Body length estimation of Neogene macrophagous lamniform sharks (*Carcharodon* and *Otodus*) derived from associated fossil dentitions

Victor J. Perez, Ronny M. Leder, and Teddy Badaut

ABSTRACT

The megatooth shark, *Otodus megalodon*, is widely accepted as the largest macrophagous shark that ever lived; and yet, despite over a century of research, its size is still debated. The great white shark, *Carcharodon carcharias*, is regarded as the best living ecological analog to the extinct megatooth shark and has been the basis for all body length estimates to date. The most widely accepted and applied method for estimating body size of *O. megalodon* was based upon a linear relationship between tooth crown height and total body length in *C. carcharias*. However, when applying this method to an associated dentition of *O. megalodon* (UF-VP-311000), the estimates for this single individual ranged from 11.4 to 41.1 m. These widely variable estimates showed a distinct pattern, in which anterior teeth resulted in lower estimates than posterior teeth. Consequently, previous paleoecological analyses based on body size estimates of *O. megalodon* may be subject to misinterpretation. Herein, we describe a novel method based on the summed crown width of associated fossil dentitions, which mitigates the variability associated with different tooth positions. The method assumes direct proportionality between the ratio of summed crown width to body length in ecologically and taxonomically related fossil and modern species. Total body lengths were estimated from 11 individuals, representing five lamniform species: *Otodus megalodon*, *Otodus chubutensis*, *Carcharodon carcharias*, *Carcharodon hubbelli*, and *Carcharodon hastalis*. The method was extrapolated for the largest known isolated upper tooth of *O. megalodon*, resulting in a maximum body length estimate of 20 m.

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INTRODUCTION

The order Lamniformes includes two of the most iconic shark species: the extant great white, *Carcharodon carcharias*, and the extinct megalodon shark, *Otodus megalodon* (Cappetta, 1987, 2012). *Carcharodon carcharias* is the largest macrophagous shark alive today, achieving maximum body lengths of arguably 6 to 7 m (Gottfried et al., 1996; Castro, 2012; McClain et al., 2015). *Otodus megalodon* represents the largest macrophagous shark that ever lived, with contentious adult estimates ranging from 10 to more than 30 m (Dean, 1909; Randall, 1973; Bendix-Almgreen, 1983; Gottfried et al., 1996; Pimiento et al., 2010; Pimiento and Balk, 2015; Trif et al., 2016; Grant et al., 2017; Razak and Kocsis, 2018; Shimada, 2002a, 2019; Herraiz et al., 2020). This longstanding interest in body size of the world’s largest marine macropredators has been shared by professional paleontologists and the public alike. Hypothetical models of *Otodus megalodon* are used as centerpiece exhibits in numerous natural history museums (e.g., the Calvert Marine Museum, Florida Museum, Smithsonian National Museum of Natural History, Western Australian Museum). Likewise, popular science fiction has featured these taxa, with vastly overestimated body sizes (e.g., *Jaws* and *The Meg*). Constraining body size estimates is imperative if we are to accurately represent these taxa and decipher the evolutionary mechanisms that produced these enormous macropredators.

The chondrichthyan fossil record is largely restricted to teeth, due to the high preservation potential of hydroxyapatite and poor preservation potential of cartilage and soft tissue (Kent, 1994). Consequently, interpretation of chondrichthyan evolutionary history is largely restricted to what we can infer from dental data and modern analogs. To estimate the size of *Otodus megalodon*, a number of researchers have attempted to relate tooth height to total body length in the extant great white shark as a proxy (Gottfried et al., 1996; Pimiento et al., 2010; Pimiento and Balk, 2015; Reolid and Molina, 2015; Trif et al., 2016; Grant et al., 2017; Razak and Kocsis, 2018; Shimada, 2002a, 2019). Gottfried et al. (1996) estimated body size based on the relationship between total tooth (root and crown) height (TH) of the second upper anterior (A2; see Figure 1A for dentition terminology) and total body length (TL), resulting in a conservative maximum adult size of 15.9 m for *O. megalodon*. Gottfried et al. (1996) speculated the TL of *O. megalodon* throughout ontogeny by scaling up TL and body mass in extant *Carcharodon carcharias* individuals from different ontogenetic stages. Pimiento et al. (2010) used this to define three life stages for *O. megalodon*: neonate (TL < 4 m), juvenile (TL = 4 to 10.5 m), and adult (TL > 10.5 m).

Shimada (2002a, 2019) suggested that growth rates in the crown and root are not isometric, and instead developed a method based on crown height (CH). Shimada (2002a) used a linear regression to determine the relationship between CH and TL for each tooth position in *C. carcharias*. It was then proposed that these equations may be an appropriate analog to estimate body size of *O. megalodon*, given that *C. carcharias* represents the largest extant macrophagous shark. All subsequent studies (Pimiento et al., 2010; Pimiento and Balk, 2015; Reolid and Molina, 2015; Trif et al., 2016; Grant et al., 2017; Razak and Kocsis, 2018; Herraiz et al., 2020) mentioned have since used the Shimada (2002a) equations to estimate the size of *O. megalodon*. However, K-12 students applied Shimada’s (2002a) equations to a 3D printed model of an associated dentition (i.e., multiple teeth from a single individual) of *O. megalodon* (UF-VP-311000), resulting in a range of body length estimates from 12 to 45 m for this single individual (Grant et al., 2017). This prompted the authors of this study to re-evaluate the Shimada (2002a) method and to develop the novel method described herein.

Since then, Shimada (2019) also re-evaluated the original linear equations proposed in 2002 and took a closer look at the relationship between CH and TL of anterior tooth positions in *Carcharodon carcharias*. In this study, Shimada determined that a power function results in a greater correlation between CH and TL than a linear regression. Further, Shimada determined that upper anterior teeth...
FIGURE 1. Premise of summed crown width method. (A) Carcharodon carcharias dentition in lingual view, with applicable terminology. The right half is an illustration of the typical dental formula for C. carcharias. The left half is from a 5.18 m female with one less posterior tooth in the lower tooth series (originally figured in Hubbell, 1996; figure 5). Scale bar equals 5 cm. (B) The most complete known associated dentition of Otodus megalodon (CH-31-46P) in lingual view. Scale bar equals 5 cm. (C) Body length of fossil taxa is calculated under the assumption that the ratio of summed crown width to total body length (TL) is proportional in ecologically and taxonomically related species. Silhouette proportions for O. megalodon are based on Cooper et al. (2020). A/a = anterior, I = intermediate, and L/l = lateral. Uppercase letters denote upper teeth and lowercase letters denote lower teeth.
resulted in more reliable estimates than lower teeth, and that the second upper anterior (A2) tooth position provides more reliable estimates than the first upper anterior (A1) tooth position. Shimada et al. (2020) analyzed dental homology in macropaghous lamniforms based on the typical “lamnoid tooth pattern.” In this dental pattern, anterior teeth are always the tallest (Applegate, 1965; Shimada, 2020). Shimada (2019) used the isolated O. megalodon tooth with the largest CH in a museum collection (NSM PV-19896; CH=120 mm) to estimate a maximum body size, assuming it represents tooth position A1-A2, resulting in a TL of 14.2 to 15.3 m.

These results have clear implications that applying the Shimada (2002a) method to a sample of isolated teeth could be biased by estimates derived from different tooth positions. Whether one attempts to apply the Shimada (2002a) or Shimada (2019) method to estimate body size, both require identification, or at least approximation, of tooth positions from isolated fossil teeth. Identifying tooth positions requires knowledge of a taxon’s dental formula, which defines the total number of teeth and how they are arranged within a dentition. Two forms of evidence have been used to infer the dental formula of the megatooth lineage, each of which has pros and cons. Extant macrophagous sharks offer a source of indirect evidence, in which the dental formula of the extinct megatooth lineage is assumed to reflect that of the closest living analogs (i.e., the lamnoid tooth pattern). The fossil record offers the only direct evidence of the megatooth lineage but consists primarily of isolated teeth (Cappetta, 2012). Rare disarticulated, associated dentitions represent the most holistic direct evidence available to reconstruct the dental formula of the extinct megatooth lineage (Uyeno et al., 1989; Purdy et al., 2001; Pimiento et al., 2010; Perez et al., 2019). Consequently, all reconstructions of the dental formula in the megatooth lineage represent hypothetical models (Welton and Farish, 1993; Shimada, 2005). The various dental models proposed are fundamentally important for understanding potential biases in body length estimates of the megatooth lineage. Proposed dental models for the megatooth lineage are intrinsically linked to their taxonomic assignment.

**Megatooth Shark Taxonomy**

The taxonomy of *Otodus megalodon* has changed frequently and even now a complete consensus has not been reached. *Otodus megalodon* was originally assigned to the genus *Carcharodon*, under the assumption that it was related to *Carcharodon carcharias*, and was placed in the family Lamnidae (Agassiz, 1833-1844; Applegate and Espinosa-Arrubarrena, 1996; Gottfried et al., 1996; Purdy, 1996; Purdy et al., 2001). Other researchers have proposed that *O. megalodon* evolved as a separate lineage derived from *Otodus obliquus*, placing it in the family Otodontidae (Casier, 1960; Glickman, 1964; Zhelezko and Kozlov, 1999; Ward and Bonavia, 2001; Nyberg et al., 2006; Cappetta, 2012; Ehret et al., 2012; Shimada et al., 2017; Perez et al., 2019). Three alternative generic names had been proposed: *Carcharocles* (Jordan and Hannibal, 1923), *Procarcharodon* (Casier, 1960), and *Megaselachus* (Glickman, 1964). Consequently, many researchers began to use the genus *Carcharocles*, given that it had precedence over *Procarcharodon* and *Megaselachus*, to describe all serrated taxa within the megatooth lineage (i.e., *C. auriculatus*, *C. angustidens*, *C. chubutensis*, and *C. megalodon*).

The presence of serrations represents a convenient and easily distinguishable character to delineate *O. obliquus* from its descendants; however, Cappetta (2012) argued that serrations were not a sufficient character to warrant a separate genus and instead used *Otodus* for the entire lineage. As somewhat of a compromise, Cappetta (2012) proposed the use of *Carcharocles* and *Megaselachus* as subgenera to describe serrated taxa within the megatooth lineage. Shimada et al. (2017) also argued for the use of *Otodus* for the entire megatooth lineage to achieve monophyly, under the assumption that *Parotodus* and *Megalolamna* were sister taxa to *Otodus* within the family Otodontidae. However, Shimada et al. (2017) acknowledged that this shifts the issue of paraphyly to the ancestor of *O. obliquus*, *Cretalamna*, and that paraphyletic taxa are inevitable when attempting to place Linnaean taxa onto a phylogenetic tree.

Kent (2018) noted that the debate between the use of *Carcharocles* versus *Otodus* hinges on whether the lineage evolved through cladogenesis versus anagenesis and argued that the fossil record of megatooth sharks may not be sufficiently detailed to demonstrate anagenesis. Whether the megatooth lineage evolved through a series of rapid branching events (i.e., cladogenesis) or through the slow accumulation of mutations over time (i.e., anagenesis) is somewhat irrelevant to this study. For now, we will adopt the popular use of *Otodus* to describe the megatooth sharks as a chronospecific lineage. Rather, what is relevant is
that we agree that *O. megalodon* evolved from *O. obliquus* and represents a separate lineage from that of *C. carcharias* (Figure 2).

**The Megatooth Dental Formula**

Lamniform sharks typically exhibit heterodonty known as the "lamnoid tooth pattern," which consists of both monognathic and dignathic heterodonty (Shimada, 2002b; Shimada et al., 2020). In macrophagous lamniforms, upper monognathic heterodonty is generally characterized by two rows of anterior teeth on either side of the jaw symphysis, followed by one or more intermediate tooth rows, and a series of lateral teeth. The anterior teeth are vertically erect and generally represent the tallest teeth in the dentition. The intermediate teeth are significantly smaller, interrupting the general pattern of the tallest teeth being mesial positions, smallest teeth being distal positions, relative to the jaw symphysis. Shimada (2002b) referred to the first three tooth positions (i.e., A1, A2, and 11) as "index tooth rows" to identify dental homologies. However, it is possible for a shark to modify their dentition by either losing or gaining a tooth row. Further, there is intraspecific variability in the lamnoid tooth pattern (Hubbell, 1996).

Applegate and Espinosa-Arrubarrena (1996; figure 1a) depicted *C. carcharias* with 46 teeth in its functional series. In this model, a complete half of the dentition has an upper tooth series with two anterior, one intermediate, five lateral, and four posterior teeth and a lower tooth series with three anterior, five lateral, and three posterior teeth. In Shimada (2002a; Figure 1), *Carcharodon carcharias* has a functional series consisting of 50 teeth. In this model, a complete half of the dentition has an upper tooth series with two anterior, one intermediate, and 10 latero-posterior teeth and a lower tooth series with two anterior, one intermediate, and nine latero-posterior teeth. Based on dentitions observed by the authors of this study, we assume that the typical dental formula for *C. carcharias* is in between both models, with 48 teeth in the first functional series (Figure 1A).

Shimada (2005) defined a spectrum of preservation states for fossil shark teeth, ranging from common isolated teeth to rare associated dentitions. Isolated tooth sets (i.e., composite or artificial dentitions) are comprised of teeth from multiple individuals from the same taxon and represent a completely fabricated dental model. Associated tooth sets (i.e., associated dentitions) represent multiple teeth belonging to a single individual, which can be found either in articulation or disarticulated. Articulated dentitions provide a true dental model. Disarticulated dentitions require inferences to reassemble the true dental model but provide context for making those inferences. Six disarticulated, associated dentitions of *O. chubutensis* and *O. megalodon* have been referenced in peer-reviewed literature (Uyeno et al., 1989; Purdy et al., 2001; Pimiento et al., 2010; Grant et al., 2017; Perez et al., 2019). To date, no one has reported a verified articulated, associated dentition of *O. chubutensis* or *O. megalodon*.

Applegate and Espinoza-Arrubarrena (1996) created artificial dentitions of *Otodus/Carcharodon* using isolated tooth sets, in which they proposed a dental formula for *O. megalodon* comprised of 58 teeth in the functional series. In this model, a complete half of the dentition consisted of an upper functional series with two anterior, one intermediate, seven lateral, and four posterior teeth and a lower functional series with three anterior, eight lateral, and four posterior teeth. Purdy et al. (2001) figured partial reconstructions of two disarticulated, associated dentitions of *Otodus chubutensis* (USNM 411881 and USNM 299832) and one disarticulated, associated dentition of *Otodus megalodon* (NCSM 13073) from the Lee Creek Mine in Aurora, North Carolina. These reconstructions all included a medially-oriented upper intermediate tooth, based on an assumed evolutionary relationship with *Carcharodon carcharias*.

Pimiento et al. (2010; figure S1) provided a dental model for *Otodus megalodon* consisting of 46 teeth in its functional series, with a complete half of the dentition comprised of an upper functional series with three anterior and nine latero-posterior teeth and a lower functional series with three anterior and eight latero-posterior teeth. This was the first dental model for *Otodus megalodon* that lacked any intermediate tooth positions. It is unclear if tooth position A3 in *O. megalodon* should...
be considered an intermediate tooth, but the proportions are certainly not analogous to I1 in C. carcharias (Figure 1). Pimiento et al. (2010) stated that this dental model was adapted from Gottfried et al. (1996), which presumably refers to the composite, artificial dentition of O. megalodon on exhibit at the Calvert Marine Museum (see Gottfried et al., 1996; figure 1B). Gottfried et al. (1996) stated that the figured O. megalodon dentition was casted from the original Smithsonian composite dentition “… prior to the discovery of important natural tooth sets at Lee Creek Mine, North Carolina and in Florida that help clarify the arrangement of teeth in C. megalodon.”

The dental model for Otodus megalodon in Pimiento et al. (2010; figure S1) originated, at least in part, from dental reconstructions based on the fossil associated dentitions that Gottfried et al. (1996) had alluded to. Pimiento et al. (2010; tables S1 and S2) referred to these specimens as “juvenile Carcharocles megalodon associated tooth set” and “adult Carcharocles megalodon associated tooth set” and reported crown height (CH) and crown width (CW) measurements for each tooth position. The juvenile individual was recovered by L. Martin and G. Hubbell in 1995 from the Bone Valley region in Florida (herein referred to as CH-31-46P, Figure 1B), and the adult individual was recovered by A. Felt in 1996 from the Lee Creek Mine in North Carolina (herein referred to as UF-VP-311000). Both disarticulated, associated dentications were originally reconstructed by G. Hubbell. Images of CH-31-46P can be found on www.elasmo.com (Bourdon, 2005; Razak and Kocsis, 2018), and a 3D model of UF-VP-311000 is figured in Grant et al. (2017; figure 1). The Bone Valley dentition (CH-31-46P) has been referenced in exhibits at the Florida Museum, Smithsonian National Museum of Natural History, and Western Australian Museum.

Pimiento et al. (2010; figure 4) used the CH:CW ratio in these associated denticions to approximate tooth positions of isolated O. megalodon teeth from the Gatun Formation. Some teeth were assigned to an exact tooth position, while others were given a range of potential tooth positions. Pimiento et al. (2010) then used these tooth positions to estimate body size with the equations reported by Shimada (2002a) and determined that the Gatun Formation represented a paleo-nursery for O. megalodon. Pimiento and Balk (2015) repeated this process to analyze spatial and temporal macroevolutionary body size trends in O. megalodon. This method has since propagated into numerous studies of O. megalodon (Reolid and Molina, 2015; Trif et al., 2016; Grant et al., 2017; Razak and Kocsis, 2018; Herraiz et al., 2020).

Given that fossil associated dentitions were influential in predicting the dental formula for Otodus megalodon, it seems logical that we should test the accuracy of body length estimates using these associated dentitions as hypothetical models. Further, preliminary results reported in Grant et al. (2017) indicated that the Shimada (2002a) method results in highly variable body length estimates. As such, this study analyzes variability in body length estimates by applying the dental model proposed by Pimiento et al. (2010) to fossil associated dentitions. In addition, a novel method for estimating TL is proposed based on this dental model.

Body Length Estimation using Jaw Width

Methods for estimating body length based on TH, CH, and CW all follow a similar rationalization that teeth grow proportionately with total body length. Reolid and Molina (2015; figure 4) compared the linear relationship between different crown measurements in the second upper anterior tooth position (A2) and total body length (TL) in Carcharodon carcharias, reporting a greater correlation for CH than CW (r=0.92 for CH and r=0.53 for CW). However, after re-calcultating the Pearson coefficient of correlation (r) from the Reolid and Molina (2015) dataset, the correlation should be r=0.77 for CW (n=58) and r=0.93 for CH (n=85). While this still indicates that CH is a better predictor of body length than CW, it does not consider other tooth positions or summed crown width (SCW) of associated teeth as a proxy for body length. In addition, there has been inconsistency in the literature regarding tooth position nomenclature. Hubbell (1996) referred to the first two anterior teeth of C. carcharias as A2 and A3, due to a hypothesis from Applegate and Espinosa-Arrubarrena (1996) that C. carcharias had lost an anterior tooth row during its evolution. Consequently, Reolid and Molina (2015) included data from Hubbell (1996) that actually refers to tooth position A1 and combined this with data from other studies for tooth position A2.

Lowry et al. (2009) studied the relationship between jaw circumference and total body length in 14 shark species, as a means of estimating body size from shark bite marks. This forensic method allows researchers to estimate the body size of individuals involved in shark attacks. Lowry et al.
used a metric referred to as the interdental distance (IDD), which is measured as the distance between the crown apex of two neighboring teeth, as a proxy for jaw circumference. A linear regression of IDD versus TL for *C. carcharias* resulted in $R^2=0.98$ for the upper jaw and $R^2=0.97$ for the lower jaw. Siversson (2012) first emphasized the potential of applying this method to estimate body size in fossil taxa, specifically the extinct Cretaceous lamniform *Cardabiodon ricki*. Newbrey et al. (2015) applied the jaw circumference method to a partially articulated, associated dentition of *C. ricki*, resulting in a body length estimate of 5.5 m. This method has a strong foundational premise but does not offer a way to account for missing teeth in fossil dentitions and requires assumptions regarding interdental spacing.

This study introduces a novel method for estimating body size in fossil lamniform sharks that builds on the concept that jaw circumference is proportional to total body length. A linear regression of IDD versus TL for *C. carcharias* resulted in an $R^2=0.98$ for the upper jaw and $R^2=0.97$ for the lower jaw. Siversson (2012) first emphasized the potential of applying this method to estimate body size in fossil taxa, specifically the extinct Cretaceous lamniform *Cardabiodon ricki*. Newbrey et al. (2015) applied the jaw circumference method to a partially articulated, associated dentition of *C. ricki*, resulting in a body length estimate of 5.5 m. This method has a strong foundational premise but does not offer a way to account for missing teeth in fossil dentitions and requires assumptions regarding interdental spacing.

This study introduces a novel method for estimating body size in fossil lamniform sharks that builds on the concept that jaw circumference is proportional to total body length. Such that, the summed crown width (SCW) in the functional series is constrained by jaw size. Based on a simple mathematical concept referred to as the rule of three, this method assumes direct proportionality between the ratio of SCW to TL in related modern and fossil taxa (Figure 1C). Herein, we apply this method to fossil associated dentitions from five macrophagous lamniform species within the families Otodontidae and Lamnidae: *Otodus megalodon*, *Otodus chubutensis*, *Carcharodon carcharias*, *Carcharodon hubbelli*, and *Carcharodon hastalis*.

**MATERIALS AND METHODS**

**Modern and Fossil Dentitions**

Data was collected from the first functional series in 11 fossil associated dentitions of lamniform sharks (Table 1) and 19 modern lamniform dentitions (Table 2). These specimens are repository in the Florida Museum (UF-VP), Gordon Hubbell Collection (GHC), and the National Museum of Natural History of the Smithsonian Institution (USNM). Specimens in the GHC are gradually being acquisitioned into the Florida Museum vertebrate paleontology collection and will remain accessible for research throughout the process. Provenance information for these specimens was derived from each museum’s collections database and/or personal communication with the museum’s respective collections manager (i.e., R. Hulbert, G. Hubbell, and D. Bohaska).

Four associated dentitions and a single isolated tooth of *Otodus megalodon* were included in this study. UF-VP-311000 consists of 37 teeth (22 from the first functional series) collected by A. Felt in 1996 within the Nutrien Corp phosphate mine (commonly known as the Lee Creek Mine) in Aurora, North Carolina (Figure 3A). 3-D files of UF-
TABLE 2. Specimen data for 19 modern dentitions. SCW = summed crown width. TL = total body length. Measurements are separated for the different regions preserved: upper left (UL), upper right (UR), lower left (LL), and lower right (LR).

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<th>LL</th>
<th>LR</th>
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VP-311000 are freely available on MorphoSource.org. GHC 1 consists of 92 teeth (36 from the first functional series) found by M. Marinela-rena in 2003 within the phosphate mining region of the Atacama Desert in northern Chile (Figure 3B). CH-31-46P consists of 95 teeth (44 teeth from the first functional series) found by L. Martin and G. Hubbell in 1995 within the Four Corners phosphate mine in Polk County, Florida (Figure 1B and 3C). The physical specimen resides in the GHC; although, reproductions can be purchased from Bone Clones under the code CH-31-46P (we use this as a temporary identifier until the specimen is assigned a globally unique identifier). UF-VP-460000 consists of 43 teeth (30 from the first functional series) found by J. Arias in 2003 within the phosphate mining region of the Atacama Desert in northern Chile (Figure 3D). GHC 6 represents the largest isolated tooth of *O. megalodon* known to the authors of this study (TH=165 mm, CH=122 mm, CW=133 mm). This specimen was found by V. Bertucci while diving in the Morgan River in South Carolina.

Four associated dentitions of *Otodus chubutensis* were included in this study. USNM 411881 consists of 106 teeth (38 from the first functional series) found by P. Harmatuk in 1986 within the Nutrien Corp phosphate mine in Aurora, North Carolina (Figure 4A). USNM 299832 consists of 32 teeth (20 from the first functional series) found by P. Harmatuk in 1981 within the Nutrien Corp phosphate mine in Aurora, North Carolina (Figure 4B). GHC 3 consists of 65 teeth (31 from the first functional series) found by G. Hubbell in 1993 on the western bank of the Ica River in Peru (Figure 4C). UF-VP-312864 consists of 53 teeth (35 from the first functional series) collected in 2002 from the Ica River in Peru (Figure 4D).

Three associated dentitions of *Carcharodon* spp. were included in this study, all from Sacaco, Peru. GHC 4 is a *C. carcharias* dentition consisting of 65 associated teeth (32 from the first functional series) collected by C. Martin in 1988 (Figure 5A).
GHC 5 is a *C. hastalis* dentition consisting of 165 teeth (40 from the first functional series) collected by C. Martin in 1988 (Figure 5B). UF-VP-226255 is a partially articulated skeleton of *C. hubbelli* consisting of 222 teeth, 45 centra, and portions of the cartilaginous mandibular arch and neurocranium (see Ehret et al., 2009, 2012).

To date, no completely articulated dentitions of *O. megalodon* nor *O. chubutensis* have been reported. Consequently, reconstructions of the fossil dentitions represent hypothetical models that can only be verified by the discovery of fully articulated dentitions of each species being analyzed. As such, the dental patterns utilized reflect a major assumption that is fundamental to the body length method described in this study.

Among the 19 modern lamniform dentitions, measurements were recorded from 17 *Carcharodon carcharias* individuals, one *Isurus oxyrinchus* individual, and one *Isurus paucus* individual. All modern specimens originated from the GHC collection. Specimens that were physically available were measured using calipers, whereas specimens that were figured in Hubbell (1996) were measured digitally using ImageJ. See Table 2 for additional specimen information.

**Dental Measurements**

Crown width (CW) and crown height (CH) were measured for every tooth position in the 19 modern dentitions (Appendices 1 and 2) and 11 fossil dentitions (Appendices 3 and 4). CW is defined as a line segment between the crown-root contact on the mesial and distal cutting edges. CH is defined as a perpendicular line segment from the crown apex to the CW line segment. For nine of the 19 modern analogs, authors VJP and RML independently measured CH and CW. These measure-
Dentitions were segmented into four regions (upper left, upper right, lower left, and lower right). Summed crown width was measured for each region of the dentitions. The summed crown width of all teeth present in the dentition is referred to as SCW\textsubscript{ms}. To account for missing teeth in the fossil dentitions, correction factors (CF) were determined by calculating the percent of the total summed width attributable to each tooth position in the modern and fossil dentitions (Figure 6 and Appendix 5). The mean percentage for each tooth position of modern *Carcharodon carcharias* was applied as a correction factor to estimate the width of missing teeth in the fossil dentitions of *Carcharodon* spp. Missing teeth in the *Otodus* spp. dentitions were corrected using percentages from the most complete, comparable fossil *Otodus* dentition (Appendix 5). Corrected summed crown width (SCW\textsubscript{c}) is calculated to account for missing teeth by dividing the measured summed crown width (SCW\textsubscript{ms}) by one minus the sum of all correction factors (Eq. 1). Summed crown width, measured and corrected, are provided in Table 3 for all fossil dentitions.

\[
SCW_c = \frac{SCW_{ms}}{1 - \sum CF}
\]

*Eq. 1*

**Body Length Estimation**

Body length estimates were calculated under a basic assumption that the ratios of summed
crown width (SCW) to body length (TL) are proportional in taxonomically and ecologically related species (Eq 2). The rule of three method assumes direct proportionality between the ratio of summed crown width \((a = SCW_m)\) to body length \((b = TL_m)\) in modern taxa (e.g., *C. carcharias*, *I. oxyrinchus*, and *I. paucus*) and the ratio of summed crown width \((c = SCW_f)\) to body length \((x = TL_f)\) in fossil taxa (e.g., *O. megalodon*, *O. chubutensis*, *C. carcharias*, *C. hubbelli*, and *C. hastalis*). Such that,

\[
\frac{a}{b} = \frac{c}{x}
\]

**FIGURE 5.** Associated dentitions of *Carcharodon* spp. in lingual view. (A) *Carcharodon carcharias*, GHC 4. (B) *Carcharodon hastalis*, GHC 5. Scale bars equal 5 cm. See Ehret et al. (2009) for images of the *Carcharodon hubbelli* dentition, UF-VP-226255.

**FIGURE 6.** Portion of summed crown width (SCW) for each tooth position. (A-B) Proportions from 10 modern *Carcharodon carcharias* dentitions. (C-D) Proportions from all eight fossil *Otodus* associated dentitions, after applying correction factors. (A, C) upper and (B, D) lower. Correction factors were derived from these proportions (see Appendix 5).
there are four variables, of which three can be measured \((a, b, \text{ and } c)\) and the fourth solved for \((x)\):

\[
\frac{a}{b} = \frac{c}{x} \Rightarrow \frac{SCW_m}{TL_m} = \frac{SCW_i}{TL_i}
\]

**Eq. 2**

**Table 3.** Body length estimates from 11 fossil dentitions of *Otodus* and *Carcharodon*. Summed crown width measured (SCWms) and corrected (SCWc), and total body length (TL). * denotes the best estimate. UL = upper left, UR = upper right, LL = lower left, and LR = lower right.

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<th>SCWc (mm)</th>
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interspecific and intraspecific variation in the relationship between SCW and TL. The rule of three method was applied for each region of the fossil dentitions preserved (upper left, upper right, lower left, and/or lower right) with all 19 modern analogs. Among the 19 modern analogs, eight only had half of the dentition preserved (left or right). Thus, for a single fossil dentition with all four quadrants preserved, there were a total of 120 different body length estimates. The range and mean body length estimates from each region of the jaw were calculated (Eq. 3). This equation is similar to a linear regression, where the calculated mean approximates the linear trendline and the range captures the extent of variance around the mean. The benefit of calculating TL with this equation is the ability to quickly compare estimates derived from specific, individual modern analogs and better explain the source of variance around the mean.

\[ TL = \text{mean}(x_1 \cdot \frac{b_1}{a_1}, x_2 \cdot \frac{b_2}{a_2}, \ldots, x_n \cdot \frac{b_n}{a_n}) \]  

Eq. 3

Extrapolation for Isolated Teeth

The correction factors that were calculated using the modern dentitions of Carcharodon carcharias and the most complete fossil associated dentitions of Otodus spp. were used to estimate body size from isolated teeth. To extrapolate the SCW method in this manner, one must first determine which associated dentition to use as an analog to approximate the appropriate tooth position. For example, if you wish to estimate the size from an isolated O. megalodon tooth, then you could use either the adult individual (UF-VP-311000) or the juvenile individual (CH-31-46P) as your analog to do so. If you are uncertain which is the best analog, then body length should be calculated using both models and an average estimate should be reported. If a tooth cannot be identified to a specific tooth position, then a range of potential positions and corresponding body lengths should be reported.

This extrapolated version of the SCW method offers the opportunity to re-assess body size esti-
mates in previous paleoecological studies (i.e., Pimiento et al., 2010; Pimiento and Balk, 2015; Herraiz et al., 2020) and compare them with estimates derived from the Shimada (2002a, 2019) isolated CH method. Body length estimates were re-calculated using a subset of the Pimiento and Balk (2015) sample. The SCW method was applied by using proportions from CH-31-46P and UF-VP-311000 separately. The average of these two estimates was compared to those reported by Pimiento and Balk (2015) to determine if there was a predictable variation in body length estimates based on the tooth position assigned.

The SCW method was also extrapolated to estimate the maximum body size of *Otodus megalodon*, using the isolated tooth with the widest known crown width (GHC 6; CW=133 mm). The tooth position was assigned based on a comparison of the CH:CW ratios in the largest associated dentition of *O. megalodon*, UF-VP-311000. The corrected SCW was calculated based on the CW proportions in UF-VP-311000.

**RESULTS**

Segmenting the dentitions into four regions (upper left, upper right, lower left, and lower right) allowed for the inclusion of partial associated dentitions, comparison of estimates derived from the lower versus upper dentitions, and comparison of estimates derived from the left versus right sides of the dentitions. However, this also required determination as to which of the four regions provided the most reliable body length estimate. In all taxa, bilateral symmetry is generally observed; however, this symmetry is often imperfect. Body length estimates derived from the left and right sides of a single individual varied from 0 to 0.4 m (Table 3). In terms of selecting left versus right, the more complete side of the fossil dentitions is considered the more reliable estimate.

For all individuals, the mean body length estimates derived from the lower dentition were greater than estimates derived from the upper dentition (Table 3 and Figure 7). For the upper dentition, estimates using *Isurus* as the modern analog were greater than estimates derived from *Carcharodon*. For the lower dentition, estimates using *Isurus* as the modern analog were within the range of estimates derived from *Carcharodon*. These differences between the upper and lower dentitions and different modern analogs can be explained by varying tooth morphologies attributed to different functional roles and feeding ecologies (see Discussion below). Thus, the mean TL from the most complete upper half of the dentition, with *Carcharias* as the modern analog, is considered the most reliable estimate (Table 3).

For the four individuals of *O. megalodon*, mean body lengths estimates were 10.9, 12.4, 17.3, and 17.3 m, respectively. Using the same modern comparative dataset, the body length estimates for the four individuals of *O. chubutensis* were 5.9, 9.0, 10.9, and 11.0 m, respectively. Only a single individual of *Carcharias, C. hubbelli*, and *C. hastalis* were used in this study, which resulted in body length estimates of 4.0, 4.9, and 5.1 m, respectively.

The single individual of *Carcharodon hubbelli*, UF-VP-226255, was the only fossil specimen found as an articulated skeleton, comprised of 222 teeth and 45 vertebral centra. Ehret et al. (2009) applied body length equations based on regressions of vertebral diameter and vertebral radius, proposed by Cailliet et al. (1985), Gottfried et al. (1996), Wintner and Cliff (1999), and Natanson (2001). These body length estimates based on vertebral measurements ranged from 4.7 to 5.2 m, with a mean estimate of 4.9 m. Ehret et al. (2009) also applied the Shimada (2002a) method, resulting in a range of estimates from 3.0 to 7.0 m, with a mean estimate of 5.1 m. The SCW method resulted in a range of estimates from 4.1 to 5.8 m, with a mean estimate of 4.9 m.

The Shimada (2002a, 2019) methods were applied to the 11 fossil dentitions for comparison (Figure 7 and Table 4). Body length estimates based on the Shimada (2002a) method resulted in a wider range of estimates than the SCW method described herein. For *Otodus* individuals, body length estimates based on the Shimada (2002a) method resulted in a range of estimates from 3.0 to 7.0 m, with a mean estimate of 5.1 m. The SCW method resulted in a range of estimates from 4.1 to 5.8 m, with a mean estimate of 4.9 m.

For all individuals, the mean body length estimates derived from the lower dentition were greater than estimates derived from the upper dentition (Table 3 and Figure 7). For the upper dentition, estimates using *Isurus* as the modern analog were greater than estimates derived from *Carcharodon*. For the lower dentition, estimates using *Isurus* as the modern analog were within the range of estimates derived from *Carcharodon*. These differences between the upper and lower dentitions and different modern analogs can be explained by varying tooth morphologies attributed to different functional roles and feeding ecologies (see Discussion below). Thus, the mean TL from the most complete upper half of the dentition, with *Carcharias* as the modern analog, is considered the most reliable estimate (Table 3).

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Extrapolating the SCW method for the largest known isolated tooth of *O. megalodon* (GHC 6) resulted in a corrected summed crown width ranging from 1266.7 to 1303.9 mm. Based on a SCW of 1266.7 mm, the SCW method results in a TL of 17.4 to 23.5 m, with a mean estimate of 20.0 m. Based on a SCW of 1303.9 mm, the SCW method results in a TL of 18.0 to 24.2 m, with a mean estimate of 20.6 m.
### TABLE 4. Comparison of different methods for estimating body length: Shimada (2002a), Shimada (2019), and this study. All estimates are in meters. * denotes the best estimate. UL = upper left, UR = upper right, LL = lower left, and LR = lower right.

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DISCUSSION

Accounting for Error

There are a few different potential sources of error that could impact body length estimates using the SCW direct proportionality method. These sources can be broadly grouped into three categories: natural variation, human observer error, and mathematical assumptions. Natural variation in tooth proportions occurs within individuals, within species, and between species, which could introduce error to our body length estimates. Human error could be introduced while measuring teeth or through assumptions related to the dental count and pattern of fossil dentitions. Assumptions regarding the optimal mathematical equation introduce yet another avenue for error. Accounting for each source of error offers constraints on the potential accuracy of body length estimates proposed in this study and in previous studies.

Natural variation. Variation within individuals was accounted for by comparing estimates between the four quadrants in the dentition. Given that all individuals exhibit bilateral symmetry, there is minimal variation between the left and right sides of the dentition (i.e., < 0.5 m difference in TL estimates). This was accounted for by simply selecting the side of the fossil dentition that was most completely preserved as the superior estimate.

Greater variability was observed in estimates derived from the upper versus the lower dentitions, in which the lower dentition always resulted in larger TL estimates than the upper dentition (Figure 7). This consistent difference in body length estimates derived from upper and lower dentitions can be explained by two factors: function and phylogeny. There is a difference in shape and corresponding functionality between the lower and upper dentition (Kent, 1994; Cappetta, 2012). Lower teeth tend to be narrower and have a more robust root relative to upper teeth, which aids in penetrating and grasping prey. After the lower teeth have anchored the prey, the upper teeth then tear and cut the flesh. While both upper and lower teeth are predominantly designed for cutting, there is greater grasping functionality in lower teeth than in upper teeth. Further, this grasping functionality of the lower teeth is more apparent in representatives of the genus *Carcharodon* than those in the genus *Otodus*. *Carcharodon* spp. tend to have more symmetrical, vertically erect lower teeth than *Otodus* spp.

Within the extant comparative dataset of *C. carcharias*, SCW versus TL was plotted for the upper and lower dentitions separately (Figure 8). Both showed a strong correlation; however, the coefficient of determination between upper SCW and TL was stronger than that of the lower SCW and TL (upper: $R^2=0.9326$ and lower: $R^2=0.8977$). Thus, we concluded that the most reliable body length estimates are from the most complete upper half of the dentition. The determination that the upper dentition is more reliable than the lower dentition is in agreement with the TH method employed by Gottfried et al. (1996) and the CH method proposed by Shimada (2019). It seems plausible that the offset in TL estimates from the upper and lower jaws may be predictable and, thus, a correction factor could be calculated. Although, determining if this is the case is a task for a future research endeavor and beyond the intended scope of this manuscript.

In addition to segmenting the jaw, three modern analogs within the family Lamnidae were con-
Considered: *C. carcharias*, *Isurus oxyrinchus*, and *Isurus paucus*. Body length estimates derived from the upper jaw of *Isurus* spp. resulted in larger body length estimates than those derived from *C. carcharias*. For the lower dentition, estimates using *Isurus* spp. as the modern analog were within the range of estimates derived from *C. carcharias*. While all three are macrophagous lamniform sharks, they express very different dentition types that correspond to different feeding ecologies. *Isurus* spp. have narrower, non-serrated teeth that serve to feed on fast swimming fishes, as opposed to the more broadly triangular, serrated teeth of *C. carcharias* that are better adapted to feeding on fleshier marine mammals (Kent, 1994; Cappetta, 2012). Similar to the functional argument that explains the differences between upper and lower estimates, the narrower teeth of *Isurus* spp. also result in relatively larger body length estimates. All the fossil taxa analyzed in this study have more broadly triangular teeth that are functionally more similar to *C. carcharias* than *Isurus* spp.

These differences in body length estimates between different parts of the jaw and different modern analogs highlight the importance of selecting an ecologically similar modern analog to estimate body size. This is particularly important when attempting to estimate the size of taxa that are even more distantly related (e.g., *Otodus obliquus*). Cooper et al. (2020) stated that “*Lamna nasus* is considered the best dental analogue for both *Cretalamna* and *Megalolamna*, *mako sharks* (*Isurus* spp.) have similar dental morphology to *Otodus*, and *C. carcharias* has similar dentition to *Otodus* (*Carcharocles*) and *Otodus* (*Megaselachus*).” However, we feel that the typical tri-cusped morphology of teeth in *O. obliquus* is more akin to the teeth of the extant *Lamna nasus*. Future research that aims to expand this method to other taxa should take this into consideration when selecting their modern analogs.

Aside from heterodonty, ontogeny is often referenced as the greatest source of natural intraspecific variation in lamniform dentitions (Hubbell, 1996; Tomita et al., 2017). In *C. carcharias*, teeth tend to be narrower in juveniles and become progressively broader throughout ontogeny (Hubbell, 1996; Tomita et al., 2017). Interestingly, Tomita et al. (2017) noted that the intermediate tooth position is proportionately similar to adjacent teeth in early-

**FIGURE 8.** Correlation between summed crown width and total body length in 17 modern *Carcharodon carcharias* individuals, comparing the upper versus lower dentition. (A) linear function (upper dentition: $R^2 = 0.93$); (B) power function (upper dentition: $R^2 = 0.97$); (C) linear function (lower dentition: $R^2 = 0.90$); and (D) power function (lower dentition: $R^2 = 0.95$).
term embryos and that the lamnoid tooth pattern is achieved in mid- to full-term embryos. Consequently, the CH and CW proportions vary throughout ontogeny. This potential ontogenetic variation is difficult to account for in the extinct megatooth lineage, particularly with the small sample of associated dentitions available. Among the four associated dentitions of *Otodus megalodon*, they can be grouped into two size classes: juvenile (CH-31-46P and UF-VP46000) and adult (UF-VP-311000 and GHC 1). Consequently, correction factors for *O. megalodon* were separated into two categories (juvenile and adult) based on the most complete associated dentitions available.

While lamniform sharks generally conform to the lamnoid tooth pattern, there is undoubtedly intra- and inter-specific variation in the dental count and pattern. The general dental formula used in this study for *Carcharodon carcharias* had 48 teeth in its functional tooth series. However, the extant individuals measured in this study had a functional series ranging from 46 to 50 teeth (see Appendices 1 and 2). TL estimates for the largest associated dentition of *O. megalodon*, UF-VP-311000, using the 17 modern individuals of *C. carcharias* as analogs, ranged from 15.1 to 20.3 m, with a mean estimate of 17.3 m. TL estimates for the smallest associated dentition of *O. megalodon*, UF-VP-460000, using the same modern analogs, ranged from 9.4 to 12.7 m, with a mean estimate of 10.9 m. The range of estimates largely reflects the natural intraspecific variation exhibited in the modern *Carcharodon carcharias* used as analogs. Thus, there is an unavoidable range of error of approximately ± 1 to 3 m that is accounted for simply by taking the mean value. This range of error is positively correlated with TL, meaning that there will always be a greater range of estimates for larger individuals.

**Human observer error.** The natural variability in a species dental count is difficult to account for in disarticulated, associated dentitions of extinct taxa and is, consequently, a potential source of human error. The dental count for all *Otodus* dentitions was based on the most complete known associated dentition (CH-31-46P), under the assumption that the functional series in the left half of the dentition was completely present (Figure 1B). Thus, it was assumed that a complete functional series in *Otodus chubutensis* and *Otodus megalodon* consisted of 46 teeth (24 upper and 22 lower), which meant that every associated dentition of *Otodus* was missing at least one posterior tooth.

It is easy to imagine how posterior teeth could be missing from these fossil dentitions. As the smallest teeth in the dentition, they are the most susceptible to being transported away through taphonomic processes and would be the easiest to simply not see while collecting the other teeth. However, there is also the possibility that some of these individuals had a dental count slightly different from that of CH-31-46P, which would change the corrected SCW and resultant TL estimates. For example, in the largest associated dentition of *O. megalodon*, UF-VP-311000, the corrected SCW was calculated under the assumption that tooth position L9 was not found but would have been present (Figure 3A). If tooth position L9 was not missing and the teeth recovered represent the complete upper half of the dentition, the SCW would be 1055.5 mm and the TL estimate would be 16.7 m (0.6 m smaller than the current estimate). Likewise, we cannot be certain that this dentition did not have a dental count greater than that of CH-31-46P.

Every method that attempts to estimate body size from teeth requires one to identify tooth positions, whether working with isolated teeth or associated dentitions. When applying the Shimada (2002a) method to samples of isolated teeth, researchers often account for this by identifying teeth to a range of potential tooth positions and reporting the mean estimate. For example, Pimiento et al. (2010) identified an isolated tooth of *O. megalodon* (UF 246002) as belonging to tooth position L7-L9, with a CH = 24.5 mm. Applying the Shimada (2002a) method results in a range of TL estimates from 16.5 to 34.2 m, with an average estimate of 25.2 m. However, Pimiento et al. (2010) reported a TL of 11.5 m for this specimen. Under further inspection, Pimiento et al. (2010; figure S8) incorrectly reported that the TL equation for tooth position L9 from Shimada (2002a) as TL = -10.765 + 17.616(CH). This is actually the equation for tooth position i1 in Shimada (2002a; table 1). The correct equation for tooth position L9 in Shimada (2002a) was TL = -62.050 + 142.142(CH). This obviously has a drastic effect on the body length estimate and potentially major implications for the conclusion that the Gatun Formation represented a paleonursery for *O. megalodon*. Further, even with the correct equations from the Shimada (2002a) study, the range of error on the body length estimate is approximately ± 9 m for this specimen, which is much larger than any of the potential sources of error discussed so far.
To develop the SCW method, we had to infer which tooth positions were present in the disarticulated, associated dentitions. Generally speaking, it should be relatively easy to distinguish anterior vs lateral teeth and upper vs lower teeth, based on common trends observed in the typical lamnoid tooth pattern, but it is extremely difficult to identify a single tooth to an exact tooth position without context. However, when you have multiple teeth preserved together, there is context for making that determination. With a collection of associated teeth, one can assume that the taller, more symmetrical teeth will be more anteriorly positioned and the smaller, more asymmetrical teeth will be more posteriorly positioned. Even if multiple teeth were misplaced, as long as they are identified to the correct quadrant, their CW will still be included in the SCW. However, this could result in inaccurate correction factors used to estimate CW of missing tooth positions. At present, we consider these reconstructions as a hypothetical model and accept this as a source of error that cannot yet be fully accounted for.

Yet another potential source of human observer error is introduced while measuring specimens. Linear measurements (CH and CW) were recorded for teeth from 19 modern dentitions, 11 fossil associated dentitions, and one isolated fossil tooth. Despite these measurements being clearly defined, there will always be some degree of inconsistency between individuals. This source of error is often disregarded, yet it is present. For example, Shimada (2019) noted that Applegate and Espinosa-Arrubarrera (1996) incorrectly reported the TH for the largest known specimen of *O. megalodon* as 168 mm and stated that the correct TH is 162 mm. This changes the maximum TL estimate reported by Gottfried et al. (1996) from 15.9 to 15.3 m. Therefore, using the TH linear regression, measurement error may account for more than 0.5 m difference in TL.

To account for this potential source of error, authors VJP and RML independently measured CH and CW in nine of the 19 modern analogs. Body lengths were calculated using measurements from both subsamples separately. TL estimates using the SCW method varied by less than 0.2 m for all individuals. Admittedly, this measurement error likely also exists in the total body lengths reported from the extant comparative individuals; however, we cannot account for this variability, as the measurements were not taken by the authors of this study.

Mathematical equations. Methods for estimating body length based on TH, CH, and SCW all follow a similar logic that teeth grow proportionately with total body length. In contrast with the results of Reolid and Molina (2015), based on the sample of modern *C. carcharias* from this study, for the A2 tooth position, there is comparable correlation for CH and TL ($r=0.99$) versus CW and TL ($r=0.97$). Likewise, the correlation between upper SCW and TL ($r=0.97$) is comparable (Figure 8A).

Shimada (2019) determined that a power function has a greater coefficient of determination ($R^2$) than a linear function when analyzing the relationship between CH and TL. The same is true when analyzing the relationship between SCW and TL (Figure 8). This can likely be attributed to the greater variability in tooth proportions throughout ontogeny (Hubbell, 1996; Tomita et al., 2017). However, $R^2$ values may not be appropriate for determining which model is superior (Cornell and Berger, 1987). $R^2$ values are calculated differently for linear and nonlinear functions, so they are not necessarily directly comparable. Further, $R^2$ values are influenced by factors such as sample size and sample distribution. By segmenting the dentition into multiple regions to analyze variation within individuals, we effectively increased our sample size, which has an inverse effect on $R^2$ values. Thus, the higher $R^2$ produced by the Shimada (2019) power functions do not necessarily indicate that the model is better than SCW power function in this study.

To illustrate the variability associated with different mathematical assumptions, we estimated TL using the SCW direct proportions equation, SCW linear regression equation, and the SCW power regression equation (Table 4). As is to be expected, the SCW direct proportions and linear regression result in nearly identical TL estimates. Among the *Otodus* spp., in the upper dentition, TL estimates vary by at most 0.1 m; however, for the lower dentition, TL estimates vary by up to 0.8 m. This further supports that there is greater variance in the lower dentition. For the *Otodus* spp., the SCW power function always results in a larger TL estimate than the SCW direct proportions or linear regression methods. Again, as one would expect, the difference in TL estimates increases with larger individuals; however, all estimates still fall within the range of estimates produced by the SCW direct proportions method. Among the *Carcharodon* spp., there is less than 0.1 m difference in estimates derived from the SCW direct proportions, linear regression, and power regression methods for both
the upper and lower dentition. As such, we suggest that either the SCW direct proportions or SCW linear regression should be used to estimate TL of Otodus spp., whereas the SCW direct proportions, linear regression, or power function can be used to estimate TL of Carcharodon spp.

Paleoecological Implications

Body size estimates based on CH of isolated Otodus megalodon teeth have been utilized to identify paleo-nurseries (Pimiento et al., 2010; Herraiz et al., 2020) and as evidence for geographically discrete communities (Pimiento and Balk, 2015). These studies all employed the CH linear regressions developed by Shimada (2002a) to estimate body size, which result in widely variable estimates when applied to Otodus spp. (Figure 7 and Table 4). While applying the Shimada (2002a) linear equations to the associated dentitions in this study, it became apparent that the variation in TL estimates followed a predictable trend according to tooth position, in which body length estimates generally increased from anterior to posterior tooth positions (Figure 9A-B). Further, estimates from the upper and lower dentition differed the greatest in lateral tooth positions. This variability in estimates derived from different parts of the dentition has implications for the potential of sampling bias affecting interpretations of body size trends in O. megalodon.

In the Pimiento and Balk (2015) study, the total sample consisted of 544 O. megalodon teeth. Of which, 355 (65.2%) were identified as upper teeth and 189 (34.8%) were identified as lower teeth. Using a subsample of the 355 upper teeth, body lengths were re-estimated by extrapolating the SCW method. In general, the SCW method resulted in smaller TL estimates than the CH method. Negative values indicate that the SCW method results in smaller TL estimates than the CH method. Teeth are subdivided based on the tooth positions assigned by Pimiento and Balk (2015); however, there is some overlap between categories given that some teeth were identified as antero-lateral (e.g., A1-L1) or latero-posterior (e.g., L3-L7).
tooth positions and explains why the Shimada (2019) method always results in lower estimates than the SCW method.

As for the Pimiento et al. (2010) study, we pointed out that the TL equation for tooth position L9 was incorrectly transcribed from the Shimada (2002a) manuscript. Among the sample of 28 *O. megalodon* teeth from the Gatun Formation, 20 were assigned to upper tooth positions. Three of these teeth were identified as potentially belonging to tooth position L9. After re-calculating TL with the correct equation from Shimada (2002a), the estimates for these three specimens are 16.9, 17.1, and 25.2 m (Figure 10A). If we instead re-calculate TL using the SCW method, with a subsample of the 20 upper teeth, the range of estimates is much lower (Figure 10B). Using the SCW method, the same three specimens identified as possibly belonging to tooth position L9 result in TL estimates of 13.7, 13.8, and 11.5 m, respectively.

As such, the SCW method would actually provide better support for the hypothesis that the Gatun Formation represented a paleo-nursery for *O. megalodon* than the CH method. However, there are two main caveats to this conclusion. First, the life stages derived from Gottfried et al. (1996) should be re-assessed, as it was stated that “estimated lengths and mass for megatooth fetuses and neonates … are speculative.” Shimada (2021) reported a neonate TL of 2 m based on growth bands in a disarticulated, associated vertebral column of *Otodus megalodon* (IRSNB 3121); however, this single individual is not sufficient evidence for determining the size range of all *O. megalodon* neonates. Second, if one were to use the SCW method, the conclusion regarding the presence of a paleo-nursery would be drawn from a sample of only 20 teeth. That is a fairly small sample size to support an inference about the reproductive and distribution habits of an extinct species.

**Maximum Body Length**

As was described in the introduction, the maximum body size of *O. megalodon* has been revised on numerous occasions. Gottfried et al. (1996) reported a conservative maximum estimate of 15.9 m based on a maximum TH = 168 mm (FMNH PF 11306). However, Shimada (2019) noted that the TH for FMNH 11306 was incorrectly reported and is actually 162 mm, which would result in a TL of 15.3 m using the Gottfried et al. (1996) TH linear regression. Alternatively, Shimada (2019) reported a conservative maximum estimate of 14.2 m based on a maximum CH = 120 mm (NSM PV-19896).

Larger TL estimates have been reported by researchers that have attempted to employ the Shimada (2002a) CH linear regressions to isolated teeth (e.g., Pimiento et al., 2010: 16.8 m; Pimiento and Balk, 2015: 17.9 m; Reolid and Molina, 2015: 20.3 m). However, these estimates are all deemed unreliable based on errors in the data reported and the inconsistent results that are produced by the Shimada (2002a) method.

GHC 6 represents the largest tooth of *Otodus megalodon* known and physically accessible to the authors of this study (Figure 11A; TH=165 mm, CH=122 mm, and CW=133 mm). This specimen has a CH:CW of 0.917, which corresponds best with tooth positions L1-L2 based on the two largest associated dentitions of *O. megalodon* (UF-VP-311000 and GHC 1). In UF-VP-311000, tooth position L1 accounts for 10.5% of the SCW and tooth position L2 accounts for 10.2% of the SCW (see Appendix 5). The SCW direct proportions method was extrapolated using these proportions, resulting in a corrected SCW of 1266.7 to 1303.9 mm.
Based on a SCW of 1266.7 mm, the TL estimates ranged from 17.4 to 23.5 m, with a mean estimate of 20.0 m. Based on a SCW of 1303.9 mm, the TL estimates ranged from 18.0 to 24.2 m, with a mean estimate of 20.6 m.

One can quickly calculate the mean estimate by directly comparing the proportions in UF-VP-311000 and GHC 6 (Figure 11). This follows the same principle that was used to develop the SCW method. Since we are estimating body size based on the tooth with the widest known CW, it seems reasonable to assume that this tooth also represented the widest tooth in the dentition from which it originated. In which case, the proportions of tooth position L1 from UF-VP-311000 should result in the best estimate. For this tooth position, the SCW direct proportions method results in a TL of 20.0 m, the SCW linear regression results in a TL of 20.1 m, and the SCW power regression results in a TL of 21.6 m. We assume that the power regression overestimates the TL and that the SCW direct proportions and linear regression produce the more reliable estimate. As such, we predict a maximum body size for *O. megalodon* of approximately 20.0 m.

**CONCLUSION**

Through an evaluation of body length estimates using fossil associated dentitions, it is evident that estimates based on crown height of isolated teeth are highly variable when applied to different tooth positions. Further, this variability follows a predictable trend, in which anterior teeth result in smaller estimates than posterior teeth. This has clear implications for previous studies that
have employed the Shimada (2002a) CH method to analyze paleoecological and macroevolutionary trends based on body size of *Otodus megalodon*.

Herein, a novel method was developed based on summed crown width using fossil associated dentitions. By segmenting the associated dentitions into different regions, we were able to assess different sources of error that can affect body size estimation of fossil lamniform sharks. Specifically, estimates based on upper teeth are likely more accurate than estimates based on lower teeth. This is largely attributed to differences in the functional role of upper versus lower teeth. This method provides significantly greater constraint on the resultant body length estimate than the isolated CH method but requires assumptions about the dental pattern and count in the extinct megatooth lineage. These associated dentitions represent the most holistic direct evidence available to reconstruct the dental and cranial anatomy of the megatooth lineage but should still be treated as hypothetical models until an articulated dentition is recover.

Based on a sample of 17 modern *C. carcharias* dentitions, there was an approximately ± 1 to 3 m range of error in TL estimates for the eight associated dentitions of *Otodus* spp. that can be attributed to natural intraspecific variation. This range of error increases for larger individuals. While the sample size of fossil associated dentitions is not large enough to determine the size range of the taxa analyzed in this study, we did extrapolate the SCW method for the tooth of *O. megalodon* with widest crown width (GHC 6) to provide a maximum body size estimate. Assuming this tooth represents position L1, the SCW direct proportionality method results in a maximum body length estimate of 20.0 m for *O. megalodon*, with a range of error of approximately ± 3.5 m. Again, this range of error reflects the natural variation in the modern sample of *Carcharodon carcharias* used to calculate TL.

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APPENDICES

APPENDIX 1.

Crown width measurements in mm from modern *Carcharodon carcharias* (n=10) and *Isurus oxyrinchus* (n=1). Measurements are separated for the different regions preserved: upper left (UL), upper right (UR), lower left (LL), and lower right (LR). (All appendices are available for download in one zipped file at https://palaeo-electronica.org/content/2021/3284-estimating-lamniform-body-size.)

APPENDIX 2.

Crown height measurements in mm from modern *Carcharodon carcharias* (n=10) and *Isurus oxyrinchus* (n=1). Measurements are separated for the different regions preserved: upper left (UL), upper right (UR), lower left (LL), and lower right (LR). *pathologic or broken.* (All appendices are available for download in one zipped file at https://palaeo-electronica.org/content/2021/3284-estimating-lamniform-body-size.)

APPENDIX 3.

Crown width measurements in mm of 11 fossil dentitions of *Otodus* and *Carcharodon*. Measurements are separated for the different regions preserved: upper left (UL), upper right (UR), lower left (LL), and lower right (LR). (All appendices are available for download in one zipped file at https://palaeo-electronica.org/content/2021/3284-estimating-lamniform-body-size.)

APPENDIX 4.

Crown height measurements in mm of 11 fossil dentitions of *Otodus* and *Carcharodon*. Measurements are separated for the different regions preserved: upper left (UL), upper right (UR), lower left (LL), and lower right (LR). (All appendices are available for download in one zipped file at https://palaeo-electronica.org/content/2021/3284-estimating-lamniform-body-size.)

APPENDIX 5.

Correction factors applied for missing teeth in fossil *Carcharodon* and *Otodus* dentitions. Correction factors for *Carcharodon* spp. are based on the mean percentage of the total summed crown width from a sample of 10 modern *C. carcharias* dentitions (see Figure 6A-B). Correction factors for adult *Otodus megalodon* are based on crown width proportions in UF-VP-311000. Correction factors for juvenile *Otodus megalodon* are based on crown width proportions in CH-31-46P. Correction factors for *Otodus chubutensis* are from GHC 3. (All appendices are available for download in one zipped file at https://palaeo-electronica.org/content/2021/3284-estimating-lamniform-body-size.)