

## Performance and Macrophage Activity of Broilers Fed with a Sorghum Meal with Different Yeast Wall Levels

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### ABSTRACT

*The aim of this work was to evaluate the effect of broiler breeders' age and yeast wall (YW) levels on broilers' performance and macrophage activity. The experiment consisted in a completely randomized design and a 2 X 5 factorial arrangement and two controls of two broiler breeders age (34 and 57 weeks of age) and five YW levels (zero, one, two, three or four kg of YW/ton of diet). They received sorghum diet compared to a control corn/soybean meal diet. The age of the broiler breeders influenced the performance, but did not affect macrophage activity. At 34 weeks age broiler breeders' progeny, all diets were similar considering the average weight. For 57 weeks age broiler breeders' progeny, at 21 days, only average weight of sorghum diet supplemented with 4 kg/ton was similar to corn diet results. The optimum level of YW for maximal macrophage activity was 2.06 kg/ton of meal.*

**Key words:** carcass yield, mananoligossacarydes, phagocytosis

### INTRODUCTION

The poultry production is a dynamic activity that involves many areas, which depend on the technological innovations for results in field improvement, such as weight gain, feed conversion, or egg production. It is a complex chain that transforms the one-day old chicken, weighing about 42-45 g to broiler chickens weighing 2.1-2.5 kg at 42 days. This fast production of meat depends on genetic, environment, nutrition, and health status. These factors are interdependent among which nutrition and health status are basic factors to attain the productivity goals. Several studies have been conducted for better understanding of the

physiological necessities of the birds to obtain performance indices and the physiology of the immune system, including under adverse sanitary conditions (Qureshi et al., 1994; Ferket et al., 2002; Soares, et al., 2004; Pucciarelli and Benassi, 2005; Leblanc, et al., 2006).

Studies have been made on developing the products and/or substances that act as modulators and/or stimulators of the immune system for the birds, for examples, the manna oligosaccharides and the  $\beta$ -glucan, derived from the yeast cellular wall (Leblanc, et al., 2006). The macrophages, acting in the innate immunity, through the destruction of pathogens agents and adaptative antigens presentation (Qureshi, 2003; Abbas et al., 2007) are an important defense mechanism of the

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birds. Related studies on the function of these cells add important knowledge on the immune response of the birds (Qureshi et al., 1994).

The objective of this study was to evaluate the effect of increasing levels of yeast wall on the growth performance and the fagocytic activity of macrophages of the broilers fed with sorghum diets instead of corn.

## MATERIAL AND METHODS

The birds were placed in a brick house with concrete floor, a lateral wall of 40 cm and wire mesh roof. It has a mobile curtain system, a temperature control system with fans and sprinklers, and an oven heating system. There were 48 boxes of 3.90 X 1.50 meters, located in the central region of the house (aviary), with capacity of 70 birds each. The feeders were the tubular type and the water drinkers were the pendulum type. In this experiment, 3,360 male chickens Cobb breed

were used, half of which were from 34 weeks old broiler breeders headquarters and the other half from half 57 weeks old broiler breeders.

Two basal, isoproteic, isoaminoacidic, and isoenetic meal diets were formulated as described by Rostagno et al. (2005). The first one was a traditional diet based on corn and soybean, and the second diet was formulated with 100% of sorghum. Both the diets were divided in the initial phase (from 0 to 21 days old) (Table 1) and growth phase (from 22 to 42 days old) (Table 2). The experimental diets had been produced by the unit Rações do Abatedouro Coroaves LTDA and the feed ingredients used were submitted to the quality control adopted from the company, involving the classification of grains, crude protein analyses, soluble protein, ether extract, crude fiber, ashes (ANFAR, 1998), tannin analysis (Nutron Alimentos LTDA), and aflatoxin levels dosage through commercial kit (RIDASOFT®) with a 450 nm filter density optic reader (ELX-800).

**Table 1** - Centesimal and calculated composition of experimental diets for broilers from 1 to 21 days.

Ingredients %	Ration					
	Corn	Sorghum and Yeast wall (YW - %)				
		0 YW	0.1 YW	0.2 YW	0.3 YW	0.4 YW
Corn	53.07	-	-	-	-	-
Sorghum	-	53.93	53.93	53.93	53.93	53.93
Soybean meal	28.63	32.43	32.43	32.43	32.43	32.43
Full-Fat soybean	13.80	9.17	9.17	9.17	9.17	9.17
Aderex	0.25	0.30	0.20	0.20	0.20	0.20
Caulin	0.40	0.40	0.30	0.20	0.10	-
Limestone	0.97	0.97	0.97	0.97	0.97	0.97
Dicalcium phosphate	1.63	1.60	1.60	1.60	1.60	1.60
Salt	0.49	0.50	0.50	0.50	0.50	0.50
L-lysine HCl 30%	0.38	0.40	0.40	0.40	0.40	0.40
Methionine 99%	0.23	0.25	0.25	0.25	0.25	0.25
Mineral supplement	0.05	0.05	0.05	0.05	0.05	0.05
Vitamin supplement	0.10	0.10	0.10	0.10	0.10	0.10
Yeast wall	-	-	0.10	0.20	0.30	0.40
Total	100	100	100	100	100	100
Calculated composition						
Crude protein %	23.05	23.26	23.26	23.26	23.26	23.26
Metabolizable energy kcal/kg	3,050	3,050	3,050	3,050	3,050	3,050
Digestible lysine %	1.19	1.19	1.19	1.19	1.19	1.19
Digestible met + cys %	0.84	0.84	0.84	0.84	0.84	0.84
Digestible tryptophan %	0.26	0.27	0.27	0.27	0.27	0.27
Digestible threonine %	0.77	0.77	0.77	0.77	0.77	0.77

**Mineral supplement (1 kg):** Copper: 20,000 mg; Iodine: 2,000 mg; Iron: 100,000 mg; Manganese: 160,000 mg; Zinc: 100,000 mg. **Vitaminic supplement (1 kg):** vit A: 8,000,000 IU; vit D3: 2,000,000 IU; vit E: 14,500 IU; vit K3: 1,900 mg; vit B1: 1,333 mg; vit B2: 5,750 mg; vit B6: 2,380 mg; vit B12: 11 mg; Biotin: 30 mg; Folic acid: 760 mg; Nicotinic acid: 23,800 mg; Pantothenic acid: 11,400 mg; Selenium: 220 mg.

**Table 2** - Centesimal and calculated composition of experimental diets for broilers from 22 to 42 days.

Ingredients %	Ration					
	Corn	Sorghum and Yeast wall (YW - %)				
		0 YW	0.1 YW	0.2 YW	0.3 YW	0.4 YW
Corn	56.53	-	-	-	-	-
Sorghum	-	58.57	58.57	58.57	58.57	58.57
Soybean meal	19.67	27.73	27.73	27.73	27.73	27.73
Full-Fat soybean	19.40	9.33	9.33	9.33	9.33	9.33
Aderex	0.25	0.23	0.23	0.23	0.23	0.23
Caulin	0.40	0.40	0.30	0.20	0.10	-
Limestone	0.93	0.93	0.93	0.93	0.93	0.93
Dicalcium phosphate	1.50	1.47	1.47	1.47	1.47	1.47
Salt	0.47	0.48	0.48	0.48	0.48	0.48
L-lysine HCl 30%	0.48	0.47	0.47	0.47	0.47	0.47
Methionine 99%	0.22	0.24	0.24	0.24	0.24	0.24
Mineral supplement	0.05	0.05	0.05	0.05	0.05	0.05
Vitamin supplement	0.10	0.10	0.10	0.10	0.10	0.10
Yeast wall	-	-	0.10	0.20	0.30	0.40
Total	100	100	100	100	100	100
Calculated composition						
Crude protein %	21.39	21.61	21.61	21.61	21.61	21.61
Metabolizable energy kcal/kg	3,150	3,150	3,150	3,150	3,150	3,150
Digestible lysine %	1.10	1.10	1.10	1.10	1.10	1.10
Digestible met + cys %	0.79	0.79	0.79	0.79	0.79	0.79
Digestible tryptophan %	0.23	0.25	0.25	0.25	0.25	0.25
Digestible threonine %	0.71	0.71	0.71	0.71	0.71	0.71

**Mineral supplement (1 kg):** Copper: 20,000 mg; Iodine: 2,000 mg; Iron: 100,000 mg; Manganese: 160,000 mg; Zinc: 100,000 mg. **Vitaminic supplement (1 kg):** vit A: 8,000,000 IU; vit D3: 2,000,000 IU; vit E: 14,500 IU; vit K3: 1,900 mg; vit B1: 1,333 mg; vit B2: 5,750 mg; vit B6: 2,380 mg; vit B12: 11 mg; Biotin: 30 mg; Folic acid: 760 mg; Nicotinic acid: 23,800 mg; Pantothenic acid: 11,400 mg; Selenium: 220 mg.

The experimental birds had been vaccinated on the 1<sup>st</sup> day at hatchery against Marek Disease and Infectious Bursal Disease (combined), and Infectious Bronchitis in spray (MASS-I<sup>®</sup>). On the 7<sup>th</sup> and 14<sup>th</sup> day, they were vaccinated against the Infectious Bursal Disease in water (Bursine Plus<sup>®</sup>), and on the 14<sup>th</sup> day they were vaccinated against NewCastle Disease in water (Poulvac NDW<sup>®</sup> enterotropic). All vaccines were alive and attenuated.

The experiment was totally randomized with a 2x5 design, with two broiler breeders ages (34 and 57 weeks) and five yeast wall supplementation levels (zero, one, two, three or four kg/ton), and two control diets (corn and soybean meal). Each treatment had four repetitions, with 70 birds each. The average composition of the yeast wall was 30% of protein (maximum), 3.0% of crude fiber (maximum), 6.9% of ashes (maximum), aflatoxin absence and 25% of manna oligosaccharides and 30% of  $\beta$ -glucans.

For the evaluation of zootechnical parameters, the birds were weighed at the time of delivery in the

aviary and then weighed weekly at 7 am before feeding until 42<sup>th</sup> day. The diets left overs were weighed on day 7, 14, 21, 28, 35 and 42. Based on these data, average weight gain and feed conversion were calculated.

On day 42, three broilers of each repetition were weighed and sent in cages to the slaughterhouse. In the pre-slaughter period, the broilers were kept in a platform with a forced ventilation and humidification systems to minimize the stress conditions. The slaughter process was followed by hanging, stunning, automatic bleeding (the company worked with humanitarian slaughter control), picking, and manual evisceration. After the evisceration, the birds were directed to the cutting room, where they underwent to an evaluation of carcass yield and were considered the weight of the eviscerated carcass (without viscera, neck, head and feet).

Two broilers of each repetition were used for the evaluation of the phagocytic activity of macrophages, totalizing four birds for the treatment. The broilers were inoculated on the 38<sup>th</sup>

day of life, through Sephadex G-50<sup>®</sup> (Sigma) 3% intra-abdominal injection, (one mL/100g of live weight) (Qureshi et al., 1986; Gore and Qureshi, 1997; Konjufca, 2004). Intravascular catheters G-14 were used for this. Forty two hours after the inoculation, the broilers were stunted by eletronarcosis and sacrificed through cervical disconnection. Before collecting the abdominal fluid, the broilers skin was washed with neutral detergent and decontaminated with 70% alcohol. At the laboratory, each abdominal fluid sample was washed with 20 mL of sterile PBS + heparine solution (contends 0,5 UI/mL of Liquevine<sup>®</sup> 25,000 UI/5mL - Roche).

Approximately 15 mL of abdominal liquid of each broiler was collected with Pasteur pipettes and immediately conditioned in Falcon pipes in ice bath. All the samples were centrifuged at 1500 x g for 10 minutes and the pellet was resuspended in 2 mL of RPMI 1640<sup>®</sup> solution (Sigma). One hundred and fifty microliter of this suspension was added to each well of the culture plate (24 wells), with a 13 mm diameter glass slide. The culture plates were incubated at room temperature for one hour. Each plate was washed by sterile cooled PBS to remove the not adherent cells. After that, 150 µL of RPMI 1640<sup>®</sup> solution (Sigma) with 3% of sheep erythrocytes was added and the plates were incubated at room temperature in a 5% of carbonic gas atmosphere for one hour. Once again the cooled PBS solution was used to remove the not adherent sheep erythrocytes. After that, the staining was performed using a commercial kit (Panótico Fast LB<sup>®</sup> - Laborclin). After 24 h, the glass slides were embedded using Permount<sup>®</sup>. Three hundred macrophages were counted in each glass slide and also the number of the cells that contained engulfed sheep erythrocytes. The phagocytic activity was calculated considering the number of macrophages with engulfed erythrocytes divided by the total number of counted macrophages.

The average weight, feed conversion, average weight and yield of the cuts were submitted to polynomial regression analysis admitting normal distribution and identity link function (Interactive Data Analysis – SAS, 2000). The phagocytic activity was submitted to the regression analysis admitting gamma distribution and identity link

function (Interactive Dates Analysis – SAS, 2000). The Dunnett test was adopted for comparison of the control diet (corn) with the other treatments (SAS, 2000).

## RESULTS AND DISCUSSION

The age of the broiler breeder affected the weight of commercial cuts ( $P \leq 0.05$ ); however, considering the cuts yield, this effect was not observed ( $P > 0.05$ ). Regression analysis showed no influence of the levels of inclusion of yeast wall ( $P > 0.05$ ) on the average weight of the broilers and had also no improvement in the feed conversion using sorghum and yeast wall and the corn control diet (Table 3). There was no difference among the treatments for the young chickens derived from breeders of 34 weeks old for average weight at 21 and 42 days ( $P > 0.05$ ).

The feed conversion was better for the corn based diet ( $P \leq 0.05$ ) at 21 days. However, at 42 days, the feed conversion of the corn based diet showed no difference ( $P > 0.05$ ) with the treatments with sorghum and 2 or 4 kg/ton of yeast wall. Treatments of the sorghum based diet and 1 or 3 kg/ton of yeast wall showed worse feed conversion (Table 03).

For the progeny arrays of 57 weeks old, it was observed that chickens at 21 days with corn based diet had better average age ( $P \leq 0.05$ ) than those with sorghum based diet and yeast wall, except for the treatment with the supplementation of 4 kg/ton of yeast wall ( $P > 0.05$ ) (Table 3). There was no difference among the treatments for average weight at 42 days ( $P > 0.05$ ).

The feed conversion at 21 days was better for the corn based diet when compared with sorghum and yeast wall ( $P \leq 0.05$ ). However, at 42 days, the corn diet and the sorghum diet with 4 kg/ton of yeast wall showed similar results ( $P > 0,05$ ) and were better ( $P \leq 0.05$ ) than the other treatments. In the studies carried out by Rutz et al. (2006), improvements in the feed conversion had been observed when the broilers received 2% of yeast extract at the first week and between 38 and 42 days, which was attributed to the inositol and nucleotides present in the yeast extract.

**Table 3** - Means and standard errors of average weight (AW) and feed conversion (FC) at 21 and 42 days for broilers hatched from breeders with 34 and 57 weeks old fed with different levels of yeast wall (YW) supplementation in a sorghum diet compared to a controlled diet.

Diet	30 weeks broiler breeders			
	AW 21 days (kg)	AW 42 days (kg)	FC 21 days (kg)	FC 42 days (kg)
Corn	0.864 <sup>a</sup> ± 0.005	2.491 <sup>a</sup> ± 0.025	1.33 <sup>a</sup> ± 0.005	1.94 <sup>a</sup> ± 0.033
Zero YW	0.833 <sup>a</sup> ± 0.008	2.518 <sup>a</sup> ± 0.019	1.39 <sup>b</sup> ± 0.009	1.99 <sup>a</sup> ± 0.023
1 kg/ton YW	0.840 <sup>a</sup> ± 0.006	2.473 <sup>a</sup> ± 0.010	1.42 <sup>b</sup> ± 0.008	2.04 <sup>b</sup> ± 0.017
2 kg/ton YW	0.830 <sup>a</sup> ± 0.007	2.494 <sup>a</sup> ± 0.038	1.39 <sup>b</sup> ± 0.010	2.00 <sup>a</sup> ± 0.005
3 kg/ton YW	0.839 <sup>a</sup> ± 0.007	2.470 <sup>a</sup> ± 0.014	1.42 <sup>b</sup> ± 0.029	2.02 <sup>b</sup> ± 0.016
4kg/ton YW	0.827 <sup>a</sup> ± 0.015	2.486 <sup>a</sup> ± 0.013	1.41 <sup>b</sup> ± 0.011	2.00 <sup>a</sup> ± 0.011
	57 weeks broiler breeders			
Corn	0.938 <sup>a</sup> ± 0.005	2.655 <sup>a</sup> ± 0.014	1.31 <sup>a</sup> ± 0.032	1.97 <sup>a</sup> ± 0.018
Zero YW	0.883 <sup>b</sup> ± 0.010	2.536 <sup>a</sup> ± 0.018	1.38 <sup>b</sup> ± 0.008	2.05 <sup>b</sup> ± 0.027
1 kg/ton YW	0.896 <sup>b</sup> ± 0.010	2.575 <sup>a</sup> ± 0.035	1.41 <sup>b</sup> ± 0.015	2.09 <sup>b</sup> ± 0.013
2 kg/ton YW	0.893 <sup>b</sup> ± 0.012	2.581 <sup>a</sup> ± 0.006	1.39 <sup>b</sup> ± 0.013	2.05 <sup>b</sup> ± 0.011
3 kg/ton YW	0.875 <sup>b</sup> ± 0.014	2.541 <sup>a</sup> ± 0.010	1.40 <sup>b</sup> ± 0.018	2.04 <sup>b</sup> ± 0.008
4kg/ton YW	0.909 <sup>a</sup> ± 0.009	2.629 <sup>a</sup> ± 0.030	1.39 <sup>b</sup> ± 0.010	2.02 <sup>a</sup> ± 0.020

Different words at same column differ from control (Dunnett) ( $P \leq 0.05$ ).

The average weight results of this study differed from Hooge (2004) who found positive effect of the manna oligosaccharides on the weight and feed conversion of the broilers when compared with a traditional diet without antibiotics. When compared with the diet with growth-promoter antibiotics, the author observed that the diet containing manna oligosaccharides had similar results. The findings of Zhang et al. (2005) showed that broilers at 35 days old that had received diets with 0.3% of yeast wall and 0.5% of yeast had higher weight than the control diet.

In the studies carried out with turkeys, Ferket et al. (2002) compared the use of the manna oligosaccharides and antibiotics and did not observe distinct effect between the tested antibiotics and the manna oligosaccharides. The effects observed could be considered a bit conclusive. However, under the experimental conditions, where the environment is quite favorable and the challenge to the digestive tract of the broilers is minimized, the use benefits of the yeast wall can be masked (Ferket et al., 2002).

Waldroup et al. (2003) reported that the manna oligosaccharides derived from *Saccharomyces cerevisiae* were promising in the suppression of gut pathogens, modulation of the immune system, promotion of the intestine integrity and improvement of the performance (feed conversion and weight gain) of turkeys and broilers. Spring et al. (2000) also observed reduction of *Salmonella sp* in the digestive tract of birds treated with 4g of

oligosaccharides/kg. This effect is possibly related with the adsorption of the bacteria.

Fernandes et al. (2002) and Garcia (2005) had not observed disadvantages in substituting the corn by sorghum in the feed conversion, effect justified between its nutritional values, which led to assume that the variations observed in the values of feed conversion occurred randomly. Iji and Tievy (1998) demonstrated that the supplementation with manna oligosaccharides (1g/kg of feed) resulted in increased weight gain and improved the feed conversion. However, when the inclusion levels increased (5g/kg of feed), the same happened to the viscosity of the ingesta, showing no benefit in the performance of the broilers.

The increasing levels of yeast wall exhibited no effect in the values for gross weight of cuts, nor in its yield in relation with the carcass for the descending lineages of the progenies offspring of 34 and 57 weeks ( $P > 0.05$ ). Comparing the sorghum based diets, supplemented with or without yeast wall, with the corn based diet, there were no differences ( $P > 0.05$ ) in the weights of the cuts (chest, whole leg, whole wing and back – Table 04). These results were obtained even for gross weight and cut yield in relation to the carcass weight. Similar results were found by Rutz et al. (2006) that used yeast extract (inclusion of 2%). The equivalence observed in the results for gross weight and cut yield for the broilers of the same breeders line and gender were in agreement with the results of Murakami et al. (1995) who

observed differences in the yield of cuts between the strains (Cobb and Ross). In the case of gender, Stringhini et al. (2003) found greater carcass weight in the males.

The production of cuts, particularly the noble cuts, chest, thigh, and leg, is considered quite important in the poultry industry, and it is evidenced by Brazil's export of 260,630,815 kg of chicken cuts

between January and August of 2005 to the Japanese market (APA, 2006). This clearly shows the potential of the export market in absorbing the production. In this context, the possibility to use the yeast wall for the weight gain and consequently increasing the production of more meat should be studied and optimized (Ferket et al., 2002; Hooge, 2004).

**Table 4** - Means and standard errors for commercial cuts average weight (g) and carcass yield (%) for broilers hatched from breeders with 34 and 57 weeks old fed with different levels of yeast wall (YW) supplementation in a sorghum diet compared to a controlled diet

Diet	Carcass		Breast		Whole Legs		Whole Wing		Back	
	Kg	%	g	%	g	%	g	%	g	%
34 weeks broiler breeders)										
Corn	1.893	75.99	678 ± 22	35.81	581 ± 19	30.69	224 ± 07	11.83	410 ± 17	21.66
Zero YW	1.871	74.31	696 ± 22	37.20	562 ± 19	30.04	224 ± 07	11.97	389 ± 17	20.79
1 kg/ton YW	1.846	74.65	666 ± 22	36.08	547 ± 19	29.63	221 ± 07	11.97	412 ± 17	22.32
2 kg/ton YW	1.771	71.01	671 ± 22	37.89	520 ± 19	29.36	212 ± 07	11.97	368 ± 17	20.78
3 kg/ton YW	1.958	79.23	704 ± 22	35.95	577 ± 19	29.47	238 ± 07	12.15	439 ± 17	22.42
4kg/ton YW	1.825	73.41	655 ± 22	35.89	550 ± 19	30.14	218 ± 07	11.94	402 ± 17	22.03
57 weeks broiler breeders)										
Corn	1.952	73.52	714 ± 27	36.58	595 ± 20	30.48	231 ± 08	11.83	412 ± 15	21.11
Zero YW	1.995	78.67	732 ± 27	36.69	606 ± 20	30.37	237 ± 08	11.88	420 ± 15	21.05
1 kg/ton YW	1.994	77.44	730 ± 27	36.61	589 ± 20	29.54	238 ± 08	11.93	437 ± 15	21.91
2 kg/ton YW	1.871	72.10	695 ± 25	37.34	573 ± 18	30.79	222 ± 07	11.93	371 ± 14	19.93
3 kg/ton YW	1.883	74.10	695 ± 25	36.91	567 ± 18	30.11	225 ± 07	11.95	396 ± 14	21.03
4kg/ton YW	1.985	75.50	760 ± 25	38.29	587 ± 18	29.57	236 ± 07	11.89	402 ± 14	20.25

Some authors have described that the component of the yeast wall, such as the manna oligosaccharides, act by the mechanisms that involve the adsorption of pathogens in the alimentary system, in particular those that present type 1 fimbriae (Spring et al., 2000), and would help to uniform and improve the villus integrity. Lowry et al. (2005) had observed that the  $\beta$ -glucan, present in the yeast walls, reduced the invasion capacity of enteric *Salmonella* serovar *Enteritidis* in young chickens tissues (one week old), and as a consequence provided better immunity, better weight gain and, possibly, better exploitation of cuts. However, in the present study, such observations were not made.

The age of the breeders did not show any effect on the fagocytic activity of macrophages, characterized by the phagocytosis of sheep erythrocytes, stimulated through the abdominal

injection of Sephadex G-50<sup>®</sup> (Sigma) ( $P > 0.05$ ). It was considered that, under the aspect innate immunity, the breeders age did not benefit the offspring. However, factors related to the genetic lines had demonstrated differences between the macrophages functions (Qureshi and Miller, 1991).

The results demonstrated that the activity of the macrophages was influenced in a quadratic form ( $Y = 0.1297 + 0.1143 PL - 0.0278 PL^2$ ) with a maximum point for inclusion of 2.06 kg of yeast wall/ton (Table 5). In comparison (Dunnnett test) with the corn based diet (control), the treatment with sorghum and 2 kg of yeast wall/ton presented greater number of macrophages in active fagocytosis process (engulfed erythrocytes). The other treatments showed no difference from the corn based diet.

**Table 5** - Means and standard errors of macrophage activity.

Diet	Macrophages with erythrocytes/Counted macrophages
Corn	14.88 ± 5.29 <sup>b</sup>
Zero YW <sup>1</sup>	12.19 ± 8.98 <sup>b</sup>
1 kg/ton YW	23.71 ± 4.23 <sup>b</sup>
2 kg/ton YW	31.33 ± 8.58 <sup>a</sup>
3 kg/ton YW	23.09 ± 8.66 <sup>b</sup>
4kg/ton YW	17.83 ± 3.57 <sup>b</sup>

1 YW: Yeast Wall; Different words at same column differ from control (Dunnett) ( $P \leq 0.05$ ). Quadratic response in sorghum treatments  $Y = 0,1297 + 0,1143 YW - 0,0278 YW^2$

The macrophages are cells of the monocuclear fagocytic system (McCorkle, 1998), integrant of the innate immunity of the broilers, and also important for the adaptative immunity (Qureshi, 2003; Abbas et al., 2007), acting in the organism defense processes by the destruction of antigens and secreting cytokines that act in the inflammatory process. The main functions executed for this cell are fagocytosis, destruction of bacteria (Qureshi et al., 1986), secretion of prostaglandins and cytokines, and antigen presentation for development of the immune response (Abbas et al., 2007). The  $\beta$ -glucans, present in the composition of the yeast wall, have revealed importance in the stimulation of macrophages in fish by the action of specific receptors presents in these cells (Bartelme, 2006). In general, the purified  $\beta$ -glucans demonstrate good activity in stimulating the cellular defense system, especially the neutrophils in the humans (LeBlanc et al., 2006) and fishes (Palic et al., 2006), and macrophages in fishes (Falcon, 2007) and mammals (Adachi, 2004). These composites bind to unspecific immune system cells (macrophages) through portion receptors of the  $\beta$ -1-3 or  $\beta$ -1-6. As they are considered the first line of defense (Qureshi, 1998), the activation and fagocytic activity of the macrophages, if adequately stimulated, tends to supply better conditions for the broilers to engulf and to destroy the pathogens.

In the current situation of the poultry industry and looking towards the near future, the market scenraion requires reduction in the use of the antibiotics. This requires search for immune modulators that do not leave residues in the meat, promote good growth performance and works in adverse conditions so that the broilers can defend themselves against the pathogens (Dietert and Golemboski, 1998).

## CONCLUSIONS

In conclusion, the breeders age influenced the growth performance; however it did not influence the macrophages activity. The sorghum could be used without any losses to the weight of the commercial cuts. However, the 57 weeks old progeny showed reduced performance on sorghum based diets. The best level of yeast wall for attaining the maximum macrophage activity of was 2.06 kg/ton of food.

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