1 **Growth parameters of the clupeids *Limnothrissa miodon* and *Stolothrissa tanganicae* in**
2 **northern Lake Tanganyika (Bujumbura sub-basin)**

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SUMMARY

The analysis of growth parameters in fish stocks, such as the asymptotic length (L_∞) and the curvature parameter (K), is crucial for the estimation of production rates and total mortality rates in fisheries management. Here, we estimated the growth parameters of the clupeids *Limnothrissa miodon* and *Stolothrissa tanganicae* in the northern part of Lake Tanganyika (Bujumbura sub-basin). Both species are important targets of pelagic fisheries, but the available estimates are based on scarce size frequency data, and rarely on size as a function of age data. We provide new estimates of L_∞ and K based on size as a function of age data, with age being inferred from otolith weights. Furthermore, we reanalyze several existing length frequency datasets using advanced statistical procedures to test if growth parameters estimated by length frequency analysis are consistent with those obtained from size as a function of age. We found that length frequency analysis consistently underestimates L_∞ and overestimates K as compared to length-at-age data. We recommend that length-at-age data are collected at a much broader spatial and temporal scale than currently available to advance our understanding of the growth dynamics of these two sardine species. This will improve the assessment of stock productivity of Lake Tanganyika's pelagic fisheries and the conservation of these economically important fish species.

Key words: growth, size frequency analysis, daily increments of otoliths, *Stolothrissa tanganicae*, *Limnothrissa miodon*, Lake Tanganyika, Bujumbura sub – basin

INTRODUCTION

Lake Tanganyika is about 10 million years old (Cohen *et al.* 1993) and is the second deepest and the fifth largest freshwater body in the world (Langenberg, 2008). So far, 1200 animal and plant species have been identified in this lake, ranking it among the most diverse lake ecosystems in the world. Among the main groups in this lake, fishes show a high degree of biodiversity (Cohen *et al.* 1993; Van Steenberge *et al.* 2011). Lake Tanganyika's pelagic fisheries is a major source of food and economic activity of the human population living around the lake. These fish feed over 10 million people in the areas adjacent to Lake Tanganyika (Mölsä et al. 1999). Two small-sized clupeids, *Stolothrissa tanganicae* and *Limnothrissa miodon*, commonly known as "freshwater sardines", along with the latid perch *Lates stappersii*, account for 60 % to 90 % of the total commercial catch in the pelagic environment (Mölsä et *al.* 1999). In the mid-1990s, annual harvests from Lake Tanganyika were estimated to range between 165 000-200 000 tons, quantities that translate into annual revenues of tens of millions of US dollars (Mölsä et *al.* 1999). More recent estimates are missing, because data coverage is spotty and there are long time gaps between monitoring projects (Van der Knaap 2014; Plisnier et al. 2023).

As for other African Great Lakes, the lack of a long-term monitoring programme for Lake Tanganyika creates uncertainty about the status of the lake and its pelagic fish stocks (Plisnier et al. 2023). Fisheries statistics published from the North of Lake Tanganyika suggest that annual catch rates (measured as catch per unit effort; CPUE) have been declining since the early 1980s (Mulimbwa 2006; Mulimbwa 2020; LTA Secretariat, 2012; Mushagalusa et al. 2015; Okito et al. 2017; Van der Knaap et al. 2014). The declines have been attributed to fishing pressure (Mulimbwa 2006; Mulimbwa 2020; LTA Secretariat, 2012; Mushagalusa et al. 2015; Okito et al. 2017; Van der Knaap et al. 2014), as well as to global warming (Plisnier, 1997; O'Reilly et al. 2003; Verburg et al. 2003). However, overfishing can only be assessed based on declines in the total annual catch, for which no reliable data exist (Van der Knaap et al. 2014). Interestingly, while fisheries stakeholders in the North of the Lake do indicate a noticeable decrease in fish abundance and size, they do not attribute these fishery problems to overfishing (De Keyzer et al. 2020). Yet another recent study found that CPUE has been stable in the North of Lake Tanganyika since 1993 (Sarvala & Mulimbwa 2023), indicating that not all fisheries statistics show the same trend. The impact of the effects of global warming on the pelagic fisheries has been questioned as well (Sarvala et al. 2006a, Sarvala et al. 2006b; Sarvala & Mulimbwa 2023). These uncertainties confirm that better fisheries monitoring is required to more accurately forecast fishing yields, assess fisheries potential, and understand the impact of fishing activities (Kolding et al. 2019).

One cornerstone of adequate fisheries management concerns the analysis of growth. Growth parameters such as the asymptotic length (L_∞) and the curvature parameter (K) form the basis to estimate production rates and total mortality rates, which enable the prediction of fishing yields (Beverton & Holt 1957). Growth curves are used where age data are not available, as in most tropical fish stock assessments (Sparre & Venema 1989). L_∞ and K can either be determined based on changes in length frequency distributions over time (length-frequency

analysis; LFA) or based on length as a function of age data (length-at-age analysis; LAA).
LFA has the advantage that size data are easy to obtain. However, the method requires young
fish and frequent catch data to identify subsequent cohorts and track their modal growth
progression. In environments and species with multiple periods of reproduction throughout
the year, multiple overlapping cohorts and multimodal size frequency distributions may
complicate cohort identification. Here, LAA might be more powerful. A commonly used
technique for age determination in fish involves counting natural growth rings on the otoliths
(Beamish & McFarlane 1983). In several tropical fishes, the diurnal transition from light to
dark leaves a mark on the otoliths, and these daily increments can be used to estimate fish age
in days (Kaningini, 1995; Kumura, 1995; Meisfjord et al. 2006; Chifamba 2019). This
method is suitable for tropical environments where the temperature is almost constant
throughout the year. While counting daily otolith rings can be time consuming, otolith radius
and otolith weight tend to correlate well with the number of daily increments and are
therefore also suitable to establish length-age relationships (Lou et al. 2005).
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In this study, we estimated growth parameters for *S. tanganicae* and *L. miodon*. Various
studies have gained useful insights in the biology of both clupeids, including their
evolutionary origin (Milec et al. 2022), population genetic structure (De Keyzer et al. 2019;
Junker et al. 2020), position in the food web (Sarvala et al. 1999; Ehrenfels et al. 2023),
parasites (Kmentová et al. 2020), reproductive biology (Mulimbwa et al. 2014a; Mulimbwa
et al. 2022), and responses to limnological changes (Kimirei & Mgaya, 2007; Plisnier et al.
2009; Mulimbwa et al. 2014b). While *S. tanganicae* is fully pelagic, *L. miodon* makes more
use of the littoral areas, also for reproduction and as nursery grounds (Mulimbwa et al. 2022).
Both fish species are the target of artisanal fisheries in the pelagic, but in shallow littoral
waters, juveniles of *L. miodon* are also subjected to the relatively low selectivity of beach
seines (Mölsä *et al.* 2002). Basic information for fisheries management, such as age at
maturity and size at maturity are available for both species, although there are no in-depth
studies assessing the variability in those parameters. In addition, the data are patchy in both
time and space. Age at maturity in Lake Tanganyika is usually estimated at 8.5 months
corresponding to a total length of 78 mm for females and 77 mm for males in *S. tanganicae*,
and at 8.6 months corresponding to a total length of 101 mm for females and 95 mm for
males in *L. miodon* (Mannini 1998). Growth parameters have been obtained for both species
as well (Chapman & van Well, 1978; Roest, 1978; Coulter, 1991; Moreau et al. 1991;
Mulimbwa & Shirakihara, 1994; Kimura 1995; Ahonen 2001; Mulimbwa 2006). Most
studies have inferred growth parameters using LFA on length–frequency distributions from
catch data. The few exceptions using LAA include Kimura (1991b, c), Kimura (1995) and
Ahonen (2001). Especially in *S. tanganicae*, the most abundant cohort is often the one
recruited during the dry season (Mulimbwa et al. 2014b). Yet, multiple partially overlapping
cohorts do occur in both species, and pose challenges of cohort delineation for the assessment
of growth.
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To address the need for more accurate growth data and validate existing growth data for the
two clupeid fish species, we provide new estimates of L_∞ and K based on length-at-age data,
with age inferred from otolith weight. Furthermore, we reanalyze three published datasets

with length frequency data using more recent and advanced statistical procedures, and test if
growth parameters estimated by size frequency analysis are consistent with those obtained
from length-at-age data.

MATERIAL AND METHODS

Study design

This study combines datasets from different surveys on the two clupeids that were all
conducted in the northern end of Lake Tanganyika (03° 28' S, 29° 17' E; Fig. 1) in the
Bujumbura sub-basin. We included (1) length frequency data for the period 1987-1989 from
Mulimbwa and Shirakihara (1994), (2) length frequency data for the period 1999-2001 from
Mulimbwa (2006), (3) length as a function of age data for the period 2004-2005 from
Mulimbwa et al. (2014b), and length frequency data for the period 2007-2008 from
Mulimbwa et al. (2014b) (Table 1). Sampling and data collection procedures for each data
type are described in more detail below.

As *S. tanganicae* is entirely pelagic, all data for this species originate from samples collected
in the pelagic environment. In contrast, *L. miodon* occurs in the pelagic environment as well
as in littoral areas of the lake. The data for *L. miodon* were obtained from both the littoral and
the pelagic in 1987 – 1988 and in 1999 – 2001, while in 2004 – 2005 and in 2007-2008, the
species was only sampled in the pelagic.

Because the available length frequency data (1987-1989, 1999-2001, 2007-2008) and length
at age data (2004-2005) originated from different survey periods, our study design is not
optimal to compare growth parameters based on the two data types. Nevertheless, since the
length-at-age survey period is embedded within the two decades spanning the three length
frequency surveys, it should still be possible to distinguish effects of data type from temporal
patterns, especially when temporal variation is minor.

Length-at-age data

Briefly, 260 specimens of *S. tanganicae* and 244 specimens of *L. miodon* were collected in
the pelagic from the study area in 2004-2005 (Table 1). First, the total length (TL) of each
specimen was measured to the nearest mm, and otoliths were extracted and dried at room
temperature at the CRH laboratory in Uvira (DR Congo). Subsequently, the otoliths were
weighed with a microbalance (µg; dry weight at 20 °C) at the University of Turku (Finland).
Finally, the age of these fish was estimated using the relationship between otolith dry weight
(ODW) in µg and age in days based on counts of daily rings following the formula age =
(ODW + a)/b, where a = 99.4, b = 2.42 for *S. tanganicae* and a = 120.8, b = 2.53 for *L.*
miodon (Ahonen 2001). Ahonen (2001) indicated that the relationship between ODW and age
was linear or near-linear, with a high Pearson correlation coefficient in both species (*S.*
tanganicae: $R^2 = 0.88$; *L. miodon*: $R^2 = 0.96$). Simple linear regressions of ODW versus age

for the two species, as originally established by Ahonen (2001), are visualised in
Supplementary Figure 1.
186
187 *Length frequency data*
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For all three studies, samples were taken from artisanal fishing units using lift nets with a
mesh size of 4.4 mm (knot to knot) in the upper four-fifths of the net and 3.3 mm (knot to
knot) for the rest of the net. The samples were collected twice a week during the investigation
period. After collection, samples were sorted by species at the CRH laboratory in Uvira (DR
Congo). Their total lengths (TL) were determined to the nearest mm, and the specimens were
grouped into size classes of 2 mm for *S. tanganicae*, and 4 mm for *L. miodon*, in 1999-2001
and 2007-2008. The data from 1987-1989 were collected in size classes of 5 mm for both
species. Total sample sizes for each period are shown in Table 1.
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For 1987 – 1988, the original data from Mulimbwa and Shirakihara (1994) could not be
retrieved from the authors. Instead, we digitized the length frequency plots in Figure 2 (*S.*
tanganicae) and Figure 4 (*L. miodon*) in Mulimbwa and Shirakihara (1994) using
WebPlotDigitizer version 4.6 (Rohatgi 2022). Since length in the 1987-1988 study was
measured in mm as standard length (SL), size classes were converted to total length (TL) in
mm following the formula TL = a + (b * SL), where a = 0.8729, b = 1.1562 for *S. tanganicae*
and a = 1.6658, b = 1.1873 for *L. miodon* (Mannini *et al.* 1996).
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206 *Data analysis*
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208 1) Length-at-age analysis
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210 A Von Bertalanffy (1938) growth function (VBGF) was fitted to the otolith derived length-
211 age data, and its parameters L_∞ (cm) and K (year^{-1}) were determined using the non-linear
212 least squares method, available within the function "growth_length_age()" in the TropFishR
213 package (Mildenberg et al. 2017).
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215 2) Length frequency analysis
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The growth parameters to fit a VBGF to length frequency data have historically been
estimated using several methods (e.g. the Power-Wetherall or Ford-Walford method), most of
which have been questioned for containing intrinsic biases from different sources
(Schwamborn 2018). However, recent improvements to ELEFAN (electronic length
frequency analysis, Pauly & David 1981), the most widely used method for fitting a growth
curve to length frequency data, provide a promising new path for length-based studies of
aquatic populations. These advancements include the estimation of both L_∞ and K with fewer
assumptions than previous models (Taylor & Mildenberg 2017), as well as the calculation of
their associated uncertainties using bootstrapping (Schwamborn et al. 2019). We applied
these new methods using the TropFishR and fishboot packages in R (Mildenberg et al. 2017;
Taylor & Mildenberg 2017; Schwamborn et al. 2019) with the parameters shown in Table 2.

ELEFAN relies on attributing different scores to restructured different length bins, depending on how much each bin deviates from a moving average across a user-defined number of length bins. This was done by grouping the size classes into bins of 0.8 cm each, and a moving average of 5-7 size classes (Table 2), depending on the size of the smallest cohort, following the general rule of thumb (Taylor & Mildenberg 2017). Afterwards, a score is given to different growth curves, depending on whether they intersect with positively or negatively scored length bins (Gayanilo & Pauly 1997). The curve with the highest score is considered the best fitting curve to the data. Finding the best fitting curve can be computationally costly. Two optimization approaches ("simulated annealing" and "genetic algorithm") have been implemented for growth function fitting in R (Taylor & Mildenberg 2017; Mildenberg et al. 2017). Both optimization approaches can find high-scoring solutions, but because several models often score high, different solutions can be reached depending on the initial settings, which can create an intrinsic source of bias. To overcome this limitation, bootstrapping methodology can be applied, fitting a large number of growth curves with similar scoring solutions, and finally calculating the uncertainty of the mean value of the estimations (Schwamborn et al. 2019). We estimated L_∞ and K of both species from length frequency data by bootstrapping both algorithms, using the package "fishboot" in R (Schwamborn et al. 2019), and we report the probability distribution of the estimations. The total parameter space considered ranged from 1 to 10 for K for both species (Table 2). For L_∞, we considered values from 6 cm to 12 cm for *S. tanganicae*, and from 8 to 22 cm for *L. miodon* (Table 2). Here, the upper values are above the maximum total length reported for each species in Fishbase (*S. tanganicae*: 10 cm SL ~ 11.65 cm TL; *L. miodon*: 17 cm SL ~ 20.35 cm TL; Froese & Pauly 2017). We manually set a maximum age to each species (Table 2; 18 months for *S. tanganicae* and 30 months for *L. miodon*; Mannini et al. 1996), which is a key step for the calculation of the growth parameters in this method.

3) Consistency between methods

To assess the consistency between length-at-age-based and size frequency-based methods, the agreement of growth parameters (L_∞ and K) between different sampling periods was tested by comparing the 95 % confidence intervals of the estimates based on LAA with the 95 % credibility intervals of the estimates based on LFA.

RESULTS

S. tanganicae

Length frequency data across time periods ranged between 2.1 and 11.1 cm TL. Recruitment was continuous throughout the year, as indicated by small individuals (2.1 - 4.0 cm) appearing all year round in the length frequency data of the three survey periods. Length-at-age data comprised individuals ranging between 3.3 cm and 10.6 cm in TL, and between 56

and 372 days in age. The length distribution of the length frequency data and the length-at-
age data was comparable (Fig. 2).

Based on length-frequency data analysis (LFA), estimates of L_∞ (range: 8.38 - 9.56 cm) were
consistent between estimation methods with narrow confidence intervals, and showed
substantially lower values in 1987-1989 and 1999-2001 than in 2007-2008 (Table 3;
Supplementary Figure 2). In contrast, the annual K (range: 5.48 – 8.46) showed high
uncertainty which led to an overlap in confidence intervals between most study periods and
estimation methods (Table 3). Based on the length-at-age data (LAA) from 2004-2005, we
obtained an estimate of 10.64 cm for L_∞ and 3.18 for K (Table 3). Non-overlapping
credibility and confidence intervals indicated that all estimates of L_∞ and most estimates of K
were respectively significantly lower and higher for LFA than for LAA. According to the
LAA curve, *S. tanganicae* grows rapidly during the first five months of its life and growth
becomes almost constant after ten months (Fig. 3; Fig. 4; Supplementary Figure 3).

L. miodon

Length categories of the length-frequency data across time periods ranged between 2.4 and
15.2 cm TL. Recruitment occurred less frequently throughout the year than in *S. tanganicae*,
as small cohorts (2.4 - 6.0 cm) were less frequent across the three survey periods. Length-at-
age data comprised individuals ranging between 2.5 cm and 13.6 cm in TL, and between 62
and 571 days in age. The length distribution of the length frequency data and the length-at-
age data was comparable (Fig. 2).

Based on length-frequency data, estimates of L_∞ (range: 9.52 - 10.97 cm) were consistent
across estimation methods, but were lower in 2007-2008 than in 1987-1989 and 1999-2001
(Table 3; Supplementary Figure 2). Ranging between 2.16 and 5.07, the annual K was more
consistent across the three LFA survey periods and estimation methods than estimations for
S. tanganicae, although the uncertainty was generally also high (Table 3). Based on the
length-at-age data from 2004-2005, we obtained an estimate of 12.82 cm for L_∞ and 1.73 for
K (Table 3). Similar to *S. tanganicae*, non-overlapping credibility and confidence intervals
intervals indicated that all estimates of L_∞ and half of the estimates of K were respectively
significantly lower and higher for LFA than for LAA. According to the LAA curve, *L.*
miodon grows rapidly during the first ten months of its life and growth becomes almost
constant after 18 months (Fig. 3; Fig. 4; Supplementary Figure 3).

DISCUSSION

The objective of this study was to examine the growth patterns of *S. tanganicae* and *L.*
miodon in the North part of Lake Tanganyika. We compared growth estimates derived from
observed length frequency data and indirectly estimated length-at-age data, collected over
different survey periods. This comparison informs us about the accuracy of size frequency-
based estimation methods for growth parameters, and the necessity of complementing

fisheries management with more complete life history information using age-based data. For
both species, we consistently found lower estimates of L_∞ and higher estimates of K for
length frequency data than for length at age data. Below we discuss the reliability of this
result, and its implications for the management of the artisanal fisheries.

Because length frequency and length at age data were collected in different survey periods,
estimation procedures and survey periods are confounded in our study, and therefore do not
allow us to directly compare the results. Nevertheless, because the differences in estimates
between the two procedures largely exceed the differences between survey periods within
LFA, it is unlikely that temporal variation alone can account for the observed differences
between the two methods. It is indeed more likely that the disparities between data types and
estimation methods stem from methodological nuances that influence the estimation of
growth parameters. Otolith-based methods are believed to provide accurate estimations of
age, which together with direct measurements of length, should provide accurate growth
curves (Schwamborn et al. 2023). Ahonen (2001) demonstrated that the age of the two
sardine species can be reliably estimated from otolith dry weight (Pearson correlation
between otolith dry weight and age based on counts of daily otolith rings: $r^2 = 0.88$ for *S.*
tanganicae; $r^2 = 0.96$ for *L. miodon*; Supplementary Figure 1). While this relationship was
established using individuals from the central part of Lake Tanganyika (Kigoma, Malagarasi
and Kipili; Ahonen 2001), we applied it here to individuals from the North of Lake
Tanganyika. Nevertheless, this should yield acceptable results, and overall, the accuracy of
our LAA-based estimates most likely exceeds the accuracy of our LFA-based estimates.

In general, LFA is highly sensitive to sampling bias, including the lack of large, old
individuals (Schwamborn 2018; Schwamborn et al. 2019; Wang et al. 2021), as well as the
absence of young cohorts. Other potential sources of sampling bias include the concentration
of fishing efforts on commercially valuable areas, and differences in fishing gears and
methods. These potential issues with LFA likely apply to the two Lake Tanganyika sardine
species, as notified previously by Mulimbwa & Shirakihara (1994). In the present study, the
discrepancy between LFA-based and LAA-based estimates may be due to an
underrepresentation of the largest individuals in the length-frequency data of both species.
The relatively low number of young cohorts in *L. miodon* may also have lowered the
precision of LFA in this species. Avoiding sampling bias for size is particularly challenging
in *L. miodon* because large, mature individuals are more commonly observed in littoral areas
than in the pelagic zone, so the sampling effort in each area can directly affect the results
(Mulimbwa & Shirakihara 1994; Mulimbwa et al. 2022). Since no *L. miodon* were collected
in the littoral areas during the most recent survey period (2007-2008), part of the variation in
LFA-based estimates of L_∞ and K between years might be due to sampling bias. However,
similar fluctuations in LFA-based estimates of L_∞ and K were observed in *S. tanganicae*,
which was always sampled in the pelagic. Considerable variation in growth curves based on
LFA is thus expected. Currently, it remains unclear if LAA data generate more consistent
results than LFA. Our LAA-based estimates of L_∞ for both *S. tanganicae* and *L. miodon* from
catches in 2004-2005 were substantially lower than the LAA-based estimate for both species
from catches in 1990 by Kimura (1995) (Figure 5). However, both studies were conducted at

opposite ends of the lake, and spatial variation in environmental conditions may also affect
growth trajectories. Overall, better data are required to assess the robustness of growth
parameters.

Another challenge of LFA is the aggregation of individuals into length bins, which has the
potential to obscure the age structure within each bin (Taylor & Mildenberger 2017;
Schwamborn, 2018). This can manifest as well in lower L_∞ values and higher K values,
particularly with bigger bin sizes (Wang et al. 2020). However, a recent study directly
comparing LFA and LAA in lane snapper (*Lutjanus synagris*) found no difference in L_∞ and
K between the two methods (Schwamborn et al. 2023). Interestingly, Kimura (1995)
concluded that the growth parameters obtained from length-at-age data (based on daily counts
of otolith rings) generally agreed with those previously estimated from analyses of length–
frequency distributions. However, close inspection of Kimura's (1995) results reveals that the
author's otolith-based estimates for L_∞ almost always exceeded the estimates for L_∞ from
length-frequency studies, and *vice versa* for K, and thus seemed to be subjected to the same
bias we report here.

Finally, both our LAA-based and LFA-based estimates for the growth trajectory of the two
sardine species from the North of Lake Tanganyika deviated substantially from estimates of
previous studies (Table 1; Figure 5). This includes LAA-based and LFA-based studies from
various regions of Lake Tanganyika (Figure 5), but also the length frequency datasets from
the North of Lake Tanganyika for which we re-estimated the growth parameters (Figure 5;
Table 1; Mulimbwa & Shirakihara 1994; Mulimbwa 2006). Almost all studies reported
significantly larger values of L_∞, and therefore lower values of K than the present study
(Figure 5). This was true for both species, although the discrepancy for K between previous
studies and the present study was more pronounced in *S. tanganicae*, while the discrepancy
for L_∞ was more pronounced in *L. miodon* (Figure 5). For the LFA-based estimates, this
discrepancy could be explained by the fact that previous studies are largely based on outdated
methods such as the Ford-Walford plot, which are known to generate inaccurate results
(Schwamborn 2018). Indeed, the improved ELEFAN methods we used here are based on
fewer assumptions and allow for screening a large range of parameter values, which should
lead to more precise and easily reproducible results (Mildenberger et al. 2017; Schwamborn
et al. 2019). Still, the subjective choice of suitable parameter space for L_∞ can often result in
biased results with narrow confidence intervals (Schwamborn et al. 2023). For this reason, we
opted for upper parameter limits above the maximum length reported in Fishbase for each
species, which should result in unbiased estimations with realistic uncertainty estimates. As a
result, some of our LFA-based estimates, especially for K, come with very high uncertainties,
which may indicate that the quality of the available length frequency data is insufficient to
achieve robust conclusions. This leaves little room for scientific advice beyond improving
monitoring efforts to obtain better and more accurate growth parameters based on LFA.

Conclusions and implications for fisheries management

Growth parameters link age and size and are thus key to all length-based stock assessment
models where data on catches by age are not available. The lower L_∞ and higher K values

obtained with length frequency data relative to length-at-age data for the two sardine species emphasizes the need for future research aimed at advancing our understanding of growth dynamics and their implications for the pelagic fisheries of Lake Tanganyika. We recommend that length-at-age data are obtained at a much broader spatial and temporal scale than is currently available. However, as length frequency data can be more easily collected, such data will remain our primary source of information and should continue to be obtained. Furthermore, both length frequency data and length-at-age data should be collected for the same study periods to test for a systematic bias between the two methods. In addition, it is important that all samples in a time series come from the same landing site and fishing area to rule out that changes in length frequencies over time are due to sampling bias. As we have shown, it is straightforward to reanalyze existing length frequency data whenever the original data can be retrieved, but it is also possible to regenerate the data after digitizing published length frequency plots.

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DATA AVAILABILITY

Datasets including the two files with length-at-age data and the six files with length frequency data, as well as the script to run the analysis with the TropFishR package in R are available on Zenodo (https://doi.org/10.5281/zenodo.12812204) and Github (https://github.com/CescGV/Parameters_Limno_Stolo_Tanganyika-).

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  TABLES

  Table 1. Overview of datasets used in this study for the analysis of growth parameters
  including survey period, total sample size (N), data type (LF: length frequency data; LA:
  length-at-age data), published estimates of L_∞ (TL in cm) and K, and reference. NA indicates
  that no information is available from the original publications. Estimates of L_∞ marked with
  an asterisk are converted from SL to TL following Mannini *et al.* (1996).

Survey period (months)	N	Data type	L_∞	K	Reference
S. tanganicae					
Nov 1987 - Apr 1989 (18)	24,167	LF	12.34*	2.14	Mulimbwa & Shirakihara 1994
May 1999 - Nov 2001 (30)	84,503	LF	11.70	2.6	Mulimbwa 2006
2004 - 2005	260	LA	NA	NA	Mulimbwa et al. 2014b
Feb 2007 - May 2008 (16)	NA	LF	NA	NA	Mulimbwa et al. 2014b
L. miodon					
Nov 1987 - Apr 1989 (18)	3,834	LF	16.12*	1.15	Mulimbwa & Shirakihara 1994
May 1999 - Nov 2001 (27)	12,272	LF	17.90	0.74	Mulimbwa 2006
2004 - 2005	244	LA	NA	NA	Mulimbwa et al. 2014b
Feb 2007 - May 2008 (16)	NA	LF	NA	NA	Mulimbwa et al. 2014b

649 **Table 2.** Parameters used in the VBGF fitting using ELEFAN bootstrapping in each of the
650 three surveys based on length frequency data. GA: Genetic algorithm; SA: Simulated
651 annealing; MA: moving average; L_∞: asymptotic total length; K: annual growth coefficient;
652 t_anchor: peak spawning month; nresamp: number of permutations.

Species	Algorithm	Parameter (unit)	1987-1988	1999-2001	2007-2008
S. tanganicae	GA, SA	MA (bins)	5	7	5
		L_∞ (cm)	6-12	6-12	6-12
		K (year^{-1})	1-10	1-10	1-10
		t_anchor (year)	0-1	0-1	0-1
		nresamp	400	400	400
		Agemax (year)	18/12	18/12	18/12
		bin_size (cm)	0.8	0. 8	0. 8
L. miodon	GA, SA	MA (bins)	5	7	7
		L_∞ (cm)	8-22	8-22	8-22
		K (year^{-1})	1-10	1-10	1-10
		t_anchor (year)	0-1	0-1	0-1
		nresamp	400	400	400
		Agemax (year)	30/12	30/12	30/12
		bin_size (cm)	0. 8	0. 8	0. 8

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Table 3. Growth parameters of clupeids (*S. tanganicae*, *L. miodon*) estimated for different
time periods and using different estimation procedures for length frequency analysis (LFA)
and length-at-age analysis (LAA). Estimation methods include two optimization approaches
in the case of LFA (SA: ELEFAN with simulated annealing; GA: ELEFAN using genetic
algorithm), and the non-linear least-squares method (LSM) in the case of LAA. L_∞:
asymptotic total length (cm), K: annual growth coefficient. CI: Bayesian credibility intervals
(in the case of LFA) or confidence intervals (in the case of LAA).
663

Survey period	Analysis	Estimation	L_∞ [95 % CI]	K [95 % CI]
S. tanganicae				
1987-1989	LFA	GA	8.82 [7.97-8.90]	5.48 [4.49-7.59]
		SA	8.38 [7.65-8.99]	8.27 [4.66-9.91]
1999-2001	LFA	GA	8.67 [8.47-8.80]	7.86 [6.81-8.79]
		SA	8.71 [8.06-8.84]	8.46 [5.01-10.42]
2004-2005	LAA	LSM	10.64 [10.32-11.05]	3.18 [2.76-3.63]
2007-2008	LFA	GA	9.41 [9.20-9.60]	6.64 [3.72-9.35]
		SA	9.56 [9.01-9.64]	5.89 [3.57-9.99]
L. miodon				
1987-1989	LFA	GA	10.93 [10.72-11.18]	2.55 [2.12-3.63]
		SA	10.83 [10.46-11.31]	2.49 [1.30-9.42]
1999-2001	LFA	GA	10.96 [10.69-11.17]	2.36 [1.69-6.34]
		SA	10.97 [10.65-11.30]	2.16 [1.51-6.13]
2004-2005	LAA	LSM	12.82 [12.18-13.64]	1.73 [1.47-2.01]
2007-2008	LFA	GA	9.61 [9.26-9.70]	3.30 [2.44-9.84]
		SA	9.52 [8.87-10.07]	5.07 [2.10-10.09]

665 **FIGURE LEGENDS**
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667 **Figure 1.** Geographic position of Lake Tanganyika bordered by DR Congo, Burundi,
668 Tanzania and Zambia. The shaded area marks the Bujumbura sub-basin.
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670 **Figure 2.** Total length distribution (cm) for both species and the four survey periods.
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Figure 3. VBGF estimated from total length frequency data of 1987-1988, 1999-2001 and
2007-2008 for (A) *Stolothrissa tanganicae* with Genetic Algorithm (GA); (B) *Stolothrissa*
tanganicae with Simulated Annealing (SA); (C) *Limnothrissa miodon* with GA; and (D)
Limnothrissa miodon with SA. Each plot is superimposed with VBGF estimates from otolith-
derived length at age data of 2004-2005, along with its 95 % confidence interval. Bayesian
credibility intervals (in the case of LFA) or confidence intervals (in the case of LAA) were
obtained from growth curve estimations for the upper and lower limits of the 95 % credibility
or confidence intervals of parameters L_∞ and K.
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Figure 4. Total length frequency data and corresponding Von Bertalanffy growth curves for
the three survey periods for *Stolothrissa tanganicae* (left panels) and *Limnothrissa miodon*
(right panels). The red and blue dashed lines mark growth curves estimated with the genetic
algorithm and the simulated annealing algorithm, respectively. For comparison, the growth
curves estimated from length-at-age data are superimposed (black dashed lines with t_0 set to
1st of January). The length frequency data are plotted in dark blue, and blue shades reflect the
frequency of each length class.
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Figure 5. Top: annual growth coefficient (K) against asymptotic total length (L_∞ in cm) for
S. tanganicae (red) and *L. miodon* (blue) based on length-frequency analysis (LFA; circles)
or length-at-age analysis (LAA; triangles). Data points with black edges and error bars (95 %
confidence intervals) are generated by this study and are listed in Table 3. Data points
without error bars are derived from the overview provided in Table 9 and Figure 47 in
Mannini et al. (1996), in addition to estimates by Ahonen (2001) and Mulimbwa (2006).
Labeled data points are based on LFA data by Mulimbwa & Shirakihara (1994) (As: *S.*
tanganicae; Al: *L. miodon*), LFA data by Mulimbwa (2006) (Bs: *S. tanganicae*; Bl: *L.*
miodon), LAA data by Mulimbwa et al. (2014b) (Cs: *S. tanganicae*; Cl: *L. miodon*), LAA
data by Kimura (1995) (Ds: *S. tanganicae*; Dl: *L. miodon*), and LAA data by Ahonen (2001)
(Es: *S. tanganicae*; El: *L. miodon*). **Bottom**: Von Bertalanffy growth curves for the present
study and previous studies by species (*Stolothrissa tanganicae* vs. *Limnothrissa miodon*) and
method (LAA vs. LFA). Each curve corresponds to a data point of the top panel.
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703 **FIGURES**
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706 **Figure 1.**
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709 **Figure 2.**

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711   **Figure 3.**

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713 **Figure 4.**

716   **Figure 5.**

 SUPPLEMENTARY FIGURES

Figure 15. Relationship between otolith weight and age of *L. miodon* (top) and *S. tanganicae* (bottom). Markers represent areas (O= Kigoma,◻=Malagarasi, Δ=Kipili) and sexes (+ =males, •= females). Solid lines represent predicted otolith weights and dotted lines 95 % confidence intervals.

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720 **Supplementary Figure 1.** Figure 15 from Ahonen (2001).

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Supplementary Figure 2. Probability distribution of estimated parameters using genetic and
simulated annealing algorithms for *Stolothrissa tanganicae* and *Limnothrissa miodon* in each
study period.

Supplementary Figure 3. Von Bertalanffy growth functions (VBGF) for *Stolothrissa tanganicae* (in red) and *Limnothrissa miodon* (in blue), estimated from otolith-derived length-age relationships. Confidence intervals were obtained from growth curve estimations for the upper and lower limits of the 95 % confidence intervals of parameters L_∞ and K.