

Quality of the food in its raw or unprocessed state, the sanitary conditions under which the product is handled and processed and the adequacy of subsequent packaging, handling, storage conditions of the products [11]. The broad objective of this work are to Isolate, estimate, identify and characterize microorganisms (bacteria, fungi) from ready-to-eat fried chicken, obtained from different fast food centers and street vended spots in Benin City, Nigeria. It also compares the microbial load of ready-to-eat fried chicken collected from these fast food centers and street vended spots.

Materials and Methods

Study area

The study areas are the eateries and vendors at the roadside using three different locations (Local Government Areas) in Benin City, Edo State. They are: Uselu Lagos road (Egor LGA), Lagos street (Oredo LGA) and Cara market (Ikpoba Okha LGA). These areas were selected because fried chicken parts are mainly sold there and there is high patronage of street vended foods.

Sample collection

A fried chicken part (Drumstick) was bought from each vending site, making a total of two replicate samples collected from each Local Government Areas at two weeks interval. From each of the sites, ready-to-eat fried chicken samples were purchased from an eatery and a vendor at the road side and transported to the laboratory as being sold and wrapped with transparent nylon by the handlers and kept in a deep cold storage at 4°C and analyzed within 12 hours of collection. The complete process of microbiological analysis was performed under a sterile condition to avoid contamination from environment. The samples were subjected to microbiological analysis according to standard procedures.

Control sample (Laboratory prepared)

Chicken was purchased from the farm house of the University of Benin. Utensil used for killing and dressing the chicken were properly washed. The chicken was killed, properly scalded, cut into desirable sizes and properly seasoned, boiled in 500 ml of water and afterwards deep fried in hot vegetable oil. After frying, fried parts were kept in the refrigerator properly packaged. The drumstick of the chicken was used as the control.

Sterilization of media

The experiment was conducted at the main laboratory of the Faculty of Agriculture, University of Benin. The glassware's such as test tubes, glass rod, pipette, measuring cylinder, beakers and conical flasks required for this research work were soaked and washed with detergent and rinsed with distilled water. They were wrapped with aluminum foil paper and dried in the oven in inverted position at 80°C for 45-60 mins. The working area was swabbed with ethanol. Contamination by microorganisms from the external environment was reduced by closing windows and putting off fans in the laboratory.

Microbiological analysis

The samples studied were analyzed using standard microbiological methods according to Buchanan et al. [12] for the bacteria samples; while the identification and characterization of the fungi was done according to Barnett and Hunter [13].

Preparation of culture media

Nutrient agar (NA) for bacterial evaluation: 28 g of Nutrient Agar was weighed with a top loading balance and dissolved in 1 liter of distilled water in a conical flask covered with cotton wool and aluminum foil paper.

It was mixed thoroughly and sterilized using a pressure cooker at a temperature of 121°C for 15 mins. The medium was cooled to 45-59°C and then dispensed aseptically into sterile petridishes. The petridishes were covered and allowed to solidify.

Potato dextrose agar (PDA) for fungi evaluation: 39 g of Potato Dextrose Agar was weighed with a top loading balance and dissolved in 1 liter of distilled water in a conical flask covered with cotton wool and aluminum foil paper. It was mixed thoroughly and sterilized using a pressure cooker at a temperature of 121°C for 15 min. The medium was cooled to 45-59°C and then dispensed aseptically into sterile petridishes. The petridishes were covered and allowed to solidify.

Culturing and isolation of fried chicken samples: 1 g of each fried chicken sample was homogenized in 10 ml of sterile distilled water.

1ml was pipetted from it and added to next blank distilled water, making a dilution of 10^{-2} . Dilutions were made by mixing 1.0ml of the homogenate in 9.0 ml of sterile distilled water to obtain 10^{-3} dilution. The dilution was then made to 10^{-4} , 10^{-5} and 10^{-6} . Total viable counts of bacteria were determined by enumerating the colony forming units (cfu/g) by pour plating 1.0 ml of 10^{-5} diluents on nutrient agar plates supplemented with anti-fungimixture to inhibit fungi growth and incubated at 37°C for 48 hrs.

Total fungal counts were determined by pour plating on Potato Dextrose Agar plates supplemented with antibiotic mixture to inhibit bacterial growth, and incubated at room temperature ($26 \pm 2^\circ\text{C}$) for 72 hrs and the presence of *Escherichia coli* was detected by pour plating on Eosine Methylene Blue (EMB) agar plates and incubated at 37°C for 48 hrs. Samples that were positive showed a greenish metallic sheen which is the appearance of *Escherichia coli*. The experiments were carried out in triplicates. Pure cultures of bacterial and fungal isolates were obtained on the Nutrient Agar and Potato Dextrose Agar plates respectively.

Enumeration of bacterial isolates

At the end of the incubation period, the plates were read and observed for growth. The population of the bacterial isolates was estimated by counting the number of discrete colonies in the plate. Morphological and cultural characteristics were observed. Discrete colonies were sub cultured by streak plate method in order to obtain pure culture. Stock cultures of each colony were prepared in the agar slants which were incubated for 24 hrs to allow for growth after which they were stored at 4°C in the refrigerator for further experimentation.

Gram staining

Gram staining was carried out as described by AOAC [14]. A drop of sterile distilled water was placed on a clean grease free slide. The inoculating wire loop was flamed until red hot. The loop was allowed to cool and a small portion of the organism to be gram stained was picked and smeared in the drop of water on the slide. The slide was then air dried. It was heat fixed by passing it gently over flame. The smear was stained with 1% crystal violet for 1 minute and washed with distilled water. Gram iodine was added as a mordant for 1 min. this was drained off and 70% alcohol was added for 30 seconds. This acted as a

decolorizer. The above is termed primary staining. The slide was then rinsed with distilled water. The slide was finally flooded with counter stain, safranin for 1min and washed off with distilled water and air dried. The slide was observed under the microscope oil immersion $\times 100$ objective lens. The Gram positive organisms appeared purple while the Gram negative organisms appeared red.

Biochemical test

Coagulase test: This test is used to determine the enzyme coagulase that differentiates between pathogenic strains of staphylococcus. A colony of the test organisms was emulsified with sterile normal saline solution on a clean slide using a sterile wire loop. A drop of rabbit plasma was added and mixed with emulsion. The positive coagulase organisms showed clumping while negative coagulase organisms showed no clumping.

Catalase test: This test is used to demonstrate the presence of enzyme catalase, which catalyzes the release of oxygen from hydrogen peroxide. The pure culture of the test organism was placed and added to a drop of 3% hydrogen peroxide solution on a clean slide. The production of gas bubble from the surface indicates positive result.

Oxidase test: This test was carried out to detect the production of the enzyme oxidase by the bacteria isolates. A piece of filter paper was soaked in a few drop of oxidase reagent (tetramethyl-p-phenyl-diamine-dihydrochloride). A colony of the test organism was then smeared on the soaked filter paper. An oxidase producing organisms on the filter paper oxidized the phenyl-diamine in the reagent to deep purple colour. This change in colour to deep purple within 10seconds indicates positive result.

Citrate test: This test detects the ability of an organism to use citrate as the sole source of carbon and energy and ammonium salt as the sole source of nitrogen. Simmon citrate agar is used for the preparation of the media. The test is usually used as an aid in the differentiation of organisms in the *Enterobacteriaceae* and most other genera. The medium used for this test was the Simon's citrate agar was inoculated within 24 hrs culture of the isolates. The inoculation was done by stabbing the medium using sterile straight inoculating wire. The inoculated medium was then incubated at 3°C for about 24 hours. Change in colour from green to blue after about 24 hours of incubation indicated positive result.

Identification of fungal isolates

Smears of the pure culture of the fungal isolates were made on clean slide and Lactophenol cotton blue reagent was added for 1minute and after which it was rinsed with distilled water. Then 95% ethanol was added for 30 s and rinsed again and viewed under the microscope.

Proximate composition

The Proximate composition for moisture, protein, fat and ash was determined according to AOAC [14]. The crude protein content was determined by the Kjeldahl method and the crude fat content was determined by solvent extraction using the Soxhlet technique. The fried chicken sample was first hydrolyzed with boiling HCl before extraction with petroleum ether. Each extracted group was then dried in a forced air oven at 105°C for 3 hrs (to constant mass), cooled in desiccator and then weighted to obtain the dry solids content. To determine ash content, fried chicken sample (1 g) were minced, weighed and ignited in the crucible. Then it was transferred in the muffle furnace held at dark red at a rate of 550°C for 6-8 hrs until the residue was white.

Moisture content was determined by drying the samples overnight at 105°C.

Calculation of moisture:

$$\text{Moisture (\%)} = \frac{\text{Weight loss} \times 100}{\text{Original weight of the sample}}$$

Calculation of crude protein:

$$N_2 (\%) = \frac{0.00014 \times \text{Vol of acid} \times \text{Filtration flask for preparation of digest} \times 100}{\text{Weight of sample} \times \text{Amount of digest introduced into the flask}}$$

$$\text{Protein (\%)} = \% \text{ of total } N_2 \times 6.25$$

Calculation of crude fat:

$$\text{Fat (\%)} = \frac{\text{Weight of the oil} \times 100}{\text{Weight of the samples taken}}$$

Calculation of ash:

$$\text{Ash (\%)} = \frac{\text{Weight of ash} \times 100}{\text{Original weight of the sample taken}}$$

Statistical Analysis

All data collected from the experiment were subjected to the Analysis of Variance (ANOVA) using GenStat package (12.1 versions). The separation of means was carried out using Duncan's Multiple Range test at 5% level of probability ($P < 0.05$).

Results and Discussion

Proximate analysis

The proximate composition of fried chicken samples prepared in the laboratory (Control) and Fried chicken bought in different locations in Benin Metropolis are shown in Figure 1. For the moisture content of fried chicken there was no significant difference ($P > 0.05$) in the eateries and street vended chicken of Oredo and Egor LGAs. However there was significant difference ($P < 0.05$) in the street vended chicken and eatery of Ikpoba Okha. The control differed significantly ($P < 0.05$) from the three Local Government Areas; with moisture content value of 55.67% which means that the control which is the laboratory prepared had the least moisture content. Frying reduces the moisture in food. As reported by Varela et al. [15], when frying food, the hot frying fat that has penetrated into it, replaces part of the water it contains, making the food considerably more palatable. The reason why it appears that the eateries and the street vended meat had high moisture content as compared to the laboratory prepared is obvious, the commercial one would not like to fry their meat to that point where it reduces yield and dry; but would rather want more moisture in the meat in order to increase yield (more weight). The high moisture content of fried chicken sold in the eateries was also probably due to the use of microwave to warm the meat before serving.

There was significant differences ($P < 0.05$) in the Crude protein of street vended chicken and eateries of Ikpobaokha and Oredo LGAs, except in Egor LGA where the street vended chicken and eatery were not significantly different ($P > 0.05$). The control differed significantly ($P < 0.05$) from the eateries and street vended meat of the three Local Government Areas. It had the highest crude protein value of 30.52%. Crude protein content is a factor of the diet of the chicken and these chickens are sought from different locations. The variation in crude protein may be as a result of differences in species, production, sex and age of the animal etc. The crude fat content of street vended chicken and eatery of Ikpobaokha did not differ significantly ($P > 0.05$) from the street vended meat of Oredo. It was however, different from the chicken sold in the eatery of Oredo. There was no significant difference

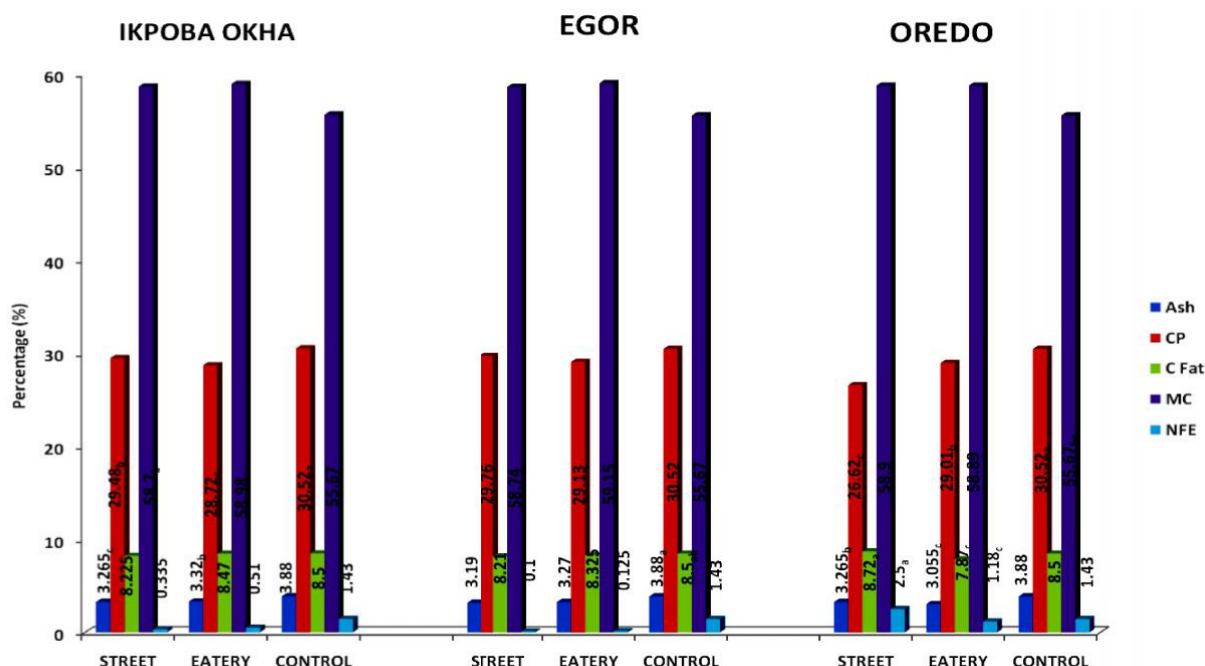


Figure 1: Proximate composition of fried chicken tissues sold in eatery and street with the control in Benin Metropolis. Mean value with subscript shows significant difference (P < 0.05) within the local government area.

(P>0.05) in the street vended chicken of Egor and the laboratory prepared. The composition of chicken meat as edible raw carcass meat has a crude fat content of 6.10% as documented by Olomu [16].

The average high fat content (value) of fried chicken might have been as a result of the absorption of fat during frying in oil [17] or what is usually practiced by most commercial vendors, refrying of leftover meat for subsequent day sales, which is in agreement with the findings of Pinthuset al. [18] who stated that during repeated frying, the polarity of oil increases and the interfacial tension decreases.

Ash content was highest in the control (3.88%), it differed significantly (P<0.05) from the meat sold in Ikpoba Okha, Oredo and Egor. The ash content of Street vended meat in Ikpoba Okha differed significantly (P<0.05) from the street vended meat in Oredo and Egor LG As the eatery in Ikpoba Okha did not differ significantly (P>0.05) from the eatery in Egor; but differed significantly (P<0.05) from the eatery in Oredo. The mineral content of meat (shown as 'ash' in Figure 1) include calcium, phosphorus, sodium, potassium and some trace elements like zinc, iron etc. The high values of the ash content correspond to the findings of Saguy and Dana [19]. They stated that during deep frying, the time required is generally short and the temperature inside the product remains below 100°C, but there is less loss of water soluble vitamin. However, nutrients such as minerals, carbohydrate and proteins are retained. The variable results of these parameters are in line with the findings of Huss [20]; who observed that the chemical composition of animal tissue varies from species to species as well as from individual animals depending on age, sex, environment and season. Figure 2, shows the graphical representation of the statistical analysis of the proximate composition of fried chicken samples. The subscript shows the significant difference (P<0.05) within each Local Government Area.

Mean value with subscript shows significant difference (P < 0.05) within the Local Government Area.

Microbial analysis

Thirteen genera of bacteria and seven genera of fungi were isolated from fried chicken (Table 1). Microbiological quality of fried chicken samples on the bases of the mean colony forming unit of the total bacteria and fungi per gram of fried chicken sample from both the street vendor and the standard eatery is shown in Table 2. The frequency of occurrence (%) of each microbial isolate was also determined (Tables 2 and 3). The highest number of isolates was also determined from fried chicken sold in the eatery in Oredo with a percentage diversity of 21.1% (Table 4). *Aspergillus niger* had a frequency of (9.9%). It was the only fungi isolated from all the fried chicken samples. Figure 2 is a graphical representation of the statistical analysis of the mean bacterial and fungi colonies per plate in the two vending sites in the different locations in Benin Metropolis. The superscripts show significant difference across the Local Government Areas; while the subscript shows significant difference within the LGAs.

The microbiological analysis of fried chicken samples obtained from the different vending sites in IkpobaOkha, Oredo and Egor LGAs in Benin City, Edo State is shown in Table 1. Laboratory prepared fried chicken served as control, had mean bacteria count of 4.670×10^6 cfu/g; the street vended chicken range from 5.8×10^6 cfu/g to 1.12×10^7 cfu/g and the chicken from the eatery ranged between 6.985×10^6 to 1.20×10^7 cfu/g. Statistical analysis of the mean bacteria load showed no significant difference (P>0.05) between the eatery and street fried chicken of IkpobaOkha, Oredo and Egor. However, the street vended fried chicken in the three LGAs differ significantly (P<0.05) from the control. It was only the eatery from Oredo that differed significantly (P<0.05) from the control sample. The results obtained were nearly

Sampling point	Vending site	Mean number of Bacterial colonies per plate. (Cfu/g ± SEM)	Average number of Fungi colonies per plate. (Cfu/g ± SEM)
IkpobaOkha	Street	9.730 X 10 ⁶ _a ± 3.76	2.615 X 10 ⁶ _a ± 2.31
	Eatery	6.985 X 10 ⁶ _{ab} ± 1.47	1.7 X 10 ⁶ _a ± 1.08
Oredo	Street	5.8 X 10 ⁶ _a ± 1.88	1.385 X 10 ⁶ _a ± 1.44
	Eatery	1.2 X 10 ⁷ _a ± 3.15	1.9 X 10 ⁶ _a ± 2.50
Egor	Street	1.12 X 10 ⁷ _a ± 1.19	1.75 X 10 ⁶ _a ± 2.39
	Eatery	1.00 X 10 ⁷ _{ab} ± 1.19	7.85 X 10 ⁵ _a ± 0.23
Control		4.67 X 10 ⁶ _b ± 1.40	4.70 X 10 ⁵ _b ± 0.19

Values with different subscripts differ significantly

Table 1: Fungi and bacteria counts, in colony forming units per gram (cfu/g) of fried chicken sample.

Isolates	No	%	IkpobaOkha		Oredo		Egor		Control
			Street	Eatery	Street	Eatery	Street	Eatery	
<i>Proteussp.</i>	7	9.9	✓		✓	✓	✓	✓	✓
<i>Micrococcussp.</i>	6	8.5	✓	×	✓	✓	✓	✓	✓
<i>Staphylococcusaureus</i>	7	9.9	✓	✓	✓	✓	✓	✓	✓
<i>Streptococcussp.</i>	3	4.2	✓	✓	✓	×	×	×	×
<i>Escherichiacoli</i>	3	4.2	✓	✓	✓	×	✓	×	×
<i>Klebsiellasp.</i>	3	4.2	✓	×	✓	×	✓	×	×
<i>Acinetobactersp.</i>	1	1.4	✓	×	×	×	×	×	×
<i>Enterobactersp.</i>	6	8.5	✓	✓	✓	✓	✓	✓	×
<i>Staphylococcusepidermidis</i>	3	4.2	×	×	✓	✓	✓	×	×
<i>Bacillussp.</i>	1	1.4	×	×	×	✓	×	×	×
<i>Staphylococcussp.</i>	1	1.4	×	×	×	✓	×	×	×
<i>Pseudomonassp.</i>	3	4.2	×	×	✓	×	✓	✓	×
<i>Streptococcusmutans</i>	2	2.8	×	×	✓	×	✓	×	×
TotalBacteria	46								
Overalltotal	46	64.8							
Diversity	46		8	7	9	11	4	5	3
%diversity	46	100.2	17.4	15.2	19.6	23.9	8.7	10.9	6.5

✓Means Present; X Means Absent

Table 2: Frequency of occurrence of bacterial isolates in fried chicken tissues sold in Benin Metropolis.

similar to those reported by Saad which recorded that the mean values of aerobic plate count examined samples of fast foods were $3.92 \times 10^4 \pm 0.57 \times 10^4$ cfu/g for grilled beefkofta; $6.33 \times 10^4 \pm 1.84 \times 10^4$ cfu/g for fried beef burger; $4.78 \times 10^5 \pm 0.96 \times 10^5$ cfu/g for fried chicken meat; $7.26 \times 10^5 \pm 2.01 \times 10^5$ cfu/g for fried chicken kofta and $1.89 \times 10^6 \pm 0.27 \times 10^6$ cfu/g for fried chicken Pattie. FAO [21,22], stated that bacteria counts that is greater than or equal to 10^6 cells per gram and *Enterobacteriaceae* counts greater than or equal to 10^5 cells per gram are dangerous and could result to some health problems like food poisoning and intoxication. From the results obtained, the values of bacterial counts were above stated dose i.e. infective dose that will cause or pose a health

risk. Laboratory prepared fried chicken served as control, had mean fungi count of 4.70×10^5 cfu/g; the street vended chicken range from 1.385×10^6 to 2.615×10^6 cfu/g and the fried chicken from the eatery ranged between 7.85×10^5 to 1.9×10^6 cfu/g. Statistical analysis of the mean fungi load showed no significant difference ($P > 0.05$) between the street vended chicken and eatery in IkpobaOkha, Oredo and Egor Local Government Areas. The results obtained in this study are nearly similar to the findings of Woguet al. [23]. Their findings indicated that the microbial load of RTE rice from both local fast food centers (Local restaurants) and standard fast food centers (High class restaurants) in Benin City, had Total colony count range from 2.0×10^4 to 1.2×10^6 for

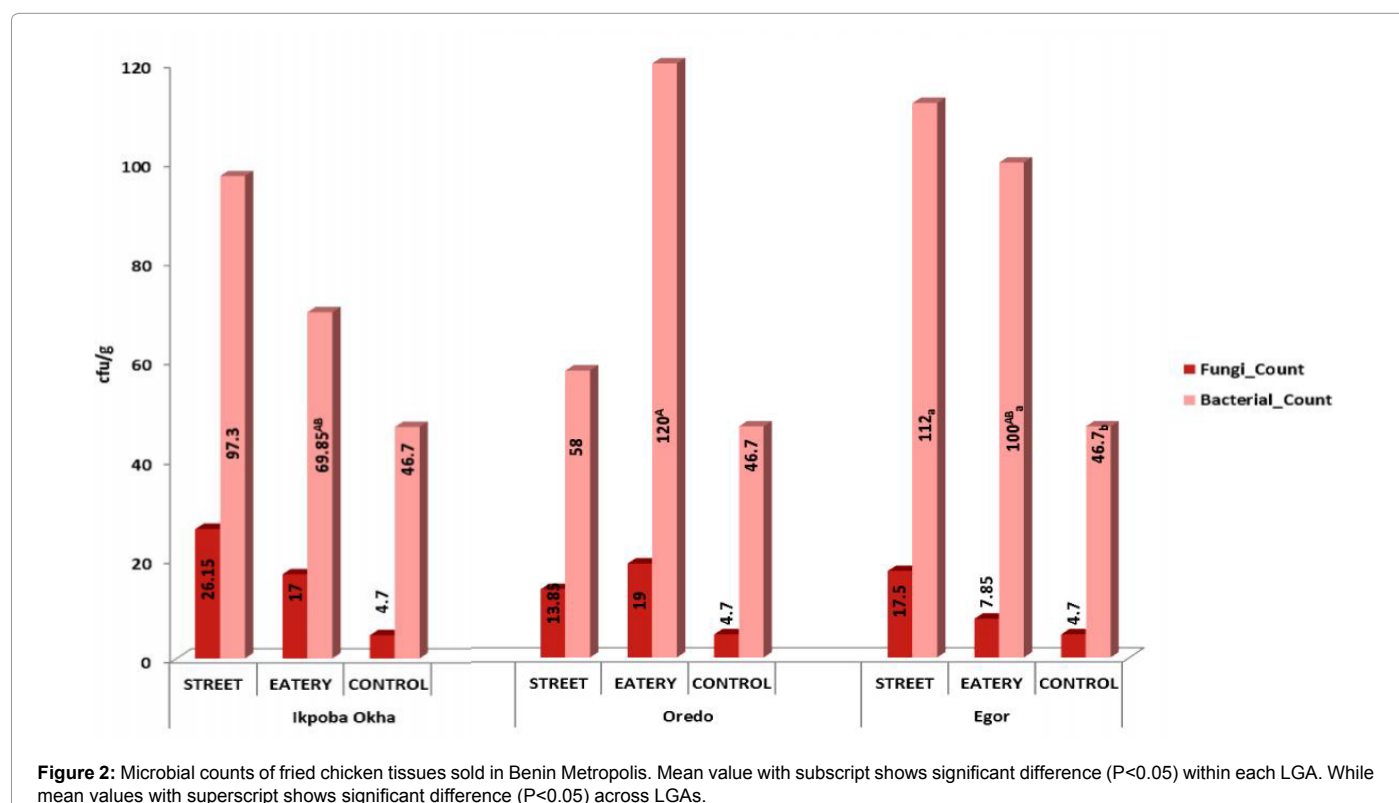
Isolates	No	Percent	IkpobaOkha		Oredo		Egor		Control
			Street	Eatery	Street	Eatery	Street	Eatery	
<i>Aspergillusniger</i>	7	9.9	✓	✓	✓	✓	✓	✓	✓
<i>Mucormucedo</i>	5	7	✓	✓	×	✓	×	✓	✓
<i>Cladosporium sp.</i>	2	2.8	✓	×	✓	×	×	×	×
<i>Penicilliumoxalicum</i>	4	5.6	✓	×	✓	✓	✓	×	×
Yeast	4	5.6	×	×	×	✓	✓	✓	✓
<i>Aspergillusnidulans</i>	1	1.4	×	×	✓	×	✓	×	×
<i>Penicilliumitalicum</i>	2	2.8	×	✓	×	×	✓	×	✓
Total fungi	25								
Overall total	25	35.1							
Diversity	25		4	3	4	4	3	3	4
% diversity	25	100	16	12	16	16	12	12	16

✓ Means Present; X Means Absent

Table 3: Frequency of occurrence of fungi isolates in fried chicken tissues sold in Benin Metropolis.

	No	Percentage	IkpobaOkha		Oredo		Egor		Control
			Street	Eatery	Street	Eatery	Street	Eatery	
Total microbial	71								
Overall total	71	99.9							
Diversity	71		12	7	15	11	12	8	7
Diversity Percent	71	101.2	16.9	9.8	21.1	15.5	16.9	11.2	9.8

Table 4: Sum total of microbial isolates in fried chicken tissues sold in Benin Metropolis.



bacteria and 8.0×10^4 to 2.0×10^5 for fungi. Their result also showed that the standard fast food centers had more microbial load and more microorganisms compared to RTE rice from local fast food centers.

The frequency of occurrence (%) of each microbial isolate was also determined (Tables 2 and 3). The highest number of isolates was obtained from fried chicken sold in the eatery in Oredo with a percentage diversity of 21.1% (Table 4). This was followed by the street fried chicken sold in Ikpobaokha and Oredo (16.9%), eatery in Ikpoba Okha (15.5%), eatery in Egor (11.2%) and street vended chicken in Egor and the control (9.8%). The frequency of occurrence of bacteria isolates was higher than the fungi isolates. For example, *Proteus sp* and *Staphylococcus aureus* (9.9%) were the most frequently isolated organisms from all the fried chicken samples. *Aspergillus niger* had a frequency of (9.9%). It was the only fungi isolated from all the fried chicken samples. This was followed by *Micrococcus sp.* and *Enterobacter sp.* (8.5%), *Streptococcus sp.*, *Escherichia coli*, *Klebsiella sp.*, *Staphylococcus epidermidis*, and *Pseudomonas sp* (4.2%), *Streptococcus mutans* (2.8%) and *Acinetobacter sp.*, *Bacillus sp.* and *Staphylococcus sp.* (1.4%). While *Aspergillus niger* (9.9%) was the most frequently isolated fungi followed by *Mucormucedo* (7%), *Penicillium oxalicum* and Yeast (5.6%), *Cladosporium sp.* and *Penicillium italicum* (2.8%) and *Aspergillus nidulans* (1.4%) which were the least frequent fungi isolates implicated in this study.

In this study, thirteen genera of bacteria were isolated from fried chicken. The isolates were identified as *Proteus sp.*, *Micrococcus sp.*, *Staphylococcus aureus*, *Streptococcus sp.*, *Escherichia coli*, *Klebsiella sp.*, *Acinetobacter sp.*, *Enterobacter sp.*, *Staphylococcus epidermidis*, *Bacillus sp.*, *Staphylococcus sp.*, *Pseudomonas sp.* and *Streptococcus mutans*. The fungi isolated include *Aspergillus niger*, *Mucormucedo*, *Cladosporium sp.*, *Penicillium oxalicum*, yeast, *Aspergillus nidulans* and *Penicillium italicum*. This is in agreement with Clarence et al. [24] that isolated six genera of bacteria which include *Staphylococcus sp.*, *E. coli*, *Klebsiella sp.*, *Pseudomonas sp.*, *Bacillus sp.* and *Enterococcus sp.* This result is also in agreement with the work of Oluwafemi and Simisaye [25] and Okonko et al. [26] where they isolated almost similar organisms from sausages and seafood respectively. Oranusi and Braide [5], isolated *Aspergillus niger* from ready-to-eat foods sold along Onitsha- Owerri highway. He stated that mishandling of food products could lead to proliferation of fungi beyond acceptable limits. The presence of these organisms in ready-to-eat food (Fried chicken) depicts a deplorable state of poor hygiene and sanitary practices employed in the processing and sales of these food products. Also, the presence of fungi such as *Aspergillus niger* in the food signals danger to public health since many of these fungi are toxin-producing organisms.

It is apparent from this investigation that the presence of coliforms in the street meat as compared to the eateries and the control could be a reflection of the level of exposure, and the handling processes in these two vending sites. In the open market the products are left exposed without any form of packaging or covering as often as the customers' demand. This open display of products is to attract the customers and this encourages sporadic visits by flies. The dusty, unhygienic market environments coupled with the poor handling by the vendors are factors contributing to the high microbial load. The common practice of using same utensils for picking meat (cross contamination) could also be responsible for the microbial loads obtained even from fried chicken gotten from the eateries [27]. Vendors of ready-to-eat fried chicken usually make use of simple facilities like wheel barrows, trays, tables and make-shift stalls, thus further increasing the risk of food contamination. Pathogens may invade the exterior parts of the

meat during processes like handling and packaging [28]. The chances of contamination both in the street vended fried chicken and the eateries may be further heightened by the fact that, fried chicken is sold without adequate storage conditions. This practice exposes the fried meat to flies as for the street meat and maintains the product at optimum temperatures for microbial invasion and proliferation of contaminants such as pathogenic mesophiles and other disease-causing agents as for the eateries. The fried chicken are processed and sold by poorly educated and untrained vendors in food hygiene [9,28]. Poorly processed street vended product (insufficient frying) has been identified as an important cause of diseases in developing countries [29,30]. *Bacillus sp.*, *Penicillium sp.* and *Aspergillus sp.* were present in most of the samples. This could be due to the fact that these organisms are spore formers and are known common environmental contaminants. However these organisms have been implicated as food borne pathogens [31-33]. The growth of *Bacillus spp* in fried chicken was ascribed to its presence in the raw materials in spore form which survived cooking; possibly coupled to its reintroduction into the fried meat through post-cooking contamination. *Staphylococcus aureus* is a normal body flora of humans found on nasal passages, skin and mucous membranes and could have been introduced through unclean hands and mouth of the preparer, vendor and/or customers [34,35]. The packaging materials (usually cellophane) are normally opened by squeezing/blowing air into it. The presence of *Staphylococcus aureus* could also be attributed to the raw ingredients used (raw tomato sauce, raw sliced onion), unclean utensils and poor environment in which the food is prepared. *Staphylococcus aureus* is an opportunistic pathogen; and the *Enterotoxigenic* strains of it are known to cause serious food-borne disease [32,36]. The presence of *Escherichia coli*, *Klebsiella sp.*, *proteus sp.* and *Enterobacter sp.* calls for concern as these organisms are frequently associated with poor sanitary practices and could be a pointer to the danger of possible food-borne infection. *Escherichia coli* are especially of fecal origin and have been implicated in numerous food-borne diseases [11,33]. The vendors, water, environmental pollution and inadequate washing of hands and utensils appear to be the major causes of the hazard associated with this fried chicken and must be addressed properly. It is interesting to note that high temperature through boiling and frying is commonly involved in the preparation of fried chicken, which is sufficient to eliminate most of the microorganisms. However, post-processing contamination may occur when the product is not appropriately handled and packaged; this affects the quality of the final product. Since *Escherichia coli* which is one of the most important food-borne pathogen, was isolated from the street vended fried chicken samples, the consumption of this food should be discouraged especially since the food is left exposed without any form of heating. Otherwise, the food should be reheated before consumption. However, in spite of the hygienic processing done to the laboratory prepared fried chicken (control) the microbial counts were high. This clearly and /or probably was due to the challenges encountered in the laboratory. The contamination might have come from the working environment.

Conclusion

The result of this study indicated that the fried chicken sample obtained from all the sampled areas in this study were microbiologically (bacteria) unsafe for consumption. Proper processing of the food under hygienic conditions so as to reduce the risk of contamination is required

References

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