



Natural Product Research

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Subject Area and Category [Agricultural and Biological Sciences](#)
[Plant Science](#)

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[Chemistry](#)
[Analytical Chemistry](#)
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Publisher [Taylor & Francis](#)

Publication type Journals

ISSN 14786419

Coverage 2003-ongoing

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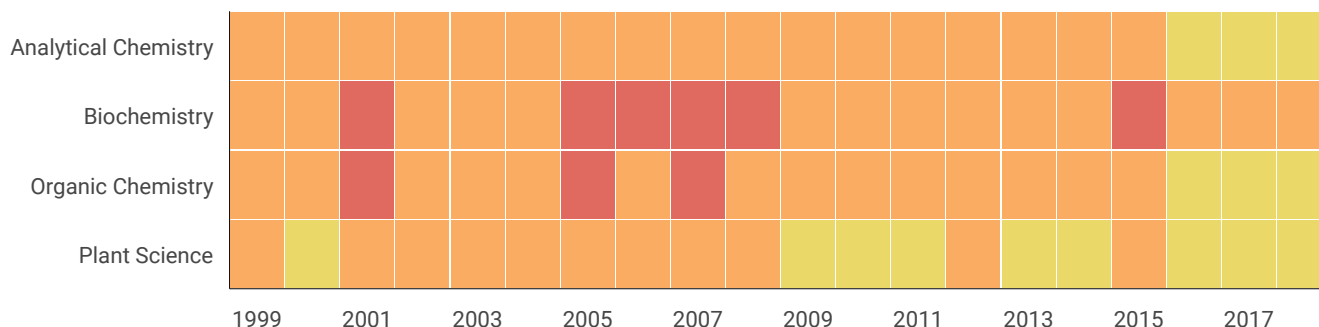


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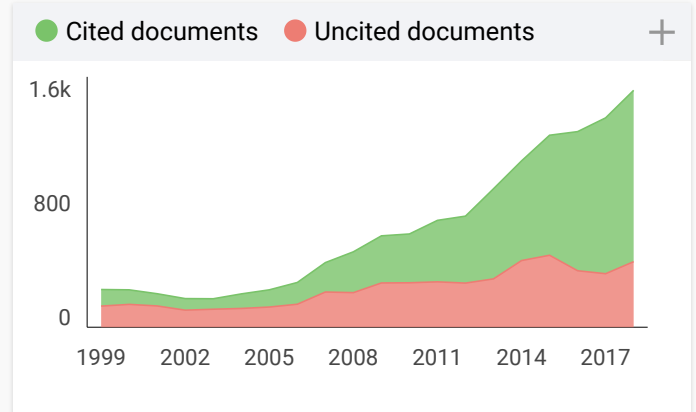
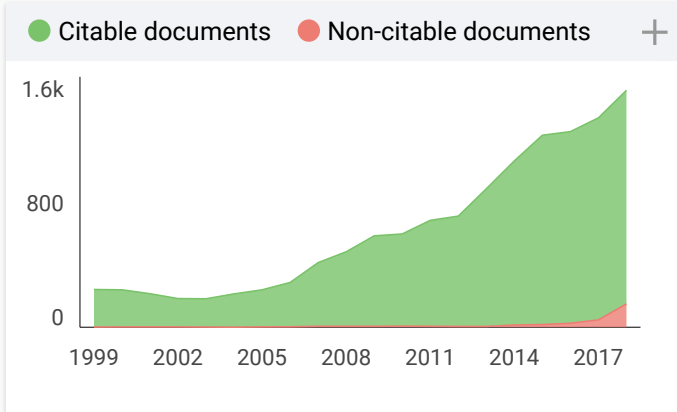
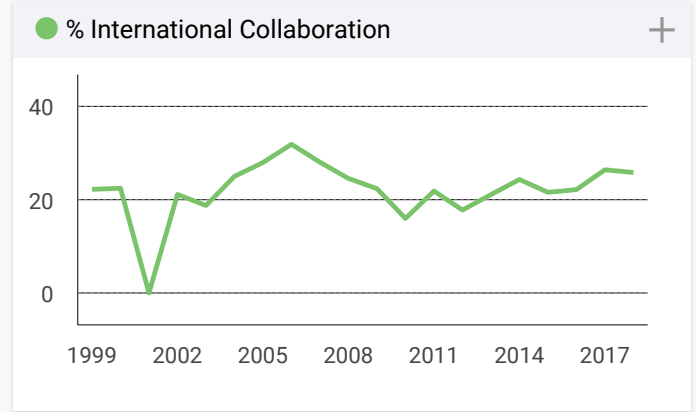
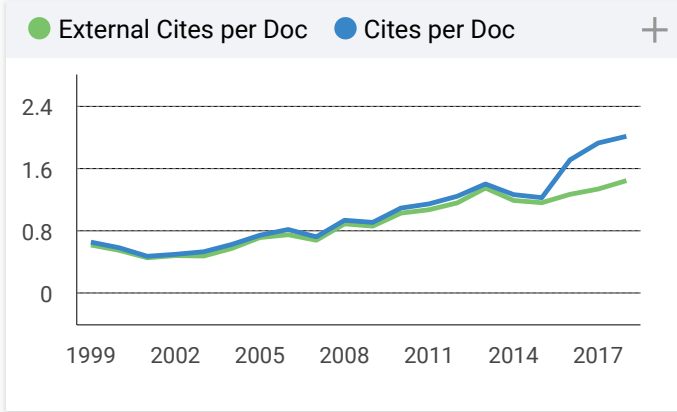
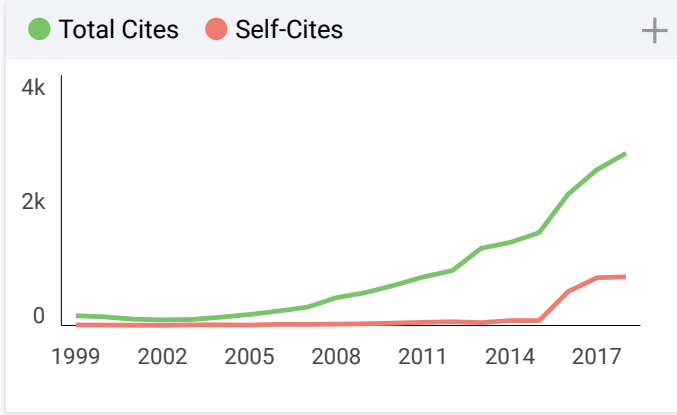
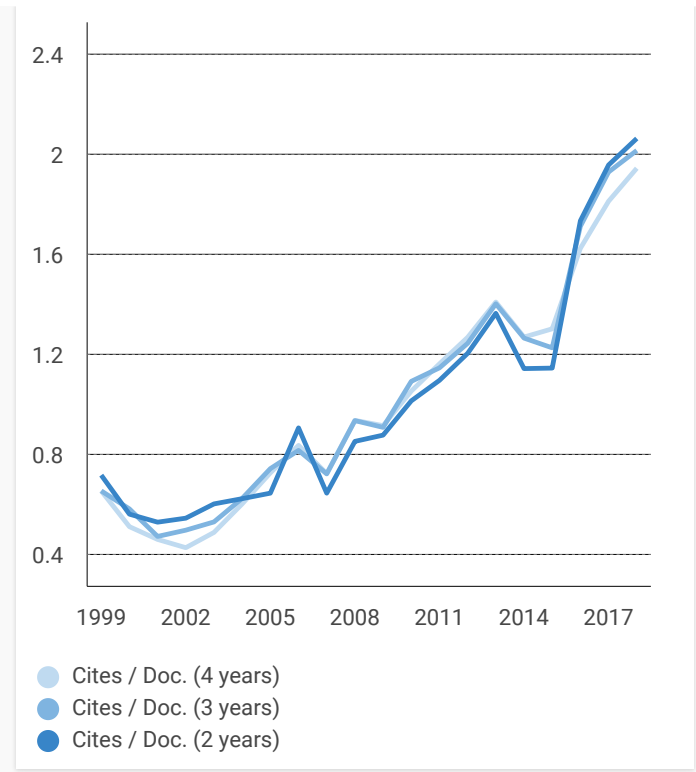
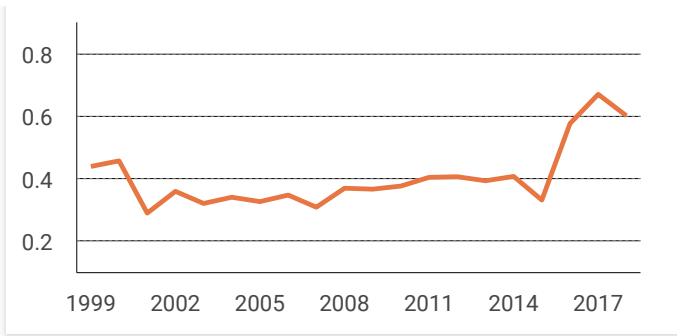


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Aims and scope

The aim of **Natural Product Research** is to publish important contributions in the field of natural product chemistry. The journal covers all aspects of research in the chemistry and biochemistry of naturally occurring compounds.

The communications include coverage of work on natural substances of land and sea and of plants, microbes and animals. Discussions of structure elucidation, synthesis and experimental biosynthesis of natural products as well as developments of methods in these areas are welcomed in the journal. Finally, research papers in fields on the chemistry-biology boundary, eg. fermentation chemistry, plant tissue culture investigations etc., are accepted into the journal.

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Publication history

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Formerly known as

Natural Product Letters (1992 - 2002)



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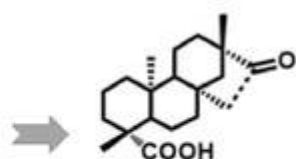
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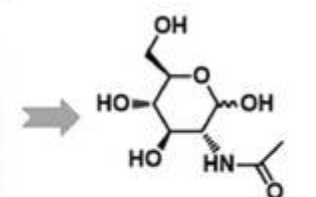
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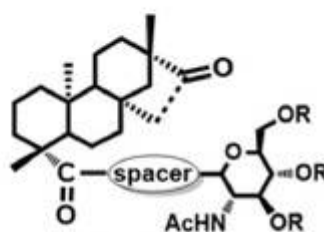
isosteviol



Chitin



N-acetyl-glucosamine



R = Ac or H

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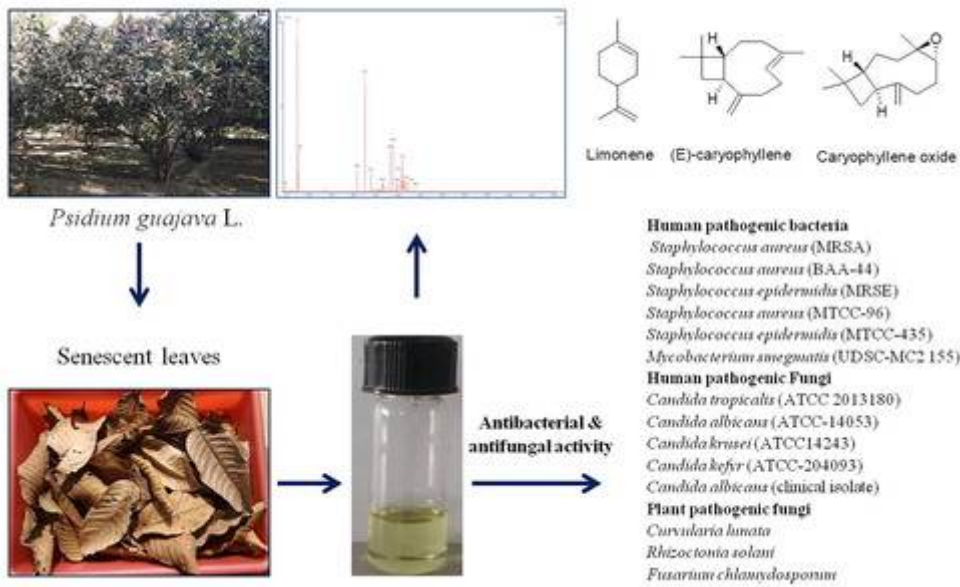
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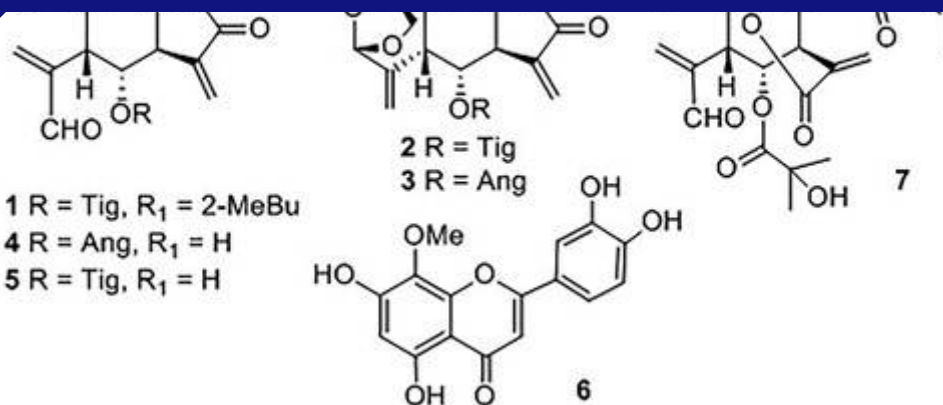
Elemenolides from *Zinnia peruviana* and evaluation of their antibacterial and α -glucosidase inhibitory activities >

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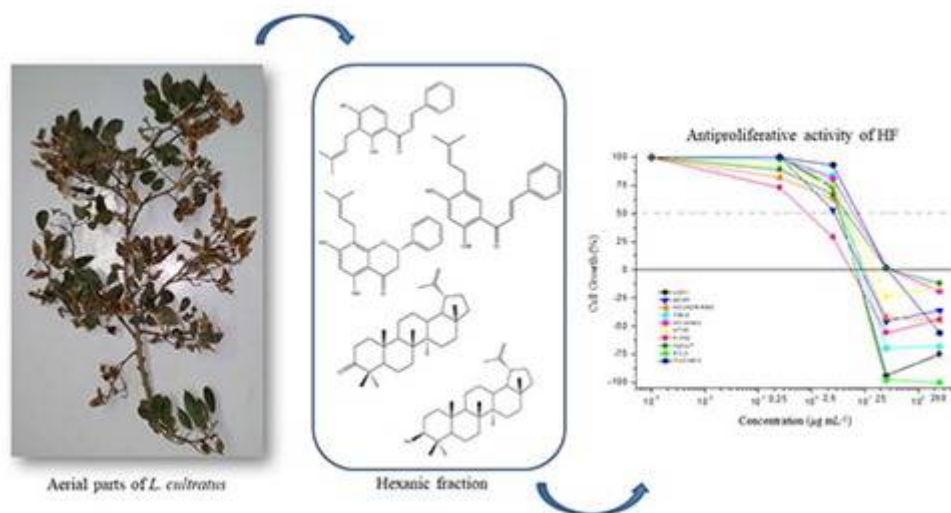
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Emanuelle M. B. M. da Silva Landim, Ana Lúcia T. G. Ruiz, João E. de Carvalho, Armando M. Pomini, Lindamir H. Pastorini & Silvana M. Oliveira Santin

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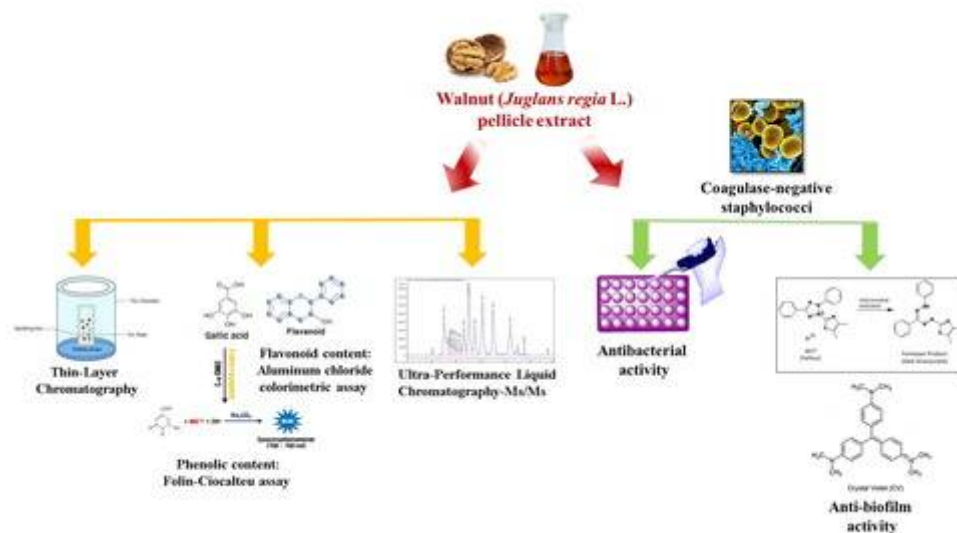
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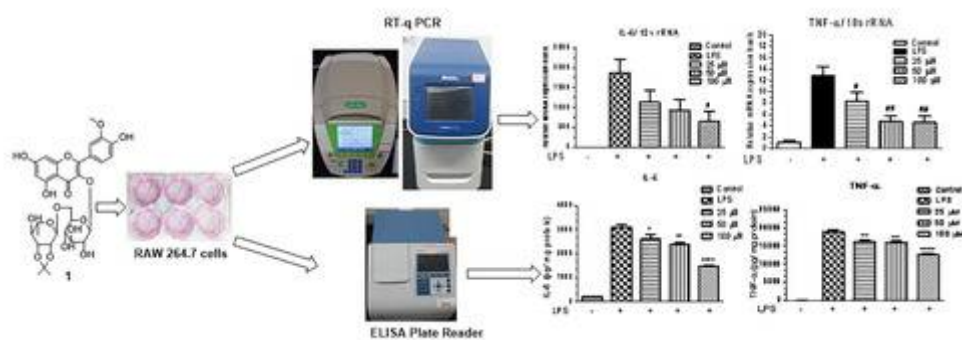
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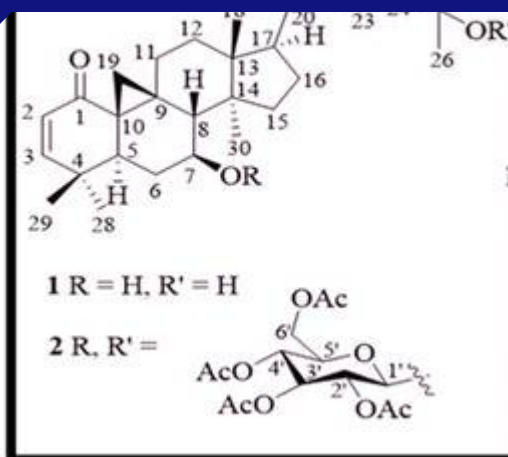
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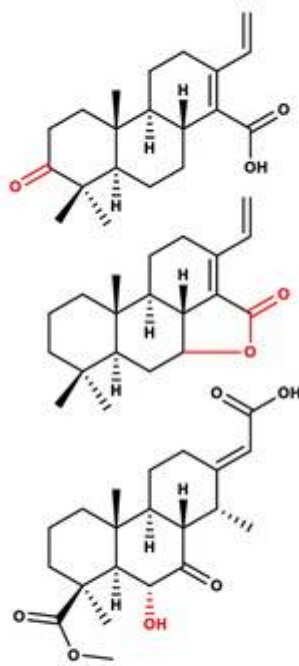
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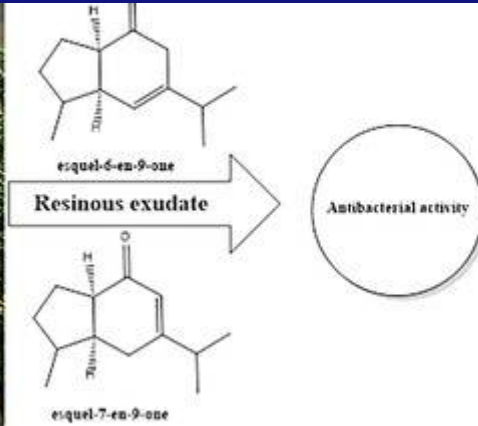
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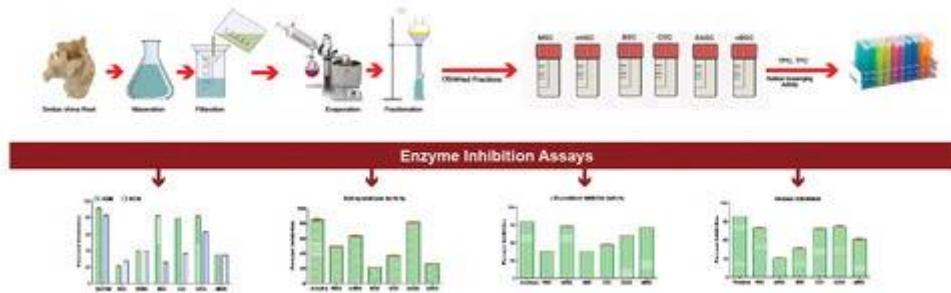
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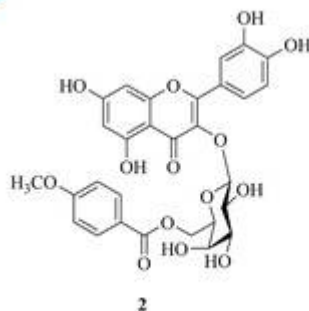
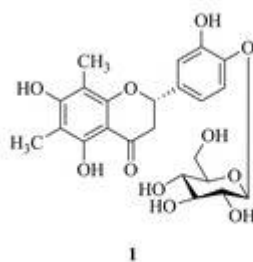


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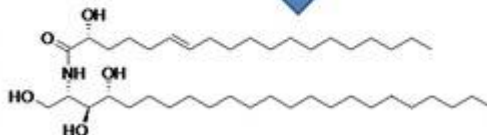
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Jean Emmanuel Mbosso Teinkela, Xavier Siwe Noundou, Simone Fannang, Achille Mbem Song, Jules Clément Assob Nguedia, Heinrich C. Hoppe & Rui Werner Maçedo Krause

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antimicrobial, antiplasmodial, antitrypanosomal and cytotoxic activities

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Chimengul Turghun, Mahinur Bakri, Ge-Yu Liu, Khayrulla Bobakulov & Haji Akber Aisa

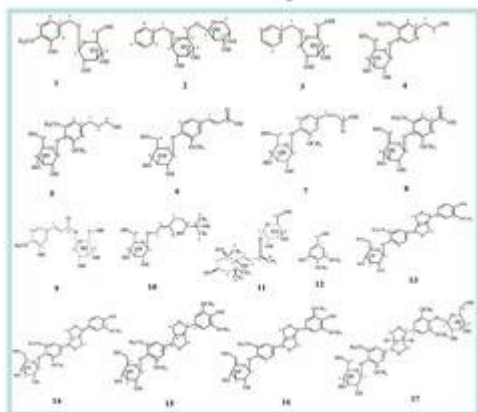
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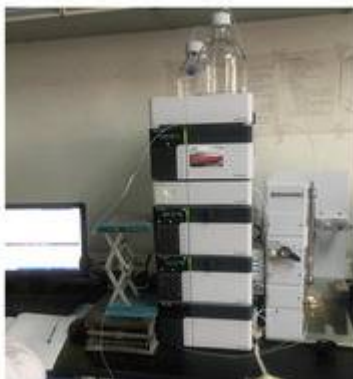
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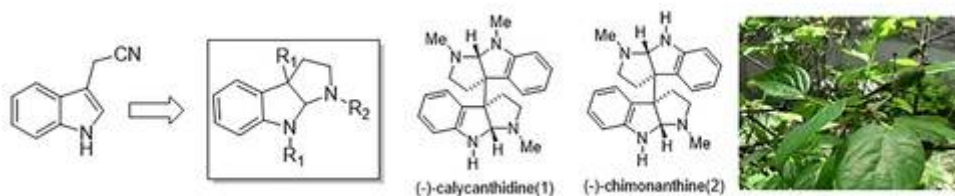
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Shaojun Zheng, Rui Zhu, Bing Tang, Lizhuang Chen, Hongjin Bai & Jiwen Zhang

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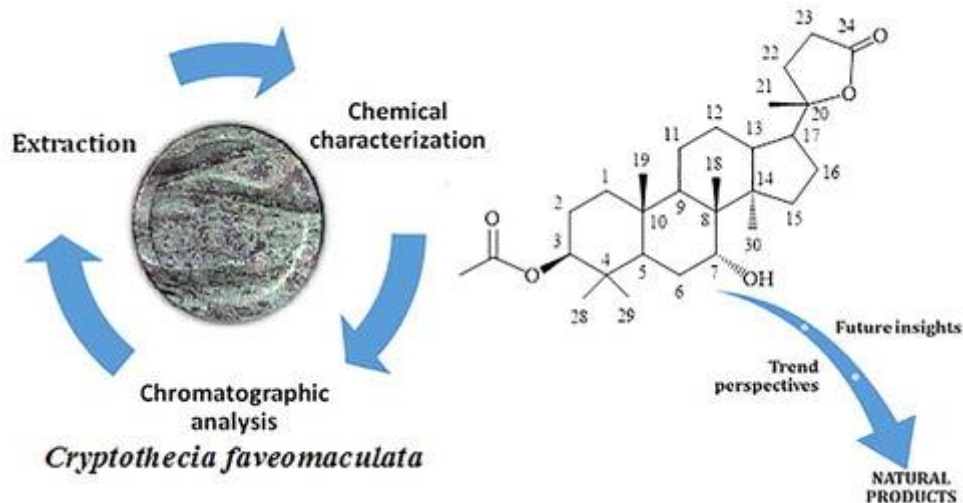
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Cryptothecia faveomaculata Makhija & Patw >

Nguyen Ngoc Tuan, Ping-Chung Kuo, Tran Trung Hieu, Le Nguyen Tuong Vi, Quach Tong Hung, Le Tien Dunge, Nguyen Duy Trinh, Nguyen Quang Trung, Nguyen Cuu Khoa, Ha Viet Hai & Tran Dinh Thang

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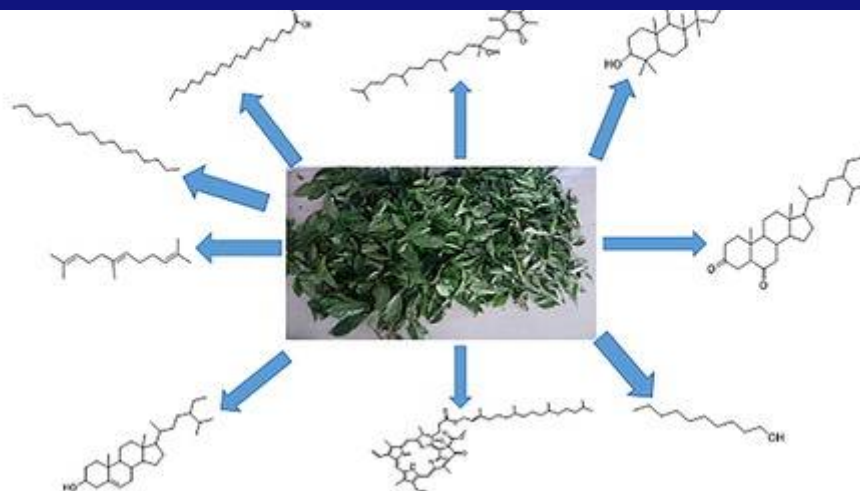
Chemical constituents from the leaves of *Elaeocarpus floribundus* >

Ayorinde Victor Ogundele & Archana Moni Das

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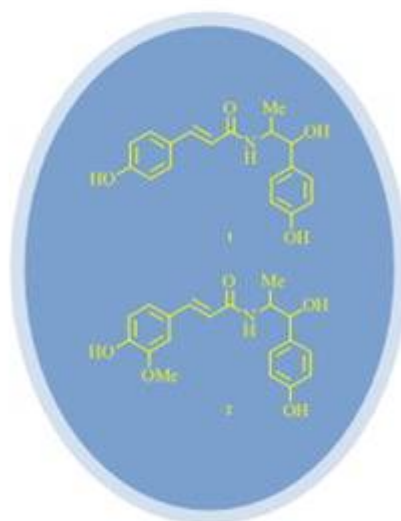
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Stixilamides A and B, two new phenolic amides from the leaves of *Stixis suaveolens* >

Quoc Anh Ngo, Thi Yen Tran, Thuy Hang Nguyen, Van Tuyen Nguyen, Hong Anh Duong & Hung Viet Pham

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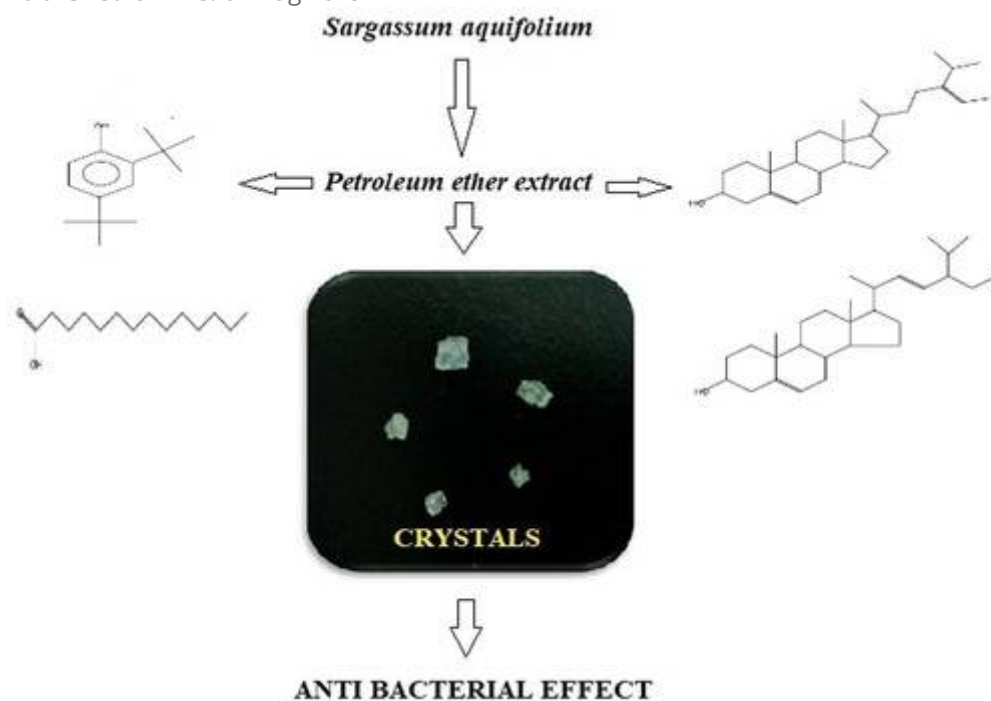


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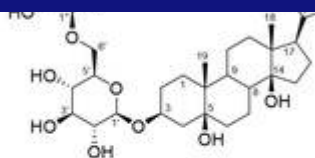
A new cardenolide glycoside from the roots of *Streptocaulon juvenas* (lour.) merr. (Asclepiadaceae) >

Xuan-Hao Bui, Phu Hoang Dang, Tuan Trong Vo, Nhi Y Thi Nguyen, Minh-Duc Nguyen & Quan Le Tran

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Periplogenin 3-O- β -gentiobioside

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1	71.87	60.16	47.72

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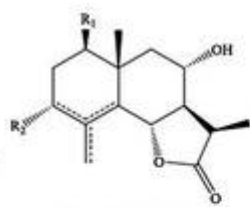
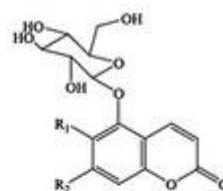
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New antimicrobial metabolites from the medicinal herb *Artemisia herba-Alba* >

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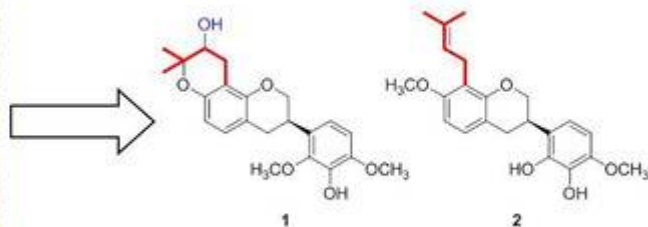
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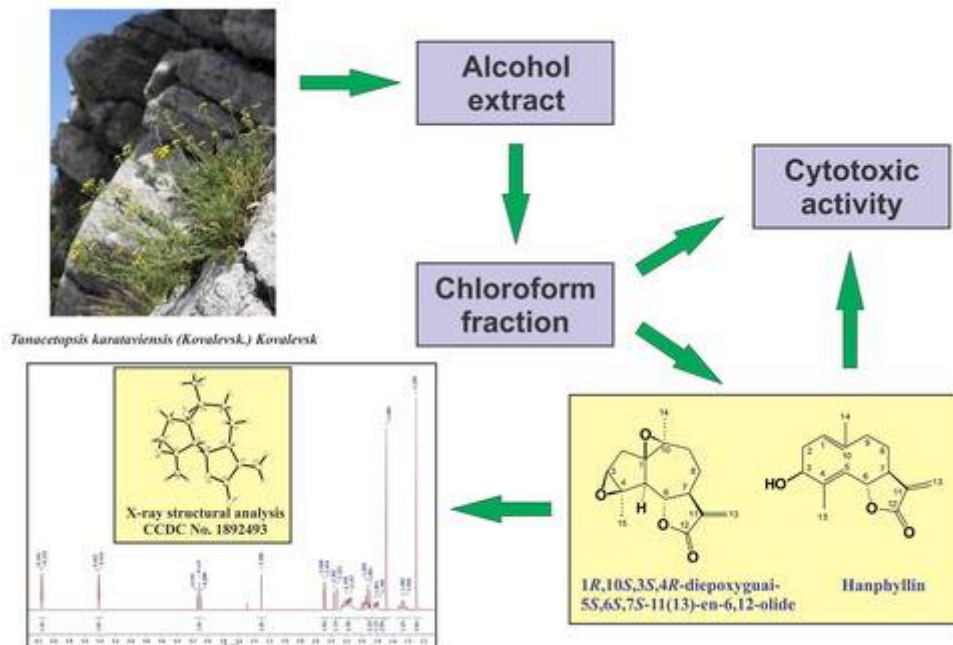
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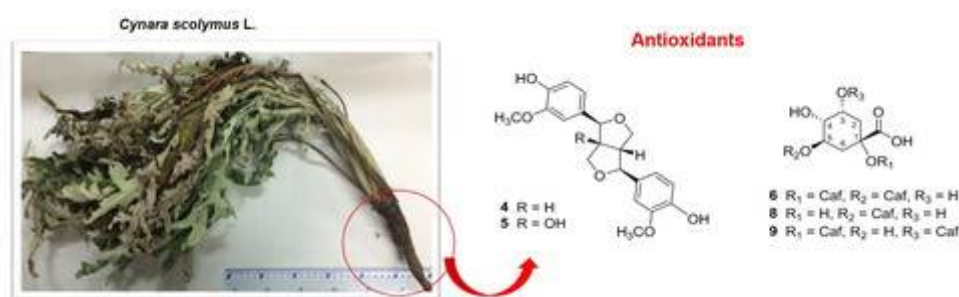
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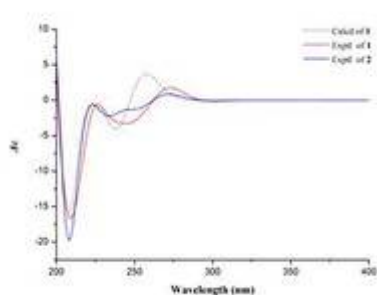
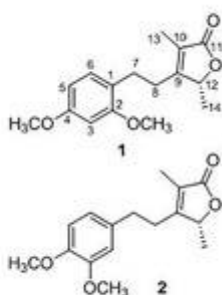
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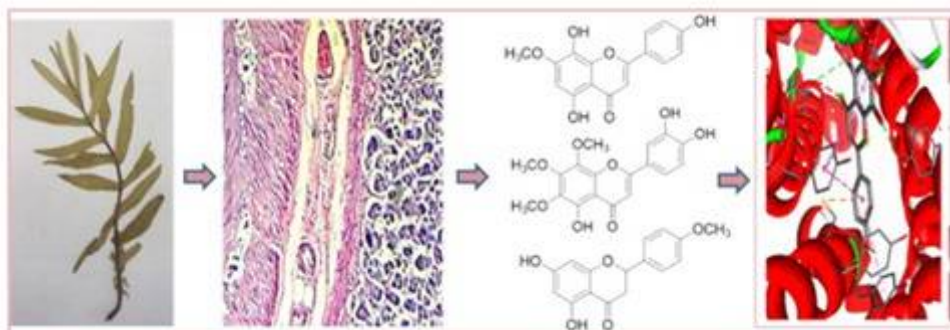
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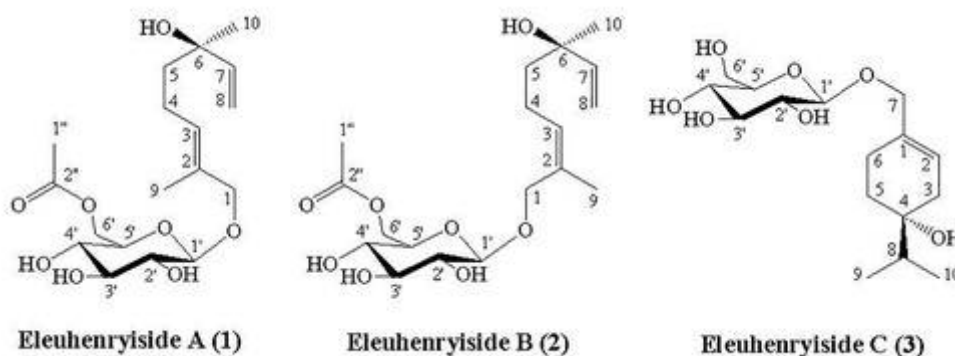
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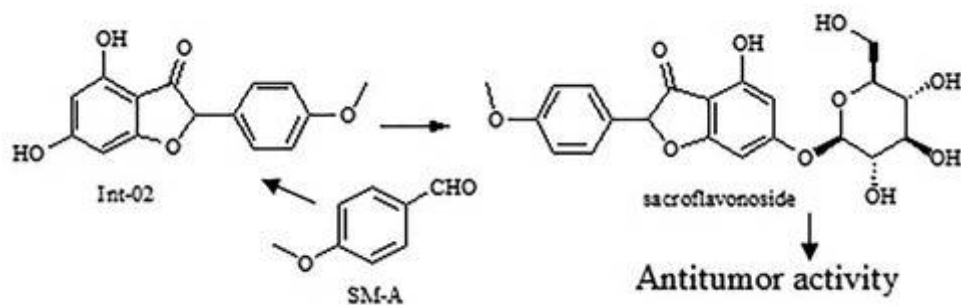
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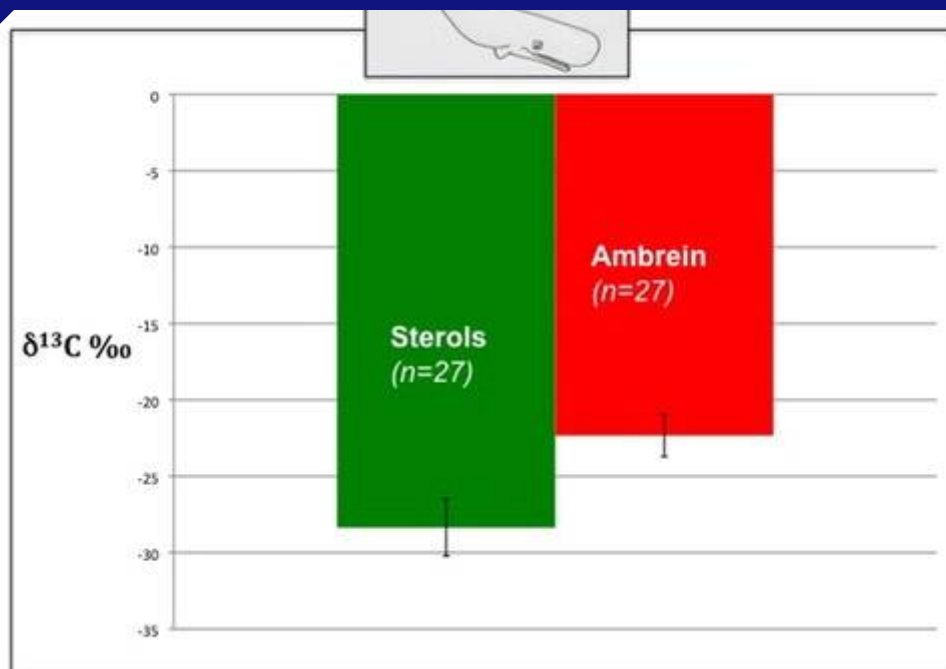
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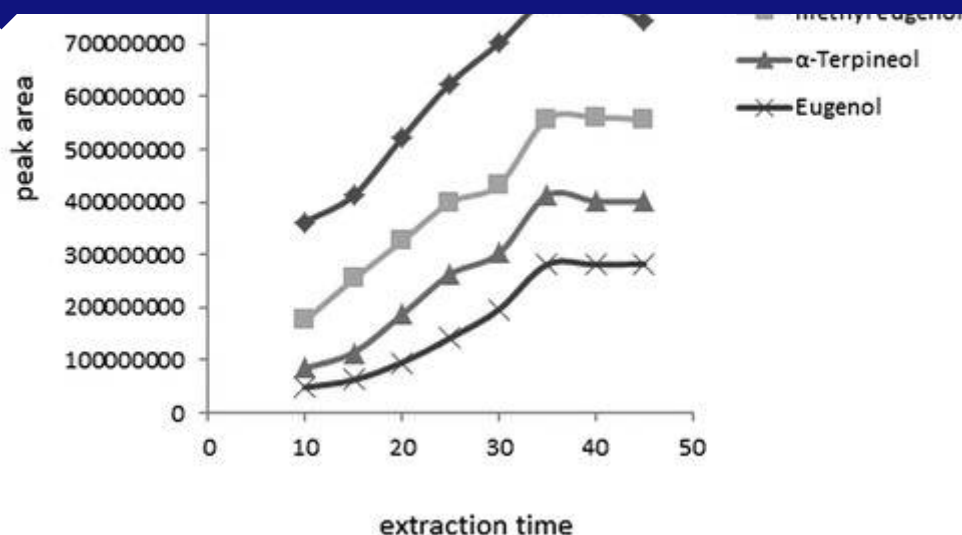
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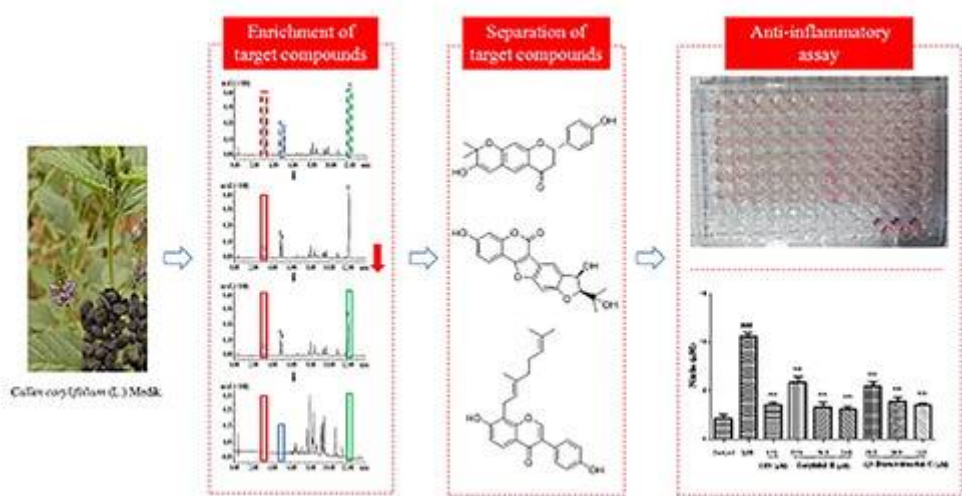
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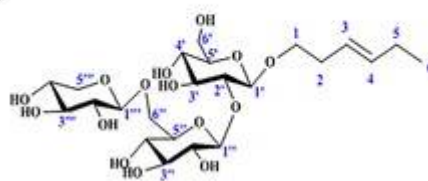
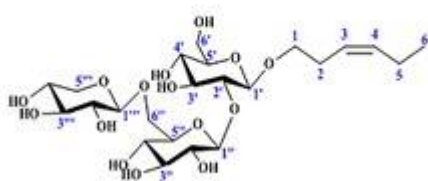
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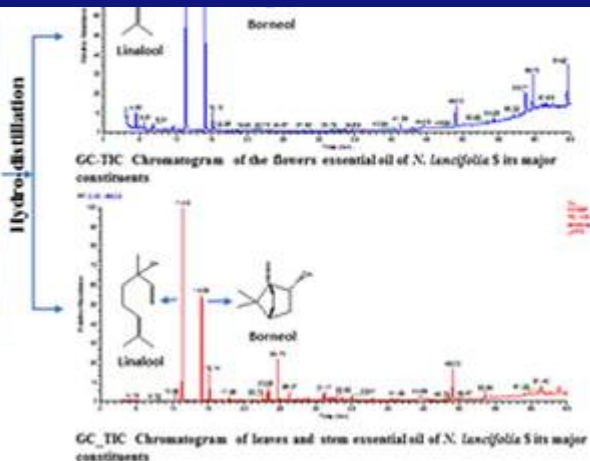
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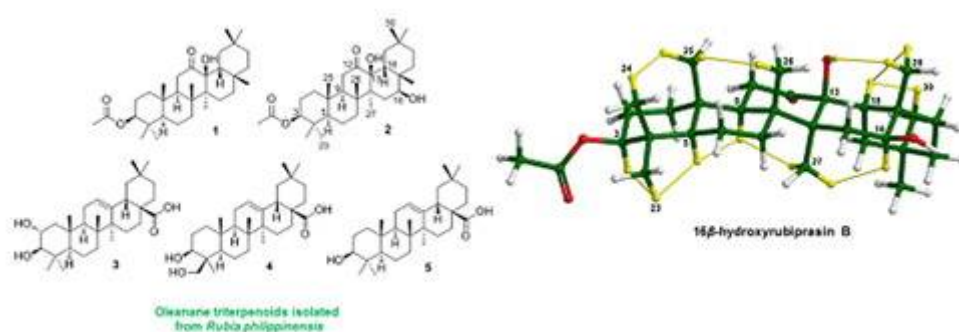
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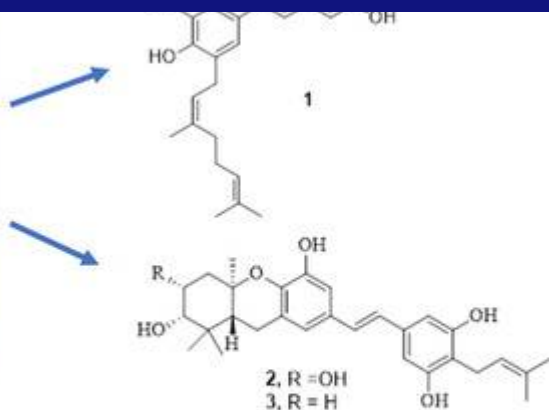
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Compound	MNTC	CC ₅₀	IC ₅₀			SI	
			E7	E13	E19	E7	E19
Mappain (1)	10 μM	3.24 μM	1.23 μM	NA	0.24 μM	2.6	13.5
Vedelianin (2)	10 nM	0.78 nM	0.025 nM	NA	0.0036 nM	31.2	216.7
Schweinfurthin G (3)	10 nM	0.82 nM	0.043 nM	NA	0.018 nM	19.1	45.6

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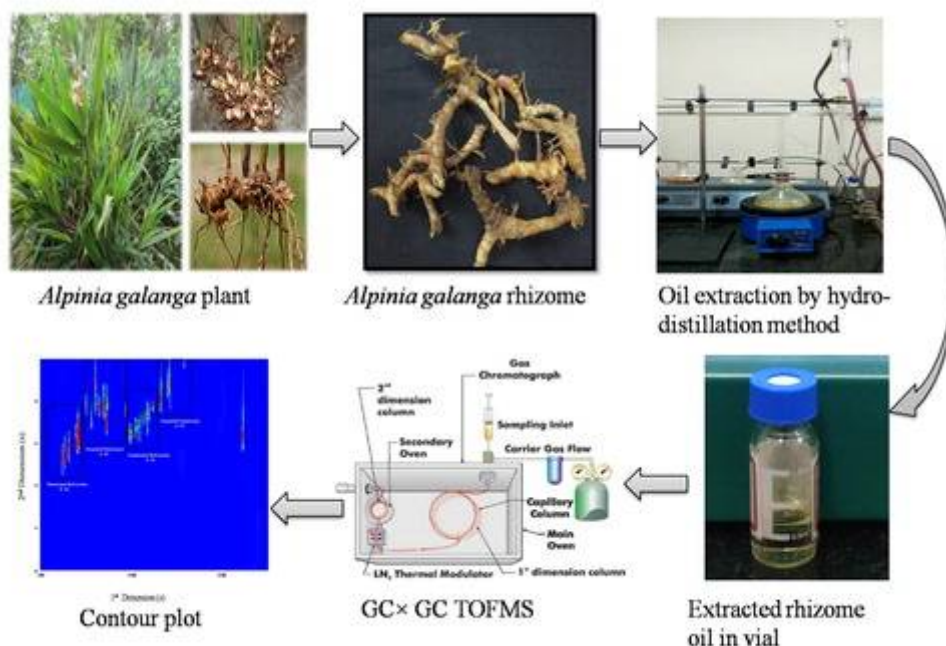
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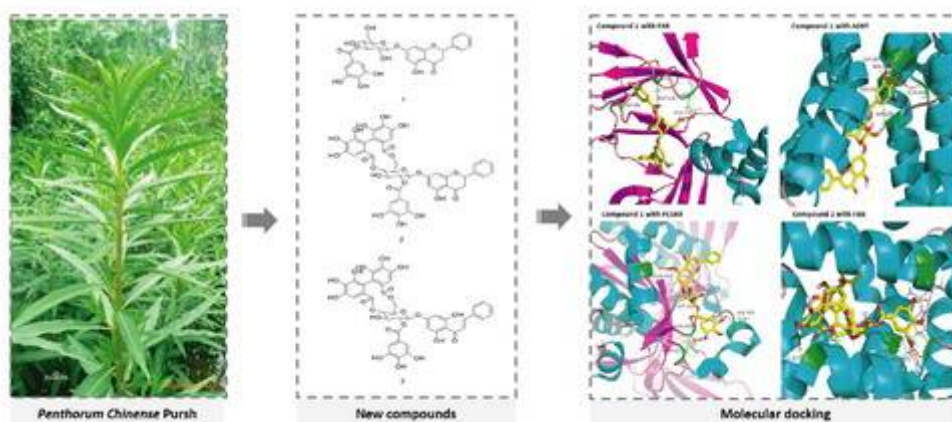
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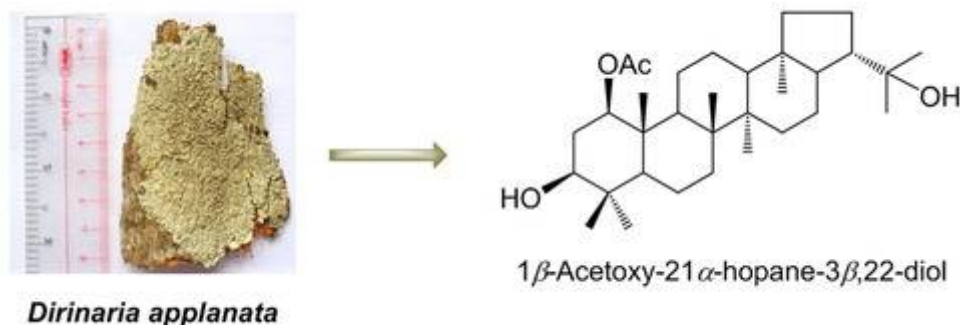
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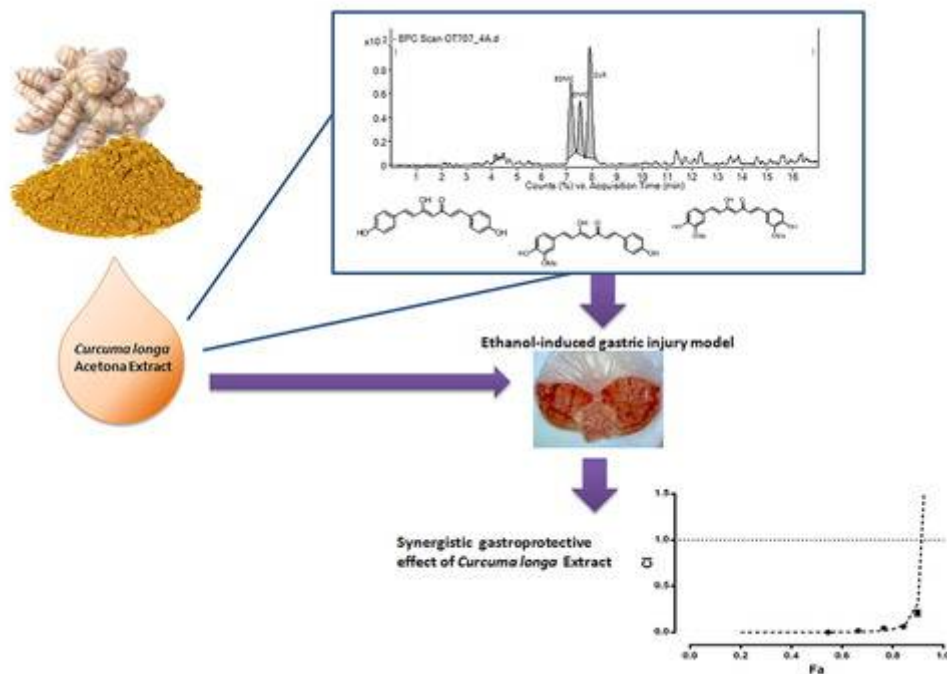
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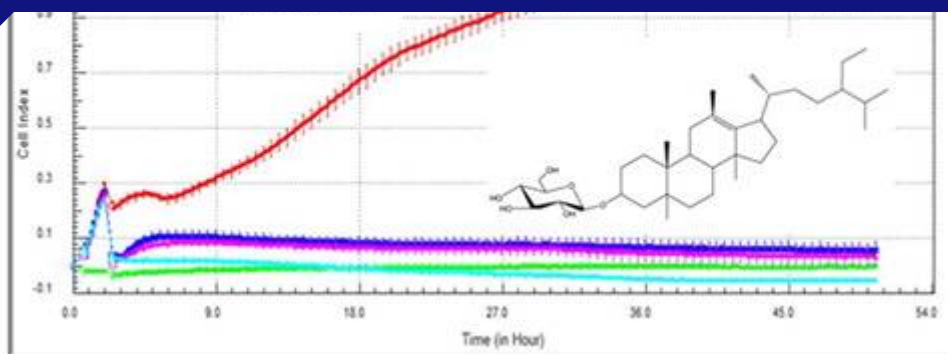
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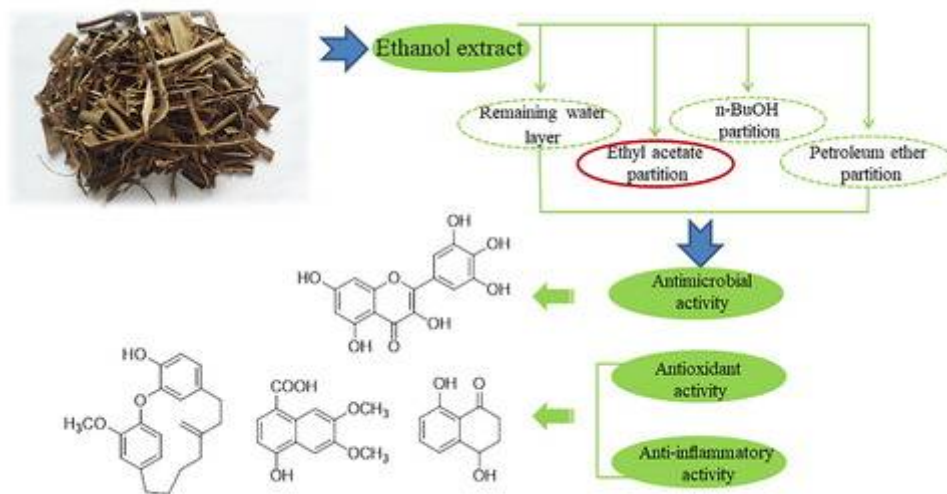
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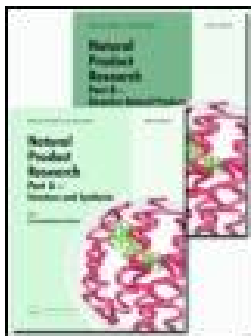
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Ratih Dewi Saputri, Tjitjik Srie Tjahjandarie & Mulyadi Tanjung


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Two novel coumarins bearing an acetophenone derivative from the leaves of *Melicope Quercifolia*

Ratih Dewi Saputri, Tjitjik Srie Tjahjandarie and Mulyadi Tanjung

Natural Products Chemistry Research Group, Organic Chemistry Division, Faculty of Science and Technology, Department of Chemistry, Universitas Airlangga, Surabaya, Indonesia

ABSTRACT

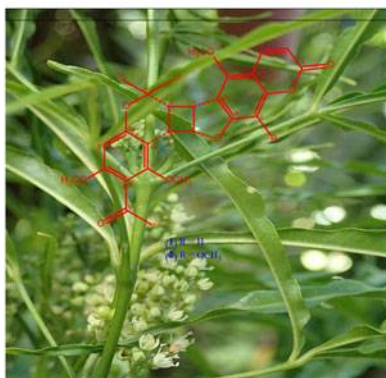
Meliquercifolins A (**1**), and B (**2**), two new coumarins bearing an acetophenone derivative were isolated from the leaves of *Melicope quercifolia* along with three known compounds, melicodenines E (**3**), F (**4**) and I (**5**). Structures of two new compounds were identified based on spectroscopic analyses (UV, HR-ESI-MS, 1D and 2D NMR). Cytotoxic activities of compounds (**1–5**) towards three human cancer cells (HeLa, MCF-7, P-388), compounds **1**, **4** and **5** showed very potent activity against HeLa cells with IC₅₀ values 2.6; 0.8; 1.1 μM, respectively.

ARTICLE HISTORY

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KEYWORDS


Cytotoxicity; *Melicope quercifolia*; Meliquercifolins A and B



1. Introduction

Melicope quercifolia (Rutaceae) is a small tree and found as an endemic plant in West Java, Indonesia. The leaves of *M. quercifolia* have been used to treat skin diseases (Appelhans et al. 2018). Acetophenones (Nguyen et al. 2016), alkaloids (George et al. 2017), coumarins (Xu et al. 2016), flavonoids (Saputri et al. 2018), and phenylpropanoids (Nakashima et al. 2012) are phenolic compounds from the *Melicope* plants. Some of the phenolic compounds showed various Diels-Alder adduct and [2 + 2]

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cycloaddition (Nakashima et al. 2012). Melicodenines E (3), F (4) and I (5) formed a [2 + 2] cycloaddition type, and a Diels-Alder adduct type. In the ongoing phytochemical investigation of *M. quercifolia*, we report two new compounds, meliquercifolins A (1), and B (2) from the leaves of *M. quercifolia*. These structures of two new compounds are coumarins bearing an acetophenone moiety. Heterodimer compounds (1 and 2) are the first time found skeleton structure from *Melicope*. The cytotoxic properties against three human cancer cells (P-388, MCF-7, and HeLa) of isolated compounds will also report.

2. Result and discussion

Meliquercifolin A (1) obtained a colorless oil in which showed an optical rotation was inactive. Based on the HRESIMS measurement of 1 gave a positive ion peak $[M + H]^+$ at m/z 479.1729 correspondings for a molecular formula $C_{27}H_{27}O_8$. The IR absorptions of 1 displayed vibrations the presence of conjugated C=O (1636 cm^{-1}), aromatic (1608 and 1579 cm^{-1}) and ether (1119 cm^{-1}) groups. Two maximum absorptions at λ_{max} 280 (3.86) and 332 (3.88) nm very like with benzoyl and cinnamoyl chromophore. Four signals of a 1,2,3,4-tetrasubstituted cyclobutane ring showed at δ_{H} 3.05 (1H, dd, $J=9.8$; 5.8 Hz, H-3), δ_{H} 3.96 (1H, t, $J=2.2$ Hz, H-4), δ_{H} 4.71 (1H, t, $J=6.9$ Hz, H-6) and δ_{H} 5.42 (1H, ddd, $J=8.2$; 6.2; 2.2 Hz, H-7). The signal at δ_{H} 5.42 implied an oxymethine attached in the cyclobutane ring. Compound 1 also demonstrated the presence of two proton signals of two aromatics at δ_{H} 5.98 (1H, s, H-8), δ_{H} 6.21 (1H, s, H-9) and a pair of *cis* vinylic of chromen-2-one ring at δ_{H} 6.01 (1H, d, $J=9.8$ Hz, H-3), δ_{H} 7.82 (1H, d, $J=9.8$ Hz, H-4). Based on the ^1H NMR spectrum, compound 1 likewise showed the presence of three methyl signals at δ_{H} 2.17 (3H, s, H-10), δ_{H} 1.19 (3H, s, H-11), δ_{H} 1.61 (3H, s, H-12) as well as three methoxyl signals at δ_{H} 4.04 (3H, s, 5-OCH₃), δ_{H} 3.67 (3H, s, 5'-OCH₃), and δ_{H} 3.59 (3H, s, 7'-OCH₃). The ^{13}C NMR analysis (APT experiment), compound 1 showed 27 carbon signals that are completely separated, consisting of six methyl carbons, eight methine carbons, and 13 quaternary carbons. Three of them (δ_{C} 161.9; δ_{C} 139.5 and δ_{C} 109.7) are characteristic of C-2, C-3, and C-4 of the chromen-2-one ring (coumarin). Furthermore, two signals of the acetyl group (δ_{C} 32.3; δ_{C} 202.5) are characteristic of the acetophenone structure (Nakashima et al. 2012). The signal of a *cis* vinylic of chromen-2-one at δ_{H} 7.82 (H-4) in the HMBC spectrum exhibited correlation with two oxyaryl carbons, C-5 (δ_{C} 152.7), C-9a (δ_{C} 156.7) and a lactone carbonyl at C-2 (δ_{C} 161.9) as well as the methoxyl signal at δ_{H} 4.04 correlated to C-5 (δ_{C} 152.7) unambiguously located the methoxyl group at C-5. Another proton signal of a *cis* vinylic at δ_{H} 6.01 (H-3) correlated to C-2 (δ_{C} 161.9) and a quaternary carbon at C-4a (δ_{C} 108.1). Furthermore, a signal of aromatic at δ_{H} 6.21 correlated to two oxyaryl carbons at C-9a (δ_{C} 156.7); C-8a (δ_{C} 168.5) and two quaternary carbons at C-4a (δ_{C} 108.1); C-5a (δ_{C} 104.7) showed that the signal of δ_{H} 6.21 at H-9. A signal of an oxymethine at δ_{H} 5.42 correlated to C-8a (δ_{C} 168.5), and C-4' (δ_{C} 34.7) revealed that a part of the heterodimer is bergapten (coumarin) (Nakashima et al. 2012). The HMBC spectrum, a signal of methine at δ_{H} 4.71 (H-6) correlated to C-3' (δ_{C} 45.5), C-4' (δ_{C} 34.7) and C-8a (δ_{C} 168.5) reinforced the structure of the bergapten experiencing a [2 + 2] cycloaddition reaction on furano ring (Nakashima et al. 2012). A methine signal (cyclobutane ring) at

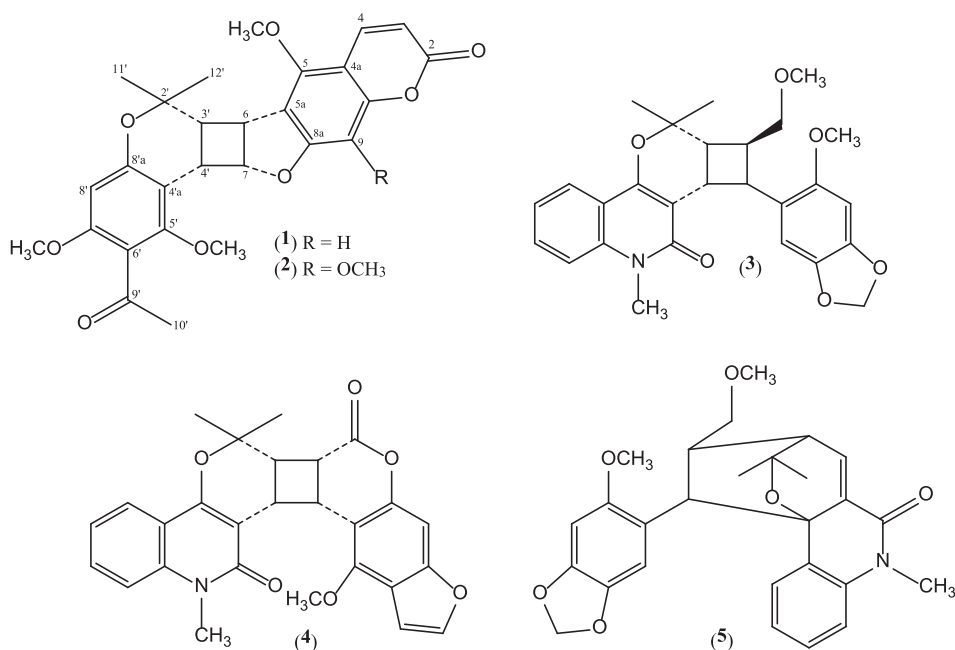


Figure 1. Chemical structures of isolated compounds from *M. quercifolia*.

δ_{H} 3.96 (H-4) correlated to δ_{C} 44.1 (C-6), δ_{C} 84.9 (C-7), δ_{C} 156.4 (C-8a) and δ_{C} 106.6 (C-4a). Two methyl signals of pyrano ring at δ_{H} 1.19 (H-11) and δ_{H} 1.61 (H-12) correlated to a methine carbon at C-3' (δ_{C} 45.5) and an ether cyclic carbon at C-2' (δ_{C} 74.0). Besides, a signal of aromatic isolated at δ_{H} 5.98 (H-8) correlated to an oxyaryl carbon at δ_{C} 156.8 (C-7), and two quaternary carbons at δ_{C} 106.6 (C-4a); 117.8 (C-6). A methoxyl signal at δ_{H} 3.67 (7'-OCH₃) correlated to δ_{C} 156.8 (C-7). Another a methoxyl signal at δ_{H} 3.59 (5'-OCH₃) correlated to δ_{C} 155.7 (C-5). A methyl signal of acetyl at δ_{H} 2.17 (H-10) correlated to δ_{C} 202.5 (C-9). These correlations showed that a part of evodionol (Nakashima et al. 2012). This result implied that compound **1** is a reaction of [2 + 2] cycloaddition between bergapten-evodionol. Therefore, the structure of meliquercifolin A established as **1**.

Meliquercifolin B (**2**) also acquired as a colorless oil. The UV absorption bands (λ_{max} 269, 335), and IR (1620, 1601, 1571 and 1110 cm^{-1}) absorptions very semblable with **1**. The molecular formula of compound **2** was assigned to be C₂₈H₂₉O₉ exhibited positive ion peak [M + H]⁺ at m/z 509.1802 (calcd for 509.1812) by HRESIMS measurement. The ¹H and ¹³C NMR spectrum of **2** had very semblable with **1**. The main difference, the ¹H and ¹³C NMR of **2** showed a methoxyl group at C-9 (δ_{H} 3.98; δ_{C} 58.6). The HMBC and HMQC experiments confirmed the location of a methoxyl group at C-9. With the addition, a methoxyl group attached in the aromatic ring of coumarin skeleton indicated is [2 + 2] cycloaddition between isopimpinellin-evodionol. Based on UV, IR, HRESIMS, 1 D, and 2 D NMR measurements, the structure of meliquercifolin B assigned as **2**. The chemical structures of isolated compounds shown in (Figure 1).

Three known compounds, melicodenine E (**3**), melicodenine F (**4**) and melicodenine I (**5**) from HRESIMS, 1 D and 2 D NMR data showed chemical formula, and the chemical

Table 1. Cytotoxic activities of isolated compounds (1–5).

Compound	IC ₅₀ (μM)		
	HeLa	MCF-7	P-388
1	2.6 ± 0.10	>100	78.9 ± 1.20
2	>100	>100	>100
3	>100	>100	11.9 ± 0.42
4	0.8 ± 0.01	>100	38.3 ± 0.87
5	1.1 ± 0.05	>100	>100
Artonin E	–	–	1.33 ± 0.07
Doxorubicin	0.9 ± 0.04	0.8 ± 0.02	–

shift very resemblant with published previously (Nakashima et al. 2012; George et al. 2017). Compound **3** is a form of a [2 + 2] cycloaddition reaction of *N*-methylflindersin-melicodin A, and compound **4** is *N*-methylflindersin-bergapten. Compound **5** is a [4 + 2] cycloaddition (a Diels-Alder adduct type) two *N*-methylflindersins (George et al. 2017; Nakashima et al. 2012).

Those cytotoxic activities of isolated compounds, compounds **1**, **4** and **5** showed high activity against HeLa cells while compounds **2** and **3** were inactive. However, all of the isolated compounds were inactive against MCF-7 and P-388 cells (Table 1). For coumarin bearing an acetophenone derivative, compound **1** more active than compound **2** against HeLa cells. The influence of a methoxyl group at C-9 suggested as a critical element to decrease the cytotoxic effect towards HeLa cells. The melicodenine E (**4**) more active than meliquercifolins A (**1**). However, compound **4** slightly active than compound **5**. The *N*-methylflindersin factor is a crucial element to enhance the cytotoxic effect of HeLa cells. In addition to the *N*-methylflindersin factor tend to increase cytotoxic activity against human colon cancer cells (DLD-1) (Nakashima et al. 2012).

3. Experimental

3.1. Plant material

The fresh leaves of *M. quercifolia* obtained from Cianten Farm, Cigudek District, Bogor, West Java, Indonesia on Nov. 2017. The plant identified by Mr. Ismail Rachman, a botanist senior from the Herbarium Bogoriense, Bogor, Indonesia. A specimen (MQ 20171104) deposited as a reference.

3.2. Extraction and isolation

The air-dried leaves of *M. quercifolia* (2.0 kg) were extracted with MeOH three times (6L, each for two days) at room temperature. Evaporation of the solvent with evaporator gave a MeOH extract (450 g) and then partitioned with *n*-hexane three times. Furthermore, the MeOH extract was added with H₂O (4:1 v/v) and partitioned with EtOAc three times gave the crude extract (5 g). The EtOAc extract (4.8 g), fractionated by CC on silica gel, eluted with *n*-hexane-EtOAc (from 9:1 to 3:7) providing two fractions, A and B. Fraction A (1.3 g) was fractionated by CC chromatography on polyamide, eluted with *n*-hexane-CHCl₃ (from 9:1 to 1:4) gave two subfractions, A₁ and A₂. Compounds **3** (13 mg) and **4** (15 mg) were isolated from subfraction A₂ (655 mg) using radial planar chromatography with the same eluent. Fraction B (800 g) was

separated by CC chromatography on polyamide, and eluted *n*-hexane-EtOAc (from 4:1 to 1:1) gave subfractions B₁-B₂. Compounds **1** (5 mg) and **2** (7 mg) was obtained from subfraction B₁ (265 mg) by radial planar chromatography, eluted with *n*-hexane-diisopropylether (from 3:9 to 7:3), and diisopropylether as a solvent system. By the same methods, compound **5** (16 mg) isolated from subfraction B₂ (148 mg).

3.3. Spectral data

Meliquercifolin A (**1**): yellow solid, UV/Vis (MeOH) λ_{\max} (nm) (log ϵ): 217 (3.03), 269 (3.08), 289 (3.20) and 325 (2.81) nm. IR (KBr) ν_{\max} (cm⁻¹): 3411, 2972, 2925, 2852, 1649, 1604, 1579 and 1159. ¹H-NMR (400 MHz, CDCl₃) δ_{H} ppm: 6.01 (1H, *d*, *J* = 9.8 Hz, H-3), 7.81 (1H, *d*, *J* = 9.8 Hz, H-4), 4.04 (3H, *s*, 5-OCH₃), 4.71 (1H, *t*, *J* = 6.9 Hz, H-6), 5.42 (1H, *ddd*, *J* = 8.2; 6.2; 2.2 Hz, H-7), 6.21 (1H, *s*, H-9), 3.05 (1H, *dd*, *J* = 9.8; 5.8 Hz, H-3), 3.96 (1H, *t*, *J* = 2.2 Hz, H-4), 3.67 (3H, *s*, 5'-OCH₃), 3.59 (3H, *s*, 7'-OCH₃), 5.98 (1H, *s*, H-8), 2.17 (3H, *s*, H-10), 1.19 (3H, *s*, H-11) and 1.61 (3H, *s*, H-12). ¹³C-NMR (100 MHz, CDCl₃) δ_{C} ppm: 161.1 (C-2), 109.7 (C-3), 139.5 (C-4), 108.1 (C-4a), 152.7 (C-5), 104.7 (C-5a), 44.1 (C-6), 84.9 (C-7), 168.5 (C-8a), 91.7 (C-9), 156.7 (C-9a), 74.0 (C-2), 45.5 (C-3), 34.7 (C-4), 106.6 (C-4a), 155.7 (C-5), 62.6 (5'-OCH₃), 117.8 (C-6), 156.8 (C-7), 55.8 (7'-OCH₃), 96.8 (C-8), 156.4 (C-8a), 202.5 (C-9), 32.3 (C-10), 25.3 (C-11) and 25.7 (C-12). HRESIMS: *m/z* [M + H]⁺ calcd. for C₂₇H₂₂O₈ 479.1706, found 479.1729.

Meliquercifolin B (**2**): yellow solid, UV/Vis (MeOH) λ_{\max} (nm) (log ϵ): 217 (3.33), 234 (3.12), 255 (3.02), 268 (2.98) and 299 (2.69) nm. IR (KBr) ν (cm⁻¹): 3448, 2966, 2923, 2854, 1651, 1627, 1460 and 1186. ¹H-NMR (400 MHz, CDCl₃) δ_{H} ppm: 6.02 (1H, *d*, *J* = 9.6 Hz, H-3), 7.79 (1H, *d*, *J* = 9.6 Hz, H-4), 3.68 (3H, *s*, 5-OCH₃), 4.75 (1H, *t*, *J* = 6.8 Hz, H-6), 5.50 (1H, *dd*, *J* = 6.8; 3.9 Hz, H-7), 3.98 (3H, *s*, 9-OCH₃), 3.09 (1H, *dd*, *J* = 9.9; 5.8 Hz, H-3), 3.99 (1H, *dd*, *J* = 5.8; 1.8 Hz, H-4), 3.68 (3H, *s*, 5'-OCH₃), 3.57 (3H, *s*, 7'-OCH₃), 5.92 (1H, *s*, H-8), 2.21 (3H, *s*, H-10), 1.12 (3H, *s*, H-11), and 1.66 (3H, *s*, H-12). ¹³C-NMR (100 MHz, CDCl₃) δ_{C} ppm: 160.9 (C-2), 109.9 (C-3), 139.6 (C-4), 104.4 (C-4a), 126.0 (C-5), 106.9 (C-5a), 44.4 (C-6), 85.3 (C-7), 159.8 (C-8a), 148.1 (C-9), 149.1 (C-9a), 74.1 (C-2), 45.7 (C-3), 34.5 (C-4), 106.2 (C-4a), 156.9 (C-5), 62.7 (5'-OCH₃), 117.3 (C-6), 156.8 (C-7), 55.8 (7'-OCH₃), 96.8 (C-8), 163.8 (C-8a), 209.2 (C-9), 32.3 (C-10), 25.4 (C-11), and 25.7 (C-12). HRESIMS: *m/z* [M + H]⁺ calcd. for C₂₈H₂₉O₉ 509.1812, found 509.1802.

3.4. Cytotoxic assay

Effect cytotoxic of compounds (**1-5**) against HeLa (human cervical cancer cells), MCF-7 (human breast cancer cells), P-388 (human murine leukemia cells) were evaluated using the MTT colorimetric methods (Mah et al. 2015; Tanjung et al. 2017, 2018; Segun et al. 2019). Doxorubicin used as a control positive for HeLa, and MCF-7 as well as artonin for P-388 cells.

4. Conclusions

Meliquercifolins A (**1**) and B (**2**), two new coumarins bearing an acetophenone derivative were isolated from the leaves of *M. quercifolia* together with three known

compounds, melicodenines E (3), F (4) and I (5). Compounds (1 and 2) are the first time coumarin bearing an acetophenone derivative found on *Melicope*.

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Disclosure statement

No potential conflict of interest reported by the authors.

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