

Inter-individual variability in reproductive success and somatic growth in *Cichlasoma dimerus* (Heckel, 1840)

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ABSTRACT. Environmental factors and social interactions are known to affect somatic growth and reproduction in teleost fish. It has been described for *Cichlasoma dimerus* that only one pair is formed under a wide range of laboratory conditions. However, this was not observed in tanks composed of three males and three females, where multiple pair formation occurred. Thus, our objective was to evaluate somatic growth and reproductive performance in *C. dimerus* under this particular condition, in which more than one pair is expected to be formed. A clear sexual growth dimorphism, with males growing faster than females, and multiple pair formation, sometimes simultaneously, were observed. Both features were absent in previous studies with other aquaria structures. Additionally, there was a significant association between reproductive events and body size, where the bigger the fish, both male and female, the higher the number of reproductive events. Despite the sexual growth dimorphism, no differences were observed between males and females in IGF-I and GHR2 mRNA levels. The results obtained for this social species show a high inter-individual variability in the aquaria in regard to reproductive success and growth. This may have implications on experimental design, where a low level of heterogeneity between fish is desirable. If this variability is not taken into account, possible treatment effects may not be detected.

KEY WORDS: Body size, cichlids, growth dimorphism, reproductive events.

INTRODUCTION

In teleost fish somatic growth is affected by environmental factors such as variations in the photoperiod (VERA CRUZ & BROWN, 2009), temperature (GABILLARD et al., 2005), social interactions and food availability (VERA CRUZ & BROWN, 2007, 2009). Somatic growth is defined as a change in size over time. This can be assessed by measuring differences in length and/or weight, implying different physiological processes (BECKMAN, 2011). Therefore, somatic growth is presented in literature by diverse parameters such as difference between final and initial length or body weight, somatic growth rate and condition factor among others.

Growth is regulated by the growth hormone (GH)/ insulin-like growth factor-I (IGF-I) axis (DUAN, 1998; BJÖRNSSON et al., 2002; WOOD et al., 2005; REINECKE, 2010). GH acts directly on certain tissues and indirectly via endocrine or local IGF-I production (ROUSSEAU & DUFOUR, 2007). GH exerts its actions by binding to specific receptors (GHRs). Two GHR types have been described: GHR1, which has been proposed to be the somatolactin receptor, and GHR2, the GH receptor (FUKADA et al., 2004, 2005). The liver is the primary source of circulating IGF-I, but its expression is well documented in other extra-hepatic tissues with a possible local function (REINDL & SHERIDAN, 2012). BECKMAN (2011) discusses the use of IGF-I as a possible indicator

of somatic growth in fish and concludes that before using IGF-I as a growth index in a given situation, the concordance between IGF-I and growth should be tested in that particular situation.

Somatic growth heterogeneity is a characteristic present in many teleost fish (FERNANDES & VOLPATO, 1993; STEFÁNSSON et al., 2000; VERA CRUZ & BROWN, 2007). Social hierarchies, differences in feeding, and gender are factors that could affect somatic growth rates (JOBLING & REINSNES, 1986; RILEY et al., 2002a; MANDIKI et al., 2004; VERA CRUZ & BROWN, 2007; JI et al., 2011). Many vertebrate species establish social hierarchies with dominant and subordinate individuals. Several studies carried out in fishes show that dominant individuals aggressively defend their territory and are reproductively active (ELLIS, 1995; FERNALD & HIRATA, 1977; SAPOLSKY, 2005; FERNALD, 2009; RYDER et al., 2009), with some social species showing a relationship between agonistic interactions and size (ABBOT et al., 1985; BEAUGRAND et al., 1996). Besides, it has been observed that social interactions and feeding hierarchies result in individual growth rate variations (JOBLING & REINSNES, 1986; RILEY et al., 2002a; MANDIKI et al., 2004; VERA CRUZ & BROWN, 2007; JI et al., 2011). In addition, it is well-known that certain fish species exhibit sexual dimorphism in somatic growth. For instance, in some species males grow faster than females, such as in the tilapia *Oreochromis mossambicus* (PETERS, 1852) (RILEY et al., 2002a), and in others females grow faster than males, such as half-smooth tongue sole *Cynoglossus semilaevis* (GÜNTHER, 1873) (JI et al., 2011) or Eurasian perch *Perca fluviatilis* (LINNAEUS, 1758) (MANDIKI et al., 2004). Several studies suggest that gonadal steroid hormones modulate the GH-IGF-I axis in fishes (RILEY et al., 2002b, ARSENAULT et al., 2004; LARSEN et al., 2004; MANDIKI et al., 2004; DAVIS et al., 2007, 2008).

Cichlasoma dimerus (HECKEL, 1840) is a South American cichlid fish that exhibits high growth rates under laboratory conditions. As

many cichlids do, *C. dimerus* displays highly organized breeding activities, as can be observed in the laboratory. The dominant pair will aggressively defend the prospective spawning site (usually a flat stone) and will start to display stereotyped pre-spawning activities (PANDOLFI et al., 2009). In a pair, the male is bigger than the female. The somatic growth rate and the reproductive performance in a time period depend on the condition in which the fish are maintained. For example, in tanks composed of four males and four females only one pair is formed between a female and the biggest male in the aquaria (preliminary results of our laboratory and ALONSO et al., 2011). In these conditions we did not observe differences in somatic growth rate between genders. On the other hand, in tanks composed of three males and three females, multiple pairs are established (previous observation). Thereby, the aim of the present study was to evaluate somatic growth and sexual performance in fish maintained in tanks where more than a pair is expected to be formed.

MATERIALS AND METHODS

Animals

Adult *C. dimerus* were collected from “Esteros del Riachuelo”, Corrientes, Argentina (27°12'50”S, 58°11'50”W), transferred to the laboratory and acclimated in fresh water tanks (400l) under stable conditions of temperature (25 ± 3)°C and photoperiod (14 hours light:10 hours dark). Pairs of *C. dimerus* established in these tanks were withdrawn and maintained together in individual aquaria until their use in the experiment. During this time, fish were fed daily with commercial pellets (Tetra Pond Variety Blend, Tetra). A pair was considered formed after a reproductive event occurred. Suitable actions were followed to minimize pain or discomfort to fish. Principles of laboratory animal care (Guidelines on the care and the use of fishes in research, teaching and testing, Canadian Council on Animal Care, 2005) were adopted. Experiments comply with the approval

of Comisión Institucional para el Cuidado y Uso de Animales de Laboratorio, Facultad de Ciencias Exactas y Naturales, Buenos Aires, Argentina (protocol number 26).

Experimental protocol

The experiment was carried out from November 2010 to May 2014. Each replication ($n = 5$) consisted of three pairs (the ones previously obtained) placed in a 400L aquarium (density = 15 fish/m³) containing gravel, artificial plants and some slabs, under the same temperature and photoperiod conditions as described above. Initial body mass (BM, males: 18.57 ± 1.1 g; females: 14.77 ± 1.08 g) and total length (L_T , males: 9.87 ± 0.19 cm; females: 9.20 ± 0.24 cm) were measured ($P = 0.07$ after MANOVA for differences in initial L_T and BM) immediately preceding the experiment. Animals were kept for two months under these experimental conditions and daily fed with the same commercial pellets. During this period, they were individually identified by their sizes and the particular spot pattern in their fins. Every day during 30 minutes, between 11-12 AM, feeding behaviour, reproductive events and hierarchy status were visually registered. In addition, several times during the course of each experiment, a video recording was performed. Feeding behaviour, rather than food intake, was evaluated as the intention of the fish to catch a pellet. When these fish take a pellet into their mouths, they spit it out many times. Therefore it is not possible to directly quantify the amount of food actually ingested.

In order to evaluate a possible relationship between the number of reproductive events (defined as the egg spawning and the posterior fertilization by the males' sperm release) and initial body size, each fish per tank was classified according to its sex and initial BM or L_T . The smallest fish in a tank was ranked as number 1, the next one as number 2 and the biggest one as number 3. This classification was carried out for males and females separately.

Tissue sample and transcripts quantification

After a two months span, BM and L_T were measured. Fish were anesthetized with benzocaine 0.1%, euthanized by decapitation, and livers collected. As indicators of the nutritional status, hepatosomatic index ($I_H = \text{organ mass/body mass} \times 100$) and condition factor ($CF = \text{BM} \times (L_T)^{-3} \times 100$) were calculated. For evaluating growth in each fish during the two months experiment ΔL_T (final L_T - initial L_T), ΔBM (final BM - initial BM) and specific growth rate (SGR) were calculated. SGR is defined as $\text{Ln}(X_f/X_i) \times t^{-1} \times 1000$, where t is time measured in days and X_f and X_i denote BM or L_T at final and initial moment of the experiment, respectively.

Total liver RNA of 16 fish from three tanks was obtained by TRIzol® (Invitrogen) following manufacturer's instructions, and the corresponding cDNA was synthesized in order to quantify the different transcripts by quantitative real-time polymerase chain reaction (RT-qPCR). RT-qPCRs for IGF-I, GHR2 and acidic ribosomal phosphoprotein P0 (ARP) (reference gene) were performed by using FastStart Universal SyBR green Master (ROCHE) with a mixture of forward and reverse specific primers (Table 1) and 2.5 μl of cDNA template per tube. The RT-qPCR protocol was as follows: 10min of denaturation at 95°C and 40 cycles of 95°C for 15 sec, 58°C for 30 sec and 72°C for 20 sec.

TABLE 1

Primers used for Real Time qPCR.

Name	Sequence (5' to 3')
GHR2 forward	ACTGCTCTCCACTCTCAATTG
GHR2 reverse	AAAGGTGATGGTTCTGGGTC
IGF-I forward	CTCCCAAGATTTCTCGCTCTG
IGF-I reverse	CCCTTCTCCGCTTTACTAACC
ARP forward	ACTGTGGGAGCAGACAATG
ARP reverse	TCCAGTGCAGGATTGTTCTC

Statistical analysis

Differences in mean total length (ΔL_T) and body mass (ΔBM) between genders were evaluated by using a Randomized Block Design (RBD) ANOVA considering the fixed factor gender and the random factor experimental replication (*i.e.*, each tank). In order to discard any influence of the initial body size on growth performance between genders, Pearson's correlations between initial L_T against ΔL_T and BM against ΔBM for each sex were performed. For statistical analysis a single value per sex was used by averaging three fish per tank to avoid pseudo-replication. A Spearman's correlation was run to determine the relationship between reproductive events and body size ranks.

Raw qPCR data were submitted to LinRegPCR software processing in order to obtain initial fluorescence values per sample for subsequent analysis (RAMAKERS *et al.*, 2003; RUIJTER *et al.*, 2009). Pearson's correlations were used to evaluate possible associations between IGF-I mRNA levels and ΔBM , ΔL_T and SGRs. IGF-I and GHR2 mRNA levels between genders were analyzed by using a RBD ANOVA as it was done for morphometric parameters.

Values are expressed as means \pm S.E.M. Statistical tests were considered significant when p-values were less than 0.05 by using InfoStat/L software. Sequential Bonferroni's correction was applied to keep the type I family-wise error rate at 0.05 level.

RESULTS

Morphometric parameters

ΔL_T and ΔBM were significantly higher in males than in females (0.60 ± 0.09 cm *vs.* 0.16 ± 0.07 cm, $P = 0.0034$) and (4.32 ± 0.95 g *vs.* 0.90 ± 0.26 g, $P = 0.0285$), respectively (Fig. 1a,b). For the sizes assessed in this experiment, we did not find any significant correlations between initial body size and ΔL_T or ΔBM (Pearson's correlation, $P > 0.05$). In addition, $SGR(L_T)$ was significantly higher in males than in females (1.36 ± 0.32 *vs.* 0.32 ± 0.15 respectively, $P = 0.0194$). Although ΔBM was higher in males than in females, the differences for the $SGR(BM)$ were not statistically significant (3.49 ± 1.46 *vs.* 0.96 ± 0.31 respectively, $P = 0.1298$). Even though females had a higher I_H , the difference between genders was not significant (males: $1.37 \pm 0.11\%$

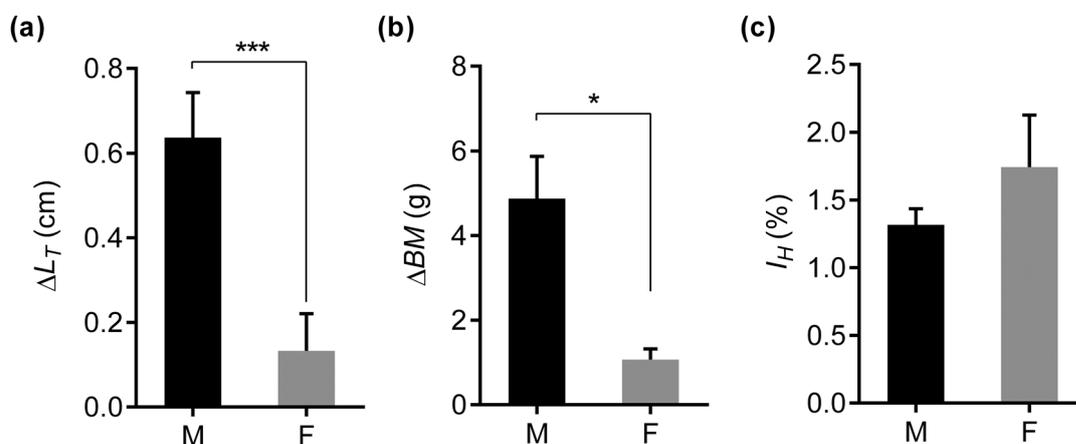


Fig. 1. – Morphometric parameters in three-pair tanks. Vertical bars represent the mean gain in (a) total length (ΔL_T), (b) body mass (ΔBM) and (c) hepatosomatic index (I_H). M: males; F: females. Results are presented as mean \pm SEM. *** $P < 0.001$.

vs females: $1.78 \pm 0.30\%$, $P = 0.3146$) (Fig. 1c) nor was it for CF (males: 1.81 ± 0.08 vs females: 1.87 ± 0.11 , $P = 0.7579$).

Reproductive performance

As expected for this species, we observed aggressive behaviour of the reproductive pair defending the spawning site. This behaviour continued during the reproductive event and the period of parental care.

In all the replications, as was expected from previous observations, more than one pair was observed during the experimental period. Those pairs were not stable, which means that after the reproductive event a new pair was formed, including one or both of these individuals. On some occasions, two pairs were present simultaneously. There was a strong positive monotonic correlation between reproductive events and the initial body size ranking for males (Spearman's correlation, L_T ranking: $r_s = 0.78$, $P = 0.0011$; BM ranking: $r_s = 0.85$, $P = 0.0001$)

and females (Spearman's correlation, L_T ranking: $r_s = 0.69$, $P = 0.0048$; BM ranking: $r_s = 0.69$, $P = 0.0048$). The greater number of reproductive events was associated with the larger fish present in every tank for both genders (Fig. 2). In all the replications, the first pair involved the biggest male. Initial pairs, established before the beginning of the experiment, were not necessarily maintained.

Additionally, video recordings suggested no difference in feeding behaviour among fish of the same tank. Even during the reproductive event and parental care, the pair showed similar feeding behaviour to the other fish in the tank.

Hepatic IGF-I and GHR2 mRNA expression

There was no difference in hepatic IGF-I mRNA levels ($P > 0.05$, Fig. 3a) nor in GHR2 mRNA levels ($P > 0.05$, Fig. 3b) between genders. No correlations between hepatic IGF-I mRNA levels and ΔBM , ΔL_T and SGR were observed (Pearson's correlation, $P > 0.05$).

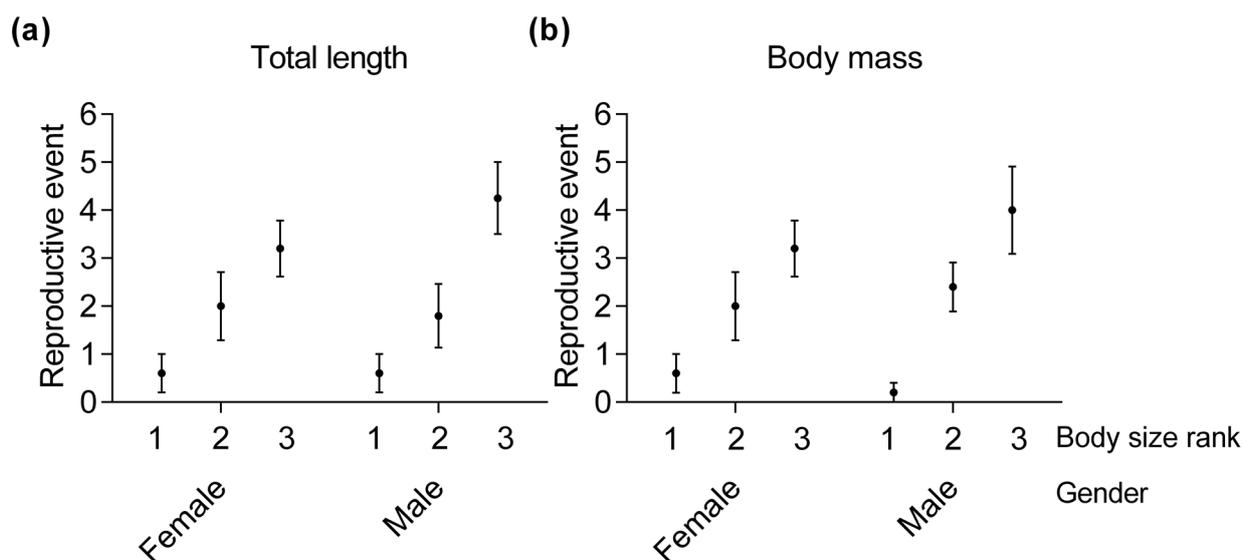


Fig. 2. – Relationship between reproductive events and the body size ranks. For both genders a greater number of reproductive events was associated with the larger fish present in every tank. Spearman's correlation was significant for total length (a) and body mass (b) both in males and females. Fish were ordered according to body size: 1: small; 2: intermediate; 3: big. Results are presented as mean \pm SEM

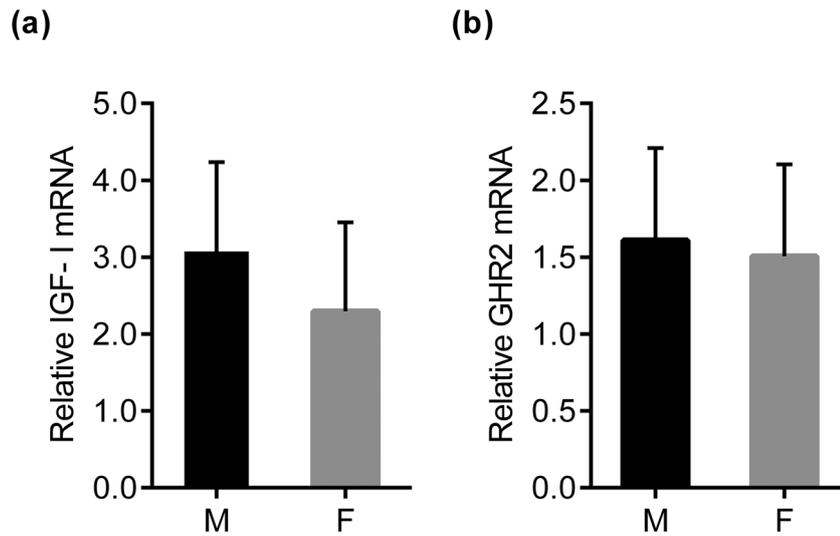


Fig. 3. – Hepatic IGF-I (a) and GHR2 (b) mRNA levels in males and females from three-pair tanks. Data were normalized using ARP as a reference gene and presented as a mean \pm SEM. No differences were observed in IGF-I and GHR2 mRNA levels between genders after an RBD ANOVA test. M: males; F: females.

DISCUSSION

Heterogeneous somatic growth is a common characteristic in fish populations (FERNANDES & VOLPATO, 1993; STEFÁNSSON et al., 2000; VERA CRUZ et al., 2007). For instance, differences in somatic growth between dominants and subordinates (KOEBELE, 1985; VERA CRUZ & BROWN, 2007) or between genders (RILEY et al., 2002a; DAVIS et al., 2008) have been described for many cichlid fish species. In this study, sexual growth dimorphism was clearly demonstrated by evaluating the difference between sexes in body mass and total length gain after two months of experiment. This comparison showed that males grew more than females. However, as Δ BM and Δ L_T can be influenced by their values at the beginning of the experiment, we also compared their SGR values. These results are in line with those obtained with Δ BM and Δ L_T, showing that males display higher specific growth rates, statistically significant at least regarding length, compared to females. Additionally, we demonstrated that for the fish sizes used in this study, there was no relationship between

the initial body size and Δ BM or Δ L_T for each sex, which therefore excludes the possibility of any influence of the initial body size on growth performance. Taken together, these results give strong evidence for sexual growth dimorphism in *C. dimerus*.

A role of steroids in the promotion of somatic growth has been studied in several species. In fishes in which males grow more than females, administration of androgens promotes growth and an increment in IGF-I (RILEY et al., 2002a, b) whereas estradiol (E2) decreases it. In contrast, in those fishes in which females grow more than males, E2 would promote growth (MALISON et al., 1985; MALISON et al., 1988; HAYWARD et al., 2001) and also an increase in IGF-I (GOETZ et al., 2009). Additionally, in Eurasian perch *P. fluviatilis*, androgens produced a decrease in growth (MANDIKI et al., 2004). In the present study fish were in different reproductive states at the end of the experiment; gonadal steroids were therefore not measured due to the high variability expected. The question is, why was a

clear sexual growth dimorphism observed in our experimental design? One possibility is that the sexual growth dimorphism was due to differences in food intake between males and females. However, this scenario seems to be unlikely as no apparent differences in feeding behaviour were observed. Moreover, the nutritional status parameters, hepatosomatic index and condition factor, were not different between genders. In our experiments fish were fed daily and food availability was never restricted, which should exclude the possibility of inequality in food availability. Another possibility could be a differential feed efficiency conversion between males and females and/or a bigger requirement of metabolic energy for reproduction in females, which would impact negatively on growth rate (MANDIKI et al., 2004). Specific experiments would be necessary to elucidate if the sexual growth dimorphism is due to differences in food intake and/or feed efficiency conversion.

Additionally, reproductive performance also depends on the tank conditions. Multiple pair formation was observed when three pairs were placed in the aquaria, a feature absent in four-pair tanks (preliminary results of our laboratory and ALONSO et al., 2011). The first pair formed always included the biggest male. Interestingly, in all tanks a strong correlation between reproductive events and body size was observed, where the bigger the fish, both male and female, the higher the number of reproductive events. To our knowledge this is the first report describing this association. In other reports, a positive correlation was observed between body size and fecundity success, but in those cases fecundity success was inferred as testis mass (PUJORAL et al., 2012), number of eggs per ovary (HOSSAIN et al., 2012) and sperm content (WATANABE et al., 2008). While gonadal parameters are widely used as indicators for fecundity, this does not necessarily imply new offspring because it depends on other events such as pair establishment and quality of the gametes among

others. Unlike those approaches, our results show strong evidence on this issue.

In view of the somatic growth dimorphism observed, we investigated if there were changes in somatic growth axis that could explain this morphometric result. Unfortunately we could not measure GH expression in all the pituitaries due to technical issues during RNA extraction. In order to test if in *C. dimerus* a positive relationship exists between hepatic IGF-I mRNA and somatic growth, these parameters were compared in females and males. No correlation was observed between growth and IGF-I levels. Our results are in accordance with those obtained in chinook salmon *Oncorhynchus tshawytscha* (WALBAUM, 1792) and barramundi *Lates calcarifer* (BLOCH, 1790) (SILVERSTEIN et al., 1998; NANKERVIS et al., 2000). As summarized by Beckman (2011, p. 236-237), there are different relationships between IGF-I levels and somatic growth. In the present work hepatic IGF-I mRNA was measured at the end of the experiment *i.e.* at a specific time point. Unfortunately, it was impossible to measure IGF-I plasma levels due to the small plasma sample obtained. A possible scenario would have been to measure plasma IGF-I during the experiment (longitudinal design) but this kind of experimental design would have impacted negatively on the reproductive behaviour. Also, another component of the somatic growth axis, GHR2 was measured. No differences were observed in hepatic GHR2 mRNA levels, in agreement with no variations in IGF-I between genders. This result differs from that obtained in the Mozambique tilapia *O. mossambicus* (DAVIS et al., 2008) where males, which present greater somatic growth rates than females, showed higher level expression for both GHR1 and GHR2 mRNA receptors in the liver. Although in *C. dimerus* there were no differences at the expression level, possible differences at the protein level cannot be ruled out. More studies including measurements of plasma IGF-I and analysis of hepatic GHR2 content are necessary to check this out.

CONCLUSIONS

In summary, this work shows clear sexual growth dimorphism in fish held in tanks containing three males and three females, and a significant association between reproductive events and body size. These results have implications in the design of laboratory experiments because it is a common practice to sample fish from a tank irrespective of social interactions that could be affecting critical parameters such as somatic growth or reproductive performance.

ACKNOWLEDGEMENTS

This work was supported by Agencia Nacional de Promoción Científica y Tecnológica (grant number: PICT 2008-2005. P.V.) and CONICET (grant number: PIP: 0276. P.V.).

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Received: 19 June 2014

Accepted: 15 September 2014

Branch editor: Roger Huybrechts