LIFE-HISTORY TRAITS OF CHLOROSCOMBRUS CHRYSURUS (ACTINOPTERYGII: PERCIFORMES: CARANGIDAE) IN TROPICAL WATERS OF THE ATLANTIC OCEAN

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Background. In fishery biology, information on life-history traits is extremely useful for species conservation and the monitoring, management and sustaining of fish stocks. Nevertheless, detailed biological information is very limited for tropical fishes, especially for those with low economic value, such as the Atlantic bumper, Chloroscombrus chrysurus (Linnaeus, 1766). This species is commonly captured by artisanal fisheries in tropical waters and the lack of information makes difficult the development of strategies for the proper management of the species. Therefore, the presently reported study intended to provide new data on the biological parameters of C. chrysurus to fill the gap in the existing knowledge.

Materials and methods. Life-history traits of the Atlantic bumper, C. chrysurus, were estimated from 335 fish collected in a western Atlantic region during fishery surveys carried out between 2010 and 2012. Age and growth were determined by annuli counts of sagittal otoliths. Growth parameters were calculated by adjusting the back-calculated length-per-age to the von Bertalanffy growth model. Empirical equations and data on maturity were used to estimate the size and age at first maturity, as well as the optimum size and age. Theoretical longevity and natural mortality were also estimated for the species.

Results. Six age classes were identified and asymptotic length ($L_\infty$), growth rate ($k$), and the theoretical age when the specimen was at zero length ($t_0$) for the whole population assumed the following values: $L_\infty = 25.49$ cm, $k = 0.32$ year$^{-1}$, and $t_0 = 0.058$ years. No evidence of sexual size dimorphism was found, with no significant differences in growth characteristics and in the length frequency distribution between sexes. The principal vital parameters were as follows: size ($L_{50} = 15.5$ cm) and age ($T_{50} = 2.9+$) at sexual maturity, optimum length ($L_{opt} = 13.08$ cm), and optimum age ($T_{opt} = 2.1+$), as well as its theoretical longevity ($A_{0.95} = 9.17$) and natural mortality ($M = 0.92$).

Conclusion. The determined size and age at sexual maturity, optimum length and age, theoretical longevity, and natural mortality indicate that the studied population of C. chrysurus is overexploited. The Atlantic bumper is vulnerable to fishing pressure and should be managed with caution.

Keywords: age determination, fish growth, longevity, natural mortality, otolith

INTRODUCTION

Global changes have impacted ecosystems in many different ways by changing food web structures (Kolar and Lodge 2000), and even allowing species to extend their distribution range throughout the globe (Júarez et al. 2008, Tavera et al. 2008). A good example, though not the only one, is the occurrence of the Atlantic bumper, Chloroscombrus chrysurus (Linnaeus, 1766), in Spanish waters, Gulf of Cádiz (Acosta et al. 2009), and in the Mediterranean Sea (Peña Rivas et al. 2013). The extension of the distribution range of this species may be linked to many factors, but especially to the current warming of the North Atlantic (Stebbing et al. 2002). In fact, some authors have predicted that warming conditions are likely to increase chances of survival and future naturalization of C. chrysurus in these regions (Acosta et al. 2009, Peña Rivas et al. 2013), which may result in several ecological and economic impacts (e.g., reducing and impacting native wildlife populations).
Faced with this prediction, it is extremely concerning that available information on biological parameters of *Chloroscombrus chrysurus* is still very scarce, especially for tropical regions where the species is widely distributed (Cunha et al. 2000, Campos et al. 2010). For instance, studies on growth parameters of *C. chrysurus*, up to now, have only been carried out with larvae in subtropical regions (Leffler and Shaw 1992, Sánchez-Ramárez and Flores-Coto 1998), and with data based on length frequency (García and Duarte 2006, Sossoukpe et al. 2017). Analyses of structures that can provide more consistent results, such as otoliths, have not been incorporated into these studies so far.

The correct estimation of life-history traits of this species is not only important for its conservation, but also for the development of management strategies, especially with the current extension of its distribution range (Peña Rivas et al. 2013). Therefore, the presently reported study aims to contribute with current knowledge on the biology of *C. chrysurus* by estimating its life-history traits in tropical waters using a combination of otolith analysis and empirical equations to provide enough information on this species’ age, growth, population structure, maturity, longevity, and natural mortality.

**MATERIAL AND METHODS**

Fish were sampled in a western Atlantic region at the central coast of Alagoas, Brazil (centroid at 9°67′40″S and 35°72′31″W, Fig. 1), during fishery surveys carried out between May 2010 and April 2012. Samples were collected monthly using two different fishing gears: a beach seine net (mesh size: 140 to 254 mm) used to catch juveniles (total length: 2–14 cm) and a gillnet (25 to 170 mm) to collect adults (15–25 cm). In the laboratory, fish individuals were identified to species level following Menezes and Figueiredo (1980), measured to the nearest 1 cm (total length), weighted (in grams), sexed, and assigned to a maturity stage using macroscopic gonadal examination following Vazzoler (1996).

The sagittae otoliths of each individual were removed, cleaned, dried, and stored in labelled vials for subsequent examination. Prior to analysis, to accomplish a better visualization of growth rings, otoliths were immersed in pure alcohol for 24 h until decalcification. Subsequently, a picture of otoliths submerged in pure alcohol and sulcus placed downwards was taken using a binocular stereoscope microscope Leica S8-APO (25–40 times magnification) with reflected light against a dark background. Age determination was achieved by counting growth rings, which were considered to be formed by one opaque band and one translucent band together (Fig. 2). Each otolith was read independently at least twice for two different readers with no prior information regarding length or sex. Later, differences in age estimation between readers were tested by a Student’s *t*-test.

The periodicity of annulus formation was verified by the relative marginal increment analysis (MIA)

$$MIA = \frac{R_t - R_{nc}}{R_{nc} - R_{nc-1}}$$

where *R*<sub>t</sub> is the total radius of the otolith; *R*<sub>nc</sub>, the distance between the nucleus and last ring; and *R*<sub>nc-1</sub>, the distance between the nucleus and next-to-last ring formed (Branstetter and Musick 1994). Analysis of variance (ANOVA) was carried out to test significant differences in the MIA throughout the year, and monthly MIA values were compared to rainfall data [mm] by a linear regression to determine trends in bands formation.

To estimate the growth parameters of *Chloroscombrus chrysurus*, we first used the back-calculation model of Morita and Matsuishi (2001) to remove bias due to age effects (Wilson et al. 2009). This model assumes that otolith radius results from a linear combination of both fish length and age (Wilson et al. 2009) and it is calculated as follows

**Fig. 1.** Map of the study area, located in the coast of Alagoas, Brazil

**Fig. 2.** Otolith of *Chloroscombrus chrysurus* caught in the tropical Atlantic Ocean; arrows indicate growth rings
Biological parameters of *Chloroscombrus chrysurus*

\[ L_t = \frac{\alpha}{\beta} + \left( \frac{L_\infty}{\beta} \right) \left( 1 - e^{-\frac{t}{\beta}} \right) \]

where \( L_t \) is the back-calculated fish body length at age \( t \); \( L_\infty \) is the fish body length at time of capture; \( \alpha \), \( \beta \), and \( \gamma \) are the constants obtained from the multiple linear regression analysis. Growth in length was then modelled by the von Bertalanffy growth model (VBGM) using a nonlinear regression

\[ L_t = L_\infty \left( 1 - e^{-k(t-t_0)} \right) \]

where \( L_t \) is the length at age \( t \); \( L_\infty \) is the theoretical asymptotic total length; \( k \) is the growth coefficient describing the rate of growth \([\text{year}^{-1}]\) towards \( L_\infty \), and \( t_0 \) is the hypothetical age (in years) at a total length of zero. The VBGM was estimated for males, females and the whole population (both sexes combined and specimens with indeterminate sex).

Length at first maturity (\( L_{m0} \)) for the whole population was estimated by fitting a logistic model using the percentage of mature individuals per 1 cm size class (King 2007)

\[ Y = \frac{1}{1 + e^{-(a + bX)}} \]

where, \( Y \) is the percentage of mature individuals at a length \( X \), and \( a \) and \( b \) are the resulted constants. The model was fitted using a log-transformed abundance of individuals per size class and the size where 50% of individuals were mature was assessed by \(-a/b\). Age at sexual maturity (\( t_{m0} \)) was known by using the inverse version of the growth equation as suggested by King (2007)

\[ t_{50} = t_0 - \left( \frac{1}{k} \right) \ln \left( \frac{1-L_{50}}{L_\infty} \right) \]

Estimated growth parameters were also used to calculate natural mortality (\( M \)), length at maximum yield per recruit (\( L_{opt} \)), optimum age (\( T_{opt} \)), and longevity, defined as the time that individuals take to achieve 95% of the asymptotic length (\( A_{0.95} \)). Natural mortality (\( M \)) was calculated by Pauly’s empirical equation

\[ \log M = -0.0066 - 0.279 \cdot \log L_\infty + 0.6543 \cdot \log k + 0.4634 \cdot \log \text{Temp} \]

where \( L_\infty \) and \( k \) are the von Bertalanffy growth parameters, and Temp is the mean water temperature (°C) (Pauly 1980). Data on annual sea surface temperature was retrieved from the Brazilian National Institute of Meteorology (INMET) and the mean value of 29.87°C was incorporated into the equation.

The length at maximum yield per recruit (\( L_{opt} \)), which expresses the length at which natural mortality equals to growth rate was calculated following Beverton (1992)

\[ L_{opt} = 3L_\infty \left( 3 + Mk^{-1} \right)^{-1} \]

where \( M \) is the natural mortality and \( L_\infty \) and \( k \) are the estimated growth parameters from the VBGM. Optimum age (\( T_{opt} \)), the age where fish reaches its highest biomass, was estimated by the equation of Krishnan Kutty and Qasim (1968)

\[ T_{opt} = t_0 + \frac{\ln(3k + M) - \ln M}{k} \]

Theoretical longevity, here defined as the time that individuals take to achieve 95% of the asymptotic length, was calculated using the equation described by Taylor (1958)

\[ A_{0.95} = t_0 - \frac{\ln(1-L_\infty)}{k} \]

where, \( L_\infty \), \( k \), and \( t_0 \) are the estimated growth parameters. Maximum and minimum values (standard errors) of \( t_0 \) and \( k \) (growth parameters) were used to estimate longevity confidence intervals.

We also examined sexual size dimorphism in species total length by analysis of variance (ANOVA). Data was previously tested for normality and homoscedasticity using Shapiro–Wilk and Levene’s tests, respectively. Additionally, length–weight relations (LWR) for females, males, and the whole population were estimated by linear regressions using the equation

\[ \log W = \log a + b \cdot \log TL \]

where \( W \) is the body weight, \( TL \) is the total length, \( a \) is the intercept, and \( b \) is the slope (Le Cren 1951, Froese 2006).

Statistical analyses were all carried out with the software R statistics (version 3.1.3) at a significance level of \( P < 0.05 \).

**RESULTS**

In this study, a total of 335 fish were included in the estimation of population parameters of *Chloroscombrus chrysurus*. Out of all individuals, 133 were females, 100 were males, and in 102 sex could not be determined. Total length ranged from 2.5 to 26.4 cm (considering all specimens, including the individuals of undetermined sex), and no evidence of sexual size dimorphism was found (ANOVA, \( P > 0.05 \), Fig. 3). LWR parameters and related statistics are given in Table 1. All regressions were highly significant \((P < 0.01)\) with correlation coefficients \((r^2)\) of 0.76, 0.80, and 0.98 to males, females, and the whole population, respectively.
No suitable samples were collected in June for calculating the MIA. Yet, marginal increments were significantly different among months (ANOVA, $P < 0.01$), with increments presenting a trend of increasing monthly until reaching its peak in May, followed by a decrease in their value starting in July (Fig. 4). Furthermore, changes in marginal increment were significantly correlated to seasonal fluctuations in rainfall rates ($P < 0.01$, $r^2 = 0.44$).

Both readers’ age estimates were symmetrical ($P < 0.05$). Estimated ages for Atlantic bumper ranged from 0’ to 6’, though the majority of fish sampled belonged to 3’ and 4’ age classes (60.89%). Observed growth parameters did not differ between sexes (Table 2), hence the von Bertalanffy growth curve was recalculated using all specimens, including the individuals with indeterminate sex (Fig. 5, $r^2 = 0.88$, $n = 335$). Information on growth parameters, theoretical longevity ($A_{90}$) and natural mortality ($M$) for {	extit{C. chrysurus}} are summarized in Tables 2 and 3.

The size at first maturity ($L_{50}$) for {	extit{C. chrysurus}} was 15.5 ± 2.8 cm (Fig. 6), and the age at sexual maturity ($T_{50}$) was 2.9’ years$^{-1}$. Optimum length and age were smaller than the $L_{50}$ and $T_{50}$, being 13.08 cm and 2.1’ years$^{-1}$, respectively.

**DISCUSSION**

This study represents the first comprehensive reference on age and growth estimation for {	extit{Chloroscombrus chrysurus}} based on otolith analyses. The use of otoliths in age estimation for tropical fish was previously considered infeasible due to complex interactions between physiological and environmental factors (Blaber 2002), as well as miscounts of growth rings (Brothers et al. 1976).

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**Fig. 3.** Length frequency distribution of females (A) and males (B) of {	extit{Chloroscombrus chrysurus}} in the tropical Atlantic Ocean

**Fig. 4.** Annual variation in the marginal increment analysis (MIA) of {	extit{Chloroscombrus chrysurus}} and the relation between monthly values and rainfall levels; number of otoliths used to get mean values are presented above error bars

**Fig. 5.** Relation between observed total length and age for {	extit{Chloroscombrus chrysurus}} in the tropical Atlantic Ocean ($n = 335$), showing von Bertalanffy growth function

**Fig. 6.** Size at first maturity for {	extit{Chloroscombrus chrysurus}} in the tropical Atlantic Ocean
Biological parameters of *Chloroscombrus chrysurus* (Shaw, 1803)  

| Parameter | Value          | 95% CI        |  
|-----------|----------------|---------------|
| L∞        | 26.438 ± 1.218 | 95% CI of 0.005–0.037 |  
| L∞ ± 0.046 | 0.95          | 95% CI of 2.84–2.88 | 0.98 |

\( L∞ \) is the time that individuals take to achieve 95% of the theoretical asymptotic total length, \( k \) is the growth rate, \( t_0 \) is the hypothetical age at a total length of zero.

**Table 1** Descriptive statistics and estimated parameters of length–weight relations for *Chloroscombrus chrysurus* caught in tropical waters of the Atlantic Ocean between May 2010 and March 2011, and from August 2011 through April 2012

| Sex     | \( n \) | \( L_{\text{a}} [\text{cm}] \) | \( k [\text{year}^{-1}] \) | \( t_0 \) | \( r^2 \) |
|---------|--------|-------------------------------|-----------------------------|----------|---------|
| Female  | 133    | 25.619 ± 0.805                | 0.323                       | 0.032 ± 0.045 |  
| Male    | 100    | 26.438 ± 1.218                | 0.316                       | 0.057 ± 0.064 |  
| Whole population | 355 | 25.494 ± 0.640                | 0.328                       | 0.058 ± 0.035 |  

\( n \) = number of individuals, \( L_{\text{a}} \) = theoretical asymptotic total length, \( k \) = growth rate, \( t_0 \) = hypothetical age at a total length of zero.

**Table 2** Estimated theoretical longevity (\( A_{95} \)) and natural mortality (\( M \)) for *Chloroscombrus chrysurus* caught in tropical waters of the Atlantic Ocean between May 2010 and March 2011, and from August 2011 through April 2012

| Sex               | \( A_{95} \) [\text{year}] | 95% CI | \( M \) [\text{year}^{-1}] | 95% CI |
|-------------------|-----------------------------|--------|-----------------------------|--------|
| Females           | 9.30 ± 0.50                 | 0.926 ± 0.030 |                     |        |
| Males             | 9.53 ± 0.76                 | 0.905 ± 0.046 |                     |        |
| Whole population  | 9.17 ± 0.44                 | 0.938 ± 0.027 |                     |        |

\( A_{95} \) is the time that individuals take to achieve 95% of the asymptotic length.

**Table 3** LWR parameters for *Chloroscombrus chrysurus* caught in tropical waters of the Atlantic Ocean between May 2010 and March 2011, and from August 2011 through April 2012

| Sex     | \( a \) | 95% CI of \( a \) | \( b \) | 95% CI of \( b \) | \( r^2 \) |
|---------|--------|-----------------|--------|-----------------|---------|
| Female  | 0.004  | 0.002–0.011     | 3.19   | 2.92–3.46       | 0.80    |
| Male    | 0.014  | 0.005–0.037     | 2.81   | 2.50–3.12       | 0.76    |
| Population | 0.013 | 0.012–0.015     | 2.84   | 2.80–2.88       | 0.98    |

Nevertheless, the results presented herein, such as the symmetrical counting of growth rings between different readers and the marginal increment analysis, show that these structures can provide consistent information on growth characteristics for *C. chrysurus*.

Marginal increment analysis (MIA) indicated a clear trend in bands formation (see Fig. 4), demonstrating that growth rings of *Chloroscombrus chrysurus* are formed once a year and are strongly correlated to seasonal changes in rainfall rates. Increment bands may be associated with both environmental conditions and cyclical biological process (Morales-Nin and Panfili 2005), depending on life-history requirements of species (Sousa et al. 2015). For many tropical species, a strong relation between rainfall and growth has been previously reported (Bwanika et al. 2007, Ong et al. 2015, Sousa et al. 2015, Efifre et al. 2016). This relation results, mainly, from changes caused by rainfall in water physicochemical parameters (i.e., turbidity, salinity, and light penetration) and productivity levels (Barletta et al. 2005, Pereira et al. 2015), as these changes may provide better feeding condition for fish (Bergenius et al. 2005).

The von Bertalanffy growth curve fitted to our observed data adequately. Growth parameters estimated herein (\( L_{\text{a}} = 25.4 \text{ cm}, k = 0.32 \text{ year}^{-1}, t_0 = 0.05 \)) differed from those calculated for species using length-frequency data in south-eastern and southern Brazil (\( L_{\text{a}} = 45.1 \text{ cm}, k = 0.22 \text{ year}^{-1}, t_0 = -0.65 \)) by Cergole et al. (2006), and in Caribbean Sea (\( L_{\text{a}} = 30.5 \text{ cm}, k = 0.63 \text{ year}^{-1} \)) by García and Duarte (2006). These differences may be associated with many factors (i.e., climate type, latitudinal differences) (Tarkan and Vilizzi 2015), but especially to the size range of individuals once growth parameters are very sensitive to samples—larger individuals tend to increase asymptotic length as growth rate decreases (Espino Barr et al. 2008). For example, in south-eastern/southern Brazil, larger individuals of *C. chrysurus* have been reported, with species growing up to 42 cm (Cergole et al. 2006), while in the studied region the maximum length registered for the species was 30 cm (Lessa and de Nóbrega 2000). Furthermore, it is important to be noticed that methodological differences in age determination may unduly influence parameters estimation and, consequently, provide false impressions of the growth potential of fish species.

The lack of significant differences in growth characteristics between sexes found for Atlantic bumper appears to be very common among Carangidae species in tropical regions, such as *Trachinotus botla* (Shaw, 1803) (see Parker and Booth 2015) and *Trachurus picturatus* (Bowdich, 1825) (see Garcia et al. 2015). Although many factors may regulate sexual size dimorphism in species, mechanisms underlying these differences between sexes remain poorly understood (Fairbairn 2005, Young 2005). For example, even though studies have shown that environmental conditions may affect sexes differently, it appears that these changes are more consistent among larger taxa (Estlander et al. 2017). For smaller species, such as *C. chrysurus*, sexual size dimorphism is more understated and studies in larger scales (encompassing...
different latitudes) are necessary to a better understanding on patterns of body growth and size variation.

Both, length ($L_{\text{opt}}$) and age ($T_{\text{opt}}$) at maximum yield per recruit for *Chloroscombrus chrysurus* were smaller than the length ($L_{\text{opt}}$) and age ($T_{\text{opt}}$) at first maturity, which is a typical reproductive strategy of small fishes (Beverton 1992). As small fishes have high predation risk, and once maximum possible yield is determined by the ratio between growth and mortality, smaller species tend to reach maximum yield during younger ages and smaller sizes in a trade-off of reproduction and survival (Jensen 1996, Froese and Binohlan 2000).

Estimated natural mortality ($M = 0.938$ year$^{-1}$) was slightly smaller than the one reported for the species in the Caribbean Sea ($M = 1.29$ year$^{-1}$) (Garcia and Duarte 2006). Yet, this value was expressively high when compared to other tropical carangid species, such as *Trachurus declivis* (Jenyns, 1841) ($M = 0.63$ year$^{-1}$) (Stevens and Hausfeld 2005). While similarities in mortality rates are expected among species in the same family (Pascual and Iribarne 1982), adaptations to local conditions and exploitation levels may cause these rates to vary. In the particular case of *C. chrysurus*, though not being considered a commercial target species, this species is frequently caught by artisanal fisheries in tropical regions, especially as by-catch (Ambrose et al. 2005, Alves et al. 2012, Cunha 2015). Indeed, our estimated theoretical longevity suggests that the Atlantic bumper may be vulnerable to fishing pressure, showing a high longevity ($A_{\text{opt}} = 9.17$ years) despite its relatively small body size, which is a typical feature of overexploited species (Wiedmann et al. 2014). In summary, *Chloroscombrus chrysurus* was found to be a small, fast growing, and long-lived fish. Such characteristics indicate that the species may be undergoing high levels of exploitation, and, therefore, should be managed with caution. We suggest that further studies of population structure of *C. chrysurus* incorporate geographic, climate, and fishery variables and hope that information provided herein may be used for developing management strategies for this species, as well as its conservation.

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