

Research Article

Newcastle Disease in Local Chickens of Live Bird Markets and Households in Zamfara State, Nigeria

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Received 30 November 2013; Accepted 18 December 2013; Published 16 January 2014

Academic Editors: Q. Chen and M. Lancellotti

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Newcastle disease constitutes a major constraint to rural poultry production system in Nigeria. This study used serological method to estimate the level of circulating antibodies against ND in nonvaccinated village chickens, raised under traditional management system in Zamfara State, Nigeria. Competitive Enzyme Linked Immunosorbent Assay was used to analyze 504 chicken sera for Newcastle disease virus antibodies from randomly selected households and live bird markets. Higher seroprevalence rate of Newcastle disease virus antibodies was detected in both household and live bird markets. Overall, seropositive rate was found to be 32.5% (164/504). About 35.8% (115/321) sero-positive rate was obtained from live bird markets while 26.8% (49/183) seropositive rate was found in households. Comparison was made between the sero prevalence of house hold and live bird markets as well as between sexes. Live bird markets show a statistically significant higher prevalence rate ($P < 0.05$) when compared with chickens sampled from households (OR 1.53; 95% CI, 1.024–2.275). The prevalence of ND indicated the presence of the virus amongst the population, and hence there should be an improvement in the vaccine campaign against ND for rural poultry especially the use of thermostable vaccine to reduce the chances of vaccine failure.

1. Introduction

Newcastle disease (ND) is a major viral disease of economic importance in poultry [1] and rated as one of the greatest constraints to the development of rural poultry production in Nigeria and in most developing countries, causing serious threats [2]. All ages of different species of birds are susceptible to ND, although being substantially less with advancement to maturity [3]. The acute and virulent form may result in 90% mortality or more in affected flocks [4]. It is an acute, rapidly spreading, contagious, nervous and respiratory disease of birds of all ages [5]. The clinical signs of ND are known to vary based on the virulence and tropism of the ND virus involved, species of the bird, age of the host, immune status, and environmental condition [6].

Nigeria poultry population is estimated to be 137.6 million, with backyard poultry population constituting 84% (115.8 million) and 16% (21.7 million) of exotic poultry, with a higher percentage of this poultry raised for subsistence

production [7]. Village poultry production provides an important source of high quality protein, is reserved for times of celebrations, and is a good source of income for rural families [8]. This category of birds represents a significant part of the Nigerian rural economy in particular and of the national economy as a whole and is kept under the extensive management system [9]. The resources derivable from the chickens cannot be fully utilized unless the disease is controlled particularly in the village poultry flocks that are believed to keep the virus in circulation and act as reservoirs and carriers to themselves and the more susceptible exotic breeds in commercial farms [10].

2. Materials and Methods

2.1. Study Area. The study was carried out in six local governments (Bungudu, Gusau, Talata-Mafara, Bakura, Kaura-Namoda, and Zurmi) of Zamfara State, Nigeria. Zamfara State is located between latitudes 10°50'N and 13°58'N and

TABLE 1: Prevalence of Newcastle disease from selected local governments of Zamfara State.

S/N	Local governments	Number of samples	Number of positives	Number of negatives	Prevalence (%)
1	Gusau	100	56	44	56.0
2	Bungudu	70	34	36	48.5
3	Zurmi	75	26	49	34.7
4	Kaura-Namoda	99	14	85	14.1
5	Bakura	70	12	58	17.1
6	Talata-Mafara	90	22	68	24.4
Total		504	164	340	32.5

Chi-square value = 58.94.

P value = 0.001.

TABLE 2: Prevalence of Newcastle based on location in Zamfara State.

S/N	Location	Number of samples	Number of positives	Number of negatives	Prevalence (%)	Odds ratio	95% CI
1	LB markets	321	115	206	35.8	1.53*	1.024–2.275
2	Households	183	49	134	26.8	1.00	
Total		504	164	340	32.5		

Chi-square value = 3.946.

P value = 0.0470.

*Statistical significance between variables when using odd ratio.

longitudes 4°16 E and 7°13 E. It has a warm tropical climate between March and May. The vegetation of the state consists of Sudan and Northern Guinea Savannah. It has a total human population of 3,278,873 [11] and an estimated poultry population of 5 845,508 [7]. The state has 14 local governments and has 3 senatorial districts.

2.2. Sera Collection. Multistage sampling procedure was applied to divide the state into three stages corresponding to three senatorial districts. Two local governments were then randomly selected from each district, making a total number of six local governments that were considered in this study. A total of 504 sera samples were collected from live bird markets and households from all the six selected local governments across the state. Two milliliter of chicken blood was collected through the brachial vein using a 21-gauge sterile hypodermic needle and 2 mL syringes. The syringes were labeled and kept in slanting position till clot formation. Sera were harvested, transferred into a sterile serum bottles, and stored at -20°C . All sera at the end of the sampling were taken to the viral zoonosis laboratory of the Department of Veterinary Public Health and Preventive Medicine, Ahmadu Bello University, Zaria, for serology.

2.3. Serology. Competitive ELISA diagnostic kit was obtained from ID Vet, Innovative Diagnostics, France. The kit procedure is based on blocking Enzyme Linked Immunosorbent Assay. 100 μL of positive control was added to positive control wells A1 and B1 and 100 μL of negative control to negative control wells C1 and D1 using micropipette. 80 μL of dilution buffer 2 was added to each of the wells. Using multichannel pipette, 20 μL of undiluted serum was then added to the remaining selected wells and mixed thoroughly

with the buffer. The plates were then sealed and incubated at room temperature for 30 minutes. Anti-NDV conjugate 10x was prepared by diluting anti-NDV-HRP conjugate to 1/10 dilution buffer 5. After incubation the plates were washed 3 times with wash solution. Using multichannel pipette, 100 μL of anti-NDV conjugate 1x was added to each well. The plates were rinsed three times with wash solution. 100 μL of the substrate solution was added to each well. And this was incubated at room temperature for ten minutes in the dark. The reaction was stopped by adding 50 μL of stop solution to each well which is mixed thoroughly. Optical density was read at 450 nm by using microplate reader.

2.4. Data Analysis. The data obtained was subjected to SPSS package version 16. Categorical variables (sex, location) were evaluated using Chi-square to check for independence, and odds ratio at 95% confidence interval was used to measure the strength of association between variables and prevalence of Newcastle disease. Values of $P < 0.05$ were considered significant.

3. Results

The overall seroprevalence of Newcastle disease in the study area was 32.5%. The highest prevalence was found in Gusau local governments (56.0%) while the least was detected in Kaura-Namoda local governments (14.1%) (Table 1). Table 2 indicates that chickens from live bird markets show a higher prevalence rate (35.8%) when compared to those sampled from households (26.8%). Prevalence of 35.7% was recorded in male chickens while 28.9% was obtained from female chickens as indicated in Table 3.

TABLE 3: Sex-specific prevalence of Newcastle disease in Zamfara State.

S/N	Sex	Number of samples	Number of positives	Number of negatives	Prevalence (%)	Odds ratio	95% CI
1	Male	272	97	175	35.7	1.37	0.936–1.990
2	Female	232	67	165	28.9	1.0	

Chi-square value = 2.324.

P value = 0.1274.

4. Discussion

The present serological study revealed the presence of circulating antibodies of Newcastle disease among sampled village chickens from selected local governments of Zamfara State. An overall seroprevalence of 32.5% from the six local governments of Zamfara State is an indication of endemicity. Antibodies detected may be a result of natural infection since vaccination of the village poultry is rarely undertaken in Nigeria [12]. In similar studies, [13] reported a variable sero prevalence of 25–81.5% in Tanzania, in Ethiopia, 43.68% seropositive rate of NDV was reported in the cool central highlands [14] while [15] reported 19.78%, [16] reported 14% sero prevalence in non-vaccinated village chickens in Niger. Reference [17] reported a prevalence of 63% in south eastern Nigeria. Reference [18] reported a prevalence of 46% in village chickens in Borno State and [19] reported an ND prevalence of 54.67% in Nasarawa State. This finding of NDV sero prevalence in these apparently healthy birds suggests that the birds have either recovered from clinical ND or are having subclinical infections [20]. A similar study was conducted in Ibadan using indirect ELISA and an overall prevalence rate of 73.3% was obtained [21]. Although there is a statistically significant ($P < 0.05$) higher prevalence in Gusau compared to other local government areas (LGAs), the higher circulating antibodies in Gusau may be due to higher concentration of live bird markets in this LGA. The lower prevalence rate in Bakura, Talata-Mafara, and Kaura-Namoda may be due to vaccination activities of commercial poultry that may contribute to mild infection due to the spread of vaccine virus to local birds through commercial poultry workers [19].

Live bird markets show a statistically significant higher prevalence rate ($P < 0.05$) when compared with chickens sampled from households (OR 1.53; 95% CI, 1.024–2.275). This may be due to the fact that live bird markets contribute to the persistence and spread of ND virus. These birds are exposed to birds from multiple sources having a higher tendency of circulating the virus and may serve as a source of infection to house hold chickens when introduced [22]. Similar studies conducted by [23] show 25.5% prevalence in live bird markets of Kogi State, Nigeria.

This study also shows a higher prevalence rate among the male chickens. It was however observed that this difference is statistically insignificant ($P > 0.05$). In a similar study conducted by [15] in Ethiopia, ND shows a higher prevalence rate among males (21.74%) than among females (19.16%). However, a slightly higher prevalence of 32.63% among female chickens was obtained by [24] when compared with a prevalence of 31.63 among male chickens.

Acknowledgments

The authors would like to thank all the farmers that allowed them to take samples from their chickens and to all staff of the department of Veterinary Public Health and Preventive Medicine, Ahmadu Bello University, for their technical support.

Conflict of Interests

There is no competing interest whatsoever that could have influenced the results of this study in any manner.

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