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## N-Cinnamoyltetraketide Derivatives from the Leaves of *Toussaintia orientalis*

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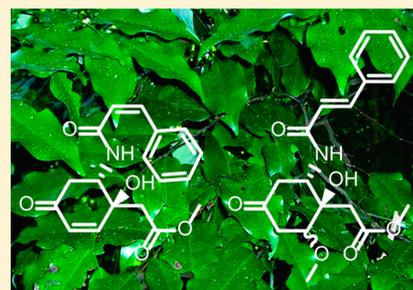
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### Supporting Information

**ABSTRACT:** Seven N-cinnamoyltetraketides (**1**–**7**), including the new Z-toussaintine E (**2**), toussaintine F (**6**), and toussaintine G (**7**), were isolated from the methanol extract of the leaves of *Toussaintia orientalis* using column chromatography and HPLC. The configurations of E-toussaintine E (**1**) and toussaintines A (**3**) and D (**5**) are revised based on single-crystal X-ray diffraction data from racemic crystals. Both the crude methanol extract and the isolated constituents exhibit antimycobacterial activities (MIC 83.3–107.7  $\mu$ M) against the H37Rv strain of *Mycobacterium tuberculosis*. Compounds **1**, **3**, **4**, and **5** are cytotoxic (ED<sub>50</sub> 15.3–105.7  $\mu$ M) against the MDA-MB-231 triple negative aggressive breast cancer cell line.



*Toussaintia orientalis* Verdc. (Annonaceae) is endemic to Tanzania<sup>1</sup> and is one of the four *Toussaintia* species that are found only in tropical Africa.<sup>2</sup> It occurs in the Tanzanian coastal forests and in the Eastern Arc Mountains and is classified as an endangered species.<sup>3</sup> Controversially, *T. orientalis* is used in traditional medicinal. Its roots mixed with lion skin are boiled to produce a concoction that is applied as a remedy for the treatment of asthma and skin rashes arising from abdominal worms, whereas its leaves mixed with chicken feathers are burnt, with the resulting ashes being mixed with coconut oil to treat irritating rashes.<sup>2</sup>

Recent phytochemical investigations of *T. orientalis* serendipitously uncovered myriads of new bioactive metabolites,<sup>1,4,5</sup> among others bioactive N-cinnamoyltetraketides having  $\alpha,\beta$ -unsaturated cyclohexenoid, tetrahydrobenzofuranoid, or indolidinoid skeletons amidated with a cinnamoyl moiety.<sup>1,6</sup> These structural units are commonly found in antituberculous,<sup>7</sup> cytostatic,<sup>8</sup> antimalarial, and cardioprotective compounds.<sup>9</sup> As part of our ongoing work on the phytochemical characterization of East African medicinal plants, we report the isolation, identification, and the cytotoxic and antimycobacterial activities of three new and four known N-cinnamoyltetraketide derivatives isolated from the leaves of *T. orientalis*. Moreover, the configurations<sup>1</sup> of the known toussaintines A, D, and E are corrected based on X-ray crystallographic data.

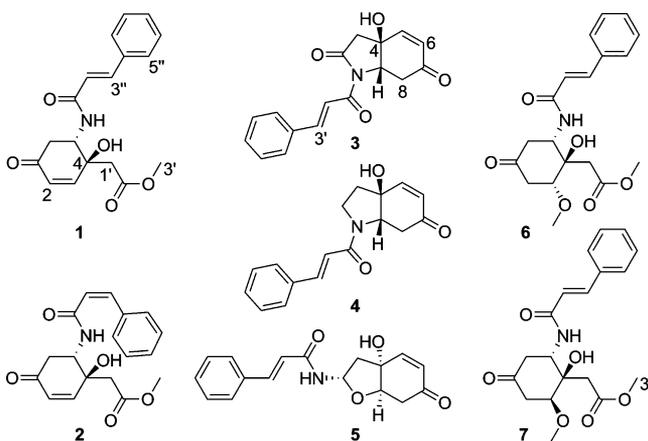
## ■ RESULTS AND DISCUSSION

Vacuum liquid chromatographic (VLC) fractionation of the methanol extract of *T. orientalis* leaves, followed by repeated gravitational column chromatographic separation on silica gel 60 and Sephadex LH-20, afforded E-toussaintine E (**1**),<sup>1</sup> Z-toussaintine E (**2**), and toussaintines A (**3**), C (**4**), D (**5**), F (**6**), and G (**7**). Of these, **2**, **6**, and **7** are new natural products. The compounds were identified using NMR, MS, and single-crystal X-ray studies as well as by comparison of their observed and reported spectroscopic data.

E-Toussaintine E (**1**) and Z-toussaintine E (**2**) were obtained as white solids and showed similar <sup>1</sup>H and <sup>13</sup>C NMR spectra, with the main difference being the chemical shifts and the <sup>3</sup>J<sub>HH</sub> values of the olefinic protons of their cinnamoyl groups (Table 1, Supporting Information). Compound **1** was identified as E-toussaintine E by comparison of its experimental and published spectroscopic data.<sup>1</sup> Previously, it was obtained as a minor component of the leaves of *T. orientalis*,<sup>1</sup> and its relative configuration was proposed based on NMR data. Good-quality single crystals were obtained from a 1:1 MeOH/MeCN solvent mixture. The X-ray crystallographic analysis confirms the previously established main conformation of its cyclohexenoid

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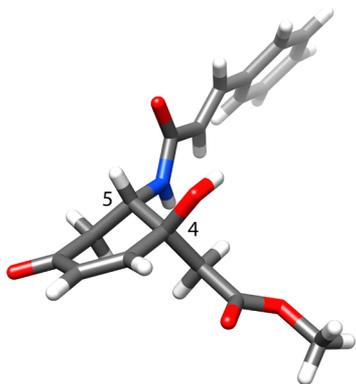
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**Table 1.**  $^1\text{H}$  and  $^{13}\text{C}$  NMR Spectroscopic Data for *Z*-Toussaintine E (2) Acquired in  $\text{CDCl}_3$  at  $25^\circ\text{C}$  ( $\delta_{\text{H}}$ , Multiplicity (*J* in Hz))

position	$\delta_{\text{C}}$ , type	$\delta_{\text{H}}$	<i>m</i> ( <i>J</i> in Hz)
1	195.3, C=O		
2	128.0, CH	5.96	d (10.5)
3	152.4, CH	6.92	d (10.5)
4	72.1, C–O		
5	54.6, CH	4.53	ddd (4.8, 8.0, 11.2)
6	39.7, CH <sub>2</sub>	2.58, 2.18	dd (4.8, 17.6), dd (11.2, 17.6)
1'	40.6, CH <sub>2</sub>	2.62	d (15.1)
		2.49	d (15.1)
2'	171.4, C=O		
3'	52.4, OCH <sub>3</sub>	3.73	s
1''	168.5, C=O		
2''	124.1, CH	6.06	d (12.7)
3''	138.9, CH	7.01	d (12.7)
4''	134.7, C		
5''	128.8, CH	7.39	m
6''	129.0, CH	7.39	m
7''	129.4, CH	7.37–7.36	m
NH		5.60	d (8.0)

core and the relative configuration of C-5 (Figure 1); however, it did not support the previously suggested<sup>1</sup> C-4 relative configuration. On the basis of the X-ray diffraction structure, the correct relative configuration of racemic *E*-toussaintine E (1) is methyl 2-[6 $\alpha$ -(1-*E*-cinnamido-1 $\beta$ -hydroxy-4-oxacyclohex-2-en-1-yl)]acetate.



**Figure 1.** X-ray crystal structure of racemic *E*-toussaintine E (1) with its C-4 and C-5 stereocenters being highlighted.

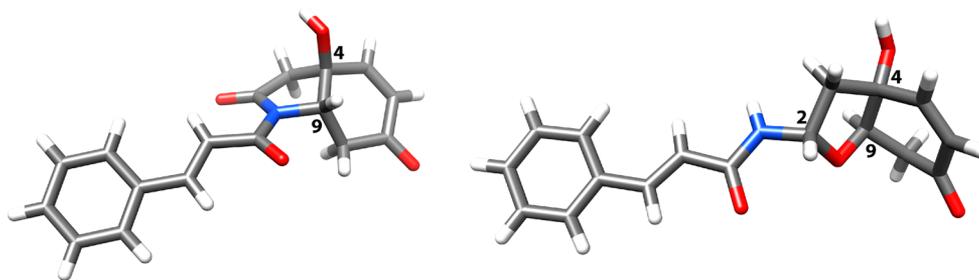
The vicinal coupling constant of the protons of the cinnamoyl double bond indicates a *Z*-configuration for 2 ( $^3J_{\text{HH}} = 12.7$  Hz), but an *E*-configuration for 1 ( $^3J_{\text{HH}} = 15.8$  Hz). The shielding of the olefinic protons of 2 ( $\delta_{\text{H-2}''} 6.06$ ,  $\delta_{\text{H-3}''} 7.01$ ) relative to those of 1 ( $\delta_{\text{H-2}''} 6.44$ ,  $\delta_{\text{H-3}''} 7.71$ ) is likely due to the difference in their relative orientation as compared to the carbonyl moiety that possesses a strong magnetic anisotropy. The chemical shifts of the proximal H-1', H-5, and H-6ab are also influenced by the *E*–*Z* geometrical isomerism, yet to a lesser degree. The 2D NMR and MS data, shown in the Supporting Information, are consistent with the proposed structures of 1 and 2. Thus, on the basis of its spectroscopic data 2 was characterized as the *Z*-isomer of the known 1, i.e., methyl 2-[6 $\alpha$ -(1-*Z*-cinnamido-1 $\beta$ -hydroxy-4-oxacyclohex-2-en-1-yl)]acetate, and was assigned the trivial name *Z*-toussaintine E.

Toussaintines A (3), C (4), and D (5) were isolated as white solids and were identified by comparison of their NMR spectroscopic data (Supporting Information) with reported data.<sup>1</sup> Single crystals were obtained for 3 and 5 by recrystallization from MeOH/MeCN (1:1). On the basis of the X-ray structures (Figure 2), the relative configurations at C-4 of toussaintine A and at C-2 and C-9 for toussaintine D are here unambiguously determined. As for toussaintine E (1), the X-ray diffraction structures of toussaintine A (3) and toussaintine D (5) were based on racemic crystals.

Toussaintine F (6) was obtained as a white solid and was assigned the molecular formula  $\text{C}_{19}\text{H}_{23}\text{NO}_6$  based on HRMS (ESI) data ( $[\text{M} + \text{H}]^+$  *m/z* obs 362.1616, calcd 362.1604). It was identified as a cinnamide derivative upon observation of the *E*-olefinic protons H-3'' ( $\delta_{\text{H}} 7.57$ , *J* = 15.7 Hz) and H-2'' ( $\delta_{\text{H}} 6.27$ , *J* = 15.7 Hz), the H-5''/9'' ortho ( $\delta_{\text{H}} 7.51$ –7.47), H-6''/8'', and H-7'' ( $\delta_{\text{H}} 7.40$ –7.34) aromatic protons, and the amido proton at  $\delta 6.67$  (1H) (Table 2) by  $^1\text{H}$  NMR data. The  $^{13}\text{C}$  NMR signals of C-1'' ( $\delta_{\text{C}} 165.0$ ), CH-3'' ( $\delta_{\text{C}} 141.6$ ), CH-6''/8'' ( $\delta_{\text{C}} 129.0$ ), C-4'' ( $\delta_{\text{C}} 134.8$ ), C-5''/9'' ( $\delta_{\text{C}} 128.0$ ), CH-7'' ( $\delta_{\text{C}} 129.9$ ), and CH-2'' ( $\delta_{\text{C}} 120.7$ ), giving HSQC and HMBC correlations to the above protons, further confirmed this assumption (Table 2, Table S1).

Unlike the recently reported *N*-cinnamoyltetraketide derivatives,<sup>1</sup> the  $^1\text{H}$  NMR spectrum of 6 does not show olefinic signals typical of a cyclohexenoyl unit, but instead aliphatic resonances corresponding to a trisubstituted cyclohexanone ring, as confirmed by  $^1\text{H},^1\text{H}$  COSY (Figure S43). The cinnamide amido proton ( $\delta_{\text{H}} 6.67$ ) couples to methine H-5 ( $\delta_{\text{H}} 4.78$ ), which in turn couples to the diastereotopic methylene H-6a and H-6b ( $\delta_{\text{H}} 3.20$ , 2.40) and to the oxymethine H-3 ( $\delta_{\text{H}} 3.81$ ) through a *W*-coupling, frequently observed for 1,3-diequatorial protons of cyclohexanes. Thus, this  $^4J$  coupling reveals the equatorial orientation of both H-3 and H-5 in the preferred conformation of the cyclohexanone ring of 6 (Figure 3). Vicinal coupling is observed between H-3 ( $\delta_{\text{H}} 3.81$ ) and the diastereotopic methylene H-2a ( $\delta_{\text{H}} 3.08$ ) and H-2b ( $\delta_{\text{H}} 2.69$ ).

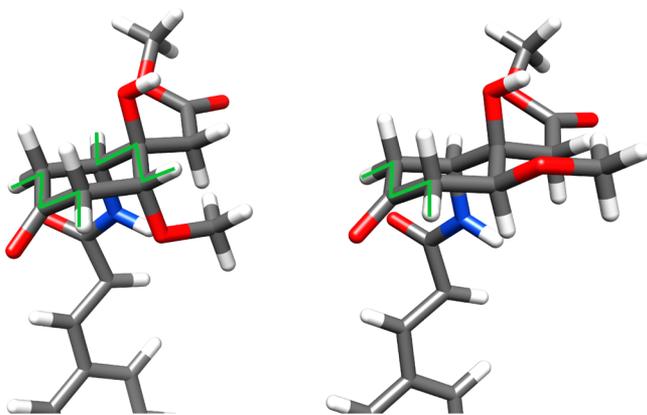
The methylene H-2b ( $\delta_{\text{H}} 2.69$ ) shows a *W*-coupling to H-6b ( $\delta_{\text{H}} 2.40$ ), revealing the equatorial orientation of both protons (Figure 3). The position of OCH<sub>3</sub>-3 ( $\delta_{\text{H}} 3.41$ ) is revealed by its  $^3J_{\text{CH}}$  correlation to C-3 ( $\delta_{\text{C}} 85.9$ ), whereas that of the methoxycarbonyl methyl substituent is indicated by the HMBC correlations of the diastereotopic CH<sub>2</sub>-1' ( $\delta_{\text{H}} 2.88$  and 2.80) to C-4 ( $\delta_{\text{C}} 71.2$ ). The composition of this substituent is indicated by the HMBC cross-peaks of OCH<sub>3</sub>-3' ( $\delta_{\text{H}} 3.75$ ) and of H-1' ( $\delta_{\text{H}} 2.88$  and 2.80) to the ester carbonyl C-2' ( $\delta_{\text{C}}$



**Figure 2.** X-ray crystal structures of racemic toussaintines A (3) and D (5), to the left and to the right, respectively, with their stereocenters being highlighted.

**Table 2.**  $^1\text{H}$  and  $^{13}\text{C}$  NMR Spectroscopic Data for Toussaintines F (6) and G (7) Acquired in  $\text{CDCl}_3$  at 25 °C ( $\delta_{\text{H}}$ , Multiplicity ( $J$  in Hz))

position	6			7		
	$\delta_{\text{C}}$ , type	$\delta_{\text{H}}$	$m$ ( $J$ in Hz)	$\delta_{\text{C}}$ , type	$\delta_{\text{H}}$	$m$ ( $J$ in Hz)
1	207.8, C=O			207.0, C=O		
2	40.3, CH <sub>2</sub>	3.08, 2.69	dd (2.9, 14.7), ddd (2.8, 2.8, 14.7)	42.3, CH <sub>2</sub>	2.85, 2.77	dd (9.7, 14.0), ddd (1.9, 4.8, 14.0)
3	85.9, CH–O	3.81	ddd (2.5, 2.8, 2.9)	80.4, CH–O	3.50	dd (4.8, 9.7)
4	71.2, C–O			74.5, C–O		
5	54.7, CH	4.78	dddd (2.4, 2.5, 5.8, 10.2)	52.2, CH	4.66	ddd (4.3, 5.8, 9.0)
6	43.8, CH <sub>2</sub>	3.20	dd (5.8, 14.6)	43.1, CH <sub>2</sub>	3.24	dd (5.8, 14.8)
		2.40	ddd (2.4, 2.8, 14.6)		2.31	ddd (1.9, 4.3, 14.8)
3-OMe	57.9, OCH <sub>3</sub>	3.41	s	57.5, OCH <sub>3</sub>	3.38	s
1'	37.8, CH <sub>2</sub>	2.87	d (16.7)	38.3, CH <sub>2</sub>	2.79	d (15.5)
		2.80	d (16.7)		2.71	d (15.5)
2'	173.8, C=O			173.4, C=O		
3'	52.4, OCH <sub>3</sub>	3.75	s	52.4, OCH <sub>3</sub>	3.76	s
1''	165.0, C=O			166.1, C=O		
2''	120.7, CH	6.27	d (15.7)	119.7, CH	6.33	d (15.5)
3''	141.6, CH	7.57	d (15.7)	142.7, CH	7.62	d (15.5)
4''	134.8, C			134.5, C		
5''	128.0, CH	7.51–7.47	m	128.1, CH	7.49	7.4, 2.2
6''	129.0, CH	7.40–7.34	m	129.1, CH	7.40–7.36	m
7''	129.9, CH	7.40–7.34	m	130.2, CH	7.40–7.36	m
NH		6.67	d (10.2)		5.71	d (8.9)

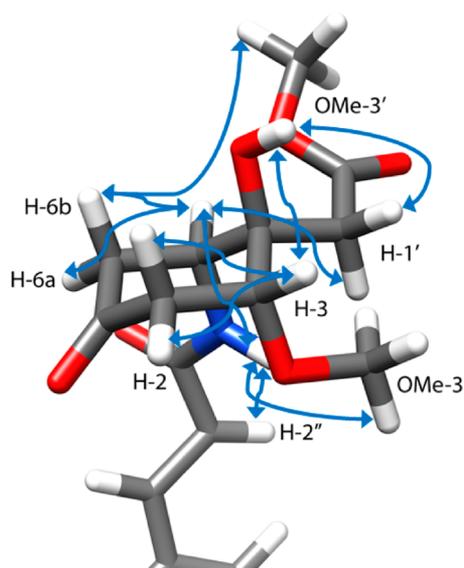


**Figure 3.** W-Couplings ( $^4J$ ) observed for the equatorial protons of toussaintines F (6) and G (7) reveal the preferred conformation of their cyclohexanone cores.

173.8). The substitution pattern of the cyclohexanone core of **6** is further indicated by the  $^3J_{\text{CH}}$  HMBC correlations of OH-4 ( $\delta_{\text{H}}$  4.70) to C-3 ( $\delta_{\text{C}}$  85.9) and C-5 ( $\delta_{\text{C}}$  54.7) and the very weak  $^3J_{\text{CH}}$  to C-4 ( $\delta_{\text{C}}$  71.2). The resonance of C-1 ( $\delta_{\text{C}}$  207.8) is confirmed by its  $^2J_{\text{CH}}$  correlation to H-6a ( $\delta_{\text{H}}$  3.20), for

instance. The *E*-configuration of the C-2''–C-3'' olefinic bond is indicated by the  $^3J_{\text{H-2''},\text{H-3''}}$  value of 15.7 Hz. NOESY correlations (Figure 4) confirm the proposed constitution and relative configuration. Hence, the cinnamide ( $\delta_{\text{H}}$  6.67) shows NOEs to H-6ab ( $\delta_{\text{H}}$  3.20 and 2.40), H-5 ( $\delta_{\text{H}}$  4.78), OCH<sub>3</sub>-3 ( $\delta_{\text{H}}$  3.41), CH<sub>2</sub>-1'ab ( $\delta_{\text{H}}$  2.80 and 2.87), and H-2'' ( $\delta_{\text{H}}$  6.27). In turn, H-5 ( $\delta_{\text{H}}$  4.78) shows further NOEs to CH<sub>2</sub>-6ab ( $\delta_{\text{H}}$  3.20 and 2.40) and CH<sub>2</sub>-1'ab ( $\delta_{\text{H}}$  2.80 and 2.87), whereas the latter protons give NOEs to OH-4 ( $\delta_{\text{H}}$  4.70) and the equatorial H-3 ( $\delta_{\text{H}}$  3.81) as well. Overall, the relative configuration and the main conformation of the cyclohexanone core of **6** are revealed by the  $^3J_{\text{H-6b},\text{H-5}}$  (2.4 Hz),  $^3J_{\text{H-6a},\text{H-5}}$  (5.8 Hz),  $^4J_{\text{H-5},\text{H-3}}$  (2.5 Hz),  $^3J_{\text{H-3},\text{H-2a}}$  (2.9 Hz),  $^3J_{\text{H-3},\text{H-2b}}$  (2.8 Hz), and  $^3J_{\text{H-2b},\text{H-6b}}$  (2.8 Hz) gauche couplings and are corroborated by the corresponding set of NOEs shown in Figure 4. No crystals for X-ray analysis were obtained; however, based on NOE interactions and presuming a common biosynthetic route to **1**, **2**, and **6**, the relative configuration is presumably that shown in Figure 4. On the basis of the above data, **6** was characterized as methyl 2-[6 $\alpha$ -(1-*E*-cinnamido-1 $\beta$ -hydroxy-2 $\alpha$ -methoxy-4-oxacyclohex-1-yl)]acetate and was assigned the trivial name toussaintine F.

Toussaintine G (**7**) was obtained as a white solid and was assigned the molecular formula C<sub>19</sub>H<sub>23</sub>NO<sub>6</sub> based on HRMS (ESI) data ( $[\text{M} + \text{H}]^+$   $m/z$  obs 362.1429, calcd 362.1604). It



**Figure 4.** Key NOE correlations (blue arrows) observed for compound **6** (mixing time 700 ms,  $\text{CDCl}_3$ , 25 °C, 800 MHz, spectrum S44, Supporting Information).

exhibited spectroscopic features similar to those of **6**, suggesting a closely related cinnamide-based structure, with significant chemical shift differences observed only for their C-3 to C-5 region (Table 2, Supporting Information). The  $^3J_{\text{H-5,H-6a}}$  (5.8 Hz) and  $^3J_{\text{H-5,H-6b}}$  (4.3 Hz) gauche couplings along with the  $^4J_{\text{H-2b,H-6b}}$  (1.9 Hz, Figure 3) indicate that the cyclohexanone ring of **7** prefers a main conformation that is comparable to that of **6**. However, its H-3 is axial, as revealed by its trans  $^3J_{\text{H-2a,H-3}}$  (9.7 Hz) and cis  $^3J_{\text{H-2b,H-3}}$  (4.8 Hz) values, in contrast to **6**, whose H-3 is equatorial and exhibits two gauche vicinal couplings. The absence of  $^4J_{\text{H-3,H-5}}$  *W*-coupling (Figure 3) and the observation of an NOE between the axial H-3 ( $\delta_{\text{H}}$  3.50) and the amido proton ( $\delta_{\text{H}}$  5.71) of the axial substituent further corroborate the above conclusion. The similar  $^3J_{\text{HH}}$  value and NOE pattern of the  $\text{CH}_2\text{-6ab-H-5-H-1'}$  region of **6** and **7** suggest their comparable relative configurations at C-4 and C-5 as well as their similar overall conformation. On the basis of the NMR data and presuming a common biosynthetic route to **1**, **2**, and **6**, compound **7** was characterized as methyl 2-[6 $\alpha$ -(1-*E*-cinnamido-1 $\beta$ -hydroxy-2 $\beta$ -methoxy-4-oxacyclohex-1-yl)] and was given the trivial name toussaintine G.

Motivated by the use of *T. orientalis* in traditional medicine and by the previously reported antimycobacterial<sup>6</sup> and cytostatic activities<sup>8</sup> of cinnamoyl derivatives, compounds **1**–**7** and the crude leaf extract of *T. orientalis* were evaluated for in vitro activity against *Mycobacterium tuberculosis* (H37Rv strain) and for cytotoxicity against the aggressive human breast cancer cell line (MDA-MB-231, Table 3). Both the crude methanol extract and its isolated constituents exhibit weak antimycobacterial activities (MIC 83.3–107.7  $\mu\text{M}$ ) against the H37Rv strain of *M. tuberculosis*. Compounds **1**, **3**, **4**, and **5** are cytotoxic, with  $\text{ED}_{50}$  values of 15.3–105.7  $\mu\text{M}$ . Sample limitations precluded assessment of **2**, **6**, and **7** for cytotoxicity.

Thus, seven compounds were isolated from the leaves of *T. orientalis*. Three of these are new natural products, whereas the relative configurations of three additional compounds are revised here. The crude extract and its isolated constituents exhibit weak antitubercular activity and weak to moderate toxicity against MDA-MB-231 breast cancer cells. The above

**Table 3.** Antimycobacterial (MIC  $\mu\text{M}$ ) and Cytotoxic ( $\text{ED}_{50}$   $\mu\text{M}$ ) Activities of Selected Toussaintines and of the Crude Leaf Extract of *Toussaintia orientalis*

sample	MIC <sup>a</sup>	ED <sub>50</sub> <sup>a</sup>
<i>T. orientalis</i> leaf extract	>32	53.7
<i>E</i> -toussaintine E ( <b>1</b> )	96.9	15.3
<i>Z</i> -toussaintine E ( <b>2</b> )	96.9	n.d.
toussaintine A ( <b>3</b> )	107.7	105.7
toussaintine C ( <b>4</b> )	83.3	23.6
toussaintine D ( <b>5</b> )	107.0	70.6
toussaintine F ( <b>6</b> )	88.4	n.d.
toussaintine G ( <b>7</b> )	88.4	n.d.
isoniazide (control)	0.3	
rifampicin (control)	0.9	

<sup>a</sup>MIC and  $\text{ED}_{50}$  are given in  $\mu\text{g}/\text{mL}$  for crude and in  $\mu\text{M}$  for pure compounds; n.d., not determined.

observations provide useful information for the understanding of the phytochemistry of *Toussaintia* species and may provide insight into the antitubercotic and cytostatic properties of cinnamoyltetraketides.

## EXPERIMENTAL SECTION

**General Experimental Procedures.** Column chromatography (CC) was carried out using silica gel 60 (230–400 mesh). NMR spectra were acquired on Varian VNMR-S 500 and Bruker Avance III HD 800 MHz spectrometers and were processed using MestReNova-10.0. Structural assignment was based on  $^1\text{H}$  and  $^{13}\text{C}$  NMR, COSY,<sup>10</sup> NOESY,<sup>11</sup> HSQC,<sup>12</sup> and HMBC<sup>13</sup> spectra. The solvent residual peak was used for chemical shift referencing ( $\text{CHCl}_3$ ,  $\delta_{\text{H}}$  7.26 and  $\delta_{\text{C}}$  77.16). LC-MS (ESI) chromatograms were acquired on a PerkinElmer PE SCIEX API 150 EX instrument equipped with a Turbolon spray ion source and a Gemini 5 mm RPC<sub>18</sub> 110 Å column, applying a  $\text{H}_2\text{O}/\text{MeCN}$ , 80:20–20:80 (0.2% formic acid), gradient solvent system with a separation time of 8 min. HRMS (ESI) was obtained with a Q-TOF-LC/MS spectrometer (Stenhagen Analys Lab AB, Gothenburg, Sweden) using a  $2.1 \times 30$  mm, 1.7  $\mu\text{m}$ , RPC<sub>18</sub> column and a  $\text{H}_2\text{O}/\text{MeCN}$  gradient system (5:95–95:5 gradient and 0.2% HCOOH). Analytical thin-layer chromatography (TLC) was performed on silica gel 60 F<sub>254</sub> (Merck) precoated aluminum plates, which after development with an appropriate solvent system were evaluated under UV light (254 and 366 nm) and then sprayed with 4-anisaldehyde reagent (prepared by mixing 3.5 mL of 4-anisaldehyde with 2.5 mL of concentrated  $\text{H}_2\text{SO}_4$ , 4 mL of glacial HOAc, and 90 mL of MeOH) followed by heating for identification of UV-negative compounds and detection of the color change of the UV positive spots. Gel filtration was carried out over Sephadex LH-20 (Pharmacia) suspended in  $\text{CH}_2\text{Cl}_2/\text{MeOH}$  (1:1) or 100% MeOH. Preparative HPLC was performed on a Waters 600E system using the Chromulan (Pikron Ltd.) software and an RP-C<sub>8</sub> Kromasil column (250 mm  $\times$  25 mm) with the solvent system  $\text{H}_2\text{O}/\text{MeOH}$  (gradient, 70:30 to 100% MeOH, for 20–40 min, flow rate of 8–15 mL/min).

**Plant Material.** The leaves of *T. orientalis* were collected on January 15, 2013, from the Gongo village at the edge of Saadan National Park, Bagamoyo District, Pwani Region, Tanzania. The plant was identified in the field by Frank M. Mbago and confirmed in the Herbarium of the Botany Department, University of Dar-es-salaam, where a voucher specimen is preserved as FMM 3330.

**Extraction and Isolation.** The air-dried and pulverized leaves (1 kg) were extracted with MeOH for 48 h ( $2 \times 500$  mL) at room temperature. The extracts were concentrated using a rotary evaporator at 40 °C to give 113 g of crude extract. Vacuum liquid chromatography (silica gel, 30–100% v/v EtOAc/petroleum ether gradient) of the leaf extract (80 g) gave 10 fractions, ca. 250 mL each, which were concentrated and analyzed by TLC using 4-anisaldehyde spray reagent. The combined fractions 3–4, obtained by eluting with 50%

EtOAc/petroleum ether, were passed through an activated charcoal column using  $\text{CH}_2\text{Cl}_2$  eluent to remove chlorophyll, providing 10 fractions, ca. 50 mL each. Subfractions 3–10 were subjected to CC on silica gel using a 50–75% EtOAc/petroleum ether gradient to yield 30 subfractions, ca. 20 mL each. The contents of subfractions 9–13 were white solids following evaporation. These were combined and recrystallized from a 1:1 mixture of MeOH/MeCN, yielding *E*-toussaintine E (1, 87.2 mg, anisaldehyde: yellow-orange), whereas subfractions 18–21 yielded *Z*-toussaintine E (2, 9.3 mg, anisaldehyde: yellow-orange) following CC using Sephadex LH-20 and MeOH as eluent, followed by HPLC (C-8 column, 70:30 to 100:0  $\text{H}_2\text{O}/\text{MeOH}$  gradient for 20 min, flow rate of 8 mL/min). The combined VLC fractions 6 and 7, upon removal of chlorophyll using activated charcoal, were subjected to repeated CC purification on silica gel 60, with 75% EtOAc/petroleum ether as eluent, followed by HPLC (C-8 column, 70:30 to 100:0  $\text{H}_2\text{O}/\text{MeOH}$  gradient for 20 min, flow rate of 8 mL/min), affording toussaintine F (6, 8.5 mg, anisaldehyde: yellow) and toussaintine G (7, 6.4 mg, anisaldehyde: yellow). Following the removal of chlorophyll using activated charcoal and  $\text{CH}_2\text{Cl}_2$ , the combined VLC fractions 7–8 were subjected to repeated column chromatographic purification using silica gel 60 and 75% EtOAc/petroleum ether eluent, followed by gel filtration on Sephadex LH-20, with MeOH as eluent, yielding toussaintine C (4, 22.4 mg, anisaldehyde: orange). Semipurified fractions<sup>1</sup> containing 3, 4, and 5 were purified by CC using silica gel 60 with 30–50% EtOAc/isohexane as eluent, to afford toussaintine A (3, 30 mg, anisaldehyde: yellow), whereas with 50–75% EtOAc/isohexane as eluent a mixture yielding toussaintine C (4) and toussaintine D (5) was obtained. Fractions containing 4 and 5 were further purified by HPLC using a C-8 column with a 70:30 to 100:0  $\text{H}_2\text{O}/\text{MeOH}$  gradient for 20 min with a flow rate of 8 mL/min, providing toussaintine C (4, 12 mg, anisaldehyde: orange) and toussaintine D (5, 22 mg, anisaldehyde: pink). Compounds 3 and 5 were crystallized from a 1:1 mixture of MeCN/MeOH.

*E*-Toussaintine E (1) (ref 1): white solid, MS (ESI)  $m/z$  330.3 [M + H]<sup>+</sup>; <sup>1</sup>H and <sup>13</sup>C NMR data, see Supporting Information.

*Z*-Toussaintine E (2): white solid,  $[\alpha]_D = -8$  ( $\text{CH}_3\text{OH}$ ,  $c$  0.0015); HRESIMS  $m/z$  330.1134 [M + H]<sup>+</sup> (calcd for  $\text{C}_{18}\text{H}_{19}\text{NO}_5$ , 330.3472); <sup>1</sup>H and <sup>13</sup>C NMR data, see Table 1 and Supporting Information.

Toussaintine A (3) (ref 1): white, amorphous solid; MS (ESI)  $m/z$  298.6 [M + H]<sup>+</sup>; <sup>1</sup>H and <sup>13</sup>C NMR data, see Tables S2 and S3, Supporting Information.

Toussaintine C (4) (ref 1): white, amorphous solid; MS (ESI)  $m/z$  284.4 [M + H]<sup>+</sup>; <sup>1</sup>H and <sup>13</sup>C NMR data, see Tables S2 and S3, Supporting Information.

Toussaintine D (5) (ref 1): white, amorphous solid; MS (ESI)  $m/z$  300.5 [M + H]<sup>+</sup>; <sup>1</sup>H and <sup>13</sup>C NMR data, see Tables S2 and S3, Supporting Information.

Toussaintine F (6): white, amorphous solid;  $[\alpha]_D = -14$  (MeOH,  $c$  0.001); HRESIMS [M + H]<sup>+</sup>  $m/z$  362.1616 (calcd for  $\text{C}_{19}\text{H}_{23}\text{NO}_6$ , 362.1604); <sup>1</sup>H and <sup>13</sup>C NMR data see Table 2.

Toussaintine G (7): white, amorphous solid;  $[\alpha]_D = -10$  (MeOH,  $c$  0.001); HRESIMS [M + H]<sup>+</sup>  $m/z$  362.1429 (calcd for  $\text{C}_{19}\text{H}_{23}\text{NO}_6$ , 362.1604); <sup>1</sup>H and <sup>13</sup>C NMR data see Table 2.

**Cytotoxicity Assay.** The cytotoxic activity of the methanol extract and of its constituents was evaluated against human breast cancer cells (MDA-MB-231) as previously reported by Irungu.<sup>14</sup> MDA-MB-231 human breast cancer cells were cultured in Dulbecco's modified Eagle medium (DMEM) supplemented with 10% (v/v) fetal bovine serum, 2 mM L-glutamine, 100 units/mL penicillin, and 100  $\mu\text{g}/\text{mL}$  streptomycin at 37 °C in humidified 5%  $\text{CO}_2$ . For cytotoxicity assays, cells were seeded in 96-well plates at optimal cell density (10 000 cells per well) to ensure exponential growth for the duration of the assay. After a 24 h preincubation growth, the medium was replaced with experimental medium containing the appropriate drug concentrations or vehicle controls (0.1% or 1.0% v/v DMSO). After 72 h of incubation, cell viability was measured using Alamar Blue reagent (Invitrogen Ab, Lidingö, Sweden) according to the manufacturer's instructions. Absorbance was measured at 570 nm with 600 nm as a reference wavelength. Results were expressed as the mean  $\pm$  standard

error for six replicates as a percentage of vehicle control (taken as 100%). Experiments were carried out three independent times with six replicates each time. Statistical analyses were performed using a two-tailed Student's *t*-test.  $p < 0.05$  was considered to be statistically significant.

**Antimycobacterial Activity Assay.** Antimycobacterial activity of the MeOH extract and its purified constituents was evaluated against the H37Rv strain of *Mycobacterium tuberculosis* using rifampicin (RIF) and isoniazide (INH) as positive controls. The minimum inhibitory concentration against replicating *M. tuberculosis* was determined by the microplate Alamar Blue assay (MABA). Compound stock solutions and the range of final testing concentrations were 64  $\mu\text{g}/\text{mL}$  and 32–0.5  $\mu\text{g}/\text{mL}$ , respectively. *M. tuberculosis* (H37Rv) was grown to late log phase (70 to 100 Klett units) in Difco Middlebrook 7H9 broth (catalog no. 271310) supplemented with 0.2% (v/v) glycerol, 0.05% Tween 80, and 10% (v/v) albumin–dextrose catalase (BBL Middlebrook ADC Enrichment, catalogue no. 212352, 7H9-ADCTG). The cultures were centrifuged, washed twice, and then resuspended in phosphate-buffered saline. Suspensions were passed through an 8  $\mu\text{m}$  pore-size filter to remove clumps, and the aliquots were frozen at –80 °C. Twofold dilutions of the studied compounds were prepared in 7H9-ADC-TG in a volume of 100  $\mu\text{L}$  in 96-well, black, clear-bottom microplates (BD Biosciences, Franklin Lakes, NJ, USA). *M. tuberculosis*, 100  $\mu\text{L}$  containing  $2 \times 10^5$  CFU, was added, yielding a final testing volume of 200  $\mu\text{L}$ . The plates were incubated at 37 °C. On the seventh day of incubation, 12.5  $\mu\text{L}$  of 20% Tween 80 and 20  $\mu\text{L}$  of Alamar Blue were added to all wells. Following incubation at 37 °C for 16–24 h, the fluorescence was read at excitation  $\lambda = 530$  nm and emission  $\lambda = 590$  nm. The MIC was defined as the lowest concentration effecting a reduction in fluorescence of  $\geq 90\%$ , relative to the mean of replicate bacterium-only controls.

## ■ ASSOCIATED CONTENT

### 📄 Supporting Information

1D and 2D NMR, MS spectra, X-ray crystallography. The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.jnatprod.5b00356.

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### Notes

The authors declare no competing financial interest.

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