

Original Research Article

Characteristics and Antibioqram Studies of Bacteria Associated with Vegetables Stored In Raffia Baskets in Nigeria

ABSTRACT

Raffia baskets are used in rural communities to preserve fresh vegetables for a given period of time. This method of storage is aimed at extending the shelf life of the vegetables until they are used. Thus, this study was aimed at determining the microbial succession during storage and susceptibility pattern of bacterial isolates from stored leafy vegetables to various antibiotics. Five (5) leafy vegetables namely, Bitter leaf (*Vernonia amygdalira*), Water leaf (*Talinum triangulare*), Fluted pumpkin leaf, (*Telfairia occidentalis*), Okazi leaf (*Gnetum africana*) and Scent leaf (*Ocimum gratissimum*) were obtained from the Nigerian Stored Products Research Institute (NSPRI) farm, Port Harcourt. These vegetables were stored in a raffia basket for fourteen (14) days while another set of the vegetables were left in the open air as control. This set up was monitored for changes to occur. Standard microbiological techniques were employed for the various analyses of the vegetable samples. Both cultural and molecular characterizations of the isolates were done. The disc diffusion method was used in determining the susceptibility pattern of the bacterial isolates. The results of the predominant microorganisms identified were of the genus; *Bacillus*, *Pseudomonas*, *Enterococcus*, *Enterobacter*, *Bordetella*, *Staphylococcus*, *Myroides*, *Escherichia*, *Serratia*, *Micrococcus* and *Acetobacter*. Also, *Bacillus* species occurred in all the vegetables while *Enterococcus faecalis*, *Acetobacter orientalis*, *Bordetella pertussis*, *Myroides xuanwuensis* and *Bacillus flexus* were isolated on the third day of storage from the vegetables. The total heterotrophic bacterial counts ranged from 1.8×10^6 cfu/g to 1.25×10^7 cfu/g (bitter leaf), 9×10^5 cfu/g to 9.0×10^6 cfu/g (Scent leaf), 7×10^5 cfu/g to 1.88×10^7 cfu/g (Okazi), 3.2×10^6 cfu/g to 1.05×10^7 cfu/g (Fluted pumpkin leaf) and 8.4×10^6 cfu/g to 2.04×10^7 cfu/g (water leaf). Antibioqram of bacterial isolates revealed that they were resistant to Augmentin and Ceftazidime according to Clinical Laboratory Standard Institute schemes. Combination of the antibiotics showed that most of the isolates were 100% susceptible to ciprofloxacin + erythromycin, and ciprofloxacin + ceftriaxone. The study revealed that for all the vegetables, those stored in the basket lasted longer and were fresher compared to those kept in the open space. This study revealed that the raffia baskets are suitable for storage and preservation of leafy vegetables, but microbial succession occurred as a result of increase in the period of storage leading to the proliferation of microorganisms. This could be from aerosols or by chance inoculation, through handling or materials used or other microorganisms which are inherent in the vegetables because of their contact with the soil or water used in the washing of the leaves

Keywords: Bacterial isolates, Raffia baskets, Antibiotic susceptibility, Leafy vegetables, NSPRI,

1. INTRODUCTION

Vegetables generally are good sources of nutrients such as minerals, oil, vitamins and carbohydrates [1]. Fresh vegetables contribute immensely in human nutrition as sources of vitamins (A, C, B6, thiamin, niacin, and thiamin), minerals and dietary fiber. Vegetables lower the risk of diseases such as cancer because of its composition as it

contains flavonoids, carotenoids, polyphenols and phytonutrients [2]. Vegetables exist as either leaves or fruits. Commonly consumed leafy vegetables in Nigeria include; *Vernonia amygdalifera* (Bitter leaf), *Talinum triangulare* (Water leaf), *Telfairia occidentalis* (Fluted pumpkin leaf), *Gnetum africana* (Okazi leaf), *Ocimum gratissimum* (scent leaf), *Piper guineense* (Uziza leaf) and *Gongronema latifolium* (Utazi) while fruity vegetables include carrots, tomatoes, ginger, garlic etc. [3]. They are used in preparing different delicacies such as soups and salads. Other vegetables include those of the *Alliaceae* family such as shallots, leek, garlic and chives and the cruciferous vegetables that belong to the *Brassicaceae* family which include cabbage, cauliflower, kale and broccoli, and those that belong to *Cucurbitaceae* family include squash, pumpkin, cucumber, melon and bitter gourd. They are all rich in carotenoids and tocopherols [4]. They are more nutritive compared to cereals because of the mineral content [5].

Vegetable production has been on-going in Nigeria for a long time, providing sources of income for a teeming population especially in the rural/peasant farming communities. All fresh vegetables, like most horticultural produce, is-are high in water content and are subject to desiccation (wilting or shrinking) and mechanical injury. They are also prone to bacterial and fungal attack followed by pathological breakdown during harvesting and storage [6]. Physiological processes associated with vegetable spoilage and quality losses include respiration rate, ethylene production, rate of compositional changes (which cause changes in color, texture, flavours and nutritive values), mechanical injuries, water stress, sprouting and rooting [7]. The rate of biological deterioration in vegetables depend on several environmental factors such as temperature, relative humidity (RH), air velocity, atmospheric composition (oxygen, carbon dioxide and ethylene) as well as handling. These spoilage factors impact on vegetable at post-harvest or during storage when they are not protected or subjected to conditions that are not good enough for safety of the products

Vegetables spoilage can result to changes or hydrolysis or decomposition of matrices by microorganisms resulting in unpleasant disorder of the vegetable due to toxic substances released in the vegetable material resulting in pigmentation, degeneration and discoloration of the vegetable materials. These physical abnormalities can make the vegetables inedible for humans or- and unacceptable to the public or in some cases may pose a public health risk to consumers due to the high microbial load or pathogens associated with spoiled vegetables. In Nigeria, vegetables are chewed raw or heated lightly in order to retain its organoleptic properties, and this has in most cases been responsible for food-borne infections [8], because in the process of harvesting, disease-causing microorganisms may contaminate these vegetables through contact with organic wastes such as faecal matter, organic manure, sewages, untreated irrigation water or surface waters used as nutrients or composts to improve soil quality [9].

The water sprinkled on the vegetables to keep them fresh may also be a source of microbial contamination [10] depending on the source of the water used. Antinutritional factors such as hydrogen cyanide, phytates and oxalates are

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50 able to influence the metabolic processes needed for the increase and bioavailability of nutrients in vegetables [11]. Most
51 vegetables are used as nutrients in flavoring soups and sauces. Vegetables have pleasant aroma when added to soups
52 and sauces, also contribute to the protein and essential minerals. Subjecting them to heat in the form of cooking in soups
53 and sauces increases their digestibility. Their presence or use in foods preparation introduces a variety in the flavors
54 obtained from the same foodstuff and enhances nutritional value of the food. Therefore, vegetables have great potential
55 as key protein and mineral sources which are basic ingredients for food supplementation.

56 In most rural areas and some urban centers, leafy vegetables are stored in covered baskets made with bamboo sticks
57 and wrapped with jute bags. Most times the vegetables are exposed to dews overnight for preservation and when they are
58 left exposed for long time, spoilage occurs. The wetting of the basket and evaporation from the basket gives a cooling
59 effect to the inside of the basket making it conducive for survival of the leaves for a longer period of time compared with
60 when they are left in the open [12]. This study therefore aims to determine the bacterial characteristics and antibiotic
61 susceptibility of associated isolates from some selected leafy vegetables stored in raffia basket as a means of
62 preservation. This artisanal method using rudimentary materials will enable consumers and farmers preserve their
63 vegetables in local communities to avoid post-harvest losses.

65 **Materials and Methods**

66 **Collection of Vegetable**

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68 Five (5) leafy vegetables which include; *Vernonia anydalira* (Bitter leaf), *Talinum triangulare* (Water leaf), *Telfairia*
69 *occidentalis* (Fluted pumpkin leaf), *Gnetum africana* (Okazi leaf), *Ocimum gratissimum* (scent leaf) were obtained from the
70 Nigerian Stored Product Research Institute (NSPRI) farm in Port Harcourt. The vegetables were transported to the
71 Microbiology laboratory for analyses.

72 **Preparation of Vegetable Samples**

73 The vegetables were prepared and preserved in raffia baskets; the baskets were occasionally sprinkled with about one
74 liter of sterile water with the aid of a water sprinkler. This was done to keep the inner part cold without the vegetables
75 being wet. This provides a humid environment for the leaves keeping them fresh for longer than when left in the open. The
76 setups are illustrated below in Plates 1-15;

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78 **Plate 1: Day 1 Bitter Leaf**

79 **Plate 2: Day 3 Bitter leaf**



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81 **Plate 3: Day 6. Bitter leaf (Spoiled)**

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Plate 4: Day 1 Stored pumpkin leaf



Plate 5: Day 3 Stored pumpkin leaf



Plate 6: Day 7 Stored pumpkin leaf

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Plate 7: Day 1 Stored Scent leaf



Plate 8: Day 3 Stored Scent leaf



Plate 9: Day 7 Stored Scent leaf

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Plate 10: Day 1 Stored Okazi leaf



Plate 11: Day 3 Stored Okazi leaf



Plate 12: Day 7: Stored Okazi leaf

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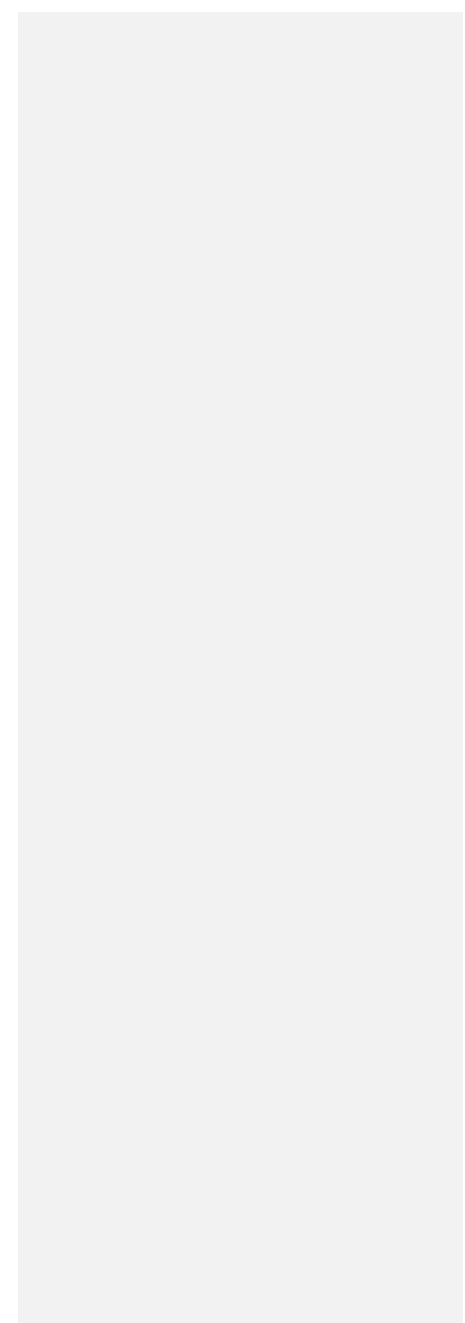




Plate 13: Day 1: Stored water leaf

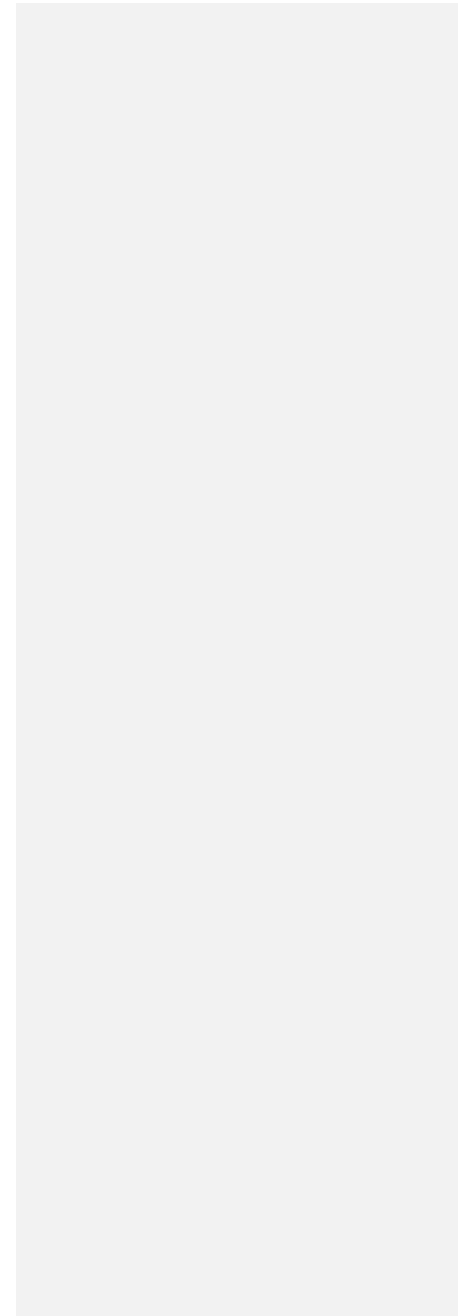


Plate 14: Day 3: Stored water leaf



Plate 15: Day 7: Stored water leaf

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134 **Microbiological Analysis**

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136 Ten grams (10g) of each vegetable leaf was homogenized in 90ml of sterilized normal saline solution in 200ml conical
137 flasks, after which the homogenized samples were diluted serially to achieve the 10^{-5} dilution [13].

138 **Inoculation and Enumeration of colonies on plates**

139 Aliquots (0.1 ml) of appropriate dilutions were transferred to plates of surface dried Nutrient agar (NA) media in duplicates
140 and inoculated by spreading with flamed glass spreader and incubated at 37°C for 24 hours. After incubation, colonies
141 that appeared on the plates were counted and the mean expressed as cfu/g (or as \log_{10}/g) for the samples. This was
142 carried out on each of the vegetables at two days interval until spoilage occurred. Counts for each sample were then
143 calculated using the formula below [14]

$$144 \text{ THC (cfu/g)} = \frac{\text{Number of Colonies}}{\text{Dilution (10}^{-6} \times \text{Volume plated (0.1ml))}}$$

146 **Maintenance of pure cultures**

147 Discrete bacterial colonies that developed on the Nutrient agar plates were sub-cultured using streaked plate method by
148 transferring each colony of same morphology onto freshly prepared NA plates under strict aseptic conditions. This was
149 incubated at 37°C for 24 hours. The bacterial cultures were then maintained according to the method reported by Amadi
150 *et al.* [15] using 10% (v/v) glycerol suspension at -4°C.

151 **Biochemical Characterization and Identification of Isolates**

152 The isolates were characterized based on their appearance on the culture media which include; shape, colour, wetness,
153 and Gram's reaction. Biochemical tests such as; Indole production, motility, coagulase, Methyl Red, citrate utilization,
154 sugar fermentation (sucrose, glucose, lactose, and maltose) were carried out based on the method of Cheesbrough [16].
155 To further confirm the bacterial isolates, molecular identification techniques were carried out. The molecular techniques
156 were carried out as described by Wemedo and Robinson [17].

157 **Preparation of Standard Bacterial Inoculum**

158 Loop-full of 24-hour old pure culture of the test organisms from nutrient agar plates, were transferred into sterile diluents
159 tubes and adjusted to 0.5 McFarland Turbidity Standard [16,17]. This was used as the standard inoculum.

160 **Antimicrobial susceptibility testing**

161 The antibiotic sensitivity of the bacterial isolates was determined using the disc diffusion method [18]. Multiple antibiotic
162 sensitivity testing was also carried out using the method adopted by Ogbonna and Inana [19]. Standardized inoculum of

163 24-hour old cultures were spread on Mueller-Hinton agar plates using sterile swab sticks. The plates were dried at room
164 temperature for 1 hour before placing the antibiotic discs at the centre. The plates were then incubated at 37°C for 24
165 hours and the diameter of zone of inhibition was measured in millimetre. Organisms were classified as sensitive,
166 intermediate or resistant, based on the Clinical and Laboratory Standard Institutes.

167 RESULTS AND DISCUSSION

168 Bacterial counts

169 The results of total heterotrophic bacterial counts of the stored vegetable leaves from day one to the day that spoilage
170 occurred are presented in Table 1. The result shows that water leaf had the highest bacterial load followed by pumpkin
171 leaf, Okazi Leaf while bitter leaf had the lowest bacterial counts after sixteen days of storage. The total heterotrophic
172 bacterial counts from day 1 to the day spoilage occurred in the five selected vegetable ranged from 1.8×10^6 cfu/g to 1.25
173 $\times 10^7$ cfu/g (bitter leaf), 9×10^5 cfu/g to 9.0×10^6 cfu/g (Scent leaf), 7×10^5 cfu/g to 1.88×10^7 cfu/g (Okazi), 3.2×10^6 cfu/g to
174 1.05×10^7 cfu/g (Pumpkin leaf) and 8.4×10^6 cfu/g to 2.04×10^7 cfu/g (water leaf). The bacterial counts increased as the
175 spoilage occurred with increase in the number of days respectively. For example, the highest microbial counts for
176 pumpkin leaf was observed on day five which was the day spoilage began while for scent leaf spoilage occurred on day 7,
177 bitter leaf was on day 5 while water leaf spoiled on day 5 and Okazi spoiled on day 12 (Figs 1-5) respectively.

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Table1: Total Heterotrophic Bacterial counts of the Vegetables (cfu/g)

	PUMKIN LEAF		SCENT LEAF		WATER LEAF		OKAZI LEAF		BITTER LEAF	
	Sample	control	Sample	control	Sample	control	Sample	Control	sample	control
Day1	2.34×10 ⁶	2.45×10 ⁶	1.78×10 ⁵	7.59×10 ⁵	1.55×10 ⁶	2.19×10 ⁵	2.04×10 ⁶	2.34×10 ⁶	6.03×10 ⁵	7.94×10 ⁵
Day3	4.36×10 ⁶	4.36×10 ⁶	2.19×10 ⁵	1.78×10 ⁶	1.48×10 ⁷	3.72×10 ⁸	1.74×10 ⁶	4.37×10 ⁶	3.98×10 ⁵	1.48×10 ⁶
Day5	5.25×10 ⁶	1.26×10 ⁸	3.39×10 ⁵	1.58×10 ⁷	1.82×10 ⁷	Sp	1.55×10 ⁶	5.25×10 ⁷	3.47×10 ⁶	1.82×10 ⁷
Day7	8.32×10 ⁶	Sp	4.47×10 ⁵	Sp	7.94×10 ⁷	Sp	2.24×10 ⁶	8.32×10 ⁷	2.88×10 ⁶	7.94×10 ⁷
Day9	Sp	Sp	7.59×10 ⁵	Sp	Sp	Sp	8.32×10 ⁶	2.34×10 ⁸	1.48×10 ⁷	sp
Day12	Sp	Sp	Sp	Sp	Sp	Sp	1.38×10 ⁷	Sp	4.37×10 ⁷	sp
Day14	Sp	Sp	Sp	Sp	Sp	Sp	2.82×10 ⁷	Sp	sp	sp
Day 16	Sp	Sp	Sp	Sp	Sp	Sp	sp	Sp	sp	sp

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Table 2: Biochemical Characteristics of Isolates

Colonial/cell characteristics	Gram reaction	Coagulase	Tests								Sugar fermentation					Probable organism
			Catalase	Oxidase	Indole	MR	VP	Citrate	Motility	Sucrose	Mannitol	Lactose	Glucose	Galatose	Fructose	
Creamy, Round, Opaque, Entire Small, Elevated, Moist, rods	+ve	-	+	-	-	+	-	+	-	+	-	+	+	-	-	<i>Bacillus</i> sp.
Creamy, Circular, Flat, Opaque, Entire, Small, moist, rods	-ve	-	-	+	+	-	+	-	+	-	-	+	-	-	+	<i>Acetobacter</i> sp.
Creamy, Circular, Flat, Opaque, Entire, Moist, rods	-ve	-	-	+	+	-	+	-	+	-	-	+	-	-	+	<i>Enterobacter</i> sp.
Greenish yellow, Smooth rods	-ve	-	+	+	-	-	-	+	+	-	+	-	+	-	+	<i>Pseudomonas</i> sp.
Red, Round, flat, Muroid, Large, Translucent, Serrated rods	-ve	-	+	-	-	-	+	+	+	+	+	-	+	-	-	<i>Serratia</i> sp.
Golden yellow, Raised, Translucent, Moist cocci	+ve	-	+	-	-	+	+	+	-	+	+	+	+	+	+	<i>Staphylococcus</i> sp.
Smooth, Creamy, Raised, Translucent rods	-ve	-	+	-	+	+	-	-	+	+	+	+	+	+	-	<i>Escherichia coli</i>
Yellow, Round, Opaque-, Small	+ve	-	+	-	-	+	+	-	-	+	+	+	-	+	+	<i>Micrococcus</i> sp.
yellow, Round, Smooth rods	-ve	-	+	+	-	-	+	+	-	-	-	-	-	-	-	<i>Myroides</i> sp.
Small, Rough, Flat, rods	-ve	-	+	+	-	-	-	-	-	-	-	-	-	-	-	<i>Bordetella</i> sp.
Smooth, Creamy, Entire, cocci	+ve	-	-	-	-	-	+	-	-	+	+	+	+	-	+	<i>Enterococcus</i> sp.

Key: -ve = Gram negative, +ve =Gram positive, - = Negative, + = Positive.

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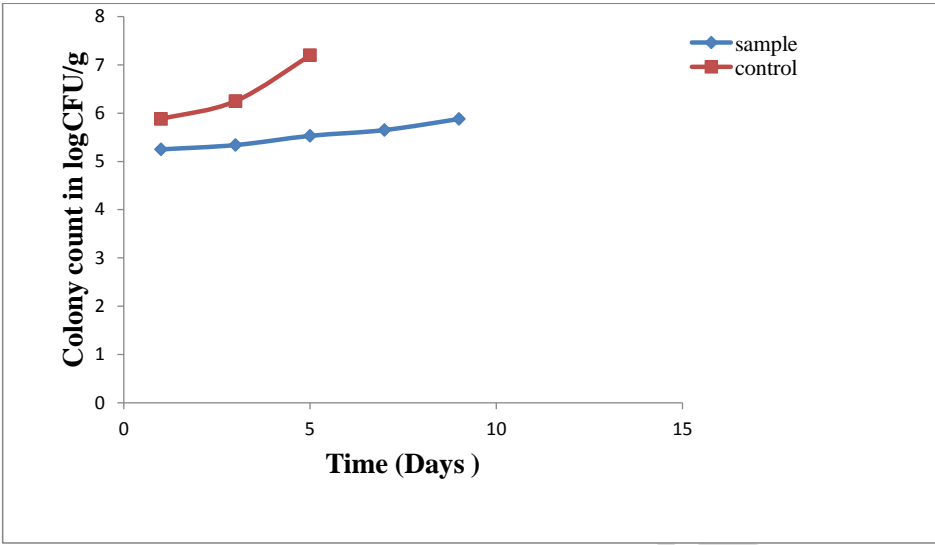


Fig. 1: Bacterial counts and shelf life for pumpkin leaf (*Teferia occidentalis*).

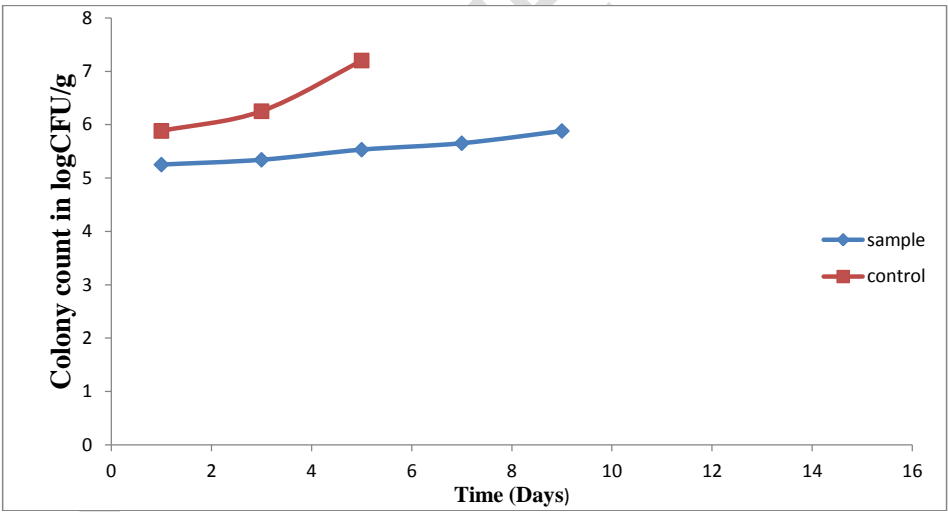


Fig. 2: Bacterial counts and shelf life for Scent (*Ocimum gratissimum*)

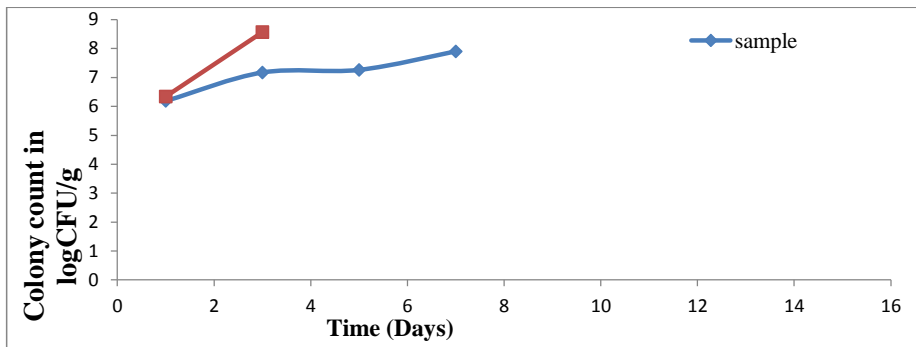


Fig.3: Bacterial counts and shelf life for Waterleaf (*Talinum triangulare*).

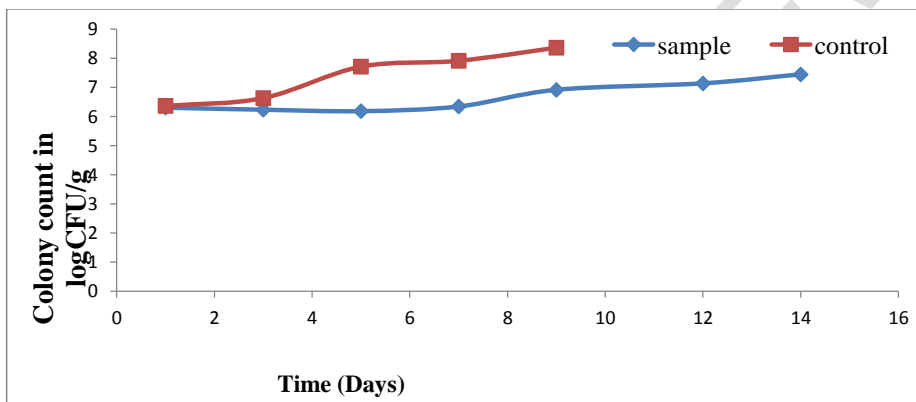
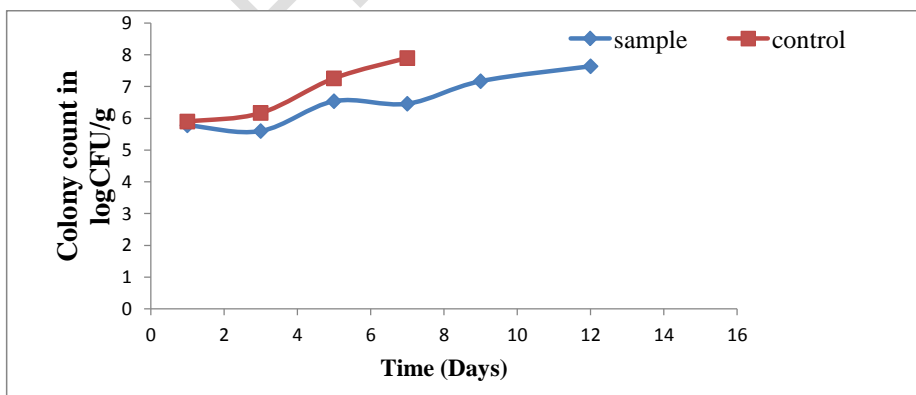


Fig.4: Bacterial counts and shelf life for Okazi (*Gnetum africanum*)



230 **Fig.5: Bacterial counts and shelf life for Bitterleaf (*Veronia amygdalina*).**

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232 The results of total heterotrophic bacterial counts obtained in this study are in agreement with the counts reported by [7].

233 The results obtained show that the bacterial counts increased as the storage in the baskets was extended to the day

234 spoilage occurred. It was also observed that the control samples had a higher bacterial load and lasted for few days

235 before spoilage occurred compared to the vegetables stored in the vegetable basket. The storage in this circumstance

236 depends on chance inoculation from the environment. The storage conditions are inconsistent and were not optimized so

237 the bacterial isolates could be distinguished as contaminants and could cause spoilage within a short period as against

238 the vegetables preserved in a raffia basket [20].

239 **Table 3: Bacterial Succession during the period of storage in raffia baskets**

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Sample	Day 1	Day 3 to Day 15.
Pumpkin	<i>Bacillus</i> sp.	<i>Acetobacter orientalis</i>
	<i>Micrococcus</i> sp	<i>Bordetella pertussis</i>
	<i>Escherichia coli</i>	<i>Staphylococcus</i> sp
Water leaf	<i>Bacillus</i> spp.	<i>Myroides xuanwuensis</i>
	<i>Micrococcus</i> sp	<i>Enterobacter xiangfangensis</i>
	<i>Escherichia coli</i>	
Scent leaf	<i>Bacillus</i> sp	<i>Enterococcus faecalis</i>
	<i>Micrococcus</i> sp	<i>Escherichia coli</i>
Okazi leaf	<i>Bacillus</i> sp	<i>Myroides</i> sp
	<i>Pseudomonas</i> sp	<i>Bacillus flexus</i>
		<i>Bacillus subtilis</i>
Bitter leaf		<i>Serratia</i> sp.
	<i>Bacillus</i> sp	<i>Pseudomonas</i> sp
	<i>Staphylococcus</i> sp	<i>Micrococcus</i> sp
		<i>Enterococcus faecalis</i>

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242 The predominant bacterial genera identified were *Bacillus*, *Pseudomonas*, *Enterococcus*, *Enterobacter*, *Bordetella*,

243 *Staphylococcus*, *Myroides*, *Escherichia*, *Serratia*, *Micrococcus* and *Acetobacter* (Table 2). Also, *Bacillus* sp occurred in all

244 the vegetables during the period of the study. This organism is a normal flora of the soil environment, with the potential to

245 remain in the environment for a longer time because of its ability to produce endospores [21]. However, *Enterococcus*

246 *faecalis*, *Acetobacter orientalis*, *Bordetella pertussis*, *Myroides xuawensis* and *Bacillus flexus* were isolated in the stored

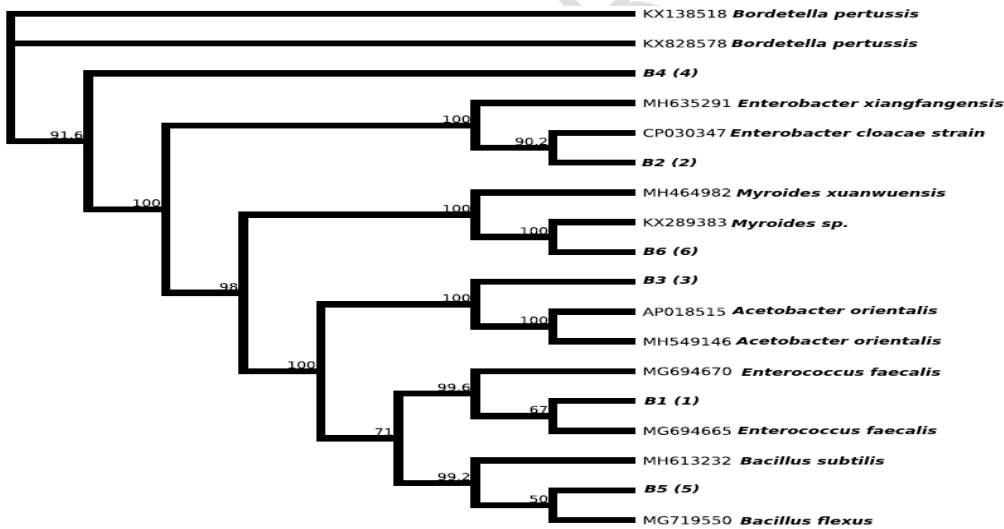
247 vegetables from the third day, in most of the vegetables. The molecular characterization of the bacterial isolates showed

248 that they have relatedness to the bacterial presented in the phylogenetic tree (Fig 6). This result is in agreement with the

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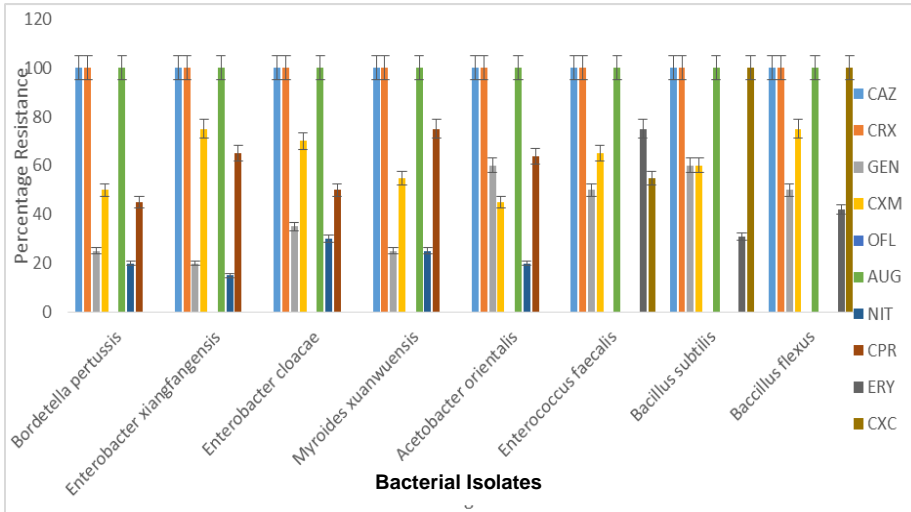
249 report of Mandrell *et al.*, [22] who reported similar bacterial isolates in their research. Bacteria, belonging to the genus
 250 *Bacillus* are spore formers and often heat-resistant. They are able to cause opportunistic infections such as endocarditis,
 251 bacteremia, wound abscesses ~~ete~~ among others [23,24]. *Myroides* are able to cause skin infections in patients with
 252 diabetes as well as urinary tract infections [25]. *Bordetella pertussis* is the causative agent of whooping cough while
 253 *Enterobacter* causes meningitis, pneumonia, bacteremia as well as urinary tract infection. *Enterococcus faecalis* are
 254 emerging as hospital pathogens and are mostly found in the gut. They are able to produce a cytolyisin (streptolysin) that
 255 affects the gut as well as has been implicated in multi-drug resistance [26]. Hence it is important to improve the quality
 256 and processing by optimizing the process conditions or even manipulating process factors such as temperature, humidity,
 257 aeration and pH [20]. Against the background of the significance and benefits of this process in this study, it is desirable to
 258 improve the art of the storage conditions to improve the shelf life of vegetables. As of now, the process is still largely a
 259 traditional art and the characteristic features of preservation is lacking. This practice predisposes the process of
 260 preservation to contamination because the raffia baskets have tendencies to accumulate dust, although presumably
 261 provides warmth and humid environment.

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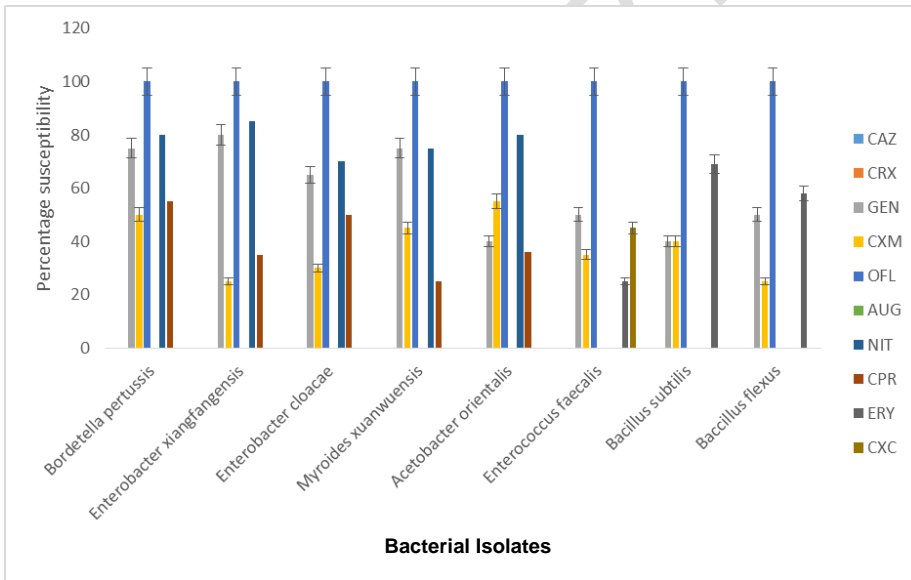


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Fig 6: Phylogenetic Tree showing the evolutionary distance between the bacterial Isolates



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Figure 7: Percentage resistance of the isolates to different antibiotics



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Fig 8: Percentage susceptibility of the isolates to different antibiotics

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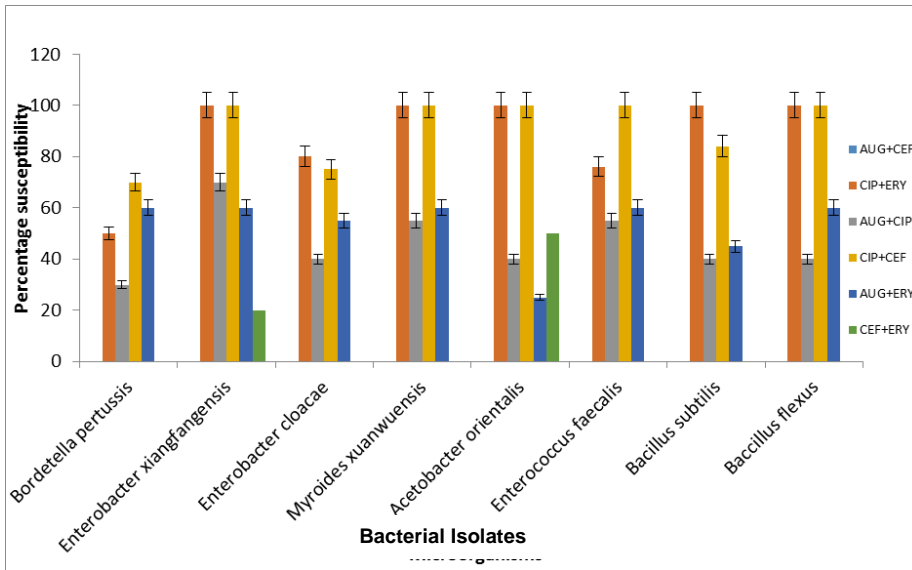


Fig 9: Combined Antibiotic Sensitivity pattern of Isolates from the five selected Vegetables

All the bacterial species were susceptible to Ofloxacin, Ciprofloxacin + Erythromycin, Augmentin + Ciprofloxacin, and Ciprofloxacin+ Ceftriaxone while all were resistant to Augmentin and Ceftazidime. The antibiotic profile of the isolates and identified bacteria revealed that ciprofloxacin, ofloxacin, were more susceptible, hence, they were considered as drugs of choice for infections caused by bacteria isolated in this study. Comparatively, the results of antibiotic profile tests against the isolates and bacteria reported in this study is in agreement with observations by Wemedo and Robinson [17] who reported the resistance of some bacterial isolates against Augmentin. The resistance of the microbes to Augmentin and Ceftazidime may be due to a regular abuse of the drugs against the isolates *in vitro*.

Conclusion

The results of the bacterial counts obtained in this study revealed that, bacterial load of vegetables preserved in the Raffia baskets were lower than those kept in an open air and as spoilage began to occur, more bacteria were isolated and the bacterial load increased with increase in storage time. It was also observed that, the control samples had a higher bacterial load and lasted for fewer days before spoilage occurred compared with the vegetables stored in the vegetable basket that had fewer organisms and lasted longer. The antibiogram of the bacterial isolates revealed that all the

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297 bacterial species were susceptible to Ofloxacin, Ciprofloxacin + Erythromycin, Augmentin + Ciprofloxacin, and
298 Ciprofloxacin+ Ceftriaxone while all were resistant to Augmentin and Ceftazidime.

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303 of fungi associated with the spoilage of sweet orange (*Citrus sinensis*) fruits in Sokoto State. *Nigerian*
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