

Molluscicidal Activity of Compounds Isolated from *Euphorbia conspicua* N. E. Br.

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O latex de *Euphorbia conspicua* foi fracionado nas frações triterpênic e irritantes I e II. Da fração triterpênic foram isolados 15 compostos já conhecidos e um novo triterpeno denominado 3β-(*E*)-cinamoileuforbol. A fração irritante II forneceu o 20-*O*-acetil-3-*O*-angeloil-ingenol. A atividade moluscicida dos compostos eufol, 3β-acetoxieufa-8,24-dieno, 3β-(*E*)-cinamoileuforbol e 20-*O*-acetil-3-*O*-angeloil-ingenol foi avaliada. O 20-*O*-acetil-3-*O*-angeloil-ingenol apresentou uma LC₁₀₀ de 1 μg mL⁻¹, a qual foi equivalente ao moluscicida padrão niclosamida. Os compostos eufol, 3β-acetoxieufa-8,24-dieno e 3β-(*E*)-cinamoileuforbol apresentaram uma fraca atividade moluscicida. O 3β-(*E*)-cinamoileuforbol foi submetido a testes de mutagenicidade (teste de Ames com TA 98, 100 e 102) na presença e ausência de ativação metabólica (mistura S9). Foram também realizados os ensaios de citotoxicidade (teste MTT) e genotoxicidade (teste dos micronúcleos, CBMN) com e sem mistura S9, em células V79 de Hamster chinês. O 3β-(*E*)-cinamoileuforbol revelou-se fracamente citotóxico e sem atividade mutagênica ou genotóxica.

Euphorbia conspicua latex was fractionated into triterpenic and irritant fractions I and II. The triterpenic fraction afforded 15 known compounds and a new triterpene, 3β-(*E*)-cinnamoyleuphorbol. 20-*O*-Acetyl-3-*O*-angeloylingenol was isolated from irritant fraction II. The compounds euphol, 3β-acetoxyeupha-8,24-diene, 3β-(*E*)-cinnamoyleuphorbol and 20-*O*-Acetyl-3-*O*-angeloylingenol were evaluated for molluscicidal activity. 20-*O*-Acetyl-3-*O*-angeloylingenol presented LC₁₀₀ value of 1 μg mL⁻¹, equivalent to that of the standard molluscicide niclosamide. Compounds euphol, 3β-acetoxyeupha-8,24-diene and 3β-(*E*)-cinnamoyleuphorbol showed low molluscicidal activity. Mutagenic assays (Ames test with strains TA 98, 100 and 102) were performed with 3β-(*E*)-cinnamoyleuphorbol in the presence and absence of metabolic activation (S9 mix). In V79 cells, the cytotoxicity of 3β-(*E*)-cinnamoyleuphorbol was evaluated using the MTT assay and the genotoxicity was assessed using the cytokinesis-block micronucleus assay (CBMN) with or without S9 mix. Mutagenic or genotoxic activity was not detected, and no significant cytotoxicity was observed for 3β-(*E*)-cinnamoyleuphorbol at lower doses.

Keywords: *Euphorbia conspicua*, molluscicidal activity, mutagenic activity, cytotoxic activity, genotoxicity, 3β-(*E*)-cinnamoyleuphorbol, 20-*O*-acetyl-3-*O*-angeloylingenol

Introduction

Euphorbia conspicua N. E. Br.¹ (Euphorbiaceae) is a succulent tree endemic to Angola and is traditionally used as a treatment for dermatitis and leprosy wounds.² The genus *Euphorbia* is the largest in the spurge family, with more than 1000 species divided into many subgenera and

sections. Many studies have investigated the use of several of these plants in folk medicine to treat cancerous conditions.³⁻⁵ Some *Euphorbia* species have been studied for molluscicidal properties against schistosomiasis or bilharzia, leading to the discovery of milliamines, the most potent molluscicides identified so far.⁶

Schistosomiasis is a major source of morbidity and mortality in developing countries. Three schistosome species infect humans: *Schistosoma mansoni*, *S. haematobium* and

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S. japonicum. Because freshwater snails are intermediate hosts for these parasites,⁷ the use of molluscicides is desirable in the integrated control of schistosomiasis.⁷ Currently, only niclosamide is widely used in control programs, and it is highly active at all stages of the snail life cycle and on schistosome larvae. Natural molluscicidal compounds isolated from a large number of plants have received much attention in hope that they might provide cheap, biodegradable and effective control agents in rural areas where schistosomiasis is endemic.^{8,9}

As part of our ongoing study of the plants of Angola,^{2,10,11} we observed that *Euphorbia conspicua* latex exhibits strong molluscicidal activity² and may be a potential source of bioactive compounds. This work intended to isolate and describe the constituents of the active fractions and to evaluate their effects. As a result, sixteen compounds were characterized, including a new natural product named 3 β -(*E*)-cinnamoyl euphorbol (**3**). Compounds **1-3** and **16** were evaluated for their molluscicidal activity. The cytotoxicity, mutagenic and genotoxicity activities of compound **3** were evaluated using the MTT test, Ames test and cytokinesis-block micronucleus assay (CBMN), respectively.

Results and Discussion

Compounds euphol (**1**),¹² 3 β -acetoxyeupha-8,24-diene (**2**),¹³ cembrene-A (**4**),¹⁴ germacrene-B (**5**),¹⁵ 3 β -acetoxyeuphorbol (**6**),¹⁶ 3 β -acetoxycycloart-24-ene (**7**),¹⁷ β -amyrin (**8**),^{18,19} 3 β -(*E*)-cinnamoyl- β -amyrin (**9**),¹⁹ 3 β -(*E*)-cinnamoyloxyeupha-8,24-diene (**10**),¹² cycloart-24-en-3 β -ol (**11**),¹⁷ 24-methylenecycloartan-3 β -ol (**12**),¹² boeticol (**13**),²⁰ 3 β -acetoxylphenol (**14**),²¹ cholesterol (**15**)²² and 20-*O*-acetyl-3-*O*-angeloylingenol (**16**)²³ (Figure 1) were identified by comparison of experimental ¹H and ¹³C NMR results with spectral data in the literature.

Compound **3** was obtained as a white amorphous solid, and its molecular formula C₄₀H₅₈O₂ was established by HREIMS, showing a molecular ion peak *m/z* of 570.4417 [M⁺] (calc. 570.4436) and 12 degrees of unsaturation. The IR spectrum revealed absorption bands for an ester (1720 and 1153 cm⁻¹), a terminal methylene group (890 cm⁻¹), a fully substituted double bond (1640 cm⁻¹) and an aromatic ring (1580, 850, 820 and 679 cm⁻¹). The ¹H NMR spectrum (Table 1) exhibited a signal for a terminal methylene group [δ_{H} 4.66 and 4.72 (s, 1H, each)], a secondary (*E*)-cinnamoyloxy group [δ_{H} 6.44 and 7.67 (d, 1H, *J* 16.8 Hz, each)] and eight methyl groups (five singlets [δ_{H} 0.77, 0.88, 0.93, 0.96 and 1.01, (3H each)] as well as three doublets [δ_{H} 1.02 and 1.03 (3H, *J* 6.6 Hz each) and 0.94 (3H, *J* 6.0 Hz)]. The methyl doublets δ_{H} 1.02 and 1.03 (*J* 6.6 Hz) and the methine septet δ_{H} 2.23 (*J* 6.6 Hz)

suggested the existence of an isopropyl group in the molecule; moreover, the multiplicity of the methine as a septet implied that the isopropyl group was bonded directly to a quaternary carbon. The ¹³C NMR spectrum (Table 1) showed signals of two sp² carbons of a tetrasubstituted double bond (δ_{C} 133.6 and 134.0) along with signals of a terminal methylene group (δ_{C} 156.9 and 106.0) and a cinnamate moiety (δ_{C} 166.9, 144.3 and 118.9).

Table 1. ¹H NMR and ¹³C NMR data and HMBC correlations of compound **3** in CDCl₃^a

Position	δ_{C}	δ_{H}	HMBC
1	35.0	1.35 m H- α ; 1.78 m H- β	2, 5, 10, 19
2	24.3	1.70 m H- α ; 1.62 m H- β	1, 3, 4, 10
3	81.0	4.66 dd (11.4, 5.4) H- α	2, 4, 28, 29, 1'
4	38.1		
5	51.1	1.27 br d (12.0) H- α	1, 3, 4, 6, 7, 9, 10, 19, 28, 29
6	18.7	1.71 m H- α ; 1.44 m H- β	4, 5, 7, 8, 10
7	27.5	1.95 m H- α ; 2.09 m H- β	5, 8, 9
8	133.6		
9	134.0		
10	37.2		
11	21.4	2.08 m H- α ; 1.94 m H- β	8, 9, 13
12	30.8	1.69 m	9, 11, 13, 17, 18
13	44.1		
14	50.0		
15	29.8	1.55 m H- α ; 1.21 m H- β	13, 16, 17, 30
16	28.0	1.35 m H- α ; 1.94 m H- β	13, 20, 22
17	50.0	1.51 m H- β	13, 14, 16, 18, 20, 21
18	15.4	0.77 s H- α	12, 13, 14, 17
19	20.2	1.01 s H- β	1, 5, 9, 10
20	36.5	1.44 m H- α	22, 23
21	18.8	0.94 d (6.0) H- β	17, 20, 22, 23
22	35.1	1.15 m; 1.56 m	17, 21, 23
23	31.3	1.88 m; 2.12 m	22, 24, 25, 24'
24	156.9		
25	33.8	2.23 sept (6.6) ^b	23, 24, 26, 27, 24'
26	22.0	1.02 d (6.6)	24, 25, 27
27	21.8	1.03 d (6.6)	24, 25, 26
28	28.0	0.93 s H- α	3, 4, 5, 29
29	16.8	0.96 s H- β	3, 4, 5, 28
30	24.3	0.88 s H- β	8, 13, 14, 15
24'	106.0	4.66 s; 4.72 s	23, 25
1'	166.9		
2'	118.9	6.44 d (16.8)	1', 1''
3'	144.3	7.67 d (16.8)	1', 1'', 2'', 6''
1''	134.6		
2'', 6''	128.0	7.52 m	2'', 3'', 4'', 5'', 6''
3'', 5''	128.8	7.37 m	1'', 2'', 3'', 5'', 6''
4''	130.1	7.37 m	2'', 3'', 5'', 6''

^aSpectra were recorded at 600 MHz for ¹H NMR and 150.9 MHz for ¹³C NMR; 2D NMR experiments recorded in accordance; coupling constants (*J* Hz) are in parenthesis; ^bpartially overlapped.

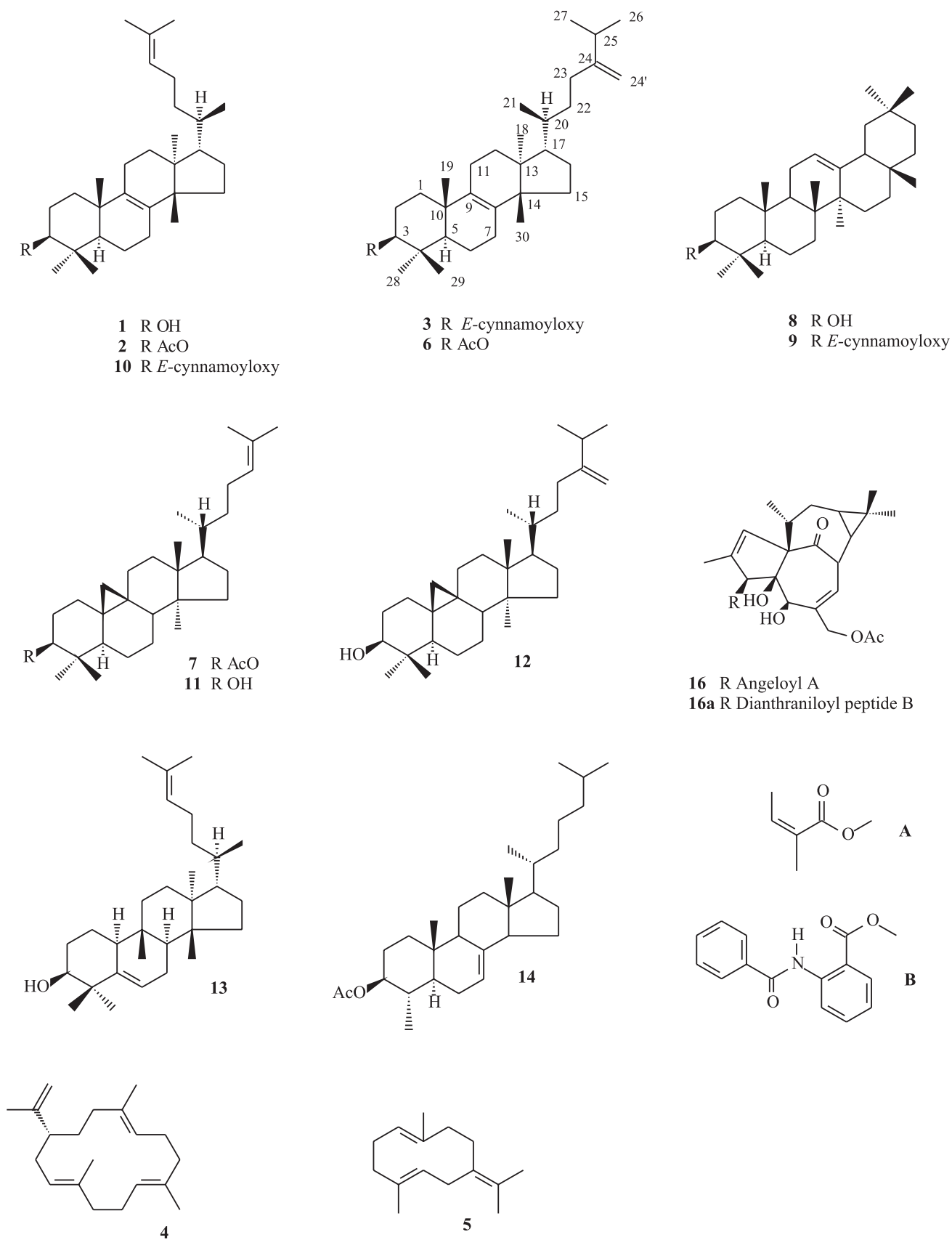


Figure 1. Chemical structures of the isolated compounds.

The EI mass spectrum revealed fragment ion peaks at m/z 555 ($[M]^+ - \text{Me}$), 407 ($[M]^+ - \text{Me} - \text{HO}_2\text{CCH}=\text{CHC}_6\text{H}_5$), 297 (loss of C_9H_{17} and $\text{HO}_2\text{CCH}=\text{CHC}_6\text{H}_5$), 255 ($297 - \text{C}_3\text{H}_6$) and 241 ($255 - \text{CH}_2$), which corroborated the structure as a Δ^{8-9} tetracyclic triterpene bearing a cinnamoyl group at the C-3 position and a side chain containing nine carbon atoms, including isopropyl and C-24 methylene groups.

Analysis of ^1H and ^{13}C NMR spectra along with data from COSY (homonuclear correlation spectroscopy), HSQC (heteronuclear single-quantum correlation) and HMBC (heteronuclear multiple bond coherence) experiments (Table 1) suggested a euphane- or tirucallane-type triterpene. The NOESY (nuclear Overhauser enhancement spectroscopy) data (Figure 2) revealed ^1H - ^1H long-range correlations between Me-29 and Me-19,

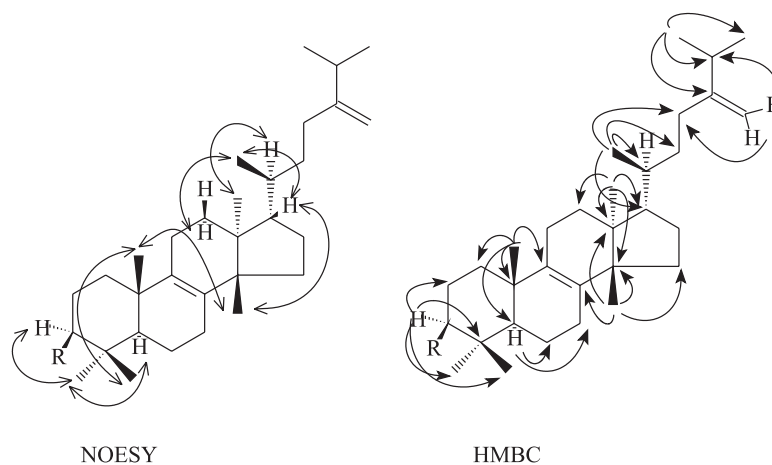


Figure 2. Relevant NOESY and HMBC correlations of **3**.

Table 2. Molluscicidal activity of terpenes from *Euphorbia conspicua* N. E. Br. latex on *Biomphalaria glabrata* Say

Compound	Snail diameter / mm	Concentration / ($\mu\text{g mL}^{-1}$)	Dead snails / %		
			After 24 h of exposure	After 24 h of recovering	After 48 h of recovering
1	13-20	100	0	20	20
	13-20	50	0	20	20
	13-20	20	0	0	0
	15-21	10	0	0	0
	15-21	1	0	0	0
2	15-20	100	0	40	40
	13-20	50	0	40	40
	13-20	20	0	0	0
	15-21	10	0	0	0
	15-21	1	0	0	0
3	15-20	100	0	20	20
	13-20	50	0	0	0
	13-20	20	0	0	0
	15-21	10	0	0	0
	15-21	1	0	0	0
16	15-17	100	100	–	–
	15-20	10	0	100	–
	15-20	1	0	100	–
	13-21	0.1	0	60	60
	13-21	0.01	0	0	20
	13-21	0.001	0	0	20
	Dechlorinated water	15-20		0	0
Dechlorinated water with 1% DMSO	15-21		0	0	0
Cupric sulfate	15-20	50	100	–	–

Temperature: 25 ± 1 °C.

Table 3. Effect of compound **3** on cell viability of V79 Chinese hamster cells using the MTT assay

Dose / ($\mu\text{g per well}$)	Viability / % ^a
25	115.4 \pm 5.1
50	123.0 \pm 8.1
250	59.7 \pm 37.1

^aViability is expressed as percentage values relative to control cells; results are expressed as mean value % viability \pm standard deviations (SD) (n = 3); in each independent experiment four replicate cultures were used.

Me-30, H-17 and Me-21 on the β face of the structure, H-3 and H-5 with Me-28, H-20 with Me-18 and H-12 α with Me-21. These data were consistent with a tirucallane-type structure.^{16,20,23} Thus, compound **3** was identified as a new natural product and named 3 β -(E)-cinnamoyl euphorbol.

Compounds **1-3** and **16** were evaluated for molluscicidal activity against *Biomphalaria glabrata*, a vector of *S. mansoni* (Table 2). The molluscicidal activity of

compound **16**, with an LC₁₀₀ of 1 $\mu\text{g mL}^{-1}$, was equivalent to that of niclosamide, the synthetic compound used for the control of mollusks (LC₁₀₀ 1.5 $\mu\text{g mL}^{-1}$).²⁴ Compound **16** presented a dose dependent and continuous effect on adult snails after 24 h of exposure, while it was completely inactive against the egg masses. In contrast, the triterpenic compounds displayed weak activity. Compounds **1** and **3** caused 20% mortality, while compound **2** cause 40% of mortality at a concentration of 100 $\mu\text{g mL}^{-1}$. Due to a shortage of sample materials, the other triterpenic compounds were not tested.

The survival values in V79 cells are given in Table 3, highlighting that only a slight decrease of survival was observed for 250 $\mu\text{g per well}$. Ames assay outcomes (Table 4) revealed compound **3** as a non-mutagenic agent on the strains tested at doses up to 250 $\mu\text{g per plate}$. The results obtained for the induction of micronuclei in Chinese hamster cells (V79 cells) at concentrations up to 100 $\mu\text{g mL}^{-1}$ revealed no significant increase compared to the negative control in the absence or presence of S9 mix (Table 5).

Table 4. Mutagenic activity of compound **3** in the Ames assay, revertants in three strains of *Salmonella typhimurium* (TA 98, TA100 and TA102) treated with different concentrations of compound **3** in the presence and absence of metabolic activation (S9)

Dose / ($\mu\text{g per plate}$)	Revertants per plate					
	TA 98		TA 100		TA 102	
	-S9	+S9	-S9	+S9	-S9	+S9
0	17.5 \pm 3.5	27.5 \pm 9.2	130.5 \pm 29.0	123.5 \pm 20.5	268.0 \pm 56.6	342.0 \pm 31.1
5	15.0 \pm 1.4	18.5 \pm 3.5	127.5 \pm 33.2	128.5 \pm 23.3	259.5 \pm 36.1	296.5 \pm 30.4
25	19.5 \pm 0.7	18.0 \pm 7.1	129.0 \pm 14.1	124.0 \pm 14.1	231.0 \pm 0	309.5 \pm 51.6
50	16.5 \pm 0.7	28.0 \pm 12.7	116.5 \pm 14.8	107.5 \pm 26.2	288.5 \pm 55.9	312.5 \pm 88.4
250	14.5 \pm 2.1	20.5 \pm 13.4	92.5 \pm 4.2	87.5 \pm 13.4	257.5 \pm 96.8	260.5 \pm 132.2
Quercetin						
10	284.0 \pm 77.7	1314.5 \pm 102.5				
4-NQO ^a						
10			1432		2842	

Values are presented as the mean \pm standard deviation (SD) (n = 2); dose 0 as negative control; quercetin and 4-NQO as positive controls; ^a4-NQO: 4-nitroquinoline-1-oxide.

Table 5. Effect of compound **3** on the frequency of micronucleated binucleated cells (% MNBN) in V79 Chinese hamster cells in the presence (+S9) and absence (-S9) of metabolic activation

Test compound	% MNBN		% BN	
	-S9	+S9	-S9	+S9
Compound 3 / ($\mu\text{g mL}^{-1}$)				
0	0.3 \pm 0.1	0.2 \pm 0.2	41.0 \pm 14.2	37.4 \pm 2.3
20	0.4 \pm 0.2	0.2 \pm 0.0	32.5 \pm 5.1	36.1 \pm 1.0
100	0.3 \pm 0.0	0.1 \pm 0.1	44.2 \pm 1.6	35.2 \pm 1.8
Mytomicin C				
2.5 $\mu\text{g mL}^{-1}$	10.5 \pm 1.8	-	25.0 \pm 4.2	-
Cyclophosphamide				
2.0 $\mu\text{g mL}^{-1}$	-	2.1 \pm 0.3	-	38.6 \pm 0.1

Results are expressed as mean values \pm standard deviations (SD) (n = 2); in each experiment 1000 binucleated cells were analyzed for the presence of micronuclei; % of binucleated cells (% BN) was used as index of cell proliferation; mytomicin C and cyclophosphamide as positive controls, dose 0 as negative control.

Conclusions

E. conspicua, a succulent tree endemic to Angola, was previously evaluated for its molluscicidal activity,² but the latex chemical composition was not determined. The present study revealed that *E. conspicua* latex is composed mainly of triterpenes with euphane and cycloartane skeletons along with other triterpenes and ingenane diterpenes and contains a new compound with a tirucallane skeleton, 3 β -(*E*)-cinnamoyl euphorbol (**3**). Four of the isolated compounds, **1-3** and **16**, were evaluated for their molluscicidal activity. Compound **16** was found to be the most active, with an LC₁₀₀ value of 1 $\mu\text{g mL}^{-1}$, which is similar to that for milliamine L, the most powerful molluscicide of plant origin characterized thus far.²⁵ Although compound **16** and milliamine L are structurally differing from each other at the position C-3, where milliamine L bears a dianthraniloyl peptide group and compound **16** an angeloyl group, molluscicidal activity remains. At this point compound **16** seems to be the major responsible for *E. conspicua* molluscicidal activity but further investigation is needed. Compound **16** was found to be totally inactive against the egg masses of *B. glabrata*. Compound **3** was evaluated for its cytotoxicity, mutagenic activity and genotoxicity using the MTT test, the Ames test and the cytokinesis-block micronucleus assay (CBMN), respectively. No mutagenic or genotoxic activity was detected and little or no cytotoxicity was found concluding that this particular compound as no potential risk regarding their future use as bioactive compound.

The irritant properties of the latex can explain the ethnopharmacological use against leprosy wounds and dermatitis in general, but more studies are needed to validate this idea.

Biological and chemical evaluation of other components of *E. conspicua* latex will require large amounts of latex and will be the subject of a future study.

Experimental

General experimental procedures

Optical rotations were obtained with a Bellingham+Stanley Ltd ADP 220 polarimeter. HREIMS measurements were conducted on a VG Autospec M and recorded at 70 eV. The mass spectrum of **3** was obtained from a GC-MS (Hewlett-Packard 5989 A) spectrometer. The IR spectra were recorded in a Unicam Mattson 5000 FTIR. NMR spectra of **3** were recorded in a Bruker Avance II 600 MHz (¹H NMR) and 150.9 MHz (¹³C NMR) spectrometer in CDCl₃. Spectra of compounds **1-2** and **4-16** were

recorded in a Bruker AC 250P 250 MHz (¹H NMR) and 62.9 MHz (¹³C NMR) spectrometer in CDCl₃. Chemical shifts are given in δ ppm and are referenced to residual CHCl₃, 7.26 ppm for the ¹H and 77.0 ppm for ¹³C. Two-dimensional experiments were performed with standard Bruker software. Column chromatography was conducted on silica gel 60 (70-230 mesh, Merck, Darmstadt, Germany).

Plant material

Plant material was collected at Cacuaco, Luanda and was identified by Professor Esperança da Costa from Agostinho Neto University (Biology Department). A voucher specimen (No 4498) has been deposited at the Luanda Herbarium, Luanda, Angola.

Extraction and isolation

The latex (900 g) was collected and frozen ($-20\text{ }^{\circ}\text{C}$). Defrosted latex was dissolved in MeOH at $50\text{ }^{\circ}\text{C}$, and then maintained at room temperature ($22-23\text{ }^{\circ}\text{C}$) until a solid formed; the solid material was separated by filtration and exhaustively extracted with Me₂CO at $40\text{ }^{\circ}\text{C}$.² The Me₂CO fraction was evaporated and redissolved in MeOH/H₂O (1:1) and subsequently extracted with hexane. The hexane fraction (30 g) was analyzed by ¹H and ¹³C NMR and identified as the triterpenic fraction.¹⁷ The metanolic filtrate, obtained from the defrosted latex, was evaporated to dryness and extracted with hexane to yield fraction I (1 g); the insoluble portion was dissolved in MeOH-H₂O (1:1) and extracted with Et₂O (fraction II, 2 g).²⁶ Due to their irritant properties to the eyes and skin of the operator, fractions I and II were named irritant fractions.²⁶ The triterpenic fraction (30 g) was submitted to column chromatography (CC) over silica gel 60 (Merck) using a gradient elution from 100% hexane, hexane/toluene (9:1, 8:2, 7:3, 1:1), toluene/EtOAc (95:5, 9:1, 7:3) to 100% EtOAc. A total of 140 fractions of ca. 100 mL each were collected and pooled into ten fractions. Fraction 1 afforded cembrene-A (**4**) (6 mg). Fraction 2 yielded germacrene-B (**5**) (10 mg). Column chromatography of fraction 5 on 10% AgNO₃/silica gel 60 (Merck) (m/m) using a hexane/EtOAc gradient elution, from 100% hexane to 100% EtOAc, afforded 3 β -acetoxy eupha-8,24-diene (**2**) (35 mg), 3 β -acetoxy euphorbol (**6**) (5 mg), 3 β -acetoxy cycloart-24-ene (**7**) (12 mg) and β -amyrin (**8**) (11 mg). Column chromatography of fraction 6 on silica gel 60 (Merck) using hexane, hexane/toluene (95:5, 9:1, 8:2, 7:3, 1:1) and toluene/EtOAc (99:1, 95:5, 9:1, 8:2, 7:3, 6:4, 1:1) yielded euphol (**1**) (65 mg), 3 β -(*E*)-cinnamoyl euphorbol (**3**) (60 mg), 3 β -(*E*)-cinnamoyl- β -amyrin (**9**) (18 mg) and 3 β -(*E*)-cinnamoyloxy eupha-8,24-diene (**10**)

(15 mg). Column chromatography of fraction 7 on 10% AgNO₃/silica gel 60 (Merck) (m/m) using hexane/EtOAc (95:5, 9:1, 8:2, 7:3, 6:4, 1:1) afforded cycloart-24-en-3 β -ol (**11**) (11 mg) and 24-methylenecycloartan-3 β -ol (**12**) (14 mg). Column chromatography of fraction 8 on silica gel 60 (Merck) using a hexane/EtOAc gradient yielded boeticol (**13**) (7 mg), 3 β -acetoxylofenol (**14**) (10 mg) and cholesterol (**15**) (33 mg). Irritant fraction II (2 g) was submitted to column chromatography (CC) over silica gel 60 (Merck) using a gradient elution from 100% hexane to hexane/EtOAc (95:5, 9:1, 8:2 7:3, 1:1). A total of 76 fractions of ca. 50 mL each were collected and pooled into six fractions. Column chromatography of fraction 5 on 10% AgNO₃/silica gel 60 (Merck) (m/m) with hexane/diethyl ether (95:5, 9:1, 8:2, 7:3) yielded 20-*O*-acetyl-3-*O*-angeloylingenol (**16**) (22 mg).

3 β -(*E*)-Cinnamoyleuphorbol (**3**)

White amorphous solid; $[\alpha]_D^{20} - 19.0^\circ$ (*c* 1.33, CHCl₃); IR (KBr) ν_{\max} /cm⁻¹: 3040, 3015, 3021, 1720, 1640, 1580, 1153, 890, 850, 820, 679; ¹H NMR (CDCl₃, 600 MHz) and ¹³C NMR (CDCl₃, 150.9 MHz): see Table 1; HREIMS *m/z* 570.4417 [M]⁺ (calcd for C₄₀H₅₈O₂, 570.4436).

Molluscicide activity

The snail colony (*B. glabrata*) was sustained as described.^{27,28} Molluscicidal activity for adults and egg masses of *B. glabrata* was evaluated according to established procedures.²⁷⁻³⁰

MTT reduction assay.

MTT assay is a cell viability assay based on conversion of MTT dye by mitochondrial enzymes of viable cells into a formazan which can be spectrophotometrically measured. In this assay the absorbance is proportional to the number of viable cells.³¹

The MTT assay was conducted on V79 Chinese hamster cells. Approximately 10⁴ cells were grown at 37 °C in a 5% CO₂ atmosphere for 24 h in 96-well plates in 200 μ L of Ham's F-10 medium supplemented with 10% newborn calf serum and 1% penicillin/streptomycin solution. Different doses of the compound (25, 50 and 250 μ g *per well*) were added, and the cells were incubated for 3 h. The medium was removed, and cells were incubated for 3 h with MTT (0.5 mg mL⁻¹). The cells were washed carefully with PBS, and then 200 μ L of DMSO was added to each well. The absorbance of the converted dye was measured at 595 nm in a Zenith 3100 microplate reader. Cell viability was assessed by comparing the absorbance values of treated

cells with that of the control. Absorbance values presented by V79 cell cultures without the addition of compound **3**, i.e. control cultures, correspond to 100% of cell viability. Three independent experiments were performed. In each independent experiment four replicate cultures were used.

Ames assay

Mutagenicity testing was conducted by the plate incorporation assay described by Maron and Ames³² using *Salmonella typhimurium* strains TA 98, TA 100 and TA 102 in the presence or absence of S9 mix.³² At least two independent experiments were performed for each assay. Quercetin and 4-nitroquinoline-1-oxide were used as positive controls and dose 0 as negative control.

Cytokinesis-block micronucleus assay (CBMN)

Approximately 5 \times 10⁵ V79 Chinese hamster cells were cultured for 24 h in 25 cm² culture flasks and then exposed to compound **3** at concentrations of 20 and 100 μ g mL⁻¹. Mitomycin C (2.5 μ g mL⁻¹) or cyclophosphamide (2.0 μ g mL⁻¹) was used as a positive control with and without S9 mix and dose 0 as negative control, respectively. 24 h after the genotoxic treatment, the cells were washed with fresh culture medium, and cytochalasin-B (Cyt-B) was added to a final concentration of 4.5 μ g mL⁻¹. The cells were incubated for additional 16 h, harvested by trypsinization, rinsed and submitted to a mild hypotonic treatment as described elsewhere.³³ The centrifuged cells were placed onto dry slides, and smears were made. After air-drying, the slides were fixed with cold methanol for 30 min. One day later, the slides were stained with Giemsa (4% (v/v) in 0.01 mol L⁻¹ sodium phosphate buffer, pH 6.8) for 10 min. For each experimental point, 1000 binucleate V79 cells (BN) with well-preserved cytoplasm were scored. Micronuclei were identified under a light microscope using a magnification of 1250 \times according to the criteria proposed by Caria *et al.*³⁴ We evaluated MN/BN (data not showed), which represents the average number of micronuclei per binucleated cell, and the frequency of micronucleated binucleated V79 cells (% MNBN), which represents the fraction of cytokinesis blocked (binucleated) cells with micronuclei, regardless of the number of micronuclei per BN cell.³⁵ At least two independent experiments were performed for each assay.

Cell proliferation

The decrease in cell proliferation in the experiments described above was assessed by determining the frequency

of binucleate cells (% BN).³⁵ For this index 1000 cells with well-preserved cytoplasm were analyzed according to number of nuclei at a magnification of 500 ×. Two independent experiments were conducted.

Supplementary Information

¹H NMR, ¹³C NMR, ¹H-¹H COSY, HSQC, HMBC and NOESY NMR spectra of compound **3** and physical data of compounds **1**, **2** and **16** are available free of charge at <http://jbc.org.br> as a PDF file.

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References

- Brown, N. E.; *In Fl. Trop. Afr.* **1912**, *6*, 600.
- dos Santos, A. F.; Lopes, L. A.; Mata, R. C. S.; de Mendonça, D. I. M. D.; Sant'Ana, A. E. G.; *Bioresour. Technol.* **2007**, *98*, 135.
- Uemura, B.; Katayama, C.; Luno, E.; Sasaki, K.; Hirata, Y.; *Tetrahedron Lett.* **1975**, 1703.
- Appendino, G.; Jakupovic, S.; Tron, C. G.; Jakupovic, J.; Milon, V.; Ballero, M.; *J. Nat. Prod.* **1998**, *61*, 749.
- Alberto, J. M.; Sanz-Cervera, J. F.; Alberto, Y.; Jakupovic, J.; *J. Nat. Prod.* **1999**, *62*, 110.
- Hecker, E.; *Pure Appl. Chem.* **1977**, *49*, 1423.
- <http://www.who.int/topics/schistosomiasis/en/> accessed in June 2011.
- Hostettman, K.; Marston, A.; Ndjoko, K.; Wolfender, J. L.; *Curr. Org. Chem.* **2000**, *4*, 973.
- Queiroz, E. F.; Ahua, K. M.; Hostettman, K.; *Chimia* **2005**, *59*, 299.
- Borges, C. M. P.; Diakanawma, C.; de Mendonça, D. I. M. D.; *J. Braz. Chem. Soc.* **2010**, *21*, 1121.
- Sebastião, N'S. N.; Cordeiro, I. J. S.; dos Santos, A. F.; Gaspar, J. F.; Martins, C.; Rueff, J.; Diakanamwa, C.; Sant'Ana, A. E. G.; de Mendonça, D. I. M. D.; *Phytochemistry* **2010**, *71*, 798.
- Gewali, M. B.; Hattori, M.; Tezuka, Y.; Kikuchi, T.; Namba, T.; *Phytochemistry* **1990**, *29*, 1625.
- Akihisa, T.; Kimura, Y.; Koike, K.; Shibata, T.; Yoshida, Z.; Nikaido, T.; Tamura, T.; *J. Nat. Prod.* **1998**, *61*, 409.
- Bai, B.; Jai, M.; *Magn. Reson. Chem.* **2008**, *46*, 791.
- Brown, E. D.; Sam, T. W.; Sutherland, J. K.; Torre, A.; *J. Chem. Soc., Perkin Trans. 1* **1975**, 2326; Adio, A. M.; *PhD Thesis*, Institute of Organic Chemistry, University of Hamburg, Germany, 2005; http://deposit.ddb.de/cgi-bin/dokserv?idn=974207071&dok_var=d1&dok_ext=pdf&filename=974207071.pdf accessed in June 2011.
- Akihisa, T.; Wijeratne, E. M. K.; Tokuda, H.; Enjo, F.; Toriumi, M.; Kimura, Y.; Koike, K.; Nikaido, T.; Tezuka, Y.; Nishino, H.; *J. Nat. Prod.* **2002**, *65*, 158.
- Teresa, J. P.; Urones J. G.; Marcos, I. S.; Basabe, P.; Cuadrado, M. J. S.; Moro, R. F.; *Phytochemistry* **1987**, *26*, 1767.
- Chen, H.; Jia, Z.-J.; *Indian J. Chem. B.* **1996**, *35B*, 1308.
- Atta-Ur-Rahman; *Pentacyclic Triterpenoids*, vol. 2; Elsevier Science B.V.: Amsterdam, 1994, pp. 21, 443.
- Ferreira, M. J. U.; Ascenso, J. R.; Tavares, O. S.; *J. Nat. Prod.* **1995**, *58*, 275.
- Bonerji, R.; Misra, G.; Nigam, S. K.; *Phytochemistry* **1987**, *26*, 2644.
- Kovganko, N. V.; Kashkan, Zh. N.; Borisov, E. V.; *Chem. Nat. Compd.* **2000**, *36*, 595.
- Akihisa, T.; Kimura, Y.; Kokke, W. C. M. C.; Takase, S.; Yasukama, K.; Tamura, T.; *J. Chem. Soc., Perkin Trans. 1* **1996**, *1*, 2379.
- Perret, S.; Whitfield, P. J.; *Parasitol. Today* **1996**, *12*, 156.
- Zani, C. L.; Marston, A.; Hamburger, M.; Hostettmann, K.; *Phytochemistry* **1993**, *34*, 89.
- Urones, J. G.; Barcala, P. B.; Cuadrado, M. J. S.; Marcos, I. S.; *Phytochemistry* **1988**, *27*, 207.
- Santos, A. F.; Sant'Ana, A. E. G.; *Phytother. Res.* **1999**, *13*, 660.
- Santos, A. F.; Sant'Ana, A. E. G.; *Phytomedicine* **2000**, *6*, 431.
- Thiilborg, S. T.; Christensen, S. B.; Cornet, C.; Olsen, C. E.; Lemmich, E.; *Phytochemistry* **1994**, *36*, 753.
- World Health Organization (WHO); *Bull. WHO* **1965**, *33*, 567.
- Carmichael, J.; DeGraff, W. G.; Gazdar, A. F.; Minna, J. D.; Mitchell, J. B.; *Cancer Res.* **1987**, *47*, 936.
- Maron, D. M.; Ames, B. N.; *Mutat. Res.* **1983**, *113*, 173.
- Van Hummelen, P.; Kirsch-Volders, M.; *Mutagenesis* **1990**, *5*, 203.
- Caria, H.; Chaveca, T.; Laires, A.; Rueff, J.; *Mutat. Res.* **1995**, *343*, 85.
- Oliveira, N. G.; Castro, M.; Rodrigues, A. S.; Gonçalves, I. C.; Cassapo, R.; Fernandes, A. P.; Chaveca, T.; Toscano-Rico, J. M.; Rueff, J.; *Mutagenesis* **2001**, *16*, 369.

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