Annual Bands in Vertebrae Validated by Bomb Radiocarbon Assays Provide Estimates of Age and Growth of Whale Sharks

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Conservation and management strategies for endangered and threatened species require accurate estimates of demographic parameters such as age and growth. The whale shark, *Rhincodon typus*, is the largest fish in the world and is highly valued in the eco-tourism sector. Despite conservation concerns and advances in our understanding of their life history, basic demographic parameters for growth, longevity and mortality are of questionable accuracy; previous growth studies could not agree whether the vertebral growth bands were formed annually or biannually. Here, we provide the first validation of the annual formation of growth bands within the vertebrae of the whale shark using bomb radiocarbon assays. Ages of up to 50 years were estimated from sectioned vertebrae of sharks collected in Taiwan and Pakistan. There was no cessation of the formation of growth bands in the vertebrae of older sharks and our study provides the oldest observed longevity for this species. Initial estimates of growth ($k = 0.01–0.12$) and natural mortality rates ($M = 0.09–0.14$) are consistent with those expected of long-lived sharks, which highlights their sensitivity to fishing pressure and conservation concerns.

Keywords: whale shark, vertebrae, age determination, radiocarbon, longevity, growth bands

INTRODUCTION

Accurate and reliable estimates of the age and growth of individuals in a population are central to effective strategies for the management and conservation of any species. For teleost marine fishes, estimates of age are usually obtained from counts of the annual growth bands formed within otoliths, which are calcified structures within the skull case (Campana, 2001). For elasmobranchs such as sharks, skates and rays, which lack otoliths, age estimates have been calculated from growth bands formed in the vertebrae (Cailliet, 1990).

It is critical that age estimates provided by otoliths and vertebrae are accurate, since uncertainty or underestimates surrounding these ages can lead to stock collapses
of exploited species (e.g., orange roughy *Hoplostethus atlanticus*; Smith et al., 1995), or compromise the effectiveness of recovery programs for species that are threatened or endangered. For this reason, many studies have sought to validate the timing of the production of growth bands (Campana, 2001). A common approach is to tag individuals with a chemical marker such as oxytetracycline (OTC) that is laid down within an otolith or vertebrae. Individuals are released and when recaptured at some time in the future, the tag acts as a time stamp that allows the rate of deposition of subsequent growth bands to be determined. For large fishes and sharks that are relatively long-lived and difficult to tag and recapture, validation of annual banding patterns can also be obtained through an analysis of bomb radiocarbon in vertebrae. Above-ground testing of thermonuclear weapons in the 1950s and 60s increased the ratio of carbon 14 isotopes in the atmosphere that were then mixed into the ocean, passed up food webs and incorporated into marine organisms. As a result, the timing of the deposition of bands can be validated by comparing carbon isotope values within vertebrae, with an isotope baseline chronology of known age (Campana, 2001; Campana et al., 2002; Goldman et al., 2012).

The whale shark, *Rhincodon typus*, is a huge (up to 18 m length; Mcclain et al., 2015), highly migratory, filter-feeding shark found in all tropical and warm temperate seas (Compagno, 2001; Chen et al., 2002; Stevens, 2007). It forms aggregations in productive coastal areas and is a highly valued target for marine eco-tourism (e.g., Huveneers et al., 2017). However, the whale shark has recently been classified as Endangered (IUCN Red List; Pierce and Norman, 2016) and there is now an urgent need for reliable and accurate information on age and growth of the species in order to develop effective conservation and management strategies. At present, there is relatively little demographic data available, especially for large or mature individuals. Using X-radiography, Wintner (2000) analyzed the growth bands in whole vertebrae of juveniles that had stranded on the coast of South Africa to develop an initial growth curve for the species. More recently, Hsu et al. (2014) provided growth and age estimates for individuals collected from a fishery in Taiwan and used marginal increment ratios and centrum edge analysis to conclude that growth bands were deposited biannually in both whole and sectioned vertebrae. The reliability of the age estimates of these studies remains unknown and is of concern, because other studies show that whole vertebrae are known to provide underestimates of age and longevity, and thus overestimates of growth rate in slow-growing sharks (Cailliet and Goldman, 2004; Harry, 2018; Natanson et al., 2018). To our knowledge, only one study has attempted to validate an aging method for whale sharks, which involved a captive immature shark reared in an aquarium after being fed OTC. When the animal died 2 years later, two growth bands were observed following the OTC mark (Wintner, 2000).

Here, we provide the first age validation of whale sharks using bomb radiocarbon assays. We then used sectioned vertebrae from a small sample of sharks to provide initial estimates of growth, longevity and mortality data that can be used in support of current conservation and management efforts.

**METHODS**

**Sample Collection**

A subset of vertebral samples were taken from 92 vertebral samples that were previously published in Hsu et al. (2014). These were dead individuals that had been landed by the Taiwanese fishery, before the whale shark fishery was closed in November 2007 (Hsu et al., 2012). The vertebral sample from Pakistan was obtained from a dead stranded whale shark.

**Sample Preparation and Age Interpretation**

The vertebral samples from Hsu et al. (2014) were sectioned with a single cut using paired blades separated by a spacer on an Isomet low-speed diamond-bladed saw. Sections were digitally photographed at 2048 × 1536 resolution using a digital color Leica camera DFC295 mounted on a stereo microscope Leica M205C (Leica Microsystems, Germany), while immersed in ethanol. Age interpretation was based on images enhanced for contrast using Adobe Photoshop CS6, following the interpretation criteria of Natanson et al. (2002). The precision of the age determinations was quantified with both average percent error (APE) and coefficient of variation (CV) (Campana, 2001).

**Bomb Radiocarbon Analyses**

Vertebrae used for bomb radiocarbon age validation were taken from two specimens that had died after becoming entangled in fishing gear. A 10 m total length (TL) female with an estimated weight of 7000 kg was landed in Karachi, Pakistan in February 2012. One of the cervical vertebrae was cleaned of tissue and then stored frozen. A second individual, a mature male weighing 8500 kg with TL of 9.9 m was landed in Taiwan in April 2005. A cervical vertebra over the gills was extracted and stored in ethanol until assay. Vertebral growth bands from both sharks were isolated from 1 mm thick longitudinal sections of the vertebrae. All sections were prepared using the same procedure outlined above. Sections were digitally photographed at 2048 × 2048 resolution under a binocular microscope at 16–40X magnification using reflected light while immersed in ethanol.

Multiple samples from each of the vertebral sections (*N* = 11 samples; 5–13 mg each) were extracted from growth bands visible in the corpus calcareum region while working at 16X magnification under a binocular microscope. For the shark landed in Taiwan, the first three growth bands were extracted as a single sample from the vertebral section. For the shark landed in Pakistan, the first-formed growth band (distal to the birth band) was extracted, as were individual growth bands corresponding to later growth. Extracted samples were isolated as solid pieces using a Gesswein high-speed hand tool fitted with steel bits < 1 mm in diameter. The presumed date of sample formation was calculated as the year of shark collection minus the annulus count from the birth band to the mid-point of the sample. After sonification in Super Q water and drying, the sample was weighed to the nearest 0.1 mg in preparation for $^{14}$C assay with accelerator mass spectrometry (AMS). AMS assays also provided $\delta^{13}$C (%) values,
which were used to correct for isotopic fractionation effects. Radiocarbon values were subsequently reported as $\Delta^{14}C$, which is the per mil ($\delta$) deviation of the sample from the radiocarbon concentration of 19th-century wood, corrected for sample decay prior to 1950 according to methods outlined by Stuiver and Polach (1977). The mean standard deviation of the individual radiocarbon assays was about $4\delta$.

To assign dates of formation to an unknown sample, it is necessary that the $\Delta^{14}C$ of the unknown sample be compared with a $\Delta^{14}C$ chronology based on known-age material (a reference chronology). Since whale sharks are surface planktivores, we assumed that a reference chronology for dissolved inorganic carbon (DIC) in surface waters was most appropriate for our analysis. Therefore, we used a reference chronology developed from young known-age otoliths (calcium carbonate) in the northwest Atlantic (Campana et al., 2008), which has a period of increasing bomb radiocarbon values nearly identical to that of surface waters off of both Pakistan and Taiwan (Andrews et al., 2011a). We also included another reference chronology based on corals from the Mentawai Islands in Sumatra, Indonesia (Grumet et al., 2004).

**Growth Models**

Preliminary growth estimates were obtained from length-at-age data using two types of growth model. The first was a conventional 3-parameter von Bertalanffy growth function (Von Bertalanffy, 1938) and the second was a logistic growth function (Smart et al., 2013) with length-at-birth fixed at 60 cm (Chang et al., 1997).

3-parameter von Bertalanffy growth function:

$$L_t = L_0 + (L_\infty - L_0)(1 - e^{-kt})$$

Logistic growth function with fixed length-at-birth:

$$L_t = \frac{L_\infty L_0 e^{kt}}{L_\infty + L_0 (e^{kt} - 1)}$$

where $L_t$ is length-at-age $t$, $L_0$ is length-at-age 0, $L_\infty$ is asymptotic length and $k$ is the growth coefficient.

Longevity estimates generally require either a precisely defined growth model or an estimate of mortality rate, neither of which were available here. Therefore, only the observed maximum age is reported here. Natural mortality was estimated from two equations. The first was based on the linear regression equation of observed maximum age (Hoenig, 1983):

$$\ln(M) = 1.44 - 0.982 \times \ln(t_{\text{max}})$$

The second natural mortality estimate was based on the non-linear least squares equation of observed maximum age, with a prediction error of 0.32 (Then et al., 2015):

$$M = 4.899 \times t_{\text{max}}^{-0.916}$$

where $M$ is the estimated instantaneous rate of natural mortality and $t_{\text{max}}$ is the observed maximum age.

**RESULTS**

**Counts of Growth Bands in Vertebrae Samples**

All vertebrae exhibited distinct growth band (annulus) patterns (Figure 1). The birth mark was identified as the most pronounced first band. Subsequent annuli consisted of a pair of alternating opaque and translucent bands that crossed the entire centrum, except in the oldest sharks. Band width decreased with age, narrowing substantially in the oldest individuals (Figure 1). Counts of growth bands in 20 sharks ranged from 15 to 50 (Table 1). Aging precision was acceptable across both readers, with an APE of 5.5% and CV of 8.2%.

**Bomb Radiocarbon Assays and Age Validation**

The date of formation of the vertebral samples was estimated in two ways: (1) through age determination of the shark based on growth band (annulus) counts; and (2) through comparison...
TABLE 1 | Summary details for vertebral samples.

| ID  | Date of collection | Sex | TL (cm) | Weight (kg) | Growth bands |
|-----|--------------------|-----|---------|-------------|--------------|
| RT01 | 18 March 2001      | F   | 425     | 760         | 23           |
| RT02 | 26 May 2005        | F   | 345     | 400         | 18           |
| RT03 | NA                 | M   | 596     | 2000        | 24           |
| RT04 | NA                 | NA  | NA      | NA          | 18           |
| RT05 | NA                 | M   | 423     | 750         | 18           |
| RT06 | NA                 | M   | 391     | 600         | 21           |
| RT07 | 18 May 2005        | M   | 421     | 700         | 25           |
| RT08 | 30 September 2005  | M   | 372     | 350         | 20           |
| RT09 | 26 April 2005      | M   | 358     | 360         | 16           |
| RT10 | 3 July 2005        | M   | 494     | 1100        | 16           |
| RT11 | 6 October 2005     | F   | NA      | NA          | 15           |
| RT12 | 2 July 2005        | M   | 503     | 905         | 18           |
| RT13 | 13 September 2005  | M   | 300     | 200         | 18           |
| RT14 | 10 July 2005       | M   | 430     | 450         | 19           |
| RT15 | 18 June 2005       | M   | 386     | 450         | 17           |
| RT16 | NA                 | NA  | NA      | NA          | 19           |
| RT17 | 10 October 2005    | M   | 526     | NA          | 19           |
| RT18 | 3 May 2005         | M   | 268     | 123         | 17           |
| RT19 | April 2005         | M   | 990     | 8500        | 35           |
| RT20 | February 2012      | F   | 1000    | 7000        | 50           |

TL, total length (cm). RT19 and RT20 were samples used in the bomb radiocarbon analyses. All samples were collected in Taiwan except for RT20, which was collected in Pakistan.

TABLE 2 | Details for bomb radiocarbon assays.

| ID   | Growth bands sampled | Date of formation | δ13C (%) | Δ14C (%) |
|------|-----------------------|-------------------|----------|----------|
| RT19 | 1–3                   | 1972.0            | −15.1    | 81.3     |
| RT20 | 0–1                   | 1962.5            | −13.3    | 15.1     |
| RT21 | 1                     | 1962.5            | −13.3    | 15.1     |
| RT22 | 2                     | 1963.5            | −13.9    | 59.4     |
| RT23 | 3                     | 1964.5            | −13.1    | 70.0     |
| RT24 | 4                     | 1965.5            | −12.9    | 70.0     |
| RT25 | 5                     | 1966.5            | −15.5    | 40.0     |
| RT26 | 5                     | 1966.5            | −15.6    | 40.0     |
| RT27 | 6                     | 1967.5            | −12.8    | 20.6     |
| RT28 | 10                    | 1971.5            | −12.1    | 63.4     |
| RT29 | 15                    | 1976.5            | −11.9    | 45.0     |

RT19 was collected in Taiwan and RT20 was collected in Pakistan. Each row is a separate assay, any replicates were from different assays.

Preliminary Growth, Longevity and Mortality Estimates

Both sexes were combined for estimation of growth (Figure 2B). The von Bertalanffy growth function produced an asymptotic length $L_\infty = 2189$ cm and a growth coefficient $k = 0.014 \text{ year}^{-1}$. The logistic growth function with a fixed length-at-birth $L_0 = 60$ cm produced estimates of $L_\infty = 1071$ cm and $k = 0.122 \text{ year}^{-1}$. We caution, however, that the estimates of asymptotic length and growth coefficients are uncertain because of low sample size. The maximum observed age was 50 years based on vertebral aging and bomb radiocarbon assays. The Hoenig (1983) estimated rate of instantaneous natural mortality was 0.09 year$^{-1}$, while the estimate from Then et al. (2015) was 0.14 year$^{-1}$.

DISCUSSION

Our study used bomb radiocarbon assays to provide the first validated age estimates for whale sharks. We showed that growth bands in sectioned vertebrae can provide accurate estimates of sharks aged up to 50 years old. These results confirm the use of sectioned vertebrae as age indicators for these sharks, as is also the case for other large species, such as white (Hamady et al., 2014), shortfin mako (Natanson et al., 2006), sandbar (Andrews et al., 2011b), and porbeagle (Natanson et al., 2002) sharks. We found no evidence that vertebral counts underestimated the age of older individuals, as can be the case for porbeagle and white sharks (Francis et al., 2007; Hamady et al., 2014), presumably because the much larger asymptotic body size means that there is no cessation of vertebral growth in the older sharks we sampled in our study.

Although our understanding of the movements, behavior, connectivity and distribution of whale sharks have improved dramatically over the last 10 years (Schmidt et al., 2009; Sequeira et al., 2012, 2013), basic life history traits such as age, longevity and mortality remain unknown and are frequently inferred (e.g., Bradshaw et al., 2007). This lack of basic demographic information has been consistently highlighted in multiple reviews.

of annulus $\Delta^{14}C$ values with the values known to be present in surface marine waters at the time (the NWA reference chronology). Agreement between the annulus- and $\Delta^{14}C$ -based dates would confirm that the annuli were interpreted correctly for age estimation, at least on average. Under- or over-aging of annuli would be apparent as a left or right phase-shifting of the reference curve relative to the assay values.

Eleven samples from two whale sharks, aged 35 and 50 years based on growth band counts, were analyzed for $\Delta^{14}C$ (Table 2). assay values ranged between 15.1 and 70.0. Two of the samples, including one with the earliest date of formation (1962.5), were too depleted in $\Delta^{14}C$ (15.1 and 20.6) to have formed post-bomb (Figure 2), but no pre-bomb samples were identified. The remaining $\Delta^{14}C$ values all ranged between 40 and 70, which is consistent with a post-bomb year of formation. All samples were characterized by $\delta^{13}C$ values consistent with typical shark vertebrae of metabolic origin (mean = −13.6; SE = 0.4; Table 2).

All of the assay values aligned well with the reference chronologies (Figure 2A), with no obvious bias to one side or the other. Since errors in growth band counts would result in misalignment of the reference and assay values, the assay results indicate that the two sharks must have been aged correctly, at least on average. The 35-year old shark from Taiwan was least informative in this respect, since its post-bomb assay value indicated only that the shark could not have been over-aged by more than 10 years. Conversely, aging error of more than about 5 years would have been apparent as an obvious misalignment in the 50-year old Pakistan shark.
validation, but is often the only technique available to researchers. This approach has been criticized as a problematic form of age of marginal increment analysis, which gave inconsistent results. off the coast of Taiwan. Age validation was based on two forms analyzed the vertebrae of 92 whale sharks collected by a fishery. A second, more recent study by Hsu et al. (2014) assumed that two growth bands were formed each year. In their study, the oldest specimen (a male, 988 cm total length) had 42 growth bands and was thus assumed to be 21 years old. Perhaps more importantly, counts of growth bands by Hsu et al. (2014) were made in the intermedialia region, which contrasts to most other studies that use the corpus calcareum region of shark vertebrae for age interpretation (e.g., Campana et al., 2002; Christiansen et al., 2016). Our results, based on growth bands visible in the corpus calcareum region of sectioned vertebrae and validated with bomb radiocarbon assays, confirmed that growth bands must have formed annually, suggesting that the study of Hsu et al. (2014) overestimated growth rates of the species.

Our estimate of $k = 0.014$ year$^{-1}$ for whale sharks from the von Bertalanffy growth model was lower than the estimates provided by both Wintner (2000) and Hsu et al. (2014). The study by Wintner (2000) reported linear growth for 15 individuals, all of which were less than 8 m in length and under 30 years old. To constrain the growth curve, Wintner (2000) added two theoretical data points (60 and 100 years with 14 m TL) to obtain $k = 0.032$ or 0.021 year$^{-1}$, respectively, with $L_\infty$ of 13.7 m TL. The more recent study by Hsu et al. (2014) included the lengths of 3 full-term embryos and used a modified 2-parameter von Bertalanffy growth model to obtain two growth curves that were based on either biannual or annual deposition of growth bands. For annual growth bands, they reported estimates of $k = 0.021$ year$^{-1}$ with $L_\infty$ of 15.3 m TL. In our study, the predicted $L_\infty$ (21.9 m TL) was close to the largest maximum length ever recorded in the wild, estimated at 20 m from Taiwan in March 1987 (Chen et al., 2002) and close to maximum sizes recorded in other locations (Mcclain et al., 2015). This suggests that the growth models of both Wintner (2000) and Hsu et al., 2014 underestimated maximum sizes of whale sharks. We did, however, find a large difference between the growth coefficients of the von Bertalanffy and the logistic growth models, with the latter having higher growth coefficients but seeming to underestimate $L_\infty$. In this context, it is important to note that our dataset represents a small sample of individuals and only included two mature individuals, hence it is likely that the asymptotic length estimated in the von Bertalanffy growth model was poorly constrained and thus unrealistic. Actual growth parameters are probably bracketed by the results of the two growth models. Given the closure of the fishery in Taiwan and the protection of whale sharks in the waters of many of the countries where they occur (Rowat and Brooks, 2012), increased sample sizes are likely to rely on unfortunate but opportunistic events such as stranding (Wintner, 2000; Speed et al., 2009) to provide new vertebrae for analysis. Alternatively, photo-identification and imagery techniques may now offer a means to estimate in-situ growth rates for whale sharks, for at least the individuals and size classes participating in nearshore aggregations (e.g., Perry et al., 2018).

Our estimates of natural mortality for whale sharks, ranging from 0.09 to 0.14 year$^{-1}$ was close to those of other large species.
of sharks, such as the filter-feeding basking shark \((0.07 \text{ year}^{-1})\); Pauly, 2002; Campana et al., 2008), white \((0.08 \text{ year}^{-1})\;\text{ Mollet and Cailliet, 2002})\), and scalloped hammerhead \((0.10 \text{ year}^{-1})\;\text{ Cortés and Brooks, 2018})\) sharks. These estimates are generally considered low, however, for smaller whale sharks \((< 3 \text{ m TL})\), mortality rates may be higher, since the early juvenile stage is likely to be the most vulnerable to predators \((\text{ Rowat et al., 2008})\). Information on this life history stage is difficult to gather, because neonatal and very young whale sharks are only rarely encountered and are assumed to reside in the open ocean away from coasts \((\text{ Rowat et al., 2008})\).

Our estimates of slower growth and greater observed longevity have important implications for conservation of whale sharks. Underestimation of longevity and overestimation of growth is a serious concern for management strategies for fisheries, because it has led to population crashes due to overharvesting \((\text{ e.g., orange roughy; Smith et al., 1995})\). The case for whale sharks is somewhat different from other species that are targeted in fisheries, in part because they are protected across most of their distribution \((\text{ Bradshaw et al., 2008; Hsu et al., 2012})\). This status reflects the continuing rise and value of eco-tourism in sites where they aggregate, such as Ningaloo Reef in Western Australia \((\text{ Meekan et al., 2006})\). Although the harvesting of whale sharks has been reduced for over a decade, the sizes and abundances of populations have declined in multiple regions \((\text{ Theberge and Dearden, 2006; Bradshaw et al., 2008})\), which is reflected in the recent upgrade of the species from Threatened to Endangered by the IUCN Red List \((\text{ Pierce and Norman, 2016})\). Given the slow growth rates, extended longevity, late maturity and global connectivity of this species \((\text{ Bradshaw et al., 2007; Graham and Roberts, 2007; Sequeira et al., 2013})\), this species is likely to be highly susceptible to sources of anthropogenic mortality such as ship-strike \((\text{ Bradshaw et al., 2007; Speed et al., 2008})\). We are hopeful that the demographic data we have provided in this study will help to improve the accuracy of population models \((\text{ e.g., persistence, survival})\) and hence, better inform management and conservation efforts for this iconic species.

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**DATA AVAILABILITY STATEMENT**

All datasets generated for this study are included in the article/supplementary material.

**ETHICS STATEMENT**

This study was carried out in accordance with the guidelines of use of animal tissue/cadaver and the protocol was approved by the University of Western Australia Institutional Biosafety Committee.

**AUTHOR CONTRIBUTIONS**

MM and SC conceived of the idea, HH and LF provided the vertebral samples. JO did the sample preparation and analyses for vertebral aging. SC contributed to bomb radiocarbon results and verified aging results. JO wrote the manuscript, with critical feedback and help from SC, MM, LF, and HH.

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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.