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## Cytotoxicity of triterpenes from the leaves of *Garcinia prainiana* King (Guttiferae)

Shamsul ON, Wan Mohd Nuzul Hakimi Wan SALLEH, Farediah AHMAD, Hasnah Mohd SIRAT, Muhammad TAHER

### ABSTRACT

Phytochemical investigation of the leaves of *Garcinia prainiana* resulted in the isolation of five triterpenes identified as squalene **1**, friedelin **2**, lupeol **3**, 3 $\beta$ -hydroxyeupha-5,22-diene **4**, and 3 $\beta$ -acetoxyeupha-5,22-diene **5**. The structures of these compounds were established by analysis of their spectroscopic data, as compared to that of reported compounds. All

compounds were tested for cytotoxicity against H1299 and A549 lung cancer cell lines using MTT assay. 3 $\beta$ -Acetoxyeupha-5,22-diene **5** exhibited the most significant activity against human non-small cell lung carcinoma cell lines, H1299 and A549 with IC<sub>50</sub> values 18.0  $\mu$ g/mL and 36.3  $\mu$ g/mL, respectively.

**Keywords:** Triterpene, *Garcinia prainiana*, Cytotoxicity

### INTRODUCTION

Triterpenes are naturally occurring alkenes of vegetable, animal and also fungal origin, classified among an extensive and structurally diverse group of natural substances, referred to as triterpenoids. Their structure includes 30 elements of carbon and they are constituted by isoprene units (1-4). Taking into consideration the structure, triterpenes may be divided into linear ones mainly derivatives of squalene, tetracyclic and pentacyclic, containing respectively four and five cycles, as well as two- and tricyclic ones (5). Representatives of those show anticancer (6), anti-inflammatory (7), antioxidative (8), antiviral (9), antibacterial (10) and antifungal activities (11).

The genus *Garcinia* (Guttiferae) is commonly distributed in tropical and subtropical countries of South East Asia, West and East Africa, in addition to Central and South America (12-13). It comprises over 400 species in Asia and reported with 49 species from Malaysia region (14). The genus is known for the production of phenolic compounds including xanthenes, benzophenones, biflavonoids and depsidones (15). Xanthenes are well recognized as chemotaxonomic markers for plants of *Garcinia* species and many of which have interesting pharmacological effects, including antioxidant, antimicrobial, anti-inflammatory and cytotoxic activities (16-17). Several *Garcinia* species are used in traditional medicine to cure various ailments such as abdominal pain, dysentery, diarrhoea, infected wound and gonorrhoea (18).

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*Garcinia prainiana* King (Guttiferae), locally known as *cherupu*, *kechupu* or *menchupu*, is a small tree up to 10 m tall. The bark produces white exudates and it is scattered in lowland forest throughout Peninsular Malaysia. It grows in similar agroecological tropical conditions as the common mangosteen. It thrives in areas with annual rainfall above 1500 mm and occurs in many soil types but prefers soils that are porous, deep, wet but well-drained. It occurs wild in lowland forests and on hill sides and ridges up to 900 m altitude (19). Herein, we report the cytotoxicity of phytochemicals isolated from the leaves *G. prainiana*.

## MATERIAL AND METHODS

### Plant material

Leaves of *G. prainiana* was collected from Agricultural Garden, Kuantan, Pahang in November 2009 and identified by Dr. Muhammad Taher. The voucher specimens (MT27) were deposited at the herbarium of Kulliyah of Pharmacy, International Islamic University, Malaysia.

### General experimental procedures

Silica gel 60 F<sub>254</sub> precoated aluminium plates (0.2 mm, Merck) were used for TLC analysis. The TLC and PTLC spots were visualized under UV light (254 and 366 nm) followed by spraying with 5% H<sub>2</sub>SO<sub>4</sub> in methanol and 1% vanillin in MeOH, followed by heating at 120°C for 5 min. Vacuum Liquid Chromatography (VLC) was carried out on silica gel 230-400 mesh (Merck) while Column Chromatography was carried out on silica gel 70-230 mesh (Merck). The UV spectra were obtained in methanol on a Shimadzu UV 1601PC spectrophotometer. The IR spectra were obtained on a Perkin-Elmer 1600 FTIR spectrophotometer. The mass spectra were recorded on Bruker Mass Spectrometry Services, obtained from National University of Singapore (NUS). The <sup>1</sup>H-NMR and <sup>13</sup>C-NMR spectra were recorded in deuterated chloroform on a Bruker Avance 400 MHz spectrometer, chemical shifts are reported in ppm on  $\delta$  scale, and the coupling constants are given in Hz. All solvents were analytical reagent grade.

### Extraction and isolation procedures

The dried and powdered leaves of *G. prainiana* (300 g) were extracted using a Soxhlet extractor for 9 h each with *n*-hexane, dichloromethane (DCM) and MeOH (2.5 L each) successively at room temperature. Concentration of each solvent under reduced pressure afforded a crude *n*-hexane (10 g), DCM (7 g) and MeOH extract (18 g), all as gummy

dark green liquid. The *n*-hexane extract (8 g) was submitted to VLC on silica gel using a gradient of *n*-hexane:CHCl<sub>3</sub>:EtOAc to yielded thirteen fractions. Fraction 3-7 (180 mg) was combined and separated using silica gel CC (diameter 3.0 cm, length 50 cm) using a gradient of *n*-hexane:CHCl<sub>3</sub> to give seventy sub-fractions. Sub-fraction 5-30 (75 mg) showed the existence of two major components on TLC (hexane:CHCl<sub>3</sub>) and was submitted for further purification by CC (diameter 1.5 cm, length 40 cm) eluted with a gradient hexane:CHCl<sub>3</sub> to produced **1** (14 mg) and **2** (30 mg). Sub-fraction 33-45 (100 mg) was applied to silica gel CC (diameter 1.5 cm, length 40 cm) eluted with *n*-hexane:CHCl<sub>3</sub> to furnish **3** (15 mg). The MeOH extract (10 g) was submitted to VLC on silica gel using gradient of CHCl<sub>3</sub>:EtOAc (95:5, 90:10, 80:20, 70:30, 60:40, 50:50, 40:60, 20:80, 0:100) to furnish nine fractions. Fraction 3-5 (150 mg) was combined and applied to silica gel CC (diameter 2.5 cm, length 45 cm) eluted with CHCl<sub>3</sub>:MeOH led to the isolation of **4** (15 mg) and **5** (20 mg). The identification of isolated compounds was achieved by UV, IR, MS, <sup>1</sup>H, <sup>13</sup>C NMR and 2D NMR methods. Their structures were confirmed by comparisons with reported data.

### Cytotoxic assay

The assay against human non-small cell lung carcinoma cell lines H1299 and A549 were conducted using the 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) test according to Mosmann et al. (20). The monolayer cell lines were cultured in DMEM supplemented with 10% (v/v) foetal bovine serum and antibiotic (1% penicillin-streptomycin) in a 96-well plate at a density of 2×10<sup>4</sup> cells/mL. After reaching confluence (2×10<sup>5</sup> cell/mL), then the cells were treated with the samples. The tested compounds were dissolved in DMSO (1 mg/mL). Each compounds were treated in triplicates with serial dilution of sample (0.01-1 mg/mL) for 24 h. Doxorubicin was used as a commercialized anticancer standard. To reduce the interference of residue of supernatant, the monolayer cell lines was washed two times with phosphate buffer saline (PBS), the 20 ml of MTT stock solution (5 mg/mL) was added and the plates were further incubated overnight at 37°C. DMSO (100 mL) was added to each well to dissolve the purple formazan crystal. After 1 h, the solubilized formazan was measured at 570 and 630 nm (reference) with a UV microplate reader. The half reduction in cell number relative to control or IC<sub>50</sub> was established by extrapolation from linear regression of experimental data.

### Scanning Electron Microscopy

In brief, about 2.5 mL of confluent cell was transferred into a

6-well plate, which contained sterile cover slips at the bottom of the wells, and incubated overnight in CO<sub>2</sub> incubator at 37°C. The seeded cell was treated with sample/standard using the IC<sub>50</sub> value to make sure that just 50% of the treated cell will inhibit and incubated with media again for overnight. Then the media and sample/standard were discarded and washed with phosphate buffer. Then, McDowell and Trump fixative was added and it was kept overnight. Then the fixative was removed gently and washed with cacodylate buffer for two times. Then, a 1 mg/mL of osmium tetroxide was added in small volume just to cover the surface for 1 h. After that, a double-distilled water was used for washing to remove the excess of osmium tetroxide for 3 times. Afterward, the dehydration process was started by washing with 50% ethanol for 15 min once, followed by 75% ethanol for 15 min once, followed by 95% ethanol for 20 min for 3 times, and then followed by absolute ethanol for 20 min for 3 times. Finally, 0.5 mL HMDS was added and left uncovered overnight. After that, the cover slip was taken out and coated with gold in a Baltec-CED 030 and viewed by Carl Zeiss Evo 50 Scanning Electron Microscopy (21-22).

### Statistical analysis

Data obtained from cytotoxicity was expressed as mean values. The statistical analyses and significance of the results was determined using a one way ANOVA ( $p > 0.05$ ). A statistical package (SPSS version 11.0) was used for the data analysis.

## RESULTS AND DISCUSSION

The structures of the triterpenes were elucidated on the basis of 1D and 2D NMR data and by comparisons with published data. The compounds were characterized as squalene **1**, friedelin **2**, lupeol **3**, 3 $\beta$ -hydroxyeupha-5,22-diene **4** and 3 $\beta$ -acetoxyeupha-5,22-diene **5** (Figure 1). Mawa and Said, (23) have reported the isolation of friedelin, eupha-8,24-diene-3 $\beta$ -ol, stigmasterol, teraxerone and teraxerol from the twigs of *G. prainiana*, while Susanti et al. (24) isolated friedelin and lanosterol from the same part. Previous studies also reported that the methanol leaf extract of *G. prainiana* exhibited an inhibition against nitric oxide production (25), while the leaf and fruit extracts were found to exhibit antioxidant capacity (26). In addition, the twig extracts have shown significant DPPH radical scavenging with IC<sub>50</sub> value 2.9 (MeOH extract) and 1.9 (EtOAc extract)  $\mu\text{g/mL}$  (26). With the interest in the chemotaxonomy and biomedical potential of *Garcinia*, the present study was conducted to search for potential compounds which are cytotoxic to human lung cancer. The

isolated compounds were subjected to cytotoxicity screening against human lung cancer H1299 and A549 cell lines. Table 1 shows the cytotoxic activity of the isolated compounds. Compounds **1** and **5** were found to have significant activity towards both cell lines. Compound **5** shown the best activity with IC<sub>50</sub> values of 18.0 and 36.3  $\mu\text{g/mL}$ , against H1299 and A549 cell lines, respectively. Compound **1** gave IC<sub>50</sub> values of 23.2  $\mu\text{g/mL}$  (H1299) and 74.8  $\mu\text{g/mL}$  (A549). The most active compounds (**1** and **5**) showed more activity against H1299 than A549 cell lines. Scanning electron microscope has been done on the cells after treatment with **1** and **5** as shown in Figure 2. Both compounds induced apoptosis on H1299 and A549 indicated by blebbing phenomenon, a specific pattern on apoptosis (27). Cytotoxic activity has also been observed in derivatives of linear squalene. Apart from the inhibitory role, squalene also shows the capability of inducing apoptosis in many neoplastic lines: leukemia, melanoma, colon cancer, prostate cancer, ovarian carcinoma, liver cancer, breast cancer, lung cancer and peripheral nervous system carcinoma (28). Triterpenes are natural compounds showing a wide spectrum of biological effects. They proved to have antibacterial, antiviral, antifungal, antioxidative and anti-inflammatory properties, as well as anticancer and chemopreventive ones (29).

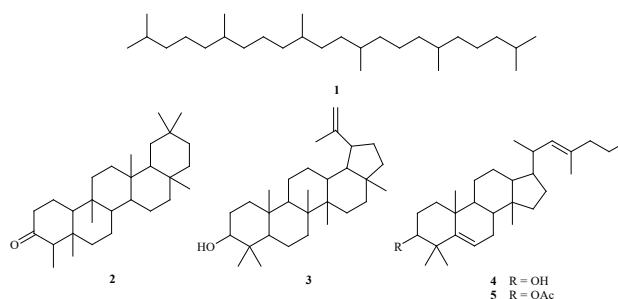
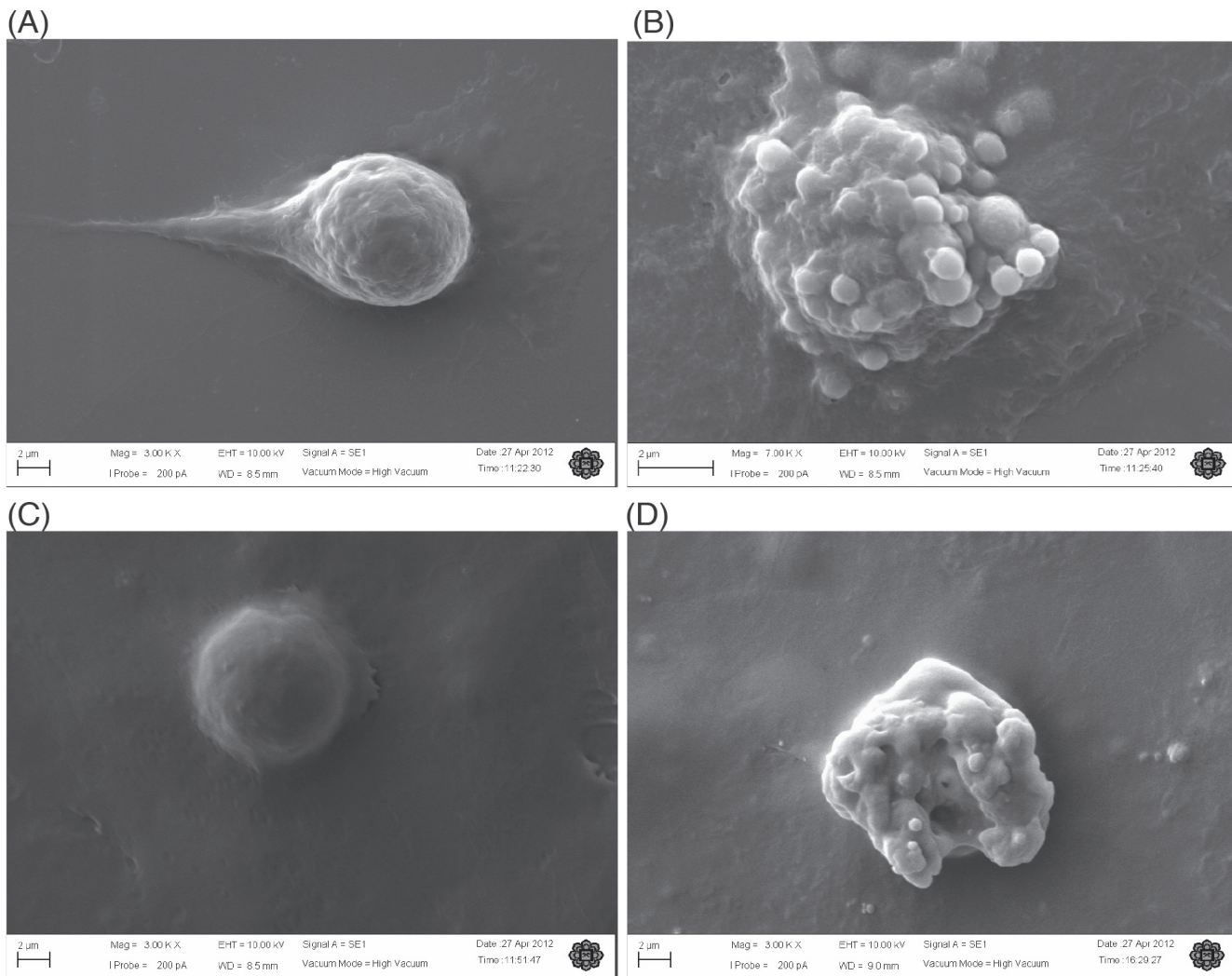


Figure 1. Chemical structures of triterpenes isolated from *G. prainiana*

Table 1. Cytotoxicity activity of triterpenes isolated from *G. prainiana*

| Compounds   | IC <sub>50</sub> ( $\mu\text{g/mL}$ ) |                 |
|-------------|---------------------------------------|-----------------|
|             | H1299                                 | A549            |
| <b>1</b>    | 23.2 $\pm$ 2.1                        | 74.8 $\pm$ 3.4  |
| <b>2</b>    | 163.9 $\pm$ 1.9                       | 254.7 $\pm$ 2.1 |
| <b>3</b>    | 126.1 $\pm$ 1.3                       | 110.2 $\pm$ 0.2 |
| <b>4</b>    | 103.2 $\pm$ 1.6                       | 156.5 $\pm$ 0.4 |
| <b>5</b>    | 18.0 $\pm$ 1.7                        | 36.3 $\pm$ 2.6  |
| Doxorubicin | 10.0 $\pm$ 0.1                        | 2.38 $\pm$ 0.2  |

Each value represents the mean  $\pm$ SD.



**Figure 2.** Cell membrane blebbing during apoptosis after treated with 100 µg/mL of **1** and **5**. (A) H1299 (untreated with **1**) (B) H1299 (treated with **1**) (C) A549 (untreated with **5**) (D) A549 (treated with **5**).

## CONCLUSION

This study investigated the isolated triterpenes from *G. prainiana* and their cytotoxicity. The cytotoxicity evaluation of isolated pure compounds provides valuable information regarding the prospective use of medicinal plants as sources

of new drugs. Also, it emphasizes the rationale for using medicinal plants in folk medicine. The cytotoxicity of isolated compounds may validate the traditional use of the plant as an anticancer agent. Therefore, more directed research is needed to explore the ability of plants to enhance the discovery and development of new chemical entities.

### *Garcinia prainiana* King (Guttiferae) Yapraklarından Elde Edilen Triterpenlerin Sitotoksik Özellikleri

#### ÖZ

*Garcinia prainiana* yapraklarını konu alan fitokimyasal araştırmalar sonucunda triterpen yapısında beş bileşik izole edilmiştir; skualen **1**, friedelin **2**, lupeol **3**, 3β-hidroksiöfa-5,22-dien **4**, and 3β-asetoksiöfa-5,22-dien **5**. İzole edilen

bileşiklerin yapıları spektroskopik yöntemler kullanılarak aydınlatılmış ve elde edilen sonuçlar yukarıda belirtilen bileşiklere ait spektroskopik verilerle karşılaştırılmıştır. Tüm bileşiklerin, H1299 ve A549 akciğer kanseri hücre hatları üzerindeki sitotoksik etkileri MTT yöntemiyle çalışılmıştır. 3β-Asetoksiöfa-5,22-dien **5**, H1299 ve A549 hücre hatlarına karşı sırasıyla  $IC_{50} = 18.0$  µg/ml ve  $IC_{50} = 36.3$  µg/ml derişimde etki göstererek serisindeki en etkili bileşik olarak öne çıkmıştır.

**Anahtar kelimeler:** Triterpen, *Garcinia prainiana*, Sitotoksiste.

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