

## Effect of dietary Lacto cel-con probiotic on growth performance and hematology indices of fingerlings mono-sex Nile tilapia (*Oreochromis niloticus*)

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### ABSTRACT

A 120 day's feeding study was designed to study the effect of Lacto cel-con probiotic on the growth performance, hematology indices and body composition of Nile tilapia. *Zea mays* was substituted by sorghum in five formulated diets with inclusion levels of (0,25,50,75 and 100%). Fingerlings of mono-sex Nile tilapia with an initial weight of (1.94±0.13g) were distributed in 1m<sup>3</sup> hapa fixed in earthen pond at stocking rate of 40 fish/replicate. The highest growth performance parameters in terms of (weight gain and specific growth rate) and nutrient utilization (feed conversion ratio and net protein utilization) were recorded by replacing 75 and 100% of *Zea mays*. On the other hand, blood indices include, Red blood Cell (RBCs) count, Lymphocyte, Hemoglobin concentration and Hematocrit value) displayed non-significant different among treatments. However, the values of White Blood Cell count (WBCs) were shown remarkable significance (P<0.05) with DS75 and DS100% diets. No significance differences were obtained in carcass composition of tilapia fed in different sorghum levels. Results showed that sorghum meal supplemented with (0.3% Lacto-cel con) can be entirely substituted up to 75% of corn meal in fingerlings of mono-sex Nile tilapia diets without negative effects on growth performance, nutrient utilization, some blood parameters and body composition. It's conclude that (DS75) is a potential protein-rich with low cost feeding diet for mono-sex Nile tilapia (*Oreochromis niloticus*).

### INTRODUCTION

Tilapia is commonly cultured specie in the world, where its high source of animal proteins, essential amino acid, vitamins and minerals. Also fish can comfortably culture, highly adapt to environmental changes and is a vital species in developed countries (FAO,2010). Global tilapia production is expected to rise 3-4 % in 2018, reaching around 6.3 million tones (FAO, 2018). About 30% from this production from China, although the highly production is intensified in little producing countries (FAO, 2018).

Sorghum (*Sorghum bicolor* L.) is the fifth most important cereal crop in the world commerce and grown in over 100 countries with different uses as animal feed, food, in brewery, and recently as a potential source of biofuel (Ali and Warren,1987

and Araújo *et al.*,2017). Global sorghum production in 2017 is projected to be 60.46 million metric tons, and the U.S.A has been the highest producer with 9.42 million metric tons, then Nigeria, 6.55 million metric tons, and Ethiopia, 3.77 million metric tons (Araújo *et al.*,2017). United States of America was the top producer in 2009, with a harvest of 9.7 million metric tons. The next main sorghum producers with low quantities were India, Nigeria, Sudan and Ethiopia, respectively. Otherwise, the major regions for sorghum producing in the world from harvested quantities were Australia, Brazil, China, Burkina Faso, Argentina, Mali, Cameroon, Egypt, Niger, United Republic of Tanzania, Chad, Mozambique, Venezuela and Ghana. The world sorghum harvested was 55.6 million tons in 2010 (FAO, 2010).

Sorghum is similar in chemical composition to corn (*Zea mays*) and has a nutritional value similar to other cereals (Aderolu *et al.*,2009). However, including anti-nutritional factors like tannins, phytates and cyanogenic glucosides among others could probably have effect on nutrient utilization and growth of fish. Processing of sorghum by different treatments such as soaking and wet-milling can removes high percent of these anti-nutrients.

The optimum inclusion level of carbohydrate led to improving growth parameters in fish and decreasing nitrogen waste with least cost diet (Mohanta *et al.*, 2007). The use of carbohydrates is important in the fish diet so, protein or lipids are not catabolized for energy. Economically, it is more reasonable that protein can be utilized for muscle tissue synthesis not for produce metabolic energy (Oriro *et al.*, 2014). Sparing protein by non-protein energy sources has been evident in a high range of species (Hemre, *et al.*, 2002 ,Castro, *et al.*,2016 and Wu *et al.*, 2016).

Until now, probiotics, point out as specific stimulant and a potent method, have highly received essential interest due to its benefits for limitation use of antibiotics, antimicrobial products and protection against infectious diseases as well as improving growth performance in aquaculture (Sahu *et al.*,2008a and Gao *et al.*, 2013).

Recently more studies illustrated that diet supplementation with probiotic may have many beneficial effects on health status, immunostimulation and disease control (Panigrahi *et al.*,2009 and Oliva-Teles,2012). In the same manner, extensive studied on probiotics revealed a positive enhancement on growth and feed utilization in Nile tilapia *Oreochromis niloticus* (Lara-Flores *et al.*,2010). These benefits possibly as a result of that probiotic had capacity to stimulate and/or produce digestive enzymes, feed digestion and, in consequence, feed utilization efficiency. Indeed, some probiotics, like *Bacillus* species, produce a wide range of exoenzymes (amylase, protease, lipase and cellulase) that are very efficient in degradation of carbohydrates, proteins and lipids (Ray *et al.*,2012).

Alternatively, probiotics are now used in aquaculture as simple and safe additive to improve the health of the host. Probiotics have many advantages in aquaculture, such as modulating microbial colonization, enhancing growth, providing nutrients, improving immune responses, increasing digestive enzyme activities, improving feed utilization and digestibility, controlling diseases and improving water quality (Qi *et al.*, 2009 and Merrifield *et al.*, 2010a).

The present study was designed to detect the effect of different inclusion levels of sorghum supplemented with Lacto cel-con probiotic on the growth performance, feed efficiency, body proximate analysis and some blood parameters in fingerlings of mono-sex Nile tilapia (*Oreochromis niloticus*).

## MATERIALS AND METHODS

### Fish culture system

Fingerlings of Nile tilapia (*Oreochromis niloticus*) were obtained from private tilapia hatchery at Fayoum Governorate, Egypt. They transferred to Shakhshouk Research Station at Fayoum Governorate by using fish transport car held with pure oxygen source. The trial was conducted in earthen ponds of the station. The system consisted of 15 rectangular hapa designed as small ponds from small size nets (maga 60) and fixed with wood column from all sides. The volume of each unit was 1m<sup>3</sup> as (1m×2m×0.5m). The earthen pond filled with agriculture drain water through cement channel. The running drain water were connected to the filled pond and changed as (100% twice/week). Physicochemical characteristics of water earthen ponds were examined every two weeks according to (APHA, 2005). During the experimental study water quality values measured within the optimal ranges for this specie, where dissolved oxygen ( $7.2 \pm 0.2$  mg dL<sup>-1</sup>), temperature ( $28.42 \pm 1.2^\circ\text{C}$ ), salinity ( $4.14 \pm 0.5$ ) and pH ( $7.5 \pm 0.2$ ) as recorded by (Bergheim, 2007).

This experiment was performed for a period of 120 days (from April 1<sup>st</sup>, 2014 till July 30<sup>th</sup>, 2014). Two weeks of acclimatization period of fish on commercial diet containing (30% crude protein) before starting the experimental. Fish were randomly distributed as density of 40 fingerlings in each pond with initial mean weight of ( $1.94 \pm 0.13$ g) in three replicates for each treatment.

### Experimental diets

Sorghum meal was obtained from local farm at Shakhshouk-Fayoum, while the other ingredients purchased from El-Asil feed company-Dakahlea Governorate, Egypt. Sorghum was soaked with tap water along three days and then sun-dried before crushing. Ingredients were grinding into fine powder through a 150- $\mu\text{m}$  mesh before pelleting through California pelleted machine with 2mm diameter. Five tested diets were prepared as: DC100 (which considered to be the control without sorghum meal (100% corn), then sorghum meal was partially and totally replaced corn (*Zea mays*) as follows: DS25 (25% sorghum meal replacement), DS50 (50% sorghum meal replacement), DS75 (75% sorghum meal replacement) and DS100 (100% sorghum meal replacement). Lacto cel-con was incorporated as 0.3% with the control and the four inclusion levels. The formulation and proximate composition of the practical diets are shown in (Table,1).

### Probiotics

In the present experimental we used a commercial Lacto cel-con probiotic produced in China and their contents was as follows:

Lacto cel-con is mixture of *Saccharomyces cerevisia* in purified mature live cells with the most beneficial intestinal micro flora-*Lactobacillus acidophilus* and *Enterococcus faecium* and is a dried natural product each 1 Kg contains:

- *Saccharomyces cerevisia* ( $2 \times 10^{12}$  CFU)
- *Lactobacillus acidophilus* ( $100 \times 10^9$  CUF)
- *Enterococcus faecium* ( $70 \times 10^9$  CUF)
- Carrier with mixture of ground yellow corn and gluten meal with a ratio of 3:1.

Guaranteed analysis:

Crude protein, minimum	18.00%
Crude fat, minimum	3.00%
Crude fiber, maximum	5.00%
<i>Saccharomyces cerevisiae</i> , minimum	1 Trillion CFU's per lb.
<i>Lactobacillus acidophilus</i> , minimum	50 Billion CFU's per lb.

*Enterococcus faecium*, minimum

35 Billion CFU's per lb.

Table 1. Formulation and proximate composition of experimental diets (%DM basis).

Items Ingredients	Experimental diets%				
	DC100	DS25	DS50	DS75	DS100
Yellow-Corn	33.0	24.75	16.5	8.25	zero
Sorghum	Zero	8.25	16.5	24.75	33.0
Soya bean meal	20.0	20.0	20.0	20.0	20.0
Wheat bran	25.6	25.6	25.8	26.3	26.6
Glutein	10.0	10.0	10.0	10.0	10.0
Fish meal	5.7	5.7	5.5	5.0	4.7
Molasse	1.0	1.0	1.0	1.0	1.0
Lactocel-con g/kg	0.3	0.3	0.3	0.3	0.3
Fish Oil	3.0	3.0	3.0	3.0	3.0
Vit,Min Mix. <sup>1</sup>	1.2	1.2	1.2	1.2	1.2
DL.Methionine	0.2	0.2	0.2	0.2	0.2
Total	100	100	100	100	100
Proximate composition of experimental diets (%DM basis)					
Dry matter (DM)	91.17	90.90	90.65	90.77	91.01
Crude protein (CP)	25.08	25.08	25.05	25.00	25.00
Ether extract (EE)	10.80	9.50	10.60	9.90	9.40
Crude fiber (CF)	6.27	6.59	6.55	6.89	6.56
NFE <sup>2</sup>	40.29	40.97	40.76	41.50	42.27
Ash	17.56	17.71	16.79	16.36	16.37
Tannin <sup>3</sup>	-	0.15	0.25	0.35	0.40
GE(Kcal/kg) <sup>4</sup>	4089	3995	4088	4050	4034
C/P %ratio	66.91	67.50	66.80	67.53	66.94

1-Vitamin-mineral premix supplied the following (vitamin IU/kg<sup>-1</sup> diet and mineral mg/ Kg<sup>-1</sup> mixture); retinyl acetate 0.67; ascorbic acid 120; cholecolciferol 0.1; tocopheryl acetate 34.2; menodione 22; thiamin 5.6; riboflavin 12; pyridoxine 4.5; calcium panthothenate 14.1; p-aminobenzoic acid 40; cyanocobalamin 0.03; niacin 30; biotin 0.1; choline chloride 350; folic acid 1.5; inositol 50; canthaxanthin 10; butylated hydroxytoluene 1.5; butylated hydroxyanisol 1.5.; CaHPO<sub>4</sub>·2H<sub>2</sub>O 29.5; Ca (H<sub>2</sub>PO<sub>4</sub>)<sub>2</sub> H<sub>2</sub>O 217; NaHCO<sub>3</sub> 94.5; Na<sub>2</sub>SeO<sub>3</sub>·5H<sub>2</sub>O 0.011; Kci 100; Nacl 172.4; Ki 0.2; Mgcl<sub>2</sub> 63.7; MgSO<sub>4</sub> 34.3; MnSO<sub>4</sub> 2; FeSO<sub>4</sub>·H<sub>2</sub>O 10; CuSO<sub>4</sub>·5H<sub>2</sub>O 0.4; Zn SO<sub>4</sub> 10.

2-Calculated by differences, Nitrogen free extract (NFE) = [100-(CP+EE+CF+Ash)].

3-Tannin = percent tannin on a catechin equivalent basis.

4- Estimated according to (NRC, 2011) by using these values, 5.64, 9.44 and 4.11 Kcal/g for protein, lipid and NFE, respectively.

### Chemical analysis

Before the experiment, 30 fish from the initial fish were randomly chosen to determine initial body composition. After the end of the experimental, fish were starved for 24hrs prior to sample collection. Fish in each hapa weighed, counted and determine growth performance, feed efficiency and survival rate. Finally, 3 fish samples from each replicate were collected to determine the final carcass proximate composition. Analyses of diets and carcass composition were done by (AOAC, 2006); dry matter determined by drying samples in an oven at 105 °C until constant weight; crude protein was measuring nitrogen by (N×6.25) after acid digestion (Kjeldahl method); crude lipid was determined through petroleum ether extraction using the (Soxhlet method); ash was detected by incineration in a furnace muffle at 550 °C for 16h, while nitrogen free extract (NFE) was calculated by difference. Gross energy in feed samples was calculated in chemical composition and estimated according to (NRC, 2011) by using these values: 5.64, 9.44 and 4.11 Kcal/g for protein, lipid and NFE, respectively.

Tannin content of sorghum meal determined using a modified version of (Price *et al.*, 1978).

### Growth parameters

Growth indices and feed utilization were measured by using the following equations:

Body weight gain BWG = (FW–IW).

FW=Final weight (g).IW=Initial weight(g).

Specific growth rate SGR (%/day) = (Ln FW-Ln IW)/T×100.

Ln = Natural logarithm T = period (days).

Condition factor (K)=W/L<sup>3</sup>×100.

Where: W=fish weight (g). L=fish standard length (cm).

Feed conversion ratio (FCR)=Feed intake (g)/Weight gain (g).

PER (%)=Weight gain (g)/Protein intake (g).

Net Protein utilization (NPU%) =(Final body protein-Initial body protein/protein intake)×100.

Hepatosomatic index (HSI %)=(liver weight)/(fish weight)×100.

### Blood analysis

Three fish from each replicate were collected from the experimental hapa, and anesthetized by MS222 overdose and immediately bled. The blood was collected by using Vacutainer tubes (Becton-Dickinson, Rutherford, NJ, USA) provide with lithium heparin as anticoagulant to collect blood from the caudal artery and vein, then mixing and prepared to analysis. Standard haematological parameters were measured by (Blaxhall and Daisley,1973), as followed: Haemoglobin (Hb) with the cyanomethaemoglobin method, packed Cell Volume (PCV) by micro Haematocrit method, White Blood Cell (WBCs) with the improve Neubauer Counter, differential counts and lymphocytes) were done on blood film stained with may Grumwald-Giensa stain. Red blood cell (RBCs) was determined using the relationship between Hb and PCV (Miale,1982).

### Statistical analysis

The results were presented as means ±SD for three replicates. All data were exhibited one-way analysis (ANOVA) to test the effect of different sorghum levels and probiotic on various performance, feed utilization, and blood values of experimental fish according to (Snedecore and Cochran,1987). Tukey multiple range tests were applied to detect the significant differences between the means of treatments (Zar,1999). All analysis were performed using SPSS version 20,(2016) SPSS Institute, Cary, NC, USA).

## RESULTS

### Chemical composition of diets

Our results show that the tested diets were similar in their protein contents ranging from (30.04 to 30.27%) and growth energy contents ranging from (19.73 to19.96 MJ kg<sup>-1</sup> diets) as presented in (Table, 1).

### Growth performance

The growth performance values revealed that mean of initial weights of the experimental ranged between (1.75 ±0.1-2.09 ±0.11) for weight and (1.50 ±0.11-2.60±0.12) in length as shown in (Table, 2, Figs.1&2). The growth values show no significant differences among groups point to the complete homogeneity among the experimental fish at the onset of the experiment.

Table 2: Growth performance and feed utilization of tilapia fed on different experimental diets (Mean±SD n=9).

Parameters	Experimental diets%				
	DC100	DS25	DS50	DS75	DS100
Init. fish weight (g)	1.90 <sup>a</sup> ±0.11	2.09 <sup>a</sup> ±0.11	1.90 <sup>a</sup> ±0.12	1.75 <sup>a</sup> ±0.11	2.01 <sup>a</sup> ±0.12
Final fish weight (g)	56.00 <sup>d</sup> ±4.19	60.25 <sup>c</sup> ±5.05	61.85 <sup>c</sup> ±4.35	69.20 <sup>a</sup> ±5.20	66.95 <sup>b</sup> ±5.25
Init. fish length(cm)	2.00 <sup>a</sup> ±0.11	2.6 <sup>a</sup> ±0.12	2.50 <sup>a</sup> ±0.12	1.50 <sup>a</sup> ±0.10	2.50 <sup>a</sup> ±0.13
Final fish length (cm)	15.70 <sup>a</sup> ±0.32	16.00 <sup>a</sup> ±0.34	16.10 <sup>a</sup> ±0.32	16.50 <sup>a</sup> ±0.34	16.56 <sup>a</sup> ±0.32
Total weight gain (g)	54.10 <sup>d</sup> ±4.75	58.16 <sup>d</sup> ±4.66	59.95 <sup>c</sup> ±4.55	67.45 <sup>a</sup> ±4.26	64.94 <sup>b</sup> ±4.54
SGR (%/ day)	1.22 <sup>b</sup> ±0.02	1.21 <sup>b</sup> ±0.02	1.26 <sup>ab</sup> ±0.04	1.33 <sup>a</sup> ±0.04	1.27 <sup>ab</sup> ±0.05
Condition factor(K)	1.44 <sup>b</sup> ±0.02	1.47 <sup>ab</sup> ±0.02	1.48 <sup>ab</sup> ±0.04	1.54 <sup>a</sup> ±0.01	1.47 <sup>ab</sup> ±0.03
Feed intake (g/fish)	115.65	122.14	123.46	124.37	122.53
Feed conversion ratio	2.14 <sup>b</sup> ±0.11	2.10 <sup>b</sup> ±0.12	2.06 <sup>b</sup> ±0.04	1.84 <sup>a</sup> ±0.13	1.89 <sup>a</sup> ±0.12
Protein efficiency ratio	1.86 <sup>b</sup> ±0.11	1.89 <sup>b</sup> ±0.12	1.93 <sup>b</sup> ±0.12	2.17 <sup>a</sup> ±0.11	2.12 <sup>a</sup> ±0.13
Net protein utilization (%)	29.24 <sup>b</sup> ±1.2	29.96 <sup>b</sup> ±1.4	28.41 <sup>b</sup> ±1.2	29.31 <sup>b</sup> ±1.6	30.78 <sup>a</sup> ±1.4
Hepatosomatic index (HSI %)	2.1 <sup>a</sup> ±0.02	2.2 <sup>a</sup> ±0.04	2.3 <sup>a</sup> ±0.02	2.2 <sup>a</sup> ±0.04	2.4 <sup>a</sup> ±0.05

Means in the same row with different super script letters are significantly different (P<0.05).

However, the final weight gain, specific growth rate and condition factor showed significance difference (P<0.05) between treatments. Furthermore, the highest values in previous parameters were obtained with the fish fed up to 75% sorghum meal (DS75%) compared with corn diet (control) and other experimental diets. In the same manner, feed utilization from feed intake, feed conversion ratio, protein efficiency ratio and net protein utilization were significant difference between treatments (P<0.05). The elevated parameters of feed conversion ratio and other values of feed utilization were observed with fish fed (DS75%) as illustrated in (Fig.,3). However, the hepatosomatic index (HSI) show no significant differences between dietary treatments.

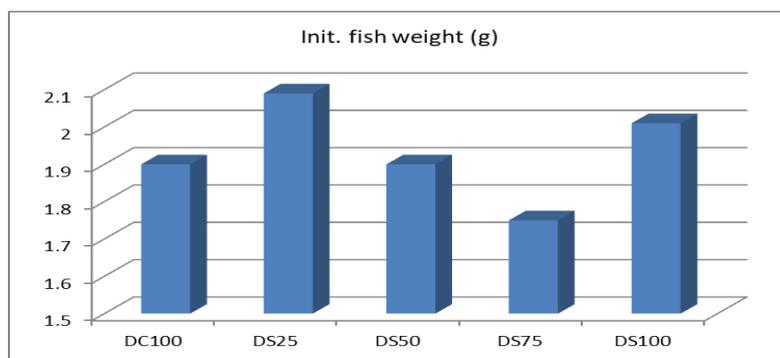


Fig. 1: Initial fish weight of experimental fish.

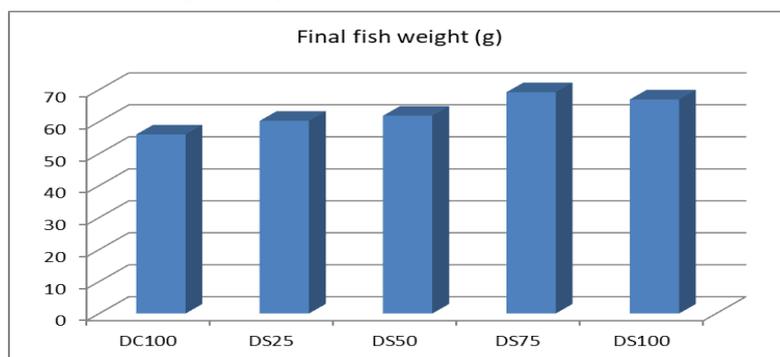


Fig. 2: Final average weight of fish fed experimental diets.

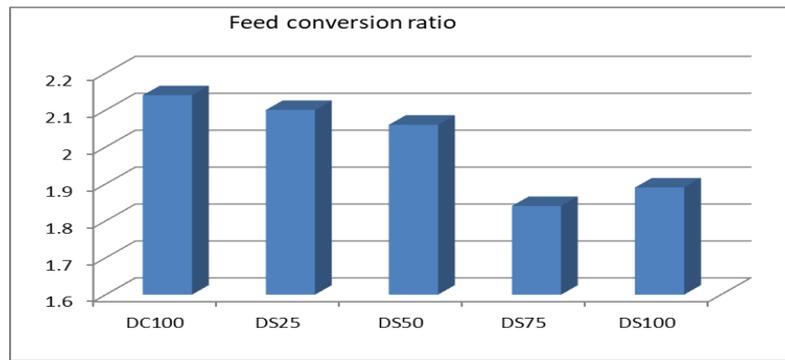


Fig. 3: Feed conversion ratio of experimental diets.

### Blood parameters

As presented in (Table, 3), there was a significance increased ( $p < 0.05$ ) in white blood cells (WBCs) with increasing sorghum meal (DS75 and DS100) and the lowest value was obtained with 25% sorghum meal diet. However, RBCs, Lymphocyte, Hemoglobin and Hematocrit ratios were not showed significant differences between experimental diets.

Table 3: Blood parameters of tilapia fed on different experimental diets (Mean±SD n=9).

Parameters	Experimental diets%				
	DC100	DS25	DS50	DS75	DS100
WBCs (10 <sup>6</sup> /mm)	74.20 <sup>b</sup> ±0.88	73.40 <sup>b</sup> ±0.82	76.34 <sup>a</sup> ±0.74	78.72 <sup>a</sup> ±0.71	78.12 <sup>a</sup> ±0.76
RBCs (10 <sup>3</sup> /ml <sup>3</sup> )	2.56 <sup>a</sup> ±0.06	2.42 <sup>a</sup> ±0.05	2.49 <sup>a</sup> ±0.04	2.67 <sup>a</sup> ±0.05	2.46 <sup>a</sup> ±0.04
Lymphocyte	91.84 <sup>a</sup> ±1.40	91.20 <sup>a</sup> ±1.42	92.22 <sup>a</sup> ±1.35	93.46 <sup>a</sup> ±1.34	91.43 <sup>a</sup> ±1.34
Hemoglobin (g/dL)	9.33 <sup>a</sup> ±0.42	9.26 <sup>a</sup> ±0.52	9.12 <sup>a</sup> ±0.45	9.20 <sup>a</sup> ±0.35	9.32 <sup>a</sup> ±0.45
Hematocrit (%)	34.25 <sup>a</sup> ±0.67	34.66 <sup>a</sup> ±0.57	34.34 <sup>a</sup> ±0.23	34.57 <sup>a</sup> ±0.43	34.44 <sup>a</sup> ±0.43

Means in the same raw with different super script letters are significantly different ( $P < 0.05$ ).

### Carcass composition

The whole body composition of fish was presented in (Table 4). As shown no significance difference ( $P > 0.05$ ) was detected in dry matter, crude protein ether extract and ash between treatments.

Table 4: Whole body analysis (Mean±SD n=9) of tilapia fed on the experimental diets (% w/w basis).

Chemical analysis	Experimental diets%				
	DC100	DS25	DS50	DS75	DS100
Dry matter	27.33±0.40	27.35±0.20	27.56±0.25	28.02±0.15	27.82±0.25
Crude protein (CP)	15.49±0.03	15.68±0.05	15.46±0.10	15.45±0.10	15.32±0.10
Ether extract (EE)	5.32±0.20	5.51±0.30	5.97±0.25	6.47±0.10	6.46±0.20
Ash	6.52±0.16	6.16±0.10	6.13±0.07	6.10±0.18	6.04±0.17

## DISCUSSION

The effects of probiotics have been extensively investigated since its onset use in aquaculture, and their beneficial on growth and immunity for aquatic animals have been confirmed.

The utilization of probiotics in aquaculture has increased during recent years. Newly, a specific type, heat-killed or inactivated probiotics has been applied as a promoter or an immuno-stimulant/immuno-biotic to promote growth performance, improve feed utilization, enhance none-specific immune responses, resistance to stress and protection against bacterial pathogens of aquatic animals (Dawood *et al.*, 2016b).

In the current study, the dietary supply of Lacto cel-con significantly enhanced the growth and feed efficiency of tilapia in terms of (weight gain, specific growth rate, condition factor, feed conversion ratio, protein efficiency ratio and net protein utilization) compared to those fed without Lacto cel-con diets. Similar results were detected in tilapia *Oreochromis niloticus* diet supplemented with Lacto cel-con by (El Zayat, 2014 and Hussein *et al.*, 2016) and other type of probiotic (Khattab *et al.*, 2004, EL-Haroun *et al.*, 2006, Eid and Mohamed, 2008, Olmos *et al.*, 2011, Chiu and Liu, 2014, Ridha and Azad, 2016, Lopez *et al.*, 2016, Dawood *et al.*, 2016b, Dawood *et al.*, 2016c, Dawood *et al.*, 2017 and Nguyen *et al.*, 2019).

The above observations were generally demonstrated that the growth promotion of fish was likely attributed to the positive effects of inactivated and/or dead (killed) probiotic bacteria. Inactivated compounds enhanced the concentration of extracellular enzymes secreted by the gut microflora and improved the digestibility (Dawood *et al.*, 2015c). Otherwise, enhanced the immune responses, resulting an improvement in health status (Rodriguez-Estrada *et al.*, 2013 and Zhou *et al.*, 2010). Other researchers suggested that the use of these compounds, induce the secretion of the intestinal brush border enzymes by intestinal epithelial cells together with extracellular enzymes synthesized by intestinal microbes (Kesarodi-Watson *et al.*, 2008). However, the probiotic roles in synthesis and utilization nutrients in fish have not been completely elucidated (Abumourad *et al.*, 2013 and Dawood *et al.*, 2016a), and the metabolism of Lacto cel-con related to elevated growth and feed efficacy in tilapia, which takes place in the gastro intestinal tract, is possibly complicated and need further studies.

In the present trial it's clear that the sorghum meal can be used up to 33% to replace 75% of corn *Zea mays* without negative effect on growth parameters and feed efficiency of fingerlings tilapia, concept the lowering of corn from 100% to 25 % by substituting with sorghum meal. The high inclusion level of sorghum in the different diets was not affected by its palatability. This might be attributed to the soaking and heating during pelting of sorghum seeds prior to use in tilapia diets. This finding is in agree with (Fagbenro, 1999, Francis *et al.*, 2001, Siddhuraja *et al.*, 2003 and Obe, 2014), they reported that the reduction in anti-nutrient by various processing techniques resulted to better palatability and growth in fish. The present results are in agree with the previous results in tilapia, which revealed the ability of this specie to utilize up to 40% of dietary starch sources (maize or sorghum) as recorded by (Yones, 2005, Yones and Metwalli, 2016, Abd-El-Azem, 2016 and Hussein *et al.*, 2016). However, Wilson *et al.*, (2003) reported a high inclusion up to (44% of diet) from low tannin sorghum 0.4% tannin in feeding of Nile tilapia. In the present study, the normal relative liver weight (HSI) revealed that the fish was well being utilized this source of carbohydrate to provide energy, probably as a consequence of the good utilization of CHO coupled with the using of Lacto cel-con in feeding strategy.

Data showed that the blood parameters of fingerlings tilapia from (Hematocrit, Hemoglobin, Red blood cell (RBCs), and Lymphocyte) were not affected with sorghum inclusion levels and similar with the recorded values in tilapia (Akinrotimi *et al.*, 2012 and Hussein *et al.*, 2016). In contrast significance differences and high levels of WBCs were obtained by using Lacto cel-con in different groups as a result of immunity effect. The data obtained are in line with the findings of using probiotics as immuno-stimulants to enhance the non-specific immune system of the host and protected against stress (Hussein *et al.*, 2016).

The whole chemical composition of fish was not affected by probiotic and inclusion levels of sorghum and these values were in accordance with the previous

results (Anderson *et al.*, 1984, Al-Asgah and Alli, 1994, Al-Ogaily and Alli, 1996, Solomon *et al.*, 2007, Yones and Metwalli, 2016 and Hussein *et al.*, 2016).

## CONCLUSION

The use of sorghum meal to replace up to 75% of corn in diet supplemented with 0.3% Lacto cel-con probiotic didn't shown any adverse effects on growth performance, feed utilization, blood parameters and chemical composition of mono-sex Nile tilapia fingerlings. It's also clear the benefit effect of probiotic in the previous parameters. The present results confirm that sorghum meal can be substituted instead of corn meal in diet formulation as alternative and cheap carbohydrate source in tilapia feeding. Otherwise, more studies on sorghum digestibility will be effective to detect the adequate inclusion level in tilapia diet. In the same manner, further research is still needed to detect the role of probiotics (mode of action) on digestibility, immune response and stress resistance. It is also important to define the optimum levels of probiotic, which can be used in tilapia diets to avoid any adverse effects on fish culture.

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## ARABIC SUMMARY

تأثير تغذية الاكتو-سيلكون بروبيوتك على معدلات الأداء ومؤشرات الدم لإصبعيات أسماك البلطي النيلي

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إجريت هذه التجربة لفترة ١٢٠ يوم لتقييم تأثير استخدام الاكتو-سيلكون بروبيوتك على النمو، مؤشرات الدم وتركيب الجسم لإصبعيات البلطي النيلي. تم احلال الذرة الصفراء بالذرة الرفيعة في خمس علائق تجريبية حيث كانت نسب الاحلال (٠، ٢٥، ٥٠، ٧٥ و ١٠٠%). وزعت إصبعيات البلطي وحيد الجنس بمتوسط وزن أولى (١.٩ ± ٠.١٣، جم) في هابة ١ متر مكعب مثبتة داخل حوض أرضى وبمعدل تخزينى ٤٠ سمكة / متر مكعب/ مكررة. أظهرت قياسات النمو متمثلة في (عائد النمو ومعدل النمو النوعى) وكفاءة استخدام الغذاء مثل (معدل التحويل الغذائى و كفاءة استخدام البروتين) نتائج أعلى عند احلال الذرة الصفراء بمستوى ٧٥ و ١٠٠% بالذرة الرفيعة. وعلى الجانب الآخر لم تظهر قياسات الدم مثل محتوى (كرات الدم الحمراء، نسبة الهيموجلوبين و الهيماتوكريت) إختلافات معنوية بين المعاملات المختلفة بينما ارتفعت نسبة كرات الدم البيضاء واختلفت معنويا لكل من المعاملة التى أدخلت الذرة الرفيعة فى العليقة عند نسبتى ٧٥ و ١٠٠. لم يظهر تركيب جسم الأسماك إختلافات معنوية مع مستويات الإحلال المختلفة من الذرة الرفيعة. أظهرت نتائج الدراسة إمكانية إحلال الذرة الرفيعة محل ٧٥% من الذرة الصفراء دون تأثيرات معاكسة على معدل الأداء، بعض صفات الدم وتركيب جسم الأسماك كما أن العليقة المحتوية على ٧٥% ذرة رفيعة كانت اقتصادية فى تغذية إصبعيات البلطي النيلي وحيد الجنس.