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ANTIMICROBIAL RESISTANCE IN ENTEROBACTERI-ACEAE FROM INTENSIVELY-REARED APPARENTLY HEALTHY AND DISEASED POULTRY IN ABEOKUTA, NIGERIA

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ABSTRACT

The emergence and wide-spread dissemination of antimicrobial resistant bacteria strains is a global phenomenon of great public health and economic implications. Antimicrobial resistance was investigated in enterobacteriaceae isolated from apparently healthy and diseased poultry birds using the broth micro-dilution method to determine antimicrobial minimum inhibitory concentration (MIC). In all, 504 bacterial isolates including Escherichia coli (471), Klebsiella spp (28) and Salmonella enterica isolates (5) were studied. The isolates were resistant to ampicillin (88.5%), chloramphenicol (62.3%), ciprofloxacin (74.8%), enrofloxacin (81.0%), neomycin (83.9%), norfloxacin (78.8%), streptomycin (91.3%) and tetracycline (83.3%). The geometric mean MIC (µg/µL) of tested antimicrobials for enterobacteriaceae is as follows: ampicillin (102.5), chloramphenicol (48.4), ciprofloxacin (19.1), enrofloxacin (34.5), neomycin (47.7), norfloxacin (24.5), streptomycin (142.2) and tetracycline (62.5). Although rates of resistance to ampillin, streptomycin and tetracycline were similar among isolates from apparently healthy and diseases birds, resistance to chloramphenicol, ciprofloxacin, enrofloxacin, neomycin and norfloxacin were significantly higher (p<0.05) in isolates from diseased chickens than in those from apparently healthy chickens. The high rates of antimicrobial resistance in bacteria may contribute to the persistence of pathogens in poultry flock and ineffectiveness of antimicrobial chemotherapy during disease outbreaks.

Keywords: Antimicrobial resistance, apparently healthy chickens, diseased chickens, diseased turkeys, *enterobacteriaceae*

INTRODUCTION

Antimicrobials are important drugs for the prevention and treatment of bacterial infections in humans and animals (Schwarz and Chaslus-Dancla 2001; DANMAP, 2011). The introduction and use of antimicrobials

have contributed remarkably to the sustenance and growth of the livestock industry (Schwarz and Chaslus-Dancla 2001). As a result of the benefits derivable from antimicrobials usage, these drugs have been used without restriction in livestock production.

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Many livestock producers depend on antimicrobials to cover-up for unhygienic and inadequate management practices that expose animals to potential pathogens and increase their susceptibility to infections (Soulsby, 2007; Silbergeld *et al.*, 2008). Oftentimes, antimicrobials are administered without due consideration for the possible deleterious effects they exert on the microand macro- ecosystem (WHO, 2007).

Over the years, the continuous use of antimicrobials has boomerang into a situation where the continued efficacy of these drugs is under threat due to the occurrence of highly resistant bacteria strains which are refractory to antimicrobial therapy (Barbosa and Levy, 2000). The increasing widespread emergence and dissemination of these multi -drug resistant bacteria is a result of the combined effects of overdependence on antimicrobials, inadequate management practices, climate change, globalization and international trade (Harbarth and Samore, 2005; MacPherson et al., 2009). Globally, there is an increase in reports of resistant bacteria of human and animal origins. The socio-economic consequences associated with increased morbidity and mortality from refractory infections have reached such a magnitude that calls for concerted efforts by all local and international stakeholders in tackling the problem of antimicrobial resistance in bacteria (WHO, 2001). *Escherichia coli* is an important pathogen in humans and animals. Pathogenic E. coli is capable of causing devastating intestinal and extra-intestinal diseases in infected hosts (Nataro and Kaper, 1998). *Escherichia coli* is a major cause of morbidity and mortality in poultry and can be transmitted to humans through the consumption of contaminated poultry products (van den Bograad et al., 2001; Stordeur et al., 2002; Kabir, 2010).

The organism is also used as an indicator bacterium for the surveillance of antimicrobial resistance in the ecosystem and also for tracing faecal contamination of food products, hence, the possible presence of other pathogenic bacteria (Momtaz et al., 2012; Bergeron et al., 2012). Antimicrobial resistance in commensal E. coli plays important roles in the maintenance and dissemination of resistant traits in the community (Kijima-Tanaka et al., 2003). Drug-resistant E. coli may serve as important reservoirs of resistant genes for pathogenic and nonpathogenic recipient bacterial species (Osterloh, 2004; Sunde and Norström 2006). Surveillance programmes for monitoring antimicrobial resistance in bacteria are important in the development of strategies for the prevention and control of antimicrobial resistance. However, in the developing countries, scarcity of data complicates attempts to assess the magnitude of threat to the livestock industry public health by resistant bacteria. Inadequate documentation of observable trends in antimicrobial resistance hampers risk assessment and development of suitable interventions to mitigate the menace of antimicrobial resistance in developing countries.

The present study investigates the incidence of antimicrobial resistance in *E. coli, Klebsiella* spp and *Salmonella* serotypes bacteria isolated from apparently healthy and diseased intensively-reared chickens and turkeys in Abeokuta, Nigeria.

MATERIALS AND METHODS Sampling information

Between March 2008 and December 2011, samples were collected from seven poultry farms for bacteria isolation and determination of antimicrobial susceptibility.

Sampling from apparently healthy chickens:

Faecal samples were collected from apparently healthy, intensively raised commercial layer chickens from five farms. The birds were in battery cages and had history of previous vaccination against Infectious Bursal Disease, Newcastle Disease, Marek's Disease and Fowl Pox. In addition, regular prophylactic antimicrobial and booster Newcastle Disease vaccine administrations were common practices in all the farms. Pooled cloacal swabs were collected from live birds on the farms. Five cloacal swabs were pooled as one sample. Sixty pooled samples (300 cloacal swabs) were collected from each farm. Pooled faecal sampling was used because it increases the chance of inclusion of faecal materials from infected birds which may contain high numbers of organisms and thus compensate for the possible low level present in other birds (Carrique-Mas and Davies, 2008; Varga et al., 2008). Chickens sampled were randomly selected among the flock. A total of 300 pooled faecal samples were thus collected from five farms.

Sampling from sick birds:

Clinical samples (diarrhoeic faeces and tissue samples) from two farms with history of diarrhoea were examined. One of the farms was a commercial layer farm with adult laying chickens. The other farm was a broiler farm with young chicks and turkey poults of four to six weeks old. Bacterial infections were suspected in both cases (after ruling out viral and protozoan involvement) and samples submitted for bacteriology. Cloacal swabs were collected from individual live sick birds. Post mortem tissue samples from liver, lung and spleen samples were aseptically collected for bacteriological examination. From the commer-

cial layer farms, 78 cloacal swabs and 60 tissue samples (20 each of liver, lung and spleen) were examined. From the broiler farm, 62 cloacal swabs and 33 tissue samples (11 each of liver, lung and spleen) were collected from diarrhoeic and dead chicks while 96 cloacal swabs and 60 tissue samples (20 each of liver, lung and spleen) were collected from diarrhoeic and dead poults.

Bacteria isolation and identification:

Faecal samples were each inoculated directly onto MacConkey agar (CM 0115 Oxoid® Basingstoke, UK) while tissue samples were first inoculated into Tryptic Soy Broth (TSB) for enrichment before being transferred onto MacConkey agar and 5% blood agar. Cultures were incubated at 37°C for 18 to 24 hours. After incubation, agar plates were examined for bacterial growth. Discrete colonies of bacteria were identified and selected. Selected colonies were purified on MacConkey agar and blood agar, Gram-stained for microscopy and tested for catalase and cytochrome oxidase production. Colonies that yielded oxidase negative, catalase positive, Gram-negative rods were subjected to further identification using biochemical tests kits (Oxoid Microbact GNB 24E[®]) and reactions interpreted by using accompanying computer software package (Oxoid Microbact[®] 2000 version 2.03).

Antimicrobial susceptibility testing

The bacteria isolated from samples were tested for susceptibility to antimicrobial agents. Susceptibility to ampicillin (Amp), chloramphenicol (Chl), ciprofloxacin (Cip), enrofloxacin (Enr), neomycin (Neo), norfloxacin (Nor), streptomycin (Str) and tetracycline (Tet) were determined by the broth micro-dilution technique to determine the minimum inhibitory concentration (MIC) using antimicrobial concentrations ranging

from 0.25-512 $\mu g/\mu L$ according to the standard guidelines by Clinical and Laboratory Standards Institute (2008). The antimicrobial MIC were determined with reference to the respective antimicrobial breakpoint concentrations for bacterial isolates (ampicillin, 32 $\mu g/\mu L$; chloramphenicol, 32 $\mu g/\mu L$; ciprofloxacin, $4 \mu g/\mu L$; enrofloxacin, $4 \mu g/\mu$ μ L; neomycin, 16 μ g/ μ L; norfloxacin, 4 μ g/ μ L; streptomycin, 64 μ g/ μ L and tetracycline, 16 μ g/ μ L) (CLSI, 2008). Isolates with minimum inhibitory concentrations (MIC) higher than the breakpoint for the respective antimicrobial agents were regarded as resistant while those with MIC equal to or lower than the breakpoint were regarded as susceptible.

Statistical Analysis:

Data were expressed in absolute values and in percentages. The geometric mean of MIC values were determine using Microsoft Office Excel 2007 software package. Rates of antimicrobial resistance were compared between isolated from apparently healthy and diseased birds by Chi-square test at p<0.05 probability level using Statistical Software Package for Social Sciences (SPSS, version 16, 2007).

RESULTS

A total of 504 bacterial isolates belonging to three genera in the family *Enteroobateriaceae* were obtained in this study. The isolates comprised of *E. coli* (471), *Klebsiella spp* (28) and *Salmonella enterica* (5) (Table 1). Overall, the isolates showed resistance to ampicillin (88.5%), chloramphenicol (62.3%), ciprofloxacin (74.8%), enrofloxacin (81.0%), neomycin (83.9%), norfloxacin (78.8%), streptomycin (91.3%) and tetracycline (83.3%) (Table 1). Rates of resistance to chloramphenicol, ciprofloxacin, enrofloxacin, neomycin and norfloxacin were signifi-

cantly higher (p < 0.05) in bacterial isolates from diseased chickens than in those from apparently healthy chickens. However, there was no significant difference (p > 0.05) in the rates of antimicrobial resistance in isolates from diseased chickens and turkeys.

Antimicrobial resistance in bacterial isolates from apparently healthy commercial chickens

The rates of antimicrobial resistance in *E. coli* isolates from commercial chickens is ampicillin 74.4%, chloramphenicol 37.8%, ciprofloxacin 36.1%, enrofloxacin 48.8%, neomycin 57.6%, norfloxacin 46.5%, streptomycin 76.7% and tetracycline 57.6%. The geometric mean MIC was highest (121.5 μ g/ μ L) for streptomycin and lowest (3.0 μ g/ μ L) for norfloxacin (Table 2).

Klebsiella isolates from apparently healthy commercial chickens showed resistance to ampicillin (75.0%), chloramphenicol (25.0%), ciprofloxacin (12.5%), enrofloxacin (37.5%), neomycin (50.0%), norfloxacin (12.5%), streptomycin (62.5%) and tetracycline (62.5%). The geometric mean MIC was highest (152.2 μ g/ μ L) for ampicillin and lowest (0.5 μ g/ μ L) for ciprofloxacin (Table 3).

All the five *Salmonella* isolates from apparently healthy commercial chicken were resistant to ampicillin, four (80.0%) were resistant to streptomycin, two (40.0%) showed resistance to each of chloramphenicol, enrofloxacin and neomycin while only one (20.0%) was resistant to each of ciprofloxacin, norfloxacin and tetracycline. The geometric mean MIC was highest (168.9 μ g/ μ L) for ampicillin and lowest (0.5 μ g/ μ L) for ciprofloxacin (Table 4).

Table 1: Antimic Abeokut	obial resistan a, Nigeria	ice in bacter	ial isolates fro	m apparently	healthy and o	diseased chicl	kens and turk	eys in
Bacterial isolates by source (number	Number (%) o	of resistant isol	ates					
tested)	Ampicillin	Chloram- phenicol	Ciproflox- acin	Enrofloxacin	Neomycin	Norfloxacin	Streptomycin	Tetracycline
Escherichia coli								
Apparently healthy chickens (172)	128.0 (74.4)	65.0 (37.8)	62.0 (36.1)	84.0 (48.8)	99.0 (57.6)	80.0 (46.5)	132.0 (76.7)	99.0 (57.6)
Diseased chickens	194.0 (100.0)	146.0 (75.3)	194.0 (100.0)	194.0 (100.0)	194.0 (100.0)	194.0 (100.0)	194.0 (100.0)	194.0
Diseased Turkeys	94.0 (89.5)	83.0 (79.1)	101.0 (96.2)	105.0 (100.0)	105.0 (100.0)	101.0 (96.2)	105.0 (100.0)	101.0 (96.2)
Subtotal (471)	416 (88.3)	294 (62.4)	357 (75.8)	383 (81.3)	398 (84.5)	375 (79.6)	431 (91.5)	394 (83.7)
Apparently healthy	6.0 (75.0)	2.0 (25.0)	1.0 (12.5)	3.0 (37.5)	4.0 (50.0)	1.0 (12.5)	5.0 (62.5)	5.0 (62.5)
Diseased chickens	14.0 (93.3)	12.0 (80.0)	14.0 (93.3)	15.0 (100.0)	14.0 (93.3)	15.0 (100.0)	15.0 (100.0)	15.0 (100.0)
Diseased Turkeys	5.0 (100.0)	4.0 (80.0)	4.0 (80.0)	5.0 (100.0)	5.0 (100.0)	5.0 (100.0)	5.0 (100.0)	5.0 (100.0)
Subtotal (28)	25 (89.3)	18 (64.3)	19 (67.9)	23 (82.1)	23 (82.1)	21 (75.0)	25 (89.3)	25 (89.3)
Salmonella serotypes Apparently healthy chickens (5) Diseased chickens	5.0 (100.0) -	2.0 (40.0) -	1.0 (20.0) -	2.0 (40.0) -	2.0 (40.0) -	1.0 (20.0) -	4.0 (80.0) -	1.0 (20.0)
Diseased Turkeys	ı	ı	ı		,	,	ı	1
Subtotal (5)	5.0 (100.0)	2.0 (40.0)	1.0 (20.0)	2.0 (40.0)	2.0 (40.0)	1.0 (20.0)	4.0 (80.0)	1.0 (20.0)
Overall total (504)	446 (88.5)	314 (62.3)	377 (74.8)	408 (81.0)	423 (83.9)	397 (78.8)	460 (91.3)	420 (83.3)

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Table 2: Mir healthy	imum inh	libitory concent	tration of	antimicrok	oial agents	for Escher	richia coli	i isolated fro	om apparently
Antimicrobial agents	Number of iso- late tested	Range of tested antimicrobial concentration	Lowest MIC (µg/ µL)	Highest MIC (µg/ µL)	Geometric mean MIC (µg/ µL)	MIC50 (µg/ µL)	MIC90 (µg/ µL)	Number (%) sensitive	Number (%) resistant
Ampicillin	172	0.25-512	0.5	>512.0	73.1	128.0	512.0	44.0 (25.6)	128.0 (74.4)
Chloram- phenicol	172	0.25-512	≤0.25	>512.0	22.3	16.0	256.0	107.0 (62.2)	65.0 (37.8)
Ciprofloxacin	172	0.25-512	≤0.25	>512.0	4.3	2.0	64.0	110.0 (63.9)	62.0 (36.1)
Enrofloxacin	172	0.25-512	≤0.25	>512.0	5.4	2.0	64.0	88.0 (51.2)	84.0 (48.8)
Neomycin	172	0.25-512	≤ 0.25	>512.0	17.7	32.0	256.0	73.0 (42.4)	99.0 (57.6)
Norfloxacin	172	0.25-512	≤0.25	>512.0	3.0	2.0	32.0	92.0 (53.5)	80.0 (46.5)
Streptomycin	172	0.25-512	1.0	>512.0	121.5	256.0	512.0	40.0 (23.3)	132.0 (76.7)
Tetracycline	172	0.25-512	≤0.25	>512.0	16.1	16.0.	512.0	73.0 (42.4)	99.0 (57.6)
Table 3: Mini inter	imum inhil Isively-rear	bitory concentra ed chickens in	tion of ant Abeokuta,	timicrobial Nigeria	agents for K	(lebsiella s _l	pp isolate	d from appar	ently healthy
Antimicrobial	Numbe	r Range of tester	d Lowest	Highest	Geometri	c MIC50	MIC90	Number	Number (%)
agents	of isola	te antimicrobial	MIC (μ	3/ MIC (με T)	5/ mean MIC	Iμ/gμ) C	Iμ /gμ) (.) (%) sonsitivo	resistant
	ורארת		(m	(m)	(mg/ hm)			3011311140	
Ampicillin	~ ~ ~	0.25-512	512.0	512.0	152.2 22 2	512.0	512.0	2.0 (25.0)	6.0 (75.0)
Chloramphenic	8 0 0	0.25-512	16.0 0.7	256.0	26.9	16.0 0.01	256.0	(0.9/) 0.9 (0.2 L)	2.0 (25.0)
Ciprolioxacin E proflovacin	000	0.25-512 0.35 513	۲.0 ۵ ر	0.4 0.0	0.0 0 c	0.2.U	0.4	(C.18) U.1 (3 C4) 0 3	(2.21) (1.1)
Neomycin	0 00	0.25-512	2.0 16.0	32.0	2.0 10.4	2.0 16.0	32.0	4.0 (50.0)	4.0 (50.0)
Norfloxacin	8	0.25-512	1.0	4.0	1.2	1.0	4.0	7.0 (87.5)	1.0 (12.5)
Streptomycin	8	0.25-512	128.0	128.0	69.8	128.0	128.0	3.0 (37.5)	5.0 (62.5)
Tetracycline	8	0.25-512	128.0	128.0	26.9	128.0	128.0	3.0 (37.5)	5.0 (62.5)

	Antimicrobial agents	Number of isolate tested	Range of tested antimicrobial concentration	Lowest MIC (µg/ µL)	Highest MIC (µg/ µL)	Geomet- ric mean MIC	MIC50 (µg/ µL)	MIC90 (µg/ µL)	Number (%) sensitive	Number (%) resistant
	Ampicillin	5	0.25-512	64.0	>512.0	168.9	128.0	512.0	0.0 (0.0)	5.0 (100.0)
	Chloram- phenicol	ß	0.25-512	0.5	128.0	13.9	8.0	128.0	3.0 (60.0)	2.0 (40.0)
	Ciprofloxacin	5	0.25-512	≤0.25	8.0	0.5	0.25	8.0	4.0 (80.0)	1.0 (20.0)
	Enrofloxacin	5	0.25-512	0.5	256.0	5.3	1.0	256.0	3.0 (60.0)	2.0 (40.0)
110	Neomycin	2	0.25-512	4.0	>512.0	16.0	4.0	512.0	3.0 (60.0)	2.0 (40.0)
	Norfloxacin	2	0.25-512	≤0.25	16.0	0.9	0.5	16.0	4.0 (80.0)	1.0 (20.0)
	Streptomycin	5	0.25-512	16.0	>512.0	147.0	256.0	512.0	1.0 (20.0)	4.0 (80.0)
	Tetracycline	5	0.25-512	0.5	128.0	6.1	4.0	128.0	4.0 (80.0)	1.0 (20.0)
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Antimicrobial resistance in bacterial isolates from diseased chickens:

All 194 *E. coli* isolates from diseased chickens were all resistant to tested antimicrobials except chloramphenicol to which 146 (75.3%) of the isolates were resistant. The geometric mean MIC was highest (156.4 $\mu g/\mu L$) in ampicillin and least (38.1 $\mu g/\mu L$) in ciprofloxacin (Table 5).

Klebsiella isolates from diseased chickens showed 100% resistance to enrofloxacin, norfloxacin, streptomycin and tetracycline; 93.3% resistance to ciprofloxacin, ampicillin and neomycin and 80.0% resistance to chloramphenicol. The geometric mean MIC was highest (406.4 μ g/ μ L) in ampicillin and least (46.3 μ g/ μ L) in ciprofloxacin (Table 6).

Antimicrobial resistance in bacterial isolates from diseased turkeys:

Escherichia coli isolates from diseased turkeys were all resistant to enrofloxacin, neomycin

and streptomycin. The rate of resistance to ciprofloxacin, norfloxacin and tetracycline was 96.2% each while resistance was 89.5% and 79.1% for ampicillin and chloramphenicol respectively. The geometric mean MIC was highest (146.1 μ g/ μ L) in streptomycin and least (95.7 μ g/ μ L) in norfloxacin (Table 7).

All five (100%) *Klebsiella* isolates from diseased turkeys were resistant to ampicillin, enrofloxacin, neomycin, norfloxacin and tetracycline while four (80.0%) were resistant to chloramphenicol and ciprofloxacin. The geometric mean MIC was highest (512.0 μ g/ μ L) in ampicillin and least (55.7 μ g/ μ L) in ciprofloxacin (Table 8).

Table 5: Minii intens	mum inhil sively-rear	bitory concentra ed chickens in A	tion of anti Nbeokuta, r	imicrobial Nigeria	agents for E	scherichi	a coli isola	ted from dis	eased
Antimicrobial agents	Number of iso- late tested	Range of tested antimicrobial concentration	Lowest MIC (µg/ µL)	Highest MIC (µg/ µL)	Geometric mean MIC (µg/ µL)	MIC50 (µg/ µL)	MIC90 (µg/ µL)	Number (%) sensitive	Number (%) resistant
Ampicillin	194	0.25-512	128.0	>512.0	156.4	128.0	128.0	0.0 (0.00)	194.0 (100.0)
Chloram- phenicol	194	0.25-512	8.0	>512.0	67.3	8.0	128.0	48.0 (24.7)	146.0 (75.3)
Ciprofloxacin	194	0.25-512	16.0	64.0	38.1	16.0	64.0	0.0 (0.00)	194.0 (100.0)
Enrofloxacin	194	0.25-512	64.0	>512.0	111.4	64.0	128.0	0.0 (0.00)	194.0 (100.0)
Neomycin	194	0.25-512	16.0	256.0	66.3	16.0	128.0	0.0 (0.00)	194.0 (100.0)
Norfloxacin	194	0.25-512	16.0	>512.0	83.4	16.0	128.0	0.0 (0.00)	194.0 (100.0)
Streptomycin	194	0.25-512	128.0	>512.0	139.5	128.0	128.0	0.0 (0.00)	194.0 (100.0)
Tetracycline	194	0.25-512	64.0	>512.0	112.6	64.0	128.0	0.0 (0.00)	194.0 (100.0)

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seased	er Number (%) resistant e) 14.0 (93.3)	12.0 (80.0)) 14.0 (93.3)) 15.0 (100.0)) 14.0 (93.3)) 15.0 (100.0)) 15.0 (100.0)) 15.0 (100.0)	
from dis	Numbe (%) sensitiv	1.0 (6.7)	3.0 (20 0)	1.0 (6.7)	0.0) 0.0	1.0 (6.7)	0.0) (0.0)	0.0) (0.0)	0.0) 0.0	
pp isolated	MIC90 (µg/ µL)	512.0	128.0	64.0	128.0	128.0	512.0	512.0	512.0	
ebsiella s	MIC50 (µg/ µL)	512.0	128.0	64.0	64.0	64.0	64.0	256.0	256.0	
agents for Kl	Geometric mean MIC (µg/ µL)	406.4	67.0	46.3	67.0	55.7	70.2	222.9	185.3	
imicrobial a Nigeria	Highest MIC (µg/ µL)	>512.0	128.0	64.0	128.0	128.0	>512.0	>512.0	>512.0	
ition of anti Abeokuta,	Lowest MIC (µg/ µL)	16.0	0.5	2.0	32.0	8.0	16.0	64.0	16.0	
itory concentra ed chickens in	Range of tested antimicrobial concentration	0.25-512	0.25-512	0.25-512	0.25-512	0.25-512	0.25-512	0.25-512	0.25-512	
num inhib sively-rear	Number of isolate tested	15	15	15	15	15	15	15	15	
Table 6: Minin intens	Antimicro- bial agents	Ampicillin	Chloram- nhenicol	Ciprofloxacin	Enrofloxacin	Neomycin	Norfloxacin	Streptomycin	Tetracycline	
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nia coli isolated from diseased	MIC90 Number Number) (µg/ µL) (%) (%) resistant sensitive	128.0 11.0 (10.5) 94.0 (89.5)	128.0 22.0 (20.9) 83.0 (79.1)	128.0 4.0 (3.8) 101.0 (96.2)	128.0 0.0 (0.0) 105.0 (100.0)	128.0 0.0 (0.0) 105.0 (100.0)	128.0 4.0 (3.8) 101.0 (96.2)	128.0 0.0 (0.0) 105.0 (100.0)	128.0 4.0 (3.8) 101.0 (96.2)	
scherich	MIC50 (µg/ µL)	16.0	16.0	64.0	64.0	128.0	64.0	128.0	128.0	
agents for E	Geometric mean MIC (µg/ µL)	107.1	9.66	97.0	104.3	117.5	95.7	146.1	124.7	
microbial ligeria	Highest MIC (µg/ µL)	>512.0	>512.0	128.0	256.0	>512.0	256.0	>512.0	>512.0	
tion of anti beokuta, N	Lowest MIC (µg/ µL)	16.0	16.0	2.0	32.0	16.0	2.0	128.0	8.0	
itory concentra I chickens in A	Range of tested antimicrobial concentration	0.25-512	0.25-512	0.25-512	0.25-512	0.25-512	0.25-512	0.25-512	0.25-512	
um inhibi vely-reared	Number of isolate tested	105	105	105	105	105	105	105	105	
Table 7: Minir intensi	Antimicrobial agents	Ampicillin	Chloram-	Ciprofloxacin	Enrofloxacin	Neomycin	Norfloxacin	Streptomycin	Tetracycline	

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Antimicrobial agents	Number of isolate tested	Range of tested antim- icrobial con- centration	Lowest MIC (µg/ µL)	Highest MIC (µg/ µL)	Geometric mean MIC (μg/ μL)	MIC50 (µg/ µL)	MIC90 (µg/ µL)	Number (%) sensitive	Number (% resistant
Ampicillin	5	0.25-512	512.0	>512.0	512.0	512.0	512.0	5.0 (100.0)	5.0 (100.0)
Chloram-	5	0.25-512	8.0	256.0	128.0	256.0	256.0	4.0 (80.0)	4.0 (80.0)
Ciprofloxacin	Ŋ	0.25-512	2.0	128.0	55.7	128.0	128.0	4.0 (80.0)	4.0 (80.0)
Enrofloxacin	5	0.25-512	4.0	128.0	64.0	128.0	128.0	5.0 (100.0)	5.0 (100.0)
Neomycin	2	0.25-512	32.0	256.0	168.9	256.0	256.0	5.0 (100.0)	5.0 (100.0)
Norfloxacin	5	0.25-512	64.0	128.0	111.4	128.0	128.0	5.0 (100.0)	5.0 (100.0)
Streptomycin	5	0.25-512	256.0	>512.0	445.7	512.0	512.0	5.0 (100.0)	5.0 (100.0)
Tetracycline	Ъ	0.25-512	128.0	256.0	147.0	128.0	256.0	5.0 (100.0)	5.0 (100.0)

DISCUSSION

In the present study, E. coli was the predominant isolate from apparently healthy and diseased poultry birds. This agrees with earlier report by Kilonzo-Nthenge et al. (2008) that E. coli is a major cause of morbidity and mortality in poultry worldwide (Kilonzo-Nthenge et al., 2008). Apart from being a primary cause of diseases, E. coli is also implicated as an opportunistic pathogen capable of complicating infections caused by other pathogens. Other bacteria identified in the present study (Salmonella enterica isolates and Klebsiella spp.) are also known to induce clinical diseases in poultry (Kilonzo-Nthenge et al., 2008). Salmonella species was the least encountered and was detected only in apparently healthy chickens. Although a major avian pathogen of high economic importance, the presence of Salmonella in apparently healthy birds showed that birds may harbour Salmonella without clinical manifestations (Agbaje et al., 2010). Apparently healthy carriers may thus serve as sources of persistent Salmonella infection in the flock.

The present study showed varying degrees of antimicrobial resistance in bacterial species isolated from apparently healthy chickens in the study area. In all the bacteria species, the highest rates of antimicrobial resistance and geometric mean MIC were recorded in ampicillin and streptomycin. Ampicillin and streptomycin are first generation antimicrobials which are commonly used antimicrobials in the livestock industry. This may account for the higher rates of resistance to these drugs. Generally, there was moderate level of neomycin resistance which did not exceed 50.0% except in E. coli (57.6%). Among the antimicrobials from different classes represented in this study, resistance to the fluoroguinolones was rela-

tively low. However, it is evident that bacteria are developing resistance to the fluoroquinolone which are considered the drug of choice for the treatment of gastroenteritis in humans (Guerrant et al., 2001). Among fluoroguinolones (ciprofloxacin, enrofloxacin and norfloxacin) resistance rate and geometric mean MIC were observed to be highest for enrofloxacin and lowest for ciprofloxacin. Enrofloxacin resistance rate was as high as 48.8% in *E. coli* and ciprofloxacin resistance as low as 12.5% in *Klebsiella* spp. A previous study in Nigeria showed that enrofloxacin is the most commonly administered fluoroguinolones in poultry production in the study area (Ogunleye et al., 2008). This may be responsible for the higher rates of resistance to enrofloxacin than to other fluoroquinolones. Fluoroquinolone resistant avian E. coli has been reported in other regions of the world (White et al., 2000; Thorsteinsdottir et al., 2010; Chen et al., 2011) and may be transmitted to humans through the food chain (Warren et al., 2008). The continued efficacy of fluoroguinolone therapy in the treatment of human diseases can be achieved by regulating the use of these drugs in humans and disallowing their use in food animals (Cheng et al., 2012).

The presence of drug resistant bacteria in apparently healthy chicken as observed in the present study has implications for poultry production and public health. Nonpathogenic resistant bacteria resident in apparently healthy birds may share their resistant trait and confer resistance on virulent pathogens or acquire virulent traits from pathogenic bacteria (Yaron *et al.*, 2000; Osterloh, 2004). Exchange of resistance and virulence genes is common among enteric bacteria especially the enterobacteriaceae (Balis *et al.*, 1996; Yaron *et al.*, 2000). Close contact between humans and birds, consumption of contaminated poultry products and environmental contamination may increase the possible transmission of resistant bacteria more so it has been reported that transmission of resistant clones and plasmids from poultry to humans is a common occurrence (Van dan Boggard *et al.*, 2001).

The overall high rates of antimicrobial resistance observed among the bacterial isolates in this study were due largely to high resistance observed in isolates from diseased birds. When considered separately, resistance rates were significantly lower (p < 0.05) in isolates from apparently healthy birds than in isolates from diseased birds. The present study suggests that antimicrobial resistant bacteria may predominate in disease outbreaks. Antimicrobial resistant Salmonella spp. have been reported to be more invasive than susceptible strains thereby producing more severe and fatal infections (Helms et al., 2004). The present study investigated outbreaks of diarrhoea refractory to antimicrobial therapy accompanied by high mortality of over 60%. Escherichia coli was isolated from all the clinical samples submitted for bacteriological examination. Few isolates of *Klebsiella* spp were also obtained from the samples. High levels of antimicrobial resistance of between 80.0% and 100% (100% in most cases) were observed among the bacterial isolates. Involvement of multi-drug resistant bacteria in disease outbreak as observed in the present study could undermine the efficacy of therapeutic intervention in the control of bacterial infections. The direct effect of orally administered drug on enteric bacteria may alter the integrity of gastrointestinal microflora leading to the eradication of susceptible strains and proliferation of resistant ones (Zhoa et al., 2001). This may lead to an increase in the prevalence of antimicrobial resistance in

isolates recovered from animals while on antimicrobial therapy. In the present study, antimicrobial agents had been administered to the sick birds before sample collection and as such may account for the high level of antimicrobial resistance in the bacterial isolates (Boothe and Debavalya 2011). Antimicrobial usage during an outbreak may therefore eliminate competing susceptible bacteria co-habiting the gut with resistant pathogens. This will aid the proliferation of the resistant pathogen and increase the damage done to the host. The high rates of antimicrobial resistance in bacteria may also contribute to the persistence of pathogens in poultry flock because of the ineffectiveness of chemoprophylactic eradication approach.

CONCLUSION

The present study showed high level of antimicrobial resistance in clinical and nonclinical bacterial isolates from intensively reared birds in the study area. The major factors selecting for antimicrobial resistance in bacteria are antimicrobial use, overcrowding and poor sanitation. These factors are typical of many intensive poultry farming and may explain the high prevalence of antimicrobial resistance in bacteria as encountered in this study. Antimicrobial resistance in avian bacterial pathogens is a threat to profitable poultry production, protein availability and public health.

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