ethyl acetate root afforded two known flavonoids namely pinocembrin (**1**) and pinostrobin (**2**) (Figure 1)

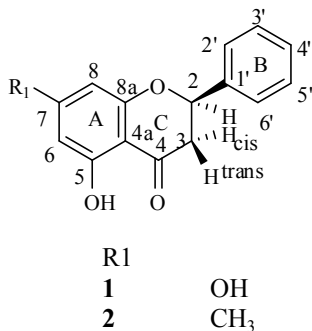


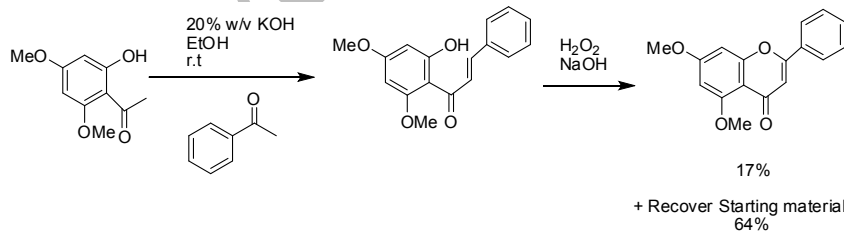
Figure 1. Structure of compounds (**1-2**) isolated from *A. odoratissimus*

Synthesis of pinocembrin and pinostrobin: Synthesis of 2-hydroxy-4,6-dimethoxychalcone via Claisen-Schmidt condensation

20% of aqueous KOH (2 mmol) was added to a solution of benzaldehyde (4 mmol) and 2-hydroxy-4,6-dimethoxyacetophenone (4 mmol) in C₂H₅OH (25 mL). The reaction mixture was stirred at room temperature until completion of reaction (monitored by TLC). The reaction mass was then poured into ice water and neutralized with aqueous 10% HCl solution. The precipitate was filtered, washed with excess of water, dried and recrystallized from methanol to obtain pure 2-hydroxy-4,6-dimethoxy chalcones.

Synthesis of flavonol

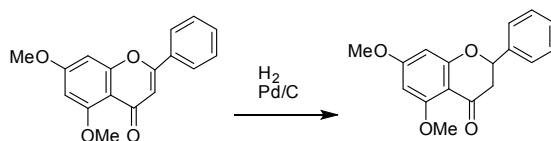
A suspension of a mixture of powdered 2-hydroxy-4,6-dimethoxychalcone (0.10 g, 0.45 mmol) in EtOH, an aq. NaOH solution (8 M, 1.0 ml) and a 30% hydrogen peroxide solution (0.25 g) was stirred at room temperature for 2 hours. The crude product was filtered off, washed with water to give flavonol (scheme 1). Recrystallization of the crude product from MeOH gave pure compound.



Scheme 1. Formation of flavonol

Formation of flavanone via hydrogenation reaction

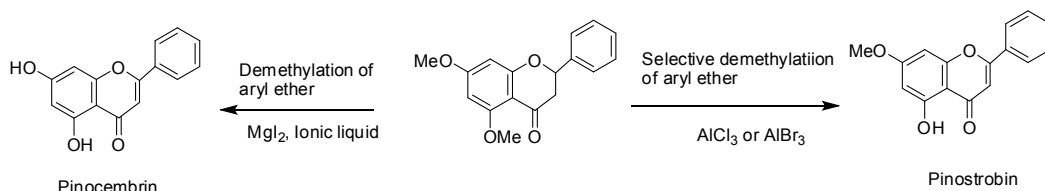
The reaction involves hydrogenation of flavonol to form flavanone (scheme 2).



Scheme 2. Formation of flavanone

Demethylation of aryl methyl ethers

To a solution of aryl methyl ether (1.0 mmol) in [BMIM]BF₄ (2 mL) was added magnesium iodide (0.42 g, 1.5 mmol). The reaction mixture was stirred at 50 °C for 4 – 7 hours. After addition of 5% aqueous sodium thiosulfate (10 mL), the product was extracted with ether (2 × 10 mL), washed with brine (5 mL), and extracted with ether (2 × 10 mL), washed with brine (5 mL), and dried over MgSO₄. The solvent was removed in vacuo and the crude mixture was purified by silica gel chromatography using ethyl ether: hexane (1:3) as eluent to give the desired phenolic product (scheme 3).



Scheme 3. Formation of phenolic product

Results and Discussion

The extraction and separation from the root extract of *Artocarpus odoratissimus*, afforded two flavonoids, which were identified and characterized as pinocembrin (**1**) and pinostrobin (**2**) based on data comparison literature.

Characterization study: Pinocembrin (**1**)

White yellowish crystalline; mp 185-186 °C; IR (KBr) $\nu_{\max}$ cm⁻¹: 3435 (Ar-H stretching), 1744 (C=O ester), 1497, 1578 (C=C Aromatic), 1248 (C-O) and 976 (=C-H) cm⁻¹; EI-MS m/z (% rel. int.): 256 (100) [M⁺ C₁₅H₁₂O₄], 238 (14), 179 (70), 152 (80), 124 (45), 104 (18), 77 (20), 69 (20) and 51(9); ¹H NMR (400 MHz; CDCl₃): δ 2.86 (3-cis), 3.10 (3-trans), 5.43 (2H, t, J=6.7), 6.03 (H-6), 6.03 (H-8), 7.41-7.49 (H-2'-H6'), 12.07 (C-5-OH). ¹³C NMR (100 MHz; CDCl₃): δ 43.37 (C-3), 79.18 (C-2), 95.56 (C-8), 96.81 (C-6), 102.97 (C-4a), 126.16 (C-2', C-6'), 128.87 (C-3', C-4', C-5'), 138.42 (C-1'), 163.11 (C-8a), 164.34 (C-5), 195.65 (C-4).

Pinostrobin (**2**)

Colorless crystal; mp 100-102 °C; IR (KBr) $\nu_{\max}$ cm⁻¹: 3467 (Ar-H stretching), 1741 (C=O ester), 1497, 1578 (C=C Aromatic), 1538, 1360 (C-O) and 968 (=C-H) cm⁻¹; EI-MS m/z (% rel. int.): 270 (90) [M⁺ C₁₆H₁₄O₄], 269 (50), 252 (8), 193 (100), 166 (80), 138 (50), 110 (18), 95 (35), 78 (15) and 51 (20); ¹H NMR (400 MHz; CDCl₃): δ 2.81 (3-cis), 3.86 (C-7-OMe), 3.10 (3-trans), 5.43 (2H, t, J=6.7), 6.03 (H-8), 6.04 (H-6), 7.28-7.49 (H-2'-H6'), 12.07 (C-5-OH). ¹³C NMR (100 MHz; CDCl₃): δ 43.36 (C-3), 55.60 (C-7-OMe), 79.25 (C-2), 95.49 (C-6, C-8), 103.20 (C-4a), 126.17 (C-2', C-6'), 128.92 (C-3', C-4', C-5'), 138.30 (C-1'), 163.16 (C-5), 164.36 (C-7), 164.63 (C-8a), 195.79 (C-4).

Compound (**1**) was obtained as white yellowish crystalline with melting point 185 – 186 °C. The mass spectrum showed the molecular ion peak at m/z 256, indicated the formal formula C₁₅H₁₂O₄. In the ¹H NMR, one proton singlet at δ 12.07 sited the free hydroxyl at position 5 with H-bridge to O at the carbonyl at position 4. Two doublets at δ 6.03 attributed to meta H-6 and H-8. A five-proton multiplet at δ 7.41 – 7.49 confirmed the non-substitution of the B-ring. Two doublet-doublet of one proton each at δ 2.86 and δ 3.10 attributable to H-3 (*cis*) and H-3(*trans*), respectively. The ¹³C NMR spectrum showed 15 carbons signals including one methylene, eight methine and six quaternary carbon atoms.

Compound (**2**) was obtained as colorless crystal with melting point 100 – 102 °C. The mass spectrum showed the molecular ion peak at m/z 270 indicating that the compound had the formula C₁₆H₁₄O₄. The IR spectrum showed the presence of aromatic stretch at 3467 cm⁻¹, carbonyl group, ester (C=O) at 1741 cm⁻¹ as well as an olefin functional group (C-O) at 1538 cm⁻¹. In the ¹H NMR, a methoxy group was seen as a singlet at δ 3.86. A one-proton singlet at δ 12.07 sited the free hydroxyl at position 5 with H-bridge to O at the carbonyl at position 4. The ¹³C NMR

spectrum showed 16 carbons signals including one methyl, one methylene, 8 methine and 6 quaternary carbon atoms. Among these signals the carbon signal of methoxy group is clearly assigned due to the presence of chemical shift 55.60.

The Claisen-Schmidt condensation of 2-hydroxy-4,6-dimethoxyacetophenone with benzaldehyde under basic condition (KOH 50%) in ethanol proceeded smoothly to furnish 2-hydroxy-4,6-dimethoxychalcone in excellent yield, (97.1%). The cyclisation of 2-hydroxy-4,6-dimethoxychalcone in ethanol and aqueous sodium hydroxide solution and a 30% hydrogen peroxide solution gave flavonol and the hydrogenation of flavonol produced flavanone. The reaction of aryl methyl ether and magnesium iodide gave pinocembrin and with aluminum trichloride gave pinostrobin as product.

Conclusion

The isolation of *Artocarpus odoratissimus* root extract had successfully led to two known flavonoids, namely pinocembrin and pinostrobin. These compounds were then successfully synthesized with 2-hydroxy-4,6-dimethoxyacetophenone and benzaldehyde as starting materials. The identification of compounds was carried out using spectroscopic methods: Infrared (IR) spectroscopy, Nuclear Magnetic Resonance (NMR) spectroscopy and Gas Chromatography Mass Spectroscopy (GC-MS) and a comparison with the literature data of compound.

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