Effect of photoperiod on locomotor activity, growth, feed efficiency and gonadal development of Nile tilapia

Galileu Crovatto Veras, Luis David Solis Murgas, Priscila Vieira Rosa, Márcio Gilberto Zangeronimo, Matheus Soares da Silva Ferreira, Jonathan Antonio Solis-De Leon

ABSTRACT - The objective of this study was to evaluate the effect of photoperiod on locomotor activity, growth and gonadal development in Nile tilapia (Oreochromis niloticus) fingerlings. A completely randomised design was used, with five treatments (0L:24D, 6L:18D, 12L:12D, 18L:6D and 24L:0D) and four replicates, with the aquarium as the experimental unit. One hundred and sixty fingerlings of tilapia weighing 3.21±0.05 g and measuring 4.35±0.07 cm each were distributed among 20 aquaria of 20 L in a recirculation system with the temperature controlled to 27 °C, with eight fish per aquarium. Feeding was carried out twice daily for 75 days, with extruded feed containing 40% crude protein. The fingerlings subjected to a photoperiod of 12L:12D as well as those under 18L:6D and 24L:0D showed the greatest locomotor activity, whereas those under 6L:18D and 0L:24D showed the lowest activity. Fish subjected to a photoperiod of 18L:6D and 24L:0D showed the highest levels of performance. However, manipulation of the photoperiod did not influence the gonadal development, survival or the appearance of deformities in juvenile Nile tilapia. Under long photoperiods (18L:6D and 24L:0D), fish direct their energy to somatic growth and induce best feed efficiency.

Key Words: feed intake, fish, locomotor rhythm, Oreochromis niloticus, performance, sexual maturation

Introduction

The influence of environmental factors on fish has been well studied, especially those related to effects on reproduction and growth (Boeuf & Bail, 1999). Among these factors, photoperiod acts as a synchroniser of the endogenous rhythm, influencing locomotor activity, growth, metabolic rate, body pigmentation, sexual maturation and reproduction of teleost fish (Boeuf & Bail, 1999; Biswas et al., 2002; Biswas & Takeuchi, 2002; Trippel & Neil, 2003; El-Sayed & Kawanna, 2004).

Furthermore, photoperiod is one of the most important factors that affects the fish feeding strategy (Reynalte-Tataje et al., 2002) and in most species, feeding occurs in a non-random way, following certain standard biorhythms, i.e., the circadian rhythms that are influenced by the photoperiod. Thus, diurnal fish are most active in daylight and less active during the dark, whereas the reverse is true for nocturnal fish (Boeuf & Bail, 1999).

For some species, long photoperiods might indirectly modify growth by eliciting an increased feed intake, developing muscle mass through increased locomotor activity (Boeuf & Bail, 1999), enhancing nutrient use efficiency (Biswas et al., 2006) and/or redirecting energy from gonadal development into somatic growth (Boeuf & Bail, 1999; Ginés et al., 2004; Rad et al., 2006).

However, the effect of photoperiod on somatic growth and sexual maturation has been little studied during the early stages of fish development (Rad et al., 2006). To date, long photoperiods have been used to stimulate somatic growth and/or delay sexual maturity in some teleost fish such as Atlantic salmon (Salmo salar), Atlantic halibut (Hippoglossus hippoglossus), sea bream (Sparus aurata), sea bass (Dicentrarchus labrax) and Nile tilapia (Oreochromis niloticus) (Boeuf & Bail, 1999; Oppedal et al., 1999; Jonassen et al., 2000; Simensen et al., 2000; Kissing, et al., 2001; Randall et al., 2001; Gines et al., 2004; Rad et al., 2006). However, the manipulation of photoperiod does not always cause benefits to fish performance and survival. In the long term, changes in the light regime might lead to negative effects on the metabolism and development of fish, especially when extreme photoperiods are used (24L:0D and 24D:0L), which differs considerably from conditions in the wild (Villamizar et al., 2011). The Nile tilapia (Oreochromis niloticus) is noted for its rapid growth under intensive culture and for being a rustic as well as one of the most cultivated species in the world. Currently, some knowledge is available concerning the effects of photoperiod on the growth, feeding efficiency, locomotor activity and sexual maturation of tilapia. However, some information is contradictory, which limits conclusions.
The objective of study was to evaluate the locomotor activity, growth and sexual maturation of Nile tilapia subjected to different photoperiods.

**Material and Methods**

The experiment was conducted at the Laboratory of Photoperiod Fish of Sector Physiology and Pharmacology of the Veterinary Medicine Department, at Universidade Federal de Lavras, MG, Brazil, over 75 days.

The design utilized was randomised with five treatments, i.e., different photoperiods (0L:24D; 6L:18D; 12L:12D; 18L:6D and 24L:0D) and four replicates, with the aquarium as the experimental unit. One hundred and sixty fingerlings of Nile tilapia with initial weight and length of 3.21±0.05 g and 4.35±0.07 cm, respectively, were distributed randomly at a stocking density of eight fish per aquarium.

The experiment was conducted in twenty 20 L tanks with a water recirculation system and the temperature was controlled by a thermostat. The daily parameters of water quality measured were: temperature (27.2±0.99 °C) and dissolved oxygen (7.35±0.41 mg/L) using oximeter AT 150 – Alfakit, and pH (6.65±0.05) was measured using pH meter AT 315. The ammonium concentration (0.12 mg/L) was monitored once a week via a card kit ammonia indotest – Alfakit.

Groups of four aquariums were kept in isolation under a controlled lighting system with individual timers and a 20 W fluorescent lamp with a constant intensity of 1200 lx at the water surface. All lamps for the respective photoperiods were lit at 07h00, except the lighting programmes in which fish were kept under 24 h light or dark, where the lamps remained on or off, respectively, throughout the experimental period. According to the methodology of Larson et al. (2004), for illumination of the external environment of the laboratory, a 60 W red lamp was used during feeding, biometrics and cleaning the aquaria.

The fish were fed twice daily at 09h00 and 17h00, with a commercial extruded feed containing 400 g/kg crude protein and pellets 2 mm in diameter. Thirty minutes after the feed was supplied, any leftover was removed, frozen and dried at 55 °C to determine intake. The amount of feed was 5% body weight during the first 15 days and thereafter, and 3% of body weight by the end of the experiment. The weight and length were measured every 15 days to correct the feed supply.

To assess the locomotor activity of fish, the aquaria were equipped with photocells (Omron, mod E3S-AD62, Kyoto, Japan), which were fixed and centralized to the aquarium. The photocells were connected to a channel board (USB-1024HLS, Measurement Computing, Norton, Massachusetts, USA) and connected to a computer. These photocells continuously emitted a beam of infrared light and every interruption caused by a fish was counted and recorded on a data sheet by a specialized computer program (DIO98USB, University of Murcia, Spain) at 10 min intervals. The locomotor activity data were imported into Microsoft Office Excel® 2007 and processed and averaged for each treatment.

At the end of the experiment, the fish were deprived of feed for a period of 24 hours. The animals were then anaesthetised in 2-phenoxyethanol (0.6 mL.L⁻¹), weighed and measured to determine the following performance variables: weight gain (WG, g) = final weight – initial weight; gain in length (GL, cm) = final length – initial length; survival rate (S, %) = (number of dead fish/total fish) × 100; specific growth rate (SGR, % day⁻¹) = [(ln final weight – ln initial weight)/75 days] × 100; protein efficiency ratio (PER) = weight gain/protein intake; daily feed intake (FI, g day⁻¹) = average feed intake/75 days; feed conversion ratio (FCR) = average feed intake/weight gain. Lastly, all fish from each photoperiod were euthanised in 2-phenoxyethanol (1.0 mL.L⁻¹) to measure the weight of the gonads.

The analysis was performed using the statistical program Statistica version 7.0. Data were checked for normality using the Shapiro-Wilk normality test (P<0.05) and the Cochran homogeneity test (P<0.01) and subjected to ANOVA. Differences (P<0.05) between means were compared by the Tukey test at 5% significance. The locomotor activity showed a normal distribution, but the variance of the data was not homogeneous, even after transformation. However, if the size of each sample (i.e., the number of repetitions) are the same, ANOVA is robust with regard to the homogeneity of variances. The survival rate did not show a normal distribution, even after transformation of the data. Thus, for the analysis of this variable, the Kruskal-Wallis test (P<0.05) was applied and in the case of significance, the test of Dunn at 5% significance was used.

**Results and Discussion**

The photoperiod did not affect the survival rate of Nile tilapia fingerlings (Table 1).

Similar to the present study, other studies with tilapia (El-Sayed & Kawanna, 2004) and tambaqui (Colossoma macropomum; Mendonça et al., 2009) have also demonstrated no effect of photoperiod on the survival rate of these fish. Shan et al. (2008) showed a lower survival rate in a 0L:24D photoperiod, with 100% mortality after the seventh day of life, in studies with larvae of miuuy
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croaker, *Miichthys miuy*. According to the authors, this high mortality is attributable to the low ability of these larvae to find food in environments with no light. However, Adewolu et al. (2008) showed that fingerlings of African catfish, *Clarias gariepinus*, had better survival rates when reared under a photoperiod of 0L:24D. The longer survival in this condition can also be attributed to the feeding habit of this species, which feeds comfortably in the dark (Adewolu et al., 2008), dwelling in general, in river bottoms where the incidence of light is low (Feiden et al., 2006). These differences in the survival of different species can be attributed to the extreme variation in the preferred photoperiod, which is species-specific and depends on the stage of development (Britz & Pienaar, 1992; Silva-Garcia, 1996; Boeuf & Bail, 1999; Adewolu et al., 2008).

Increased activity was observed in fish subjected to a photoperiod of 12L:12D, followed by fish exposed to 24L:0D and 18L:6D photoperiods. These photoperiods, in turn, elicited a higher locomotor activity than 6L:18D and 24L:0D and 18L:6D photoperiods. These photoperiods, in turn, elicited a higher locomotor activity than 6L:18D and 24L:0D and 18L:6D photoperiods. These photoperiods, in turn, elicited a higher locomotor activity than 6L:18D and 24L:0D and 18L:6D photoperiods. These photoperiods, in turn, elicited a higher locomotor activity than 6L:18D and 24L:0D and 18L:6D photoperiods.

![Photoperiod effects on locomotor activity](image)

Values followed by different letters are significantly different according to the Tukey test (P<0.05). The coefficient of variation (CV) of activity within each photoperiod was: 0L:24D (CV = 12.00%), 6L:18D (CV = 22.65%), 12L:12D (CV = 5.00%), 18L:6D (CV = 3.00%) and 24L:0D (CV = 12.00%).

Table 1 - Means (±SD) for measurements and performance of Nile tilapia fingerlings subjected to different photoperiods.

<table>
<thead>
<tr>
<th>Variables</th>
<th>Photoperiod</th>
<th>Survival (%)</th>
<th>Feed intake (g day⁻¹)</th>
<th>Feed conversion ratio</th>
<th>Protein efficiency ratio</th>
<th>Final length (cm)</th>
<th>Length gain (cm)</th>
<th>Final weight (g)</th>
<th>Length gain (cm)</th>
<th>Specific growth rate (% day⁻¹)</th>
<th>Gonad weight (g)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0L:24D</td>
<td>87.5±0.21</td>
<td>0.52±0.02b</td>
<td>1.23±0.11ab</td>
<td>2.05±0.17ab</td>
<td>9.87±0.38b</td>
<td>5.42±0.49b</td>
<td>35.12±3.05b</td>
<td>1.84±3.12b</td>
<td>3.03±0.17b</td>
<td>0.15±0.03</td>
</tr>
<tr>
<td></td>
<td>6L:18D</td>
<td>84.6±15.27</td>
<td>0.60±0.02a</td>
<td>1.32±0.06b</td>
<td>1.90±0.08b</td>
<td>10.24±0.2ab</td>
<td>5.95±0.66ab</td>
<td>37.54±8.4ab</td>
<td>3.27±8.1ab</td>
<td>3.15±0.05b</td>
<td>0.16±0.05</td>
</tr>
<tr>
<td></td>
<td>12L:12D</td>
<td>93.7±7.22</td>
<td>0.61±0.3a</td>
<td>1.21±0.07ab</td>
<td>2.08±0.12ab</td>
<td>10.52±3.9ab</td>
<td>6.18±0.46ab</td>
<td>41.33±1.87a</td>
<td>38.16±1.85a</td>
<td>3.32±0.10a</td>
<td>0.15±0.02</td>
</tr>
<tr>
<td></td>
<td>18L:6D</td>
<td>100.0±0.00</td>
<td>0.59±0.01a</td>
<td>1.12±0.03a</td>
<td>2.57±0.07a</td>
<td>10.74±0.29a</td>
<td>6.47±0.36a</td>
<td>41.75±2.80a</td>
<td>38.52±2.70a</td>
<td>3.31±0.12a</td>
<td>0.15±0.02</td>
</tr>
<tr>
<td></td>
<td>24L:0D</td>
<td>96.8±6.25</td>
<td>0.59±0.04a</td>
<td>1.13±0.05a</td>
<td>2.21±0.09a</td>
<td>10.51±4.2ab</td>
<td>6.14±0.43ab</td>
<td>40.86±2.78a</td>
<td>37.70±2.86a</td>
<td>3.31±0.18a</td>
<td>0.16±0.02</td>
</tr>
</tbody>
</table>

Means±SD in the same row with different letters are significantly different according to the Tukey test (P<0.05).
studies with salmon, growth hormones have been shown to be released under continuous light, which is thought to have a positive effect on appetite, which becomes stronger with longer periods of light exposure (Johnson & Björnsson, 1994; McCormick et al., 1995).

Feed conversion and protein efficiency ratio were influenced by the different photoperiods. Fish subjected to 18L:6D and 24L:0D showed the best conversion rates and the protein efficiency ratio did not differ between them. Moreover, fingerlings exposed to 6L:18D showed the worst conversion value and protein efficiency ratio. Fish subjected to 0L:24D and 12L:12D showed a similar value of feed conversion and protein efficiency ratios among them and among fingerlings under a photoperiod of 6L:18D, as well as fish under photoperiods of 18L:6D and 24L:0D (Table 1).

The best feed conversion and protein efficiency ratios found in fish subjected to long photoperiods could be attributed to the time during which the animals were exposed to light after the second feeding. In this case, the fish possibly had the highest appetite, as demonstrated by Biswas et al. (2005, 2006). As described in a review by Boeuf & Bail (1999), Gross et al. (1965) were the first to demonstrate that growth is influenced by photoperiod by not only stimulating consumption, but also by improving the feed conversion ratio. Biswas et al. (2005, 2006) showed that long intervals between feeding fish during a long and constant photoperiod might allow for more efficient digestion, which probably improved nutrient retention. The same effect might have occurred in the tilapia fingerlings exposed to 18L:6D and 24L:0D in this study, since these fish showed better protein efficiency ratios. The long period of light between the first and second feeding (16 h) might mean that these fish have better efficiency of dietary protein use due to improvement in the digestive process.

The final length and gain in length of fish kept under 18L:6D was higher than for those exposed to the other photoperiods. However, the fish under a 0L:24D photoperiod showed the smallest increase in length. The fingerlings under photoperiods of 6L:18D, 12L:12D and 24L:0D showed a final length and gain in length similar to those under a 18L:6D photoperiod (Table 1).

The greatest final weight and weight gains were shown by fish under photoperiods of 18L:6D, 24L:6D and 12L:12D, and were not significantly different between treatments. Fish under the 0L:24D treatment had the lowest weight gain and fingerlings in the 6L:18D photoperiod showed a final weight and weight gain similar to those kept under 12L:12D, 18L:6D and 24L:0D photoperiods (Table 1).

Fish in the 12L:12D, 18L:6D and 24L:0D treatments had the highest growth rates, which did not differ significantly among these photoperiods. However, fingerlings in the 0L:24D and 6L:18D treatments had the lowest specific growth rates (Table 1).

The growth data obtained in this study corroborate those in a study of post-fry Nile tilapia. El-Sayed & Kawanna (2004) showed that post-fry subjected to a long photoperiod (18L:6D and 24L:0D) showed the highest specific growth rate and also showed a trend of increased growth of fingerlings of the same species when they were subjected to photoperiods of 18L:6E and 24L:0D. Bezerra et al. (2008) also showed that Nile tilapia subjected to a long photoperiod (16L:8D and 24L:0D) possessed higher growth and survival rates. Rad et al. (2006) demonstrated that Nile tilapia grew best under a photoperiod of 24 h light when compared with 20L:4D and 18L:6D photoperiods. Long photoperiods also promote higher growth in other species such as juvenile red sea bream (Biswas et al., 2005), striped knifejaw (Oplegnathus fasciatus, Biswas et al., 2008), larvae and juveniles of croaker miuy (Miichthys miuy, Shan et al., 2008), fingerlings of Persian sturgeon (Acipenser persicus, Zolfaghari et al., 2011) and tambaqui (Mendonça et al., 2012), among others.

Long photoperiods might indirectly modify fish growth via the development of muscle mass due to increased locomotor activity (Boeuf & Bail, 1999). In fishes subjected to photoperiods of 12L:12D, 18L:6D and 24L:0D, some energy might be redirected to meet the energy demand due to an increased locomotor rhythm. Increased swimming activity probably stimulated the deposition of amino acids for body protein formation, thus leading to a higher growth of these fish (Biswas et al., 2005). This might be because body proteins are responsible for the majority of growth in terms of weight gain (Biswas et al., 2005). In addition to increased protein deposition, fish exposed to long and continuous photoperiods might still have a low body lipid concentration, indicating that some lipids might have been used to supply the high energy demand of increased activity (Ginés et al., 2004; Biswas et al., 2005).

In some species, the increase in photoperiod and temperature led to rapid increases in GH and IGF-1, which are particularly potent stimulators of muscle growth (Taylor & Migaud, 2009). According to Taylor et al. (2005), juvenile rainbow trout (Oncorhynchus mykiss) subjected to a long photoperiod (18L:6D) demonstrate a direct stimulation in growth due to increased plasma levels of IGF-1 compared with fish subjected to natural photoperiod or a 6L:18D photoperiod.

The manipulation of photoperiod to improve the growth of fish has become increasingly common in the production of several commercial species (Taylor & Migaud, 2009).
However, recently, long photoperiods or constant light have been shown to have a negative effect on the early development of some fish species (Villamizar et al., 2011). According to Villamizar et al. (2009), European sea bass larvae developed fins and teeth faster in constant light (24L:0D), than under 0L:24D or 12L:12D; however, their well-being was compromised, as demonstrated by low bladder inflation 17 days after hatching, as well as by the presence of larvae with malformed jaws. A similar result was shown by larvae of Senegalese sole, Solea senegalensis, kept in constant light (Blanco-Vives et al., 2010). However, in this study, manipulation of the photoperiod showed no influence on malformation of Nile tilapia.

The manipulation of photoperiod had no influence on the gonadal development of Nile tilapia. This result differs from others, in which the manipulation of photoperiod influenced the gonadsomatic index (GSI) of fishes (Boeuf & Bail, 1999; Ginés et al., 2003; Ginés et al., 2004; Rad et al., 2006). However, comparisons of the GSI between fishes of statistically different weight do not represent actual gonadal development. For example, fish of different sizes have a similar weight of gonads. Due to weight differences, heavier fish might have a lower GSI than lighter ones, as the GSI is inversely proportional to the final weight; this was the case when the GSI was determined in the present study. The same appeared to occur in a similar study with Nile tilapia subjected to different photoperiods (Rad et al., 2006), where fingerlings under longer photoperiods showed the highest growth and a lower GSI, compared with fish in a natural photoperiod (12L:12D), which showed the lowest growth.

Conclusions

Long photoperiods (18L:6D and 24L:0D) induce the best growth, feed conversion ratio and protein efficiency ratios in Nile tilapia fingerlings. However, manipulation of the photoperiod does not influence survival, the appearance of body deformations or gonadal development.

References


