# Antimicrobial Resistance Evaluation of Organisms Isolated from Liquid Herbal Products Manufactured and Marketed in South Eastern Nigeria. Ujam NT<sup>1</sup>, Oli AN<sup>1\*</sup>, Ikegbunam MN<sup>1</sup>, Adikwu MU<sup>2</sup>, Esimone CO<sup>1</sup>

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# 16 **ABSTRACT**

### 17

**Objective:** To determine the susceptibility and resistance pattern of bacteria and fungi isolates obtained from herbal anti-infective liquid preparations manufactured and marketed in South-East Nigeria to conventional antibiotics.

# Study Design: Experimental

**Place and Duration of the study:** Pharmaceutical Microbiology and biotechnology Laboratory, Faculty of Pharmaceutical Sciences, Nnamdi Azikiwe University, Agulu Campus between October 2011 and March 2012

**Methodology:** Isolation and characterization of contaminating microorganisms were carried out using standard procedures. A total of forty-nine (49) bacteria and forty (40) fungi isolated from the herbal products were examined for susceptibility to conventional antibiotics using the disc diffusion method. The bacterial isolates were tested against ciprofloxacin, ofloxacin, amoxicillin-clavulanic acid, gentamicin, cefotaxime, ceftraidime, ceftriazone, sulphamethoxazole, tetracycline and ampicillin were employed while fungi isolates were tested against five common antifungal-griseofulvin, nystatin, ketoconazole, fluconazole and clotrimazole. The Multiple Antibiotic Resistance Index (MARI) of each of the isolated bacteria was obtained following the standard method.

**Result:** The antimicrobial susceptibility-resistance profile of the bacteria isolates revealed that most of the bacteria were sensitive to ciprofloxacin, ofloxacin, gentamicin, and ceftriaxone, On the other hand, a good number of the isolates demonstrated high level of resistance to common antibiotics like Ampicillin, amoxycillin-clavulanic acid, trimethoprim-sulphamethoxazole, and moderate level of resistance to Tetracycline, and some of the third generation cephalosporins - ceftazidime and cefotazime. Multiple Antibiotic Resistance Index (MARI) evaluation revealed that most of the isolates were resistance to more than fifty percent (50%) of the number of antibiotics used. The fungal isolates were susceptible to nystatin, ketoconazole and clotrimazole, resistance to fluconazole and high resistance recorded against griseofulvin.

**Conclusion:** The results of this study revealed that the herbal medications can serve as a trail of spread of antibiotic-resistance genes.

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Keywords: {Susceptibility, antibiotic resistance, herbal anti-infectives}

# 21 **1. INTRODUCTION**

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The use of herbal medicine has always been part of human culture, as some plants possess important therapeutic properties, which can be used to cure human and other animal diseases [1]. Herbal medicine is becoming increasingly

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25 popular in both developing and developed countries [2]. A World Health Organization survey indicates that about 70–80% 26 of the world population, particularly in developing nations; rely on non-conventional medicines mainly of herbal sources in 27 their primary health care [3]. Medicinal plant materials normally carry a large number of microbes originating from the soil. 28 Microorganisms of various kinds are normally adhered to leaves, stems, flowers, roots and seeds. Additional 29 contaminants may also be introduced during harvesting, handling and production of various herbal remedies since no conscious efforts are made to decontaminate the herbs other than by washing them. [4]. Herbal medicines are therefore 30 vulnerable to attack by microorganisms and as such are disposed to spoilage. Accordingly, gross microbial contamination 31 32 of herbal medicinal products commonly consumed in Nigeria has been severally demonstrated [5, 6,7]. The presence of antibiotic resistant microbial isolates in the Herbal Medicinal Products (HMPs) could lead to transfer of antibiotic 33 34 resistance traits to hitherto sensitive out or oral micro flora of consumers [8].

The emergence of multiple drug resistant bacteria (MDR) has become a major cause of failure of the treatment of infectious disease [9]. As a result, society is facing one of the most serious public health dilemmas over the emergence of infectious bacteria displaying resistance to many and in some cases, effective antibiotic [10] much like the situation in human medicine. Bacteria and fungi resistance to antimicrobial drugs has continued to grow in the last decades [11]. The increased prevalence of their resistant is due to extensive use and misuse of antimicrobials. This has rendered the current available antimicrobial agents insufficient to control microbial infections and create major public health problem.

Resistant bacteria strains may develop almost anywhere particularly in a pressurized environment containing previously non-resistant bacteria strains as contaminants. One of such environments can be created by widespread use of HMP. HMPs have been previously implicated as a pool for such contaminations [12,13]. It is of utmost importance to both monitor and ascertain the microbial purity of HMPs given the huge medical and economic implications of any such microbial contamination especially with multiple drug resistant strains. Such surveillance will both help to identify microbial contamination of herbal products and slow down and prevent the emergence of drug-resistant strains. The present study evaluated the presence of contaminating organisms and the susceptibility-resistance pattern of the isolated organisms.

# 49 2. EXPERIMENTAL DETAILS

#### 50 2.1 MATERIALS: HERBAL SAMPLES

A total of twenty liquid herbal anti-infectives were purchased randomly from different shops and herbal outlets located within the five states that make up the south-east, Nigeria and were used in this study. The samples which were within their shelf lives and were kept at room temperature (as indicated by their manufacturers) were used within two weeks of collection.

#### 55 2.2 METHODS

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#### 56 **2.2.1 Isolation and Identification of Microbial Contaminants in the Herbal**

57 The herbal anti-infectives were serially diluted and plated on nutrient agar and sabouroud dextrose agar plates in triplicate 58 and incubated at 37°C for 18-24 hours and 20°C- 27 °C for 72-168hours for bacteria and fungi respectively. The resultant 59 colonies were further purified, isolated and characterized using standard methods [14].

#### 60 2.2.2 Characterization of Microorganisms Isolated From the Herbal Preparations

The bacteria isolates were characterized using the morphological appearance (macroscopy) of their colonies, their Gram stain reaction and confirmatory biochemical tests. The fungi isolates were also identified on the basis the morphological characteristics (macroscopy) of their colonies, microscopy, staining with ordinary stain and the appearance of their mycelia [15].

#### 65 2.2.3 Antibiotics Susceptibility Testing

The susceptibility tests were performed following the method M2-A6 disc diffusion method recommended by the National 66 Committee for Clinical Laboratory Standards [16] using Mueller Hinton and Sabouraud Dextrose Agar. The bacterial 67 isolates from the samples were reactivated by sub-culturing from agar slant onto nutrient agar plate and was incubated for 68 18-24 hours. The inoculum was standardized by transferring three distinct and separate colonies of the pure culture of the 69 70 test organism using sterile wire loop into 3mls of sterile nutrient broth. The suspension was incubated for 3 hours at 37°C 71 to allow for the growth of test organism till the density was equivalent to the turbidity of 0.5 McFarland. The standardized 72 inocula were swabbed onto Mueller-Hinton agar and Sabouraud Dextrose Agar plate and the discs were placed on the 73 inoculated plates and pressed firmly onto the agar plate for complete contact. The bacterial strains were tested against 74 the following discs:, ofloxacin (OFX, 5µg) ; ciprofloxacin (CIP,5µg) ; amoxicillin/clavulanic acid (AMC,20/10µg) ; gentamicin, (GN ,10µg) ; ceftazidime (CAZ,30µg) ; cefotaxime (CTX,30µg) ; trimethoprim-sulfamethoxazole 75 (SXT,1.25/23.75µg); Ampicillin (AMP,10µg); tetracycline (TE, 30µg); ceftriaxone (CRO, 30µg). The fungal strains were 76 tested against the following discs: nystatin (N.20µg) ; clotrimazole (C.20µg) ; griseofulvin (G.20µg) ; ketoconazole 77 78 (K,20µg) and fluconazole (F,20µg). The Plates were inverted and left on the work table for 30 minutes to allow for pre-79 diffusion of antibiotics into the agar. The plates were incubated at 37°C for 18-24 hours and at 25°C 24-48hours for 80 bacteria and fungi respectively. The susceptibility of each isolate to each antibiotic was shown by a clear zone of growth inhibition and this was measured using a meter rule in millimeters and the diameter of the zones of inhibition was then 81 82 interpreted using standard chart [17].

# 8384 2.2.4 Determination of Multiple Antibiotics Resistance Index (MARI)

The Multiple Antibiotics Resistance Index (MARI) of ten antibiotics (ofloxacin, ciprofloxacin, gentamicin, amoxycillinclavulanic acid, sulphamethoxazole-trimethoprim, ceftriazone, ceftazidime, cefotazime, tetracycline and ampicillin) were determined using the formula, MARI = a/b

- 88 Where; a = the aggregate resistance of antibiotics to all isolates and b = the total number of antibiotics that was used
- 89 3. RESULTS AND DISCUSSION
- 90 3.1 RESULTS
- 91
- Table 1: Microorganisms Isolated from the Herbal anti-infective Products.
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Samples Code	<mark>Shelf</mark> life	Contents	Therapeutic Claims	Identity Of Bacteria Isolated	Identity Of Fungi Isolated
1	(48 Months)	Carica papya, Magnifera indica, Newbouldlia, leavis, Azadricha indica, Jaminum officionili, Aloe, barbedensis, Ginseng, T <u>reated</u> water 60cl.	Antibacterial, Antimalarial, Ant rheumatic, infertility, Antiviral.	a)Staphylococcus aureus b)Proteus spp	a) Microsporuim spp. b) Aspergillus spp c) Nigrospora spp
2	<mark>(36</mark> Months)	38 African Roots, Herbs, Fruits, Barks plus ginseng, Aloevera and Garlic.	Antibacterial, Antirheumatic, Antifungal and Antiviral.	a)Escherichia coli b)Staphylococcus aureus c)Staphylococcus epidermidis d)Pseudomonas aeruginosa e)Bacillus spp f)Proteus spp	a) Candida tropicalis b) Microsporum canis
3	<mark>(30</mark> Months )	60% herbs, 25%flower, 10% leaves, 5% roots.	Antibacterial, Antirheumatic, Antifungal, Earlier Menopause, Painful and irregular menstruation.	a)Staphylococcus aureus b)Bacillus subtilis c)Bacillus cereus d)2 Salmonella spps	a) Candida albicans b) Candida tropicalis c) Trichosporon spp
4	(48 Months)	Aloe vera plus 31 roots and herbs ,fruits and barks	Antibacterial, Antifungal.	a)Escherichia coli b)Staphylococcus areus c)Streptococcus spp d)Bacillus spp	a) Coccidioides immitis b) Microsporuim audounii
5	(36 Months)	Water, herbs, root and fruits.	Antibacterials, Antimalarial, Antiparasitic, Internal heat, pile, and reduces sugar.	a)Staphylococcus epidemidis b)Streptococcus spp c)Yersinia spp	-
6	(24 Months)	•	Antibacterial, Treatment of all form of eye infections.	a)Escherichia.coli b)Bacillus spp c)Proteus spp	a)C.topicalis b) Microsporuim audounii c)Aspergillus niger
7	<mark>(12</mark> Months)	Nauclea diiderchi 10%,Hippocrates pallens 20%,Alluim sativum12.5%,Cochios permum planchoni 5.5%,Uvaria chame 5%,Punica granatum 47%.	Antibacterial, Antimalarial.	a)Escherichia coli b)Staphyloccus spp c)Pseudomonas d)Bacillus spp	a) Blastomyces b) Microporuim canis

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8	(36 Months)	Aloe vera 40%, Olong tea 20%, Flower and roots 40%, Saracin.	Antibacterial.	a)Staphylococcus spp b)Salmonella spp	a) Blastomyces
9	<mark>(36</mark> Months)	Aloe Vera	Antibacterial Anti- malarial, HBP, Cough Antirheumatism, e.t.c.	a)Staphylococcus areus	a) Blastomyces spp b) Cryptococcus neoformans c) Histoplasma
10	<mark>(36</mark> Months	Aloe Vera, Flowers, Fruits seed barks.	Antibacterial, Hypertension, Antiviral, fibroid, stroke.	a)Pseudomonas aeruginosa	a)Penicillum spp b)Aspergillus spp
11	<mark>(24</mark> Months)	Honey (natural), Lime juice, Zingiber officillinar, Herbal seeds and roots.	Antibacterial and Asthmatic cough.	a)Escherichia coli b)Staphylococcus	a)Candida spp
12	<mark>(31</mark> Months)	•	Anti-bacterial Antiviral, Diabetes, Reduces cholesterol.	a)Staphylococcus areus b)Streptocoloccus spp	a) C.albicans b)Blastomyces
13	<mark>(24</mark> Months)	25 different types of roots, herbs, seeds and flowers.	Anti-bacterial, Anti- malaria, Antirheumatic. Antifungal	a)Escherichia coli b)Corynebacteruim diphtheria	a)Aspergillus spp b) Nigrospora spp c) C.tropicalis
14	(60 Months)	Herbs, water, root and fruits.	Antibacterial, Antiviral, Antirheumatic, Antifungal, Antiparatic, internal heat, pile.	a)Escherichia coli b)Streptococcus spps c)Yersinia spp	a)C.albicans b) Aspergillus spp
15	(29 Months)	Magnifera indica, Carica papaya leaves, Psiduim guajava, Breadfruit bark, Masularia acuminate roots, Citrus lemon leaves, Zingiber Officinale roots, Cymbopogon spp.	Antibacterial, Antimalarial, Antirheumatic Antiviral.	a)Escherichia coli	a)Microsporum spp b) Coccidioides spp c)Aspergillus spp.
16	(48 <mark> Months)</mark>	Awapa bark, white lotus, Golden seal, Mahogany, Ukor root, Aloe barbaders, Mistletoe, Osisika Aguru, Uda roots, Uvuru ilu, Lemon grass.	Antibacterial, Antirheumatic and Arthritis, Veneral diseases.	a)Escherichia coli b)Pseudomonas aeruginosa	a)C.albicans b) Mucor spp
17	<mark>(15</mark> Months)	Aloe Vera, Cadeperi salt, Lime	Antibacterial, Treatment and prevention of toothache.	a)Proteus spp	a) Aspergillus spp
18	(12 Months)	Lymbopogon citrates, Carica papaya leaves, Magnifera indica, bark, Treculia Africana, Citrus, Limonia, Psiduim guajava, Zingibar officinale root, Alluim sativum,	Antibacterial, Antirheumatism reduces sugar and cholesterol.	a)Escherichia coli	a)Mucor spp
19	<mark>(24</mark> Months)	Natural roots and barks.	Antibacterial, Antiviral, Purifies	a) Bacillus spps	a) Yeast/Blastomyces

			toxins, Builds immune system, Stops dizziness, weakness.		c) Microsporum s <sub>l</sub>
20	<mark>Months)</mark> sati bitte Pre Hibi	lealatifolia, Allium vum, Aloe Vera er, Chick weed, clina nitida, scus sabdrrifa, a, Ethanol.	Antibacterial, Antiparasitic, ulcer, constipation, fibroid, internal heat heart burn and diabetes	a) Staphylococcus epidermidis	a)Aspergillus spp
Table Or Dares	untage of microh	ial inclates from th	o Llorbol onti infontiv	ve Dreducte	
BACTERIA	<pre>%</pre>	FUNGI ISOLATE	e Herbal anti-infectiv	e Products	
SOLATES	70 Occurrence	IUNOIDOLAI	Occurrence	<u>.</u>	
E.coli	20.4	<mark>Aspergillus spp</mark>	22.5		
S.aureus	24.5	Microsporuim sp			
P.aeruginosa	8.2	Candida spp	22.5		
Strep.spp	<mark>10.2</mark>	Trichosporon sp			
Bacillus	<mark>16.3</mark>	Coccidiodes spp		)	
Salmonella	<mark>6.1</mark>	Blastomyces sp	<mark>) 12.5</mark>	5	
Proteus spp	<mark>8.2</mark>	Cryptococcus sp	p 2.5	5	
<mark>Yersinia spp</mark>	<mark>4.1</mark>	Histoplasma spp	2.5	5	
	<mark>2.0</mark>	Penicilluim spp	<mark>2.5</mark>	5	
		Nigrospora spp	<mark>5.0</mark>	)	
C.diphtheria					

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A total of 89 microbial strains (49 bacterial and 40 fungal strains) were isolated from the herbal preparations. The
identified microbial isolates consists of nine (9) bacterial genera and eleven (11) fungal genera which include *Staphylococcus, E. coli, Bacillus, Streptococcus, Pseudomonas, Proteus, Salmonella, Yersinia, Corynebacteruim diphtheria* and *Aspergillus, Candida, Microsporuim, Trichosporon , Coccidiodes, Blastomyces, Cryptococcus, Histoplasma, Penicilluim, Nigrospora*, Mucor respectively. The most frequently isolated bacteria and fungi specie were *Staphylococcus* spp (24.5%) and *Aspergillus* spp/*Candida* spp (22.5%) respectively. The least frequently isolated bacteria
specie was *Corynebacteruim diphtheria* (2.0%) and that of fungi were *Trichosporon* spp, *Cryptococcus* spp, *Histoplasma*and *Penicillium* spp (2.5%). See Tables 1 and 2 above.

105 Table 3: The Antibiotic Susceptibility-Resistance Profile of the isolated bacteria

Drugs and Strength (µg)		OFX-5	CIP-5	SXT- 1.25/ 23.75	AMC- 20/10	GN-10	CTX- 30	CAZ- 30	TE-30	AMP- 10	CRO- 30	
			N (%)	N (%)	N (%)	N (%)	N (%)	N (%)	N (%)	N (%)	N (%)	N (%)
	E. coli	S	8 (80)	10 (100)	0 (0)	0 (0)	9 (90)	0 (0)	3 (30)	1 (10)	0 (0)	5 (50)
		T	1 (10)	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)	2 (20)
TES		R	1 (10)	0 (0)	10 (100)	10 (100)	1 (10)	10 (100)	7 (70)	9 (90)	10 (100)	3(30)
ISOLATES	P. aeruginosa	S	3 (75)	3 (75)	0 (0)	0 (0)	3 (75)	0 (0)	2 (50)	0 (0)	0 (0)	2(50)
		T	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)	3 (75)	0 (0)	0 (0)	0 (0)	1 (25)
TER		R	1 (25)	1 (25)	4 (100)	4 (100)	1 (25)	1 (25)	2 (50)	4 (100)	4 (100)	<b>30</b> <b>N (%)</b> 5 (50) 2 (20) 3(30) 2(50) 1
BACTERIA	Staphylococcus spp	S	8 (67)	10 (83)	0 (0)	0 (0)	10 (83)	0(0)	2 (17)	2 (17)	0 (0)	7(58)
		L	4 (33)	2 (17)	0 (0)	0 (0)	0 (0)	2 (17)	1 (8)	2 (17)	N (%)         N           0 (0)         5           (5)         (5)           0 (0)         2           10         3(           (100)         0           0 (0)         2(           0 (0)         2(           0 (0)         1           (2         10           0 (0)         1           (100)         0           0 (0)         1           (100)         7(           0 (0)         4           (3)         (3)	
		R	0 (0)	0 (0)	12 (100)	12 (100)	2 (17)	10 (83)	9 (75)	8 (67)		1(8)

	0		4 (00)	4 (00)	0 (0)	0 (0)		0(07)	0 (07)	4 (00)	0 (0)	4(00)
	Salmonella spp	S	1 (33)	1 (33)	0 (0)	0 (0)	3 (100)	2(67)	2 (67)	1 (33)	0 (0)	1(33)
		I	1 (33)	2 (67)	0 (0)	0 (0)	0 (0)	0(0)	0 (0)	0 (0)	0 (0)	1(33)
		R	1 (33)	0 (0)	3 (100)	3 (100)	0 (0)	1 (33)	1 (33)	2 (67)	3 (100)	1(33)
	Streptococcus spp	S	3 (60)	4 (80)	0 (0)	0 (0)	4 (80)	2 (40)	2 (40)	1 (20)	0 (0)	1(20)
		1	1 (20)	1 (20)	0 (0)	1 (20)	1 (20)	0 (0)	1 (20)	1(20)	0 (0)	0(0)
		R	1 (20)	0 (0)	5 (100)	4 (80)	0 (0)	3 (60)	2 (40)	3 (60)	5 (100)	4 (80)
	Bacillus spp.	S	3 (38)	5 (63)	0 (0)	0(0)	6 (75)	0 (0)	1 (13)	4 (50)	0 (0)	5(63)
		T	5 (63)	3 (38)	0 (0)	1(13)	0 (0)	1 (13)	1 (13)	1 (13)	1 (13)	1(13)
		R	0 (0)	0 (0)	8 (100)	7(88)	2 (25)	7 (88)	6 (75)	3 (38)	7 (88)	2(25)
	Proteus spp	S	2 (50)	1 (25)	0 (0)	0 (0)	2 (50)	1 (25)	2 (50)	2 (50)	0 (0)	1 (25)
		I.	2 (50)	3 (75)	0 (0)	0 (0)	0 (0)	0 (0)	1 (25)	0 (0)	0 (0)	2 (50)
		R	0 (0)	0 (0)	4 (100)	4 (100)	2 (50)	3 (75)	1 (25)	2 (50)	4 (100)	1(25)
	Yersinia spp	S	0 (0)	1 (50)	0 (0)	0 (0)	2 (100)	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)
		T	2 (100)	1 (50)	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)	1 (50)
		R	0 (0)	0 (0)	2 (100)	1 (100)	0 (0)	2 (100)	1 (50)	2 (100)	2 (100)	1 (50)
	C. diphtheria	S	0 (0)	1 (100)	0 (0)	0 (0)	1 (100)	0 (0)	1 (100)	0 (0)	0 (0)	0 (0)
		L	1 (100)	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)	0 (0)	1(100)
		R	0 (0)	0 (0)	1 (100)	1 (100)	0 (0)	1 (100)	0 (0)	1 (100)	1 (100)	0 (0)
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Key: S = Sensitive, I = intermediate, R - Resistance, N = number of organisms, OFX= ofloxacin, CIP = ciprofloxacin,
 CAZ = ceftazidime, TE = tetracycline, AMP = ampicillin, SXT = trimethoprim-sulfamethoxazole, GN = gentamicin,

108 CTX = cefotaxime, CRO = ceftriaxone, AMC = amoxycillin-clavulanic acid.

109 Table 3 above shows the antibiogram, of all the bacterial strains isolated from the Herbal products - a representation of

110 the bacteria that are Susceptible, Intermediate or Resistant to the different antibiotics using the NCCLS break points [17].

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Table 4: Antibiogram of Fungi Isolated from the Herbal Anti-Infectives

Samples	Isolates	Inhibition Zo	Inhibition Zone Diameter (IZD) in millimeter (mm)						
Code		Griseofulvin	Nystatin	Ketoconazole	Clotrimazole	Fluconazole			
1	Microsporuim spp.	0	25	0	0	0			
	Aspergillus spp	0	32	10	12	0			
	Nigrospora spp	0	30	20	14	12			
2	Candida tropicalis	0	26	23	7	15			
	Microsporum canis	0	25	0	0	0			
3	Candida albicans	0	22	12	10	7			
	Candida tropicalis	0	23	11	10	8			
	Trichosporon spp	7	30	32	18	8			
4	Coccidioides immitis	0	30	0	0	0			
	Microsporuim audounii	0	29	0	0	0			
6	C.topicalis	0	27	12	11	8			
	Microsporuim audounii	0	8	0	0	0			
	Aspergillus niger	0	29	0	7	0			
7	Blastomyces	0	32	17	18	0			
	Microporuim canis	0	30	0	0	0			

8	Blastomyces	0	31	18	18	0	
9	Blastomyces spp	0	29	16	19	0	
	Cryptococcus neoformans	0	0	0	0	0	
	Histoplasma	0	30	29	12	15	
10	Penicillum spp	0	31	15	7	0	
	Aspergillus spp	0	23	0	12	0	
11	Candida spp	0	22	12	10	7	
12	Candida albicans	0	20	14	12	9	
	Blastomyces	0	30	15	19	0	
13	Aspergillus spp	0	32	17	8	0	
	Nigrospora spp	0	28	14	15	0	
	Candida tropicalis	0	35	25	20	0	
14	Candida albicans	0	22	13	10	10	
	Aspergillus spp	0	26	0	7	0	
15	Microsporum spp	0	25	0	0	0	
	Coccidioides spp	0	30	0	0	0	
	Aspergillus spp.	0	31	10	0	0	
16	C.albicans	0	24	23	8	0	
	Mucor spp	0	26	0	0	0	
17	Aspergillus spp	0	31	10	0	0	
18	Mucor spp	0	25	0	0	0	
19	Yeast/Blastomyces	0	28	18	16	0	
	Aspergillus spp	0	29	0	9	0	
	Microsporum spp	0	0	0	0	0	
20	Aspergillus spp	0	36	0	0	0	

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114 Table 5: Multiple Antibiotics Resistance Index (MARI) of the Isolated Bacteria.

Grouping	Isolates and Samples Code	Multiple Antibiotics Resistance Index (MARI)
Group A	Pa16	1
Group B	<mark>S3</mark>	0.8
Group C	E4,11,14,15,Sa1,P2	0.7
Group D	E7,15,16,13,18,Sa2,Sa5,Sa7,Sa20,St12,St4,Ba7,Ba19b,Ba3a,Ba6,Ba3b,P17,Y5	0.6
Group E	E2,6,Sa3,Sa4,Sa8,Sa11,Sa9,Sa2,Pa10,St5,St14,Ba9a,P6,Y14,C13	0.5
Group F	Sa12,Pa2,Pa7,St14,Ba2,S8,P1	0.4

<sup>115</sup> Notes for Table 5: E = *E.coli*, Sa = Staphylococcus spp, Pa = Pseudomonas aeruginosa, St = Streptococcus spp, Ba =

117 represent the product numbers

#### 118 **3.2 DISCUSSION**

119 Antimicrobial susceptibility testing of the isolated microorganisms was carried out to evaluate the activity of conventional 120 antibiotics against the isolated bacteria and fungi strains. The bacteria contaminants isolated from these herbal 121 preparations showed wide resistance to penicillins, especially ampicillin, augmentin (amoxycillin-clavulanic acid 122 combination) and cloxacillin, suggesting that they could be producers of penicillinases. The resistance to trimethoprim-123 sulphamethoxazole (co-trimoxazole) by all the isolates especially the Gram-negative isolates calls for attention. The findings of this study agree with an earlier work [12]. Staphylococci strains were the most frequently isolated bacteria 124 species and it probably originated from handlers, as its habitat is human skin. Staphylococcus showed wide resistance to 125 126 penicillins suggesting possibly that they are producers of penicillinase enzymes. Resistance to trimethoprim by S. aureus and S. epidermidis has been reported with increasing frequency [18,19, 20]. It seems probable that S. epidermidis serves 127 128 as a reservoir for resistance, which can be transferred to S. aureus. Also, inter-generic transfer of resistance among 129 different genera of Gram-positive cocci and between Bacillus species and Staphylococci and Streptococci has been reported [20, 21]. Escherichia coli were the second most frequently isolated species in these medications which is an 130 intestinal bacterium and an indicator of faecal contaminant. Presence of Escherichia coli in the sample indicates poor 131 hygiene practices and lack of adequate handling of the products. According to the European pharmacopoeia 2007 [22], no 132 Salmonella spp or Escherichia coli strain should be present in oral medicines. The presence of E. coli in herbal drugs had 133 been reported by another researcher [23]. The Escherichia coli isolates showed a wide resistance to ampicillin, 134 135 ceftazidime, sulphamethoxazole-trimethoprim, amoxycillin-clavulanic acid, and tetracycline. Bacillus spp. were the third

<sup>116</sup> Bacillus spp, S = Salmonella, P = Proteus spp, Y = Yesinia spp, C = Corynebacteruim spp. The numbers attached

136 most frequently found in these herbal medicaments because they are widely distributed in the soil, dust, air and because 137 they are resistant to environmental destructive factors [20, 24]. A number of reports have described serious human 138 infections caused by members of the genus Bacillus even though they have been regarded as non-pathogenic [25,26,27]. All the strains of *Pseudomonas* isolated were resistant to β-lactam antibiotics; Inducible β-lactamase activity is a general 139 140 property of Pseudomonas cepacia [28]. Gram negative rods usually have wide resistance against antimicrobial agents [20] (Esimone et al., 2007a). Streptococcus spp showed high resistance to sulphamethoxazole-trimethoprim and 141 142 ampicillin. Salmonella spp, were resistant to sulphamethoxazole-trimethoprim, amoxycillin clavulanic acid, and ampicillin. Proteus spp. Yersinia spp and Corynebacteruim diphtheria showed wide resistance to sulphamethoxazole-trimethoprim. 143 amoxycillin clavulanic acid, ceftazidime and ampicillin (Table 3). On the other hand, the bacterial isolates were susceptible 144 145 to some groups of the antibiotics (ofloxacin, ciprofloxacin, gentamicin and ceftriaxone).

146 Multiple Antibiotic Resistance Index (MARI) evaluation (Table 5) revealed that species of Escherichia coli showed high 147 level of multiple antibiotic resistances to the panel of antibiotics used in this study. The MARI value ranged from 0-5 -0.7, 148 with three (30%) resistant to seven antibiotic out of the ten used, six (60%) resistant to six of the antibiotics used and two 149 (20%) resistant to five. Staphylococcus spp have MARI values ranging from 0.4-0.7, with one (8.3%) resistance to seven 150 antibiotic, four (33.3%) resistant to six antibiotics, six (50%) resistant to five antibiotics and one (8.3%) resistance to four 151 antibiotics. The MARI result of Bacillus spp ranged from 0.4 - 0.6, with five (62.5%) resistant to six antibiotics, One 152 (12.5%) being resistance to five, four and three antibiotics each. Proteus spp MARI value is from 0.4 - 0.7, with one (25%) each of the four isolates resistant to seven, six, five and four respectively. Pseudomonas spp had MARI value ranging 153 from 0.4 - 1.0, one (25%) showed high resistance index, being resistance to ten of the antibiotics used in this study, one 154 (25%) resistant to five antibiotics and two (50%) resistance to four antibiotics. The three species of Streptococcus isolated 155 showed MARI values from 0.3-0.8, that is, one (33.3%) resistant to eight antibiotics, one (33.3%) to four antibiotics and 156 one (33.3%) to three out of the ten antibiotics. We had two isolates of Yersinia spp and the MARI values are 0.5 and 0.6. 157 158 Lastly, Corynebacteruin diphtheria isolate is resistance to five antibiotics out of the ten antibiotics used in this study. 159 Bacteria with high MAR index originate from the environment where antibiotics are over used [29].

- 160 Fungal infections are becoming an increasing cause of morbidity and mortality especially among immunocompromised patients. With the increased incidence of systemic fungal infections and the growing number of antifungal agents, 161 162 laboratory methods to guide and select antifungal therapy have gained greater attention. However, determining antifungal 163 susceptibilities of filamentous fungi is fraught with difficulties associated with slow growth of filamentous forms and 164 subjectivity of interpreting visual endpoints [30]. In the present study, antifungal susceptibility testing of 40 fungi isolates 165 was observed against five common antifungal agents (griseofulvin, nystatin, ketoconazole, clotrimazole, and fluconazole) using disc diffusion method, presence of inhibition zone was considered as sensitive while absence of inhibition zone was 166 recorded as resistance. The fungi isolates were very sensitive to nystatin, ketoconazole and clotrimazole. Most of the 167 168 fungi isolates were resistance to fluconazole while all are resistant to Griseofulvin (Table 5).
- The importance of surveying resistant environmental strains is that under favourable situations, they may transfer their 169 170 resistance plasmids to pathogens [31,32]. If such organisms are present in medicaments, such as herbal medicinal 171 products they could behave as opportunist pathogens and initiate an infection, particularly in immuno-compromised patients as well as lead to transfer of antibiotic resistance traits to hitherto sensitive microorganisms co-habiting within the 172 173 consumers of those products. Given the increasing rate of development of resistant bacteria strains, our challenge is to 174 slow the rate at which resistance develops and spreads. In order to decrease the spread of resistance among antibiotics, 175 physicians, pharmacists, researchers and consumers alike need to be more aware of the selective pressures driving these bacteria to decrease their susceptibility [33]. These selective pressures include the abuse, overuse and misuse of 176 177 antimicrobials in therapy, improperly manufactured and mishandled HMPs [13, 34] as well as other numerous 178 socioeconomic factors that govern the development of multi-drug resistant bacteria strains [35]. 179

#### 180 **4. CONCLUSION**:

The high rate of resistance to antimicrobial agents of microbial strains isolated from these herbal preparations may indicate a widespread antibiotic resistance among microorganisms from different sources. It is therefore mandatory that herbal medicines should not be taken indiscriminately and that current good manufacturing practices (cGMPs) must be observed by these herbal practitioners in the production of the medicines.

# 185 Authors' Contributions:

Oli Angus N - drafting of the manuscript/Corresponding author, Esimone Charles O – conceptualized and designed the work as well as revising the manuscript critically for important intellectual content, Ujam Nonye T – carried out the experiments and did analysis and interpretation of data, Adikwu Michael Umale - revised the manuscript critically for important intellectual content, Ikegbunam MN- carried out some part of the experimental work

190 **Competing Interest:** Authors have no competing interests to declare

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