



## Microalgae Biomass Application in Commercial Broilers Nutrition and Their Efficacy Against Challenge with Epidemic Newcastle Disease Virus in Egypt

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Received: 07 May 2019

Accepted: 04 June 2019

### ABSTRACT

Using microalgal biomass in animal diets has been studied recently. Many species of cultivated algae were found effective in maintaining animal growth performance, and in improving body weight. Using of microalgae collected from high rate algal ponds (HRAP) as a feed additive to broilers ration was studied. One hundred and twenty broiler chicks were divided into 6 groups of 20 birds, three of them have fed on balanced broiler ration supplied with 1% weight per weight (W/W) of microalgae biomass and have variable vaccination schemes of live attenuated and inactivated Newcastle disease virus (NDV) vaccines genotype II or either non-vaccinated control. In addition, the other 3 groups have fed on free microalgae biomass balanced ration with the same vaccination treatment. Furthermore, weight gain, antibody response, mortalities, viral shedding and normal viability of chickens were estimated in order to assess the efficiency of microalgae as a feed additive. The results showed that the microalgae have no hazard effect on feed and water intake as well as enhanced viability of chickens. And in regards to immune function and body weight, they have similar effect with the free microalgae groups in normal serological response and viral shedding post vaccination with NDV vaccines as well as similar protection rate and body weight gain. In conclusion, microalgae can be used in broiler ration with no deleterious effect on growth rate, weight gain, poultry viability and immune response. In conclusion dried microalgal biomass harvested from HRAP can be used in broiler ration with no deleterious effect on growth rate, weight gain, poultry viability and immune response. Furthermore, future studies should be applied with increasing microalgae percent in poultry feed up to 5, 10 or 20% (W/W) in order to assess better performance on poultry production.

**Key words:** Algae biomass, Immune response, Newcastle disease virus, Poultry feed

### INTRODUCTION

Microalgae considered as a good source of a wide range of metabolites that are suitable for animal feed. These metabolites include protein, carbohydrate, fat, vitamins, minerals, and other organic compounds (Abdelnour et al., 2019; Andrade et al., 2018). Microalgae are identified as microscopic, unicellular and photosynthetic organisms and can grow in saline and fresh water which are rich source of nutrients and biologically active compounds, including proteins, amino acids, polyunsaturated fatty acids (LCPUFA n-3), microelements, vitamins, antioxidants, as well as carotenoids which have a long history of application as a food for human (Belay et al., 1996). Microalgae of different species can be successfully

included into poultry diets, for example as a defatted biomass byproduct from biofuel production, which can have a beneficial influence on birds' health, performance, and the quality of meat and egg (Abdelnour et al., 2019). Especially important for the poultry industry are recent studies where microalgal biomass was efficiently used in the production of eggs containing health-promoting lipids, i.e. eggs enriched with health promoting long-chain n-3 polyunsaturated fatty acids (LCPUFAs n-3) (Wu et al., 2012). Microalgae has many advantages including high growth rates, growth potential during the year and minimal land and water requirements compared to crop plants (Rawat et al., 2011). Furthermore, the high capital costs for large scale production of microalgal biofuel currently

precludes this when coupled with wastewater treatment, it may become viable in the future (Rawat et al., 2011; Benemann, 2008).

High Rate Algal Ponds (HRAP) is an advanced open pond system that was found to offer more effective wastewater treatment technologies, since it performs high removal rates for nutrients, in addition to lower capital and operation costs, in comparison with traditional treatment ponds (Sutherland et al., 2014; Craggs et al., 2014). HRAP systems have the added advantage for resource recovery from the wastewater, via algal biomass, using as fertilizer, feed or as a feedstock for biofuel production, with the later use receiving considerable attention in recent years (Craggs et al., 2014; Park and Craggs, 2011).

Newcastle Disease Virus (NDV) causes one of the most important infectious diseases of poultry, Newcastle Disease (ND) found worldwide and causes losses from mortality and condemnation of carcasses (OIE, 2012). There are many countries that have endemic NDV, with increasing occurring year after year. Also known as avian paramyxovirus serotype-1 virus, NDV is a member of the genus Avulavirus in the Paramyxoviridae family (Mayo, 2002). Feeding of  $\beta$ -glucan derived from micro algae resulted as insignificant increase in the antibody titer against NDV and Infectious Bronchitis Virus (IBV). It was suggested that dietary  $\beta$ -glucan and BA-pro might be useful in treating against viral diseases because of immunostimulating activity as reported by An et al. (2008).

The objective of the present study was to investigate the dietary effects of micro-algae biomass on growth performance, humoral immunity and viral shedding against challenge with velogenic NDV (VDNV) genotype VII in broiler chicks.

## MATERIALS AND METHODS

### Ethical approval

The animal experiment was conducted in strict accordance with and adherence to the relevant policies regarding animal handling as mandated under international national, and /or institutional guidelines for the care of animals and was approved by the research ethical committee at the national research centre, Cairo, Egypt.

### Algal biomass dominated by *Microcystis* sp.

Race way-type pond (HRAP) made of glass-fiber reinforced plastics material, with 6.5 m<sup>3</sup> capacity without connection or separation. Its dimensions are 7m length  $\times$  3m width  $\times$  40cm depth and the effective wastewater

depth is 0.3 m. The HRAP was constructed and installed in Zenin wastewater treatment plant- Giza Company for water and wastewater (Figure 1). Algal biomass collected from HRAP (1 m<sup>3</sup>) is harvested biweekly and precipitated by cationic starch. Algal biomass was dried (using sun drier) and grinded to fine particles (0.1 mm).



**Figure 1.** High rate algal pond system used for water treatment an algal biomass production

### Identification of algal community contained in the algal biomass

Along one year, samples were collected twice a week and subsamples were dispensed into glass Sedgewick-Rafter cells and examined using OLYMPUS CX41<sup>®</sup> microscope (20 $\times$ power). Species composition and dominance in the samples were determined semi-quantitatively. Algal identification has been done according to the main references used in phytoplankton identification (Streble and Krauter, 2006).

### Toxicity study for coagulants used in the precipitation of algae

#### Animals

Thirty Swiss mice with 25g average body weight were housed in standard cages (six rats), under specific pathogen-free conditions in facilities maintained at controlled room temperature (21-24°C) with a 40-60% relative humidity and under normal environmental dark-light cycles (from November to March, 7-9 hours of light and from April to October 9-12 hours of Light in Cairo, Egypt). All animals had free access to rat chow diet and water *ad libitum* and were acclimated for two weeks prior to initiation of the experiment in the laboratory of in the national research centre.

**Acute toxicity study.** Selected thirty mice of uniform weight are taken and divided into five groups each of six. The coagulants (cationic starches 1, 2, 3 and 4) and aluminum sulfate were dissolved in distilled water then given orally to four groups of mice in graded doses (1gm, 2gm, 3gm 4gm) up to 5 g/kg (El Naggar et al., 2018). In

addition, the Non-Polar Fraction of dried Algal biomass (NPFA) was suspended in distilled water then given orally to rats in graded doses up to 5 g/kg. The control group received the same volumes of distilled water. The percentage mortality for extracts was recorded 24 hours later (Desoukey et al., 2016). Observation of rats for 14 days, for any changes in the skin and fur, respiratory, circulatory, autonomic, central nervous systems, somatomotor activity and behavior pattern. Particular observation for tremors, convulsions, salivation, diarrhea, lethargy, sleep, and coma were done (El Nagggar et al., 2018).

**Table 1.** Composition of the experimental diets applied in broiler feed

Item	Starter (1 to 21 day)	Grower (22 to 40 day)
<b>Ingredient (%)</b>		
Corn	59.2	64.65
Soybean meal	30.0	24.0
Corn gluten meal	2.5	3.0
Soybean oil	2.8	3.0
Limestone	1.2	1.2
Dicalcium phosphate	1.7	1.6
Vitamin-mineral premix*	1.0	1.0
Salt	0.3	0.3
l-Lysine	0.15	0.15
dl-Methionine	0.15	0.1
Algae	1.0	1.0
<b>Analyzed chemical composition</b>		
ME (kcal/kg)	3,122	3,188
CP (%)	20.72	18.93
Ether extract (%)	4.14	5.79
DM (%)	89.1	88.4
Lysine (%)	1.17	1.0
Methionine (%)	0.5	0.41
Methionine + cysteine (%)	0.65	0.6
Threonine (%)	0.7	0.6
Tryptophan (%)	0.17	0.16
Calcium (%)	0.9	0.84
Phosphorus (%)	0.6	0.5

\* Vitamin-mineral mix (per kilogram of diet): vitamin A: (10000 IU/g) 8000 IU, vitamin D3: (200000 IU/g) 1600 IU, vitamin E: (200000 IU/g) 11 mg, riboflavin: (53 mg/g) 9 mg,  $\alpha$  calcium pantothenate: (80 g/lb), vitamin B12: (60 mg/g) 13 mg, niacin: 26 mg; choline chloride: (50%) (74% choline) 900 mg, vitamin K: (Hetrazen 35.27 mg/g) 1.5 mg, folic acid: (13.23 mg/g) 1.5 mg, biotin: (2%) 0.2 mg, santonin: (25%) 125 mg, manganese oxide: (56% Mn) 55 mg, selenium premix: (200 mg/kg) 0.1 mg, zinc oxide: (80% Zn) 50 mg, copper sulfate: (25% Cu) 5 mg,

ferrous oxide: (69.94% Fe) 30 mg, ME: (kcal/kg) metabolizable energy (kilocalorie per kilogram), CP: crude protein, DM: Dry matter

#### Determination of carbohydrates content

Total carbohydrate content of algal biomass was determined where 0.1g of sample was treated with 25ml of 1 normal sulfuric acid (1N H<sub>2</sub>SO<sub>4</sub>) was added and the mixture was hydrolyzed for two hours on a boiling water bath. At the end of hydrolysis, a flocculent precipitate was noticed. This was freed of sulphate by precipitation with barium carbonate. The mixture was filtrated and completed to 100ml. one ml of the filtrate mixed with one ml 5% phenol and 5 ml concentrated H<sub>2</sub>SO<sub>4</sub> (sulfuric acid). The absorbance was measured spectrophotometrically at wave length at 485nm (Albalasmeh et al., 2013).

#### Determination of total protein content

Total protein content was determined by kjeldahl method and then multiplied with a factor of 6.25 to convert total measured organic nitrogen to total Protein to give the total protein (Stephen et al., 2013).

#### Viruses and vaccines

**Challenge virus.** Very Virulent (vv) NDV used in the challenge was kindly supplied by poultry diseases department, veterinary research division, National Research Centre, Dokki, Egypt, characterized by sequencing as vvNDV genotype VIIId designated as NDV/Chicken/EG-MN/NRC/2015 under accession number (MF418020.1) on gene bank. The virus challenge dose equal 6-Log<sub>10</sub> EID<sub>50</sub> given 0.5 ml / bird via Intramuscular route (IM) (OIE, 2012).

**Live NDV vaccine.** Freeze-dried vaccine containing live NDV LaSota strain (Jovac ND LaSota<sup>®</sup> 1000 doses, Jordan) supplied by local agency. The vaccinal dose equal 6-Log<sup>-10</sup> EID<sub>50</sub>/ bird was given via oculonasal route as recommended by manufacturer.

**Inactivated NDV genotype II vaccine.** Inactivated NDV genotype II vaccine (Boehringer Ingelheim<sup>®</sup> AI ND 1000 doses, Germany) supplied by local agency. The vaccinal dose equal 8.2-Log<sub>10</sub> EID<sub>50</sub> given 0.5ml / bird via subcutaneous route (SC) as recommended by manufacturer.

#### Serology

Blood was collected pre and post challenge from all birds at 14, 21, 28 and 35 days of age, serum was extracted then tested by Hemagglutination Inhibition (HI) assay. The HI assay was performed using LaSota NDV antigen

according to standard procedures with four haemagglutination units virus/antigen in 0.025 ml (OIE, 2012).

### Virus shedding

Virus Isolation (VI) was performed to identify virus shedding conducted on oral and cloacal swabs as All VI-positive swabs were titrated in 9–11 days old specific pathogen free embryonated chicken eggs. Virus titers were calculated reported as mean embryo infectious dose (EID<sub>50</sub>/ 0.1 ml) on a Log 10 scale (OIE, 2012).

**Table 2.** Experimental design for usages of microalgae biomass in Broilers diet treated with different vaccines of Newcastle disease against velogenic Newcastle disease virus challenge

Group no.	Birds no.	Vaccination regime		Microalgae Addition with 1% In feed	Challenge at age / day <sup>c</sup>
		Type	Age / days		
1	20	Live GII <sup>a</sup>	5 and 18	+	28
2	20	Live. GII Inact.NDV.GII <sup>b</sup>	5 and 18 7	+	28
3	20	None	None	+	28
4	20	Live GII	5 and 18	None	28
5	20	Live. GII Inact.NDV GII	5 and 18 7	None	28
6	20	None	None	None	28

<sup>a</sup> Live NDV vaccine genotype II (LaSota strain) . The vaccinal dose equal 6-log<sub>-10</sub> EID<sub>50</sub> / bird given via oculonasal route, <sup>b</sup> Inactivated oil emulsion NDV vaccine genotype II (LaSota strain). The vaccinal dose equal 8.2-Log<sub>-10</sub> EID<sub>50</sub> given 0.5 ml / bird via SC route, <sup>c</sup> Challenge with velogenic Newcastle disease virus (genotype VII). The virus challenge dose equal 6-Log<sub>-10</sub> EID<sub>50</sub> given 0.5 ml/bird via IM route. no: number

## RESULTS

### Algal dynamics and predominance in the high rate algal pond

The high rate algal pond operation started on 30 June 2017 (summer months) which lead to the predominance of *Microcystis flos aquae*, *Microcystis aeruginosa* and other species of *Microcystis*. Always, different *Microcystis* sp. represent the upper most layer, while near the bottom layer of the pond different green and diatoms group are present. Many species include *Scenedesmus obliquus* and *Scenedesmus quadricauda*, *Ankistrodesmus*, *Coelastrum microporum*, *Selenastrum* and *Micractinium pusillum* (green algae group), *Oscillatoria limnetica* (blue green algae group) and *Nitzschia linearis* (diatoms group) was the most pre dominant species. Since the beginning of November 2017, the community structure of HRAP completely changed where the *Microcystis* sp. disappeared and the dominance of different algal species took place. In addition, all algal species mixed and floated in pond

### Experimental design

One hundred and twenty broiler chicks (Cobb 500®) one day-old supplied from commercial hatchery, were divided into six groups of 20 birds in separate units with strict biosecurity level. Conventional animal welfare regulations and food standards were taken into account (Table 2).

water column. The predominant algal species are *Scenedesmus obliquus*, *Scenedesmus quadricauda*, *Ankistrodesmus*, *Coelastrum microporum*, *Selenastrum*, *Oocystis parva*, *Dictyosphaerium pulchellum*, *Coelastrum reticulatum*, *Pediastrum gracillimum*, *Siderocells elegans*, *Eudorina elegans*, *Clamydomonas reinhardi* and *Micractinium pusillum* (green algae group), *Euglena sp.* (Euglenophyta), *Oscillatoria limnetica* (blue-green algae group) and *Nitzschia linearis* (diatoms group) by microscopical examination.

Along one year, algal biomass (from inside the algal pond) were harvested biweekly and precipitated using cationic starch and then dried. The dried algal biomass grinded to fine particles (0.1mm) and mixed thoroughly.

### Acute toxicity study

The results obtained no mortality after 24h of oral administration of NPFA at increasing doses up to a 5 g/kg. Also, after 15 days of single oral administration of cationic starch as well as NPFA, the results revealed that no

obvious changes were detected in skin and fur, respiratory, circulatory, autonomic, central nervous systems, as well as somatomotor activity and behavioral pattern were apparently normal.

Proteins and carbohydrates content were detected in order to evaluate the possibility of using the biomass for feeding properties. Total protein content detected was ranging from 232.1 to 371.3 mg/gm. Referring to carbohydrates content, total carbohydrates content was ranging from 100-138 mg/gm. Microalgae was reported to have high percentage of proteins, lipids and carbohydrates and could be used in the manufacture of different products. Microalgae can be grown in open ponds or

closed photobioreactors (Kit *et al.*, 2017; Koopmans, 2013).

### Serological response and hemagglutination inhibition assay

Similar immunological response was detected between treated and non-treated algae groups. HI geometric means titers were ranged from 2.81 to 5.43 versus 2.83 to 5.23 HI titers in G1 and G4, respectively. While, ranged from 3.93 to 6.64 versus 3.86 to 6.61 HI titers in G2 and G5, respectively. Whereas, non-vaccinated infected controls G3 and G6 were, 1.73 to 1.91 HI titers against 1.36 to 1.86 HI titers, respectively before and after challenge (Table 3).

**Table 3.** Serological response of different vaccines against velogenic Newcastle disease virus challenge after algal biomass addition in broilers diet

Group no.	Birds no.	Vaccination regime		Microalgae Addition with 1% In feed	Challenge at age/day <sup>c</sup>	HI <sup>d</sup> titre means Log-2 at age/days (N = 5)			
		Type	age/days			14	21	28	35
1	20	Live GII <sup>a</sup>	5 and 18	+	28	2.81	3.56	4.42	5.43
2	20	Live GII Inact.NDV GII <sup>b</sup>	5 and 18 7	+	28	3.93	4.86	5.21	6.64
3	20	None	None	+	28	1.73	1.61	1.91	NT
4	20	Live GII	5 and 18	None	28	2.83	3.61	4.33	5.23
5	20	Live GII Inact.NDV GII	5 & 18 7	None	28	3.86	4.52	5.36	6.61
6	20	None	None	None	28	1.36	1.65	1.86	NT

<sup>a</sup>Live NDV vaccine LaSota strain. The vaccinal dose equal 6-log<sub>10</sub> EID<sub>50</sub>/bird given via oculonasal route. <sup>b</sup>Inactivated oil emulsion NDV vaccine genotype II. The vaccinal dose equal 8.2-Log<sub>10</sub> EID<sub>50</sub> given 0.5 ml/bird via SC route. <sup>c</sup>Challenge with velogenic Newcastle disease virus (genotype VII) and The virus challenge dose equal 6-Log<sub>10</sub> EID<sub>50</sub> given 0.5 ml/bird via IM route. <sup>d</sup>HI titre ≤ 2 Log- 2 considered negative (OIE, 2012). N: Number of tested samples. no: Number, NT: None tested.

### Protection from clinical disease and post-mortem gross lesions

The objective of this study was to estimate the effect of microalgae in poultry feed on viability, weight gain and enhancement of immune response and lesser shedding of viral infection that already circulated in field. Accordingly, chickens in all groups were monitored for one-week Post-Challenge (PCH). Vaccinated-challenged groups (G1, G2, G4 and G5 either with or without algae in feed, respectively) indicated similar degree of protection against clinical disease in the form of mild respiratory signs, dull appearance and decreased in feed and water intakes for three days of PCH. While non-vaccinated-challenged groups (G3 and G6 with or without algae in feed, respectively) showed more severe signs of foamy conjunctivitis, swollen eye lids, respiratory sounds, slight depression and marked decrease in feed and water intake. Obvious greenish diarrhea, marked depression with sleepy appearance at three days PCH. Severe respiratory sounds,

conjunctivitis, nasal discharge four days PCH, and paresis ended by nervous signs (torticollis) seven days PCH.

On the other side, all dead birds were necropsied for Post-Mortem (pm) gross lesions assays revealed that, vaccinated-challenged birds showed mottled and enlarged spleens with either hemorrhagic spots and/or petechial hemorrhages on the proventricular glands (Figure 2). While non-vaccinated infected controls had severe pm gross lesions including severe congestion in the trachea, necrosis and congestion of liver, mottled and enlarged spleen, either hemorrhagic spots and/or petechial hemorrhages on the proventricular glands and enteritis with greenish intestinal contents (Figure 2).

### Body weight value of microalgae

Since the nutritional value of microalgae varies considerably with the species used, 1% treated algae ration was not enough to add value in compare with non-treated groups (Table 4). The G1 average Body Weight (BW) in

compared with G4 was 1.43 kg versus 1.46 kg, respectively. Whereas, G2 in compared with G5 was 1.41 versus 1.42 kg, respectively. Furthermore, non-vaccinated controls G3 1.3 versus G6 1.25 kg, respectively all at 28 days of age. In addition to, viability of chickens and rate of feed and water consumption was nearly equal and in normal if comparing algae and non-algae treated groups.

**Protection against mortalities**

One of the major purposes of this study was to determine the role of micro algae treated ration in enhancement of better viability and fair protection against mortalities when challenge birds with vvNDV post vaccination regimes. Table 5 indicates 95% of non-vaccinated infected controls were dead at one week PCH in both G3 and G6. While, mortalities in G1 compared with G4 was 60% versus 55%, respectively. Furthermore, was 25% in both G2 and G5 one week PCH.

**Virus shedding**

Viral shedding is one of the most relevant parameter in judgment of vaccine efficacy in correlation with the differences in immunological response obtained from microalgae supplement. Accordingly, No obvious differences were detected in bio-algae feeded groups in compared with non-feeded ones. Viral titers (Log 10) expressed as mean embryo infectious doses per 0.1 ml in G1 against G4 were 4.1 and 5.7 versus 4.2 and 5.6, respectively at three and seven days PCH. While, in G2 and G5 were 2.9 and 4.1 versus 2.8 and 4.2, respectively at three and seven days PCH. In addition, viral titer was 6.5 in both non-vaccinated challenged controls (G3 and G6) three days PCH (Table 6 and Figure 4).

**Table 4.** Weight gain after algal biomass addition in broilers diet treated with different vaccines against velogenic Newcastle disease virus challenge

Group no.	Birds no.	Vaccination regime		Microalgae Addition with 1% In feed	Challenge at age / day <sup>c</sup>	Average BW in kilograms at 28 days of age
		Type	Age / days			
1	20	Live GII <sup>a</sup>	5 and 18	+	28	1.43
2	20	Live GII Inactivated NDV GII <sup>b</sup>	5 and 18 7	+	28	1.41
3	20	None	None	+	28	1.3
4	20	Live GII	5 and 18	None	28	1.46
5	20	Live GII Inactivated .NDV GII	5 and 18 7	None	28	1.42
6	20	None	None	None	28	1.25

<sup>a</sup> Live NDV vaccine LaSota strain. The vaccinal dose equal 6-log<sub>-10</sub> EID<sub>50</sub> /bird given via oculonasal route. <sup>b</sup> Inactivated oil emulsion NDV vaccine genotype II. The vaccinal dose equal 8.2-Log<sub>-10</sub> EID<sub>50</sub> given 0.5 ml / bird via SC route. <sup>c</sup> Challenge with velogenic Newcastle disease virus (genotype VII), The virus challenge dose equal 6-Log<sub>-10</sub> EID<sub>50</sub> given 0.5 ml / bird via intramuscular route. BW: body weight; no: Number.

**Table 5.** Mortality percent after algal biomass addition in broilers diet treated with different vaccines against velogenic Newcastle disease virus challenge

Group no.	Birds no.	Vaccination regime		Microalgae Addition with 1% In feed	Challenge at age/day <sup>c</sup>	Mortalities one week PCH	
		Type	age/days			no.	%
1	20	Live GII <sup>a</sup>	5 and 18	+	28	12	60
2	20	Live GII Inactivated NDV GII <sup>b</sup>	5 and 18 7	+	28	5	25
3	20	None	None	+	28	19	95
4	20	Live GII	5 and 18	None	28	11	55
5	20	Live GII Inact.NDV GII	5 & 18 7	None	28	5	25

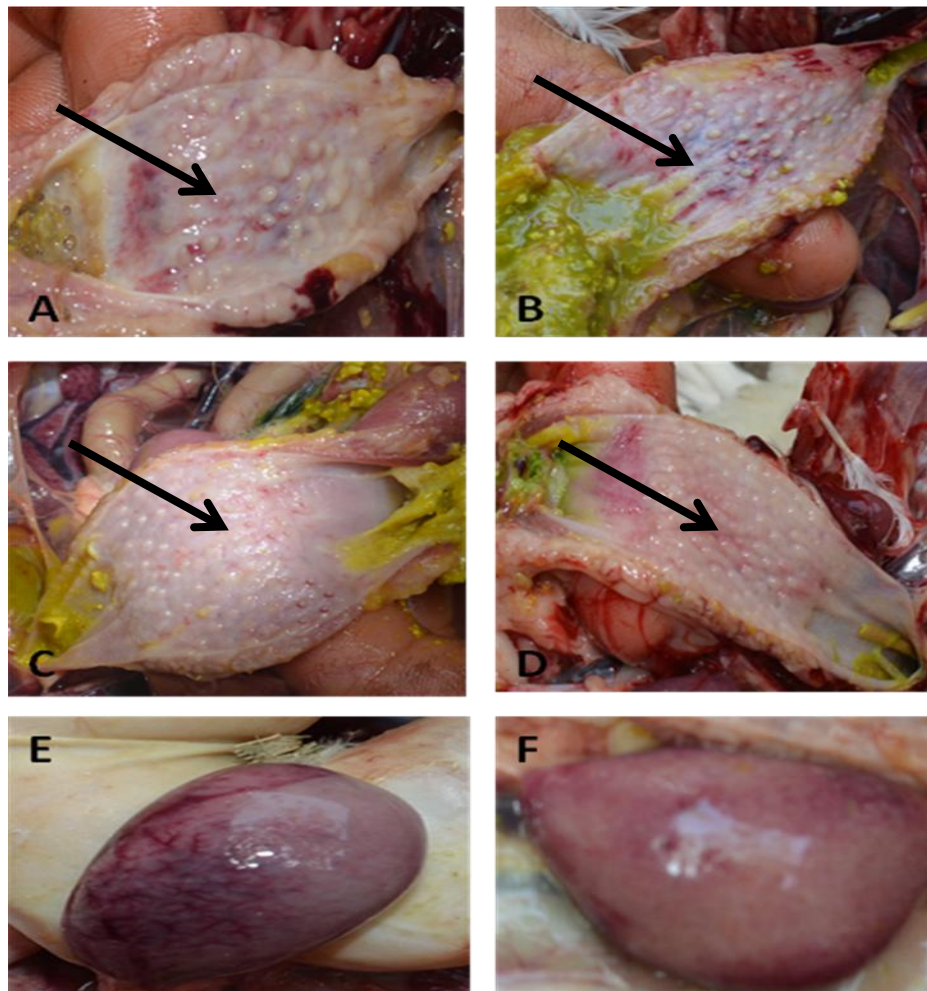
6	20	None	None	None	28	19	95
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<sup>a</sup>Live NDV vaccine LaSota strain. The vaccinal dose equal 6-log<sub>10</sub> EID<sub>50</sub>/bird given via oculonasal route. <sup>b</sup>Inactivated oil emulsion NDV vaccine genotype II. The vaccinal dose equal 8.2-Log<sub>10</sub> EID<sub>50</sub> given 0.5 ml / bird via subcutaneous route (S/C). <sup>c</sup>Challenge with velogenic Newcastle disease virus (genotype VII). The virus challenge dose equal 6-Log<sub>10</sub> EID<sub>50</sub> given 0.5 ml / bird via intramuscular route. PCH: post challenge, no: Number.

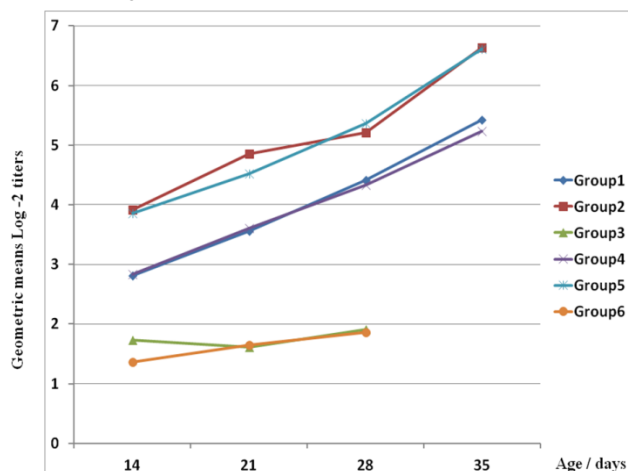
**Table 6.** Viral shedding after algal biomass addition in broilers diet treated with different vaccines against velogenic Newcastle disease virus challenge

Group no.	Birds no.	Vaccination regime		Microalgae Addition with 1% In feed	Challenge at age/day <sup>c</sup>	<sup>d</sup> Shedding at days post-challenge	
		Type	age/days			3 days (OP swabs)	7 days (CL swabs)
1	20	Live GII <sup>a</sup>	5 and 18	+	28	4.1	5.7
2	20	Live GII Inactivated NDV GII <sup>b</sup>	5 and 18 7	+	28	2.9	4.1
3	20	None	None	+	28	6.5	NT
4	20	Live GII	5 and 18	None	28	4.2	5.6
5	20	Live GII Inact.NDV GII	5 & 18 7	None	28	4.8	4.2
6	20	None	None	None	28	6.5	NT

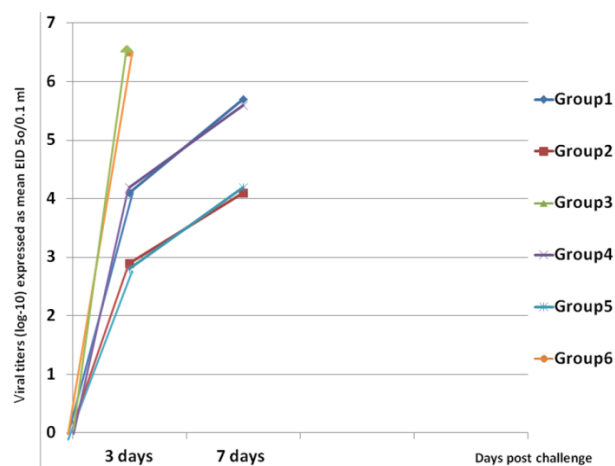
<sup>a</sup> Live NDV vaccine LaSota strain. The vaccinal dose equal 6-log<sub>10</sub> EID<sub>50</sub> / bird given via oculonasal route. <sup>b</sup> Inactivated oil emulsion NDV vaccine genotype II. The vaccinal dose equal 8.2-Log<sub>10</sub> EID<sub>50</sub> given 0.5 ml / bird via subcutaneous route, <sup>c</sup> Challenge with velogenic Newcastle disease virus (genotype VII). The virus challenge dose equal 6-Log<sub>10</sub> EID<sub>50</sub> given 0.5 ml/bird via intramuscular route, <sup>d</sup> Viral titers (log10) expressed as mean embryo infectious doses per 0.1 ml from oral and cloacal swabs taken at day 3 and 7 post-challenge from the challenged birds. OP: oropharyngeal, CL: cloacal, NT: None tested, no: number of birds in each group.



**Figure 2.** The post mortem lesions in broiler chickens treated with different vaccines of Newcastle disease against velogenic Newcastle disease virus challenge. **A** and **B**: Petechial hemorrhage on proventricular glands of infected controls Groups 3 and 6, respectively at 4 days after challenge. **C** and **D**: petechial hemorrhage on proventricular glands of infected vaccinates Groups 2 and 5, respectively at 5 days after challenge. **E** and **F**: mottled and enlarged spleen of infected vaccinates groups 1 and 4, respectively at 5 days after challenge.



**Figure 3.** Serological response evaluation after microalgae biomass application in commercial broilers feed and received different Newcastle disease virus vaccination schemes against velogenic Newcastle disease virus challenge at 14, 21, 28 and 35 days post challenge with haemagglutination titers of the same rang of both algae-treated groups or no-algae feeded ones either challenged or non-infected controls.



**Figure 4.** Viral shedding after velogenic Newcastle disease virus challenge at 3 and 7 days from oropharyngeal and cloacal swabs for evaluation of microalgae biomass application effect in commercial broilers feed treated with different Newcastle disease virus vaccination schemes. Showing viral titers of the same rang of both algae-treated groups or no-algae feeded ones either challenged or non-infected controls.

## DISCUSSION

Using microalgal biomass in feeding animals has recently applied in poultry as Microalgae improve skin color, shanks and egg yolks. Numerous nutritional and toxicological evaluations revealed the appropriateness of algal biomass as a valuable feed supplement or used instead of conventional protein sources (soybean meal, fish meal, rice bran, etc.) (Becker, 2007). In addition to, using algae as a protein source for livestock, many were used for health benefits (i.e. improved immune response, improved fertility, better weight control, healthier skin and a lustrous coat) (Pulz and Gross, 2004).

This study aims to explore the effects of dietary micro algae supplementation in broiler chicks as enhancer of BW and immune status in broilers in a challenged setting. No considerable differences were found in BW among the treatment groups with 1% bioalgae supplement (Andrade, 2018; Abdelnour et al., 2019). Nonetheless earlier studies have reported changeable results debating the effects of micro algae on performance in poultry. Zhang et al. (2008) found that  $\beta$ -glucans extracted from the algae at the percentage of 50 and 75 mg/kg in the diet increased BW. In a similar work, (Rathgeber et al., 2008) observed higher BW in broilers that fed on  $\beta$ -glucan during the growth phase. On the other hand, other researchers reported low performance after B-glucan supplementation in chickens (Huff et al., 2006). In harmony with our findings, several cases reported no significant effects of micro algae on growth performance in either non-challenged reports (Cheng et al., 2004; Chae et al., 2006; Morales-Lopez et al., 2009) or challenged settings (Chen et al., 2006; Chen et al., 2008). The conflicting results found in those studies could be due to differences in the source of the algae or the presence and type of challenge used, or both.

The use of live and inactivated vaccines has been used in the prevention and control of NDV (Zhao et al., 2014). Vaccines induce antibody production of IgA, IgG, and IgM to high levels of vaccinated chicks (Russell and Ezeifka, 1995). Because micro algae can play a role in stimulating these immune responses, the use of algae products in poultry vaccinated with NDV has been studied. In a study by An et al. (2008), algal B-glucan was fed to NDV vaccinated broilers at levels of 0, 0.025, and 0.1% for 35 days. The inclusion of algal B -glucan



significantly increased ( $P < 0.05$ ) at day 35 BW of NDV vaccinated broilers than control. Additionally, NDV specific antibody titers were elevated through the supplementation of algal  $\beta$ -glucan at 0.1%. The cell mediated immune response was suggested by (Cheng *et al.*, 2004) to be improved by algal  $\beta$ -glucan through modulating macrophage activity. These observations are supported by (An *et al.*, 2008) who also observed an increase in day 35 Newcastle virus antibody titers when feeding algal  $\beta$ -glucan at 0.05 and 0.1%, suggesting dosage level of  $\beta$ -glucan impacted the increase in antibody titers. These data confirm that an algal supplement can improve early performance parameters in a non-challenge setting, increase Newcastle virus specific antibody titers, and eliminate growth performance reductions in *Eimeria* challenged broilers. These data also confirm that dosage level of algae plays a significant role in performance effects in both non-challenged and challenged settings. These three experiments demonstrate the effectiveness of micro algae in poultry production although additional research must be conducted to further confirm proper dosage level. The results of Morales-Lopez *et al.* (2009) supporting our results, revealing that no benefits to performance or NDV specific antibody titers of a purified algae when fed to NDV vaccinated broilers.

In recent work and in agreement with our study Evans *et al.* (2015) showed that the incorporation of 16% of dried algae into a broiler diet has no negative effects on the performance of chicks. In addition, (Ross and Dominy, 1990) showed similar results in his work where found that no significant differences in performance of broilers fed a diet containing 1.5, 3, 6 or 12% dehydrated algae in feed. As well as, (Raach-Moujahed *et al.*, 2011) also reported that fed with or without 4 or 8% of algal biomass in the diet has no difference in growth performance of broilers. In contrast, (Shanmugapriya *et al.*, 2015) recently observed improvement in body weight gain and food conversion rate in broilers when fed a diet with algal biomass. Furthermore, Mariey *et al.* (2012) reported that including algal biomass in low concentration (0.02 or 0.03%) not only improved performance in broilers, but also increased dressing percentage, meat color score, blood morphology was improved and relative abdominal fat weight was decreased.

A lot of projects in many countries are going on since the algal biomass production has concerned, currently there are lot of projects going on in many countries. We concluded that, the present study has established the safety and acceptability of micro algae biomass as poultry feed up to 1% replacement and may be

incorporated into practical broiler diets that utilized by broiler chicks without causing detriment to performance, immunity or either viability and body weight gain. So as to, further research is required to identify the optimal dosage of microalgal biomass supplementation for consistent favorable results in poultry.

## DECLARATIONS

### Acknowledgments

The authors sincerely thank the academy of scientific research and technology, ministry of higher education, Egypt, for the kind support and providing infrastructure facilities to carry out this work. This work was developed within the scope of the financed project entitled “novel approach to maximize the use of stabilization pond in Egypt: A model for water and energy food nexuses” ID: 1343.

### Author’s contribution

Sayed, Sameh, Hagar, Rehab, Abeer and Mohamed Kutkat participated in design, experimental procedure, writing, revised, and reviewing the manuscript.

### Competing interests

The authors have declared that no competing interest exists

### Consent to publish

The authors guarantee that this work has not been published elsewhere and that any person named as a coauthor of this work is aware of the fact and has agreed to be so named.

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