Microbial Assessment of Foods and Currencies from Street Food Vendors and Antibiotic Resistance **Profile of Isolates**

ABSTRACT

Aims: Street food vendors handle money and food simultaneously, creating possibilities of cross contamination. This study focused on determining the microbial loads of street vended foods and currency denominations received as change; identify and determine the antibiotics resistance profile of the isolates.

19 Materials and Methods: Street vended food samples and currency (10 - 500 Naira) notes from four 20 vendors (FV) were collected. Microbial loads of food and money, biochemical (MRVP, motility, carbohydrate fermentation) and antibiotic testing of isolates were performed using standard methods. 21 Results: TVC value ranged from (1.0×10⁷ to 1.7×10⁸ CFU/g); Staphylococcal (5.8×10⁶ to 1.6×10⁷ 22 CFU/g), Salmonella-Shigella (NG to 1.2×10⁷ CFU/g) and Coliform (no growth) from FV1. TVC ranged 23 from (4.2×10⁶ to 2.0×10⁷ CFU/g); Staphylococcal (1.2×10⁶ to 2.0×10⁷ CFU/g), Salmonella-Shigella (1.6×10⁶ to 1.1×10⁷ CFU/g) and Coliform (NG to 1.5×10⁵ CFU/g) from FV2. From FV3 TVC ranged 24 25 from $(9.3 \times 10^6$ to 4.6×10^7 CFU/g); Staphylococcal $(1.9 \times 10^6$ to 5.1×10^6 CFU/g), Salmonella-Shigella $(1.3 \times 10^6 \text{ to } 5.9 \times 10^6 \text{ CFU/g})$ and Coliform $(5.0 \times 10^3 \text{ to } 1.0 \times 10^6 \text{ CFU/g})$. FV4 load ranged from $(2.8 \times 10^6 \text{ CFU/g})$. 26 27 to 1.3×10⁸ CFU/g); Staphylococcal (1.7×10⁶ CFU/g to TNTC), Salmonella-Shigella (NG to 5.7×10⁷ 28 CFU/g) and Coliform (6.5×10⁴ to 6.8×10⁵ CFU/g). Eleven genera were identified from food and 29 currencies including: Staphylococcus, Klebsiella, E. coli and Enterobacter. Organisms showed varied 30 resistance patterns to the different antibiotics screened. About (62.5 %) of isolates were resistant to 31 Ampicillin (10 µg) and (61.5 %) Ceftazidime (30 µg). 32

33 Conclusion: In conclusion, there are possibilities of cross contamination between food and 34 currencies making the safety of consuming the foods a public health concern. 35

Keywords: Street foods; food; money; simultaneous handling; safety.

1. INTRODUCTION

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Street foods are ready-to-eat foods and beverages prepared and/or sold by vendors on streets and 42 43 other public places [1]. Street foods are sources of nutrition for many, at affordable prices. Some of 44 the popular street foods sold in Nigeria include: rice, beans, akara (fried bean cake), boli (roasted plantain), roasted corn, amala (yam flour meal), garri, fufu (cassava meal), fried fish, stick or peppered 45 46 meat, suya (peppered stick meat), fruits, salads and more. The street food industry plays an important 47 role in developing countries in meeting the food demands of the urban dwellers [2]. There is a noticeable increase of food vendors in Nigeria as a result of dwindling economy and unemployment 48 [2]. Types of vending sites encompass stalls, a variety of push-carts, roadside stands, and hawkers 49 depending upon the ingenuity of the individual, resources available and type of food sold [3]. 50

Trading has been part of mankind from time immemorial. Items passed from hand to hand are likely to 51 be contaminated with disease causing microorganism especially if handled with unclean hands or 52 kept in dirty surroundings [4]. Paper and polymer currency notes may habour various deadly 53 54 pathogenic microorganisms, and this could represent a universal medium of transmission of bacteria 55 in the environment and among humans [5].

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56 Many of the street food vendors serve foods and collect money for the foods simultaneously and do 57 not wash or sanitize their hands between tasks [6, 7, 8]. In spite of numerous advantages offered by 58 street foods, there are several health hazards associated with this sector of the economy. There is a 59 possibility that currency notes might act as vehicles for the transmission of potential pathogenic 60 microorganisms from human and environment into foods especially during simultaneous handling of 61 food and money by street food vendors. Money on which pathogenic microorganisms might survive 62 represents an often over looked reservoir for enteric disease [9, 10,11].

Vendors often lack formal education and untrained in food hygiene; work under crude and unsanitary conditions and have very little knowledge about the causes of foodborne diseases [3]. Evidently, since street vended foods have been shown to have epidemiological links with illness [12, 13, 14, 15], it further suggests that street foods contribute immensely to food poisoning outbreaks.

Furthermore, the rise of street food vending and simultaneous handling of money by same vendors has created opportunities for contaminations, resulting in public health problems [16, 17]. Unsafe food is becoming an increasingly serious threat to public health in Nigeria.

70 Perpetual and indiscriminate administration of antimicrobial agents and antibiotics in developing 71 countries coupled with poor hygiene habits has led to the development of multiple drug resistant 72 microorganisms. Furthermore, the prevalence of antibiotic resistance among foodborne pathogens 73 has increased in recent times [18]. Failure of food service workers to adequately wash and or sanitize 74 hands between handling of money and serving of food could put consumers at risk of diarrheal 75 diseases. Therefore, the aim of this study was to assess the microbial quality of food and currency denominations from street food vendors for possibility of cross contaminations and determine the 76 77 antibiotic resistance profile of isolates. 78

79 2. MATERIAL AND METHODS

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81 2.1 SAMPLE COLLECTION

Four types of food samples namely: Ekuru, jollof rice, pounded yam with sauce and suya were procured from food (stationary and mobile) vendors in Iwo, Osun State. Samples were collected in sterile polyethylene bags and kept in an ice pack inside food flasks for transportation to Bowen University Food Science laboratory and isolation from samples were carried out within 24 hours of purchase. Table 1 describes the food categories. Also, the currency notes returned as change after purchase were placed in separate Ziploc bags. Both food items and the currency notes were used for analyses.

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92 Table 1: Description of food samples obtained during research 93

Food categories	Composition of Food in the category					
Ekuru	Uncoated bean paste, pepper sauce, oil and meat					
Jollof rice	Rice, pepper sauce, oil and meat					
Pounded Yam	Yam, vegetable stew, pepper sauce and meat					
Suya	Steak meat, ground pepper, oil and seasoning					

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95 2.2 Microbial analysis of food

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About 10 g of the food samples was homogenized in 90 mL peptone water and mixed properly. Then,
1 mL of sample was serially diluted in 0.85 % saline solution up to 10⁻⁵. Using the pour plate method,
1 mL of the last dilution was plated into plate count agar; Eosin methylene blue agar (LAB);
Salmonella- Shigella agar (SRL); Baird Parker agar; and Sabouraud dextrose agar (SDA) (Park
Scientific Limited) in duplicates and incubated at 37 °C overnight while SDA plates were incubated at
28 °C for 72-96 h. All plates with less than 300 colonies were enumerated after incubation.

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104 Microbial analysis of paper currencies

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106 Collected currency notes from each vendor were analyzed separately. Notes were separated into 107 denominations of polymer (10, 20, 50 Naira); 100 Naira; 200 Naira and 500 Naira. Each denomination 108 was aseptically placed into a new Ziploc bag. About 100 mL of peptone water was added and the 109 samples were shaken vigorously for 10 mins in order to dislodge adhering microorganisms. Further 110 serial dilution up to 10⁻⁴ was performed, plating, incubation and enumeration were as previously 111 described.

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113 2.3 Identification of isolated microorganisms 114

115 After enumeration, colonies with distinct morphological differences like color, size and shape were 116 randomly selected for further biochemical analyses. Isolates were purified by repeated plating and preserved on Nutrient agar slants and stored at 4 °C until characterization was carried out. Isolates 117 118 were characterized using Gram reaction, motility, sugar fermentation test, MR VP test, citrate 119 utilization test, starch hydrolysis test and catalase test [19].

121 2.4 Antibiotic resistance testing

122 123 A total of 13 (5 Gram positive and 8 Gram negative) isolates were randomly selected and tested for 124 antimicrobial resistance profile. About 5 mL of nutrient broth was inoculated with a loopful of the organism and incubated at 37 °C overnight to an inoculum density equivalent to a 0.5 McFarland 125 126 turbidity standard. Sterile cotton swab was moistened with each isolate and used to swab Muller-127 Hinton (Biomark, India) agar plates in duplicates. Then, the plates were left to dry for about 5 mins before placing the antibiotics discs (Rapid Labs, UK) on each of the plates as described by [20]. 128 129 Antibiotic discs and concentrations used were: Ceftazidime (CAZ) - 30 µg; Cefuroxime (CRX) - 30 130 μg; Gentamicin (GEN) – 10 μg; Cefriaxone (CTR) – 30 μg; Erythromycin (ERY) – 5 μg; Cloxicillin (CXC) -5 µg; Ofloxacin (OFL) – 5 µg; Amoxycillin/ Clavulanate (AUG) -30 µg; Ciprofloxacin (CPR) – 5 131 132 μg; Nitrofurantoin (NIT) – 300 μg; and Ampicillin (AMP) – 10 μg. Zoning diameter was taken as the 133 mean zone along the two directions on perpendicular lines using a ruler on the reverse side of the 134 plates and measured in millimeters. In addition, zone size was expressed as susceptible/resistant based on the recommendation of National Committee for Clinical Laboratory Standards Institute 135 136 (National Committee for Clinical Laboratory Standards [21].

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138 2.5 Statistical Analysis

139 Data collected from microbial load were analyzed using Statistical Package for the Social Sciences 140 (SPSS) (2011) [22]. Analysis of Variance (ANOVA) was used to evaluate significant differences and 141 separation of the mean values was carried out using Duncan Multiple Range Test at (p<0.05). 142

143 3. RESULTS AND DISCUSSION

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145 3.1. Microbial loads of food and currency denomination from food vendors 146

147 The results of microbial loads of the food samples and currency denominations (Naira) received as 148 change after purchase are presented in Tables 2. Overall, all of the samples analyzed in this study had microbial contamination. Ekuru had (1.7x10⁸ CFU/g); (1.3x10⁷ CFU/g); (1.2x10⁷ CFU/g) and 149 (1.9x10⁷ CFU/g) for total viable (TVC), Staphylococcal, Salmonella-Shigella and fungal counts 150 respectively. While Naira denominations from same vendor ranged from $(1.0x10^7 \text{ to } 1.7x10^7 \text{ CFU/g})$; 151 (5.8x10⁶ to 1.6x10⁷ CFU/g) and (1.0x10⁶ to 1.4x10⁶ CFU/g) for TVC, Staphylococcal and Fungal 152 counts respectively. There were no observable enterobacteriaceae growths on the currency 153 154 denominations.

155 Food vendor 2, jollof rice sample, TVC was (2.0 ×10⁷ CFU/g), Staphylococcal count was (2.0×10⁷ CFU/g), Salmonella-Shigella was (1.1×10⁷ CFU/g) and fungal count was (1.9×10⁷ CFU/g) as shown in 156 Table 3. While for the currency denominations obtained from the vendor, TVC ranged from (4.2 to 5.0 157 x 10⁶ CFU/g); Staphylococcal count ranged from (1.2 to 2.0×10⁶ CFU/g); Salmonella-Shigella count 158 ranged between $(1.6 \times 10^6 \text{ to } 3.2 \times 10^6 \text{ CFU/g})$. 159

There were no observable coliform growth in currency denominations except 200Naira notes which 160 161 had an average of (1.5×10⁵ CFU/g). However, fungal counts ranged between (5.7 and 7.3×10⁵ 162 CFU/q). Total viable count from pounded vam sample was (4.6×10⁷ CFU/q), the total Staphylococcal count was (5.1×10⁶ CFU/g) the total Salmonella-Shigella count was (5.9×10⁶ CFU/g), coliform count 163 164 (1.0×10^{6}) CFU/q) while the total fungi count was (8.5×10⁵ was CFU/q).

165 Table 2. Microbial load (CFU/g) of food and currency samples from food vendors

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Sample	TVC*	Staph	SS ¹	Coliforms	Fungi
FV1					
Ekuru	1.7±0.6 ^ª ×10 ⁸	1.3±0.7 ^a ×10 ⁷	1.2±0.2 ^ª ×10 ⁷	NG	1.9±0.2 ^a ×10 ⁷
Currency					
Polymer	1.7±0.5 ^b ×10 ⁷	1.4±0.9 ^a ×10 ⁷	NG	NG	1.1±0.4 ^b ×10 ⁶
200Naira	1.0±0.5 ^b ×10 ⁷	$5.8\pm0.4^{a}\times10^{6}$	NG	NG	1.0±0.3 ^⁵ ×10 ^⁵
500Naira	1.2±0.8 ^b ×10 ⁷	1.6±0.2 ^ª ×10 ⁷	NG	NG	1.4±0.1 ^b ×10 ⁶
FV2					
Jollof rice	2.0±0.3 ^a ×10 ⁷	2.0±0.2 ^a ×10 ⁷	1.1±0.2 ^ª ×10 ⁷	NG	9.3±0.2 ^a ×10 ⁶
Currency					
Polymer	4.5±1.2 ^b ×10 ⁶	1.2±0.2 ^b ×10 ⁶	3.0±0.3 ^b ×10 ⁶	NG	5.7±0.6 ^b ×10 ⁵
200Naira	4.2±0.8 ^b ×10 ⁶	2.0±0.3 ^b ×10 ⁶	1.6±0.1 ^b ×10 ⁶	1.5±0.3 ^b ×10 ⁵	7.4±0.3 ^b ×10 ⁵
500Naira	5.0±0.9 ^b ×10 ⁶	1.7±0.5 ^b ×10 ⁶	1.6±0.4 ^b ×10 ⁶	NG	$6.2\pm0.3^{b}\times10^{5}$
FV3					
Pounded yam	4.6±0.5 ^ª ×10 ⁷	5.1±0.4 ^ª ×10 ⁶	5.9±0.1 ^ª ×10 ⁶	1.0±0.8 ^a ×10 ⁶	8.5±0.0 ^a ×10 ⁵
Currency	_				
100Naira	9.3±0.4 ^ª ×10 ⁶	2.8±0.8 ^b ×10 ⁶	1.7±0.8 ^b ×10 ⁶	$1.0\pm0.0^{b}\times10^{4}$	1.4±0.1 [°] ×10 ⁵
200Naira	3.2±0.2 ^ª ×10 ⁷	2.1±0.2 ^b ×10 ⁶	5.6±0.8 ^b ×10 ⁵	$9.0\pm0.1^{b}\times10^{4}$	2.5±0.4 ^b ×10 ⁵
500Naira	2.4±0.1 ^ª ×10 ⁷	1.9±0.3 ^b ×10 ⁶	1.3±0.8 ^b ×10 ⁶	$5.0\pm0.0^{b}\times10^{3}$	1.3±0.0 ^c ×10 ⁵
FV4					
Suya	1.3±0.4 ^ª ×10 ⁸	TNTC	$5.7\pm0.6^{a}\times10^{7}$	3.1±0.6 ^a ×10 ⁵	NG
Currency					
Polymer	2.8±0.6 ^b ×10 ⁶	1.7±0.1×10 ⁶	NG	$3.7\pm0.2^{b}\times10^{5}$	ND
100Naira	6.0±0.6 ^b ×10 ⁶	TNTC	NG	$6.5\pm0.8^{b}\times10^{4}$	ND
200Naira	3.0±0.4 ^b ×10 ⁶	1.2±0.8×10 ⁷	$4.9\pm0.1^{b}\times10^{5}$	6.8±0.6 ^b ×10 ⁵	ND

167 FV= Food vendor; TVC^* = Total viable count; SS= Salmonella –Shigella; NG= No growth; ND= not determined; 168 TNTC= Too numerous to count; Polymer notes included: 10, 20 and 50 Naira notes. Values are mean \pm SD of

169 duplicate; Duncan separation of means with same alphabets are not different (P =< .05) in each column. For the change obtained from the vendor after purchase, the counts ranged from (9.3×10⁶ to 2.4×10⁷ 170 CFU/g); (1.9 to 2.8×10⁶ CFU/g); (5.6×10⁵ to 1.7×10⁶ CFU/g); (5.0×10³ to 9.0×10⁴ CFU/g) and (1.4×10⁵ 171 to 2.5×10⁵ CFU/g) for TVC, Staphylococcal, Salmonella-Shigella, Coliform and fungal counts 172 respectively. From food vendor 4, suya sample had total viable count of (1.3×108 CFU/g), 173 staphylococcal count was too numerous to count (TNTC), Salmonella-Shigella count was (5.7×10⁷ 174 CFU/g) and total coliform count of $(3.1 \times 10^5 CFU/g)$ while there was no observable fungal growth For 175 the currency notes obtained, the counts ranged from (2.8 to 6.0×10⁶ CFU/g) for total viable count; 176 (1.7×10⁶ CFU/g to TNTC) for Staphylococcal count; (6.5×10⁴ CFU/g to 6.8×10⁵ CFU/g) for coliform 177 178 count.

Food handling personnel play important roles in ensuring food safety throughout the food chain of 179 180 production, processing, storage and distribution. Mishandling and wanton disregard of hygienic 181 measures on the part of the food vendors have been reported to introduce contaminants and 182 pathogens that survive and multiply in sufficient numbers to cause illness in the consumers [23, 183 24.25.261. The proper preparation of foods in advance of consumption, exposure to unclean 184 environment, holding at ambient temperature conducive for microbial multiplication coupled with the 185 rich nutrient medium of these foods are factors that increase microbial loads of the samples [27]. Microbial guideline for cooked foods stipulates that "plate counts must be less than 1.0×10⁷CFU/g for 186 187 meat; 1.0×10⁴ for plant products [28]. The microbial loads of the food samples were higher than the 188 stipulated, hence, consumption of the foods constitute health risk to consumers. It can be adjudged 189 that the street foods obtained in this study are not bacteriologically fit for consumption. Thus, it calls for improvement in awareness at all levels, especially among personnel working in food 190 191 establishments, on the possible health risks associated with poor handling of paper currencies while 192 serving/preparing foods. Consequently, the combination of the widespread use of paper currencies 193 and their constant exchange make them a likely agent for various disease transmissions since 194 communicable diseases can spread through contact with fomites [29]. Furthermore, a study revealed 195 that paper currencies can serve as an ideal breeding ground for microorganisms [30]. Pieces of 196 money are in continuous circulation, passing in all environments that constitute a reservoir of various 197 bacteria some of which can survive. Paper currency absorb moisture which encourages the growth of 198 microorganisms on them. Fungi get into money especially from air, soil and where it is kept (for

instance, sacks pockets and socks). Money is contaminated as it circulates from hand to hand [31]. Besides reducing the life span of the notes, some of these fungi have the potential to cause skin, eye and gastrointestinal infections [32, 33], infections of internal organs [34, 35] as well as serious diseases of the respiratory tract [36] in man. In Nigeria, currency notes are often manhandled and mutilated by different categories of people, including traders, churches, beggars and commuter bus conductors [37].

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206 **3.2. Identification of microorganisms**

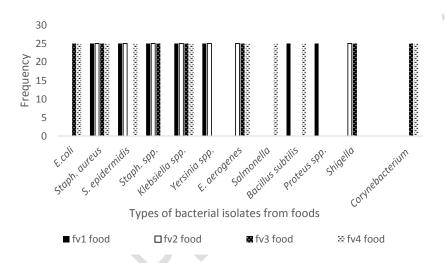
A total of 50 isolates were recovered from all the street food and currency denominations.

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209 Isolates from food samples

210 From food vendors 1 (ekuru and currency samples), 5 and 4 genera were isolated and identified from

- food and currency denominations (Figs. 1a and 1b). The genera included: Staphylococcus, Klebsiella,
- 212 *Bacillus, Proteus, Yersinia, Enterobacter and Salmonella.* At least 3 bacteria genera are found in 213 common in the food and change from same vendor.
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Fig. 1a: Types and frequency of isolation of bacteria from foods (ekuru from food vendor 1; jollof rice from food vendor 2; pounded yam from food vendor 3; and suya from food vendor 4)

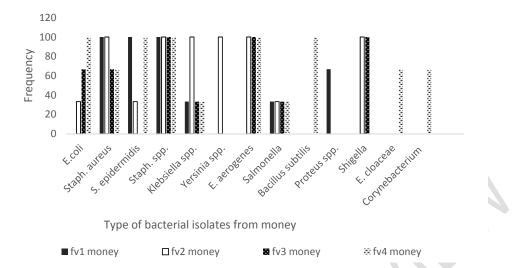


FIG. 1B: Types and frequency of isolation of bacteria from currency denominations received as change from food vendors

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225 From food vendor 2, 5 and 7 genera were identified from food and currency respectively. A total of 7 226 genera were identified from pounded yam, and 6 from the change obtained from food vendor 3. While 227 7 and 8 genera from suya and currency were identified respectively. The organisms isolated were 228 observed to be similar to those isolated by Adamu et al [38]. Shigella, Klebsiella spp. E.coli, 229 Staphylococcus aureus and Staphylococcus spp. were isolated in the food samples. According to 230 Asogwa et al [39], during preparation and vending, food items like raw meat for 'suya' (roasted pieces 231 of meat attached to a stick) and salad raw materials were prepared using the same knife without in-232 between washing. After cooking or roasting, foods are continuously exposed to open environment for 233 quick sale and are often invaded by flies [39].

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3.3. Antibiotics resistance profile of isolates

238 Microbial strains showed varied resistance patterns to the different antibiotics screened. 239 Corynebacterium, E. coli, E. aerogenes, E. cloaceae, Yersinia and Proteus were multidrug resistant, 240 since multidrug resistance was defined as resistance to ≥3 of the antimicrobial agents tested. it was 241 further observed that bacteria show resistance to ceftazidine (30 µg), cefuroxime (30 µg) and 242 amoxycillin/clavulanate (30 µg) while being susceptible to cefriaxone (CTR), ampicillin (AMP), 243 Nitrofurantoin (NIT) and Oflaxacin (OFL) as presented in Table 3. Furthermore, all Gram negative 244 organisms except Yersinia were resistant to Ceftazidime (30 µg). Five of the eight Gram negative 245 were resistant to Ampicillin (10 µg) except Pseudomonas (10 mm), Klebsiella (10 mm) and E. coli (11 246 mm). The percent resistance for Ceftazidime was (61.5 %), Amoxycillin/Clavulanate (53.8 %), 247 Cefuroxime (30.7)%). Ampicillin (62.5 %) (Fia. 3).

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Table 3. Zone inhibition (mm) of antibiotics against foodborne and currency isolates

Isolates	CAZ	CRX	GEN	CTR	ERY	CXC	OFL	AUG	AMP	NIT	CPR
(G. Rxn)											
S.epidermidis (+)	23	29	19	24	21	22	23	30	-	-	-
Corynebact. (+)	R	R	17	27	16	10	14	R	-	-	-
B. subtilis (+)	25	12	26	26	19	21	22	21	-	-	-
S. aureus (+)	13	R	17	16	R	10	18	24	-	-	-
Staph. spp. (+)	15	18	20	26	15	18	26	R	-	-	-
P. aeruginosa (-)	R	10	12	-	-	-	30	20	10	25	26
Klebsiella (-)	R	30	18	-	-	-	23	R	10	34	24
E. coli (-)	R	21	13	-	-	-	R	18	11	24	R
E. aerogenes (-)	R	R	14	-	-	-	18	R	R	25	R

E. cloacae (-)	R	R	24	-	-	-	R	R	R	21	R
Salmonella (-)	R	39	23	-	-	-	31	29	R	39	31
Yersinia (-)	20	23	R	-	-	-	R	R	R	R	R
Proteus (-)	R	21	15	-	-	-	20	R	R	23	21

CAZ= Ceftazidime; CRX= Cefuroxime; GEN= Gentamicin; CTR= Cefriaxone; ERY= Erythromycin; CXC=
 Cloxicillin; OFL= Ofloxacin; AUG= Amoxycillin/ Clavulanate; CPR= Ciprofloxacin; NIT= Nitrofurantoin; and AMP=
 Ampicillin; R=Resistant; - = Not tested.

256 Salmonella, Shigella, E.coli, Yersinia and Staphylococcus aureus are public health important 257 organisms that have been demonstrated to pose significant health risks [40]-[41]. Some of the 258 isolates were found to be multidrug resistant particularly, there were higher drugs resistance in the 259 Gram negative organisms as compared to the Gram positive organisms. Salmonellosis and 260 Shigellosis remain a major public health problem across the globe [42]-[44]. Similarly the Centers for 261 Disease Control and Prevention (CDC) estimates that 48 million cases of foodborne illnesses occur in 262 the United States annually [44]. More than 200,000 persons die of food poisoning in Nigeria every 263 year [45] 264

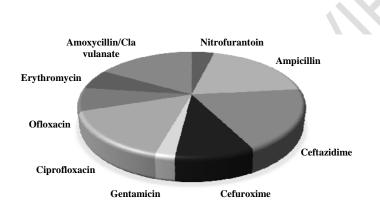


Fig. 3: Antibiotic resistance profile of bacterial isolates from Street vended foods and money

It has been previously reported that indiscriminate use of antibiotics can lead to multidrug resistance [46]. The resistance to antimicrobial agents by bacterial pathogens is a major hindrance to successful therapy and some bacterial strains have been reported to be resistant to most available antimicrobial treatments [47, 48, 49]. In this study, Gram positive organisms were observed to be less resistant compared to Gram negatives, which was also observed by Exner *et al* [50].

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4. CONCLUSION

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285 The microbial load of all the foods in this study exceed the stipulated limit for ready-to-eat foods, 286 thereby making consumption of these foods of safety and public health concern. Suva and ekuru food 287 samples had higher microbial loads, followed by pounded yam and jollof rice samples. Some of the 288 organisms isolated from the money samples were also found in the food samples, this could be an 289 indication of cross contamination between money and food. Gram negative isolates were found to 290 have higher drug resistance than the Gram positive isolates. Also, some of the organisms were found 291 to be multidrug resistant. Most of the isolated organisms in this study were resistant to Ceftazidime 292 (30µg) while Yersinia was resistant to most of the drugs.

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