



Spirostanol glucosides from the leaves of *Cestrum laevigatum* L.

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ABSTRACT

Two new steroidal saponins, (25R)-spirost-5-ene-3 β ,26 β -diol 3-O- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)-[(1 $\rightarrow$ 2)- α -L-rhamnopyranosyl]- β -D-glucopyranoside (**1**) and (25R)-spirost-6-ene-3 β ,5 β -diol 3-O- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)-[(1 $\rightarrow$ 2)- α -L-rhamnopyranosyl]- β -D-glucopyranoside (**2**), along with the known diosgenin 3-O- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- β -D-glucopyranoside (**3**), chonglouoside SL-5 (**4**) and Paris saponin Pb (**5**) were isolated from the leaves of *Cestrum laevigatum*. The structures of the compounds were determined using spectroscopic analyses including HRESI-MS, 1D and 2D NMR data, followed by comparison with data from the literature. Among them, two are particularly unique, compound **1** is the first Δ^5 -spirostanol saponin and compound **2** has an unusual C-26 hydroxyl in the Δ^5 -spirostanol skeleton. Antifungal testing showed a potent activity to formosanin C against *Candida albicans* and *Candida parapsilosis*. Evaluation of the cytotoxic activity indicated that compound **1** has a moderate activity against HL-60 and SF-295 cell lines, while compound **2** were active only against HL-60.

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1. Introduction

Cestrum is the second largest genus of the Solanaceae family, with 150 species distributed in tropical and subtropical America, 50 of which occurring in Brazil [1–2]. As particular to some Solanaceae genera, several species have been reputed as poisonous plants and the search for active compounds revealed a prolific source of bioactive steroids saponins containing spirostanol or furostanol glycoside skeletons [2–7].

Cestrum laevigatum L. is an evergreen shrub native to South America and has been introduced to South Africa. It is widespread into natural grasslands, forests, riparian habitats, and coastal dunes, where is popularly known as “coerana”, “lady of the night” and “corana”. The dried leaves are used in traditional medicine as treatment for malaria and fever [8], and smoked by the Mapuche Indian of southern Chile as a substitute for cannabis [9]. Although commonly used for ornamental purposes, *C. laevigatum* is consid-

ered the most lethal plant to mammals, among the group of common invasive species that cause liver damage. For this reason, is one of toxic plants of greater importance in Brazilian livestock for its wide distribution and economic losses, and its growth is severely controlled or eradicated into pastureland [10,11].

In the course of our search for bioactive natural compounds we report the isolation and structural characterization of two new sapogenins (25R)-spirost-5-ene-3 β ,26 β -diol 3-O- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)-[(1 $\rightarrow$ 2)- α -L-rhamnopyranosyl]- β -D-glucopyranoside (**1**) and (25R)-spirost-6-ene-3 β ,5 β -diol 3-O- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)-[(1 $\rightarrow$ 2)- α -L-rhamnopyranosyl]- β -D-glucopyranoside (**2**) from the leaves of *C. laevigatum*, in addition to the known diosgenin 3-O- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- β -D-glucopyranoside (**3**) [12], chonglouoside SL-5 (**4**) [13] and formosanin C (**5**) [14]. The screening for antimicrobial activities against *Candida parapsilosis* (ATCC[®] 22019TM), *Candida albicans* (ATCC[®] 10231TM), *Candida krusei* (ATCC[®] 14243TM), *Pseudomonas aeruginosa* (ATCC[®] 9027TM), *Staphylococcus aureus* (ATCC[®] 6538TM) and *Bacillus subtilis* (ATCC[®] 6633TM), and evaluation of cytotoxicity using human

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promyelocytic leukemia (HL-60), ovarian carcinoma (OVCAR-8), colorectal adenocarcinoma (HCT-116) and glioma (SF-295) cell lines were performed (see Fig. 1).

2. Experimental

2.1. General experimental procedures

Melting points were obtained on a MettlerToledo-FP82HT and were uncorrected. IR spectra were recorded as KBr pellets on a Perkin-Elmer FT-IR Spectrum 1000 using KBr. The NMR spectra were performed on Bruker Avance DRX 300 or on Avance DRX500 MHz equipped with an inverse detection probe head and z-gradient accessory. The ^1H and ^{13}C chemical shifts are expressed in the δ scale and were referenced to TMS through the residual solvent. High-resolution mass spectra were recorded on a Waters Acquity UPLC system coupled with a Quadrupole/Time-of-Flight system (UPLC/Qtof MSE spectrometer) in the positive mode. The TOF conditions were as follow: source temperature 120 °C; desolvation temperature 350 °C; desolvation gas flow of 350 L/h; capillary voltage 2 kV; collision Energy Ramp 20 eV. The mode was acquired from 110 to 1200 Da. Optical rotations were obtained on a Perkin-Elmer Q-2000 polarimeter, at 589 nm and 25 °C. Column chromatography was performed over Sephadex LH-20 (Pharmacia) and SPE C18 cartridge (Phenomenex), while TLC was performed on precoated silica gel aluminum sheets (Merck). The compounds were visualized by UV detection and by spraying with vanillin/perchloric acid/EtOH solution, followed by heating.

2.2. Plant material

C. laevigatum L. was collected at the Guaramiranga Mountain, Pacoti, Ceará State, Northeast of Brazil. Voucher specimens (#38643) were deposited at the Herbário Prisco Bezerra (EAC) and identified by MSc. Edson de Paula Nunes, Departamento de Biologia, Universidade Federal do Ceará, Ceará, Brazil.

2.3. Extraction and isolation

Leaves of *C. laevigatum* (1.14 kg) were pulverized and extracted with EtOH (3 × 9.0 L) at room temperature. The solvent was removed under reduced pressure to yield a dark green residue (131.4 g).

Part of EtOH extract (91.0 g) was dissolved in a mixture of MeOH:H₂O (1:1 v/v) and submitted to liquid–liquid partition chromatography with hexane, CHCl₃, EtOAc and *n*-BuOH to give four fractions: hexane (0.11 g), CH₂Cl₂ (27.1 g), EtOAc (11.3 g) and *n*-BuOH (32.0 g).

An aliquot of the *n*-BuOH fraction (2.0 g) was rechromatographed on Sephadex LH-20 (10.0 g) (column 2.5 cm × 10 cm) to afford forty-five fractions (3.0 mL) that were pooled together into five resulting sub-fractions after TLC analysis. Sub-fraction F-3 (1.1 g) was further chromatographed on a SPE C18 (5.0 g) cartridge using MeOH/H₂O 1:1 (30.0 mL), MeOH/H₂O 7:3 (3.00 mL), MeOH/H₂O 8:2 (40.0 mL), MeOH/H₂O 9:1 (30.0 mL) and MeOH (50.0 mL), yielding five fractions. Sub-fraction F-5 (0.360 g) was submitted to semi-preparative RP-18 HPLC analysis, using MeOH/H₂O (87:13) as eluent, to afford **1** (10.6 mg).

An Aliquot of the EtOAc fraction (2.1 g) was rechromatographed on Sephadex LH-20 (10.0 g) (column 2.5 cm × 10 cm) to give forty sub-fractions (3.0 mL), which were combined into four resulting sub-fractions according to TLC analysis. Sub-fraction F-3 (0.70 g) which was then chromatographed on a SPE C18 cartridge by elution with MeOH/H₂O 1:1 (30.0 mL), MeOH/H₂O 7:3 (30.0 mL), MeOH/H₂O 8:2 (40.0 mL), MeOH/H₂O 9:1 (30.0 mL) and MeOH (40.0 mL), yielding five fractions. Fraction F-3 (0.26 g) was submitted to semi-preparative RP-18 HPLC chromatography, using an isocratic mixture MeOH/H₂O 8:1 to afford **2** (10.0 mg), **3** (8.0 mg), **4** (10.5 mg) and **5** (19.5 mg).

2.3.1. (25R)-spirost-5-ene-3 β ,26 β -diol 3-O- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)-[(1 $\rightarrow$ 2)- α -L-rhamnopyranosyl]- β -D-glucopyranoside (**1**)

White amorphous powder, mp 240–242 °C; $[\alpha]_D^{20}$ -172.8 (c 0.10, MeOH); Rf 0.22 (CH₂Cl₂/MeOH 25%); ^1H and ^{13}C NMR data: see

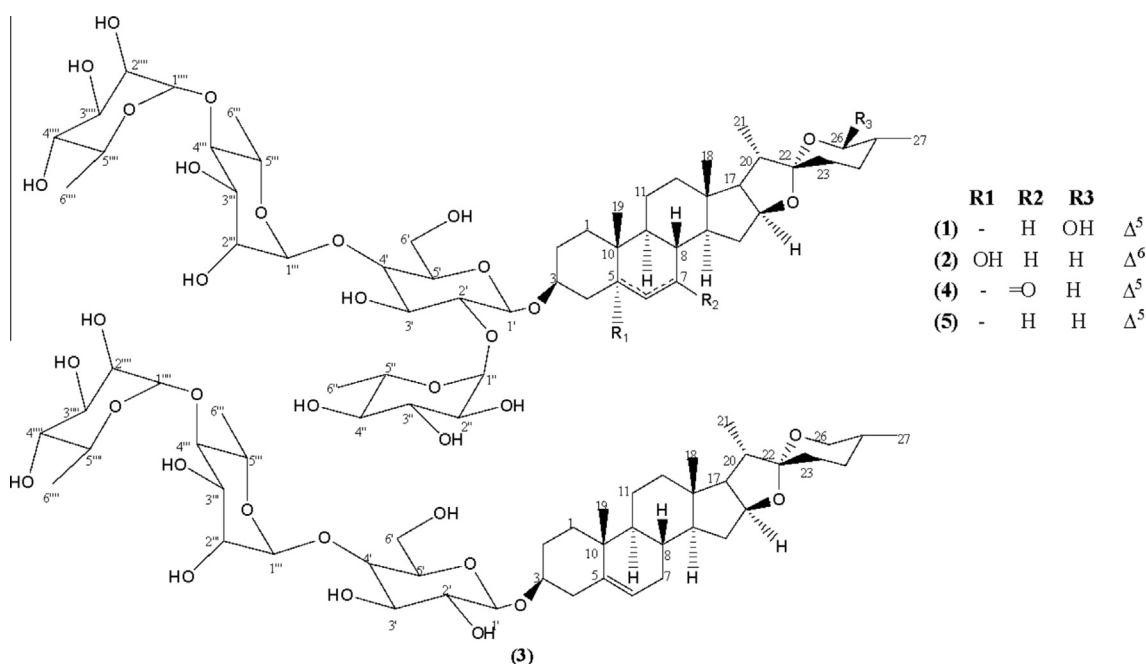


Fig. 1. Structures of compounds 1–5.

Table 1
NMR spectral data of compounds **1** and **2** (^1H – 500 MHz; ^{13}C – 125 MHz; CD_3OD).

Position	1		2	
	δ_{C}	δ_{H}	δ_{C}	δ_{H}
1	38.1	1.82 (m) 1.02 (m)	29.6	1.68 (m) 1.40 (m)
2	30.6	1.90 (m) 1.60 (m)	32.3	1.79 (m) 1.57 (m)
3	79.4	3.54 (m)	75.2	4.17 (m)
4	39.6	2.44 dd (12.5, 3.0) 2.35 t (12.5)	39.3	2.50 (m) 2.36 t (12.7)
5	142.1	–	84.2	–
6	122.8	5.38 d (4.4)	134.4	5.72 d (10.2)
7	33.3	2.00 (m) 1.58 (m)	131.7	5.62 dd (10.2, 2.1)
8	33.0	1.28 (m)	39.8	2.17 (m)
9	51.2	0.90 (m)	45.4	1.80 (m)
10	38.2	–	39.6	–
11	22.2	1.55 (m)	21.7	1.54 (m) 1.37 (m)
12	41.1	1.78 (m) 1.20 (m)	41.8	1.75 (m) 1.19 (m)
13	41.6	–	42.7	–
14	58.0	1.18 (m)	54.7	1.40 (m)
15	–	1.70 (m) 1.58 (m)	32.4	2.10 (m) 1.40 (m)
16	82.5	4.58 (m)	82.3	4.40 (m)
17	63.9	1.82 (m)	63.5	1.80 (m)
18	16.9	0.80 (s)	17.0	0.82 (s)
19	20.0	1.04 (s)	16.0	0.98 (s)
20	43.1	1.92 (m)	42.8	1.92 (m)
21	15.0	1.01 d (6.5)	14.9	0.96 d (6.9)
22	113.8	–	110.6	–
23	32.9	2.00 (m) 1.50 (m)	33.2	2.50 (m) 1.50 (m)
24	29.2	1.60 (m) 1.50 (m)	29.8	1.90 (m) 1.60 (m)
25	38.2	1.32 (m)	31.3	1.60 (m)
26	97.2	4.52 d (8.0)	67.9	3.45 (m) 3.38 (m)
27	17.3	0.93 d (8.0)	17.6	0.79 d (6.4)
β -D-glucopyranosyl(1 $\rightarrow$ 3)				
1'	100.6	4.49 d (8.0)	100.8	4.50 d (7.8)
2'	79.5	3.40 (m)	79.4	3.40 (m)
3'	78.1	3.56 (m)	78.0	3.58 (m)
4'	79.6	3.52 (m)	79.9	3.54 (m)
5'	76.8	3.30 (m)	76.6	3.32 (m)
6'	62.1	3.80 (m) 3.65 (m)	62.0	3.81 (m) 3.68 (m)
α -L-rhamnopyranosyl(1 $\rightarrow$ 2)				
1''	102.5	5.18 d (1.5)	102.3	5.20 (s)
2''	73.0	3.78 (m)	72.0	3.94 (m)
3''	72.5	3.65 (m)	72.3	3.61 dd (9.5, 3.2)
4''	74.0	3.38 (m)	73.8	3.40 (m)
5''	69.2	4.05 (m)	69.7	4.12 (m)
6''	18.7	1.29 d (6.5)	17.9	1.25 d (6.0)
α -L-rhamnopyranosyl(1 $\rightarrow$ 4)				
1'''	102.8	4.83 d (1.5)	102.6	4.83 d (1.3)
2'''	72.5	3.93 (m)	72.8	3.79 (m)
3'''	73.0	3.76 (m)	72.9	3.75 (9.1, 3.1)
4'''	81.0	3.57 (m)	80.8	3.55 (m)
5'''	70.0	4.14 (m)	69.2	4.03 (m)
6'''	18.0	1.24 d (5.5)	18.6	1.29 d (6.2)
α -L-rhamnopyranosyl(1 $\rightarrow$ 4)				
1''''	103.3	5.18 d (1.5)	103.0	5.19 (s)
2''''	72.5	3.93 (m)	72.0	3.94 (m)
3''''	72.3	3.60 (m)	72.3	3.68 (m)
4''''	74.1	3.38 (m)	73.9	3.40 (m)
5''''	70.6	3.70 (m)	70.4	3.70 (m)
6''''	18.1	1.25 d (6.5)	18.0	1.25 d (6.0)

Table 1; IR (KBr) 3355; 2928; 1640; 1452; 1036 cm^{-1} . HRESIMS m/z : 1031.5468 $[\text{M}+\text{H}]^+$ (calcd for $\text{C}_{51}\text{H}_{83}\text{O}_{21}$ $[\text{M}+\text{H}]$, 1031.5427).

2.3.2. (25R)-spirost-6-ene-3 β ,5 β -diol 3-O- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)-[(1 $\rightarrow$ 2)- α -L-rhamnopyranosyl]- β -D-glucopyranoside (**2**)

White amorphous powder, mp 178–179 °C; $[\alpha]_{\text{D}}^{20}$ -132.7 (c 0.16, MeOH); Rf 0.20 ($\text{CH}_2\text{Cl}_2/\text{MeOH}$ 25%); ^1H and ^{13}C NMR data: see Table 1; IR (KBr) 3362; 2928; 1653; 1453; 1037. HRESIMS m/z : 1031.5427 $[\text{M}+\text{H}]^+$ (calcd for $\text{C}_{51}\text{H}_{83}\text{O}_{21}$ $[\text{M}+\text{H}]$, 1031.5427).

2.3.3. (25R)-spirost-5-ene-3 β -ol 3-O- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- β -D-glucopyranoside (**3**)

$[\alpha]_{\text{D}}^{20}$ -55.5° (c 0.09; MeOH) $\{[\alpha]_{\text{D}}^{24}$ -62.0 (c 0.09; MeOH) [15]; Rf 0.50 ($\text{CH}_2\text{Cl}_2/\text{MeOH}$ 25%).

2.3.4. (25R)-7-oxo-espirost-5-ene-3 β -ol 3-O- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)-[(1 $\rightarrow$ 2)- α -L-rhamnopyranosyl]- β -D-glucopyranoside (**4**)

$[\alpha]_{\text{D}}^{20}$ -115.8° (c 0.11; MeOH) $\{[\alpha]_{\text{D}}^{15}$ -32.8° (c 0.12; MeOH) [13]; Rf 0.23 ($\text{CH}_2\text{Cl}_2/\text{MeOH}$ 25%).

2.3.5. (25R)-espirost-5-ene-3 β -ol 3-O- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)-[(1 $\rightarrow$ 2)- α -L-rhamnopyranosyl]- β -D-glucopyranoside (**5**)

$[\alpha]_{\text{D}}^{20}$ -116.7° (c 0.20; MeOH) $\{[\alpha]_{\text{D}}^{21}$ -118.6° (c 0.20; MeOH) [15]; Rf 0.23 ($\text{CH}_2\text{Cl}_2/\text{MeOH}$ 25%).

2.4. Acid hydrolysis and sugar analysis

Compounds **1** and **2** (5.0 mg) were dissolved in 2 M HCl (dioxane/ H_2O , 1:1, 2 mL) and stirred at 90 °C for 2 h. After cooling, the reaction mixture was neutralized with solution of 1 M NaOH, extracted with CH_2Cl_2 (3 $\times$ 5 mL), and the aqueous layer was evaporated to give a mixture of monosaccharides. The residue was dissolved in hexamethyldisilazane/trimethylchlorosilane/pyridine (3:1:9), and stirred at 70 °C for 60 min. The supernatants (3 μL) were analyzed by GC Agilent model GC-789/MSD-5977A (quadrupole), under the following conditions: CP-ChiraSil-L-Val column, 0.25 mm $\times$ 25 m; temperatures for detector and injector 150 and 200 °C respectively; temperature gradient system for the oven, 100 °C for 1 min and then raised to 180 °C; rate 5 °C/min kept for 5 min. The configurations of sugars were determined by comparison of the retention times of the corresponding derivatives with those of standards treated simultaneously with same silylating reagents (L-rhamnose 11.15 and D-glucose 16.57). Peaks of the hydrolysates of **1** and **2** were detected at 11.20 (L-rhamnose) and 16.57 (D-glucose). Co-injection of each hydrolysate with standard D-glucose and L-rhamnose gave single peaks.

2.5. Cytotoxic activity

The tested tumor cell lines (colorectal adenocarcinoma HCT-116, ovarian carcinoma OVCAR-8, human promyelocytic leukemia HL-60 and glioma SF-295) were kindly donated by the National Cancer Institute (Bethesda, MD, USA). Cells were maintained in RPMI 1640 medium supplemented with 10% fetal bovine serum, 2 mM glutamine, 100 U mL^{-1} penicillin, and 100 $\mu\text{g mL}^{-1}$ streptomycin at 37 °C with 5% CO_2 . The cytotoxicity of the isolated compounds was tested the against tumor cell lines using the 3-(4,5-dimethyl-2-thiazolyl)-2,5-diphenyl-2H-tetrazolium bromide (MTT) (Sigma Aldrich Co., St. Louis, MO/USA) reduction assay [16]. Cells were plated in 96-well plates (10^5 cells per well for adherent cells or 3×10^5 cells per well for suspended cells in 100 μL of medium) and compounds (0.05–25 $\mu\text{g mL}^{-1}$) were dissolved in DMSO, added to each well using the HTS – high-throughput screening-biomek 3000-Beckman Coulter (Inc. Fullerton, California, USA), and incubated for 72 h. Doxorubicin was used as

the positive control. Control groups received the same amount of DMSO. After 69 h of incubation, the supernatant was replaced by fresh medium containing MTT (0.5 mg mL⁻¹). Three hours later, the MTT formazan product was dissolved in 150 µL of DMSO, and the absorbance was measured at 595 nm (DTX 880 Multimode Detector, Beckman Coulter, Inc. Fullerton, CA, USA).

2.6. Antimicrobial activity

2.6.1. In vitro antibacterial activity

Tests were performed according to the M02-A11 (CLSI, 2012) protocol with modifications [17]. Wells with 6 mm diameter were made in the agar overlay of the Petri dish [18]. To those wells, a volume of 20 µL (1000 µg mL⁻¹) of the obtained compounds was applied. The plates were incubated for 20 h at 35 °C. The solvents and diluents used in the compounds dissolution were used as negative control.

2.6.2. In vitro antifungal activity

The broth microdilution (BMD) antifungal susceptibility test was performed according to M27-A3 protocol using RPMI broth (pH 7.0) buffered with 0.165 M MOPS [3-(*N*-morpholino)propane-sulfonic acid] (Sigma–Aldrich, St Louis, MO, USA) [19]. Compounds were dissolved in dimethyl sulfoxide (DMSO; Sigma–Aldrich) and tested at concentrations ranging from 1.95 to 1000 µg mL⁻¹. The yeasts and compounds were incubated in 96-well culture plates at 35 °C for 24 h and the results were examined visually. The minimum inhibitory concentration (MIC) of each compound was determined as the concentration that inhibited 50% of fungal growth.

3. Results and discussion

Compound **1** was isolated as a white solid. Its molecular formula was defined as C₅₁H₈₂O₂₁ by the protonated molecular ion peak at *m/z* 1031.5468 [M+H] (calcd 1031.5427) in the HRESIMS spectrum. The IR spectrum displayed an absorption band relative to hydroxyl at 3355 cm⁻¹.

The ¹H NMR spectrum revealed the presence of two singlet methyls at δ_H 0.80 (s, H-18), 1.04 (s, H-19) and five doublets at δ_H 0.93 (*J* = 8.0 Hz, H-27), 1.01 (*J* = 6.5 Hz, H-21), 1.24 (*J* = 5.5 Hz, H-6'''), 1.29 (d, *J* = 6.5 Hz, H-6'') and 1.25 (*J* = 6.5 Hz, H-6'''), besides a vinyl hydrogen attached of a trissubstituted double bond at δ_H 5.38 (H-6, *J* = 4.4 Hz), and several signals in the range at δ_H 3.30–5.18, indicating **1** to be a steroidal glycoside.

The ¹³C NMR spectrum displayed forty-four signals, some of which were oxygenated and superimposed and twenty-seven others attributable to the aglycone moiety. The presence of characteristic signals related to one spiro acetal carbon at δ_C 113.8 (C-22), besides two olefinics at δ_C 142.1 (C-5) and 122.8 (C-6), after comparison to the literature data for steroidal sapogenins containing a Δ⁵-spirostanol skeleton, showed a good match with those related to gitogenin [20]. The only difference was the replacement of the signal of the oxymethylene C-26 in gitogenin at δ_C 67.1, for one additional dioxymethylene carbon at δ_C 97.2 in **1**, suggesting hydroxylation of C-26 on the ring F.

All proton and carbon signals were fully assigned heteronuclear single-quantum coherence (HSQC) and HMBC spectra (Table 1). The four sugar units were characterized by the HSQC correlations of the signals of the acetal carbon at δ_C 100.6 (C-1') with the anomeric proton at δ_H 4.49 (d, *J* = 8.0 Hz, H-1'), the carbon at δ_C 102.8 (C-1'') with the hydrogen at δ_H 4.83 (d, *J* = 1.5 Hz, H-1''), while the two others at δ_C 102.5 (C-1''') and 103.3 (C-1''') showed correlation with the signal at δ_H 5.18 (d, *J* = 1.5 Hz). This suggestion was corroborated by sequential loss of the four hexoses moieties

showed by the fragment ion peaks at *m/z* at 885.4839, 739.4393, 557.3475 and 413.3046 in the HRESIMS spectrum. The identity of the monosaccharides as one D-glucose and three L-rhamnose units were carried out by NMR analyses and comparison with the literature data [20], whose absolute configuration of glucose was determined as D and those of the rhamnoses as L, on the basis of the acid hydrolysis and HPLC analysis as described. A β-anomeric configuration of the glucopyranosyl moiety was supported by the relatively large vicinal coupling constant value of 8.0 Hz for the anomeric proton at δ_H 4.49 (H-1'), while the α-anomeric configuration of the three rhamnopyranosyl units at δ_H 4.83 (H-1''') and 5.18 (H-1'' and H-1'''), was deduced by the value of *J* = 1.5 Hz.

The HMBC spectrum permitted to establish the sequence of all sugar units by long-range correlations between the proton signal at δ_H 5.18 (H-1''') of the first rhamnopyranose unit with the carbon at δ_C 81.0 (C-4'') of the second, whose anomeric hydrogen at δ_H 4.83 (H-1'') showed correlation with the carbon at δ_C 79.6 (C-4') of the glycopyranose. The anomeric hydrogen at δ_H 5.18 (H-1'') of third rhamnopyranose also showed correlation with the glycopyranose moiety through the carbon at δ_C 79.5 (C-2'). Finally, the anomeric hydrogen of the glycopyranose unit at δ_H 4.49 (H-1') showed correlation with the C-3 carbon of the aglycone portion at δ_C 79.3 (C-3). These findings confirmed the attachment sequence of the sugar chain at C-3 to be 3-O-α-L-rhamnopyranosyl-(1 → 4)-α-L-rhamnopyranosyl-(1 → 4)-[(1 → 2)-α-L-rhamnopyranosyl]-β-D-glucopyranoside. The locations of the 26-hydroxyl and 27-methyl group equatorially positioned were confirmed by long range correlations between the methyl group at δ_H 0.93 (CH₃-27) with the acetal carbon at δ_C 97.2 (C-26), and based on the large value of the coupling constant of 8.0 Hz observed for the dioxymethylene hydrogen at δ_H 4.52 (H-26). Thus, the structure of **1** was elucidated as the new (25*R*)-spirost-5-ene-3β,26β-diol 3-O-α-L-rhamnopyranosyl-(1 → 4)-α-L-rhamnopyranosyl-(1 → 4)-[(1 → 2)-α-L-rhamnopyranosyl]-β-D-glucopyranoside.

Compound **2** was isolated as a yellow resin. The support for a spirostanol steroid skeleton came from the comparative analysis of its MS and NMR data with those observed for compound **1**. It had also a molecular formula C₅₁H₈₃O₂₁ on the basis of HRESIMS ions peak at *m/z* 1031.5427 (calcd 1031.5427).

The ¹H NMR spectral data of **2** were also quite similar to those of **1**, except the by the absence of the dioxymethylene hydrogen at δ_H 4.52 (H-26) observed in **1**, and the presence of an additional vinyl hydrogens as a pair of doublets at δ_H 5.62 (dd, *J* = 10.2, 2.1 Hz, H-7) and 5.72 (d, *J* = 10.2 Hz, H-6) in **2**, instead of the single signal in **1** at δ_H 5.38 (H-6), indicating that the trissubstituted double bond in **1** was rearranged to a *cis* configuration in **2**.

In the ¹³C NMR spectrum of **2** was identified an additional oxygenated quaternary carbon at δ_C 84.2 (C-5), and similar signals related to the four sugar anomeric carbons at δ_C 100.8 (C-1'), 102.3 (C-1''), 102.6 (C-1''') and 103.0 (C-1''') showing HSQC correlations with corresponding hydrogens signals at δ_H 4.50 (d, *J* = 7.8 Hz, H-1'), 5.20 (d, *J* = 1.5 Hz, H-1''), 4.83 (d, *J* = 1.3 Hz, H-1''') and 5.19 (d, *J* = 1.5 Hz, H-1'''), respectively, in addition to deoxygenated monohydrogenated carbon at δ_C 110.6 (C-22). Moreover, the *cis* disubstituted double bond was confirmed by the two signals of unsaturated monohydrogenated carbons at δ_C 131.7 (C-7) and 134.4 (C-6), showing correlations with the hydrogens at δ_H 5.62 (H-7) and 5.72 (H-6), respectively.

The assignments of the signals related to the sugar carbons, as well as their sequence were established by long-range connectivities in the HMBC spectrum (Table 1). Moreover, the location of the hydroxylated carbon at C-5 and the Δ⁶-double bond was determined by the three bond correlations of the hydrogens of the methyl group at δ_H 0.98 (CH₃-19) and olefin proton at δ_H 5.62 (H-7) with the oxygenated carbon at δ_C 84.2 (C-5), besides correlations of the hydrogens at δ_H 2.17 (H-8) and 2.50 (H-4) with the

Table 2Cytotoxicity of compounds against four human cancer cell lines.^a

Compounds	HL-60	OVCAR-8	HCT-116	SF-295
1	6.5 (5.2–8.1)	10.3 (5.4–19.9)	10.1 (4.5–23.0)	7.7 (4.2–14.1)
2	7.3 (6.7–7.9)	15.3 (11.9–19.6)	11.4 (9.6–13.5)	12.9 (10.8–15.4)
3	>25	>25	>25	>25
4	8.2 (7.4–9.1)	10.8 (9.4–12.4)	8.6 (7.6–9.9)	6.9 (5.6–8.4)
5	0.6 (0.4–0.7)	2.4 (1.9–2.9)	1.01 (0.74–1.37)	1.3 (1.0–1.6)
Doxorubicin ^b	0.02 (0.01–0.02)	0.3 (0.2–0.3)	0.1 (0.1–0.2)	0.2 (0.2–0.3)

^a Results are expressed as IC₅₀ values in μM and means $\pm$ standard deviation of three independent replicates.^b Positive control.

unsaturated carbon at δ_{C} 134.4 (C-6). The shielding effects of the ¹³C NMR signals observed for the carbons δ_{C} 29.6 (C-1), 75.2 (C-3) and 45.4 (C-9) as compared with those to compound **1**, suggested that these carbons were subjected to a γ -gauche shielding effect of the C-5 hydroxyl group, that consequently was axially oriented. In addition, the methyl group equatorially-oriented was determined by the diagnostic assignments observed for the carbons at C-23, C-24, C-25 and C-26 in comparison with those reported by Agrawal et al. [19].

As observed in **1**, the glucopyranosyl moiety presented a β -anomeric configuration based on the large vicinal coupling constant value of 7.8 Hz of the anomeric proton at δ_{H} 4.50 (H-1'), while the three rhamnopyranosyl units have an α -anomeric configuration by the value of $J = 1.5$ Hz, for the hydrogens at δ_{H} 4.83 (H-1'''), 5.19 (H-1''') and 5.20 (H-1''). The absolute configuration of the glucose unit was also determined as D and that of rhamnose as L, through the acid hydrolysis and the HPLC analysis. In addition, the diagnostic signals at m/z 1013.5504, 867.4950, 721.4130, 575.3622 and 413.3011 in the HRESIMS, showed the subsequent loss of H₂O and the four sugar moieties and confirmed the proposed structure.

Thus, the structure of **2** was elucidated as the (25R)-spirost-6-ene-3 β ,5 β -diol 3-O- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)-[(1 $\rightarrow$ 2)- α -L-rhamnopyranosyl]- β -D-glucopyranoside.

The known compounds were identified as (25R)-spirost-5-ene-3 β -ol 3-O- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- β -D-glucopyranoside (**3**) [12], (25R)-7-oxo-espirost-5-ene-3 β -ol 3-O- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)-[(1 $\rightarrow$ 2)- α -L-rhamnopyranosyl]- β -D-glucopyranoside (Chonglouoside SL-5) (**4**) [13] and (25R)-espirost-5-ene-3 β -ol 3-O- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)- α -L-rhamnopyranosyl-(1 $\rightarrow$ 4)-[(1 $\rightarrow$ 2)- α -L-rhamnopyranosyl]- β -D-glucopyranoside (**5**) [14] after extensive NMR spectral data analysis and comparison of its spectroscopic data with those reported in the literature.

The isolation of steroidal saponins is already reported for several species of different genera of Solanaceae as spirostane, furostane and spirofurostane sapogenins skeletons [21–24]. Although a large number of compounds belonging to the Δ^5 -spirostane series have been reported, the structure of Δ^6 -spirostane skeleton as observed for compound **2**, is a feature never reported for this class of compounds. In addition, the uncommon hydroxylation at C-26 on the six membered pyran ring observed for **1** is noteworthy.

The structure–cytotoxicity relationship of steroidal saponins has been discussed and it has been established that aglycone played an important role in biological activity, while the type, length, linkage as well as the substituents of the glycosyl chain influenced the cell recognition and regulation of biological activity [25]. A literature survey has revealed that the cytotoxic activity of formosanin C (**5**) has been well documented and it has recently emerged as potential antitumor agent [26–28].

Compounds **1–5** were evaluated against four human cancer cell lines colorectal adenocarcinoma (HCT-116), ovarian carcinoma (OVCAR-8), human promyelocytic leukemia (HL-60) and glioma (SF-295) (Table 2). As expected, our results showed compound **5** as the most active showing effective action against all tested cell lines with IC₅₀ ranging to 0.6 to 2.4 $\mu\text{g mL}^{-1}$. Compound **1** exhibited a moderate effect against HL-60 and SF-295, compound **2** exhibited a selective activity against HL-60 with a IC₅₀ value of 7.3 $\mu\text{g mL}^{-1}$, compound **4** showed moderated activity to all cell lines, while compound **3** was inactive. The antimicrobial activities of the isolated compounds against *C. parapsilosis* (ATCC® 22019™), *C. albicans* (ATCC® 10231™), *C. krusei* (ATCC® 14243™), *P. aeruginosa* (ATCC® 9027™), *S. aureus* (ATCC® 6538™) and *B. subtilis* (ATCC® 6633™) were evaluated. The results showed that only formosanin C (**5**) presented antifungal activity against *C. parapsilosis* and *C. albicans* with values of IC₅₀ values of 3.2 and 1.9 $\mu\text{g mL}^{-1}$, respectively, while compounds **1–4** were inactive.

These findings can suggest that the (1 $\rightarrow$ 2)- α -L-rhamnopyranosyl linkage of the inner glucosyl moiety can be important for cytotoxic activity, since the compound **3** was inactive in all cancer cell lines and microorganisms. The modifications on the diosgenyl aglycone structure by the introduction of a hydroxyl group at C-26 on compound **1**, one carbonyl group at C-6 on **4**, or changes of the double bond in the Δ^6 -spirostane skeleton showed only weak cytotoxic results, and no antimicrobial activities were observed.

4. Conclusion

The isolation of spirastone glycosides in *C. laevigatum* is in accordance with the previous reports about the *Cestrum* genus and the Solanaceae family. However, the occurrence of a Δ^6 -spirostane skeleton is an unprecedented feature in the literature. The cytotoxic activity against HCT-116, OVCAR-8, HL-60 and SF-295 cancer cell lines showed formosanin C (**5**) as the most active against all cell lines, while compounds **1** and **2** showed moderated activity against HL-60. Formosanin C (**5**) also showed selective antimicrobial effects against *C. parapsilosis* and *C. albicans* while compounds **1–4** were inactive. The current results suggest that the *Cestrum* genus is an important source of structurally interesting and biologically active steroidal saponins.

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Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at <http://dx.doi.org/10.1016/j.steroids.2015.12.006>.

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