REGULAR PAPER





Life history of the river goby *Glossogobius callidus* (Teleostei: Gobiidae)

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Funding information

DSI/NRF Centre of Excellence for Invasion Biology; National Research Foundation - South African Research Chairs Initiative of the Department of Science and Innovation, Grant/ Award Number: 110507; National Research Foundation, Grant/Award Numbers: 101039, 109015, 103581, 88746; Water Research Commission, Grant/Award Number: #K5/2039

Abstract

The river goby *Glossogobius callidus* is native to freshwater and estuarine habitats in South Africa. Individuals [21.1–144.4 mm total length ($L_{\rm T}$)] were sampled from impoundments in the Sundays River Valley, Eastern Cape, from February 2014 to March 2015. The largest female was 137.2 mm $L_{\rm T}$, and the largest male was 144.4 mm $L_{\rm T}$. Length-at-50% maturity was 75.2 ± 2.1 mm $L_{\rm T}$ for males and 76.2 ± 2.0 mm $L_{\rm T}$ for females. Absolute fecundity was 1028.2 ± 131.7 oocytes per fish, and relative fecundity was 50.1 ± 18.1 oocytes per gram. The spawning season extended from October to December. Fish were aged using sectioned sagittal otoliths. The growth zone periodicity was validated using edge analysis. Longevity was more than 7 years for females and more than 6 years for males. Length-at-age was similar for the two sexes and was best described using the von Bertalanffy growth model as $L_{\rm t} = 74.7(1 - {\rm e}^{-1.0({\rm t} + 0.1)})$ mm $L_{\rm T}$ for the entire population. Using the population age structure, the mortality rate was estimated at 1.3 per year.

KEYWORDS

Age and growth, fresh water, Glossogobius callidus, length-at-50% maturity, life history

1 | INTRODUCTION

The family Gobiidae is one of the largest vertebrate groups and contains more than 2000 species that inhabit freshwater, estuarine and marine habitats (Thacker, 2003). The group has not been well studied from a life-history perspective, which constrains the understanding of their general ecology and hinders trait-based comparisons with other taxa (e.g., Winemiller et al., 2015). Nonetheless, this group of fishes has recently received increased attention, as several species have invaded rivers and lakes in Australia, Europe and North America

(Gertzen et al., 2016; MacInnis & Corkum, 2000). Globally, the main attributes for invasion successes appear to be their tolerance for a wide range of environmental conditions, a broad diet and aggressive behaviour (Charlebois et al., 2001; Hempel et al., 2019; Kotta et al., 2016).

Although only native gobies are known to occur in southern African freshwater ecosystems, the river goby *Glossogobius callidus* (Smith 1937) has been documented to rapidly colonise artificial impoundments (Woodford *et al.*, 2013). This species occurs as two reproductively isolated populations along the subtropical and temperate

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African east coast from northern Mozambique to the Cape region of South Africa (Maake *et al.*, 2013). In the temperate Eastern Cape region of South Africa, *G. callidus* is particularly abundant in both estuarine and freshwater environments, where it often dominates fish communities (*e.g.*, Wasserman *et al.*, 2011; Woodford *et al.*, 2013).

Current knowledge on the biology and ecology of the river goby is scant. What is known is that this moderately small goby is an invertivorous (Mofu et al., 2019; Wasserman, 2012), speleophil hole-nester that practices parental care with the male guarding and maintaining the egg mass until the larvae hatch (Wasserman et al., 2015). Sneaking, a spawning strategy whereby smaller males that are capable of reproduction but are less able to defend a territory or nest spawn in the nest of a guarding male when the opportunity arises, has been reported for some members of the family but not for G. callidus (see Immler et al., 2004; MacInnis & Corkum, 2000; Takegaki et al., 2012). There is, however, currently no information on other life-history traits of this species, including gonadal recrudescence, maturity, fecundity, growth and mortality. The objective of the current study was, therefore, to provide the first comprehensive description of the life history of a temperate population of G. callidus to compare its life history to that of other freshwater gobies.

2 | MATERIALS AND METHODS

This project was conducted with permission from the National Research Foundation – South African Institute for Aquatic Biodiversity Animal Ethics Committee (NRF-SAIAB reference number 2011/02), and specimens were collected with permission obtained from the Eastern Cape Department of Economic Development and Environmental Affairs (DEDEA permit number CRO 18-11CR and CRO 19-11CR).

Fish were randomly sampled monthly between August 2013 and March 2015, from 10 irrigation ponds located in the Sundays River Valley, on the eastern extremity of the Cape Fold Ecoregion (Abell *et al.*, 2008) in South Africa (Figure 1) (see Supporting Information S1 for geographical coordinates). The climate in the region is warm temperate, with water temperatures ranging from a minimum of 10° C in winter to a maximum of 31° C in summer (Woodford *et al.*, 2013). Fish were sampled using a 30×2 m seine net with 12 mm mesh (stretched) in the wings and an 8 mm stretched mesh codend. Seine hauls, typically two or three per pond, were conducted until a minimum of 100 *G. callidus* were collected. The fish were euthanised with an overdose (40 mg l⁻¹) of clove oil, and specimens were either kept on ice (if required for otolith extraction for ageing) or preserved in a 10% formalin solution (for reproductive assessment).

In the laboratory, all fish were then measured for total length L_T and standard length L_S to the nearest millimetre and total mass (M_T) to the nearest 0.1 g. Fish were subsequently dissected, and gonads were first visually assessed under a dissecting microscope to determine sex, before being categorised into one of the five macroscopic stages using the standard macroscopic criteria and terminology described by Brown-Peterson et al. (2011). The gonads were subsequently removed, patted dry and weighed (M_G nearest 0.1 g), and the eviscerated body mass of the fish was recorded (M_E nearest 0.1 g). To determine absolute fecundity (FA), all mature, yolked and unovulated oocytes in the ovaries of a minimum of two spawning-capable females were used per site, and these were selected randomly across ponds and counted under a stereomicroscope. Relative fecundity (F_R) was then calculated as a proportion of M_T . Length-at-50% maturity (L_{M50}) was calculated by fitting a logistic function to the proportion of mature fish (macroscopically staged as developing, spawning capable and regressing) collected during the reproductive season that were grouped into 5 mm L_{T} size classes.

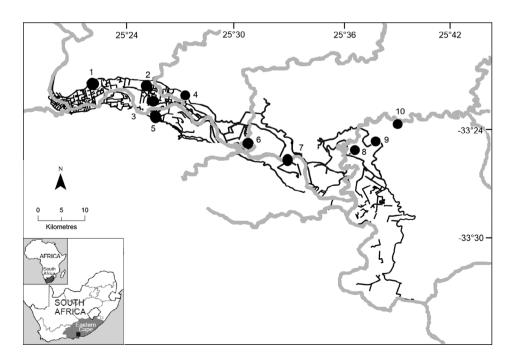


FIGURE 1 Sundays River Valley irrigation ponds in the Eastern Cape, South Africa, Pond 1; Pond 2; Pond 3; Pond 4; Pond 5; Pond 6; Pond 7; Pond 8; Pond 9; Pond 10 (a) Surveyed ponds, (—) Sundays River and tributaries, (1a) Sundays irrigation network (see Supporting Information S1 for geographical coordinated and pond names)

For ageing, a pilot study (*n* = 240 fish) demonstrated that unburnt sectioned sagittal otoliths provide the most appropriate structure for ageing as 95% of scales and 83% of burnt otolith sections were unreadable. Otoliths were set in clear polyester casting resin, sectioned transversely through the nucleus using a double-bladed diamond saw at a thickness of 0.4 mm and mounted onto microscope slides using di-*N*-butyle phthalate in xylene microscopy mountant (Merck Private Ltd., Darmstadt, Germany) (Figure 2). The sections



FIGURE 2 Unburnt otolith of a 86.1 mm L_T river goby Glossogobius callidus, showing five annuli (marked by black circles), sampled from the Sundays River Valley irrigation ponds in the Eastern Cape of South Africa

were then examined under a binocular microscope using transmitted light at variable magnifications ($\times 10$ –40) to count pairs of opaque and hyaline growth zones and to assess the appearance of the otolith margin. Precision was assessed by the average percent error (APE) and Coefficient of Variation (CV) calculated from the relative ages assigned by three separate readers as described by Beamish and Fournier (1981) and Chang (1982). The growth zone deposition rate was validated as annual using edge analysis (EA) whereby the optical appearance of the edge on the otoliths was noted as Bernoulli variables (1 = opaque zone present and 0 = absent) and plotted against time. Under the assumption that the frequency of edge zones follows a yearly sinusoidal cycle when plotted against time (Campana, 2001), a periodic logistic regression analysis was used to test a unimodal null hypothesis and bimodal alternate hypothesis (Beamish *et al.*, 2005).

Growth was estimated using a sample of 153 fish collected between November 2014 and January 2015 that were sampled from across the 10 freshwater ponds (see Figure 1). This sampling period was selected to coincide with the period when, as indicated by the validation experiment, opaque zones become optically discernible on the edge of the otoliths of the highest proportion of fish. As this sampling period was also consistent with the peak of the spawning season, the length of the newly hatched fish was measured during the spawning experiment of Wasserman *et al.* (2015). These fish averaged 21.2 mm L_T and were included as age 0 fish for modelling age. The length-at-age was subsequently modelled using the von Bertalanffy growth (VBG) function, which is described as $L_t = L_{\infty} (1 - e^{-K(t-t_0)})$, where L_t is the length at time t, L_{∞} is the theoretical asymptotic length, K is the Brody growth coefficient and t_0 is the age of a zero-

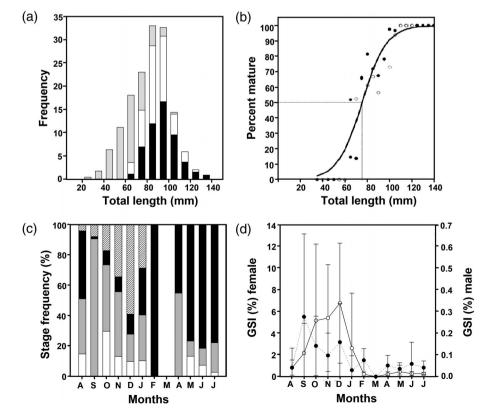
FIGURE 3 Monthly variation in (a) length-frequency distributions for (males, (

) females, (

) and juveniles; (b) logistic ogive fitted to the percentage of sexually mature male (n = 13, black circles)and female (n = 15, white circle) river goby Glossogobius callidus; red line represents L_{M50} (total length at 50% maturity) (●) male, (a) female; (c) monthly maturity stages of female (n = 431) river goby G. callidus (

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Resting, (Regenerating, (ZZZZ) Spawning capable; (d) mean (±S.D.) gonado-somatic index (GSI) for male (n = 595, white circles) and female (n = 659, black circles) river goby G. callidus, sampled from the irrigation ponds located in the Sundays River Valley, Eastern Cape, between August 2013 and March 2015 (—○—) Female, (———) Male



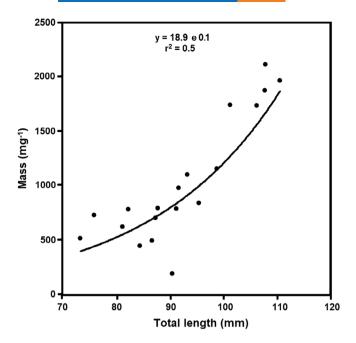


FIGURE 4 Total length (L_T)-mass (mg⁻¹) relationship for female (n = 19) river goby *Glossogobius callidus*, sampled from the Sundays River Valley irrigation ponds in the Eastern Cape, South Africa

length fish. The variance estimates were calculated using parametric bootstrap resampling with 1000 replicates drawn from a normal distribution (Efron, 1982), and standard errors and confidence intervals were constructed using the percentile method of Buckland (1984). Differences in growth parameter estimates between sexes were assessed using a likelihood ratio test (Cerrato, 1990).

The instantaneous rate of annual mortality (*Z*) was estimated by catch curve analysis (Ricker, 1975) whereby length–frequency data were converted to age–frequency data by means of an age–length key developed from the aged individuals. The annual mortality (*Z*) was estimated by the negative slope of the linear regression through the descending data points of the natural logarithm of frequency against age. As these populations are not exploited, *Z* was assumed to approximate natural mortality *M*.

3 | RESULTS

In total 1945 fish (579 female, 667 male, 699 juvenile) ranging from 21 to 144 mm $L_{\rm T}$ were examined (Figure 3a). The sex ratio did not differ from unity (χ^2 test, χ^2 = 0.02, df = 1, P > 0.05). The length-at-50% maturity was estimated at 75.1 ± 2.1 mm $L_{\rm T}$ for males and 76.2 ± 2.0 mm $L_{\rm T}$ for females. Length-at-50% maturity did not differ significantly between sexes and was calculated at 75.2 ± 2.1 mm $L_{\rm T}$ for the combined sexes (Figure 3b). Gonad development for fish exceeding $L_{\rm M50}$ followed a distinct seasonal pattern with only spawning-capable female fish recorded in the population from September (early spring) to January (mid-summer) (Figure 3c). The female gonado-somatic index followed a similar pattern observed with

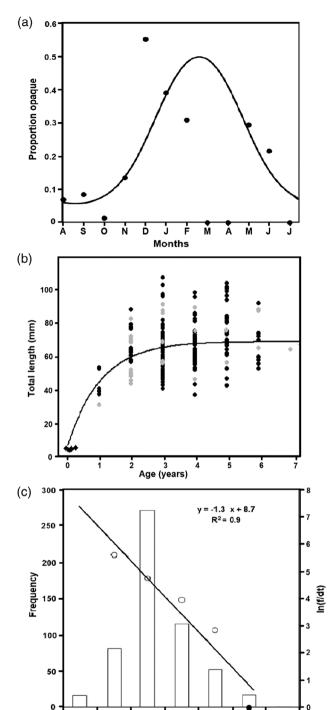


FIGURE 5 (a) Proportion of otoliths (n = 623) with an opaque zone on the margin, with a solid line representing the predicted unimodal annual cycle of opaque zone deposition using a full model periodic regression; (b) combined von Bertalanffy growth curve of total length-at-50% maturity ($L_{\rm M50}$), juvenile (n = 6, black diamond), male (n = 84, black circles), female (n = 69, grey circles) (\odot) Female, (\odot) Male, (\odot) Juvenile; (c) age–frequency histogram (n = 560) and resultant catch curves generated from length frequencies, and the instantaneous rate of mortality (Z) is the slope of the descending limb of the catch curve (closed circles). All graphs are for the river goby *Glossogobius callidus*, sampled from the Sundays River Valley irrigation ponds in the Eastern Cape, South Africa

Age (years)

	L_{∞} (mm)	S.E.	K (year ⁻¹)	S.E.	t ₀ (year)	S.E.
Entire population	74.7	1.7	1.0	0.1	-0.1	0.1

TABLE 2 Life-history comparison of the river goby Glossogobius callidus, with other goby species using von Bertalanffy growth parameters

Species	Region	Sex	L _∞ (mm)	K (year ⁻¹)	t _o (year)	Φ′(mm)	A _{max} (year)	A _{mat} (year)	A _{max} /	Mortality rate (year ⁻¹)	Main reference
Glossogobius callidus	South Africa	Р	74.7	1.0	-0.1	0.7	7.0	2.0	3.5	1.3	This study
Neogobius melanostomus	United States	М	215	0.4	-0.8	0.9	1	2	0.5	1.0	MacInnis and Corkum (2000)
	United States	F	237	0.2	-1.6	1.7	4	3	0.8	1.3	Apanasenko (1973) and Skora et al. (1999)
Gobius niger	Turkey	F	148	0.3	-1.5	1.9	9	2	4.5	0.7	Filiz and Toğulga (2009)
		М	167	0.3	-2.2	1.9	9	2	4.5	0.7	Filiz and Toğulga (2009)
Aphia minuta	Italy	М	740	1.4	-0.2	1.9	5	4	1.3	2.5	La Mesa (1999)
	Italy	F	660	1.6	-0.2	1.9	5	4	1.3	3.0	La Mesa (1999)
Neogobius fluviatilis	Ukraine	U	138	0.7	-0.3	2.1	4	1	4	1.3	Bilko (1971)
Pomatoschistus Iozanoi	Europe	U	860	0.9	-0.2	1.8	3	1	3	1.5	Froese and Binohlan (2003)
Gobius paganellus	Isle of Man	U	116	0.4	-0.2	1.7	3	1	3	1.6	Pauly (1980)

Note: A_{mat} : age at maturity; A_{max} : maximum age; F: female; K: Brody growth coefficient; L_{∞} : asymptotic length; M: male; P: entire population surveyed from all sites used in this study; Φ' (phi prime: Pauly & Munro, 1984): growth performance; t_0 : age at zero length; U: unknown; Z: mortality rate (see Materials and Methods).

macroscopic staging with an increase in October and December, whereas the male gonado-somatic index showed a marked increase in November (Figure 3d). Fecundity ranged from 100 to 2000 oocytes per fish. Oocyte diameter ranged from 107.1 to 299.1 μ m with a mean (±s.D.) of 207.2 ± 7.9 μ m. Absolute fecundity (F) is dependent on L_T whereby $F = 18.9 L_T$ (mm)^{0.1} (Figure 4).

For 1424 otoliths from the examined specimens of *G. callidus*, APE was 2.9% and CV was 3.9%. The periodic logistic regression analysis failed to reject the null hypothesis that one opaque zone was deposited annually ($\chi^2 = 0.4$, df = 3, P > 0.05) but rejected the alternative hypothesis that growth zone deposition was bimodal ($\chi^2 = 25.7$, df = 3, P < 0.05). Thus, the growth zone deposition rate was validated as annual (Figure 5a), and subsequent analyses were based on the direct interpretation that one opaque/hyaline growth zone pair is equivalent to 1 year of age. Power analysis was completed using G*Power version 3.1.9.4, and the sample size required for $\alpha = 0.05$ and a power of 95% was 111.

The oldest female fish sampled was a more than 7 year old, 69 mm L_T female, and the oldest male was a more than 6 year old, 126 mm L_T fish. VBG models fitted to male and female data did not differ significantly between sexes (log-likelihood ratio test, χ^2 = 301.1, df = 1, P > 0.05), and the best fit for the entire population is L_t = 74.7

 $(1 - e^{-1.0(t + 0.1)})$ mm L_T (Table 1; Figure 5b). The estimated total mortality Z is 1.3 per year (Figure 5c). As this species is not exploited, Z was taken to equal natural mortality M.

4 | DISCUSSION

Based on the findings of this study, *G. callidus* is best categorised as an opportunistic life-history strategist *sensu* Winemiller *et al.* (2015). The species is relatively fast growing, reaching adulthood at 2 years. Maturity is attained during the second year of life at lengths approximating 50% of maximum L_T . The species has a short spawning season from spring (October) to early summer (December). These life-history traits facilitate its prevalence in a wide range of habitats and explain why this species is able to rapidly colonise new environments (Woodford *et al.*, 2013).

The life-history of *G. callidus* is not unusual for the Gobiidae (Table 2). The males mature slightly earlier, and at smaller sizes, than females, a trait often documented in this fish family (Kovačić, 2007; Sokołowska & Fey, 2011). To compare the growth performance, the parameter Φ' takes into consideration the interaction and dependence between the VBG parameters (Pauly & Munro,

1984). The growth performance of G. callidus was lower than that of round goby Neogobius melanostomus (Pallas 1814), black goby Gobius niger (Linnaeus 1758), monkey goby Neogobius fluviatilis (Pallas 1814) and Lozano's goby, Pomatoschistus lozanoi (de Buen 1923) (Apanasenko, 1973; Bilko, 1971; Filiz & Toğulga, 2009; Froese & Binohlan, 2003; MacInnis & Corkum, 2000). In addition to slower growth, G. callidus is relatively long lived (7 years) when compared to gobies such as N. melanostomus which generally live for about 4 years (Grula et al., 2012), similar to that reported for Gobius vittatus where males attained ages of 7 years (Kovačić, 2006) but shorter lived than G. niger from Turkey (9 years) (Filiz & Toğulga, 2009). Overall, G. callidus had higher mortality rates compared with G. niger from Turkey (Filiz & Toğulga, 2009), and female Neogobius caspius (Eichwald 1831), however, had low mortality rates than male N. caspius (Aghajanpour et al., 2016) from the Caspian Sea and Aphia minuta (Risso 1810) from Italy (La Mesa, 1999). Furthermore, N. melanostomus had higher mortality rates than G. callidus (Kasapoğlu, 2016).

Although fecundity, maturity and spawning frequency are important components of understanding the life history, these need to be coupled with information regarding age and growth to allow for the interpretation of these data on the context of rates. Relatively fast growth and early maturity allow juveniles to reach maturity rapidly, a trait linked to successful colonisers of novel habitats (Corkum et al., 2004). This is similar to the traits exhibited by invasive N. melanostomus (Corkum et al., 2004). MacInnis and Corkum (2000) show that successful invaders are characterised by a short generation time, rapid growth and early maturity. Where present, these gobies have generally dominated benthic communities by highly competitive feeding strategies. A speleophil reproductive strategy, high fecundity (Corkum et al., 2004), rapid maturity (Kornis et al., 2017) and a generalist feeding strategy are traits linked to its success as an invader in the Great Lakes. Similarly, the ability of G. callidus to rapidly colonise the irrigation impoundments in the Sundays River system (Woodford et al., 2013) is most likely facilitated by its generalist diet (Mofu et al., 2019), wide environmental tolerance and similar reproductive strategy to invasive gobies. Such traits may thus represent indicators for the establishment of success in freshwater gobies introduced into novel environments.

ACKNOWLEDGEMENTS

This study was jointly funded by the Water Research Commission research grant (WRC Project number: K5/2039), the DSI/NRF Centre of Excellence for Invasion Biology (CIB) and the National Research Foundation – South African Research Chairs Initiative of the Department of Science and Innovation (Inland Fisheries and Freshwater Ecology, grant number: 110507) and NRF incentive funding (grant numbers: 109015 to O.L.F.W., 103581 to D.J.W. and 88746 to R.J.W.). We acknowledge the use of infrastructure and equipment provided by the SAIAB Research and Collections Platforms and the funding channelled through the NRF-SAIAB Institutional Support system. We extend our gratitude to P. Ndaleni for fieldwork assistance and E. Wolhuter and F. Lamont for logistical support. Fieldwork permits to sample fishes in the Sundays River system were issued by the

Eastern Cape Department of Economic Development and Environmental Affairs (DEDEA).

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SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section.

How to cite this article: Mofu L, Woodford DJ, Wasserman RJ, Weyl OLF. Life history of the river goby

Glossogobius callidus (Teleostei: Gobiidae). J Fish Biol. 2020;

1-7. https://doi.org/10.1111/jfb.14478