The color of illumination affects the stress response of jundiá (*Rhamdia quelen*, Quoy & Gaimard, Heptapteridae)

A cor da iluminação afeta a resposta ao estresse em jundiá (*Rhamdia quelen*, Quoy & Gaimard, 1824, Heptapteridae)

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**ABSTRACT**

This paper provides the first data about physiological responses to stress in jundiá (*Rhamdia quelen*) exposed to different light colours. Jundiá is a species for fish production in the southern part of South America – and suitable for any region with a temperate or subtropical climates. In order to develop a light management for jundiá fingerlings during indoor maintenance and to understand the relationship between welfare and light colour in the jundiá, fingerlings were exposed to white, blue and green light. At the 10th day of light exposure an acute stressor was imposed. One hour after the application of the stressor, fish were sampled. Stress was assessed by means of cortisol determination. Our results show that green light seems to be the worst alternative to illuminate jundiá indoor experimentation facilities, or even fish transportation. The results also suggests that colour affects the stress response of jundiá, and may be usefull for the management of this species.

**Palavras-chave:** espectro de luz, cortisol, *Rhamdia* Silver Catfish.

**INTRODUCTION**

The stress response is the reaction of an organism to a diversity of adverse factors named stressors and comprises a number of physiological processes largely coordinated by the hypothalamus – pituitary – interrenal cells (HPI) axis. Cortisol, the end product of the HPI axis, plays several physiological actions (reviewed by BARTON & IWAMA, 1991 and WENDELAAR BONGA, 1997) and is generally measured in fish blood to assess the stress response.

Jundiá (*Rhamdia quelen*, Quoy & Gaimard, 1824) is a suitable species for fish production in the southern part of South America – and probably for any region with temperate or subtropical climate.

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The study was conducted at the facilities of the University of Passo Fundo (RS, Brazil, 28°15'S / 52°24'W, 687m above sea level). Two hundred and forty 90-day old (8 ± 0.8g), mixed sex jundiá fingerlings were trannieferred to 24 40-l plastic aquaria, distributed in three isolated compartments with black plastic covers in the same experimental room, and filled with aerated fresh water (final fish density of 2g/l). Water was exchanged at a rate of 20% per day, at the same moment that food wastes were removed. During an acclimation period of five days, the fish were kept under natural photoperiod and fed twice a day (10.00 and 16.00 h) with commercial food (42% crude protein, 3,400kcal/kg DE), at 5% of body weight.

Throughout the experiment water temperature averaged 22 ± 1°C, pH ranged from 6.2 to 7.0 and dissolved oxygen ranged from 5.6 to 7.5mg/L. Total ammonia was lower than 0.5mg/L. The total hardness and alkalinity were respectively 60 and 65mg/L CaCO₃. Lights were turned on from 6:00 to 18:00h.

After the acclimation period, the three aquaria compartments were isolated, with fluorescent as the only light source. During ten days, the fish were maintained under 12 hour light: 12 hour dark, with white, blue and green light imposed in each compartment by different colour lamps (AZOO™ Super Light White AZ20013, 30W, 580nm; AZOO™ Coral Blue Light AZ20009, 30W, 436nm and AZOO™ Aquarium Green Light AZ72039, 30W, 480nm). The intensity of light reaching at the surface of each aquarium was 590 Lux measured with a Lux meter.

After 10 days, in four aquaria of each colour compartment, an acute stressor was imposed. The other aquaria were used as control groups. The acute stressor was chasing off the fish with a pen net for 60s (BARCELLOS et al., 1999). One hour after the stresseor, two fish from each aquaria were sampled, 8 in each one of the six groups (stressed and control of each light regime) totalizing 48 fish. This time was established based in our previous results (BARCELLOS et al., 2001).

For blood sampling all fish in all aquaria were anaesthetised with buffered MS222 (Finquel®, 300mg/L), previously dissolved in one liter of water and added to the aquaria to reduce the stress caused by capture. After complete immobilization and loss of orientation, the fish were captured and blood samples (0.1 to 0.30mL) were taken from the severed caudal peduncle with sterile microhematocrit tubes. The time from anaesthesia and blood collection of all fish did not exceed one minute. The tubes were centrifuged (3000g, 10min) in a microhematocrit centrifuge. The plasma was collected with a Hamilton seringe of 50μl, transferred to Eppendorf tubes, and stored at –25°C until required for analysis. Cortisol was measured in duplicate samples of unextracted plasma with a commercially available EIAgén™ CORTISOL test (BioChem ImmunoSystemS). The results were validated with the standard curve of the kit. The intrassay coefficient of variation was 6%.

Results are presented as mean ± S.E.M., and were analysed with an InSItat Sigma statistical package. The differences between cortisol levels of control and stressed fish from each light regime were compared by Student t-test. The cortisol values of all treatment groups and especially of the three control groups and stressed groups were compared by analysis of variance (ANOVA) followed by Tukey’s multiple range test. Statistical significance was accepted for P<0.05.
RESULTS AND DISCUSSION

The mean cortisol levels of control and stressed fish from each light regime are shown in table 1. The analysis of control groups of the three light regimes did not show any differences. Analysis of the differences between the post stress groups of the three light regimes showed that fish kept under white and green light had a statistically higher value than non-stressed controls. When comparing the pre- and post-stress groups in each light regime, only fish exposed to green light showed an elevation on cortisol levels.

Chasing fish with a pen net for 60s increased cortisol levels in fish kept in white and green light. In contrast, no significant effect occurred when fish were maintained under blue light, probably due to slight high control values.

An elevation of plasma cortisol concentration after acute stressors in teleost fish has been described for several species (BARTON & IWAMA, 1991; WENDELAAR BONGA, 1997) including jundiá (BARCELLOS et al., 2001, 2004a).

The un-stressed control values of cortisol in fish exposed to different light regimes are similar to those previously described for jundiá (BARCELLOS et al., 2001, 2004a). Pre-stress values of cortisol ranging from 5 – 50ng/mL are reported for other several species as reviewed by BARRETO & VOLPATO, 2004. In a review by BARTON & IWAMA (1991), few papers about fish of the family Siluridae were cited, and those cited were all of Ictalurus punctactus (Rafinesque), with pre-stress cortisol levels varying from 5 to 51ng/mL and post-stress levels varying from 30 to 309ng/mL. The cortisol peak values (the peak occurs 1h post stressor in jundiá, BARCELLOS et al., 2001) found for fingerlings in the present study varied from 30 to 50ng/mL, and are lower than the ones found for other age categories for jundiá (BARCELLOS et al., 2001, 2003 and 2004a). Several factors may contribute to differences in cortisol levels among species and among specimens from the same species such as age, sex, weight, physiological condition and also the severity and magnitude of the stressor (SUMPTER et al., 1985).

The specific effect of blue light on the stress response was first reported for fish, by VOLPATO & BARRETO (2001) for Nile tilapia. Blue light also changes fish behavior (KAWAMOTO & TAKEDA, 1951; LOUKASHKIN & GRANT, 1959; FANTA, 1995). Effects of the background colour (tanks walls) were assessed by ROTLANT et al. (2003) for Pagrus pagrus, and the authors have postulated that this effect is mediated by the hypothalamus-pituitary-interrenal cells (HPI) axis and the hypothalamus-pituitary-melanophore axis. The present study, similarly to that published by VOLPATO & BARRETO (2001), supports the idea that colour of the light influences the HPI axis.

The results presented herein showed that green light increases the acute stress response caused by handling in jundiá. The green colour of the light was reported as welfare inducer in Brycon cephalus (Günther) (VOLPATO, 2000) and Sardinops caerulea (Girard) (LOUKASHKIN & GRANT, 1959). In contrast, in our experiment the green light evoked a higher cortisol response than the white and blue light. No work relating this type of effects was found.

The activity of the R. quelen is diminished during the day and this species is more active during the night when searching for food (GOMES et al., 2000). PIAIA et al. (1999) reported that jundiá in experimental aquaria shows an aversion to light, seeking dark refuges. The green light effect on the stressor-induced increase in cortisol levels is difficult to explain, but since in natural habitats the environmental colour is a characteristic which varies according to place, and may even change over time in the same habitat, we suggest that the response to green light is associated with the predominant habitat colour. As reviewed by GOMES et al. (2000), the jundiá habitat is the bottom of rivers with low water flow where the fish spend the day in hidden places. In this environment green light may be absent. The effect of light intensity was not considered in our experiment, since the illumination of three colour lamps was the same (590Lux).

A last comment concerns the applicability of light management in aquaculture. Since stressors alter the energy metabolism and the growth in fish (WENDELAAR BONGA, 1997) and can negatively modulate the immune system (BARTON & IWAMA, 1991) as has also been shown in jundiá (BARCELLOS et al., 2004b), practices that can reduce and/or prevent the stress reponse in fingerlings are very important to increase food conversion and reduce fish losses due parasitic and bacterial pathologies. The preference of jundiá for dark environments is well known to jundiá.

Table 1 - Cortisol concentrations (ng/mL) determined 1h after acute stressor exposed fish and control fish, after maintenence by 10 days in white, blue and green light.

<table>
<thead>
<tr>
<th>Group</th>
<th>Ligth color regime</th>
<th>White</th>
<th>Blue</th>
<th>Green</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td></td>
<td>17.3 ± 4.4 aA</td>
<td>23.5 ± 6.4 aA</td>
<td>18.0 ± 7.6 aA</td>
</tr>
<tr>
<td>Post-stress</td>
<td></td>
<td>30.3 ± 6.0 bA</td>
<td>35.3 ± 8.7 aA</td>
<td>50.4 ± 8.4 bB</td>
</tr>
</tbody>
</table>

Different small letters in the columns indicates significative difference (Student’s T test, P<0.05) between control and stressed groups into the specific colour. Different capital letters in lines indicates significative difference between colours (ANOVA followed by Tukey’s multiple range test, P<0.05).
researchers and aquaculturists. However, keeping jundiá fingerlings in completely dark rooms is unfeasible in commercial practices. It is hardly possible to turn the culture facilities completely dark, due to the fact that humans working with these fish need a minimum of light.

CONCLUSION

Our results strongly suggest that colour light regime may be a modulative factor of cortisol response to stressors and that green light seems to be the worst alternative to illuminate jundiá indoor experimentation facilities or even fish transportation. At this stage, more studies, including other colours of light and regimes of illumination, are necessary for the complete elucidation of colour-induced changes. R. quelen stress response and this will be the focus of continuing studies. This study was developed in accordance with national and institutional guidelines for the protection of human beings and animal welfare.

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