Chemical constituents and antimycobacterial studies of the leaf extracts of *Pavetta crassipes* K. Schum

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Abstract: Six known compounds; β-sitosterol, ursolic acid, methyl chlorogenate, ethyl chlorogenate, rutin and mannitol were isolated from the leaf extracts of *Pavetta crassipes* (Rubiaceae), a Nigerian medicinal plant used in the indigenous treatment of tuberculosis. Separation and isolation of the compounds were achieved by chromatographic techniques and the structures of isolated compounds were established by spectroscopic and chemical methods. The isolated compounds were screened for antimycobacterial activities against *Mycobacterium tuberculosis* H₃₇Rv employing the green fluorescence protein reporter microplate assay and the broth microdilution method. Ursolic acid, methyl chlorogenate and ethyl chlorogenate were found moderately active in the broth microdilution assay with MICs of 200, 100 and 50 µg/ml, respectively while methyl chlorogenate and ethyl chlorogenate were active in the protein reporter microplate assay with MICs of 200 and 100 µg ml⁻¹, respectively. The presence of antimycobacterial terpenoids and quinate esters in leaves of *Pavetta crassipes* provides scientific evidence for the ethnomedicinal use of the plant as a traditional anti-tuberculosis remedy.

Keywords: *Pavetta crassipes* - Spectroscopy - Antimycobacterial activity - Broth microdilution assay - Green fluorescent protein reporter microplate assay.

INTRODUCTION

Tuberculosis (TB) remains a major global health concern as it is one of the world’s deadliest diseases. Approximately one third of the world’s population is infected with TB and according to a 2015 World Health Organization (WHO) report, an estimated 10.4 million people fell ill from TB and 1.8 million died from the disease. Africa has the second largest incidence of tuberculosis (26% of global total) with Nigeria as one of the six countries that accounted for 60% of new cases (WHO 2016). Current anti-tuberculosis therapy is a regimen of isoniazid, rifampicin and pyrazinamide administered over a period of six months which leads to poor adherence by patients. Inadequate, incomplete, or improperly supervised treatment regimen, poor prescriptions, and co-infection with HIV have caused the emergence of resistant strains of *Mycobacterium tuberculosis*. Natural products have played and continue to play a significant role in the drug discovery process with tremendous research all across the continents for novel anti-tuberculosis agents. Some excellent review articles on natural products have been published on different classes of compounds; exhibiting antimycobacterial activity (Cantrell *et al*. 2001, Copp 2003, Okunade *et al*. 2004, Copp & Pearce 2007).

African traditional medicine is the oldest, and perhaps the most assorted, of all therapeutic systems and the most common practice across the continent is the use of medicinal plants (Mahomoodally 2013). This practice...
plays an important role in the health care delivery of all African nations and is common especially in the rural areas due to availability, low cost and sometimes the myth associated with these herbs. The rich biodiversity and ethnomedical history of the Nigerian flora portend great possibilities in research for novel anti-tubercular compounds. This is however under-explored as only a negligible number of plants used locally in the treatment of characteristic symptoms of TB have been investigated for their antimycobacterial properties.

The leaves of Pavetta crassipes K. Schum (Rubiaceae) are traditionally used in Northern Nigeria for the management of respiratory disorders and TB related symptoms, in ethnombotanical practice. A plant widely distributed in the West African sub-region, P. crassipes is a glabrous shrub to 6 m high, trunk to 30 cm girth, of the savanna. It has stout square branchlets covered with pale corky bark which splits and falls off; leaves often in threes; flowers greenish-white and fruits black (Burkill 1997). The following pharmacological activities have been reported on the plant; anti-plasmodial/malarial activity (Sanon et al. 2003), hypotensive activity (Amos et al. 2003), inhibitory effects on gastrointestinal and uterine smooth muscles (Amos et al. 1998) and in vitro antiprotozoal, antimicrobial and antitumor activities (Balde et al. 2010). In our search for novel antimycobacterial agents from Nigerian medicinal plants, the leaves of P. crassipes were collected from four different traditional medicine practitioners (TMPs) in Nigeria, as a component of their local herbal recipes for the treatment of tuberculosis (Ibekwe et al. 2014). The ethnobotanical survey revealed P. crassipes to be an interesting plant worth further investigations and this was based on the frequency of plant in the collected recipes from TMPs, potency of the extract based on MIC values, minimal published work on the chemistry of the plant, and novelty of information on plant’s use as an anti-TB remedy, hence it was prioritized for further studies. Preliminary studies on the extracts revealed the antimycobacterial potentials of the plant (Ibekwe et al. 2012). There is presently no detailed report on the phytochemistry or antimycobacterial studies of the metabolites of the plant. We report herein, the isolation, structural elucidation of six compounds from the leaf extracts of P. crassipes and screening of isolated compounds against M. tuberculosis H37Rv.

MATERIALS AND METHODS

General Experimental Procedures

Melting points were determined with a Barnstead Electrothermal 9100 melting point apparatus and are uncorrected. IR spectra were recorded with a Perkin Elmer Spectrum One FT-IR Spectrophotometer. UV spectra of isolated compounds were run on a Varian Cary 300 Bio UV-Visible Spectrophotometer. High-Resolution Mass Spectra (HRMS) were obtained on a q-TOF Waters LCT Premier Mass Spectrometer with electrospray ionization (ESI) or atmospheric pressure ionization (API) source and in the positive or negative mode. \( ^1H \) (400 MHz), \( ^13C \) NMR (100 MHz) and 2D spectral data were recorded in deuterated CDCl\(_3\), DMSO-\(d_6\), D\(_2\)O or CD\(_3\)OD, on a Bruker AMX 400. NMR chemical shifts were expressed in parts per million (\( \delta \)) relative to tetramethylsilane (TMS) and the coupling constants were given in Hz. Mass spectra of isolated compounds were run on a Varian Cary 300 Bio UV Visible Spectrophotometer. Visible Spectrophotometer. Resolutions of isolated compounds were run on a Bruker AMX 400. NMR  chemical shifts were expressed in parts per million (\( \delta \)) relative to tetramethylsilane (TMS) and the coupling constants were given in Hz. Flash column chromatography were performed on glass columns (4 × 150 cm, 2.7 × 70 cm) using silica gel (230–400 µm mesh, Merck, Germany). Preparative TLC were carried out with pre-coated glass backed preparative TLC plates- Kieselgel 60 F\(_{254}\) (0.5 mm, 20 × 20 cm; Analtech Inc, USA). Preparative HPLC were performed on a Phenomenex system (Varian Pro Star Model 218 coupled to a Varian fraction collector model 701) using a reverse phase column 250 × 21.20 mm i.d (5µm, 100A). TLC analyses were carried out using silica 60 F\(_{254}\) pre-coated glass backed plates (0.25 mm, 20 × 20 cm; Merck, Darmstadt, Germany). Spots were detected on TLC plates under short (\( \lambda = 254 \) nm) and long (\( \lambda = 366 \) nm) UV light and/or visualized by spraying with vanillin-sulphuric acid, followed by charring at 110°C for 5 minutes. All chemicals were obtained from commercially available sources and reagents prepared according to standard procedures. The structures of the compounds were confirmed by comparing with reference data from available literature or comparing with data of commercial authentic samples, or by chemical modifications of the isolated compounds.

Plant Material

The fresh leaves of Pavetta crassipes K. Schum were collected from Suleja, Niger State, Nigeria in July 2009. The plant was identified at the Herbarium Unit of the Department of Medicinal Plant Research and Traditional Medicine, National Institute for Pharmaceutical Research and Development, Abuja (NIPRD). A labeled voucher specimen, NIPRD/H/6241 was deposited in the herbarium of the Institute.

Extraction procedure

The air dried leaves of P. crassipes (1 kg) were finely pulvserised and soaked successively with n-hexane,
EtOAc and MeOH at 24 h intervals. The hexane, EtOAc and MeOH extracts were evaporated under reduced pressure at 40°C, to give the crude hexane (green-black syrup, 10.4 g), the EtOAc (green powder, 16.8 g) and the MeOH extract (dark brownish sticky solid, 131.7 g).

Isolation of components

The hexane extract (5.0 g) was subjected to column chromatography with gradient amounts of ethyl acetate in hexane. Fractions were examined by TLC and combined to give 8 major fractions, PCH₁–PCH₈. Fraction PCH₁ (eluted with 20% EtOAc in hexane), was similarly re-chromatographed to give 5 sub-fractions, PCH₁₁–PCH₁₅. Sub-fraction PCH₁₄ (eluted with 10–15% EtOAc in hexane) was recrystallized from methanol to yield 1 (22.0 mg). Fraction PCH₆, eluted with 30% EtOAc in hexane, was re-chromatographed to obtain 6 sub-fractions, PCH₆₁–PCH₆₆. Crystallization of PCH₆₅ (eluted with 30–50% EtOAc in hexane) from ethanol, yielded 2 (34.0 mg). The ethyl acetate extract (9.0 g) was fractionated using column chromatography and eluted with gradient amounts of ethyl acetate in hexane to obtain 14 major fractions, PCE₁–PCE₁₄. Fractions PCE₄ and PCE₅, eluted with 20% and 30% EtOAc in hexane, respectively had similar TLC profiles as fractions PCH₁₄ and PCH₆, from the hexane extract. These were re-chromatographed under similar conditions to give 1 (15.0 mg) and 2 (102.0 mg). Fraction PCE₁₄, eluted with EtOAc and 10% MeOH in EtOAc, was chromatographed on a prep-HPLC system. The fractions were eluted with a MeOH-H₂O gradient in 0.01% formic acid starting from MeOH-H₂O [5:95] solvent A to MeOH-H₂O [95:5] solvent B in 50 minutes with a flow rate of 25 ml min⁻¹. Fractions with similar retention times were combined. Sub-fractions PCE₁₄₂₅–PCE₁₄₃₀ and PCE₁₄₃₁–PCE₁₄₃₂, eluted with MeOH-H₂O [80:20] and MeOH-H₂O [84:16] were pooled. This procedure was repeated to obtain sufficient amounts of the compounds. Sub-fractions PCE₁₄₃₅–PCE₁₄₃₀ and PCE₁₄₃₁–PCE₁₄₃₂ were further purified using prep TLC developed with CHCl₃-MeOH [9:1]. The separated fractions were suspended in methanol for 30 min, filtered under suction while washing with more methanol. Thereafter, the solvent was evaporated under vacuum, to yield 3 (8.0 mg) and 4 (18.0 mg). The methanol extract (10.0 g) was fractionated using column chromatography, with a gradient elution of ethyl acetate in hexane, and methanol in ethyl acetate to obtain thirteen major fractions, PCM₁–PCM₁₃. Compound 5 (56.0 mg) precipitated on standing from fraction PCM₁₀ (eluted with 20% MeOH in EtOAc) as a yellow powder and was further purified by washing with EtOAc. Fraction PCM₁₀ was further similarly re-chromatographed on a HPLC system to obtain sub-fractions PCM₁₀₂₅–PCM₁₀₃₀, eluted with MeOH-H₂O [80:20]. Similar fractions were combined and purification of PCM₁₀₂₅–PCM₁₀₃₀ was carried out using prep-TLC developed with CHCl₃-MeOH [9:1] to give compound 3 (12.0 mg). Fraction PCM₁₁, eluted by 50% MeOH in EtOAc, yielded light brown crystals on standing. Further purification by recrystallizing from methanol yielded compound 6 as a white solid (68.0 mg).

**Physical and Spectral Data of Isolated Compounds**

**β-sitosterol (1):** white crystals, mp 133-135 °C; IR ν_max (cm⁻¹): 3356 (O-H), 2959, 2933 (C-H), 1464, 1378, 1214, 1060 (C-O); TOF-ES-MS m/z: 397.4 [M - H₂O]⁺; ¹H NMR (400 MHz, CDCl₃) δ_H (ppm): 5.34 (1H, d, H-6), 3.52 (1H, ddd, H-3), 0.82 (3H, d, H-26), 0.80 (3H, d, H-27), 1.00 (3H, d, H-19), 0.91 (3H, d, J = 6 Hz, H-21), 0.84 (3H, t, J = 8.4 Hz, H-29), 0.67 (3H, s, H-18); ¹³C NMR (100 MHz, CDCl₃) δ_C (ppm): 140.7 (C-5), 121.7 (C-6), 72.0 (C-3), 56.7 (C-14), 56.0 (C-17), 50.1 (C-9), 45.9 (C-24), 42.3 (C-13), 40.4 (C-12), 39.8 (C-4), 37.2 (C-1), 36.5 (C-10), 33.9 (C-22), 31.9 (C-7), 31.6 (C-8), 29.7 (C-2), 29.1 (C-25), 28.2 (C-16), 26.0 (C-23), 24.3 (C-15), 23.0 (C-28), 21.1 (C-11), 19.8 (C-26), 19.3 (C-27), 19.0 (C-19), 18.7 (C-21), 12.0 (C-29), 11.8 (C-18).

**Ursolic acid (2):** amorphous white solid, mp 284-286 °C; IR ν_max(cm⁻¹): 3673 (O-H), 1688 (C=O); TOF-ES-MS m/z: 457.4 [M + H]⁺, 439.4 [M - H₂O]⁺; ¹H NMR (400 MHz, DMSO-d₆) δ_H (ppm): 11.92 (1H, s, COOH-28), 5.12 (1H, t, H-12), 4.27 (1H, d, OH-3), 2.99 (1H, m, H-3), 2.09 (1H, d, H-18), 1.03 (3H, s, H-27), 0.90 (3H, d, H-30), 0.89 (3H, s, H-23), 0.86 (3H, s, H-25), 0.81 (3H, d, H-29), 0.74 (3H, s, H-26), 0.67 (3H, s, H-24); ¹³C NMR (100 MHz, DMSO-d₆) δ_C (ppm): 178.2 (C-28), 138.1 (C-13), 124.5 (C-12), 76.8 (C-3), 54.7 (C-5), 52.3 (C-18), 46.9 (C-9), 46.7 (C-17), 41.6 (C-14), 38.8 (C-8), 38.47 (C-19), 38.40 (C-20), 38.3 (C-4), 38.2 (C-1), 36.5 (C-10), 36.2 (C-22), 32.6 (C-7), 30.1 (C-21), 28.2 (C-23), 27.5 (C-2), 26.9 (C-15), 23.7 (C-16), 23.2 (C-27), 22.8 (C-11), 21.0 (C-30), 17.9 (C-6), 16.9 (C-26), 16.8 (C-29), 16.0 (C-24), 15.2 (C-25).

**Methyl chlorogenate (3):** amorphous yellow solid; (MeOH) λ_max (nm): 218, 330; TOF-ES-MS, positive ion, m/z 391.1 [M + Na]⁺, 369.1 [M + H]⁺; HRESIMS (positive ion mode) m/z 369.1191[M + H]⁺ (calculated for C₁₆H₁₉O₉ + H, 369.1185); ¹H NMR (400 MHz, CD₃OD) δ_H (ppm): 7.52 (1H, d, J = 15.9 Hz, H-7), 7.03 (1H, d,
1998) and a modified Broth Microdilution Method (BMM; Coban et al. 2013) to test of antimycobacterial activity. Assays of compounds were carried out utilizing a constitutive GFP plasmid was used as a test strain. Compounds were prepared in methanol with concentrated HCl.

**Ethyl chlorogenerate (4):** amorphous yellow solid; UV (MeOH) $\lambda_{max}$ (nm): 218, 330; TOF-ES-MS, positive ion, $m/z$ 405.1 [M + Na]$^+$, 383.1 [M + H]$^+$; HRESIMS (positive ion mode) $m/z$ 383.1344[M + H]$^+$ (calculated for C$_9$H$_{12}$O$_3$ + H, 383.1342); $^1$H NMR (400 MHz, DMSO-$d_6$) $\delta$ (ppm): 7.38 (1H, d, J = 15.9 Hz, H-7), 7.01 (1H, d, J = 2.0 Hz, H-2), 6.95 (1H, dd, J = 2.0, 8.1 Hz, H-6), 6.76 (1H, d, J = 8.1 Hz, H-5), 6.10 (1H, d, J = 15.9 Hz, H-8), 5.01 (1H, m, H-3), 4.01 (2H, m, H-4), 3.86 (1H, m, H-5), 3.56 (1H, dd, H-4), 1.91-2.10 (2H, m, H-2), 1.75-2.10 (2H, m, H-6), 1.12 (3H, t, H-9); $^{13}$C NMR (100 MHz, CD$_3$OD) $\delta_c$ (ppm): 175.0 (C-7), 168.3 (C-9), 149.7 (C-3'), 147.2 (C-7), 146.9 (C-4'), 127.7 (C-1'), 123.0 (C-6), 116.5 (C-5'), 115.18 (C-7'), 75.9 (C-1), 72.6 (C-4), 72.1 (C-3), 70.4 (C-5), 53.0 (C-1), 38.1 (C-2), 37.8 (C-6)

**Rutin (5):** amorphous yellow solid; UV (MeOH) $\lambda_{max}$ (nm): 257, 358; IR $\nu_{max}$ (cm$^{-1}$): 3337 (O-H), 2943, 2833 (C-H), 1655 (C=O), 1606 (C=C; aromatic), 1448, 1361, 1306, 1202, 1170; TOF-ES-MS, positive ion, $m/z$ 633.1 [M + Na]$^+$, 611.1 [M + H]$^+$, 465.1 [M + H - 146]$^+$, 449.1 [M + H - 162]$^+$; HRESIMS (positive ion mode) $m/z$ 611.1599 [M + H]$^+$ (calculated for C$_{18}$H$_{18}$O$_{16}$ + H, 611.1611); $^1$H NMR (400 MHz, CD$_3$OD): $\delta$ (ppm): 7.66 (d, J = 2.0 Hz, H-2), 7.62 (dd, J = 2.0, 8.0 Hz, H-6), 6.85 (d, J = 8.0 Hz, H-5), 6.39 (d, J = 2.0 Hz, H-8), 6.20 (d, J = 2.0 Hz, H-6); glucosyl resonances: 5.11 (1H, d, J = 7.6 Hz, H-1'), 3.80 (1H, d, H-6), 3.44 (1H, t, J = 3.2 Hz, H-2'), 3.38 (1H, dd, H-6'); rhamnosyl resonances: 4.51 (1H, d, J = 1.6 Hz, H-1'), 1.12 (3H, d, J = 6 Hz, H-6'); other glycosidic resonances: 3.24 - 3.49 (multiplet); $^{13}$C NMR (100 MHz, CD$_3$OD) $\delta_c$ (ppm): 173.9 (C-4), 166.0 (C-7), 162.9 (C-5), 159.3 (C-2), 158.5 (C-9), 149.8 (C-4'), 145.8 (C-3'), 135.6 (C-3), 123.5 (C-6), 123.1 (C-1'), 117.6 (C-2), 116.0 (C-5), 105.6 (C-10), 99.9 (C-9), 94.8 (C-8), 93.8 (C-7), glucosyl and rhamnose: 104.7 (C-1'), 102.4 (C-2'), 75.7 (C-2'), 68.5 (C-6'), 17.8 (C-6'), other glycosidic carbons (69.7, 71.3, 72.1, 72.2, 73.9, 77.2 and 78.1)

**Mannitol (6):** white solid, mp 154-157 °C; IR $\nu_{max}$ (cm$^{-1}$): 3329 (O-H), 2989 (C-H), 2108, 1637, 1394, 1250, 1066 (C=O); TOF-AP-MS, positive ion, $m/z$ 183.1 [M + 1]$^+$, 165.1 [M + 1 - H$_2$O]$^+$, 147.1 [165.1 - H$_2$O]$^+$, 129.1 [147.1 - H$_2$O]$^+$, 111.1 [129.1 - H$_2$O]$^+$; HRESIMS (positive ion mode) $m/z$183.0872 [M + H]$^+$ (calculated for C$_6$H$_{12}$O$_6$ + H, 183.0868); $^1$H NMR (400 MHz, D$_2$O) $\delta$ (ppm): 3.75 (2H, dd, J = 2.4, 11.7 Hz, H-1, 6), 3.68 (2H, d, J = 8.6 Hz, H-3, 4), 3.64 (2H, m, H-2, 5), 3.56 (2H, dd, J = 6, 11.7 Hz, H-1, 6); $^{13}$C NMR (100 MHz, CD$_3$OD) $\delta_c$ (ppm): 70.8 (C-2, 5), 69.2 (C-3, 4) and 63.2 (C-1, 6)

**Acid hydrolysis of rutin (5):** Compound 5 (10 mg) was dissolved in 1 ml of methanol with concentrated HCl (0.5 ml) and the solution was kept under reflux for 5 h at 70°C. After removal of MeOH by rotary evaporation, the residue was partitioned between n-butanol and H$_2$O. The butanol (lower) layer was removed, dried over anhydrous Na$_2$CO$_3$ and concentrated under reduced pressure, to afford 3 mg of 5a.

**Quercetin (5a):** amorphous yellow solid, mp 297-300 °C; UV (MeOH) $\lambda_{max}$ (nm): 257, 358; TOF-MS-API, positive ion, $m/z$ 303.1 [M + 1]$^+$; $^{13}$C NMR (100 MHz, CD$_3$OD) $\delta_c$ (ppm): 176.5 (C-4'), 166.0 (C-7), 161.0 (C-5), 156.7 (C-9), 148.1 (C-4'), 147.5 (C-2), 145.7 (C-3'), 136.5 (C-3), 123.0 (C-1'), 121.0 (C-6'), 116.5 (C-2'), 116.0 (C-5'), 104.0 (C-10), 99.5 (C-6), 94.5 (C-8)

**Antimycobacterial assay**

Following the extractions, chromatographic separations and purification of compounds from the leaves of *P. crassipes*, compounds were screened against *Mycobacterium tuberculosis* and bioassays were conducted on drug sensitive H$_3$Rv ATCC 27294. *M. tuberculosis* was grown in 7H9-medium (which consisted of Middlebrook 7H9 broth base supplemented with 0.5 % (w/v) Albumin, 0.2 % (w/v) glucose, 0.2% (v/v) glycerol, 0.08% (w/v) NaCl and 0.05 % (v/v) Tween 80), to an optical density (OD) 650 nm of 0.2–0.3 after which cells were diluted 1000-fold in 7H9-medium. The Green Fluorescent Protein Reporter Microplate Assay (GFPPMA; Collins et al. 1998) and a modified Broth Microdilution Method (BMM; Coban et al. 2004) were employed in the assessment of antimycobacterial activity. Assays of compounds were carried out utilizing a constitutive Green Fluorescence Protein (GFP) expression vector direct readout of fluorescence (with excitation at 485 nm and emission at 509 nm) as a measure of bacterial growth. *Mycobacterium tuberculosis* H$_3$Rv with a constitutive GFP plasmid was used as a test strain. Compounds were prepared in 100% DMSO at an initial stock
concentration of 10 mg mL⁻¹ and serial dilutions of the compounds were prepared in the same solvent and added to the wells of a black clear-bottom 384-well microtiter plate (in order to minimize background fluorescence) at 2 µL volume compound per well. Forty eight (48) µL of H₃Rv-GFP bacterial suspension was added to the different concentrations of the compound across the wells resulting in a final volume of 50 µL. Plates were then incubated at 37°C for 5 days. Mycobacterial growth was determined by measuring GFP fluorescent intensity at 509 nm using a multilabel reader. The increase in fluorescence indicated growth of the GFP-expressing strain whereas a lack of increase of fluorescence readout or even a decrease in fluorescence relative to the day 0 fluorescence value, indicated growth inhibition. Isoniazid was used as positive control (100% growth) and DMSO as negative control (0% growth). All the bioassay experiments were done in duplicates and assay results were reported in the form of MIC values. For the broth microdilution assay, pure compounds for biological assay were prepared at a concentration of 10 mg ml⁻¹ in 100% DMSO and 40 µl of the stock solution was taken into 460 µL 7H9-medium. Serial two fold dilutions were made in the broth medium at 50 µL per well in clear 96-well round-bottom microtiter plates after which an equal volume of diluted M. tuberculosis H₃Rv was added at 10,000 cells per well with the final top concentration of DMSO at 2% and incubated at 37°C in a humid atmosphere for 7–10 days. Growth was visually scored using an enlarging inverted mirror. Isoniazid was used as positive control. The minimum inhibitory concentration (MIC) was taken as the lowest concentration that completely inhibited all visible growth.

RESULTS AND DISCUSSION

Repeated column chromatography (silica gel), prep HPLC and prep TLC of the extracts of the leaves of Pavetta crassipes yielded compounds 1–6 (Fig. 1). The chemical structures of the isolated compounds were established using spectral data obtained from UV, IR, MS and ¹H and ¹³C NMR spectra in conjunction with 2D experiments, COSY, HSQC and HMBC. These were compared with published literature and in one of the cases, with the authentic commercial specimen (Compound 2). Chemical modification and spectral analysis of reaction product were useful in establishing the structure of Compound 5.

The compounds isolated from the leaves of P. crassipes were identified as ß-sitosterol C₂₅H₃₅O (1) (Lee et al. 2003), ursolic acid (2) C₂₉H₄₈O₄ (Seebacher et al. 2003), methyl chlorogenate C₁₇H₁₅O₄ (3) (Lee et al. 2010), ethyl chlorogenate C₁₃H₂₀O₆ (4) (Lee et al. 2010), rutin C₁₇H₂₀O₁₆ (5) (Lallemand & Duteil 1977, Bello et al. 2011) and mannitol C₆H₁₂O₆ (6) (Hagiwara et al. 2005). Quercetin C₁₅H₁₀O₇ (5a), the hydrolyzed product of rutin, was also identified (Lallemand & Duteil 1977). The structures of the compounds are as shown in figure 1 and the spectroscopic and physical data are presented in the section on Material and Methods.

All compounds were screened for their inhibitory activity on the growth of Mycobacterium tuberculosis H₃Rv, with final concentrations of 200 µg mL⁻¹ (GFPM) and 400 µg mL⁻¹ (BMM) and the results are presented in table 1. The GFPM assay revealed only the quinate esters; ethyl chlorogenate (4) and methyl chlorogenate (3) with activities against M. tuberculosis H₃Rv at 100 and 200 µg ml⁻¹, respectively, while the BMM assay showed ethyl chlorogenate (4) methyl chlorogenate (3) and ursolic acid (2), with activities with MIC of 50, 100 and 200 µg/ml, respectively. Interestingly, the bioactive chlorogenate esters are structurally similar, but for the length of the alkyl substituent, with ethyl chlorogenate being less polar. The differences in biological activities of the chlorogenate esters may be owed to their differences in polarity as less polar compounds have been shown to inhibit the growth of M. tuberculosis due to the easier permeability of the cell wall of the bacterium which is lipophilic in nature (Ducati et al. 2006). The antimycobacterial activities of ethyl chlorogenate (4) and methyl chlorogenate (3) have not been reported hitherto. Ursolic acid (2) which is a pentacyclic triterpene, and its analogue, 24-hydroxyursolic acid have previously been reported as antimycobacterial agents from Valeriana laxiflora (Gu et al. 2004) and Lysysera gnaphaloides (Bamuamba et al. 2008). Some pentacyclic triterpenes with substituents in C-3 and C-17, such as oleanolic acid, oleanonic acid, and 3-epioleanol acid inhibited the growth of M. tuberculosis H₃Rv with MIC values of 50, 16, and 16 µg/mL, respectively (Caldwell et al. 2000). It has been reported that the presence of hydroxyl or keto groups in A or B rings, and a carboxylic group in D/E rings, gave the molecule a moderate antimycobacterial activity (Wächter et al. 1999, Caldwell et al. 2000). These authors also suggested that the mechanism of action of such triterpenoids depended on the lipophilicity of the compounds that allowed a rapid penetration across the lipid-rich mycobacterial cell wall. In their review article, Ducati et al. (2006) also stated that lipophilic molecules should be able to easily cross the mycobacterium membrane, dissolving in the hydrocarbon interior of the lipid bilayer, though factors such as low fluidity of the mycolic acid leaflet and the bilayer’s uncommon thickness.
may result in reduction of this process.

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\text{Figure 1. Structures of compounds (1–6) from the leaves of Pavetta crassipes.}
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\text{Table 1} \quad \text{Antimycobacterial activity of compounds against Mycobacterium tuberculosis H}_{37}\text{Rv.}
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<thead>
<tr>
<th>Compound</th>
<th>GFPMA MIC in µg mL(^{-1})</th>
<th>BMM MIC in µg mL(^{-1})</th>
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<td>1</td>
<td>NA*</td>
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<td>5a</td>
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<td>NA</td>
</tr>
<tr>
<td>6</td>
<td>NA</td>
<td>NA</td>
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</tbody>
</table>

\textbf{Note:} NA = not active at the concentrations tested. The MIC of isoniazid, the reference compound for the antimycobacterial assay was 0.2 µM

Comparing the MICs of the two bioassay methods in the antimycobacterial assay, BMM shows more sensitivity than GFPMA. However, since the intrinsically fluorescent nature of GFP precludes the need for a substrate, GFPMA offers greater simplicity and also has enhanced biosafety since the microplate need not be reopened following inoculation (Collins et al. 1998).
The phytochemistry of *P. crassipes* is also important as a chemotaxonomic study, as some of the isolated compounds may be useful as markers of the Rubiaceae family. Compounds such as triterpene acids, steroids, chlorogenic acid derivatives and flavonoids have been reported from other species of the Rubiaceae family (Martins & Nunez 2015).

**CONCLUSIONS**

This is the first detailed report of the phytochemistry of the leaves of *Pavetta crassipes*, with rutin as the only compound previously isolated (Bello *et al.* 2011). An extensive literature search carried out did not reveal much information on the phytochemistry of plants in the *Pavetta* genus. *In vitro* screening against *Mycobacterium tuberculosis* employing the microbroth dilution and the green fluorescent protein microplate assay techniques showed that the antituberculosis activities of *P. crassipes* leave were attributed to a uranes type triterpene acid, ursolic acid and two chlorogenate esters, methyl chlorogenate and ethyl chlorogenate. This is the first report of the antimycobacterial activities of these compounds from this plant and more interestingly the first report of chlorogenate esters as antimycobacterial agents. This group of phenolics may represent promising antimycobacterial agents and should be investigated further as potential leads in the drug discovery of antituberculosis agents. The antimycobacterial activities of the leaves of *P. crassipes* are relevant and it would be interesting to explore the potentials of the bioactive compounds; ursolic acid and the chlorogenate esters as drug templates by carrying out structural activity relationship (SAR) studies to synthesize new derivatives which may be highly specific to treat the disease. This study provides, at least in part, some scientific basis and a biological explanation for the ethnomedicinal use of *P. crassipes* as a traditional antituberculosis remedy in Nigeria, through a combination of indigenous knowledge and natural products chemistry.

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**REFERENCES**


