



## Feminization by 17 $\alpha$ -Ethinylestradiol of The Progeny of XY-Female Nile Tilapia (*Oreochromis niloticus*). Effects on Growth, Condition Factor and Gonadosomatic Index

Varinia Juárez-Juárez<sup>1</sup>, Juan Pablo Alcántar-Vázquez<sup>1\*</sup>, Carolina Antonio-Estrada<sup>1</sup>, José Antonio Marín-Ramírez<sup>1</sup>, Raúl Moreno-De La Torre<sup>1</sup>

<sup>1</sup>Universidad del Papaloapan (UNPA), Laboratorio de Acuicultura, Des: Ciencias Agropecuarias. Av. Ferrocarril s/n, Col. Ciudad Universitaria, Loma Bonita, Oaxaca. C.P. 68400.

\* Corresponding Author: Tel.: +52.281 8729230;  
E-mail: jupasoul@hotmail.com

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### Abstract

Production of YY-males is one of the most important alternatives to the commercial use of hormones for producing monosex populations in the culture of Nile tilapia. One of the final stages of YY-technology is the feminization of the progeny of XY-females in order to obtain YY-females, which when crossed with YY-males makes possible to obtain populations composed of 100% YY-males. The aim of the study was to evaluate the effects of four concentrations (100, 200, 300, 400 mg kg<sup>-1</sup>) of estrogen 17 $\alpha$ -ethinylestradiol (EE<sub>2</sub>) on the sex ratio, growth, condition factor (CV) and gonadosomatic index (GSI) of the progeny of XY-females. Significant differences were observed in wet weight, total length and CV during the experiment; however, final values in EE<sub>2</sub>-treated groups showed not significant differences with control group. The observed proportion of females deviated significantly from the proportion of females expected in the progeny of XY-females in all groups fed EE<sub>2</sub>; however, only 90% feminization was attained. The group fed 200 mg kg<sup>-1</sup> of EE<sub>2</sub> showed a significantly higher value of GSI in comparison to that observed in the control group. By increasing the concentration of EE<sub>2</sub> it is possible to increase the proportion of females without affecting growth or GSI.

**Keywords:** Sex-reversal, YY-females, gonadal development, YY-males.

### Introduction

In mixed-sex cultures, Nile tilapia (*Oreochromis niloticus*) (Linnaeus, 1758) reach sexual maturity long before reaching commercial size (Jiménez & Arredondo, 2000; Arboleda-Obregón, 2005). Therefore, in commercial farming of Nile tilapia, reproduction during grow-out is a major problem, leading to the presence of fry and juveniles that overpopulate ponds, and ultimately resulting in a wide range of fish sizes at harvest instead of the larger and more uniform fish expected from the original stocking (Mair, Abucay, Skibinski, & Beardmore, 1997; Tariq-Ezaz, Myers, Powell, McAndrew, & Penman, 2004). To avoid this, a series of techniques have been developed to produce a monosex, all-male population in tilapia culture. Production of all-male populations eliminates sex behavior and therefore uncontrolled reproduction, allowing the production of marketable-sized fish (Varadaraj, 1989; Ponzoni, Hamzah, Tan, & Kamaruzzaman, 2005). Additionally, all-male tilapia cultures show better growth than mixed cultures, and for many years the production of all-male populations has been recognized as the most effective technique to increase tilapia production under commercial culture

conditions (Mair *et al.*, 1997; Müller & Hörstgen 2007; Nonglak, Boonanuntanasarn, Jangprai, Yoshizaki, & Nanakorn, 2012).

Hormonal sex reversal during fry period is the most commonly used method to produce all-male populations. However, in recent years this method has been increasingly contested since there is a growing concern worldwide about the accumulation of hormones in the environment (especially in marine waters) and a growing number of consumers who do not want to eat products that have been treated with hormones (Piferrer, 2001; Müller & Hörstgen, 2007; Leet, Gall, & Sepúlveda, 2011) and demand environmentally friendly production techniques.

One strong alternative, on a commercial scale, is the production of YY-males, which when combined with normal females (XX) produce progenies composed, in theory, of 100% genetic males. The first step in YY-technology involves using hormones to feminize sexually undifferentiated male fry (XY), which allows to obtain XY-females (Vera-Cruz, Mair, & Marino, 1996; Mair *et al.*, 1997). These females will be used to produce YY-males in a percentage of 25% when combined with normal males (XY).

Production of YY-males in recent years at the

Universidad del Papaloapan has led to attempts to optimize YY-technology. One alternative for optimizing the obtention of YY-male Nile tilapia is the feminization of the progeny of XY-females. This feminization will allow the production of two genotypes; XY-females and females with two Y chromosomes (YY-females). The YY-females will be used to produce populations composed of 100% YY-males when crossed with YY-males (Figure 1).

Feminization in our laboratory of progeny of normal females (XX) using the synthetic hormone  $17\alpha$ -ethinylestradiol ( $EE_2$ ) has shown good results (>80%) in low concentrations. The application of  $EE_2$  in higher concentrations could achieve successful feminization of the YY-genotype, which has been reported to be more difficult in comparison with the process for feminizing normal progeny (Abucay & Mair, 2004).

## Material and Methods

### Broodstock

XY-female and normal male (XY) Nile tilapia used in this research were produced at the Aquaculture Station of the Universidad del Papaloapan using broodstock from locally available strains obtained from the Centro Acuicola de Temascal, Oaxaca, México and the Granja Unidad de Producción del Tesechoacan, Veracruz, Mexico. XY-females were produced by feeding fry at swim-up stage with  $120 \text{ mg kg}^{-1}$   $17\beta$ -estradiol-treated feed (53% protein) for 30 days. They were reared for 18 months in the aquaculture station of the Universidad del Papaloapan and fed twice a day with commercial pellets (32% protein, Nutripec, Agribands Purina, Irapuato Gto. Mexico).

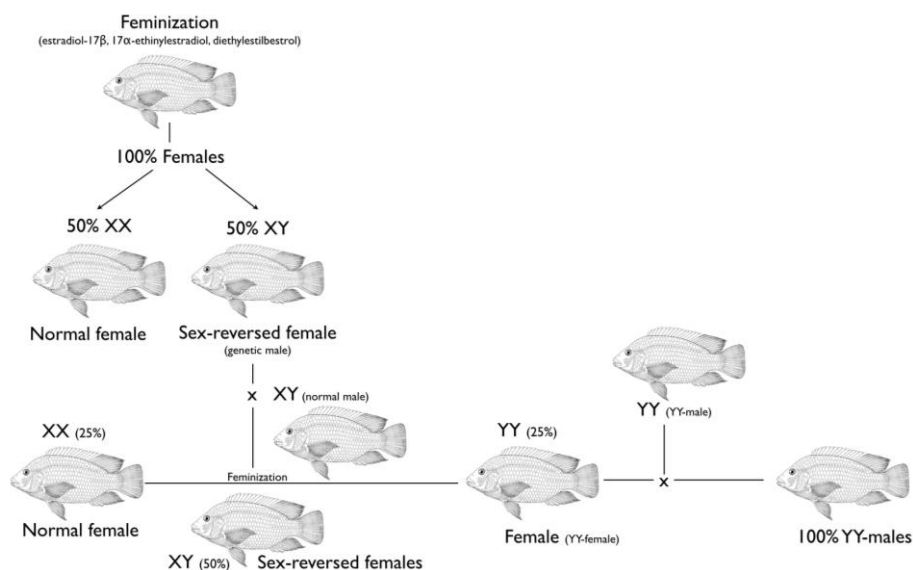
### Hormonal Treatment

The synthetic hormone  $17\alpha$ -ethinylestradiol ( $EE_2$ , Sigma Aldrich Chemical Co., St Louis, MO, USA) was added to commercial fish meal (<0.35 mm, 53% protein, 15% lipids) using the method described by Guerrero (1975). In brief, selected concentrations of  $EE_2$  were dissolved in 500 ml of 95% ethanol, sprayed over the food and maintained at room temperature for six hours to allow the alcohol to evaporate. Four levels of  $EE_2$  hormone treatments were added to the food: 100, 200, 300 and  $400 \text{ mg kg}^{-1}$ . The control group's food was handled in exactly the same manner with the exception of the added hormone.

### Fry Production and Experimental Conditions

*O. niloticus* broodstock were distributed at a male: XY-female ratio of 1:3 (Marín-Ramírez, Alcántar-Vázquez, Antonio-Estrada, Moreno-de la Torre, & Calzada-Ruiz, 2016) in two 3-m diameter outdoor concrete tanks (28-30 °C) supplied with fertilized water. Fry were collected 15 d later with a fine-mesh net after 90% of the water in the tanks had been siphoned. The recently hatched and sexually undifferentiated fry (~0.02 g wet weight and 8 mm length) were pooled, transported to a closed recirculating system and randomly divided into 15 acrylic aquaria of 85-L (three aquaria per treatment) at an initial stocking density of  $1 \text{ fry L}^{-1}$ . The water in the recirculating system was filtered using a mechanical filter (Hayward, Model S310T2, Hayward Pool Products Inc., Elizabeth, NJ, USA) and a bio-filter containing only plastic bio-balls (Aquatic Eco-System, Model CBB1, Pentair Ltd. Apopka, FL, USA).

Hormonal treatment lasted for 20 d under a



**Figure 1.** Diagram that represents the methodology applied to obtain XY-females and YY-females by feminization of XY and YY-genotypes. With information from Mair *et al.* (1997).

photoperiod of 12 L: 12 D and with water temperature adjusted thermostatically at  $26.0 \pm 0.5$  °C. Fry were fed at 1-h intervals at a feed ratio of 20% of their total body weight. Water flow was closed in all aquaria for 20 minutes after the estrogen-enriched feeds were offered in order to encourage feeding. Random samples of 25% of fry per replicate were collected every 10 d. Mean wet weight was obtained using a digital scale ( $\pm 0.01$ ), and total length was recorded to the nearest 0.01 mm from a digitized image using an imaging software (ImageJ version 1.36). Once the hormonal treatment was completed, the fry were fed with untreated commercial diet (50% protein) for 10 more days until the fry period was finished. Water temperature and dissolved oxygen were monitored daily using a multiparameter display system (YSI model 655, Yellow Springs Instrument Co., Inc., Yellow Springs, OH, USA). The aquaria were siphoned daily to remove feces and dead fry.

At the end of the fry period, all juveniles from each treatment were counted, weighed and measured for the calculation of survival rate and to determine average weight and length. Juveniles were then transferred to 3-m diameter outdoor concrete tanks (maintaining replicates per treatment) supplied with fertilized water and reared to sexual maturity (four months of age). During this period, juveniles were fed six times a day with untreated commercial diet (44 and 40% protein) for approximately 30 d, followed by a commercial diet at 35% protein three times a day for another 30 d and subsequently a commercial diet with 25% protein until the end of the experiment. Total length and wet weight were registered every 21 d for 60 random fish per treatment (20 fish per replicate).

### Fulton's Condition Factor

The value of K was calculated throughout the experiment (at each sampling procedure) following Froese (2006):

$$K = 100 * [\text{fish weight (g)} / \text{fish length (cm)}^3]$$

### Evaluation of Sex Ratio

The sex of 60 fish per treatment (20 fish per each replicate) was determined by removing the gonad. Gonads were classified as ovaries or testes.

### Gonadosomatic Index

The extracted gonads were weighed using a digital scale ( $\pm 0.01$ ) to calculate the gonadosomatic index (Sturm, 1978) using the following formula:

$$\text{GSI} = [\text{Gonad weight (g)} / \text{Fish weight (g)}] \times 100.$$

### Statistics

Differences in total length, wet weight and condition factor were analyzed using a one-way analysis of variance. The percentage of survival was arcsine transformed and analyzed using a one-way analysis of variance. The proportion of females identified in each treatment was tested against the 3:1 (male: female) expectation using a chi square test at a probability of 0.1% ( $P < 0.001$ ). Differences in gonadosomatic index between different treatments were analyzed using a Kruskal-Wallis nonparametric analysis (Zar, 1984).

### Results

#### Fry Period

Mean wet weight, total length, condition factor and the percentage of survival of control group fry and EE<sub>2</sub>-treated fry is shown in Table 1. Significant differences ( $P < 0.05$ ) were observed in wet weight and total length at 10 and 20 d of age. At 10 d of age wet weight was significantly higher ( $P < 0.05$ ) in the

**Table 1.** Wet weight (WW), total length (TL), condition factor (CF) and percentage of survival (S) ( $\pm$ S.E.) (n=60 fish) of Nile tilapia (*Oreochromis niloticus*) fed different concentrations of 17 $\alpha$ -ethinylestradiol for 20 days

Days	Hormone concentration (mg kg <sup>-1</sup> )				
	Control	100	200	300	400
0					
WW	0.02 $\pm$ 0.01 <sup>a</sup>	0.02 $\pm$ 0.01 <sup>a</sup>	0.02 $\pm$ 0.01 <sup>a</sup>	0.02 $\pm$ 0.01 <sup>a</sup>	0.02 $\pm$ 0.01 <sup>a</sup>
TL	0.80 $\pm$ 0.01 <sup>a</sup>	0.80 $\pm$ 0.01 <sup>a</sup>	0.80 $\pm$ 0.01 <sup>a</sup>	0.80 $\pm$ 0.01 <sup>a</sup>	0.80 $\pm$ 0.01 <sup>a</sup>
CF	3.90 $\pm$ 0.01 <sup>a</sup>	3.90 $\pm$ 0.01 <sup>a</sup>	3.90 $\pm$ 0.01 <sup>a</sup>	3.90 $\pm$ 0.01 <sup>a</sup>	3.90 $\pm$ 0.01 <sup>a</sup>
10					
WW	0.07 $\pm$ 0.01 <sup>c</sup>	0.09 $\pm$ 0.02 <sup>a</sup>	0.05 $\pm$ 0.01 <sup>d</sup>	0.06 $\pm$ 0.01 <sup>c</sup>	0.08 $\pm$ 0.01 <sup>b</sup>
TL	1.21 $\pm$ 0.03 <sup>ab</sup>	1.23 $\pm$ 0.04 <sup>ab</sup>	1.16 $\pm$ 0.04 <sup>b</sup>	1.20 $\pm$ 0.04 <sup>ab</sup>	1.27 $\pm$ 0.03 <sup>a</sup>
CF	4.28 $\pm$ 0.10 <sup>a</sup>	5.00 $\pm$ 0.12 <sup>a</sup>	3.20 $\pm$ 0.12 <sup>c</sup>	3.48 $\pm$ 0.09 <sup>bc</sup>	3.83 $\pm$ 0.11 <sup>b</sup>
20					
WW	0.51 $\pm$ 0.01 <sup>b</sup>	0.49 $\pm$ 0.02 <sup>bc</sup>	0.47 $\pm$ 0.01 <sup>cd</sup>	0.48 $\pm$ 0.01 <sup>bc</sup>	0.64 $\pm$ 0.02 <sup>a</sup>
TL	2.27 $\pm$ 0.05 <sup>b</sup>	2.12 $\pm$ 0.07 <sup>c</sup>	2.08 $\pm$ 0.07 <sup>c</sup>	2.12 $\pm$ 0.06 <sup>c</sup>	2.34 $\pm$ 0.05 <sup>a</sup>
CF	7.48 $\pm$ 0.10 <sup>a</sup>	5.20 $\pm$ 0.11 <sup>b</sup>	5.22 $\pm$ 0.09 <sup>b</sup>	5.11 $\pm$ 0.12 <sup>b</sup>	5.00 $\pm$ 0.11 <sup>b</sup>
S (%)	99.4 $\pm$ 0.3 <sup>a</sup>	96.1 $\pm$ 1.1 <sup>b</sup>	91.3 $\pm$ 0.8 <sup>c</sup>	95.6 $\pm$ 1.3 <sup>b</sup>	97.6 $\pm$ 0.9 <sup>b</sup>

Percentage of survival significantly different from the control group  $P < 0.05$ . Values in each row superscripted with different letters indicate significant differences between doses ( $P < 0.05$ ).

concentration of 100 mg kg<sup>-1</sup>, followed by the concentration of 400 mg kg<sup>-1</sup>, while at 20 d of age the concentration of 400 mg kg<sup>-1</sup> registered a significantly higher (P<0.05) value of wet weight, followed by the control group. A significantly higher (P<0.05) value of total length was shown for the concentration of 400 mg kg<sup>-1</sup> at 10 and 20 d of age. Significantly lower (P<0.05) values of wet weight and total length were observed in the concentration of 200 mg kg<sup>-1</sup> at 10 and 20 d of age.

Fulton's condition factor was significantly higher (P<0.05) at 10 d of age in the control group and the concentration of 100 mg kg<sup>-1</sup>. A significantly lower (P<0.05) condition factor was shown for the concentration of 200 mg kg<sup>-1</sup>. At 20 d of age, the control group showed a significantly higher (P<0.05) condition factor in comparison to the EE<sub>2</sub>-treated groups. No significant differences were observed between any of the EE<sub>2</sub>-treated groups.

The percentage of survival at the end of the hormonal treatment was significantly higher (P<0.05) in the control group compared to the EE<sub>2</sub>-treated groups. Between EE<sub>2</sub>-treated groups, a significantly lower (P<0.05) percentage of survival was detected in the concentration of 200 mg kg<sup>-1</sup>.

### Sex Ratio and Gonadosomatic Index

Sex-reversed females were produced in all EE<sub>2</sub>-treated groups. The results showed that all concentrations of EE<sub>2</sub> produced progeny with a significantly higher (P<0.001) proportion of females than the 3:1 (male: female) sex ratio predicted for XY-females (Table 2). The proportion of females was higher in the concentration of 400 mg kg<sup>-1</sup>; however, it was not possible to obtain a progeny composed of 100% females (Table 2). Significant differences (P<0.05) were observed in the gonadosomatic index, with the fish of the concentration of 200 mg kg<sup>-1</sup> showing a significantly higher (P<0.05) value than the fish of the control group. No significant differences were observed between the fish of the EE<sub>2</sub>-treated groups (Table 2).

### Survival and Growth

Final survival was significantly lower (P<0.05) in the EE<sub>2</sub>-treated groups than in the control group

(Table 2). There were significant differences in wet weight and total length between groups throughout the entire post-treatment stage of the experiment (Table 3). A significantly lower (P<0.05) wet weight was observed in the concentration of 200 mg kg<sup>-1</sup> at 41 and 62 d of age compared to the control group and the rest of the EE<sub>2</sub>-treated groups. At 83 and 104 d of age, significantly higher (P<0.05) values of wet weight were registered in the fish of the control group and the concentrations of 400 and 100 mg kg<sup>-1</sup>, respectively. At the end of the experiment (125 d of age), no significant differences were observed between the control group and all the EE<sub>2</sub>-treated groups. However, a significantly lower (P<0.05) wet weight value was observed in the fish of the concentration of 400 mg kg<sup>-1</sup> compared to the concentration of 100 mg kg<sup>-1</sup>.

Total length was significantly higher (P<0.05) in the fish of the control group at 41 and 62 d of age compared to all the EE<sub>2</sub>-treated groups. Between EE<sub>2</sub>-treated groups, the concentration of 200 mg kg<sup>-1</sup> registered significantly lower (P<0.05) values of total length. At 83 and 104 d of age, no significant differences were observed between the control group and the concentrations of 100, 300 and 400 mg kg<sup>-1</sup>. The fish of the concentration of 200 mg kg<sup>-1</sup> showed a significantly lower (P<0.05) value of total length. At the end of the experiment, similar to the findings for wet weight, no significant differences were observed between the fish of the control group and those of the EE<sub>2</sub>-treated groups. However, a significantly lower (P<0.05) value of total length was observed in the group that received the highest concentration of EE<sub>2</sub> (400 mg kg<sup>-1</sup>) in comparison to the group that received the lowest concentration of EE<sub>2</sub> (100 mg kg<sup>-1</sup>).

### Fulton's Condition Factor

Significant differences were observed in the condition factor at 41 d of age, with a significantly lower (P<0.05) value in the control group in comparison to the EE<sub>2</sub>-treated groups. Between the groups fed EE<sub>2</sub>, a significantly lower condition factor was registered for the concentration of 200 mg kg<sup>-1</sup>. At 62 d of age, a significantly higher (P<0.05) condition factor was observed in the concentration of 200 mg kg<sup>-1</sup> compared to the control group and the

**Table 2.** Percentage of survival, percentage of males and females, and gonadosomatic index (GSI) of four-months-old Nile tilapia (*Oreochromis niloticus*) fed different concentrations of 17 $\alpha$ -ethinyloestradiol for 20 days

Hormone dosage (mg kg <sup>-1</sup> )	NFE	Survival (%)	Male (%)	Female (%)	GSI
Control	60	99	81	19	0.94±0.23 <sup>b</sup>
100	60	87*	21	79 <sup>1</sup>	1.41±0.28 <sup>ab</sup>
200	60	72*	11	89 <sup>1</sup>	1.89±0.27 <sup>a</sup>
300	60	82*	16	84 <sup>1</sup>	1.39±0.21 <sup>ab</sup>
400	60	88*	10	90 <sup>1</sup>	1.11±0.21 <sup>ab</sup>

NFE = Number of fish evaluated per treatment (20 fish per replicate). \* Percentage of survival significantly different from the control group P<0.05. <sup>1</sup>Significantly different from the expected 3:1 distribution (P<0.001). Values in each column superscripted with different letters significant differences between doses (P<0.05).

**Table 3.** Wet weight (WW), total length (TL) and condition factor (CV) ( $\pm$  S.E.) (n = 60 fish) at four-months-old of Nile tilapia (*Oreochromis niloticus*) fed with 17 $\alpha$ -ethinylestradiol at different concentrations for 20 days

Days	Hormone concentration (mg kg <sup>-1</sup> )				
	Control	100	200	300	400
41					
WW	7.35 $\pm$ 0.12 <sup>a</sup>	6.12 $\pm$ 0.22 <sup>a</sup>	4.17 $\pm$ 0.25 <sup>b</sup>	5.15 $\pm$ 0.14 <sup>a</sup>	7.12 $\pm$ 0.19 <sup>a</sup>
TL	4.89 $\pm$ 0.07 <sup>a</sup>	4.16 $\pm$ 0.10 <sup>c</sup>	3.79 $\pm$ 0.12 <sup>d</sup>	3.93 $\pm$ 0.11 <sup>c</sup>	4.33 $\pm$ 0.11 <sup>bc</sup>
CF	6.28 $\pm$ 0.10 <sup>c</sup>	8.50 $\pm$ 0.15 <sup>a</sup>	7.66 $\pm$ 0.18 <sup>b</sup>	8.48 $\pm$ 0.13 <sup>a</sup>	8.77 $\pm$ 0.11 <sup>a</sup>
62					
WW	13.30 $\pm$ 0.45 <sup>a</sup>	11.82 $\pm$ 0.55 <sup>a</sup>	8.79 $\pm$ 0.55 <sup>b</sup>	11.45 $\pm$ 0.55 <sup>a</sup>	12.19 $\pm$ 0.54 <sup>a</sup>
TL	8.80 $\pm$ 0.12 <sup>a</sup>	8.03 $\pm$ 0.22 <sup>db</sup>	7.02 $\pm$ 0.13 <sup>e</sup>	8.29 $\pm$ 0.16 <sup>cb</sup>	8.18 $\pm$ 0.18 <sup>b</sup>
CF	1.95 $\pm$ 0.10 <sup>b</sup>	2.28 $\pm$ 0.14 <sup>ab</sup>	2.54 $\pm$ 0.09 <sup>a</sup>	2.00 $\pm$ 0.11 <sup>b</sup>	2.22 $\pm$ 0.14 <sup>ab</sup>
83					
WW	40.1 $\pm$ 1.56 <sup>a</sup>	34.78 $\pm$ 1.59 <sup>ab</sup>	30.89 $\pm$ 1.34 <sup>b</sup>	32.72 $\pm$ 1.37 <sup>b</sup>	36.29 $\pm$ 1.51 <sup>a</sup>
TL	12.30 $\pm$ 0.16 <sup>a</sup>	11.65 $\pm$ 0.18 <sup>a</sup>	11.39 $\pm$ 0.17 <sup>b</sup>	11.87 $\pm$ 0.16 <sup>a</sup>	12.01 $\pm$ 0.16 <sup>a</sup>
CF	2.09 $\pm$ 0.10 <sup>a</sup>	2.08 $\pm$ 0.09 <sup>a</sup>	2.02 $\pm$ 0.07 <sup>a</sup>	1.88 $\pm$ 0.09 <sup>b</sup>	2.12 $\pm$ 0.08 <sup>a</sup>
104					
WW	52.43 $\pm$ 2.58 <sup>a</sup>	50.63 $\pm$ 0.02 <sup>a</sup>	39.89 $\pm$ 2.92 <sup>b</sup>	48.38 $\pm$ 2.48 <sup>ab</sup>	44.68 $\pm$ 2.53 <sup>b</sup>
TL	13.35 $\pm$ 0.81 <sup>a</sup>	13.10 $\pm$ 0.31 <sup>a</sup>	11.85 $\pm$ 0.37 <sup>b</sup>	13.15 $\pm$ 0.26 <sup>a</sup>	12.33 $\pm$ 0.34 <sup>ab</sup>
CF	1.66 $\pm$ 0.12 <sup>b</sup>	2.07 $\pm$ 0.9 <sup>a</sup>	2.08 $\pm$ 0.10 <sup>a</sup>	2.09 $\pm$ 0.11 <sup>a</sup>	2.15 $\pm$ 0.15 <sup>a</sup>
125					
WW	82.85 $\pm$ 2.26 <sup>ab</sup>	89.10 $\pm$ 3.03 <sup>a</sup>	82.96 $\pm$ 2.40 <sup>ab</sup>	80.66 $\pm$ 2.40 <sup>ab</sup>	78.02 $\pm$ 2.26 <sup>b</sup>
TL	15.93 $\pm$ 0.15 <sup>ab</sup>	15.94 $\pm$ 0.20 <sup>a</sup>	15.45 $\pm$ 0.22 <sup>ab</sup>	15.57 $\pm$ 0.17 <sup>ab</sup>	15.21 $\pm$ 0.15 <sup>b</sup>
CF	2.03 $\pm$ 0.8 <sup>a</sup>	2.22 $\pm$ 0.11 <sup>a</sup>	2.22 $\pm$ 0.09 <sup>a</sup>	2.12 $\pm$ 0.08 <sup>a</sup>	2.19 $\pm$ 0.07 <sup>a</sup>

Values in each row superscripted with different letters indicate significant differences between doses (P<0.05).

concentration of 300 mg kg<sup>-1</sup>. At 83 and 104 d of age, a significantly lower (P<0.05) condition factor was registered for the concentration of 300 mg kg<sup>-1</sup> and the control group, respectively. Finally, at the end of the experiment no significant differences were observed between the control group and the EE<sub>2</sub>-treated groups.

## Discussion

Feminization of sexually undifferentiated fry is one of the critical stages of YY-technology since it ensures the production of XY-females (sex-reversed females) that will be used in the production of YY-males. Therefore, the feminization of the progeny of females with XY-genotype will allow us to achieve two goals simultaneously; to obtain new XY-females (product of feminizing normal males) and YY-females (product of feminizing YY-males) (Alcántar-Vázquez, Moreno-de la Torre, Calzada-Ruiz, & Antonio-Estrada, 2014). The YY-females, sometimes called neo-females, will allow us to obtain monosex populations composed of 100% YY-males when crossing them with YY-males. This is the final step in YY-technology, as it allows the obtention of a sufficient number of YY-males in order to form broodstocks that can be crossed with normal females (XX) to obtain populations composed of 100% genetic males for commercial purposes.

In this work, an increase in the proportion of females by increasing the concentration of EE<sub>2</sub> was observed. Hopkins *et al.* (1979) report a similar increase in the proportion of female blue tilapia (*Oreochromis aureus*) (Steindachner, 1864) as the

concentration of EE<sub>2</sub> increased. In the same species, Mélard (1995) achieved 100% females using a diet with 200 mg kg<sup>-1</sup> of EE<sub>2</sub>. In this experiment it was not possible to obtain a complete feminization, probably caused by the use of XY-females which normally produce a percentage of males of approximately 75% compared with 50% males produced normally by XX-females. Additionally, it has been reported that the feminization of YY-males (present in approximately 25% of the progeny of XY-females, Mair *et al.*, 1997) may be more complicated than the feminization of XY-males (Abucay & Mair, 2004). This is supported by the findings of Shved *et al.* (2007) and Lázaro-Velasco (2014) after using EE<sub>2</sub> in balanced progenies of Nile tilapia. Both experiments achieve higher feminization rates at 120 mg kg<sup>-1</sup> (83%) and 125 mg kg<sup>-1</sup> (86.5%) when compared to that observed in our work at similar concentrations of EE<sub>2</sub> applied in our experiment. Finally, Genotte, Mélard, D'Cotta, Baroiller, and Rougeot (2014) report in Nile tilapia no sex-reversal of YY-embryos exposed to 6500  $\mu$ g L<sup>-1</sup> of EE<sub>2</sub> for 4 h, whereas sex-reversal rates in XY-embryos ranged from 0-60% (2000  $\mu$ g L<sup>-1</sup>). However, in spite of the above, we cannot rule out that the sex ratios obtained could be explained by the interaction of the three components that, according to Baroiller, D'Cotta, Bezault, Wessel, and Hoerstgen-Schwark (2009) govern sex in the Nile tilapia; a complex genetic sex determination system with a major determinant locus (sex chromosomes XX/XY), some minor genetic factors (parental factors) and the influence of the temperature (environmental factors) during the fry period.

It is considered that estrogens, either natural or

synthetic, do not have a positive effect on the growth of teleosts, unlike androgens which are masculinizing hormones (Johnstone, Simpson, & Walker, 1979; Piferrer, 2001). However, a decrease in growth has been reported in several species of fish, especially when the estrogen concentration is increased (Piferrer, 2001; Marín-Ramírez *et al.*, 2016). Marín-Ramírez *et al.* (2016) reported a reduced growth after feeding Nile tilapia fry with different concentrations of diethylstilbestrol for 20 days. Using the same synthetic estrogen, Król, Poblocki, Bockenheimer, and Hliwa (2014) report in the European catfish *Silurus glanis* (Linnaeus, 1758) a decrease in growth, while the natural estrogen estradiol-17 $\beta$  does not cause such decrease in growth. However, an increase in growth has been observed when using synthetic (diethylstilbestrol) or natural (17 $\beta$ -estradiol) estrogens in several species (Varadaraj, 1989; Chiba, Iwatsukis, Hayami, & Yamauchi, 1993). In the present work, no differences in growth were observed between the control group and the groups fed with EE<sub>2</sub>, except for the highest concentration (400 mg kg<sup>-1</sup>). Although no significant differences were detected, it was observed that the group fed with the concentration of 100 mg kg<sup>-1</sup> recorded a higher wet weight compared to the control group.

These results and those observed during the fry period indicate that in Nile tilapia the synthetic estrogen EE<sub>2</sub> probably has an anabolic effect at physiological level during the sex differentiation period, which allows to achieve a higher growth rate rather than reduced as normally seen in other fish species (Piferrer, 2001). This anabolic effect has been reported in other species of the genus *Oreochromis* when using another synthetic estrogen (Varadaraj, 1989). Herman and Kincaid (1988) indicate that different metabolic pathways may be involved in both cases. However, in our case this anabolic effect was opposite (inverted) in the two growth stages, with the highest growth observed in the 400 mg kg<sup>-1</sup> concentration of EE<sub>2</sub> during the fry stage, while during the post-treatment part of the experiment the 100 mg kg<sup>-1</sup> concentration of EE<sub>2</sub> had the best results.

The growth rate obtained in this work becomes more relevant when we take into account that in the control group the proportion of males, which in Nile tilapia are characterized as having a higher growth rate than females, was over 80%, while in the groups fed with EE<sub>2</sub> the proportion of males only reached a maximum of 21%. The sex-reversed females obtained do not showed a reduction in growth rate, which could be expected in XY-females, associated with an increase of the physiological cost of producing eggs rather than producing sperm, especially in genetic males that were sex-reversed to phenotypic females (Hayward & Gillooly, 2011; Geffroy & Bardonnat, 2016). In this regard, Hayward and Gillooly (2011) report that gamete biomass in females is approximately two to four orders of magnitude higher compared to males in several animal species,

including fish. This growth rate was probably caused, as noted previously, by an anabolic effect of the EE<sub>2</sub>. However, we cannot rule out two other possibilities, the first being that the relation between growth and sex ratio is established for Nile tilapia before rather than during the sex differentiation period as observed for Diaz, Ribas and Piferrer (2013) in *Dicentrarchus labrax* (Linneo, 1758), in which case the growth rates observed in the phenotypic females (sex-reversed females) would be in accordance to that of the genetic males originally presented in the population. The second possibility is that final growth, particularly the wet weight, could be affected by the GSI obtained since female Nile tilapia normally have greater GSI than males as a result of having heavier gonads.

Condition factors including Fulton's condition factor are an accepted criterion to assess the well being or fitness of a fish (nutritional status), defined by the amount of energy possessed (available fat content) by that fish to carry on several functions, including growth and reproduction (Jones, Petrell, & Pauly, 1999; Neff & Cargnelly, 2004; Gupta, Haque, & Khan, 2012). The condition factor also can be used to assess physiological status under a potential stressor (Hoque, Yusoff, Law, & Syed, 1998) such as a feeding fry with a synthetic estrogen. In our work, condition factor was highly variable during the fry period and initial stages of the juvenile period (41 d of age). Weatherley (1972) suggests that this could be caused by fluctuations in metabolic balance, patterns of maturation and the state of fullness of the alimentary canal. However, during the time of hormonal treatment it was possible to observe in general a lower condition factor in the fish of the EE<sub>2</sub>-treated groups compared to the fish of the control group. Hoque *et al.* (1998) mention that this decrease could be interpreted as depletion of energy reserves stored as liver glycogen or body fat. A reduction in food intake could provoke such effect, especially at early stages of development. This reduction has been observed in several sex-reversal experiments carried out in our laboratory. Once hormonal treatment finished and fish reached the onset of sexual maturation (62 d of age), condition factor remained above 2.0 for almost all groups until the end of the experiment. Similar values have been reported in normally growing, healthy Nile tilapia and their hybrids of similar weight and length (Little, 1989; Fish Breeding Association, 2003; El-Saidy & Gaber, 2005; Crab, Kochva, Verstraete, & Avnimelech, 2009; Gupta *et al.*, 2012). These results indicate that once the hormonal treatment ends, a compensatory growth takes place, caused probably by an increase in food intake and accumulation of energy in the form of available fat. The fact that final growth was statistically similar for the control group and the groups fed EE<sub>2</sub> support this.

Piferrer (2001) mentions that the application of estrogen, either natural or synthetic, can adversely affect survival in the vast majority of teleosts,

especially if a certain threshold is exceeded. This is related to a number of factors, among which are: the species, the type of hormone, its concentration and the duration of the hormonal treatment. Marín-Ramírez *et al.* (2016) report a decrease in survival in groups fed with diethylstilbestrol for 20 days at concentrations from 100 to 400 mg kg<sup>-1</sup>. However, Varadaraj (1989) in the Mozambique tilapia (*Oreochromis mossambicus*) (Peters, 1852) reported no significant mortality using high concentrations of diethylstilbestrol (100 mg g<sup>-1</sup> to 1000 mg g<sup>-1</sup>). The effect of short-term exposure to different levels of EE<sub>2</sub> on survival and other parameters was investigated by Andersen, Holbech, Gessbo, Norrgren, and Petersen (2003) in the zebrafish (*Danio rerio*) (Hamilton, 1822) and by Van Aerle, Pounds, Hutchinson, Maddix, and Tyler (2002) in the Chinese minnow (*Pimephales promelas*) (Rafinesque, 1820). In both experiments, with the exception of one of all treatments evaluated, no significant decrease in survival or the appearance of malformations was observed using different water concentrations of EE<sub>2</sub> ranking from 10 to 15.4 ng L<sup>-1</sup>. In Nile tilapia, Rougeot, Kanfitine, Prignon, Genotte, and Mélard (2008) report a reduction of survival of embryos reared for five days in two concentrations of EE<sub>2</sub> (100 and 500 µg L<sup>-1</sup>). Shved *et al.* (2009) mention that exposure to EE<sub>2</sub> during early development negatively affects the IGF (insulin-like growth factor) system in Nile tilapia immune organs and could potentially interfere with the antigen presentation capacity of the immune system, thereby altering the susceptibility to infections during growth and reducing survival. In our work, it was observed that in all EE<sub>2</sub>-fed groups there was a lower survival rate than in the control group. However, increasing the concentration of EE<sub>2</sub> did not provoke a decrease in survival as was observed in other experiments using synthetic estrogens (Rougeot *et al.*, 2008; Marín-Ramírez *et al.*, 2016). This increase in mortality could be associated with the Nile tilapia's impaired ability to fight infections, which could explain the higher mortality observed in all the groups fed with EE<sub>2</sub> compared to that of the control group.

GSI indicates the sexual maturity of the fish, which also serves as an indicator of the health and nutritional status (Dadzie & Wangila, 1980; Zeyl, Love, & Higgs, 2014; Marín-Ramírez *et al.*, 2016). Several studies have shown that, based on the observed reduction in the GSI as well as the morphological and histological changes suffered by the gonads, continuous exposure to synthetic compounds will cause a decrease in gonadal development (Linderoth *et al.*, 2006; Marchand, Pieterse, & Barnhoorn, 2008; Louiz, Ben-Attiab, & Ben-Hassinea, 2009).

In the present study, GSI of fish from the control group was lower in comparison to that observed in the groups fed different concentrations of EE<sub>2</sub>. However, significant differences were obtained only when

compared to the concentration of 200 mg Kg<sup>-1</sup>. The fact that the GSI was higher in all EE<sub>2</sub>-fed groups probably indicates that EE<sub>2</sub> at concentrations used had no negative effects at physiological or morphological level on the development of Nile tilapia gonads. GSI values obtained also illustrate the change from males to females in the EE<sub>2</sub>-treated groups. In general, Nile tilapia females have heavier gonads than males. The higher GSI observed could be explained by the fact that in the control group the proportion of females was less than 20%, while in the groups fed EE<sub>2</sub> the proportion was slightly higher than 80%. Production of oocytes, especially large ones like those produced by female Nile tilapia, will result in a GSI higher than that obtained in sperm-producing males. This is important since YY-females produced during feminization with EE<sub>2</sub> are key players in the successful development of YY-males at commercial level. A YY-female with impaired gonadal growth will have a reduced probability of being selected as a breeder.

In the present study, although 100% feminization was not achieved using EE<sub>2</sub> in high concentrations, the proportion of females obtained (in addition to the GSI, growth and survival observed in the groups fed EE<sub>2</sub>) ensures an adequate production of XY- and YY-females which will be used as breeders in the production at commercial levels of YY-male Nile tilapia developed at the Universidad del Papaloapan.

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