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Phytoconstituents from *Alpinia purpurata* and their *in vitro* inhibitory activity against *Mycobacterium tuberculosis*

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ABSTRACT

Alpinia purpurata or red ginger was studied for its phytochemical constituents as part of our growing interest on Philippine Zingiberaceae plants that may exhibit antimycobacterial activity. The hexane and dichloromethane subextracts of the leaves were fractionated and purified using silica gel chromatography to afford a mixture of C_{28} to C_{32} fatty alcohols, a 3-methoxyflavone and two steroidal glycosides. The two latter metabolites were spectroscopically identified as kumatakenin (**1**), sitosteryl-3-*O*-6-palmitoyl-β-D-glucoside (**2**) and β-sitosteryl galactoside (**3**) using UV, IR, EIMS and NMR experiments, and by comparison with literature data. This study demonstrates for the first time the isolation of these constituents from *A. purpurata*. In addition to the purported anti-inflammatory activity, its phytomedicinal potential to treat tuberculosis is also described.

Keywords: *Alpinia purpurata*, fatty alcohols, kumatakenin, *Mycobacterium tuberculosis*, sitosteryl glycosides

INTRODUCTION

Several species of the genus *Alpinia* were reported to exhibit fungicidal, antioxidant and bactericidal properties [1–2]. *Alpinia purpurata* (Vieill.) K. Schum (Family Zingiberaceae) is locally known in the Philippines as *"luyang pula"* or red ginger, and is a native to the Pacific [3]. Studies on its chemical constituents revealed the presence of α-pinene, β-pinene [4], 1,8-cineole, (*E*)-methylcinnamate [5], 6-shogaol, 8-gingerol, 6-gingerol, 10-gingerol, 10-shogaol and 4-shogaol [6]. A U.S. patent reported that its total anthocyanidin, shogaol and gingerol content shows promise in the treatment of inflammatory diseases such as arthritis [6–7].

With limited literature available as to the phytochemistry and biological activity of *A. purpurata*, and with the growing interest in Philippine Zingiberaceae species that inhibit the growth of *Mycobacterium tuberculosis* $H_{37}Rv$ [8–9], we embarked on further exploration on the isolation and identification of secondary metabolites from this *Alpinia* species. In addition to the fatty alcohol mixture, we report in this paper the chromatographic purification and spectroscopic identification of a flavone and two sitosteryl glycosides namely, kumatakenin **1**, sitosteryl-3-*O*-6-palmitoyl-β-D-glucoside **2** and β-sitosteryl galactoside **3** from *A. purpurata*. The inhibitory activity against *M. tuberculosis* $H_{37}Rv$ of the extracts, fractions and the purified compounds is also presented.

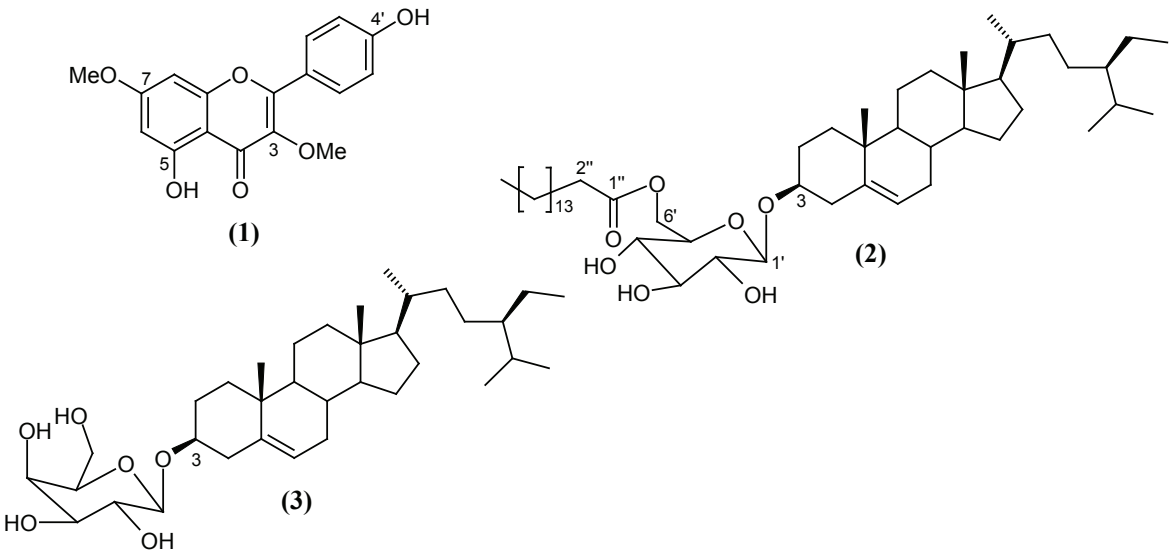

Figure 1. Flavone and sitosteryl glycosides from *A. purpurata*.

RESULTS AND DISCUSSION

The MABA assay [10] result of the crude ethanolic extract of the various parts of *Alpinia purpurata* had shown the leaf extract to possess the highest activity followed by the rhizome and flower extracts. Among the sub-extracts, the DCM sub-extract exhibited the highest activity followed by hexane and n-butanol sub-extracts. All fractions obtained from the hexane and DCM sub-extracts showed low to moderate activity (Table 1).

Further chromatographic work-up was undertaken on fraction two of the hexane sub-extract, which afforded a white amorphous solid after crystallization. This compound was distinctly identified from the ^{1}H-NMR and ^{13}C-NMR spectra to be a fatty alcohol, but the LR-EIMS spectrum otherwise showed it to be a mixture of fatty alcohols. By careful analysis of the m/z values in the mass spectrum, it could be claimed that it is composed of montanyl alcohol (C28:0), melissyl alcohol (C30:0, major component) and domelissyl alcohol (C32:0) based on several characteristic peaks due to fragment ions of $[M^{+}\text{-}H_2O]$ [11].

The dichloromethane extract was likewise subjected for further investigation owing to its interesting phytochemical profile. Nine fractions after VLC were obtained from which fractions three and four yielded three solid compounds **1**–**3**.

Compound **1**, a yellow crystalline substance (13.0 mg), was purified from fraction three after recrystallization. It was found to be a flavonoid after treating its TLC chromatograms with $FeCl_3$-$K_4Fe(CN)_6$ and 10% $SbCl_3$ in chloroform as shown by a blue-green spot and red-orange fluorescence under UV (365 nm), respectively [12]. In addition, major absorptions at 269 (Band I) and 352 nm (Band II) and a weak shoulder at 303 nm, which are typical for flavones, were observed in its UV spectrum [13–14]. The structure, and degree and pattern of oxygenation in the

Table 1. Percent inhibitory activity of the extracts, sub-extracts and fractions of *A. purpurata* at 64 µg/mL against *M. tuberculosis* H_{37}Rv

Sample	% Inhibition
Crude Ethanolic Extracts	
Flowers	30
Leaves	62
Rhizome	34
Leaf sub-extracts	
Hexane	64
DCM	72
n-Butanol	35
Hexane Fractions (APH)	
APH1	47
APH2	38
APH3	55
APH4	48
APH5	61
APH6	74
APH7	73
APH8	70
APH9	74
APH10	46
APH11	23
DCM Fractions (APD)	
APD 1	65
APD 2	58
APD 3	68
APD 4	60
APD 5	69
APD 6	66
APD 7	46
APD 8	50
APD 9	64

Control: Rifampin, 99% at 0.18 µg/mL

Table 1. Comparison of the NMR Spectral Data of 1 with the Reported values of Kumatakenin (Wang et al., 1989; Urbatsch, 1976)

Position	^{13}C NMR (1)	^{13}C NMR Kumatakenin (Wang, 1989)	^{1}H NMR (1)	^{1}H NMR Kumatakenin (Urbatsch, 1976)
	δ ppm	δ ppm	δ ppm	δ ppm
2	156.7	155.77		
3	138.4	137.71		
4	178.7	177.93		
5	161.3	160.90		
6	97.8	97.57	6.24 (d J= 2)	6.15 (d J= 2.5)
7	165.4	164.95		
8	92.2	92.09	6.37 (d J= 2)	6.43 (d J= 2.5)
9	156.8	156.16		
10	105.8	105.10		
1'	121.3	120.41		
2'	130.2	130.04	7.89 (d J= 9)	7.95 (dd J= 2, 8.5)
3'	115.5	115.52	6.82 (d J= 9)	6.85 (dd J= 2, 8.5)
4'	159.9	160.19		
5'	115.5	115.52	6.82 (d J= 9)	6.85 (dd J= 2, 8.5)
6'	130.2	130.04	7.89 (d J= 9)	7.95 (dd J= 2, 8.5)
3-OCH_3	59.9	59.51	3.69 s	3.83 s
7-OCH_3	55.6	55.86	3.78 s	3.83 s
	$CDCl_3$ with CD_3OD, 125 MHz	DMSO-d_6, 50.30 MHz	$CDCl_3$ with CD_3OD, 500MHz	DMSO-d_6, 200 MHz

flavonoid structure were examined by studying the effect of several wavelength shift reagents, NaOMe, NaOAc, NaOAc-1% aq. H_3BO_3, $AlCl_3$ and $AlCl_3$-HCl in the UV spectral region of **1**. With NaOMe, a 44 nm bathochromic shift and a significant increase in absorbance intensity were noted for Band II. This indicates the presence of a C-4' hydroxyl group. Treatment with NaOAc gave no observable change in the spectrum which shows that an alkoxy substituent is present in the C-7 position of the flavone nucleus. Addition of 1% boric acid produced no change in the spectrum which is symptomatic of the absence of *ortho*-hydroxyphenolic functionalities. This was also substantiated by the result of adding 0.1M HCl/ $AlCl_3$ in a separate experiment. The appearance of four absorption peaks (278, 304, 352 and 399 nm) after the complexation of $AlCl_3$ with **1** is a clear indication that a 5-hydroxyl moiety is present [14].

The IR spectrum of **1** showed the presence of an enone (1665 cm^{-1}) and phenolic OH's (3243 cm^{-1}). The base peak at *m/z* 314 in the LR-EIMS mass spectrum was designated the molecular-ion peak. In the 500 MHz ^{1}H NMR spectrum, two sets of AA'BB'- protons belonging to a *para*-substituted benzenoid moiety, two methoxy protons and meta-coupled protons were noted. In the proton-decoupled ^{13}C and DEPT-135 NMR spectrum, a total of 18 carbon atoms were accounted for **1** from which a conjugated ketone, six oxygenated olefinic/aromatic carbons, six aromatic methines and two methoxy carbons were deduced. The gross structure of the compound which is analogous to kumatakenin [15–16] was elucidated via an HMBC experiment. Key ^{1}H-^{13}C correlations are shown in Figure 2.

It is noteworthy to report the isolation of kumatakenin (**1**) from *A. purpurata*. This rare ethyl ether flavonol was first isolated from the seeds of *A. japonica* [17] and *A. kumatake* [18]. Hence, the identification of **1** strengthens the chemical link of *A. purpurata* with the other species of *Alpinia*.

Compound **2** was afforded as a white amorphous solid (8.4 mg). TLC chromatograms of the isolate sprayed with Liebermann-Burchard and Molisch reagents [12] suggested a steroidal glycoside structure. The IR spectrum showed the presence of hydroxyl (3439 cm^{-1}) and ester (1738 cm^{-1}) functionalities. The molecular ion peak was not observed in the LR-EIMS spectrum. Instead, fragment ions corresponding to sitosterol (m/z 414, $C_{29}H_{50}O$) and a palmitoxy group (m/z 256, $C_{16}H_{31}O_2$) were noted. Signals characteristic of sitosterol resonances i.e. C-6 olefinic proton (δ 5.36), methyls associated to the cholestane skeleton (δ 0.69–0.79), glucose (δ 4.34 - 4.38) and a palmitoyl group (δ 0.84, 1.26, 2.34) were evident in the 500 MHz ^{1}H NMR of **2**. A total of 51 carbon atoms, of which 7 CH_3, 25 aliphatic CH_2, 7 aliphatic CH, two aliphatic quaternary C, one oxygenated CH_2, 6 oxygenated CH and one each olefinic CH and C, were found in the 125 MHz ^{13}C NMR spectrum. HMBC correlations which were instrumental in finding the correct identity of **2** as sitosteryl-3-*O*-6-palmitoyl-β-D-glucoside are shown in Figure 3. The NMR values of **2** are in well agreement with those reported for sitosteryl-3-*O*-6-palmitoyl-β-D-glucoside by Pei-Wu et al. [19], Gomes et al. [20], and Shaiq Ali et al. [21].

Metabolite **3** was obtained as white crystalline flakes (4.1 mg). The partial identity of **3** was revealed to be a

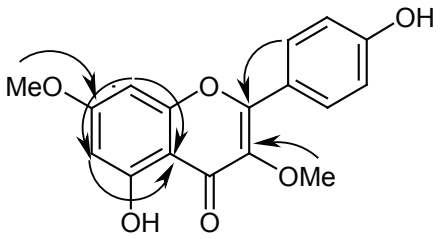

Figure 2. Key ^{1}H-^{13}C HMBC Correlations in 1.

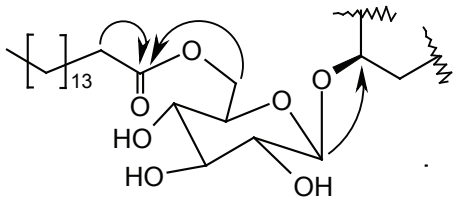

Figure 3. Key ^{1}H-^{13}C HMBC Correlations in 2.

steroidal glycoside as in **2** using the same phytochemical experiments. Only the occurrence of hydroxyl functionalities was inferred this time from the IR spectrum. The presence of a sitosterol fragment was ascertained in the LR-EIMS spectrum which was also verified by the ^{1}H and ^{13}C NMR spectra. The sugar moiety was deduced to be galactose upon comparison of the ^{1}H-NMR and ^{13}C-NMR values with those in literature [22–24]. Hence, the identity of **3** was established.

Sitosteryl glycosides **2–3** have been isolated from other several plant sources [19, 21]. These metabolites are also present in plant species belonging to the family Zingiberaceae [25].

The fatty alcohols showed an MIC value of 64 µg/mL and proved most active compared to the flavonoid kumatakenin and the steroidal glycosides previously reported to have MIC values > 128 µg/mL. Long chain alcohols show growth inhibitory activity to Gram-positive organisms including *Staphylococcus aureus* and *Propionibacterium acnes* [26]. Moreover, a study done by Togashi and co-workers in 2007 further supported the antibacterial activity of long chain aliphatic alcohols that had bactericidal activity and membrane-damaging activity on *Staphylococcus aureus*. Experimental results indicate that the antibacterial activity of long chain alcohols is mediated by damage to cell membranes that allows leakage of K^{+} ions, with subsequent reactions that induce further leakage [27]. Thus, the ability of the isolated fatty alcohols to inhibit the growth of *M. tuberculosis* H_{37}Rv gives further credence on the antibacterial activity of long chain alcohols particularly those bearing more than twenty carbon chains.

The activity of kumatakenin confirms the study by Murillo et al. [28] on its action against *M. tuberculosis* H_{37}Rv. This compound has antiviral activity against HIV [29], the virus that aggravates the problem on tuberculosis due to susceptibility to the lung pathogen. Plant sterols particularly β-sitosterol and its glucosides have been investigated as immune regulators of T-cell activity [30] and as agents in maintaining the CD_{4}^{+} count in the absence of anti-retroviral therapy in HIV-infected patients [31]. Plant sterols were effective in patients treated for pulmonary TB, causing increase in their peripheral blood lymphocytes and eosinophil counts [32].

To date, this is the first report of **2–3** from the genus *Alpinia*. More importantly, this paper cites for the first time the isolation of all compounds from *A. purpurata*. Furthermore, this study demonstrates the promise of this plant as a source of phytomedicinals that can fight TB.

METHODOLOGY

General

The leaves of *Alpinia purpurata* (Vieill.) K. Schum. were collected in Los Baños, Laguna (February 2004). Herbarium specimens (USTH 4717) were kept at the Botany Laboratory of the Research Center for the Natural Sciences, Thomas Aquinas Research Complex, University of Santo Tomas, Manila, Philippines.

Electron impact (EI) mass spectral analysis was carried out in JEOL D-300 FD mass spectrometer using *m*-nitrobenzyl alcohol/$CHCl_3$ as carrier at 60°C. Proton (^{1}H) and ^{13}C NMR measurements were recorded in JEOL GX 400 MHz NMR spectrometer using $CDCl_3$ (δ 7.26 for ^{1}H, δ 77.0 for ^{13}C) as internal reference.

Extraction and Isolation

The air-dried leaves of *Alpinia purpurata* (1.7 kg) were extracted with ethanol to give a crude extract (378 g) that was partitioned according to increasing polarity using n-hexane, dichloromethane and n-butanol.

The hexane extract (53.4 g) was subjected to VLC by gradient elution (20% increments) using hexane/ethyl acetate to give 11 fractions. Fraction 2 (8.4 g) was purified by GCC and gradient elution (5% increments) with hexane/chloroform and chloroform/acetone, to give four fractions. Sub-fraction 2 was further purified and gave 19 fractions. A solid in the eighth fraction was recrystallized in acetone to afford a white amorphous solid (10 mg) of a fatty alcohol mixture.

The dichloromethane extract (2.5 g) was subjected to vacuum liquid chromatography (VLC) (Si gel HF_{254} Merck Art. 1.07739) by gradient elution (20% increments) using chloroform, chloroform-acetone, acetone and acetone-methanol to furnish nine fractions.

After evaporation of fraction 3, yellow needle-like crystals appeared and were recrystallized in acetone to give **1** (13 mg). UV Spectral analysis was also done on compound **1** using various shift reagents to determine the aromatic substitutions of the compound. Fraction four (458 mg) was gravity column chromatographed (GCC) (Si gel 60 Merck Art. 1.07734, 25 mm I.D.) by gradient elution with benzene-acetone (5:1); (5:2); (5:3), neat acetone, acetone-methanol (1:1) and neat methanol to give nine fractions. Sub-fraction eight gave an amorphous white powder that was recrystallized in acetone to give **2** (8.4 mg). Concentration and recrystallization of sub-fraction 6 in methanol afforded **3** (4.1 mg) as white flakes.

Fatty alcohols. ^{1}H NMR (500 MHz, $CDCl_3$) 3.57 (t, J 6.6Hz, H-1), 1.49 (m, H-2), 1.18 (br s), 0.81 (t, J 7Hz). ^{13}C NMR (DEPT-135, 125 MHz, $CDCl_3$) 63.1 (C-1), 32.8 (C-2), 31.9 (C-3), 29.7, 25.7(C-3), 22.6, 14.1. LR-EIMS m/z: 392.5 $(C_{28}H_{57}OH\text{-}H_2O)^+$, 420.5 $(C_{30}H_{61}OH\text{-}H_2O)^+$, 448 $(C_{32}H_{65}OH\text{-}H_2O)^+$

Kumatakenin (1), yellow needles (13 mg), m.p. 248–249°C (uncorr., lit. 246–247°C)(14). ^{1}H NMR (500 MHz, $CDCl_3/CD_3OD$) 3.69 (3H, s, 3-OMe), 3.78 (3H, s, 7-OMe), 6.24 (1H, d, J 2Hz, H-6), 6.37 (1H, d, J 2Hz, H-8), 6.82 (2H, d, J 9Hz, H-3', H-5'), 7.89 (2H, d, J 9Hz, H-2', H-6'. ^{13}C NMR (DEPT-135, 125 MHz, $CDCl_3/CD_3OD$) 156.7 (C-2), 138.4 (C-3), 178.7 (C-4), 161.3 (C-5), 97.8 (C-6), 165.4 (C-7), 92.2 (C-8), 156.8 (C-9), 105.8 (C-10), 121.3 (C-1'), 130.2 (C-2'), 115.5 (C-3', C-5'), 159.9 (C-4'), 130.2 (C-2', C-6'), 59.9 (3-OMe), 55.6 (7-OMe). LR-EIMS m/z: 314.1 (M^+), 271.1, 256.2, 167.0, 149.0, 97.1, 57.1, 43.0.

Sitosteryl-3-O-6-palmitoyl-β-D-glucoside (2), white amorphous powder (8.4 mg). ^{1}H NMR (500 MHz, $CDCl_3/CD_3OD$) *Aglycone* 3.58 (1H, m, H-3), 5.36 (1H, m, H-6), 0.70 (3H, s, 18-CH_3), 1.03 (3H, s, 19-CH_3), 0.94 (3H, d, J 6.5Hz, 21-CH_3), 0.85 (3H, d, J 6.8Hz, 26-CH_3), 0.84 (3H, d, J 6.8Hz, 27-CH_3), 0.90 (3H, t, J 6.9Hz, 29-CH_3) *Sugar* 4.38 (1H, d, J 7.7Hz, H-1'), 3.34 (1H, m, H-2'), 3.58 (1H, m, H-3'), 3.34 (1H, m, H-4'), 3.48 (1H, ddd, J 2Hz, 5Hz, 10Hz, H-5'), 4.29 (1H, dd, J 2Hz, 12Hz, H-6'a), 4.48 (1H, dd, J 5Hz, 12Hz, H-6'b) *Fatty Acid* 2.37 (2H, t, J 7.5Hz, H-2''), 1.72 (2H, m, H3''), 1.28 (24H, broad s, H-4''-15''), 0.87 (3H, t, J 7Hz, H-16''). ^{13}C NMR (DEPT-135, 125 MHz, $CDCl_3/CD_3OD$) *Aglycone* 37.3 (C-1), 31.9 (C-2), 79.6 (C-3), 38.9 (C-4), 140.3 (C-5), 122.2 (C-6), 31.9 (C-7), 31.9 (C-8), 50.2 (C-9), 36.7 (C-10), 21.2 (C-11), 39.8 (C-12), 42.2 (C-13), 56.8 (C-14), 25.0 (C-15), 28.2 (C-16), 56.1 (C-17), 11.9 (C-18), 19.4 (C-19), 36.2 (C-20), 19.0 (C-21), 34.0 (C-22), 26.1 (C-23), 45.9 (C-24), 29.2 (C-25), 18.8 (C-26), 19.8 (C-27), 23.1 (C-28), 12.0 (C-29) *Sugar* 101.2 (C-1'), 73.6 (C-2'), 76.0 (C-3'), 70.1 (C-4'), 74.0 (C-5'), 63.2 (C-6') *Fatty acid* 174.7 (C=O), 34.2 (C-2''), 24.3 (C-3''), 28.9–29.7 (C-4''-14''), 22.6 (C-15''), 14.1 (C-16''). LR-EIMS m/z: 414.7 (aglycone), 396.4, 381.3, 368.4, 329.3, 284.3, 256.3, 241.2, 239.2, 227.0, 213.2.

β-sitosteryl galactoside **(3)**, white flakes (4.1 mg). ^{1}H NMR (500 MHz, $CDCl_3/ CD_3OD$) *Aglycone* 3.47 (1H, m, H-3), 5.25 (1H, m, H-6), 0.57 (3H, s, 18-CH_3), 0.90 (3H, s, 19-CH_3), 0.73 (3H, d, J 7Hz, 21-CH_3), 0.70 (3H, d, J 7Hz, 26-CH_3), 0.82 (3H, d, J 7.8Hz, 27-CH_3), 0.74 (3H, t, J 7.8Hz, 29-CH_3) *Sugar* 4.29 (1H, d, J 8Hz, H-1'), 3.12 (1H, dd, J 8Hz, 9Hz, H-2'), 3.32 (1H, dd, J 1Hz, 9Hz, H-3'), 3.32 (1H, dd, J 1Hz, 7Hz, H-4'), 3.18 (1H, m, H-5'), 3.63 (1H, dd, J 5Hz, 12Hz, H-6'a), 3.73 (1H, dd, J 3Hz, 12Hz, H-6'b). ^{13}C NMR (DEPT-135, 125 MHz, $CDCl_3/CD_3OD$) *Aglycone* 37.2 (C-1), 29.6 (C-2), 79.0 (C-3), 38.6 (C-4), 140.2 (C-5), 122.0 (C-6), 31.0 (C-7), 31.0 (C-8), 50.1 (C-9), 36.6 (C-10), 21.0 (C-11), 39.7 (C-12), 42.2 (C-13), 56.6 (C-14), 24.1 (C-15), 28.1 (C-16), 56.0 (C-17), 11.7 (C-18), 19.5 (C-19), 36.0 (C-20), 19.1 (C-21), 33.9 (C-22), 26.0 (C-23), 45.7 (C-24), 29.4 (C-25), 18.6 (C-26), 18.8 (C-27), 22.9 (C-28), 11.8 (C-29) *Sugar* 101.1 (C-1'), 73.4 (C-2'), 76.3 (C-3'), 70.2 (C-4'), 75.6 (C-5'), 61.7 (C-6'). LR-EIMS m/z 414.4 (aglycone), 397.4, 396.4, 381.4, 329.3, 288.3, 255.2, 213.2. Assignments were made by comparison with published data and confirmed by HMQC / COSY experiments.

Screening for antituberculosis activity

A Microplate Alamar Blue Assay (MABA) as described in the protocol of Collins and Franzblau [10] was used for anti-TB susceptibility of the extracts, fractions and purified compounds. *Mycobacterium tuberculosis* H_{37}Rv (ATCC 27294; American Type Culture Collection, Rockville MD) was grown at 37°C on a rotary shaker in Middlebrook 7H9 broth supplemented with 2% glycerol and 0.05% vv^{-1} Tween 80 until the culture density reached an optical density of 0.45–0.55 at 550 nm. Bacteria were pelleted, washed twice, resuspended in Dulbecco's phosphate-buffered saline, then filtered (8 μm) and aliquots frozen at -80°C. After a night, the stocks were thawed, sonicated and successively diluted to get the CFU. Rifampin was obtained from Sigma and stock solutions were made in accordance to the manufacturer's instructions. The assay was performed in black, clear-bottomed, 96-well microplates (Black view plates:Packard Instrument company, Meriden, Conn.) in order to reduce background fluorescence. Initial drug dilutions were prepared in either dimethyl sulfoxide or distilled ionized water, and subsequent twofold dilutions were performed in 0.1 mL

of 7H9GC (no Tween 80) in the microplates. BACTEC 12B-passaged inocula were initially diluted 1:2 in 7H9GC, and 0.1 mL was placed onto the wells. Frozen inocula were diluted 1:20 in BACTEC 12B medium followed by a 1:50 dilution in 7H9GC. Wells containing drug were used to monitor autofluorescence of compounds. Additional control wells consisted of bacteria only (B) and medium only (M). Plates were incubated at 37°C. At day 4 of incubation were added 20 µL of 10x Alamar Blue solution (Alamar Biosciences/Accumed, Westlake, Ohio) and 12.5 µL of 20% Tween 80 to one B well and one M well, and plates were reincubated at 37 °C. Detection of color change in wells was monitored at 12 and 24 h from blue to pink and for a measurement reading of > or equal to 50,000 fluorescence units (FU). Cytofluor II microplate flurometer (PerSeptive Biosystems, Framingham, Mass.) in bottom-reading setting at 530 nm for excitation and 590 nm for emission was used in fluorescence measurement. In case that a pink color was observed with B wells after 24h, the colorimetric reagent was added to the entire plate. If a blue color persisted in the well or a reading of < or equal to 50,000 FU was obtained, additional wells containing bacteria and medium were tested daily until a change in color was observed. At this point, reagents were added to other remaining wells. At 37 °C, the plates were incubated and the results were noted at 24 h post-reagent addition. Visual MIC's were defined as the lowest concentration of drug that resisted a color change. A background subtraction was performed on all wells with a mean triplicate M wells for fluorimetric MIC's. Percent inhibition was defined as 1 – (test well FU/mean FU of triplicate B wells) × 100. The lowest drug concentration exhibiting an inhibition of > or equal to 90% was assigned as the MIC.

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Supplementary data for compounds 1–3

Compound 1 (kumatakenin)

Physical Properties of **1**:
Yellow needles, Rf: 0.5 in benzene acetone (6.4:3.6); 0.72 in diethyl ether-ethyl acetate (7:3), 0.37 in benzene-chloroform-methanol (5:4:1) and 0.38 in chloroform-acetone (9:1)
Melting point: 248–249 °C (uncorrected, lit. 246–247 °C) (Kimura et al., 1967)
UV: 269 nm ($\log_\varepsilon$ = 9.32) 352 nm ($\log_\varepsilon$ = 9.44)
IR: 3243.86 cm^{-1}, 1665.45 cm^{-1}, 1600.76 cm^{-1}, 1600.79 cm^{-1}
LR-EIMS: m/z (rel. int.) m/z 314 (100)(M^+), 271 (25), 167 (38), 149 (55), 97 (24), 57 (68), 43 (67).

Compound 2 and 3

Physical Properties of (**3**)
White amorphous solid, Rf: 0.19 in benzene-acetone (6.4:3.6); 0.20 in diethyl ether-ethyl acetate (7:3), 0.15 in benzene-chloroform-methanol (5:4:1)
Melting point: not determined
UV: not determined
IR: 3419.45 cm^{-1} , 2559.3 cm^{-1}, 1073.94 cm^{-1}
LR-EIMS: m/z 396, 381, 329, 255.2

Physical properties of (**2**)
White amorphous solid, Rf: 0.13 in benzene- acetone (6.4:3.6); 0.26 in diethyl ether-ethyl acetate (7:3), 0.31 in benzene-chloroform-methanol (5:4:1) and 0.05 in chloroform-acetone (9:1)
Melting point: not determined
UV: 220 nm ($\log_\varepsilon$ = 6.43)
IR: 3439.20 cm^{-1} , 1738.78 cm^{-1}, 1600.76 cm^{-1}, 2923.87 cm^{-1}, 2852.06 cm^{-1}, 1467.34 cm^{-1}
LR-EIMS: Fragment ion peaks at m/z 414, 396, 381, 368, 329.3, 256.3, 241.2 and 227

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Table 2. Comparison of the 13C NMR Spectral Data of 2 and 3 with the Reported Data of Steryl Glycosides

Position	3 δC ppm	2 δC ppm	Pei-Wu, 1988* δ_c ppm	Della Greca, 1990** $\delta_c$ppm	Shaiq-Ali, 2002*** δ_c ppm	Crouch 2007+ δ_c ppm
Aglycone						
1	**37.2**	**37.3**	**36.75**	**37.33**	**37.0**	
2	**29.6**	**31.9**	**30.02**	**31.63**	**29.5**	
3	**79.0**	**79.6**	**79.86**	**71.73**	**79.7**	
4	**38.6**	**38.9**	**39.02**	**42.20**	**38.7**	
5	**140.2**	**140.3**	**140.48**	**140.71**	**140.2**	
6	**122.0**	**122.2**	**122.08**	**121.63**	**122.3**	
7	**31.0**	**31.9**	**32.0**	**31.96**	**32.03**	
8	**31.0**	**31.9**	**32.14**	**31.81**	**31.9**	
9	**50.1**	**50.2**	**50.25**	**51.13**	**50.1**	
10	**36.6**	**36.7**	**36.76**	**36.43**	**36.6**	
11	**21.0**	**21.2**	**22.75**	**21.09**	**21.8**	
12	**39.7**	**39.8**	**39.68**	**39.79**	**39.6**	
13	**42.2**	**42.2**	**42.41**	**42.37**	**42.2**	
14	**56.6**	**56.8**	**56.67**	**56.75**	**56.6**	
15	**24.1**	**25.0**	**25.09**	**24.15**	**24.2**	
16	**28.1**	**28.2**	**29.25**	**28.25**	**28.1**	
17	**56.0**	**56.1**	**56.31**	**56.02**	**55.9**	
18	**11.7**	**11.9**	**11.92**	**11.84**	**11.7**	
19	**19.5**	**19.4**	**19.47**	**19.46**	**19.2**	
20	**36.0**	**36.2**	**36.75**	**36.07**	**36.0**	
21	**19.1**	**19.0**	**19.10**	**18.68**	**18.21**	
22	**33.9**	**34.0**	**32.04**	**33.95**	**34.0**	
23	**26.0**	**26.1**	**26.4**	**26.10**	**26.0**	
24	**45.7**	**45.9**	**45.9**	**45.82**	**45.7**	
25	**29.4**	**29.2**	**29.25**	**29.15**	**29.2**	
26	**18.6**	**18.8**	**18.86**	**19.77**	**19.6**	
27	**18.8**	**19.8**	**19.86**	**19.21**	**18.9**	
28	**22.9**	**23.1**	**23.1**	**23.13**	**23.0**	
29	**11.8**	**12.0**	**12.02**	**11.04**	**11.8**	
Glucose						
1’		**101.2**	**101.4**		**99.6**	
2’		**73.6**	**73.76**		**71.4**	
3’		**76.0**	**73.42**		**71.8**	
4’		**70.1**	**70.59**		**68.7**	
5’		**74.0**	**73.42**		**72.9**	
6’		**63.2**	**63.77**		**61.1**	

Position	(3) $\delta_c$ppm	(2) $\delta_c$ppm	Pei-Wu, 1988* $\delta_c$ppm	Della Greca, 1990** $\delta_c$ppm	Shaiq-Ali, 2002*** $\delta_c$ppm	Ahmad, 1992**** $\delta_c$ppm	Crouch 2007+δ_c ppm
Galactose							
1’	**101.0**					**101.0**	**102.6**
2’	**73.4**					**73.4**	**75.1**
3’	**76.3**					**75.5**	**78.2**
4’	**70.2**					**70.0**	**71.6**
5’	**75.6**					**76.2**	**77.8**
6’	**61.7**					**61.8**	**62.7**
Fatty Acid							
1’’	**174.7**	**174.7**	**174.15**		**174.6**		
2”	**34.2**	**34.2**	**-**		**32.5**		
3”	**24.3**	**24.3**	**-**		**31.3,**		
4”-15”	**29.7-28.9/22.6**	**29.7-28.9/22.6**	**-**		**29.1-28.2**		
16”	**14.1**	**14.1**	**14.6**		**14.5**		

*Sitosterol-3-O-6-stearoyl-β-D-glucoside, in $CDCl_3$ (50.3 MHz)

**β-Sitosterol, in $CDCl_3$ (100 MHz)

***Sitosteryl-3-O-6-palmitoyl-β-D-glucoside, in $CDCl_3$, (75 MHz)

****sterol galactoside

+bufadienolide galactoside, in CD3OD, 100 MHz

Table 3: Comparison of Selected Protons in (2) and (3) with Reported Data of Sterol Glycosides

3 δ_H, ppm	2 δ_H, ppm	H No.	Shaiq-Ali, 2002* δ_H, ppm	Gomes, 1998** δ_H, ppm	Ahmed, 1992*** δ_H, ppm
Aglycone					
3.47 m	3.58 m	**3**	3.42 m		
5.25 m	5.37 m	**6**	5.33 distorted triplet		
0.57 s	0.70 s	**Me-18**	Unassigned		
0.90 s	1.03 s	**Me-19**	0.97		
0.73 d (⌡ = 7.0)	0.94 d (⌡ = 6.5)	**Me-21**	0.91 d (⌡ = 6.5)		
0.70 d (⌡ = 7.0)	0.85 d (⌡ = 6.8)	**Me-26**	0.82 – 0.81 d and t (⌡ = 6.5)		
0.82 d (⌡ = 7.4)	0.84 d (⌡=6.8)	**Me-27**	0.82 – 0.81 d and t (⌡= 6.5)		
0.74 t (⌡ = 7.8)	0.90 t (⌡=6.9)	**Me-29**	0.82 – 0.81 d and t (⌡ = 6.5)		
Galactose	Glucose	**Glucose**	Glucose	Glucose	**Galactose**
4.29 d (⌡ = 8)	4.40 d (⌡ = 7.7)	**1'**	4.52 d (⌡ =7.9)	4.40 d (⌡ = 8)	4.20 (d, ⌡=7.8)
3.12 dd (⌡ = 8, 9)	3.34 m	**2'**	5.01 - 4.91 m	3.31 m	3.55 (dd, J=8,9)
3.32 dd (⌡ = 1, 9)	3.58 m	**3'**	5.24 t (⌡ = 8)	3.31 m	3.15 (dd, J=3,9)
3.32 dd (⌡ = 1, 7)	3.34 m	**4'**	5.01 - 4.91 m	3.31 m	3.65 (dd, J= 1,3)
3.18 m	3.48 ddd (⌡ = 2, 5, 10)	**5'**	5.24 t (⌡ = 8)	3.70 m	3.03 (ddd,J=1,5,7)
3.63 dd (J 5. 12)	4.29 dd (⌡= 2, 12)	**6'a**	4.20 – 4.05 m	4.30 dd (⌡= 12, 2.5)	3.30 (dd, J=5,11)
3.73 dd (⌡ = 3, 12)	4.48 dd (⌡= 5, 12)	**6'b**		4.43 dd (⌡ = 12, 5.2)	3.52 (dd, J=7,11)
	2.37 t (⌡ = 7.5)	**2"**	2.20 d (⌡ = 7.1)	2.34 t (⌡= 7.7)	
		3"			
	1.72 m 1.28 br s	**4"-15"/17"**	Not mentioned	1.26 br s (very intense)	
	0.87 t (⌡ = 7)	**18"**	0.86 t (⌡ = 7)	0.90 s	

*Sitosteryl-3-O-6-palmitoyl-β-D-glucoside, in $CDCl_3$, (75 MHz)

**3-O-[6'-O-palmitoyl-β-D-glucosyl]-spinasta-7- 22(23)-diene, in $CDCl_3$, (50.3 MHz)

***3-O-β-D-galactopyranosyl-stigmasta-5,22-diene, CDCIs at 300 MHz.

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