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Determination of immune response of Newcastle disease virus vaccines in layer chickens

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Abstract

The study was conducted for the detection of persistence of Maternally derived antibody (MDA) as well as the comparative evaluation of antibody production of nine different NDV vaccines in layer chickens in the Department of Microbiology and Hygiene, Bangladesh Agricultural University, Mymensingh during the period from July to December 2008. A total of 55 layer chicks (ISA Brown breed) were divided into eleven groups each consisting of five birds of which odd number groups were vaccinated primarily with Nobilis[®] MA5+Clone 30, Avipro[®] ND-IB HB1, Cevac[®] BIL, Newcastle-Bronchitis Vaccine Fortdodge[®] and Avipro[®] ND LaSota vaccine respectively at day 5 of age and secondarily with Nobilis[®] ND Clone30, Avipro[®] ND LaSota, Cevac[®] New L, Newcastle Disease Vaccine Fortdodge[®] and Avipro[®] ND LaSota vaccine respectively at day 21 of age by single eye instillation and even number groups were vaccinated with the same vaccines respectively by double eye instillation following the same schedule. Again group 9 and group 10 were also vaccinated with RDV at 60 days of age through intramuscular route. Group 11 was kept as unvaccinated control. Sera samples were collected after 10 days of each vaccination and at day 5, 15, 20, 31 of age from unvaccinated control and subjected to HI test for the determination of antibody titres. It was observed that after primary vaccination the mean of HI titres of double eye vaccinated groups differed significantly ($P < 0.01$). Overall analysis of mean of HI titres of double eye vaccinated groups revealed that there were significant increases ($P < 0.01$) in HI titres in groups 8 (891.44 ± 228.97) and 10 (861.66 ± 140.21) compared to other groups. It was observed that secondary vaccination produced higher immune response compared to primary vaccination in case of all the vaccinated groups and double eye vaccination produced higher immune response compared to single eye vaccination in case of all the vaccinated groups. It was also observed that following vaccination with RDV in groups 9 and 10, HI titres increased significantly ($P < 0.01$) which indicated that group 10 (1204.30 ± 280.43) produced significantly higher antibody titres than group 9 (966.74 ± 144). Maternal antibody was high (483.37 ± 181.01) at day 5 of age and persisted to a minimal level (8.00 ± 0.00) until the age of day 20 and almost disappears ($\leq 4 \pm 0$) at day 31 of age. From the present research it may be concluded that LaSota strain produced higher immune response than Clone 30 and B₁ strain, Fortdose[®] and Avipro[®] vaccine produced higher immune response than all other vaccines and vaccination with lentogenic strains followed by mesogenic strain produced higher antibody titers used in this study.

Keywords: Immune response, Newcastle disease, Vaccines, Layer chickens

Introduction

Newcastle disease (ND) is a deadly viral disease of poultry all over the world since the time of first isolation in England in 1926 till today. It is also considered as one of the major economic threats to poultry population because of its high morbidity and mortality varies from 90-100% depending on the type of ND virus exposed (Brandly, 1950). It is an acute, contagious infection of pet, free living and domestic birds. Effective control of ND relies on the use of safe and good vaccines. Live vaccines prepared with lentogenic and mesogenic strain of NDV are now more commonly used in both broiler and layer birds. This is because live vaccines can be produced on a large scale at a relatively low cost. The vaccines are easy to administer on a large scale, and rapidly stimulate humoral, cell-mediated and mucosal surface immunity in the vaccinated birds. Vaccination as directed by Chowdhury *et al.* (1981) includes administration of live lentogenic vaccine (BCRDV) of F-strain by *i/o* inoculation to first week and 21 days old chicks followed by a live mesogenic vaccine (RDV) of Mukteswar strain by intramuscular route at day 60 and are repeated at every 6 months interval for the maintenance of sustainable level of neutralizing antibody in the serum of vaccinated birds. In Bangladesh, various live vaccines containing lentogenic strains of NDV such as B₁, Lasota, VG/GA, ND-clone are brought by several importer but efficacy of these vaccines in relation to climatic condition, distribution and transportation are not always investigated properly and thoroughly either by the importer or by the user. Sometimes, the farmers/ rearers are suspicious about the protective nature of those imported NDV vaccines. A number of relevant questions are faced by scientists and field Veterinarians of this country as to the immunogenicity, virus titre, stability and such other qualities of those vaccines. In order to address such query, the present study was undertaken to determine the persistence of maternally derived antibody in the layer chicks, to compare the antibody titres of chickens vaccinated with eight different imported vaccines of lentogenic strains of NDV and also to determine the antibody titres of chickens vaccinated with lentogenic and followed by mesogenic strain of NDV vaccines used in this study.

Materials and Methods

The whole experiment was conducted during the period from July to December 2008 in the Department of Microbiology and Hygiene, Faculty of Veterinary Science, Bangladesh Agricultural University, Mymensingh.

Experimental chickens

A total number of 55 day-old-chicks of ISA Brown breed with the history of parent stock vaccinated against Newcastle disease (ND) were obtained with the curtesy of C.P. Bangladesh Co. Ltd., Dhaka and carried to the experimental animal shed of the Department of Microbiology and Hygiene, BAU, Mymensingh. The birds were reared in litter supplying Nurish[®] poultry feed and water maintaining strict bio-security. Before vaccination the birds were divided into 11 groups, each group consists of five birds. The chicks of group 11 were kept as unvaccinated control.

Newcastle disease virus vaccines

Nine lyophilized (freeze-dried) NDV vaccines were used in this study. These were (i) Nobilis[®] MA5+Clone 30 (Mass and clone 30 strain) of Intervet International (ii) Avipro[®] ND-IB HB1 (Mass and B1 strain) of Lohmann Animal Health (iii) Cevac[®] BIL (Mass and B1 strain) of Ceva pharmaceuticals (iv) Newcastle-Bronchitis Vaccine Fortdodge[®] (Mass and B1 strain) of Fortdodge Animal Health (v) Nobilis[®] ND Clone30 (Clone 30 strain) of Intervet International (vi) Avipro[®] ND LaSota (LaSota strain) of Lohmann Animal Health (vii) Cevac[®] New L (LaSota strain) of Ceva pharmaceuticals (viii) Newcastle Disease Vaccine Fortdodge (LaSota strain) of Fortdodge Animal Health and (ix) RDV (M strain) of DLS. Vaccines were obtained with the courtesy of C.P. Bangladesh Co. Ltd. These vaccines were stored at -20°C until used.

Reference Newcastle disease virus

Newcastle disease virus (Komarov strain) used in microplate HI test for the determination of antibody titre were collected from the laboratory repository of the Department of Microbiology and Hygiene, BAU, Mymensingh was propagated to activate the virus into 10- day- old embryonated chicken eggs through allantoic cavity route. AF was collected aseptically using sterile syring and needle transferred to sterile containers. The presence of NDV in AF was tested by slide HA test and those manifesting HA- positive were collected (Cottral, 1978) as a source of reference ND virus and stored at -20°C until used.

Vaccination schedule

A total of 55 layer chicks (ISA Brown) were divided into eleven groups such as 1,2,3,4, 5, 6, 7, 8, 9, 10 and 11 where each group consisting of five birds. Group 1, 3, 5, 7 and 9 were primarily vaccinated with Nobilis[®] MA5+Clone 30, Avipro[®] ND-IB HB1, Cevac[®] BIL, Newcastle-Bronchitis Vaccine Fortdodge[®] and Avipro[®] ND LaSota vaccine respectively at day 5 of age and secondarily with Nobilis[®] ND Clone30, Avipro ND LaSota, Cevac[®] New L, Newcastle Disease Vaccine Fortdodge[®] and Avipro[®] ND LaSota vaccine respectively at day 21 of age by single eye instillation and 2, 4, 6, 8 and 10 were vaccinated with the same vaccines respectively by double eye instillation following the same schedule. Again group 9 and group 10 were also vaccinated with RDV at 60 days of age through intramuscular route. Group 11 was kept as unvaccinated control. Sera samples were collected after 10 days of each vaccination and at day 5, 15, 20, 31 of age from unvaccinated control and subjected to HI test for the determination of antibody titres.

Microplate haemagglutination inhibition (HI) test

Micro-plate HI test was performed (Anon, 1971) to determine the antibody level of the sera samples collected from the chicks of different groups. The test was conducted using constant 4HA unit ND virus and diluted serum (β method).

Statistical analysis

Statistical Package for social Sciences (SPSS) version 10.0 programme was used to analyze the data of the study. ANOVA Duncan's Multiple Range Test (DMRT) and Paired 't' test were performed to determine the significant differences in HI titres of chickens of a group after primary and secondary vaccination (Beri, 2005).

Results and Discussion

Maternally derived antibody (MDA) was measured on a number of occasions starting from day 5 to day 31 and the titres are presented in Table 1. It was found that the MDA level persisted at a suitable level upto 15 days and it started to decline after 15 days and reached at a negligible level after 20 days. This finding strongly supports the findings of Shil 2006; Balla 1986; Saeed *et al.*, 1988, Chowdhury *et al.*, 1982 and Islam *et al.*, 2003 who stated that the persistence of MDA in chickens were day 15 to 20 of age. But Mahud *et al.*, 2007 reported that the persistence of MDA upto 27 days of age which may be due to the high MDA titres.

Table 1. Maternal antibody of unvaccinated control of layer chickens (ISA Brown) determined by HI test

No. of serum sample	Age of birds							
	Day 5		Day 15		Day 20		Day 31	
	Titre	Mean \pm SD (n = 5)	Titre	Mean \pm SD (n = 5)	Titre	Mean \pm SD (n = 5)	Titre	Mean \pm SD (n = 5)
1	512	602.15 \pm 140.21	96	75.26 \pm 17.25	8	8.00 \pm 0.00	\leq 4	\leq 4 \pm 0
2	512		96		8		\leq 4	
3	768		64		8		\leq 4	
4	768		64		8		\leq 4	
5	512		64		8		\leq 4	

n = Number of sera samples tested.

SD=Standard deviation

HI= Haemagglutination inhibition.

The principal objective of this study was to observe the immune response of live NDV vaccines (both lentogenic and mesogenic strain) in layer chickens commonly used in Bangladesh. It was observed that the mean and Standard deviation of HI titres after primary vaccination by single eye instillation in groups 1, 3, 5, 7 and 9 were 691.69 \pm 214.18, 187.52 \pm 45.25, 388.02 \pm 140.21, 483.37 \pm 57.24 and 375.05 \pm 90.50 respectively and after 10 days of secondary vaccination 345.84 \pm 107.09, 483.37 \pm 57.24, 555.24 \pm 114.48, 602.15 \pm 114.21 and 841.60 \pm 228.97 respectively whereas birds of double eye instillation vaccinated groups 2, 4, 6, 8 and 10 the mean of the titres were 794.52 \pm 214.18, 345.84 \pm 107.09, 354.08 \pm 57.24, 555.24 \pm 114.48 and 337.79 \pm 140.21 respectively and after 10 days of secondary vaccination 397.27 \pm 107.9, 794.54 \pm 214.18, 813 \pm 114.48, 891.44 \pm 228.97 and 861.66 \pm 140.21 respectively (Table 2).

It was observed that double eyes vaccinated groups of layer chickens produced significantly ($P < 0.01$) higher immune response compare to those of single eye vaccinated chickens. This increase immune response may be due to the amount of virus present in each dose. The vaccine virus concentration is higher in double eye dose than to the single eye dose which played a vital role in the production of high immune response. For better immune response the dose of vaccine was important factor which helps to increase in the level of antibody production which is closely related to the finding of Bankowski and Hill, 1954 and Winterfield and Seadle, 1957.

After secondary vaccination it was also observed that the antibody titres increased in all groups except group 1 & 2 where the antibody level declined. So, cloned 30 might not immunogenic than other strains (B and LaSota strains). Kafi *et al.* (2003) found the highest HI titres after secondary vaccination with BCRDV. Shuaib *et al.* (2003) reported that secondary vaccination yielded HI titres that was significantly higher than the HI titres after single vaccination.

A comparative analysis of all the vaccines of different strains revealed that LaSota strain produced higher immune response than those of other strains- B₁ and Clone 30 strain. This finding is strongly supported to the findings of Almasy *et al.* (1979) and Mallick *et al.* (1969) who reported that LaSota strain provided superior antibody production after vaccination compared to B₁. Westbury *et al.* (1984) also observed that LaSota is much more immunogenic than the Hitchner B₁. However, Ibrahim (1983) did not find any significant difference in immunizing capacity, protection and stress between F strain and B₁ strain.

Table 2. Comparative HI titres of chickens between single and double eyes vaccination with different NDV vaccines

Groups	Vaccines used		Dose and route of vaccination	Serum HI titre (Mean \pm SD) (n = 5)		Level of significance
	Primary vaccination	Secondary vaccination		Post-primary vaccination ^A (15 days old)	Post-secondary vaccination ^B (31 days old)	
1	Nobilis [®] MA5+Clone30	Nobilis [®] ND Clone30	One drop in one eye	691.69 \pm 214.18	345.84 \pm 107.09	B
2			One drop in two eyes	794.54 \pm 214.18	397.27 \pm 107.09	B
3	Avipro [®] ND-IB HB1	Avipro [®] ND LaSota	One drop in one eye	187.52 \pm 45.25	483.37 \pm 57.24	C
4			One drop in two eyes	345.84 \pm 107.09	794.54 \pm 214.18	B
5	Cevac [®] BIL	Cevac [®] New L	One drop in one eye	388.02 \pm 140.21	555.24 \pm 114.48	B
6			One drop in two eyes	354.08 \pm 57.24	813 \pm 114.48	B
7	Newcastle- Bronchitis Vaccine Fortdodge [®]	Newcastle Disease Vaccine Fortdodge [®]	One drop in one eye	483.37 \pm 57.24	602.15 \pm 140.21	B
8			One drop in two eyes	555.24 \pm 114.48	891.44 \pm 228.97	A
9	Avipro [®] ND LaSota	Avipro [®] ND LaSota	One drop in one eye	375.05 \pm 90.50	841.60 \pm 228.97	AB
10			One drop in two eyes	337.79 \pm 140.21	861.66 \pm 140.21	A

n = Number of sera samples tested.

^A Primary vaccination of all the groups was done at 5 days old.

^B Secondary vaccination of all the groups was done at 21 days old.

Figures with different letter within column 7 differ significantly (P<0.01).

SD=Standard deviation

HI= Haemagglutination inhibition.

Birds of all vaccinated groups did not respond equally to vaccination in the same manner and individual variation in the humoral immune response was observed within the same group of birds at the same interval following both primary and secondary vaccination, which corresponds with the findings of Saifuddin *et al.* (1986). The variation also might be due to the presence of variable MDA in chicks or varying degree of susceptibility of immune mechanism to antigen as observed by Toth and Markovits (1964).

Overall consideration among the vaccinated groups it was observed that the HI titres of birds of groups vaccinated primarily with Newcastle-Bronchitis Vaccine Fortdodge[®] and secondarily with Newcastle Disease Vaccine Fortdodge[®] by double eyes dose and group 10 vaccinated primarily and secondarily with Avipro[®] ND LaSota by double eye dose produced higher immune response than all other vaccinated groups.

The present study also revealed that groups 9 and 10 after third vaccination with RDV (M-strain) at 60 days of age through intramuscular route showed significantly (P<0.01) higher immune response, which is closely related to the finding of Chowdhury *et al.* (1981). However Kafi *et al.* (2003) reported that administration of live lentogenic vaccine followed by inactivated killed vaccine would produce durable immunity.

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