

In-Vitro Susceptibility Profile of *Pseudomonas aeruginosa* to Selected Medicinal Plants Obtained from University of Calabar Farm, Nigeria

ABSTRACT

Antibiotic resistance represents the greatest challenges to global health. Strategies to curb the menace of antibiotic resistance include researching for new antimicrobials from plants. This study was carried out to determine the in-vitro susceptibility of *Pseudomonas aeruginosa* to *Chromolaena odorata*, *Jatropha tanjorensis*, *Ocimum gratissimum*, *Vernonia amydalina*, *Hibiscus sabdariffa* and *Aloe barbadensis* obtained within University of Calabar farm. All the test carried out were performed according to standard microbiological procedure. Ethanol was used for extraction of Phytochemicals and were purified using column chromatography. In-vitro susceptibility of selected plants against *P. aeruginosa* was in the order *C. odorata* > *J. tanjorensis* > *O. gratissimum* > *H. sabdariffa* > *V. amydalina* > *A. barbadensis* (Aloe vera). The result revealed that the leaves of *C. odorata* had the highest antimicrobial effect against *P. aeruginosa*. The first column of all the plant extracts showed no antimicrobial activity. MIC of *C. odorata*, *J. tanjorensis*, *O. gratissimum* and *H. sabdariffa* were observed in 10^{-7} , 10^{-6} , 10^{-5} and 10^{-4} dilution respectively. Statistical analysis between the zone of inhibitions and layers of column chromatography was significant (<0.05). Ethanol extract and the third column of *Chromolaena odorata*, *Jatropha tanjorensis*, *Ocimum gratissimum* have great potential antibacterial activity against *P. aeruginosa*.

Keywords: Medicinal, Extract, Plant, *Pseudomonas aeruginosa*

1. INTRODUCTION

Medicinal plants have been used for centuries to treat diseases in humans and animals [1]. This is due to the therapeutic potentials of their phytochemicals. Some of these phytochemicals exhibit a wide range of actions on pathogenic organisms [2]. The antimicrobial potential of these phytochemicals makes it a unique substance that can be used to initiate a bacteriostatic or bacteriocidal action on bacteria [3]. Phytochemicals have also been known to have some great antioxidant potentials which is of great interest as a result of their health benefits to humans [4]. Phytochemicals which include carotenoids, phytosterols, limonoids, polyphenol (i.e flavonoids, isoflavonoids anthocyanidins), Glucosinolates, phytoestrogen, terpenoids (isoprenoids), fibers, polysaccharides and saponins have been reported to have great health impact.

Flavonoids are the largest and most studied group of phytochemicals [4]. Flavonoids have been shown to have therapeutic effect as antihypertensive and antirheumatic as well as antimicrobial [3]. Glycoside have been used as a stimulant in cases of cardiac failure and disease [5]. Alkaloids can act as an antibacterial agent, anti-inflammatory, analgesic, hyporotic, local anesthetic, psychotropic and anti-tumor activity and many others. It also serves as stimulant.

Tannin is also another phytochemical that has great potential in promoting wound healing [6]. Moreso, tannin can be use as astringents against diarrhoea and as antioxidant, antiseptic and homeostatic pharmaceutical [7]. Terpenoid also refers to as isoprenoids is a naturally occurring organic chemical present in plant. It is derived from 5-carbon compound isoprene and isoprene polymer called terpenes. Terpernoid derived from plants are used for

their aromatic qualities and play a very important role in traditional herbal remedies. It can also be used as antimalarial, antimicrobial antiulcer [8,8].

Pseudomonas aeruginosa is a gram-negative bacterium that has an inherent (innate) resistant to many commonly used antibiotics such as ciprofloxacin, tobramycin and imipenem, its fast rate of developing resistant to new antibiotics has been a problem to health sector [10,11] Through *P. aeruginosa* normally pathogenic when introduce to areas devoid of normal immune defense, its resistance ability makes it a major pathogen [12,13]. Thus, this study sought to find out an alternative for treatment of infections caused by *Pseudomonas aeruginosa*.

2. MATERIAL AND METHODS

2.1 Sample collection

Stock culture of *Pseudomonas aeruginosa* was obtained from University of Calabar teaching hospital culture collection center. The isolates was taken to the microbiology laboratory for further analysis.

2.2 Identification of the isolate.

After 24 hours of incubation, Gram staining was carried out in order to confirm the suspected isolates as gram negative rods. The colonies formed were examined microscopically. Different biochemical test were employed for the presumptive identification of the isolates.

2.3 Plant collection and preparation of plant extract:

Six plant species (*Hibiscus sabdariffa*, *Chromolaena odorata*, *Jatropha tanjorensis*, *Aloe vera*, *Vernonia amygdalina* and *Ocimum gratissimum*) were collected within University of Calabar farm.

The leaves of the plant were removed from the plant, washed and then dried under room temperature. The leaves of the plant were crush into powdery form. 100g of dried powdered of each tested plants were extracted separately by maceration several times with ethanol (95%) as the solvent.

2.4 Purification of plant extract using column chromatography.

Column chromatography was use in purification of the plant extracts particularly the gravity column chromatography since the solvent was allowed to flow down the column by gravity without been forced by pressure pump. The solid phase used was silica gel and alumina.

Pieces of wire was used to add a pug of cotton to the bottom of the column, so that the silica will not fall out of the column. However, too much cotton or cotton packed too tightly will prevent the eluent from dripping at an acceptable rate.

The column was clamped to a tripod stand. Firstly, the column was filled up to 1/3 with silica gel and the alumina was filled immediately after the silica gel. A space was left on top of the column for the eluent (solvent). The eluent was drained from the column until no solvent remains above the surface of the alumina. Using a Pasteur pipet, the extracts was carefully added to the column. Eluent was added to the column to ensure that no sample remain above the surface of the stationary phase (alumina). Three different fractions were collected into separate sample bottle for further analysis.

2.5 Susceptibility testing

Two techniques (Disk diffusion and tube dilution method) were used to carry out susceptibility testing of the six selected medicinal plant extracts against *Pseudomonas aeruginosa*.

2.6 Determination of minimum inhibitory concentrations

This was determined using tube dilution method according to method described by Nwachukwu *et al.* [15]. The plant extracts with a better inhibition zone were selected and used for MIC test.

2.7 Statistical analysis.

Statistical analysis was carried out using Microsoft Excel (2007). Data obtained from the inhibition zone of ethanolic extracts and the different column of column

chromatography were analyzed using regression analysis and values for $P < 0.05$ were considered as statistically significant

3. RESULTS

The morphologically characteristics of the isolates were recorded followed by a series of biochemical test as presented in Table 1. Invitro susceptibility testing was performed on *Pseudomonas aeruginosa* species against the six medicinal plants using disc diffusion technique. Their various zones of inhibition were measured as shown in Table 2.

The result shows that *chromolaena odorata*, *Jatropha tanjorensis* and *Ocimum gratissimum* exhibited antibacterial activity on the *Pseudomonas aeruginosa* isolate. While *Vernonia amydalina* and *Aloe vera* showed no antibacterial activity on the isolates. But for *Hibiscus sabdariffa* it shows little inhibition effect.

Table 1: Morphological and biochemical characteristics of *Pseudomonas aeruginosa*

Test	Observation	Inference
Gram stain	Curve rode, which appear pink	Gram negative
Catalase	Formation of bubble	Positive
Oxidase	Colorless to dark blue color	Positive
Citrate	Green to blue	Positive
Sugar fermentation	No changes	Negative
Motility	diffused, hazy growths that spread throughout the medium rendering it slightly opaque	Motile
Urease	Color medium remained yellowish	Negative
Methyl red	Red	Positive
Voges proskauer	Yellow	Negative
Hydrogen sulphide	No change	Negative

Table 2: Inhibition zones of the medicinal plants

Plants	Ethanol extract	First column (C1)	Second column (C2)	Third column(C3)
<i>chromolaena odorata</i>	25	7	10	19
<i>Jatropha tanjorensis</i>	23	5	13	15
<i>Vernonia amydalina</i>	4	0	0	0
<i>Ocimum gratissimum</i>	20	0	5	10
<i>Aloe</i>	2	0	0	0
<i>barbadensis</i>				
<i>Hibiscus</i>	11	3	5	9
<i>sabdariffa</i>				

The antibacterial screening for *chromolaena odorata* showed that the ethanol extract exerted antimicrobial activity on *Pseudomonas aeruginosa* with inhibition zone of 25mm, column three with inhibition zone of 15mm, column two with inhibition zone of 10mm and column one with inhibition zone of 7mm.

For *Jatropha tanjorensis*, ethanolic extract showed inhibition zone of 23mm column three with inhibition zone of 15mm, column two with inhibition zone of 13mm and Column one of *Jatropha tanjorensis* showed inhibition zone of 5mm. The *Vernonia amydalina* and *Aloe vera* exhibited little or no antimicrobial activity. Ethanol extract of *Vernonia amydalina* showed inhibition zone of 4mm while for *Aloe vera*, ethanol extract showed inhibitory zone of 2m. Column three, two and one of both *Vernonia amydalina* and *Aloe vera* showed no zone of inhibition (0mm)

Ocimum gratissimum ethanol extract showed inhibition zone of 20mm, column three with inhibition zone of 10mm and column two and one with inhibition zone of 5mm and 0mm respectively. *Hibiscus sabdariffa* ethanol extract demonstrated lowered antibacterial activity with inhibition zone of 11mm, column three with inhibition zone of 9mm, column two with inhibition zone of 5mm and column one with inhibition zone of 3mm

Zones of inhibition (mm)

Furthermore, antimicrobial activities of all the plants extracts are represented in the figure 1 below

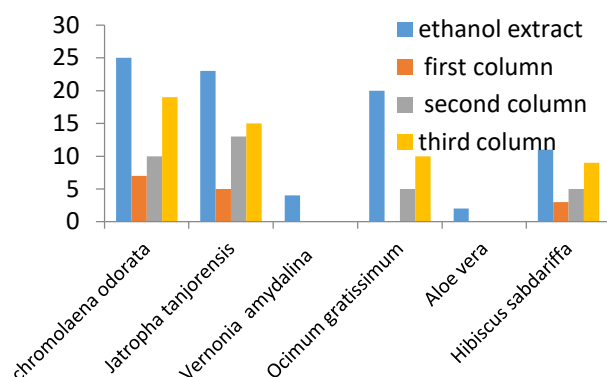


Figure 1: Antibiogram study of the ethanol extracts and the three layers or columns of the chromatography.

From the above figure, all the ethanolic extracts shows antimicrobial activity. But *chromolaena odorata* shows highest antimicrobial activity followed by *odorata*, *Jatropha tanjorensis*, *Ocimum gratissimum* and *Hibiscus sabdariffa*. Ethanolic extracts of *Vernonia amydalina* and *Aloe vera* shows least antimicrobial activity.

Extracts that gave higher zones of inhibition were used for conducting MIC using Macro tube dilution method. The result is presented in Table 3.

Table 3: Minimum inhibitory concentration

Plants extracts	ethanol	Highest dilution factor that shows no tubidity
<i>chromolaena odorata</i> ,		10^{-7}
<i>Jatropha tanjorensis</i>		10^{-5}
<i>Ocimum gratissimum</i>		10^{-6}
<i>Hibiscus sabdariffa</i>		10^{-4}

3.2 Statistical analysis.

At 95% confidence interval or 0.05 significance level. p-value of the calculated was observed to be 0.0274 which implies that $p < 0.05$. Since $p < 0.05$, the null hypothesis was rejected which stated that there is no correlation or there is no association between the independent and the dependent variable. Therefore, this variable is statistically significance.

4. DISCUSSION

In this study, *Chromolaena odorata*, *Jatropha tanjorensis*, *Ocimum gratissimum*, *Vernonia amydalina*, *Aloe vera* and *Hibiscus sabdariffa* were used to perform antimicrobial susceptibility test. Although leaf, stem and root are the three parts of plant and each has antimicrobial activity. However, the leaf of *Chromolaena odorata*, *Jatropha tanjorensis*, *Ocimum gratissimum*, *Vernonia amydalina*, *Aloe vera* and flower of *Hibiscus sabdariffa* were used in this study. Similarly, there are various solvent that can be used in the extraction of plant, but ethanol was the only solvent that was used in the extraction process.

The result of this analysis showed that the leaves of *Chromolaena odorata* ethanol extract, first, second and third column exhibited high antimicrobial effect against *Pseudomonas aeruginosa* with the zones of inhibition ranging between 7.0mm -25.0mm. This result is similar with the findings of Nurul *et al.*, [14]; Nwachukwu *et al.*, [15]; Mbajiuka *et al.*, [16] and Srisuda *et al.*, [17]. Furthermore, Srisuda¹⁸ also recorded that both ethanol and methanol solvents could extract antimicrobial compounds (flavonoids, flavones, polyphenols, terpenes and tannins) from the leaf of *C. odorata*.

Jatropha tanjorensis showed antimicrobial activity with the zones of inhibition ranging between 5mm-23mm. This is similar to the result of a study carried out by Ogunnusi¹⁸ on the activity of leaf extracts of *Jatropha tanjorensis* and found out that it had activity against *Pseudomonas aeruginosa*. The actimicrobial activity of this plant is as a result of phytochemical constituents of the plant as Ogunnusi *et al.*, [18] recorded that

the plant extracts revealed the presence of saponins, alkaloids, tannins, cardiac glycosides and flavonoids and absence of phenols and phlobatannins.

Vernonia amygdalina showed very little zone of inhibition ranging from 0mm-4mm. Recently, Kolawole *et al.*, [19] also reported that *Pseudomonas aeruginosa* was not susceptible to *Vernonia amygdalina* and stated that the reason for this might be for the low amount of steroids, flavonoids, alkaloids and tannins in *Vernonia amygdalina* (bitter leaf). This also agreed with report of Nweze [19,20].

Ocimum gratissimum showed antimicrobial activity with the exception of the first column obtained from column chromatograph. The ethanol extract, the third and second column exhibited zone of inhibition ranging between 5mm-20mm. Kolawole *et al.*, 2018 also stated that the higher presence of steroids, flavonoids, alkaloids and tannins in *Ocimum gratissimum* (scent leaf) than in *Vernonia amygdalina* (bitter leaf) could have been attributed to its higher antibacterial activity against the test organisms as this plant extract was found to be susceptible to *Pseudomonas aeruginosa*.

Priscilla *et al.*, [21] recorded that *Hibiscus sabdariffa* inhibitor effect of the antimicrobial activity of the aqueous and ethanol extracts seems to have an effect on both Gram positive and Gram-negative bacteria and suggests the broad-spectrum nature of the plant against bacteria. In this study ethanol extract showed small inhibitory effect with the zone of clearance ranging between 3mm-11mm. Gehan *et al.*, [22]; Ifeanyi *et al.*, [23] also recoded similar result, in their study *Hibiscus sabdariffa* showed lower antimicrobial activity with all the isolate of *pseudomonas* with inhibition zone ranging between 9.4mm-10.8mm.

Results obtained from this study, indicated that the *Aloe barbadensis* extracts showed lower antimicrobial activity than the commercially available antibiotics used.

Generally, the first column of all the plant extracts obtained from the column

chromatography showed little or no antimicrobial activity. The extent of antimicrobial activity between the various column and the ethanol extracts is in the order of: ethanol extract>third column>Second column> first column. This may be due to the fact that some phytochemicals constituents were washed during purification process by column chromatography

5. CONCLUSION

The study has revealed that *Chromolaena odorata*, *Jatropha tanjorensis*, *Ocimum gratissimum* plant extract have good antibacterial properties thus a potential source of new drug for treating wound infections and other ailment caused by *Pseudomonas aeruginosa*. Although, *H. sabdariffa* also showed antimicrobial activity but lower than *Chromolaena odorata*, *Jatropha tanjorensis*, *Ocimum gratissimum*. In contrast, *Vernonia amygdalina* and *aloe vera* show no antimicrobial activity.

It can be concluded from this research that, the crude ethanol extract and the third column of *Chromolaena odorata*, *Jatropha tanjorensis*, *Ocimum gratissimum*, showed the best antimicrobial activity against the isolate. This is followed by second and first column of the extracts. Therefore, they may be explored further in the field of human medicine for the production of antimicrobial agents which will significantly inhibit the growth of bacteria.

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