

## Adaptability of the Nile Tilapia, *Oreochromis niloticus* Juveniles to Water Salinity by Controlling Dietary Sodium Chloride Levels

Abdelrhman M. Abdelrhman<sup>1</sup>, Zaki Z. Sharawy<sup>1</sup>, Ashraf M. A. S. Goda<sup>1</sup>,  
Matthew J. Slater<sup>2</sup>

<sup>1</sup> Aquaculture Division, National of Oceanography & Fisheries (NIOF), Cairo, Egypt

<sup>2</sup> Alfred Wegener Institute for Polar and Marine Research, Bremerhaven, Germany

\*Corresponding Author: [abdelrhman\\_niof@hotmail.com](mailto:abdelrhman_niof@hotmail.com)

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### ABSTRACT

Increasing groundwater scarcity and salinity pose challenges for high water demand aquaculture. Adapting animals to grow optimally under saline conditions is key to future growth in many important aquaculture regions. In the current study, 10-week controlled feeding experiment was carried out to investigate dietary sodium chloride supplementation (10% NaCl) as a method of mitigating the effects of increasing water salinity (0,10 and 15 ppt) on growth and metabolism of juvenile Nile tilapia, *Oreochromis niloticus* in controlled feeding experiments. Fish fed diets supplemented with 10% NaCl recorded significantly better weight gain when compared to fish fed control when held in 10 and 15 ppt salinity water over a period of ten weeks. Blood glucose and blood lactate concentrations were both significantly higher in fish fed diets supplemented with 10% NaCl and increased with increasing water salinity. Fish fed diet containing 10% NaCl recorded the highest values of Na<sup>+</sup> concentration in blood compared to other experimental groups. Feed utilization, as indicated by feed conversion ratio decreased significantly with water salinity ( $P < 0.05$ ). Feed intake also decreased with increasing water salinity. The obtained results indicated that salinity is a key factor in controlling growth of Nile tilapia. Results clearly indicate that negative effects of increased salinity in rearing water can be partially mitigated by salt inclusion in diets.

### INTRODUCTION

Egypt is located in the subtropical belt and a major part of the country faces scarcity of rainfall, and thus significant water scarcity overall. Thus, it is extremely important to find possible alternatives for justified utilization of limited water resources. Requirements for water for agriculture and other urban activities has increased the pressure to develop aquaculture in brackish and seawater. Control of salt and water balance within a narrow limit is critical to life in all multicellular organisms, including teleost fishes. Salt tolerance is a term describing the overall fitness, or productivity, of the fish in a saline

environment. It is a combination of different quantitative traits, such as metabolism, growth, osmoregulation, immunocompetence and fecundity (**Cnaani and Hulata, 2011**).

Nile Tilapia, *O. niloticus* belongs to family *Cichlidae*, which dominates freshwater fish culture and is an excellent candidate for aquaculture in brackish water due to their ability to tolerate a wide range of water salinity. Inter-specific variation in salinity tolerance may be used to select salt-tolerant species and develop salt-tolerant hybrids. Growth of *O. niloticus* at high salinity is significantly lower than that in freshwater **Fineman, (1988)**, whereas survival is not affected by salinity. High salinity does seem to suppress, or at least delay, onset of reproduction in *O. niloticus*, thus presenting a practical method of population control to improve overall yields.

Recent studies by **Appelbaum *et al.* (2008 a, 2008b)** showed that gilthead sea bream juveniles reared in brackish water (3‰ TDS) and fed a diet supplemented with 1.5% salt grew significantly ( $P < 0.05$ ) better than those fed the control diet (no added salt). A further study (**Appelbaum and Arockiaraj, 2008 b**) using diets supplemented with higher levels of salt (8, 10 and 12%) showed that gilthead sea bream juveniles reared in brackish water of 2.9‰ (TDS) salinity grew better and had the highest survival rate when fed a diet containing 12% salt. The salinity of the culture system is known to influence metabolism and homeostatic processes in fish, meaning the organism needs for nutrients and protein particularly, may differ in freshwater and saltwater systems (**Altinok and Grizzle, 2001**).

**Yao *et al.* (2008)** investigated the best conditions for transfer of Nile tilapia from freshwater to salt water. Nile tilapia fingerlings (8 to 12 g) were transferred, either directly or gradually, from freshwater to water of variable salinities, and survival was monitored after 3 weeks. Survival of fish transferred directly to saline water was high (84.3% to 96.8%) until 17 ppt, but mortalities were significant (60-70%) above that salinity. High rate of survival (78 to 81%) was, however, achieved by gradual acclimation to salinity of 30 ppt over two days. **Larumbe-Moran *et al.* (2010)** reported that culture of Nile tilapia is feasible in saline environments at up to 25 ppt of salinity without affecting growth or survival range, and with no increases required in dietary protein if the standing biomass remains under critical maxima. Nonetheless, the tilapia do require higher protein intake at higher salinities to produce growth rates equivalent to those observed in freshwater environments. The present study was undertaken to determine the effects of water salinity and dietary supplementation of sodium chloride on growth performance, feed utilization, survival rate, and blood components of juvenile Nile tilapia.

## MATERIALS AND METHODS

### Experimental Fish and Culture Technique

The present study was conducted at the Alfred-Wegener-Institute Helmholtz Center for Polar and Marine Research, Center for Aquaculture Research (ZAF), Bremerhaven, Germany for 10 weeks (70 days, August to October 2018). Two hundred and seventy Nile tilapia, *O. niloticus* juveniles were purchased from Til-Aqua International (The Netherlands; for details on the production see [www.til-aqua.com](http://www.til-aqua.com)) with an average initial body weight of  $8.0 \pm 0.01$ g. They were then randomly stocked in 18 glass aquaria (80 X 30 X 40 cm; length X width X height) each aquaria was stocked with 15 fish. Three different water salinity levels (0, 10 and 15 ppt) and two different dietary sodium chloride NaCl supplementation levels (0 and 10%) in a factorial manner ( $3 \times 2$ ) was conducted to represented six different experimental treatments, Three replicates aquaria were randomly assigned to each treatment. Prior to the start of experiment, the fish were acclimated to the experimental conditions for two weeks.

### Experimental Diets

Two experimental diets were formulated to be isonitrogenous, 35% crude protein (CP) and isocaloric, 10.26 MJ/kg digestible energy (DE). All diets were identical except for the variation in NaCl levels. The basal experimental diet had no NaCl added. (Table 1) , Diets 2 contained sodium chloride, NaCl at levels 10%.

The diets were processed by blending the dry ingredients into a homogenous mixture. Pellets of 2 mm were made in ZAF laboratory pellet. The pelleting temperature did not exceed 40 °C and all diets were air dried for 4 hr., (moisture content of about 10%). All diets was packed in cellophane bags and cooled at 4°C prior to use.

Fish were fed the experimental diets at the rate of 3% of body weight per day and it offered two feedings at 8.00 and 13.00 hours. The fish in each aquarium were weighed every 2 weeks, and the feed weight was adjusted after each fish weighing.

Aeration of the water was continuously provided using compressed air. Salinity, temperature, dissolved oxygen, pH and ammonia were continuously monitored during the experiment as a normal routine for RAS to maintain water quality at optimum range for Nile tilapia.

**Table 1. Formulation and proximate composition of the experimental basal diet (% as-fed basis)**

<b>Ingredients</b>	<b>Percent in basal experimental diet</b>
<b>Fish meal</b>	30
<b>Soybean meal</b>	20
<b>Yellow corn</b>	10
<b>Wheat bran</b>	33
<b>Sunflower oil</b>	5
<b>Vitamin and Mineral premix<sup>a</sup></b>	2
<b>Proximate composition (%)</b>	
<b>Dry matter</b>	91.89
<b>Crude protein</b>	35.10
<b>Crude fat</b>	7.61
<b>Total carbohydrate</b>	47.36
<b>Ash</b>	9.93
<b>Gross energy<sup>b</sup> (MJ/kg)</b>	10.26

<sup>a</sup> Vitamin and mineral mixture (supplements per kg of the mixed feed): Vit. A 72000IU, Vit. B1 6 mg, Vit. B3 12000 IU, Vit. B6 9 mg, B12 0.06 mg, Vit E 60 mg, Vit. 12 mg, Pantothonic acid 60 mg, Nicotinic acid 120 mg, Folic acid 6 mg, Biotin 0.3 mg and Choline chlorids 3mg.

Each one Kg of mineral mixture contained: Zinc sulphat hepahydrate 3.0, Mg, sulphat 0.335, Coppous chloride 0.10, Calcium phosphate monobasic 135.8, Calcium Lactate 327.0, Ferric citrate 29.7, Potassium phosphate dibasic anhydrous 239.8, Sodium phosphate monobasic 87.2, Sodium chloride 43.6, Aluminium chloride anhydrous 0.15, Potassium iodide 0.15, Cobalt chloride 1.0, Sodium selenite 0.011 and L-cellulose 132.25 ( as g/Kg mineral mix) (**Gatlin and Wilson 1984**).

<sup>b</sup> Calculated using gross caloric values of 23.62, 39.52, and 17.15 kJ/g for protein, fat, and carbohydrate, respectively, according to **Brett (1973)**.

### **Growth Indices**

Mean final body weight (FBW) of each experimental treatment was determined by dividing total fish weight in each pen by number of fish. Weight gain (WG), specific growth rate (SGR), feed conversion ratio (FCR), were calculated using the following equations:

WG = Final body weight (g) - Initial body weight (g); SGR =  $(\ln \text{FBW} - \ln \text{IBW})/t \times 100$ ; where: FBW is final body weight (g); IBW is initial body weight (g); ln= natural logarithmic; t = time in days; FCR = Feed intake (g)/weight gain (g).

### Blood Samples and Analysis

Blood samples were collected at the end of the experiment. Each of the experimental treatment was sampled once for hematological indices analyses. The fish were euthanized with MS-222 and the blood samples were taken by puncturing the caudal vessels. Heparin was used as anticoagulant and the plasma separated by centrifugation at 3000 rpm for 20 min and stored at  $-20^{\circ}\text{C}$  until further analysis.

### Analytical Methods

At the beginning of the trial, a random pooled sample of 10 fish was collected, for determination of initial whole-body proximate composition. At the termination of the feeding trial, five fish were randomly selected from each pen, homogenized in a blender, to determine the final whole-body proximate composition. The fish were pooled for each pen, oven-dried, ground, and stored at  $-20^{\circ}\text{C}$  for subsequent analysis. The chemical composition of the whole fish body and diet was determined according to the procedure of AOAC, (1995). Dry matter was determined after drying the samples in an oven ( $105^{\circ}\text{C}$ ) for 24 hr. Ash by incineration at  $550^{\circ}\text{C}$  for 12 hr. Crude protein was determined by micro-Kjeldhal method, N%  $\times 6.25$  crude fat by soxhlet extraction with diethyl ether ( $40 - 60^{\circ}\text{C}$ ).

### Statistical Analysis

Data of the experiments were analyzed by two-way analysis of variance ANOVA. Significant differences were considered at  $P < 0.05$ . When significant differences were found, Duncan's multiple range tests was used to identify differences among experimental groups. All statistical analyses were performed using Duncan multiple range test at ( $P < 0.05$ ) level SPSS, (1997).

## RESULTS

Final body weight, specific growth rate and weight gain were significantly affected by NaCl supplementation, water salinity and their interaction ( $P < 0.05$ ; Table 2). Fish fed diets supplemented with 10% NaCl recorded significantly higher final body weight when compared to fish fed control either 10 ( $p < 0.05$ ) or 15 ppt ( $p \geq 0.05$ ) salinity water over a period of ten weeks. Fish fed diets supplemented with 10% NaCl recorded significantly higher weight gain when compared to fish fed control either 10 ppt ( $p < 0.05$ ) and 15 ppt ( $p < 0.05$ ) salinity water over all experimental period.

**Table (2) Growth performance of Nile tilapia juveniles at different experimental treatments**

Experimental treatments							
Dietary NaCl levels	D <sub>0</sub> %			D <sub>10</sub> %			P value
Water salinity levels, ppt	W <sub>0ppt</sub>	W <sub>10ppt</sub>	W <sub>15ppt</sub>	W <sub>0ppt</sub>	W <sub>10ppt</sub>	W <sub>15ppt</sub>	
Initial weight (g)	8.00±0.01	8.00±0.01	8.00±0.01	8.00±0.01	8.00±0.01	8.00±0.01	0.561
Final body weight (g)	43.00±0.68 <sup>a</sup>	36.00±1.30 <sup>cd</sup>	33.75±0.75 <sup>d</sup>	41.80±0.36 <sup>a</sup>	38.57±0.22 <sup>b</sup>	36.90±0.95 <sup>bc</sup>	0.024
Specific growth rate (%/day)	1.04±0.01 <sup>a</sup>	0.94±0.03 <sup>b</sup>	0.90±0.02 <sup>c</sup>	1.03±0.01 <sup>a</sup>	0.97±0.01 <sup>b</sup>	0.95±0.02 <sup>b</sup>	0.036
weight gain (g/fish)	35.00±0.68 <sup>a</sup>	28.00±1.30 <sup>cd</sup>	25.75±0.75 <sup>d</sup>	33.80±0.36 <sup>a</sup>	30.57±0.22 <sup>b</sup>	28.90±0.95 <sup>bc</sup>	0.024
Effect of dietary NaCl levels irrespective of water salinity levels on growth performance indices				Effect of water salinity levels irrespective of dietary NaCl levels on growth performance indices			
	D <sub>0</sub> %	D <sub>10</sub> %	P value	W <sub>0ppt</sub>	W <sub>10ppt</sub>	W <sub>15ppt</sub>	P value
Initial weight (g)	8.00±0.01	8.00±0.01	0.650	8.00±0.01	8.00±0.01	8.00±0.01	0.750
Final body weight (g)	37.58±1.33 <sup>b</sup>	39.09±0.78 <sup>a</sup>	0.031	42.40±0.44 <sup>a</sup>	37.29±0.63 <sup>b</sup>	35.33±0.82 <sup>c</sup>	0.0001
Specific growth rate (%/day)	0.96±0.02 <sup>b</sup>	0.98±0.01 <sup>a</sup>	0.033	1.04±0.01 <sup>a</sup>	0.96±0.01 <sup>b</sup>	0.93±0.02 <sup>c</sup>	0.0001
weight gain (g/fish)	30.17±1.33 <sup>b</sup>	31.09±0.78 <sup>a</sup>	0.031	34.40±0.44 <sup>a</sup>	29.62±0.63 <sup>b</sup>	27.867±0.34 <sup>c</sup>	0.0001

No significant effect of either dietary NaCl supplementation or water salinity and their interaction were observed on feed intake and feed conversion ratio. Irrespective of dietary NaCl levels, feed intake and feed conversion ratio affected significantly with water salinity ( $P < 0.05$ ; Table3). Feed intake decreased with increasing water salinity. No mortalities were recorded during 10 weeks .Survival (%) not affected with either dietary NaCl supplementations or water salinity and their interaction. Fish groups recorded the highest survival (%) (Table 3)

Table (3) Feed utilization of Nile tilapia juveniles at different experimental treatments

Dietary NaCl levels	D <sub>0%</sub>			D <sub>10%</sub>			P value
Water salinity levels, ppt	W <sub>0ppt</sub>	W <sub>10ppt</sub>	W <sub>15ppt</sub>	W <sub>0ppt</sub>	W <sub>10ppt</sub>	W <sub>15ppt</sub>	
Feed Intake	42.91±0.52 <sub>a</sub>	41.47±0.06 <sup>ab</sup>	40.17±0.21 <sup>b</sup>	43.06±0.11 <sup>a</sup>	41.06±0.77 <sup>b</sup>	39.74±0.59 <sup>b</sup>	0.820
Feed conversion (g feed/g gain)	1.30±0.02 <sup>d</sup>	1.49±0.07 <sup>ab</sup>	1.56±0.04 <sup>a</sup>	1.33±0.01 <sup>cd</sup>	1.39±0.02 <sup>bcd</sup>	1.46±0.07 <sup>abc</sup>	0.249
Effect of dietary NaCl levels irrespective of water salinity levels on Feed utilization				Effect of water salinity levels irrespective of dietary NaCl levels on Feed utilization			
	D <sub>0%</sub>	D <sub>10%</sub>	P value	W <sub>0ppt</sub>	W <sub>10ppt</sub>	W <sub>15ppt</sub>	P value
Feed Intake	41.52±0.4	41.29±0.56	0.613	42.99±0.24 <sup>a</sup>	41.26±0.36 <sup>b</sup>	39.96±0.01 <sup>c</sup>	0.0001
Feed conversion (g feed/g gain)	1.45±0.04 <sup>a</sup>	1.39±0.03 <sup>b</sup>	0.149	1.32±0.01 <sup>c</sup>	1.44±0.03 <sup>b</sup>	1.51±0.04 <sup>a</sup>	0.002

Irrespective of dietary NaCl supplementation, body moisture and ash contents showed significantly affect with different water salinity ( $P < 0.05$ ; Table4). No significant affect with dietary NaCl supplementation and their interaction with different water salinity, on body ether extract and ash content was recorded. Ash content insignificantly affected with dietary NaCl supplementation. Body crude protein content not significantly affected with either dietary NaCl supplementation or water salinity ( $P < 0.05$ ; Table 4).

Irrespective of water salinity, the blood serum concentrations of Na<sup>+</sup> has affected significantly ( $P < 0.05$ ) with dietary NaCl supplementation. Fish group fed diet containing 10% NaCl recorded the highest values of Na<sup>+</sup> concentration compared to other experimental groups (Table 5). The concentrations of K<sup>+</sup> in blood serum showed no significant differences either dietary NaCl supplementation or water salinity levels or their interaction ( $P < 0.05$ ; Table 5). The concentrations of Cl<sup>-</sup> in blood serum affected significantly ( $P < 0.05$ ) water salinity and their interaction but did not differ significantly with NaCl supplementation

The concentrations of Glucose in blood serum affected significantly ( $P < 0.05$ ) water salinity but did not differ significantly with NaCl supplementation and their interaction. The concentrations of Lactate in blood serum showed no significant differences with dietary NaCl supplementation, water salinity and their interaction ( $P < 0.05$ ; Table 5).

**Table (4) Proximate Body composition at different experimental treatments**

Experimental treatments							
Dietary NaCl levels	D <sub>0</sub> %			D <sub>10</sub> %			P value
Water salinity levels, ppt	W <sub>0ppt</sub>	W <sub>10ppt</sub>	W <sub>15ppt</sub>	W <sub>0ppt</sub>	W <sub>10ppt</sub>	W <sub>15ppt</sub>	
Moisture (%)	70.43±0.34	71.80±1.56	71.78±0.32	70.19±0.26	72.46±0.13	72.26±0.30	0.79
Ether extract (%)	19.47±0.14 <sup>a</sup>	18.75±0.26 <sup>ab</sup>	18.82±0.30 <sup>ab</sup>	19.01±0.39 <sup>ab</sup>	18.71±0.23 <sup>ab</sup>	18.44±0.10 <sup>b</sup>	0.682
Crude protein (%)	64.20±0.21 <sup>a</sup>	62.87±0.07 <sup>b</sup>	61.57±0.23 <sup>d</sup>	62.50±0.45 <sup>bc</sup>	61.90±0.32 <sup>cd</sup>	61.83±0.09 <sup>cd</sup>	0.035
Ash (%)	14.23±0.06 <sup>b</sup>	14.32±0.01 <sup>ab</sup>	14.62±0.22 <sup>a</sup>	14.53±0.05 <sup>ab</sup>	14.44±0.14 <sup>ab</sup>	14.28±0.09 <sup>ab</sup>	0.009
Effect of dietary NaCl levels irrespective of water salinity levels on Body composition				Effect of water salinity levels irrespective of dietary NaCl levels on Body composition			
	D <sub>0</sub> %	D <sub>10</sub> %	P value	W <sub>0ppt</sub>	W <sub>10ppt</sub>	W <sub>15ppt</sub>	P value
Moisture (%)	71.34±0.52	71.63±0.38	0.606	70.31±0.72	72.13±0.72	72.02±0.22	0.036
Ether extract (%)	19.01±0.17 <sup>a</sup>	18.72±0.16 <sup>b</sup>	0.182	19.24±0.21 <sup>a</sup>	18.73±0.16 <sup>b</sup>	18.63±0.17 <sup>b</sup>	0.074
Crude protein (%)	62.88±0.39	62.08±0.19	0.801	63.35±0.01	62.38±0.01	61.70±0.13	0.770
Ash (%)	14.39±0.09	14.41±0.06	0.803	14.38±0.08	14.38±0.07	14.45±0.13	0.801

**Table (5) The concentrations of Na<sup>+</sup>, K<sup>+</sup>, Cl<sup>-</sup>, Glucose, PH and Lactate in blood serum of Nile tilapia juveniles at different experimental treatments**

Experimental treatments							
Dietary NaCl levels	D <sub>0</sub> %			D <sub>10</sub> %			P value
Water salinity levels, ppt	W <sub>0ppt</sub>	W <sub>10ppt</sub>	W <sub>15ppt</sub>	W <sub>0ppt</sub>	W <sub>10ppt</sub>	W <sub>15ppt</sub>	
Na <sup>+</sup>	155.33±2.00 <sup>b</sup>	165.00±1.58 <sup>ab</sup>	164.22±0.59 <sup>ab</sup>	165.55±1.31 <sup>ab</sup>	166.56±2.28 <sup>ab</sup>	172.48±7.62 <sup>a</sup>	0.438
K <sup>+</sup>	6.76±0.31	5.29±0.61	6.22±0.57	4.76±0.54	5.12±0.91	5.52±0.59	0.343
Cl <sup>-</sup>	135.00±1.86 <sup>b</sup>	150.45±0.91 <sup>a</sup>	153.89±3.74 <sup>a</sup>	145.22±2.23 <sup>a</sup>	148.33±1.35 <sup>a</sup>	146.44±3.90 <sup>a</sup>	0.015
Glucose	57.44±7.16 <sup>b</sup>	81.44±4.70 <sup>ab</sup>	97.78±2.62 <sup>a</sup>	68.11±4.23 <sup>b</sup>	93.00±26.18 <sup>ab</sup>	106.33±8.21 <sup>a</sup>	0.992
PH	7.66±0.09 <sup>a</sup>	7.39±0.04 <sup>b</sup>	7.40±0.13 <sup>b</sup>	7.56±0.01 <sup>ab</sup>	7.43±0.07 <sup>ab</sup>	7.48±0.05 <sup>ab</sup>	0.462



Lactat	1.62±0.47	2.01±0.65	2.90±0.35	2.68±1.10	3.14±0.10	3.60±0.31	0.927
	Effect of dietary NaCl levels irrespective of water salinity levels on Na <sup>+</sup> , K <sup>+</sup> , Cl <sup>-</sup> , Glucose, PH and Lactate concentrations			Effect of water salinity levels irrespective of dietary NaCl levels on Na <sup>+</sup> , K <sup>+</sup> , Cl <sup>-</sup> , Glucose, PH and Lactate concentrations			
	D <sub>0%</sub>	D <sub>10%</sub>	P value	W <sub>0ppt</sub>	W <sub>10ppt</sub>	W <sub>15ppt</sub>	P value
Na <sup>+</sup>	161.52±1.73 <sup>b</sup>	168.20±2.59 <sup>a</sup>	0.033	160.44±2.52 <sup>c</sup>	165.78±1.29 <sup>b</sup>	168.35±3.92 <sup>a</sup>	0.100
K <sup>+</sup>	6.09±0.34 <sup>a</sup>	5.13±0.37 <sup>b</sup>	0.081	5.76±0.526	5.21±0.49	5.87±0.40	0.529
Cl <sup>-</sup>	146.45±3.16	146.67±1.43	0.918	140.11±2.627 <sup>c</sup>	149.39±0.87 <sup>ab</sup>	150.17±2.95 <sup>a</sup>	0.004
Glucose	78.89±6.40 <sup>b</sup>	89.15±9.78 <sup>a</sup>	0.312	62.78±4.419 <sup>c</sup>	87.22±12.17 <sup>b</sup>	102.06±4.30 <sup>a</sup>	0.020
PH	7.49±0.07	7.49±0.03	0.957	7.61±0.045	7.41±0.04	7.44±0.07	0.039
Lactat	2.18±0.32 <sup>b</sup>	3.14±0.36 <sup>a</sup>	0.068	2.15±0.584 <sup>b</sup>	2.58±0.39 <sup>b</sup>	3.25±0.26 <sup>a</sup>	0.211

**Table (6) The Viscero-somatic index (VSI), Hepato-somatic index (HSI), Spleno-somatic index (SSI) and Heart of Nile tilapia juveniles at different experimental treatments**

Experimental treatments							
Dietary NaCl levels	D <sub>0%</sub>			D <sub>10%</sub>			P value
Water salinity levels, ‰	W <sub>0ppt</sub>	W <sub>10ppt</sub>	W <sub>15ppt</sub>	W <sub>0ppt</sub>	W <sub>10ppt</sub>	W <sub>15ppt</sub>	
VSI	4.60±1.06	4.21±0.91	4.06±0.29	4.28±0.83	4.01±0.36	3.43±0.58	0.957
HSI	0.61±0.11	0.64±0.13	0.65±0.11	0.79±0.17	0.79±0.12	0.62±0.01	0.683
SSI	0.07±0.01	0.06±0.02	0.04±0.01	0.05±0.02	0.07±0.01	0.06±0.01	0.290
Heart	0.12±0.02	0.09±0.03	0.09±0.02	0.10±0.02	0.13±0.01	0.08±0.02	0.354
	Effect of dietary NaCl levels irrespective of water salinity levels on VSI, HSI, SSI and Heart			Effect of water salinity levels irrespective of dietary NaCl levels on VSI, HSI, SSI and Heart			
	D <sub>0%</sub>	D <sub>10%</sub>	P value	W <sub>0ppt</sub>	W <sub>10ppt</sub>	W <sub>15ppt</sub>	P value
VSI	4.29±0.42	3.90±0.33	0.527	4.44±0.61	4.11±0.44	3.74±0.32	0.644
HSI	0.63±0.06	0.74±0.08	0.355	0.70±0.10	0.71±0.08	0.64±0.09	0.819
SSI	0.05±0.01	0.06±0.01	0.312	0.06±0.01	0.06±0.01	0.05±0.01	0.469
Heart	0.10±0.01	0.12±0.01	0.762	0.11±0.01	0.11±0.02	0.09±0.01	0.388

The Viscero-somatic index (VSI), Hepato-somatic index (HSI) and Spleno-somatic index (SSI) insignificantly affected with dietary NaCl supplementation, water salinity and their interaction (Table 6).

## DISCUSSION

The salinity of the culture system is known to influence metabolism and homeostatic processes in fish, meaning the organism needs for nutrients and protein particularly, may differ in freshwater and saltwater systems (**Altinok and Gizzle 2001**). One strategy for increasing tolerance to adverse environments and maintaining growth rates in many species is a more efficient management of diet nutrient contents.

Since minerals absorbed from the water do not always meet the total metabolic requirements in fish, their supplementation through the diet promotes growth **Hepher, (1988)**. Fish diet is therefore an important source of salts not only to satisfy the needs for growth, but also for osmoregulation. Providing a sufficient amount of salt through feeds can spare energy that is used for osmoregulation, thereby reducing stress and allowing more energy for growth. Dietary salt has been found to be beneficial for growth in rainbow trout **MacLeod, (1978)**, common carp and mrigal **Nandeeshha *et al.*, (2000)**, Asian sea bass **Harpaz *et al.*, (2005)**, European sea bass (**Eroldogan, 2003; Eroldogan *et al.*, 2005**) and gilthead sea bream (**Appelbaum *et al.*, 2008a; Appelbaum and Arockiaraj, 2008b, 2009**).

Results of the current study show that fish fed diets supplemented with 10% NaCl recorded significantly better final body weight and weight gain when compared to fish fed control when held in 10 and 15 ppt salinity water. Feed intake and feed conversion ratio in the current study decreased with increasing water salinity reported by (**Boeuf and Payan, 2001**) reviewed the literature on salinity influence on growth in fish and concluded that salinity is also a key factor in controlling growth. They observed that the changes in growth rate that depend on salinity result from an action on metabolic rate, food intake and food conversion. Better growth at intermediate salinities (8-20 ppt) is very often but not systematically, correlated to a lower standard metabolic rate. Florida red tilapia physiologically function more efficiently in brackish and saline waters, with lower feed conversion ratios and faster weight gain than those raised in freshwater **Head *et al.*, (1994)**.

No mortalities were recorded during 10 weeks. Survival (%) not affected with either dietary NaCl supplementations or different water salinity. Fish groups recorded the highest survival (%). **Yao *et al.*, (2008)** reported that survival of Nile tilapia fingerlings transferred directly to saline water was increased (84.3% to 96.8%) up to 17 ppt, but mortalities were significant decreased above that level of salinity (60-70%). Higher survival (%) (78 to 81%) was achieved by gradual acclimation to salinity of 30 ppt over

two days. Results of **Lim *et al.*, (2006)** showed that juvenile Nile tilapia receiving dietary NaCl feeding regimens, even for a two-week period, exhibited consistently better survival values.

Salt tolerance in tilapia can be improved by optimizing acclimation protocols, adding salt to the diet. Results of the current study show moisture, ether extract and crude protein not significantly affected with NaCl supplementation. The contrast results were obtained by **Keshavanath *et al.*, (2012)**, who found that carcass composition was not affected as salt addition in fish diets.

To summarize, the results of this study confirm that the control balance of salt either in diet or water within a narrow limit is critical to life in all multicellular organisms, including teleost fishes. Salt tolerance is a term describing the overall fitness, or productivity, of the fish in a saline environment. The results of the present study revealed that juveniles of Nile tilapia has been documented as surviving in salinities up to 15 ppt with acclimation. The results demonstrating the beneficial effect that supplemental dietary salt can have beneficial effects on fish growth.

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