



# Age and growth of the blue shark (*Prionace glauca*) in the Indian Ocean

I. Andrade<sup>a,b,\*</sup>, D. Rosa<sup>a</sup>, R. Muñoz-Lechuga<sup>a</sup>, R. Coelho<sup>a,c</sup>

<sup>a</sup> Portuguese Institute for the Ocean and Atmosphere (IPMA), Avenida 5 de Outubro s/n, 8700-305, Olhão, Portugal

<sup>b</sup> University of Algarve, Campus de Gambelas, 8005-139, Faro, Portugal

<sup>c</sup> Centre of Marine Sciences (CCMAR), University of Algarve, Campus de Gambelas, 8005-139, Faro, Portugal

## ARTICLE INFO

Handled by George A. Rose

### Keywords:

Bycatch  
Elasmobranchs  
Fisheries  
Growth modelling  
Pelagic longline

## ABSTRACT

Since there is still a lack of biological information regarding *Prionace glauca* in the Indian Ocean, specifically in terms of age estimation and growth modelling, the age and growth of this species was studied by analysing vertebral samples. All samples were collected from specimens captured by pelagic longliners between March 2013 and September 2016, with sizes ranging from 82 to 301 cm fork length ( $L_F$ ). Two growth models were fitted to the age data, a three-parameter von Bertalanffy growth function (VBGF) re-parameterized to calculate  $L_0$  (size at birth) and a two-parameter VBGF with a fixed  $L_0$ . The latter was considered the most adequate to describe the growth of the species, with the estimated parameters being  $L_\infty = 283.8$  cm  $L_F$ ,  $k = 0.13$  year<sup>-1</sup> for males and  $L_\infty = 290.6$  cm  $L_F$ ,  $k = 0.12$  year<sup>-1</sup> for females. These results suggest that females have a slower growth than males. The maximum age estimated was 25 years, representing the oldest attributed age to this species so far. Further work is needed regarding *P. glauca* in the Indian Ocean, but this study adds important life-history information that can contribute for the management and conservation of the species.

## 1. Introduction

The blue shark *Prionace glauca* (L. 1758) is the only species belonging to the genus *Prionace* which belongs to the Carcharhinidae Family (Order Carcharhiniformes). *Prionace glauca* is a pelagic and oceanic species with a worldwide distribution, including both temperate and tropical waters (Compagno, 1984). In addition, it is considered by many authors as the most abundant of pelagic sharks (Compagno, 1984; McKenzie and Tibbo, 1964; Nakano and Seki, 2003; Nakano and Stevens, 2008). Therefore, these apex predators are a highly important component of pelagic ecosystems globally (IOTC, 2007). Regarding the distribution of *P. glauca* in the Indian Ocean, results of a recent study by Coelho et al. (2018) suggest that larger individuals are found in equatorial and tropical parts of this ocean, while smaller specimens seem to prefer higher latitudes. Sharks belonging to this species can be longer than 300 cm in total length ( $L_T$ ) (Pratt, 1979) and have been suggested to reach as much as 380 cm  $L_T$  (Compagno, 1984). They are placental viviparous sharks with a gestation period going from 9 to 12 months after which females give birth to 4 to 135 pups per litter (Castro and Mejuto, 1995; Compagno, 1984; IOTC, 2007; Nakano, 1994; Pratt, 1979). The young are born in the Spring and Summer (Compagno, 1984; Nakano, 1994; Pratt, 1979). Both sexes attain sexual maturity at a similar body length and age, the latter being between 4 and 6 years

for males and from 5 to 7 years for females (Cailliet et al., 1983; Lessa et al., 2004; Nakano, 1994; Pratt, 1979; Vas, 1990). Also regarding age, the longevity of *P. glauca* is thought to be of about 20 to 23 years (Cailliet et al., 1983; Manning and Francis, 2005; Stevens, 2009).

Despite being considered the most abundant of pelagic sharks (McKenzie and Tibbo, 1964; Nakano and Seki, 2003; Nakano and Stevens, 2008), *P. glauca* still faces some threats that could compromise their current populations. Specimens of this species are the most frequently caught pelagic sharks as bycatch by fisheries worldwide, in particular by pelagic longline fisheries targeting tuna and swordfish (Anderson, 1980; Bailey et al., 1996; Campana et al., 2009; Carruthers et al., 2011; Francis et al., 2001; IOTC, 2016; Pratt, 1979; Stevens, 1992). When it comes to sports fishing, *P. glauca* is one of the preferred species of sharks by who practices this activity, being one of the main targets (Anderson, 1980; Casey and Hoey, 1985; Compagno, 1984; Skomal and Natanson, 2003; Stevens, 1984, 2009; Vas, 1990). In terms of commercial fisheries, *P. glauca* was rarely a targeted species in the past (Nakano and Stevens, 2008; Stevens, 2009). However, there has been an increasing commercial interest on this species in the recent years, both as a food source and also for its fins (Aires-da-Silva et al., 2008; Dent and Clarke, 2015; Eriksson and Clarke, 2015). In the Indian Ocean, *P. glauca* is the most caught species of shark by Portuguese pelagic longlines, and it is the second most caught species following the

\* Corresponding author at: Portuguese Institute for the Ocean and Atmosphere (IPMA), Avenida 5 de Outubro s/n, 8700-305, Olhão, Portugal.

E-mail address: [inesfcandrade@gmail.com](mailto:inesfcandrade@gmail.com) (I. Andrade).

<https://doi.org/10.1016/j.fishres.2018.11.019>

Received 2 March 2018; Received in revised form 12 November 2018; Accepted 15 November 2018

Available online 26 November 2018

0165-7836/ © 2018 Elsevier B.V. All rights reserved.

main target (Muñoz-Lechuga et al., 2016). *P. glauca* is considered globally as “Near Threatened” by the International Union for Conservation of Nature (IUCN) Red List (Stevens, 2009). Regionally, it has been considered “Near Threatened” in the Northeast Atlantic (Sims et al., 2015) and “Critically Endangered” in the Mediterranean (Sims et al., 2016).

The inter-governmental organization responsible for the management of *P. glauca* in the Indian Ocean is the *Indian Ocean Tuna Commission (IOTC)*. In 2015, such organization carried out the first stock assessment for this species in the Indian Ocean, however the condition of the stock remained uncertain due to the conflicting results obtained. When it comes to stock assessment, age and growth of organisms are very important parameters to estimate growth rates, mortality rates, longevity, and other relevant aspects to evaluate the condition of stocks (Campana, 2001, 2014; Goldman et al., 2012). Since *P. glauca* is very common in pelagic ecosystems, the biology of this species has been well studied over the years, including age and growth studies. However, of the studies performed to date, most have focused on the populations of the Atlantic and Pacific Oceans with only two studies for the Indian Ocean (Jolly et al., 2013; Rabehagasoa et al., 2014).

Considering the lack of biological information about *P. glauca* in the Indian Ocean, specifically regarding age and growth, the present work aims to 1) estimate the age of *P. glauca* individuals through the reading of growth bands in vertebrae; 2) obtain growth models for both sexes in the South Indian Ocean; and finally 3) provide age and growth data of this species to IOTC for stock assessment purposes and management advice.

## 2. Materials and methods

### 2.1. Sampling

All the samples used in this study were collected by scientific fishery observers from the *Instituto Português do Mar e da Atmosfera (IPMA)* on board of Portuguese commercial longline vessels that target swordfish (*Xiphius gladius*) in the Indian Ocean. A total of 818 vertebrae were collected from March 2013 to September 2016. Vertebral samples were collected in the South Indian Ocean between 23.75°S and 34.85°S (latitude) and from 40.70°E to 92.97°E (longitude) (Fig. 1).

While on board of the fishing vessels, the sex of all individuals as well as the fork length ( $L_F$ ) were recorded. Fork lengths were measured in a straight line to the nearest lower cm. Vertebrae were removed from the region below the anterior part of the first dorsal fin of each individual. All vertebral samples were kept frozen from extraction until they were cleaned and then preserved.

### 2.2. Sample processing

All vertebrae were first cleaned and then sectioned. The cleaning process started by manually removing all the organic tissues around each vertebra using scalpels and tweezers. After that, they were immersed in a solution of 4–6% sodium hypochlorite (commercial bleach) during approximately 5–10 minutes (depending on the size of each vertebra) to remove any remaining soft tissues, and finally placed in water for a few minutes to eliminate all the sodium hypochlorite. Once cleaned, all vertebrae were stored in ethanol at 70% until further use.

To prepare the vertebrae for the sectioning process, they were first air-dried from the storing ethanol during approximately 30 min and then mounted on microscope slides, using a synthetic polymer glue (Pattex; Henkel, Düsseldorf, Germany). They were left during 24 h for the glue to air dry completely. Once the glue was fully dried, each slide was placed in a sectioning cutter, a Buehler Isomet 1000 precision low-speed saw, with two diamond wafering blades, to produce 0.5 mm longitudinal sections. The sections were cut through the centre of the vertebrae to reveal the *corpus calcareum* (bow-tie) and banding structure used to estimate age.

To enhance the band pattern, the sections obtained were stained with crystal violet solution (Sigma-Aldrich Co., St. Louis, MO), previously used in other shark ageing studies (e.g., Coelho et al., 2011; Fernandez-Carvalho et al., 2011, 2015; Rosa et al., 2017), during 2 (for small sections) or 3 min (for bigger sections) and then turned upside down and left for another 2 or 3 min to guarantee that the staining is even on both sides. Only one of the two sections obtained from each vertebra was stained, to later compare the visibility of the stained versus the non stained bow-ties of the vertebrae. After staining, both sections of each sample were covered with paper and tightly wrapped between two microscope slides, in order to maintain the original shape once fully dried. They remained wrapped for 24 h until dry.

The completed sections of each vertebra were mounted onto glass microscope slides using Neo-Mount and observed for growth band structure under a Nikon dissecting microscope with a mounted high resolution digital camera, using transmitted white light. Photographs of each observed sample were recorded and then digitally enhanced using the ImageJ software (Schindelin et al., 2015) by adjusting the contrast and brightness. The same software was used to mark the growth bands, as well as the focus and the outer edge of the *corpus calcareum* of each vertebral sample (Fig. 2a–b).

### 2.3. Age estimation and precision analysis

Age was estimated by counting the number of band pairs visible along the *corpus calcareum* in each sample using the enhanced photographs, with a band pair consisting of one wide band and one narrow band. Annual deposition of growth bands was assumed and the first distinct band (usually associated to an angle change in the outer edge of the *corpus calcareum*) was considered to be the birthmark (e.g., Francis and Ó Maolagáin, 2016; Hsu et al., 2015; Jolly et al., 2013; Rabehagasoa et al., 2014).

To develop the age reading protocol a reference set of 50 specimens (25 male and 25 female) representing the length range available was selected to be used among the age readers. These were selected to contain approximately the same number of samples corresponding to individuals of each body length class (size classes of 10 cm), in order to be representative of the total sample size. Once the reference set was complete, growth bands in these 50 vertebrae were then read by the main reader together with two other readers, reaching a consensus for the age of all of them. Then each reader carried out an independent reading. When the results from the three readers had two or three counts differing from the initial agreed age for a certain sample, those vertebrae were analysed again by the three readers together to reach a new consensus/agreed age.

From the total sample size ( $n = 818$ ), 793 samples were used for age readings, with the remaining 25 being initially excluded because of inconsistent band patterns. All 793 vertebrae were read three times and without previous knowledge of the length or sex of each specimen in order to prevent bias while counting the growth bands. To calibrate the readings (i.e., making sure the same criteria were always used when marking the growth zones), a reading of the reference set was carried out before the start of each reading. Also, to prevent familiarity with any particular vertebra, each reading was finished before starting the following one. Additionally, a fourth reading was carried out for the samples whose first three readings produced three different attributed ages, but with two of the three differing only by one year. After all the readings, only vertebrae whose band pair counts obtained three or two out of three equal readings were considered for the age and growth analysis.

In order to compare the precision between the three initial readings, the coefficient of variation (CV) (Chang, 1982), the average percent error (APE) (Beamish and Fournier, 1981), the percentage of agreement (PA) (Beamish and Fournier, 1981) and percentage of agreement within one growth band, and two growth bands ( $PA \pm 1$  year,  $PA \pm 2$  years) were calculated and compared among the readings. Additionally, age

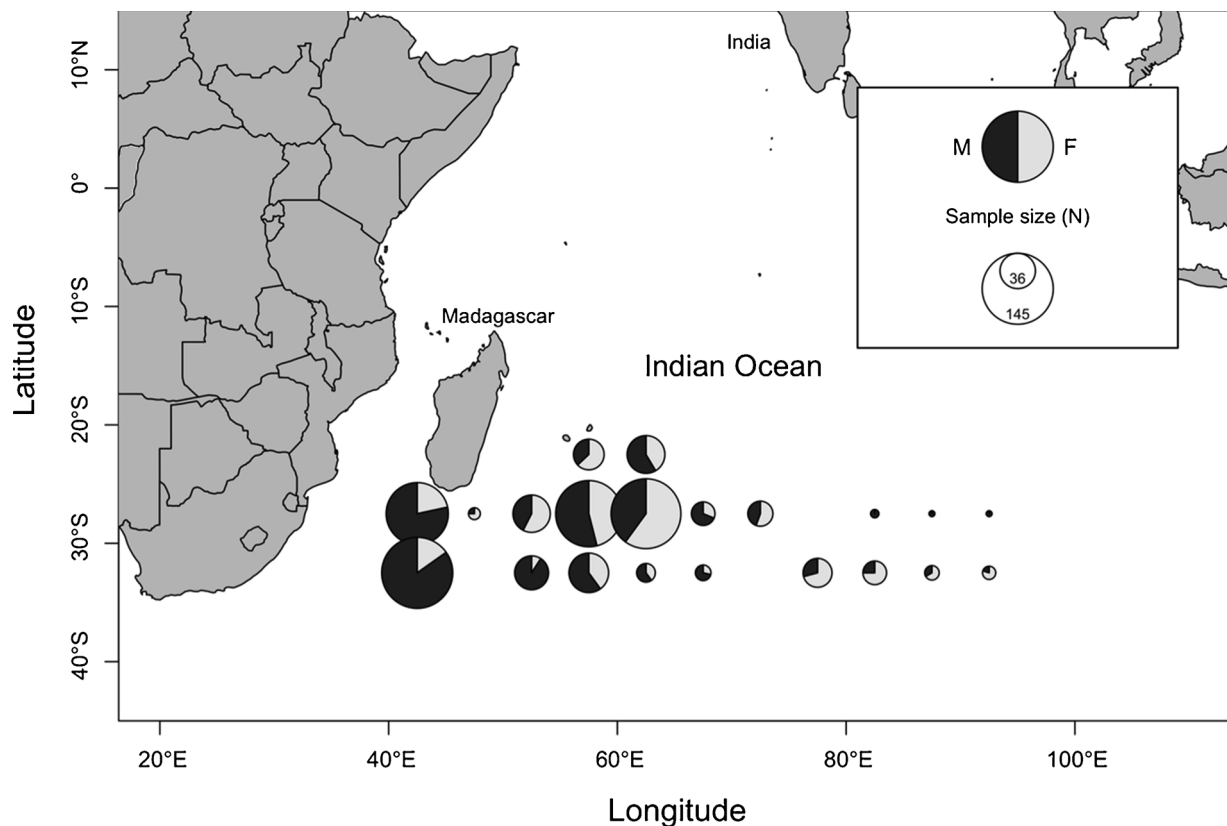


Fig. 1. Map of the area of collection of *Prionace glauca* samples (females and males represented) in the South Indian Ocean. The plots are represented in  $5 \times 5^\circ$  grids, with the sizes of the plots proportional to sample size (N).

bias plots were also used to graphically compare the accuracy of the three readings (Campana, 2001). Each of the three readings (with 95% CI) was plotted for the agreed age. The agreed age was attributed when between the three readings, at least two were identical. The precision analysis was carried out using the R statistical language (R Core Team, 2018).

#### 2.4. Growth modelling

In order to obtain the vertebral radius (VR) of each vertebra, the distance between the focus of vertebrae and the outer edge of the *corpus calcareum* was digitally measured in the photographs of each vertebral sample using the "Measure Cumulative Distances [1]" macro in the ImageJ software. This macro measures cumulative distances along a segmented line selection or between the points of a point selection. The distances were measured to the nearest pixel. Three different scales were used when taking the photos, depending on the size of the vertebrae, and all of them were adjusted to pixels in ImageJ (resulting in  $1 \text{ mm} = 298 \text{ pixels}$ ,  $1 \text{ mm} = 157 \text{ pixels}$  or  $1 \text{ mm} = 99 \text{ pixels}$ ). Since in some of the vertebral sections the tips were broken, thus not showing the focus or the complete outer edge of the *corpus calcareum*, those were not used to calculate VRs, only 714 out of the 818 were used. The relationship between the vertebral radius and fork length ( $L_F$ ) of each specimen was then obtained using a linear model following the Eq. (1):

$$L_F = a + bVR, \quad (1)$$

where,  $b$  is the slope and  $a$  is the intercept. In order to test differences in the  $L_F$  to VR relationship between sexes an ANOVA test was performed.

To obtain growth curves for the studied species, two growth models were used, both of them applied to males and females separately and to the two sexes combined. The first model used was a three-parameter von Bertalanffy growth function (VBGF) (von Bertalanffy, 1938) re-

parameterized to estimate  $L_0$  (size at birth) instead of  $t_0$  (theoretical age at which the expected length is zero) (Cailliet et al., 2006) following Eq. (2):

$$L_t = L_\infty - (L_\infty - L_0) \times \exp(-kt), \quad (2)$$

where  $L_t$  is the mean size ( $L_F$ , cm) at age  $t$  (year),  $L_\infty$  is the maximum asymptotic size ( $L_F$ ),  $L_0$  is the size ( $L_F$ , cm) at birth and  $k$  is the growth coefficient. The second model used was a two-parameter VBGF, following Eq. (2), with  $L_0$  fixed to the medium size at birth of 39.5 cm  $L_F$  described for this species considering the 35–44 cm interval (Pratt, 1979; IOTC, 2007).

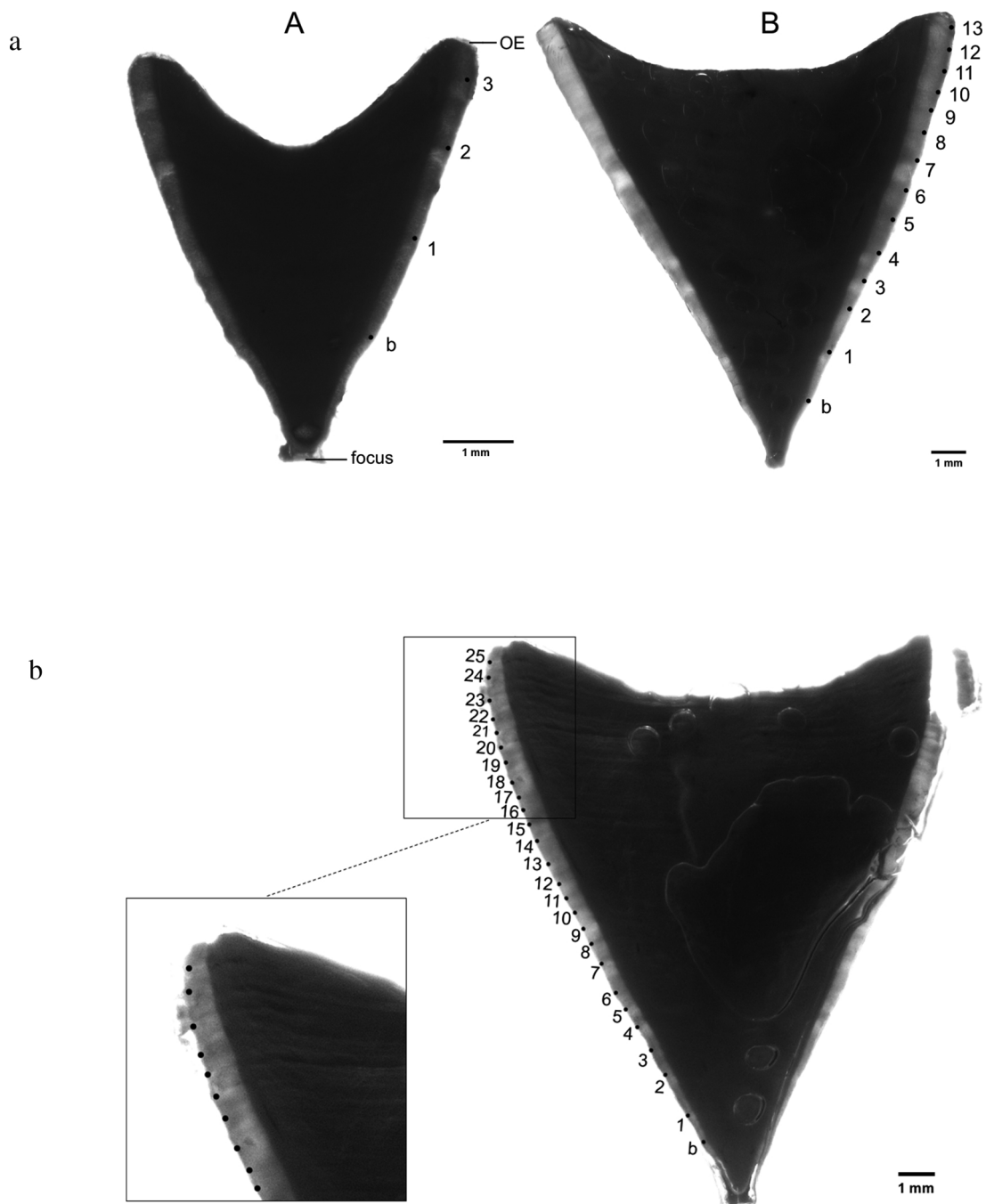
Both models were fitted to the age data, corrected to the midpoint of each age class by adding 0.5 years to the estimated ages, using non-linear least squares (nls function in R) and all plots were created with the package "ggplot2" (Wickham, 2009) in R (R Core Team, 2018). For each of the fitted models, the growth parameters were estimated, along with standard error (SE) and 95% confidence intervals (CIs).

In order to test the null hypothesis that there was no difference in growth parameters between both sexes, a likelihood ratio test (LRT) (Kimura, 1980) was performed on both the three and two-parameter VBGF. Additionally, the model goodness-of-fit was compared with the Akaike Information Criterion (AIC), as well as with the Bayesian Information Criterion (BIC) values. The model with the smallest AIC and BIC values is considered the best fit to the data.

### 3. Results

#### 3.1. Sample characteristics

A total of 818 vertebrae of *P. glauca* specimens were collected for the present study, of which 491 (60%) were from male sharks and 327 (40%) were from females. The size distribution of specimens ranged



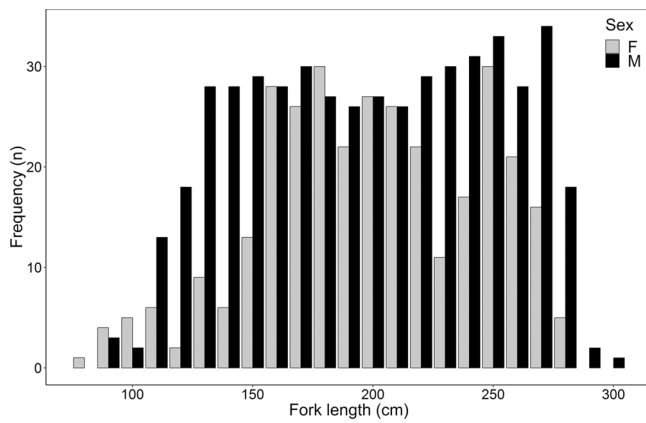
**Fig. 2.** (a)–(b) Microphotographs of three vertebral samples of *Prionace glauca* specimens collected for the present study with identification of the birthmark (b) and the growth bands (indicated by numbers), as well as the focus and the outer edge of the corpus calcareum (OE). The individual on the left (Fig. 2.a. A) has an estimated age of 3 years and the one on the right (Fig. 2.a. B) has an estimated age of 13 years. The individual in Fig. 2.b has an estimated age of 25 years, corresponding to the oldest attributed age in this study.

from 93 to 301 cm  $L_F$  for males (mean  $\pm$  SD: 203  $\pm$  50 cm) and the females ranged from 82 to 284 cm  $L_F$  (mean  $\pm$  SD: 204  $\pm$  41 cm) (Fig. 3).

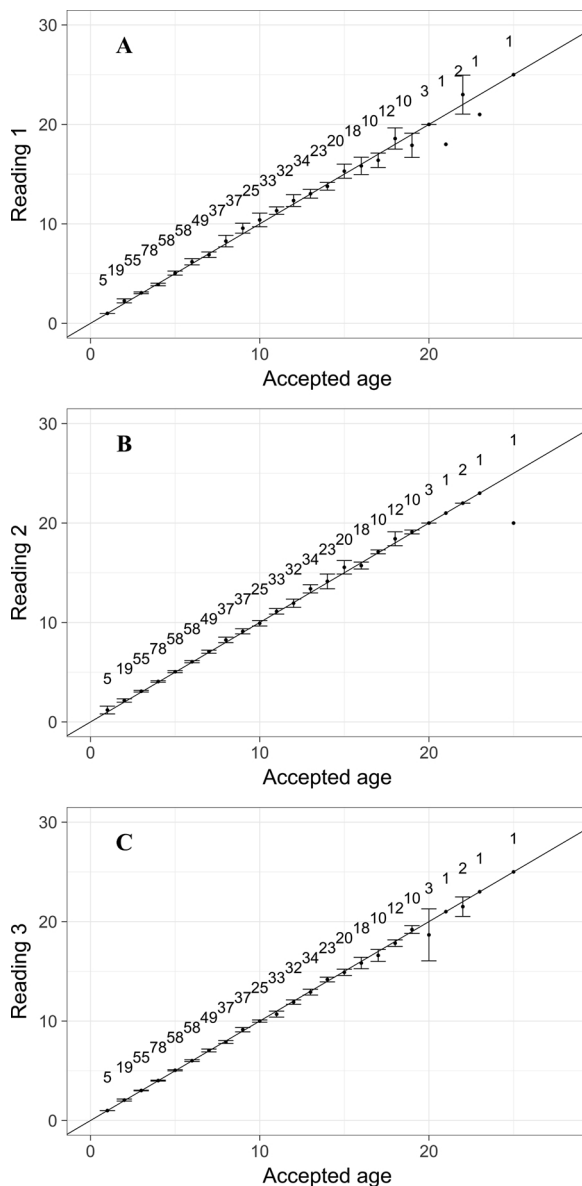
Of the 818 samples, 793 were used for age readings, with 133 of these having three different readings but at least two of them differing only by 1 year, thus a fourth reading was carried out for these 133 samples. After all readings were completed, 679 (85.6%) vertebrae (421 males and 267 females) were considered to have a valid estimated age (at least two identical readings) and were thus considered for the age and growth analysis.

### 3.2. Age estimation and precision analysis

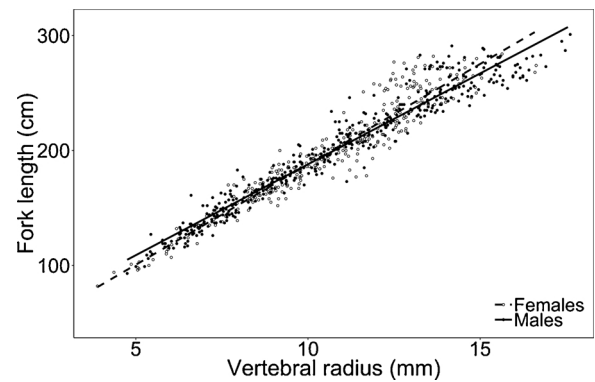
The PA between the three readings, first and the second, first and third and the second and third was 29%, 37%, 44% and 54%, respectively, suggesting a progressive improvement in the consistency of readings. The same pattern was observed with PA  $\pm$  1 and PA  $\pm$  2 years. The CV and APE between the three readings were 8.95% and 6.72%, respectively. The age bias plots (Fig. 4) between each reading and the agreed age between the three reveal a high agreement with no systematic bias.



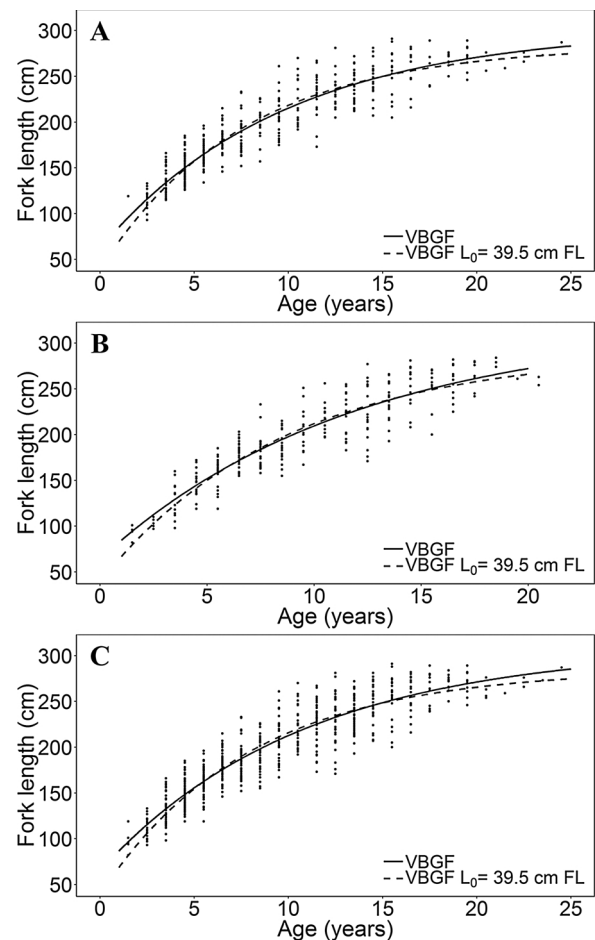
**Fig. 3.** Size ( $L_F$ , cm) frequency distribution of males ( $n = 491$ ) and females ( $n = 327$ ) vertebral samples of *Prionace glauca* individuals collected in the South Indian Ocean between March 2013 and September 2016 ( $n = 818$ ).



**Fig. 4.** Age-bias plots of pairwise age comparisons ( $n = 793$ ) between reading 1 (A), reading 2 (B), reading 3 (C) and the accepted band pair count, for vertebral samples from *Prionace glauca* collected from the South Indian Ocean.



**Fig. 5.** Relationship between the fork length (cm) and the vertebral centrum radius (mm) for *Prionace glauca* males (M) and females (F) from the South Indian Ocean ( $n = 727$ ). Dots represent individual observations and the solid lines represent the linear regressions where  $L_F = 15.82 VR + 29.82$  for males and  $L_F = 17.45 VR + 13.26$  for females.  $L_F$  = fork length and  $VR$  = vertebral radius.



**Fig. 6.** The von Bertalanffy growth function (VBGF) for *Prionace glauca* based on age estimations through counting of vertebrae growth bands ( $n = 679$ ). Circles represent observed data and the lines represent the VBGF (three-parameters VBGF and VBGF with fixed  $L_0$ ) for males (A), females (B) and combined sexes (C).



**Table 1**

Growth parameters estimated for *Prionace glauca* (males, females and combined sexes) in the South Indian Ocean with the three-parameter von Bertalanffy growth function (VBGF) and VBGF with fixed  $L_0$  at 39.5 cm fork length ( $L_F$ ). All parameter estimates for both models are presented with standard error (SE) and 95% confidence levels (95% CI).  $L_\infty$  = maximum asymptotic length,  $k$  = growth coefficient ( $\text{year}^{-1}$ ),  $L_0$  = size at birth (cm  $L_F$ ), AIC = Akaike Information Criteria and BIC = Bayesian Information Criteria. Final parameters recommended to be used are represented in bold (see discussion section for details).

Sex	Model	AIC	BIC	Parameter	Estimate	SE	95% CI	
							Lower	Upper
Males	VBGF	3543	3559	$L_\infty$	302.0	8.3	287.6	321.2
				$k$	0.100	0.009	0.084	0.121
				$L_0$	61.9	6.1	49.4	73.3
				$L_\infty$	283.8	3.6	276.7	291.5
				$k$	0.132	0.004	0.123	0.140
Females	VBGF	2350	2364	$L_\infty$	319.7	18.4	291.1	371.8
				$k$	0.084	0.013	0.058	0.111
				$L_0$	64.1	8.6	46.1	80.5
				$L_\infty$	290.6	6.9	277.9	305.6
				$k$	0.116	0.006	0.104	0.129
Combined	VBGF	5900	5918	$L_\infty$	309.5	8.3	295.0	328.6
				$k$	0.093	0.008	0.078	0.108
				$L_0$	64.9	4.9	54.9	74.2
				$L_\infty$	285.2	3.3	278.7	292.1
				$k$	0.126	0.004	0.119	0.134

### 3.3. Growth modelling

Regarding the relationship between vertebral radius (mm) of each vertebra and the  $L_F$  (cm) of the respective specimen (Fig. 5), the main effect of sex was not significant (ANOVA:  $F(1, 710) = 1.78$ ,  $P > 0.05$ ) but the interaction term between VR and sex is significant (ANOVA:  $F(1, 710) = 21.62$ ,  $P < 0.05$ ). Therefore, the regression equations between VR and  $L_F$  were calculated for females ( $L_F = 17.45 \text{ VR} + 13.26$ ;  $r^2 = 0.91$ ) and males ( $L_F = 15.82 \text{ VR} + 29.82$ ;  $r^2 = 0.95$ ) separately.

A total of 679 *P. glauca* specimens were given a final agreed estimated age, with ages ranging between 1–20 years for females and between 1–25 years for males. The LRT test revealed differences between males and females growth parameters for both the three-parameter VBGF (LRT:  $X^2(3, N = 679) = 11.85$ ,  $P < 0.05$ ) and two-parameter VBGF (LRT:  $X^2(2, N = 679) = 15.99$ ,  $P < 0.05$ ). Therefore, both the three-parameter VBGF and the VBGF with a fixed  $L_0$  were fitted for females and males separately (Fig. 6).

The estimates for growth parameters are displayed in Table 1. The estimated values of  $L_\infty$  were greater for both sexes when using the three-parameter VBGF instead of VBGF with fixed  $L_0$ . For both models,  $L_\infty$  was greater for females. The values for  $k$  were slightly greater when using VBGF with a fixed  $L_0$ . The estimates for  $L_0$  with the three-parameter VBGF were similar for males ( $L_0 = 61.9 \text{ cm } L_F$ ) and females ( $L_0 = 64.1 \text{ cm } L_F$ ) (Table 1). The three-parameter model presented a lower AIC and BIC than the model with a fixed  $L_0$ , suggesting that the first model represents a better fit to the data. However, more biologically reasonable values are likely produced with the two-parameter model with fixed  $L_0$  (see discussion section for details). The results obtained with the recommended final model suggest females reach a greater asymptotic length ( $L_\infty$ ) than males, and males have a greater growth coefficient ( $k$ ), indicating a slower growth for females (males:  $L_\infty = 283.8 \text{ cm } L_F$ ,  $k = 0.13 \text{ year}^{-1}$ ; females:  $L_\infty = 290.6 \text{ cm } L_F$ ,  $k = 0.12 \text{ year}^{-1}$ ).

## 4. Discussion

In the present work, two new growth curves were obtained for *P. glauca* in the Indian Ocean, one for males and the other for females, with new estimated growth parameters. This new information can now be used in future stock assessments, to provide more robust scientific advice on the exploitation of this stock. In addition, the maximum estimated age of 25 years that we report is the highest attributed age to this species so far.

Vertebrae of *P. glauca* are known to be difficult to read due to a poor growth band contrast (Skomal and Natanson, 2003; Manning and Francis, 2005; Rabehagasoa et al., 2014). In the present study, 25 vertebral samples of the total 818 collected were initially excluded because no band pattern was visible that could be quantified. The low number of rejections in the present study may exemplify the fact that other studies have had significant difficulties in the age reading of vertebrae. These represent a very low percentage of rejection, however, in studies with much smaller sample sizes, discarding considerable amounts of samples can represent significant age estimation problems. Some studies of other shark species have used crystal violet as a staining solution to enhance growth band structure in vertebrae (e.g., Coelho et al., 2011; Fernandez-Carvalho et al., 2011, 2015; Rosa et al., 2017), but this approach had not previously been applied to *P. glauca*. In the current study, a comparison was made between unstained and stained vertebral sections, the latter providing a significant improvement to the band structure contrast. Thus, staining sections with crystal violet as well as digitally enhancing the contrast of the growth bands seems to be a better solution to address the difficulties found with age reading of *P. glauca*.

In terms of precision analysis, CV and APE are both widely used in ageing studies. Campana (2001, 2014) suggested values of less than 7.6% for CV and 5.5% for APE, although mentioning that most shark age studies have a CV exceeding 10%. In the present study, CV and APE were of 8.95% and 6.72%, respectively, which is higher than the values suggested by Campana (2001, 2014) but lower than most shark age studies CV reported by the same author. The precision analysis together with the age bias plots, supports the consistency of age estimations and their adequacy for the studied species.

The longevity of *P. glauca* was previously thought to be of about 20–23 years (Cailliet et al., 1983; Manning and Francis, 2005; Stevens, 2009). These values are close to the maximum estimated ages in this study which were 20 and 25 years for females and males, respectively. The oldest individual was a 25 year old male with 301 cm  $L_F$  (Fig. 2.b.). The age estimates here obtained for both sexes are greater than any of the previously estimated ages of *P. glauca*.

Age validation was not performed in the current study, but other studies have verified or validated an annual periodicity of growth band deposition for *P. glauca* that could be applicable to the age readings performed in the current study (Skomal and Natanson, 2003; Lessa et al., 2004; Hsu et al., 2015; Wells et al., 2016). A preliminary age validation study using bomb radiocarbon dating on two male blue sharks from the Indian Ocean provides support for the age reading

**Table 2a**

Summary of previous age and growth studies of *Prionace glauca* in the Indian Ocean. C = combined sexes, F = female, M = male,  $L_T$  = total length,  $L_{PC}$  = precaudal length,  $L_F$  = Fork length, VBGF = von Bertalanffy growth function,  $L_\infty$  = maximum asymptotic size (in cm),  $k$  = growth coefficient,  $w$  = samples used as a whole vertebra and  $s$  = sectioned samples. Spaces filled with “-” refer to information that is not available. Note: The study of Jolly et al. (2013) also included individuals of the Atlantic Ocean.

Study	n	Measure	Length range (cm)	Sex	VBGF parameters		Max. estimated age
					$L_\infty$	$k$	
Jolly et al. (2013)	197	$L_T$	72 - 313	C	311.6	0.12	16
				M	294.6	0.14	
				F	334.7	0.11	
Rabehagasoa et al. (2014)	188	$L_F$	36 - 276	C	258	0.16	15
Present study	679	$L_F$	82 - 301	C	285.2	0.14	25
				M	283.8	0.15	
				F	290.6	0.13	

protocol used in the current study with ages of 18 and 22 years for lengths of 273 cm  $L_F$  and 270 cm  $L_F$ , respectively (E.V. Romanov, 2018, pers. comm.<sup>1</sup>). Although these results are preliminary, these specimens presented different ages despite being of the same sex and almost the same size. Therefore, age validation should not be overlooked, particularly for the largest, and presumably older, individuals. As an example, discrepancies have been observed in longevity estimates of other large pelagic sharks (e.g. sand tiger shark, Passerotti et al., 2014 and dusky shark, Natanson et al., 2014). Bomb radiocarbon dating of these large pelagic shark species have revealed a similar pattern of underestimated age for the largest sharks. In each study, the growth of the vertebrae appears to have growth bands that are missing and cannot be quantified beyond a certain size or age. In addition, another study has challenged the notion that vertebrae can reliably reflect annual growth by showing close ties to somatic growth (Natanson et al., 2018). If this is the case, then vertebrae may stop growing, as evidenced in bomb radiocarbon studies, once maximum size is approached or reached and therefore any estimates of age would be underestimated for a shark that has lived beyond this point in size and age. For *P. glauca*, the ages estimated here are supported to the early to mid 20 s, based on the findings of E.V. Romanov (2018, pers. comm.), but do not preclude the possibility of greater longevity.

Regarding the VBGF used in this study, and specifically for the three parameter equation, the model was re-parameterized to estimate  $L_0$  (size at birth) instead of  $t_0$  (theoretical age at which the expected length is zero). This was based on the fact that  $t_0$  lacks biological meaning making  $L_0$  a more robust approach with an immediate interpretation (Goosen and Smale, 1997; Carlson et al., 2003; Cailliet et al., 2006; Goldman et al., 2012). This is particularly relevant in the case of elasmobranchs, since size at birth is usually well defined (Goldman et al., 2012). The estimated  $L_0$  from our study were 61.9 cm  $L_F$  for males, 64.1 cm  $L_F$  for females and 64.9 cm  $L_F$  for the combined sexes. These values are substantially greater than the 35–44 cm  $L_F$  size at birth range described by Pratt (1979), and by IOTC (2007) in the Indian Ocean. Estimates of  $L_0$  in other *P. glauca* studies fall in this range (Cailliet et al., 1983), while others estimated slightly lower values (Megalofonou et al., 2009; Rabehagasoa et al., 2014). In the present study, the greater estimates for size at birth could be explained by the lack of samples of younger ages when comparing with the remaining ages within the total sample size. This may be due to the smaller individuals of the younger age classes not being fully selected by this gear and therefore the length-at-age for these age classes might be biased. Another reason

could be that there is a limited overlap between the area operated by the fishery and juvenile aggregation areas of the studied species. A recent paper by Coelho et al. (2018) showed that juvenile blue sharks tend to be present mainly in high latitudes, which in this specific case of the southwest Indian Ocean does not fully overlap with the fishery where the samples were collected, specifically swordfish targeting longlines that operate in temperate waters but slightly in more northern waters, not fully in those southern juveniles dominated areas.

In terms of model comparison, the AIC and BIC values for both models used in this study suggest that the three-parameter VBGF has a better statistical fit to the age data (Table 1). However, while the fits are better from a statistical perspective, in biological terms it might be more adequate to use the two-parameter VBGF with a fixed  $L_0$ . This is due to the birth size of the blue shark being already known (Pratt, 1979; IOTC, 2007) and the observed length-at-age might be biased (see discussion above), therefore the three-parameter model is projected towards an unrealistic length at  $t_0$ . As such, the inclusion of a well-known parameter in the model as a fixed value, rather than allowing for its estimation with the associated uncertainties, might be more adequate even at the expense of a somewhat poorer overall fit to the data. Therefore, the VBGF with fixed  $L_0$  was the model selected for this study and was applied to separate sexes due to significant differences in growth characteristics. Additionally, it is relevant to mention that the choice of model makes no practical difference for the majority of the data, as both the three and two parameter VBGF are nearly identical for the 4–17 years old.

When considering previous studies as well as this study (Tables 2a–2c), there are no evident trends in growth between the Atlantic, the Pacific and the Indian oceans, suggesting a similar growth for *P. glauca* among different world regions. This has been previously mentioned by Nakano and Seki (2003) and Tanaka et al. (1990), who reported that variations in the estimates between different studies are most likely due to differences in techniques used to prepare the samples, different criteria for growth zones ageing and reader precision and bias, which compromises a realistic comparison of growth between different areas, and even between studies of the same area. However, it is important to note that previous age and growth studies for *P. glauca* may not even be comparable with more recent studies such as the present one, due to flawed age protocols and very limited size ranges. The  $L_\infty$  estimates in this study are slightly greater than the ones obtained by other authors in the Indian Ocean and  $k$  values are very similar, noting however the issue previously mentioned that using a 3-parameter model with size/age data truncated at the lower range can produce very low  $k$  values (Tables 2a–c). In our study the results suggest that females reach a greater asymptotic length than males, and males have a greater growth coefficient, indicating a slower growth for females. It is of noted,

**Table 2b**

Summary of previous age and growth studies of *Prionace glauca* in the Pacific Ocean.

Study	n	Measure	Length range (cm)	Sex	VBGF parameters		Max. estimated age
					$L_\infty$	$k$	
Cailliet et al. (1983)	130	$L_T$	28- 252.1	C	265.5	0.223	9
				M	295.3	0.175	
				F	241.9	0.251	
Tanaka et al. (1990)	195	$L_T$	110 - 280	M	369	0.10	11
				F	304	0.16	
Nakano (1994)	271	$L_{PC}$	-	M	289.7	0.129	10
				F	243.3	0.144	
Blanco-Parra et al. (2008)	184	$L_T$	90 - 253	C	303.4	0.10	16
				M	299.9	0.10	
				F	237.5	0.15	
Hsu et al. (2015)	742	$L_T$	-	C	352.1	0.13	15

<sup>1</sup> E. Romanov: Centre technique d'appui à la pêche réunionnaise – CAP RUN - NEXA, Le Port, Île de la Réunion.

**Table 2c**

Summary of previous age and growth studies of *Prionace glauca* in the Atlantic Ocean. Note: the study of MacNeil and Campana (2002) strictly looked at the use of whole vs sectioned vertebrae, using samples up to 8 years old only, hence the exceptionally high  $k$  values.

Study	n	Measure	Length range (cm)	Sex	VBGF parameters		Max. estimated age
					$L_{\infty}$	$k$	
Aasen (1966)	268	$L_T$	–	C	394	0.133	8
Stevens (1975)	82	$L_T$	42–272.5	C	423	0.110	7
Henderson et al. (2001)	159	$L_T$	64–228	C	376.5	0.12	6
MacNeil and Campana (2002)	185	$L_F$	147–282	C	300 w 302 s	0.68 w 0.58 s	8
Skomal and Natanson (2003)	411	$L_F$	49–312	C M F	285.4 282.3 286.8	0.17 0.18 0.16	16
Lessa et al. (2004)	236	$L_T$	173.8–310	C	352.1	0.157	12

however, that differences in  $L_{\infty}$  seem to have little impact in this case, as growth and length-at-age are relatively similar over the range of observed data. The same was found in some of the previous studies done all over the world (Tables 2a–c), while in others the opposite results were obtained. The  $L_{\infty}$  values estimated in our study are within the range of values estimated by authors in the three oceans, which range from 198.8 cm  $L_F$  \*<sup>2</sup> to 353 cm  $L_F$  \*. The same happens for the  $k$  estimates of our study, which are within the range of 0.10 year<sup>−1</sup> to 0.18 year<sup>−1</sup> observed in nearly all the other studies.

Overall, the present study provides an improved age reading protocol and is indirectly supported by bomb radiocarbon validation data. Thus, it may provide some of the most reliable life history results to date for this species in the Indian Ocean. The maximum observed age is greater than what was previously described. These results support the fact that *P. glauca* is a long-lived, slow growth species, and provide important additional knowledge to its biology in the Indian Ocean. Nonetheless, the longevity of *P. glauca* could be even greater than estimated since as previously discussed, shark vertebrae may stop growing in some large pelagic species. If that was the case for *P. glauca*, growth parameter values could potentially change. Therefore, more work regarding age and growth of this species in the Indian Ocean should be carried out, with age validation being a priority topic that needs further exploration.

## Acknowledgments

Biological sampling for this study was conducted by the Portuguese Institute for the Ocean and Atmosphere (IPMA), within the scope of the European Data Collection Framework (PNAB/DCF). The authors thank all skippers, crews and fishery observers that contributed with data and sample collection. The authors also thank the extensive revision of the four referees of this manuscript, including Allen H. Andrews (NOAA Fisheries), Malcolm Francis (NIWA) and two anonymous reviewers, as well as the Editor, that have significantly improved the original manuscript. The authors also would like to thank Evgeny Romanov (CAP RUN - NEXA) for revising and providing valuable comments on this manuscript. R. Coelho is supported by an Investigador-FCT contract (Ref: IF/00253/2014) from the Portuguese Foundation for Science and Technology (FCT, Fundação para a Ciência e Tecnologia) supported by

the EU European Social Fund and the Programa Operacional Potencial Humano.

## References

- Aasen, O., 1966. Blåhaien, *Prionace glauca* (Linnaeus), 1758. Fisk. Hav. 1, 1–15.
- Aires-da-Silva, Alexandre, Ferreira, R.L., Pereira, J.G., 2008. Case study: blue shark catch-rate patterns from the Portuguese swordfish longline fishery in the azores. *Sharks of the Open Ocean: Biology, Fisheries and Conservation*. pp. 230–235.
- Anderson, E.D., 1980. Analysis of various sources of pelagic shark catches in the north-west and Western Central Atlantic Ocean and Gulf of Mexico. NOAA Tech. Rep. NMFS 31, 1–14.
- Bailey, K., Williams, P.G., Itano, D., South Pacific Commission, 1996. Programme, oceanic fisheries: by-catch and discards in western pacific tuna fisheries: a review of spc data holdings and literature. Oceanic Fish. Programme Tech. Rep.
- Beamish, R.J., Fournier, D.A., 1981. A method for comparing the precision of a set of age determinations. Can. J. Fish. Aquat. Sci. 38 (8), 982–983. <https://doi.org/10.1139/f81-132>.
- Blanco-Parra, M.D.P., Galvan-Magana, F., Marquez-Farias, F., Galván-Magaña, F., Márquez-Farías, F., 2008. Age and growth of the blue shark, *Prionace glauca* Linnaeus, 1758, in the Northwest coast off Mexico. Rev. Biol. Mar. Y Oceanogr. 43 (3), 513–520. <https://doi.org/10.4067/S0718-19572008000300010>.
- Cailliet, G.M., Martin, L.K., Harvey, J.T., Kusher, D., Welden, B.A., 1983. Preliminary studies on the age and growth of the blue, *Prionace glauca*, common thresher, *Alopias vulpinus*, and shortfin mako, *Isurus oxyrinchus*, sharks from California waters. *Proceedings of the International Workshop on Age Determination of Oceanic Pelagic Fishes: tunas, Billfishes, and Sharks*. NOAA Tech. Rep. NMFS 8, 179–188.
- Cailliet, G.M., Smith, W.D., Mollet, H.F., Goldman, K.J., 2006. Age and growth studies of chondrichthyan fishes: the need for consistency in terminology, verification, validation, and growth function fitting. Environ. Biol. Fishes 77 (3–4), 211–228. <https://doi.org/10.1007/s10641-006-9105-5>.
- Campana, S.E., 2001. Accuracy, precision and quality control in age determination, including a review of the use and abuse of age validation methods. J. Fish Biol. 59 (2), 197–242. <https://doi.org/10.1006/jfbi.2001.1668>.
- Campana, S.E., 2014. Age Determination of Elasmobranchs, with Special Reference to Mediterranean Species: A Technical Manual. Studies and Reviews. General Fisheries Commission for the Mediterranean. FAO, Rome.
- Campana, S.E., Joyce, W., Manning, M.J., 2009. Bycatch and discard mortality in commercially caught blue sharks *Prionace glauca* assessed using archival satellite pop-up tags. Mar. Ecol. Prog. Ser. 387, 241–253. <https://doi.org/10.3354/meps08109>.
- Carlson, J.K., Cortés, E., Bethea, D.M., 2003. Life history and population dynamics of the finetooth shark (*Carcharhinus isodon*) in the northeastern Gulf of Mexico. Fish. Bull. 101 (2), 281–292.
- Carruthers, E.H., Neilson, J.D., Smith, S.C., 2011. Overlooked bycatch mitigation opportunities in pelagic longline fisheries: soak time and temperature effects on swordfish (*Xiphias gladius*) and blue shark (*Prionace glauca*) catch. Fish. Res. 108 (1), 112–120. <https://doi.org/10.1016/j.fishres.2010.12.008>.
- Casey, J.G., Hoey, J.J., 1985. Estimated catches of large sharks by US recreational fishermen in the Atlantic and Gulf of Mexico. NMFS 31 pp. 35–19.
- Castro, J.A., Mejuto, J., 1995. Reproductive parameters of blue shark, *Prionace glauca*, and other sharks in the gulf of guinea. Mar. Freshw. Res. 46 (6), 967–973.
- Chang, W.Y.B., 1982. A statistical method for evaluating the reproducibility of age determination. Can. J. Fish. Aquat. Sci. 39, 1208–1210.
- Coelho, R., Fernandez-carvalho, J., Amorim, S., Santos, M.N., 2011. Age and growth of the smooth hammerhead shark, *Sphyrna zygaena*, in the Eastern Equatorial Atlantic Ocean, using vertebral sections. Aquat. Living Resour. 24 (4), 351–357.
- Coelho, R., Mejuto, J., Domingo, A., Yokawa, K., Romanov, E.V., Cortés, K.L.E., Liu, K., da Silva, C., Hazin, F., Arocha, F., Mwiliama, A.M., Bach, P., de Zárate, V.O., Roche, W., Lino, P.G., García-Cortés, B., Ramos-Cardelle, A.M., Forselledo, R., Mas, F., Ohshima, S., Courtney, D., Sabarros, P.S., Perez, B., Wogerbauer, C., Tsai, W., Carvalho, F., Santos, M.N., 2018. Distribution patterns and population structure of the blue shark (*Prionace glauca*) in the Atlantic and Indian Oceans. Fish. Fish. 19 (1), 90–106. <https://doi.org/10.1111/faf.12238>.
- Compagno, L.J.V., 1984. FAO Species Catalogue. Vol. 4. Part 2. Sharks of the World. An Annotated and Illustrated Catalogue of Shark Species Known to Date. FAO, Rome.
- Dent, F., Clarke, S., 2015. State of the Global Market for Shark Products. FAO Fisheries and Aquaculture Technical Paper No 590. FAO, Rome.
- Eriksson, H., Clarke, S., 2015. Chinese market responses to overexploitation of sharks and sea cucumbers. Biol. Conserv. 184, 163–173. <https://doi.org/10.1016/j.biocon.2015.01.018>.
- Fernandez-Carvalho, J., Coelho, R., Erzini, K., Santos, M.N., 2011. Age and growth of the bigeye thresher shark, *Alopias superciliosus*, from the pelagic longline fisheries in the tropical northeastern Atlantic Ocean, determined by vertebral band counts. Aquat. Living Resour. 24 (4), 359–368. <https://doi.org/10.1051/alr/2011046>.
- Fernandez-Carvalho, J., Coelho, R., Erzini, K., Santos, M.N., 2015. Modeling age and growth of the bigeye thresher (*Alopias superciliosus*) in the Atlantic Ocean. Fish. Bull. 113 (4), 468–481. <https://doi.org/10.7755/FB.113.9>.
- Francis, M.P., Griggs, L.H., Baird, S.J., 2001. Pelagic shark bycatch in the New Zealand tuna longline fishery. Mar. Freshw. Res. 52 (2), 165–178.
- Francis, M.P., Ó Maolagáin, C., 2016. Size, Maturity and Length Composition of Blue Sharks Observed in New Zealand Tuna Longline Fisheries. New Zealand Fisheries Assessment Report 2016/60.
- Goldman, K.J., Cailliet, G.M., Andrews, A.H., Natanson, L.J., 2012. Assessing the Age and Growth of Chondrichthyan Fishes. *Biology of Sharks and Their Relatives*. pp. 423–451.

<sup>2</sup> All  $L_F$  \* (fork length) measures were obtained by converting original  $L_T$  (total length) measures using the equation by Kohler et al. (1995) for *P. glauca*:  $L_F = 0.8313 \times L_T + 1.39$ .



- <https://doi.org/10.1017/CBO9781107415324.004>.
- Goosen, A.J.J., Smale, M.J., 1997. A preliminary study of age and growth of the smoothhound shark *Mustelus mustelus* (Triakidae). South Afr. J. Mar. Sci. 18 (1), 85–91. <https://doi.org/10.2989/025776197784161072>.
- Henderson, A., Flannery, K., Dunne, J., 2001. Observations on the biology and ecology of the blue shark in the North-east Atlantic. J. Fish Biol. 58 (5), 1347–1358. <https://doi.org/10.1006/jfbi.2000.1547>.
- Hsu, H.H., Lyu, G.T., Joung, S.J., Liu, K.M., 2015. Age and growth of the blue shark (*Prionace glauca*) in the South Atlantic Ocean. Collect. Vol. Sci. Pap. ICCAT 71 (6), 2573–2584.
- IOTC, 2007. Compilation of Information on Blue Shark (*Prionace glauca*), Silky Shark (*Carcharhinus falciformis*), Oceanic Whitetip Shark (*Carcharhinus longimanus*), Scalloped Hammerhead (*Sphyrna lewini*) and Shortfin Mako (*Isurus oxyrinchus*) in The Indian Ocean. IOTC-2007-WPEB-INF01. pp. 1–18.
- IOTC, 2016. Report of the 6 th Session of the IOTC Working Party on Neritic Tunas. Seychelles. IOTC-2016-WPNT06-R. pp. 1–89.
- Jolly, K.A., da Silva, C., Attwood, C.G., 2013. Age, growth and reproductive biology of the blue shark *Prionace glauca* in South African waters. Afr. J. Mar. Sci. 35 (1), 99–109. <https://doi.org/10.2989/1814232x.2013.783233>.
- Kimura, D., 1980. Likelihood methods for the von bertalanffy growth curve. Fish. Bull. 77, 765–776.
- Kohler, N.E., Casey, J.G., Turner, P.A., 1995. Length-weight relationships for 13 species of sharks from the western North Atlantic. Fish. Bull. 93 (2), 412–418.
- Lessa, R., Santana, F.M., Hazin, F.H., 2004. Age and growth of the blue shark *Prionace glauca* (Linnaeus, 1758) off northeastern Brazil. Fish. Res. 66 (1), 19–30. [https://doi.org/10.1016/S0165-7836\(03\)00193-0](https://doi.org/10.1016/S0165-7836(03)00193-0).
- MacNeil, M.A., Campana, S.E., 2002. Comparison of whole and sectioned vertebrae for determining the age of young blue shark (*Prionace glauca*). J. Northwest Atl. Fish. Sci. 22, 77–82. <https://doi.org/10.2960/J.v30.a5>.
- Manning, M.J., Francis, M.P., 2005. Age and growth of blue shark (*Prionace glauca*) from the New Zealand Exclusive Economic Zone. New Zealand Fisheries Assessment Report 26, 52.
- McKenzie, R.A., Tibbo, S.N., 1964. A morphometric description of blue shark (*Prionace glauca*) from Canadian Atlantic waters. J. Fish Res. Board Can. 21 (4), 865–866.
- Megalofonou, P., Damalas, D., De Metrio, G., 2009. Biological characteristics of blue shark, *Prionace glauca*, in the Mediterranean Sea. J. Mar. Biol. Assoc. U. K. 89 (6), 1233–1242. <https://doi.org/10.1017/S0025315409000216>.
- Muñoz-Lechuga, R., Rosa, D., Coelho, R., 2016. Depredation in the Portuguese Pelagic Longline Fleet in the Indian Ocean. IOTC–2016–WPEB12-35. pp. 1–14.
- Nakano, H., 1994. Age, Reproduction and Migration of Blue Shark in the North Pacific Ocean. Bulletin of the Natural Research Institute of Far Seas Fisheries (Japan).
- Nakano, H., Seki, M., 2003. Synopsis of biological data on the blue shark, *Prionace glauca* Linnaeus. Bull. Fish. Res. Agency 6, 18–55.
- Natanson, L.J., Stevens, J., 2008. The biology and ecology of the Blue shark, *Prionace glauca*. Sharks of the Open Ocean: Biology, Fisheries and Conservation. pp. 140–151. <https://doi.org/10.1002/9781444302516>.
- Natanson, L.J., Gervelis, B.J., Winton, M.V., Hamady, L.L., Gulak, S.J.B., Carlson, J.K., 2014. Validated age and growth estimates for *Carcharhinus obscurus* in the north-western Atlantic Ocean, with pre- and post management growth comparisons. Environ. Biol. Fishes 97 (8), 881–896. <https://doi.org/10.1007/s10641-013-0189-4>.
- Natanson, L.J., Skomal, G.B., Hoffmann, S.L., Porter, M.E., Goldman, K.J., Serra, D., 2018. Age and growth of sharks: do vertebral band pairs record age? Mar. Freshw. Res. <https://doi.org/10.1071/MF17279>.
- Passerotti, M.S., Andrews, A.H., Carlson, J.K., Wintner, S.P., Goldman, K.J., Natanson, L.J., 2014. Maximum age and missing time in the vertebrae of sand tiger shark (*Carcharias taurus*): validated lifespan from bomb radiocarbon dating in the western North Atlantic and southwestern Indian Oceans. Mar. Freshw. Res. 65 (8), 674–687. <https://doi.org/10.1071/MF13214>.
- Pratt, H.L., 1979. Reproduction in the blue shark, *Prionace glauca*. Fish. Bull. 77 (2), 445–470.
- Rabehagasoa, N., Vigliola, L., Lorrain, A., Sabarros, P.S., Romanov, E., Bach, P., 2014. Modelling Growth of Blue Shark (*Prionace Glauca*) and Silky Shark (*Carcharhinus Falciformis*) in the Southwest Indian Ocean Assessed by Back-Calculated Length From Vertebrae. IOTC–2014–WPEB10–22, (October). pp. 1–23.
- R Core Team, 2018. R: A Language and Environment for Statistical Computing. R Foundation for Statistical Computing. R Development Core Team, Vienna, Austria. [www.R-project.org](http://www.R-project.org).
- Rosa, D., Coelho, R., Fernandez-carvalho, J., Santos, M.N., 2017. Age and growth of the smooth hammerhead, *Sphyrna zygaena*, in the Atlantic Ocean: comparison with other hammerhead species. Mar. Biol. Res. 13 (0), 300–313. <https://doi.org/10.1080/17451000.2016.1267366>.
- Schindelin, J., Rueden, C.T., Hiner, M.C., Eliceiri, K.W., 2015. The ImageJ ecosystem: an open platform for biomedical image analysis. Mol. Reprod. Dev. 82 (7–8), 518–529.
- Sims, D., Fowler, S.L., Ferretti, F., Stevens, J., 2016. *Prionace glauca*. The IUCN Red List of Threatened Species 2016: e.T39381A16553182. Available at. (last Accessed on 3 October 2017). <http://www.iucnredlist.org/details/39381/3>.
- Sims, D., Fowler, S.L., Ferretti, F., Stevens, J.D., 2015. *Prionace glauca*. The IUCN Red List of Threatened Species 2015: e.T39381A48924261. Available at. (last Accessed on 3 October 2017). <http://www.iucnredlist.org/details/39381/1>.
- Skomal, G., Natanson, L., 2003. Age and growth of the blue shark (*Prionace glauca*) in the North Atlantic Ocean. Fish. Bull. 101 (3), 627–639.
- Stevens, J., 2009. *Prionace glauca*, Blue Shark. The IUCN Red List of Threatened Species 2009. Available at. pp. 8235. <https://doi.org/10.2305/IUCN.UK.2009-2.RLTS.T39381A10222811.en>. (last Accessed on 3 October 2017). <http://www.iucnredlist.org/details/39381/0>.
- Stevens, J.D., 1975. Vertebral rings as a means of age determination in the blue shark (*Prionace glauca* L.). J. Mar. Biol. Assoc. U. K. 55 (3), 657–665. <https://doi.org/10.1017/S0025315400017318>.
- Stevens, J.D., 1984. Biological observations of sharks caught by sportfishermen off New South Wales. Aust. J. Mar. Freshwater Res. 35 (5), 573–590.
- Stevens, J.D., 1992. Blue and mako shark by-catch in the Japanese longline fishery off south-eastern Australia. Mar. Freshw. Res. 43 (1), 227–236. <https://doi.org/10.1071/MF9920227>.
- Tanaka, S.H.O., Cailliet, G.M., Yudin, K.G., 1990. Differences in age and growth of the Blue Shark, *Prionace glauca*: technique or population? NOAA Tech. Rep. NMFS 90, 177–187.
- Vas, P., 1990. The abundance of the blue shark, *Prionace glauca*, in the western English Channel. Environ. Biol. Fishes 29 (3), 209–225. <https://doi.org/10.1007/BF00002221>.
- von Bertalanffy, L., 1938. A quantitative theory of organic growth. Hum. Biol. 10 (2), 181–213.
- Wells, R.J.D., Spear, N., Kohin, S., 2016. Age validation of the blue shark (*Prionace glauca*) in the eastern Pacific Ocean. Mar. Freshw. Res. 68 (6), 1130–1136.
- Wickham, H., 2009. Ggplot2: Elegant Graphics for Data Analysis. Available at. (last Accessed 15 September 2017). <http://ggplot2.org>.