



## *In vitro* antimicrobial activity of the methanol extract and compounds from the wood of *Ficus elastica* Roxb. ex Hornem. aerial roots



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### ABSTRACT

*Ficus elastica* Roxb. ex Hornem., an edible plant belonging to the family of Moraceae, is traditionally used against skin infections and allergies besides having diuretic properties. This study aimed at investigating the antimicrobial activity of the wood of *F. elastica* aerial roots against a set of bacteria (*Staphylococcus aureus*; *Escherichia coli*, *Proteus vulgaris*, *Providencia stuartii* and *Pseudomonas aeruginosa*) and a yeast (*Candida albicans*). A mixture of linear aliphatic alkanes with *n*-hexacosane as major compound,  $\beta$ -sitosterol, biochanin A, sitosteryl 3-*O*- $\beta$ -D-glucopyranoside (**1**), elasticamide (**2**), elastiquinone (**3**) and ficososide B (**4**) were purified and characterized. Antimicrobial activities, expressed as minimum inhibitory concentration (MIC), indicated that the methanol extract showed MIC of 39.1  $\mu$ g/mL; the lowest values were obtained for **3** and **4**, with MIC as low as 4.9  $\mu$ g/mL, smaller than the values of reference antibiotics (25  $\mu$ g/mL). Furthermore, as most of the studied samples exhibited Minimum Microbicidal Concentration/Minimum Inhibitory Concentration (MMC/MIC) ratios lower than 4, a microbicidal effect was clearly exhibited. The overall results provided evidence that the wood of *F. elastica* aerial roots, as well as some of its isolated components might be potential sources of new antimicrobial drugs.

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## 1. Introduction

Infectious diseases are one of the major causes of death worldwide with almost one third of all deaths in low-income countries (World Health Organisation, 2012). They remain a serious health problem worldwide, especially with the growing phenomenon of antibiotic resistance. It has been reported that about 80% of the world population is dependent (fully or partially) on plant-based drugs (Veeresham, 2012). This situation is more complicated in Africa as the population is relatively poorer and relies on medicinal plants to cure several ailments. Despite the discovery of many drugs from natural origin, the search for new antimicrobial agents is still necessary. Such researches could increase the number of drugs which are less toxic and more effective. For instance, in the years from 1981 to 2014, of the 112 antibacterial agents (small molecules) approved, ~73% are natural products

(unmodified structures) or natural product derivatives (Newman and Cragg, 2016).

The Moraceae plant family includes *Ficus* as one of the main plants with biological activities already described such as antiplasmodial (Muregi et al., 2003), antioxidant (Phan et al., 2012), anticancer (Mbosso et al., 2012, 2015, 2016a, 2016b), antimicrobial (Mbosso et al., 2012, 2015, 2016b), antiulcer (Galati et al., 2001), antidiarrhoeal (Mandal and Kumar, 2002), anti-pyretic (Rao et al., 2002), and gastroprotective (Rao et al., 2008). Note that the latex of some species of *Ficus* is exploited in traditional folk medicine for its antihelmintic activity in South and Central America (De-Amorin et al., 1999). The parasitocidal property of this plant has been ascribed to the presence of ficin (Pistelli et al., 2000) and it was also demonstrated that the latex of *Ficus elastica* Roxb. ex Hornem. (Moraceae) showed a significant antischistosomal activity (Seif el-Din et al., 2014).

In our continuous search for bioactive compounds in Cameroonian plants of the family Moraceae, we focused herein on *F. elastica* (the rubber tree) which is a widely-spread evergreen tree up to 30 m tall. Its leaf extract is used for treating skin infections and allergies, as well as a diuretic agent (Phan et al., 2012). The phytochemical investigation

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of this taxon revealed the presence of 6,10,14-trimethyl-2-pentadecanone (25.9%) and geranyl acetone (9.9%) in the leaf oil as main constituents (Ogunwande et al., 2011); emodin, sucrose, morin and rutin with antimicrobial activities from the MeOH extract of the leaves (Hassan et al., 2003), feroxidin, quercitrin, kaempferin, myricitrin, syringing, citroside B, corchoionoside C, (6S,9R)-roseoside, oleanolic acid, ursolic acid, benzyl *O*- $\beta$ -D-glucopyranoside, icariside F<sub>2</sub>, fucuselastin acid and (1'S,6'R)-8-*O*- $\beta$ -D-glucopyranosyl sodium abscisate with antioxidant activity from leaves (Phan et al., 2012), n-alkanes, friedelin, friedelinol, linear aliphatic primary alcohols, linear fatty, phytosterols, betulinic acid, ursolic acid, sitosteryl 3-*O*- $\beta$ -D-glucopyranoside, fucusamide, fucusoside and elasticoside with anticancer and antimicrobial activities from the bark of aerial roots (Mbosso et al., 2012). In our previous studies on the Cameroonian *F. elastica*, Fucusoside B endowed with anticancer activity was isolated from the bark of aerial roots (Mbosso et al., 2016a).

In the present study, we examine the antimicrobial activity of isolated molecules as well as the MeOH extract of wood of *F. elastica* aerial roots. To the best of our knowledge, the antimicrobial evaluation of the wood of *F. elastica* aerial roots is reported here for the first time against pathogenic bacteria and yeast strains.

## 2. Materials and methods

Cyclohexane (99 + %), ethyl acetate (99 + %) and MeOH (99 + %) were purchased from Chemlab and DMSO (99.7%) was from Acros.

### 2.1. Plant material

The wood of *F. elastica* aerial roots was collected from Yaoundé (Cameroon) in December 2007. The plant's identification was established by a member of the National Herbarium of Cameroon (NHC), where a voucher specimen (No. 65646 HNC) was deposited. After Air-drying, the plant material (wood of *F. elastica* aerial roots) was crushed into a fine powder by using an electric grinder.

### 2.2. Extraction and isolation

Macerate of the dried aliquot (5.50 kg) was obtained using methanol (20 L) twice for 48 h at room temperature ( $27 \pm 2$  °C) (Mohamad et al.,

2011). After filtration (Whatman Number One) and evaporation at low pressure in a rotary evaporator (bath at 40 °C), 15 g of extract was obtained (Mbosso et al., 2016a). The crude MeOH extract FBr (14.5 g) was subjected to silica gel column chromatography (CC) (cyclohexane/EtOAc/MeOH gradient of increasing polarity) to afford four fractions on the basis of TLC composition. Further purification through successive column chromatography yielded several pure molecules belonging to many classes of compounds. Products previously isolated from wood of *F. elastica* aerial roots included a mixture of linear aliphatic alkanes with *n*-hexacosane as major compound,  $\beta$ -sitosterol, biochanin A, sitosteryl 3-*O*- $\beta$ -D-glucopyranoside (**1**), elasticamide (**2**), elastiquinone (**3**) and fucusoside B (**4**) (Mbosso et al., 2016a). Molecules **1–4** were tested in the present study (Table 1).

### 2.3. Antimicrobial assay using microbroth dilution method

Six microorganisms, i.e. one Gram-positive (*Staphylococcus aureus*); four Gram-negative bacteria specimens (*Escherichia coli*, *Proteus vulgaris*, *Providencia stuartii*, *Pseudomonas aeruginosa*) and one yeast (*Candida albicans*) were tested. All these microbial strains were obtained from urine samples and vaginal swabs of in-patients suffering from urogenital tract infections at the Tiko Cottage Hospital (Cameroon). Microbial strains were identified through their growing properties using specific culture media, as well as through microscopic and biochemical characteristics described as follows. Gram positive (*S. aureus* RN4220) and Gram negative (*E. coli* JM109) reference strains were submitted to the same microscopic and biochemical characteristic procedure) as controls. We were convinced of the species determination of isolates only if those of the reference samples were confirmatory. Concerning *Candida* species: The sample was cultured on Sabouraud chloramphenicol agar, and the isolate was tested for the evidence of production of germ tube in human serum, a test which differentiates *Candida albicans* from other candida species. Fresh plates of test bacteria species were made from the isolated cultures obtained on agar slants. Discrete colonies of fresh cultures of the different isolates were then picked and suspended in 5 ml of Nutrient broth (NB, Oxoid), in well-labeled sterile bottles, and incubated for 24 h at 37 °C prior to antimicrobial susceptibility testing. Fluconazole (Sigma, USA), and gentamycin (Sigma, USA) were used as reference antibiotics against yeasts and bacteria species, respectively.

**Table 1**  
Chemical structure of isolated compounds from wood of *Ficus elastica* Roxb. ex Hornem. aerial roots.

Name	Structure	Reference
Sitosteryl 3- <i>O</i> - $\beta$ -D-glucopyranoside		
Elasticamide		
Elastiquinone		Mbosso et al. (2016a)
Fucusoside B		

The Minimum Inhibitory Concentration (MIC), considered as the lowest concentration of the sample that inhibits the visible growth of microorganisms, was determined by the microbroth dilution method (Carbannelle et al., 1987; Berghe and Vlietinck, 1991) in Mueller Hinton (for antibacterial activity) or Sabouraud broth (for anticandidal activity) supplemented with 10% glucose and 0.5% phenol red. For susceptibility testing, in a first step supplemented Mueller Hinton or Sabouraud broth (50 µL) was distributed from the first to the twelfth row on a 96 well microplate. The dry extract was initially dissolved in DMSO (20%) (100 µL) and subsequently in Mueller Hinton broth, to reach a final concentration of 10.0 mg/mL for FEBr and for **1**, **2**, **3** and **4** (the extract nomenclature is detailed on footnotes of Table 2). Solutions (50 µL) were added to the first well of each microtiter line. Successive dilutions were then carried out by transferring the mixture/solution (50 µL) from the first to the eleventh well. An aliquot (50 µL) was discarded from this eleventh well. The twelfth well served as growth control since no sample (extract, compounds, or reference antibiotics) was added. A microbial suspension (of isolates as well as of reference strains), i.e. 50 µL, 10<sup>5</sup> colony forming units/mL (CFU/mL), obtained from an overnight growth at 37 °C was added to each well. The final concentration of the FEBr extract and compounds **1**, **2**, **3** and **4** adopted to evaluate the antimicrobial activity ranged from 5000.0 to 2.4 µg/mL. Tests were incubated aerobically at 37 °C for 24 and 48 h, for bacteria and fungi cultures, respectively. The MIC was considered as the lowest concentration of the sample that prevented visible growth or changed in color from red to yellow due to the formation of acidic metabolites corresponding to microbial growth.

The minimum microbicidal concentration (MMC), considered as the lowest concentration of agent capable of causing the deaths of at least 99.99% of an inoculum was also determined. After the MIC determination, an aliquot (10 µL) from each microwell presenting no visible growth was inoculated on fresh drug-free Mueller Hinton agar (for bacteria cultures) and Sabouraud agar (for fungi) plates and incubated at 37 °C for 24 h. Plates showing no growth indicated bactericidal effect of the fraction (sensitive) (Berghe and Vlietinck, 1991).

### 3. Results and discussion

Compounds tested in this study included steroidal glucosides known as sitosterol 3-O-β-D-glucopyranoside (**1**), elasticamide (**2**), elastiquinone (**3**) and ficoside B (**4**).

They were isolated and characterized from wood of *F. elastica* aerial roots and subsequently, a notable anti-proliferative effect on 6 human cancer cell lines was highlighted (Mbosso et al., 2016a). It is noteworthy that the antimicrobial activity of β-sitosterol (Jain et al., 2001) and biochanin A was already investigated elsewhere (Sklenickova et al., 2010; Liu et al., 2011) and consequently these molecules were not tested in the present study. Compounds **1–4** as well as the crude extract from the wood of *F. elastica* aerial roots were tested for their antimicrobial activities on a panel of microbial strains and the results are reported in Table 2.

In the literature, various criteria are applied to determine the susceptibility of extracts and isolated compounds as microbial inhibitors. In our case, the antimicrobial activity of a crude plant extract has been defined as significant with MIC below 100 µg/mL, moderate with MIC between 100 µg/mL and 625 µg/mL, and low with MIC values more than 625 µg/mL (Kuate, 2010; Bueno, 2012; Siwe Noundou et al., 2016). The methanol extract from wood of *F. elastica* aerial roots inhibited the growth of tested microorganisms, with MIC value of 39.1 µg/mL except against *P. stuartii* (Table 2). The established antibiotic drugs gentamycin and fluconazole displayed MICs of 25 µg/mL on tested microorganisms. Therefore, the methanol extract from wood of *F. elastica* aerial roots could be considered as a promising herbal drug, as MIC values below 100 µg/mL were obtained against 5 out of 6 of the tested microorganisms. Interestingly, this crude extract showed also a strong activity against six human cancer cell lines (Mbosso et al., 2016a). The isolated

**Table 2**  
Antimicrobial activities (Gram-positive bacteria, Gram-negative bacteria and yeast) of total wood extract of *F. elastica* aerial roots.

Concentration (µg/mL)	Gram (-) bacteria						Gram (+) bacteria						yeast			
	<i>E. coli</i>		<i>P. vulgaris</i>		<i>P. stuartii</i>		<i>P. aeruginosa</i>		<i>S. aureus</i>		<i>C. albicans</i>		MIC <sup>c</sup>	MMC/MIC		
	MIC <sup>c</sup>	MMC	MIC <sup>c</sup>	MMC	MIC <sup>c</sup>	MMC	MIC <sup>c</sup>	MMC	MIC <sup>c</sup>	MMC	MIC <sup>c</sup>	MMC				
<b>1</b>	78.1	156.2	2	2	78.1	156.2	2	2	39.1	78.1	78.1	156.2	2	78.1	156.2	2
<b>2</b>	39.1	39.1	1	1	39.1	78.1	2	2	78.1	156.2	78.1	156.2	2	19.5	39.1	2
<b>3</b>	19.5	39.1	2	2	4.9	19.5	4	4	4.9	19.5	4.9	19.5	8	19.5	39.1	2
<b>4</b>	4.9	19.5	4	4	4.9	19.5	1	1	39.1	78.1	4.9	19.5	4	4.9	19.5	4
<b>FEBr<sup>a</sup></b>	39.1	78.1	2	2	1250.0	2500.0	2	2	39.1	78.1	39.1	78.1	2	39.1	78.1	2
Gentamycin <sup>b</sup>	25	50	2	2	25	50	2	2	25	50	25	50	2	/	/	/
Fluconazol <sup>b</sup>	/	/	/	/	/	/	/	/	/	/	/	/	/	2.5	50	2

<sup>a</sup> (FEBr) MeOH crude extract wood of *Ficus elastica* aerial roots;

<sup>b</sup> Reference antibiotics (fluconazol for yeast and gentamycin for bacteria).

<sup>c</sup> MIC is considered as the lowest concentration of the sample, that inhibits the visible growth of a microbe and MMC is considered as the lowest concentration of the sample capable of causing the death of at least 99.99% of a tested inoculum.

compounds **1–4** were further tested for their antimicrobial properties. The defined threshold values for each molecule are defined as follows: MIC below 10 µg/mL (significant activity),  $10 \leq \text{MIC} \leq 100$  µg/mL (moderate activity) and  $\text{MIC} > 100$  µg/mL (low activity) (Kuetze, 2010). Results summarized in Table 2 indicated that MICs ranged from 4.9 to 78.1 µg/mL.

More specifically, elasticamide **2** inhibited the growth of 100% of the six tested strains with MICs spanning 19.5–78.1 µg/mL and can be considered as a moderate antibacterial agent (Kuetze, 2010). In a previous report, we isolated fucosamide from the bark of *F. elastica* aerial roots. This ceramide revealed a significant activity with MIC of 3 µg/mL against *Staphylococcus saprophyticus* and low bactericidal properties against *Klebsiella pneumoniae* (MIC 380 µg/mL), *Escherichia coli* and *Enterococcus faecalis* (MIC 190 µg/mL). Note that MICs values obtained for reference gentamycin were of 980 µg/mL against *Staphylococcus saprophyticus* and *Enterococcus faecalis*; 7810 µg/mL against *Klebsiella pneumoniae* and *Escherichia coli* (Mbossso et al., 2012). Similar results were previously mentioned for structurally closed skeletons (Fischer et al., 2012; Poumale, 2012).

Elastiquinone **3** inhibited the growth the six tested microorganisms with the MICs in the range of 4.9–78.1 µg/mL and demonstrated a significant activity with a MIC of 4.9 µg/mL against *P. stuartii* and *P. aeruginosa*, 9.8 µg/mL against *P. vulgaris* and *S. aureus* and a moderate activity (MIC of 19.5 µg/mL) against *E. coli* and *C. albicans*. Results in the same order were previously obtained for similar agents (Manojlovic et al., 2000; Comini et al., 2011).

Fucoside B **4** also prevented the growth of the six strains with the same MICs range found for elastiquinone **3**. However, compound **4** was the most active against *E. coli*, *P. vulgaris*, *S. aureus* and *C. albicans* with a MIC of 4.9 µg/mL. These values were even smaller than MIC of used reference antibiotics (25 µg/mL). Molecules **3** and **4** can be considered as significant antibacterial agents as MIC values below 10 µg/mL were obtained against 4/6 of the tested microorganisms (Kuetze et al., 2010). In a previous study, we isolated spathoside from the stem bark of *Spathodea campanulata* P. Beauv., a cerebroside which showed a significant activity against *K. pneumoniae* (MIC 6.25 µg/mL), but moderate activity against *Staphylococcus aureus*, *Streptococcus faecalis*, *P. aeruginosa*, *Shigella flexneri* (MIC 12.5 µg/mL), *Bacillus subtilis* (MIC 25 µg/mL) and *Bacillus cereus*. However, no sign of growth inhibition was seen for *E. coli* and *Shigella dysenteriae* (Mbossso et al., 2008). In a previous study, we separated an elasticoside from the bark of *Ficus elastica* aerial roots, another cerebroside which showed a moderate activity against *E. faecalis* (MIC 30 µg/mL), but low activity against many microorganisms such as *Staphylococcus saprophyticus* (MIC 130 µg/mL), *K. pneumoniae* (MIC 250 µg/mL), *Trichophyton rubrum*, *C. albicans*, *E. coli*, *Salmonella typhimurium*, *S. aureus* and *Staphylococcus epidermidis* (MIC 500 µg/mL) (Mbossso et al., 2012). Other cerebrosides have also been reported to display antimicrobial activities (Catani et al., 2003; Chen et al., 2003; Shu et al., 2004).

Sitosteryl 3-O-β-D-glucopyranoside **1** inhibited the growth of the five tested bacteria with the MICs ranged from 19.5 to 78.1 µg/mL. Similar results were previously described for sugar derivative **1** which has already demonstrated antibacterial activities against *S. aureus* (Soo-Hwan et al., 2003; Phan et al., 2005), *S. aureus*, *S. flexneri* (MIC 12.5 µg/mL), *S. faecalis*, *P. aeruginosa*, *S. dysenteriae* (MIC 25 µg/mL) (Mbossso et al., 2010), *S. aureus* (MIC 12.5 µg/mL), *S. faecalis* (MIC 6.3 µg/mL), *P. aeruginosa* (MIC 3.2 µg/mL) and *S. flexneri* (MIC 12.5 µg/mL) (Mbossso et al., 2008). Compound **1** also showed a moderate antifungal activity against *C. albicans* (MIC 78.1 µg/mL) which is of the same order as other studies found in the literature: *Candida tropicalis* (MIC 100 µg/mL), *Cryptococcus neoformans* (MIC 50 µg/mL) (Tamokou et al., 2011), and *C. albicans*, *Cryptococcus neoformans*, *Aspergillus fumigatus* (MIC 125 µg/mL) (Awouafack et al., 2013).

Molecules **2–3** demonstrated a more pronounced antifungal activity (MICs 19.5 µg/mL) while Fucoside B **4** possessed a significant antifungal activity with a MIC of 4.9 µg/mL against *C. albicans*.

Data in Table 2 indicated that most MMC/MIC ratios for the crude extract were below 4, signifying the microbicidal effects on the microorganisms (Carbonnelle et al., 1987; Mbaveng et al., 2008, 2011). A keen look at the MICs and MMCs indicated that compounds **1** and **2** are bactericidal against all tested microorganisms. However, compounds **3** and **4** rather exerted bacteriostatic (MMC/MIC  $\geq 4$ ) effects (Mbaveng et al., 2008, 2011) on 50% and 66.66% of the tested microorganisms, respectively. These data suggest that the secondary metabolites of the wood of *F. elastica* aerial roots may interact synergistically to produce the observed effects. Regarding the involvement of microorganisms in treatment failures and the re-emergence of infectious diseases (Blot et al., 2007; Falagas and Bliziotis, 2007; Kuetze, 2010; Kuetze et al., 2011a), the antimicrobial activity of methanol wood extract *F. elastica* aerial roots as well as that of compounds **1–4** could be considered promising. To the best of our knowledge, the antimicrobial effect of **2–4** against *Escherichia coli*, *Proteus vulgaris*, *Providencia stuartii*, *Pseudomonas aeruginosa*, *Staphylococcus aureus*, *Candida albicans* is being reported for the first time. Data reported herein, therefore, provide additional information on the potential of various parts of this plant to fight microorganisms and more particularly, elastiquinone **3** and fucoside B **4** as the main active antimicrobial agents of the wood of *Ficus elastica* aerial roots extract.

#### 4. Conclusions

The methanol extract showed a good inhibition with the lowest MIC value (39.1 µg/mL) on the entire studied organisms except for *P. stuartii*. The most active molecules were elastiquinone **3** and fucoside B **4** whose MIC against *P. stuartii*, *P. aeruginosa* and *E. coli*, *P. vulgaris*, *S. aureus*, *C. albicans*, respectively, was 4.9 µg/mL. The results of the present study are important, taking into account the implication of the studied microorganisms in therapeutic failure. The results of our study are consistent with those reported for other *Ficus* species such as methanol extract from the stem bark of *F. ovata* (Kuetze et al., 2009), methanol extract from the roots of *F. polita* (Kuetze et al., 2011b),  $\text{CHCl}_3/\text{MeOH}$  1:1 crude extract of bark of *F. elastica* aerial roots (Mbossso et al., 2012) and methanol extract from fruits of *F. Bubu* (Mbossso et al., 2016b). These data indicate that the methanol wood extracts of *F. elastica* aerial roots as well as some of its constituents, and mostly elastiquinone and fucoside B deserve more attention in the future development of potential antimicrobial drugs to fight MDR bacterial and yeast infections.

#### Competing interest

The authors declare no conflict of interest.

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#### Appendix A. Supplementary data

Supplementary data to this article can be found online at <http://dx.doi.org/10.1016/j.sajb.2017.03.026>.

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