

Full Length Research

Antibacterial activity, phytochemicals and essential oils in ethyl acetate extract of *Cleistopholis patens*

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Accepted 27 January, 2020; Published 20 February, 2020

The ethyl acetate extract of *Cleistopholis patens* was investigated for its antibacterial activity against ten bacterial strains and compared with standard antibiotics. The extract was purified and antibacterial activities were compared with the crude extract. The multiple antibiotic resistant (MAR) index was determined using known method. The phytochemical components were examined and the essential oil components were also investigated using the GC-MS spectra analysis. The antibacterial test revealed the plant to be active against the test organisms. However, the crude extract showed larger zones of inhibition compared to the purified extract. The multiple antibiotic resistances (MAR) index ranged between 0.42 and 1.0. The phytochemical analysis revealed the presence of glycosides, steroids, phenols and tannin. Mineral content of plants are Na (24.03%), K (30.14%), Ca (32.31%), Mg (26.09%), Zn (28.09%), Fe (6.70%) and Cu (0.03%). The GC-MS analysis showed the presence of essential oils predominantly fatty acids, sterols, esters, terpenes and alcohols with fatty acid occurring in abundance. The presence of essential oil in this plant could be responsible for its biological activities and the result of this work supports the use of the plant in the traditional medicine and herbal treatment of diseases.

Key words: Essential oil, (MAR), *Cleistopholis patens*, phytochemical, standard antibiotics, GC-MS, essential oils.

INTRODUCTION

The search for alternative products and natural phytochemicals isolated from plants used as traditional folkloric medicines is on the increase as so many plants have been studied for activities against pathogens with the aim of developing active drugs against them. Chemical analysis of plant extracts showed the existence of many chemical compounds related to different classes such as alkaloids, quinines, sugar alcohols, terpenes, polyphenols, flavonoids, phenolics and many others. Flavonoids and phenolics and their derivatives are found in most plants. Such compounds are very important for both humans and plants (Rafeal *et al.*, 2012) and are

often referred to as essential oils.

Essential oils are complex systems of volatile components and have gained special attention among secondary metabolites because many of them are used as raw materials for the production of perfumes, cosmetics, pharmaceuticals and pesticides. In addition they are exploited in aromatherapy and in phyto-therapy. Furthermore they are used as spices and for nutrition (Alviano and Alviano, 2009). Volatile oils also have been the sources of natural products and used for many medical products. They are plant secondary metabolites that are known for their fragrance and food flavor properties. They consist of a complex mixture of mono and sesquiterpenes, phenyl propanoids and oxygenated compounds (Koudou *et al.*, 2005).

Essential oils are complex mixtures of volatile (around

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100 u) to semi-volatile compounds (around 300 u), usually with a strong odour, rarely colored, soluble in organic solvents but insoluble in water. They comprise volatile compounds of terpenoid and non-terpenoid origin, synthesized through different biosynthetic routes and with distinct primary metabolic precursors. Monoterpenes and sesquiterpenes are usually the main group of compounds found in essential oils. In addition, phenylpropanoids are also very frequent. Moreover, some essential oils may also contain fatty acids and their esters and more rarely nitrogen and sulfur derivatives (Krishnaveni *et al.*, 2014).

The active principles of many drugs found in plants are secondary metabolites. There is therefore, a vital need to investigate plant phytochemical for their major constituents. *Cleistopholis patens* (Benth) Engl and Diels belongs to the family Annonaceae. It is widely distributed in Senegal, Nigeria and Uganda. It is sometimes used as food preservatives (Burkil, 2004). The long narrow leaves held in one plane on slightly drooping branches give this tree a distinctive appearance. The leaves are shiny on their upper surface when fresh. Decoction of the bark is taken to treat stomach ache, diarrhea, tuberculosis, bronchitis and hepatitis. Bark pulp is applied against swelling, oedema, with low and bark sap is dropped into the nose to treat headache and rubbed in to treat rickets in children. In Uganda, the bark is used in preparation to treat malaria and measles. In Nigeria, the bark is used to treat typhoid fever and menstrual irregularities (Quattara, 2013). The root bark is used as vermifuge, leaf infusion or decoction is administered against hepatitis, fever, trypanosomiasis, and rheumatic arthritis (Atuhe, 2010).

Several Authors have identified some volatile oils from this plant to include (E)-beta-ocimene linalool, (E)- and (Z)-linalool oxides, myrcene, P-cymene, and germacrene D (Ekundayo *et al.*, 1988), α -copaene, δ -cadinene, and germacrene D, β -caryophyllene, germacrene D, and germacrene B (Boyom *et al.*, 2011), β -pinene, sabinene, (E)- β -caryophyllene, (E)- β -ocimene, germacrene D, α -pinene and germacrene B, (Quattara *et al.*, 2013). The aim of this study is to challenge the ethyl acetate extract of the stem bark against ten human pathogenic bacteria and also to identify the essential oils isolated from *C. patens*.

MATERIALS AND METHODS

Collection and identification of plant materials

Stem bark of *C. patens* was collected from private garden of a traditional healer at Ibode in Ibadan, Oyo State. The plants were identified by Mr. F.O. Omotayo of the Department of Plant Science and Biotechnology, Ekiti State University, Ekiti State. Voucher specimens with

voucher numbers WHAE2017/001 was deposited in the herbarium of the University. Plant nomenclature and uses are depicted in table 2. The plant materials were washed with distilled water, air dried at room temperature, milled into powder and kept in sterile containers until use.

Preparation of crude extracts

The ethyl acetate extracts was prepared by dissolving 200gm of powdered plant samples in 500mls solvent. The suspension was allowed to stand for 5 days (120 h) at of 32°C and then filtered using a sterile muslin cloth and a Whatmann No 1 filter. The collected filtrate was evaporated to dryness under sterile condition at room temperature after which the extracts were kept in sterile bottles until ready for use (Table 1).

Collection and preparation of microorganisms

The clinical strains of the test bacteria were obtained from the Department of Microbiology, Obafemi Awolowo University, Ile-ife, Osun State. The organisms were collected and maintained by regular sub-culturing on nutrient agar. The test organisms were subjected to biochemical test to confirm the authenticity of the organisms.

Sterilization and reconstitution of crude extract

A five gram portion of the crude extract was reconstituted in 50mL of 25% dimethyl sulfoxide, the solution was sterilized by passing them through milipore membrane filter with pore size 0.45 μ m. The filtrate was then stored in sterile bottles until use.

Standardization of inocula

One percent (1%) of solution of sulphuric acid was prepared and mixed properly. Also, 1% solution of barium chloride was prepared by dissolving 0.5 g of dehydrated barium chloride (BaCl₂.H₂O) in 50ml of distilled water. A 0.5ml aliquot of barium chloride solution was added to 99.5ml of the sulphuric acid solution and it was the mixed together. The solution was transferred into a capped tube of the same type used for both the control and the test inocula. The solution was kept at temperature of +4°C (Cheesebrough, 2000).

Table 1. Plants nomenclature and traditional uses.

English name	Botanical name	Local name	Family	Part used	Traditional uses
Salt and oil tree/ heartwood	<i>Cleistopholia patens</i>	Yor; Apako Ibo;Ojo	Anonaceae	Stem bark	Cough, antelmintics, respiratory disease, TB, purgative

Antibacterial assay of the plant extracts using the agar well diffusion method

Test organisms were suspended in Nutrient broth and incubated for 4 h to obtain a concentration corresponding to McFarlands constant (0.5×10^8 cfu/ml). The inoculums were standardized with the prepared barium sulphate. Sterile Petri dishes were inoculated by the pour plate method. One ml (1 ml) of the test inoculums was pipette aseptically into each Petri dish and about 20 ml of sterilized nutrient agar was poured into the inoculated Petri dish.

The agar plates were allowed to set. Wells of 6 mm diameter were made over the agar plates equidistant from each other using sterile cork borer and 0.5 ml of each plant extracts of different concentrations as prepared by serial dilution were added to the wells using a micropipette. The extracts were allowed to diffuse into the agar for about 20 min after which the plates were incubated for 24 h at 37°C. Thereafter, the diameter of inhibition zones formed around each well was measured in mm and recorded. The experiments were carried out in triplicates and the average values recorded.

Determination of Minimum Inhibitory Concentration (MIC) of plant extracts against test organisms.

A modified method of Weigand *et al.* (2008) was adopted in the determination of MIC. The MIC of the extract was determined by diluting the various concentrations with nutrient broth. A 1ml aliquot of a serial dilution of 100, 60, 40, 20 and 10 mg/ml of the extracts was separately added to test tubes containing specifically 0.1ml of standardized inoculums of 1 to 2×10^7 cfu/ml. The tubes were incubated aerobically at 37°C for 18-24 h. Two control tubes were prepared for each test batch. This is as follows: tube containing extracts and the growth medium without inoculums (antibiotic control) and the tube containing the growth medium and the inoculums (organism control). The inocula were then plated and inoculated at 37°C for 24 h. The MIC was determined as the lowest concentration of the extracts exhibiting no visible growth (no turbidity) when compared with the control tubes.

Determination of antibiotics sensitivity

Susceptibility testing using standard antibiotics were carried out using the disk diffusion method of Kirby-Bauer *et al.* (1966). Standard antibiotics used include; streptomycin (25µg), gentamycin (10 µg), erythromycin (5 µg), amoxilin (25 µg), ofloxacin (5 µg), chloramphenicol (10 µg), ceftriazone (30 µg), pefloxacin (5 µg), cotrimoxazole (25 µg), ciprofloxacin (10 µg), augmentin (30 µg), nitrofurantoin (200 µg) and tetracycline (30 µg).

Nutrient agar plates were prepared and seeded with bacterial inoculum already standardized with the Macfarland's constant using the pour plate method. After gelling, antibiotic disk were placed on the gelled plate and the plates were incubated at 37°C for 24 h after which the zones of inhibition were observed and recorded.

Multiple resistance index

The MAR index for the resistant bacterial strains was determined according to the procedure described by Krumpelman (1983). This is essentially to determine the degree of bacterial resistance to antibiotics. The indices were determined by dividing the number of antibiotics to which the organism was resistant to by the number of the antibiotics tested. Resistance to three or more antibiotics is taken as MAR and MAR greater than 0.2 indicates a high risk of multiple antibiotic resistant source of contamination.

Phytochemical screening of plant material

The plant was screened qualitatively for phytochemicals using the methods of Odebiyi and Sofowora (2005); Evans (2009) and Bansa and Ngbede (2006).

Test for alkaloids

A 0.2 g amount of plant extract was acidified with 1% hydrochloric acid (HCl) for 2 min and was then treated with a few drops of Dragendorff's reagent in a test tube. The formation of white precipitate indicates the presence.

of alkaloids.

Test for saponins

Sterile distilled water was used to dissolve 0.2g of plant extract. A 2ml amount of the solution was placed in different test tubes and was shaken vigorously for a few minutes. Frothing which persists on warming was taken as an evidence of the presence of saponin.

Test for tannins (Gelatin test)

To the extract, 1% gelatin solution containing sodium chloride was added. Formation of white precipitate indicates the presence of tannins.

Test for flavonoids (Shinoda's Tests)

Plant extract was dissolved in 2 ml of dilute NaOH. A yellow solution that turns faint or colourless on addition of a few drops of hydrochloric acid and a change in colour while standing indicates the presence of flavonoids.

Test for cardiac glycosides (Lieberman's Test)

The Lieberman's test was used to determine the presence of cardiac glycosides. A 5 g amount of plant extract was dissolved in 20ml of acetic anhydride and cooled with ice. Concentrated H_2SO_4 was then carefully added. A colour change from violet to blue and then to green indicated the presence of a steroidal nucleus (a glycone portion of the cardiac glycoside).

Test for steroids (Salkowski test)

A 0.5 g portion of plant extract was dissolved in 2 ml of chloroform and 0.2 ml of concentrated H_2SO_4 was carefully added to form a layer. A reddish –brown colour ring at the interface between the layers indicated the deoxy- sugar characteristic of cadenolides which indicated the presence of steroids.

Determination of anti-nutrients in plant samples

Preparation of fat free samples

A 2g amount of each sample were defatted with 100 ml of diethyl ether using a soxhlet apparatus for 2h, and then

used for the determination of anti-nutrients. Alkaloids and cyanogenic glycosides were determined by the gravimetric precipitation method as described by (Harbone, 1984). Flavonoids and anthocyanins were determined following the method of Boham and Kocipai-Abyazan (1994). While tannin was determined by the Folin-Dennis spectrophotometric method as described by Makka (1989). Spectrophotometric methods as described by Obadoni and Ochuko (2002) were used to determine the saponin content.

Determination of mineral content of plants samples

Mineral was estimated by the use of an Atomic Absorption Spectrophotometer. The sample solutions in the sample bottles were analyzed for the concentration of the individual elements. Each element has specific cathode discharge lamp and this lamp was used to determine a particular element. Discharge lamp emits radiation at a wavelength specific for each element being assayed. This specificity can be obtained only from a pure sample of the element that is excited electrically to produce an arc spectrum on that element.

Bioassay guided fractionation of plant extracts (purification).

Fractionation of the crude extract of *C. patens* was carried out using the methods of Cosa *et al.* (2006).

Purification of crude extracts

The ethyl acetate fractions were allowed to run through a short column and the fractions were collected, and the thin layer chromatography was carried out to ascertain the purity of the compounds.

Antibacterial assay of purified fractions

Fractions from column chromatography were subjected to bioassay to determine the active fraction using the disk diffusion method of Kirby- Bauer *et al.* (1966) as described.

Gas Chromatography and Mass spectra (GC-MS) analysis of purified fractions

Ethyl acetate extracts of Stem bark of *C. patens* were analyzed with the help of GC-MS analyzer (Perkin Elmer

Gas Chromatography- Mass Spectrum). On Elite-1 column the data was generated. The carrier gas helium (99.999%) was used at flow rate of 1 ml per min in split mode (10:1). 8 μ of sample was injected to column at 250°C injector temperature. Temperature of oven starts at 60°C and hold for 6 min and then it was raised at rate of 10°C per min to 300°C without holding. Holding was allowed for 6 min at program rate of 5°C per min. Temperature of ion sources was maintained at 240°C. The injector temperature was set at 250°C and detector temperature was set at 260°C. The mass Spectrum of compounds present in samples was obtained by electron ionization at 70eV and detector operates in scan mode 50 to 600Da atomic units. A 0.5 s of scan interval and fragments from 50 to 600Da was maintained. Total running time was 40 min.

RESULTS

Table 2 shows the antibacterial activity of extract of *C. patens* against test bacteria in comparison with standard antibiotics. The M100 of the CLSI (2017) was used to interpret the antibiotic test result. *Pseudomonas aeruginosa* was resistant to the extract of *C. patens*, and some of the standard antibiotics including; augmentin, ciprofloxacin, nitrofurantoin, gentamycin cotrimoxazole and amoxillin while it was sensitive to ofloxacin with a zone of inhibition of 18mm, ciprofloxacin (12 mm), tetracycline (16mm) and pefloxacin (12 mm).

Salmonella typhi was sensitive to the plant extract with zones of inhibition of 14mm but resistant to the standard antibiotics used. *Escherichia coli* was resistant to the plant extract but sensitive to gentamycin (30mm), cotrimoxazole (22 mm) and pefloxacin (24 mm).

Proteus vulgaris was sensitive to the plant extract with a zone of inhibition of 22 mm but resistant to all antibiotic disks except pefloxacin (24 mm). *Streptococcus pyogenes* was sensitive to the plant extract with a zone of inhibition of 18mm, Nitrofurantoin (14 mm), gentamycin (14 mm), cotrimoxazole (14 mm). *Klebsiella aerogenes* was sensitive to the plant extract but resistant to all antibiotics disks used. *Staphylococcus aureus* was sensitive to the plant extract (14 mm), gentamycin (28 mm) and ofloxacin (28 mm).

Escherichia coli was resistant to plant extract and some of the antibiotics but sensitive to gentamycin (30mm), cotrimoxazole (22 mm) and pefloxacin (24 mm). *Streptococcus pyogenes* was sensitive to the test extract (18 mm), nitrofurantoin (14 mm), gentamycin (14 mm) and cotrimoxazole (14 mm).

Staphylococcus aureus was sensitive to the plant extract (14 mm), gentamycin (28 mm) and ofloxacin (28 mm). *Serratia marscescens* was sensitive to the plant extract

(16 mm), gentamycin (24), ofloxacin (28 mm), ciprofloxacin (28 mm) tetracyclin (12 mm) and pefloxacin (14 mm).

Yersinia enterocolitica was sensitive to the plant extract (16mm) and cotrimoxazole (16 mm) while *Shigella dysenteriae* was resistant to all antimicrobials including the plant extract used.

MAR

The MAR (multiple antibiotic resistance index) of each test bacteria is as listed in Table 3. All test bacteria had MAR index greater than 0.2 indicating that all test organisms have a multiple drug resistant factor.

Minimum inhibitory concentrations of *C. patens* against test organisms

The MIC values of the plant extracts against test organisms are represented in Figure 1. The minimum inhibitory concentration of ethyl acetate extract against *S. aureus* was 40 mg/ml, *Y. enterocolitica* was 40 mg/ml *S. typhi*, *S. pyogenes*, *K. aerogenes* was 20 mg/ml, *P. vulgaris* was 20 mg/ml. other bacteria were resistant to the plant extract.

The antibacterial test result of purified fraction is presented in Table 4. The Cp7 fraction was active against *P. vulgaris* (7 mm), *S. aureus* (6 mm), *S. pyogenes* (6mm), *K. aerogenes* (14 mm), *S. typhi* (6 mm) and *Y. enterocolitica* (8 mm). Fraction Cp₁₂ was active against *P. vulgaris* (12 mm), *S. aureus* (10 mm), *S. pyogenes* (9 mm), *K. aerogenes* (12 mm), *S. typhi* (8 mm) and *Y. enterocolitica* (8mm). Cp 12₃ was active against *P. vulgaris* (8 mm), *S. aureus* (8 mm), *S. pyogenes* (8 mm), *K. aerogenes* (6 mm), *S. typhi* (6 mm), but no activity against *Y. enterocolitica*.

Phytochemical constituents of *C. patens*

The qualitative analysis of *Cleistopholis patens* showed the presence of glycoside, steroid, phenol, tannin and saponin. Alkaloid and anthraquinone were not detected while the quantitative analysis showed Tanin (2.20%), phenol (3.50%), phylate (17.30%), oxalate (3.69%), saponin (13.89%) flavonoid (9.89%). The mineral content of *C. patens* include Na (24.03%), K (30.14%), Ca (32.31%), Mg (26.09%), Zn (28.09%), Fe (6.70%), Cu (0.03%) while lead (Pb) was not detected.

Table 2. Antimicrobial susceptibility of test bacteria.

Organisms/zones of inhibition		Antibiotic disks																										
		C. patens at 100 mg/ml (mm)		AUG		CRO		NIT		GEN		COT		OFL		AMX		CPX		TET		PFN		STR		ERY		CHL
P. aeruginosa		-	R	-	-	R	-	R	-	R	-	R	18	S	-	R	12	S	16	S	12	S	12	S	10	S	-	R
Salmonella typhi		14	S	-	R	-	R	-	R	-	R	-	R	-	R	-	R	-	R	-	R	12	S	-	R	-	R	
Escherichia coli		-	R	-	R	-	R	-	R	30	S	22	S	-	R	-	R	-	R	24	S	12	S	20	S	24	S	
Proteus vulgaris		22	S	-	R	-	R	-	R	-	R	-	R	-	R	-	R	24	S	-	R	-	R	-	R	30	S	
Strept pyogenes		18	S	-	R	-	R	14	S	14	S	14	S	-	R	-	R	-	R	-	R	-	R	-	R	-	R	
K. aerogenes		18	S	-	R	-	R	-	R	-	R	-	R	-	R	-	R	-	R	-	R	-	R	-	R	-	R	
S.aureus		14	S	-	R	-	R	-	R	28	S	-	R	28	S	-	R	-	R	-	R	-	R	-	R	28	S	
Serratia merscences		16	S	-	R	-	R	-	R	24	S	-	R	28	S	-	R	28	R	12	S	14	S	-	R	-	R	
Y. enterocolitica		16	S	-	R	-	R	-	R	-	R	16	S	-	R	-	R	-	R	-	R	14	S	-	R	12	S	
Shigella dysenteriae		-	R	-	R	-	R	-	R	-	R	-	R	-	R	-	R	-	R	-	R	-	R	-	R	-	R	

Legend; Antibiotics; AUG- Augmentin, CRO- Ciprofloxacin, NIT- Nitrofurantoin, GEN- Gentamycin, COT- Cotrimoxazole, OFL- Ofloxacin, AMX-Amoxycillin, CPX.-Ciprofloxacin, TET-Tetracycline, PFN-Pefloxacin. - = No inhibition R--- Resistant, S –Susceptible.

Table 3. Multiple antibiotic resistance index of test bacteria.

Test bacteria	MAR index
<i>Pseudomonas aeruginosa</i>	0.47
<i>Salmonella typhi</i>	0.92
<i>Escherichia coli</i>	0.67
<i>Proteus vulgaris</i>	0.85
<i>Streptococcus pyogenes</i>	0.77
<i>Klebsiella aerogenes</i>	1
<i>Staphylococcus aureus</i>	0.77
<i>Serratia merscences</i>	0.69
<i>Yersinia enterocolitica</i>	0.77
<i>Shigella dysenteriae</i>	1.0

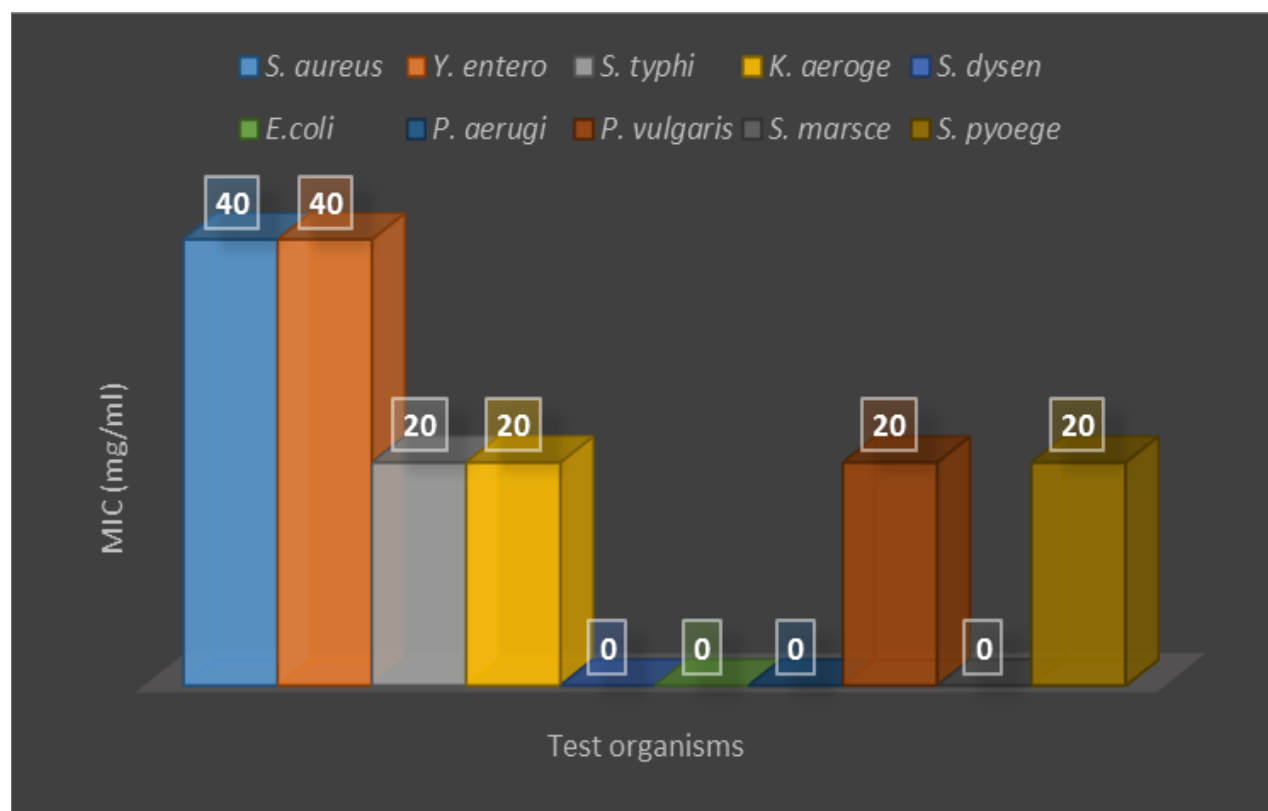


Figure 1. MIC values of plant extract against test organisms.

Table 4. Antibacterial activity of the purified fractions of *Cleistopholis patens*.

Fractions	Colour of fraction	R _f values	Antimicrobial test of purified extracts (organisms and zones of inhibition in mm)											
			<i>P. vulgaris</i>		<i>S. aureus</i>		<i>Str. pyogenes</i>		<i>Kl. aerogenes</i>		<i>Sal. typhi</i>		<i>Yer. enterocolitica</i>	
CP7	Dark brown	0.083	Crd	Pre	crd	pre	crd	pre	crd	pre	crd	Pre	crd	pre
			22	7	14	6	16	6	16	8	14	6	16	8

Legend: crd- Crude extract, pre- purified extract, R_fValue- Retention factor value.

GCMS Spectra of *C. patens* fractions

GCMS analysis of the sample contained 12 compounds that matched NIST library data. The spectra revealed the presence of predominantly α Pinene, Beta- Pinene, Azulene, Terpinolene, oxypurinol, fatty acids and esters. The identified compounds and quantities are listed in Table 5 and the spectra plate is presented in Figure 2.

DISCUSSION

Natural products, such as plant extract, either as pure compounds or as standardized extracts, provide unlimited opportunities for control of microbial growth owing to their chemical diversity. The result of the antibacterial susceptibility of the test bacteria presented a 70% susceptibility to the plant extract whereas a high degree of resistance was recorded for the standard antibiotics tested. Antibiotic resistance has been a major source of public health challenge. The systematic screening of plant-derived bioactive compounds as resistance modifying agents represents a potential approach to overcome bacterial resistance.

The MAR index of bacteria used in this work also showed a high rate of resistance to standard antibiotics. This implied that the test organisms were resistant to the antibiotics used. Multiple antibiotic resistance (MAR) in bacteria is most commonly associated with the presence of plasmids which contain one or more resistance genes, each encoding a single antibiotic resistance phenotype. High prevalence of multidrug resistance indicates a serious need for broad-based, local antimicrobial resistance surveillance and planning of effective interventions to reduce multidrug resistance in such pathogens (Osundiya *et al.*, 2013). Two principal drivers of resistance appear to be inadequate (or inappropriate) empirical antibiotic (Fish and Ohlinger, 2006) therapy and prolonged antibiotic use. The menace of antimicrobial resistance is particularly worrisome in developing countries like ours where there is a high burden of infectious disease with concomitant high rate of poverty which constrains the access to newer, more effective and (Okeke *et al.*, 2005) conversely more expensive antimicrobial agents.

The comparative antibacterial activity of the crude extracts and the purified extract showed to a great extent the synergistic effect of the different phytochemicals that make up the plant. Yuan *et al.* (2017) opined that different components in a prescription exert a synergistic effect in such ways as acting on different targets or improving the solubility of active compounds, which constitute the pharmaceutical basis of traditional medicines. The presence of different phytochemicals with

different modes of action as observed by Rasoanaivo *et al.* (2011) contribute to high efficacy of crude plant materials as compared to purified fractions.

Reduced biological activity of purified fractions have been attributed by several authors to factors such as the poor quality of ethno pharmacological studies, plant material processing, preclinical laboratory protocols which are often very different from local practices, an inadequate fractionation process, degradation of active constituents during fractionation and poor biological models to demonstrate activities (Rasoanaivo *et al.*, 2011).

Pure drugs that are isolated from plants may be chosen for their high activity against a human disease, but they have disadvantages. They rarely have the same degree of activity as the unrefined extract at comparable concentrations or dose of the active component (Wagner and Ulrich-Merzenich .2009). This phenomenon is attributed to the absence of interacting substances present in the extract. Furthermore, many plants contain substances that inhibit multi-drug resistance (MDR). This may explain why crude extracts are often more effective than isolated constituents at an equivalent dose.

The plant *Cleistopholis patens* is rich in phytochemicals such as glycosides, steroids, phenol, tannin and saponin. These phytochemicals are suspected to be responsible for the activities of the plant extract against the array of bacterial tested against it. Essential oils are complex mixtures of volatile, lipophilic and odiferous substances from the secondary metabolism of plants. They are mainly composed of monoterpenes, sesquiterpenes and their oxygenated derivatives (alcohols, aldehydes, esters, ketones, phenols and oxides).

The chemical components of the plants identified in a GC-MS analysis could be responsible for the antibacterial properties of the plant as indicated in the antibacterial test result. The analysis revealed predominantly essential oils, fatty acids, esters and terpenoids. Ouattara *et al.* (2018) in their work isolated several essential oils in the stem bark extract of *C.patens* including patchoulone , β -pinene, bornyl acetate, α -pinene, juvenile hormone III, and β -elemol among others. Several essential oils have been used as therapeutic agents since ancient times, and some of them have been scientifically proven to possess medicinal properties, including anti-inflammatory (Koudou *et al.*, 2005), antiviral (Loizzola *et al.*, 2007), antitumor (M'Barek, 2007), cytotoxic (Zarai *et al.*, 2011) and antimicrobial activities (Alviano *et al.*, 2007). Natural phytochemicals including flavonoid and phenolic compounds as well as fatty acids are major bioactive compounds known to be beneficial against many diseases and have been reported to possess a wide range of biological effects like anti-oxidant and antibacterial activities (Sofowora, 1996). *C. patens* is

Table 5. Compounds identified in fraction Cp7 of *Cleistopholis patens*.




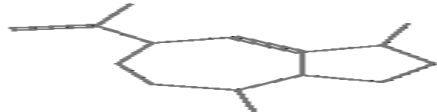

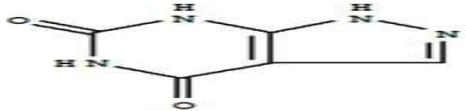
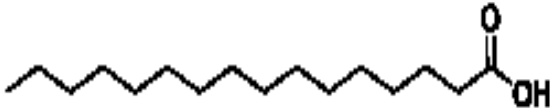
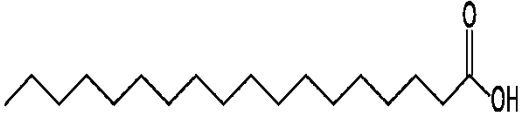


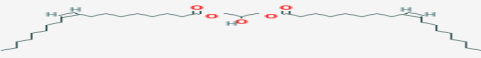
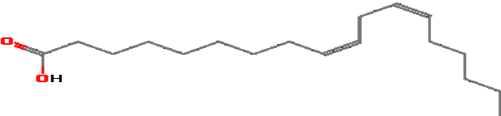
Peak	R.time.	Name of compound	Mol. formula	Mol. weight	Area	Chemical structure
1	4.988	Bicyclo[3.1.1]hept-2-ene, 2,6,6-trimethyl-alpha.-Pinene	C ₁₀ H ₁₆	136	1.65	
2	5.680	Bicyclo[3.1.1]heptane, 6,6-dimethyl-2-methylene- (beta.-Pinene)	C ₁₀ H ₁₆	136	1.43	
3	11.658	Cyclohexene,3-methyl-6-(1-methylethylidene)- (Terpinolene)	C ₁₀ H ₁₆	136	0.22	
4	14.895	1,2,3,4,5,6,7,8-octahydro-1,4-dimethyl-7-(1-methylethenyl)-, [1S-(1.alpha.,4.alpha.,7.alpha.)]- (Sesquiterpenes)	C ₁₅ H ₂₄	204	3.23	
5	15.140	1,6-Cyclodecadiene, 1-methyl-5-methylene-8-(1-methylethyl)-, [s-(E,E)]-sesquiterpenes	C ₁₅ H ₂₄	204	1.53	
6	16.611	1H-Pyrazolo[3,4-d]pyrimidine-4,6(5H,7H)-dione (oxypurinol)	C ₅ H ₄ N ₄ O ₂	152	0.75	
7	22.390	n-Hexadecanoic acid (Palmitic acid)	C ₁₆ H ₃₂ O ₂	256	4.62	

Table 5. Cont

8	24.723	Octadecanoic acid (Stearic acid)	$C_{18}H_{36}O_2$	284	21.98	
9	24.924	2,6,10-Dodecatrien-1-ol, 3,7,11-trimethyl- (Farnesol)	$C_{15}H_{26}O$	222	12.20	
10	25.724	Hexadeca-2,6,10,14-tetraen-1-ol, 3,7,11,16-tetramethyl-, (E,E,E) (Geranylgeranoil) (Diterpenes)	$C_{20}H_{34}O$	290	37.54	
11	27.184	Octadecanoic acid, 2-hydroxy-1,3-propanediyl ester (Alcoholic fatty acids)	$C_{39}H_{76}O_5$	624	5.36	
12	27.031	9,12-Octadecadienoic acid (Z,Z)- (diterpenoid)	$C_{18}H_{32}O_2$	280	6.25	

found to be rich in fatty acids including stearic acids, lenolic acids and esters which occur in copious amount in the plant and are known to have medicinal properties as antimicrobial, antifungal agents (Krishnaveni *et al.*, 2014). The biological activities of free fatty acids have roles in host defenses against potential opportunistic or pathogenic microorganisms. An important aspect

of this is growth inhibition or the quick destruction of bacteria (Desbois and Smith, 2010). Sesquiterpenes are also present in extract. Sesquiterpenes are known to have anti-inflammatory (Jeena *et al.*, 2013) and have bactericidal properties (Ishnava *et al.*, 2013). 9,12-Octadecadienoic acid (Z,Z)- a diterpenoid also present is used as anti-inflammatory,

hypocholesterolemic, cancer preventive, hepatoprotective, nematicide, insectifuge(cide), antihistaminic, antieczemic, antiacne, 5- α reductase inhibitor, antiandrogenic, antiarthritic, anti-coronary, antimicrobial (Adeoye-Isijola *et al.*, 2018).

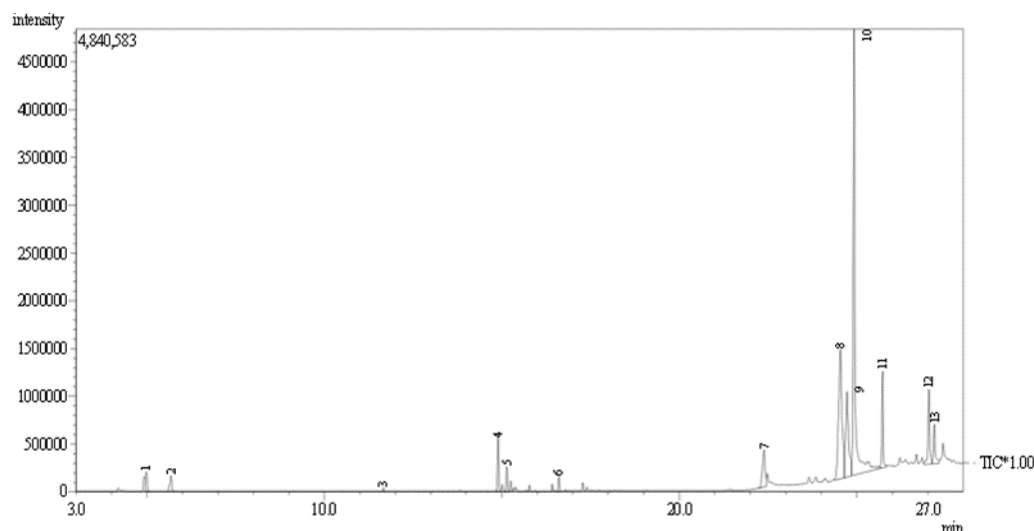


Figure 2. The GC-MS analysis of Cp7 fraction of *Cleistopholis patens*.

Conclusion

The essential oils extracted from this plant possess strong antimicrobial activity against various bacterial pathogens. The reactivity of essential oils depends upon the nature of their functional groups and orientation. Essential oils may disrupt the cell membrane of the targeted pathogens by increasing membrane permeability, inducing leakage of vital intracellular constituents, and interrupting the cellular metabolism and enzyme kinetics of the targeted pathogens. The result of this study showed plant *Cleistopholis patens* are rich in compounds that are responsible for the biological activities of the plant. The phytochemicals observed in this plant can be subjected to further studies to identify their mode of action, toxicity and spectrum of activity. However, the plant is a good candidate for drug production.

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