

Short note

Age structure of *Chionodraco hamatus* (Teleostei, Channichthyidae) samples caught in Terra Nova Bay, East Antarctica

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Received 7 August 1991; accepted 9 June 1992

Summary. The age composition of *Chionodraco hamatus* samples caught by fixed nets in Terra Nova Bay during the Austral summer 1987/88 was determined by examination of thin otolith sections. Individual ages were estimated by counting the number of annuli seen in the sections, since we postulated that annuli were laid down yearly. Fish were estimated to be 5–10 years old. Although our “apparent ages” were not confirmed by other independent estimates, our data seem to compare well with the values reported for other Antarctic fishes. Females in our samples were on average larger than males of the same age and grow somewhat faster, at least over much of the sampled size range.

Introduction

As in temperate species, many Antarctic fishes show year-rings (annuli) in their otoliths, scales and bones. This has been confirmed by length-frequency analysis (e.g. Kock, 1980, 1981; Pankhurst 1990), counts of sclerites related to total lengths (TLs; Freytag 1981) and from frequent sampling at sea to monitor the formation of the opaque/hyaline zones in otoliths (North 1988) or fast/slow growth areas in scales (Scherbich 1975).

These studies suggest that Antarctic fish may be aged by careful examination of suitable hard structures. Otolith and scale readings have been extensively used to define both the growth rates and length/age keys of many fish species of commercial interest (North and White 1987).

Little data are available on other fishes which could play an important role in the food web of some Antarctic subareas. In this paper, we report on the age composition (defined as the number of annuli counted in otolith sections) of the unexploited Channichthiid *Chionodraco hamatus* for which no similar data have been previously reported. Although our data remain unconfirmed (i.e. the concordance of annuli with year classes has not yet been

separately checked), they presumably describe the true age structure of the virgin stock from which the fishes were drawn.

Thus, our data may be regarded as a baseline to monitor future changes in the population under examination and can be compared with the age composition of other Channichthiid stocks which are extensively fished.

Material and methods

The area sampled was in Terra Nova Bay stretching between 74°41'S/164°49'E and 74°49'S/164°08'E. The specimens were obtained both by a trammel-net (length = 108 m, height = 1.80 m, mesh opening = 64 mm) and a gill-net (length = 123 m, height = 4.60 m, mesh opening = 64 mm) set in the 77–175 m depth range. A total of 23 fishing operations were carried out from January 13th through February 13th 1988, of which 19 (11 for trammel-net and 8 for the gill-net) caught *C. hamatus* specimens.

The fishes were measured, the total and eviscerated weights recorded (TL and weights being approximated, respectively, to the lower millimetre and gram), the gonads staged (Everson 1977) and the Gonado Somatic Index (GSI) values calculated. In most fishes (118 out of 180), sagittae were taken and stored dry for further study. However, 14 specimens were not “worked” at all.

In the laboratory, the resulting sagittae were brittle and almost completely opaque if observed as a whole, so we adopted the sectioning technique developed by Bedford (1983) on gadoid otoliths. In short, the sagittae were embedded in a polyester casting resin, cut in 0.5 mm thick slices, and then mounted on slides (see Bedford, loc. cit., for further details).

The otolith sections (one for each specimen aged) were then observed at low magnification (up to 50x) both in reflected and transmitted light. The slides were read separately on two occasions by two different readers.

In some cases, numbers differed by one (rarely more) unit(s), because the very central part of the sections appeared opaque so that one or more annulus/i could hardly be seen (Fig. 1). Problems in numbering the annuli of the inner part of the otoliths are apparently rather common when dealing with Antarctic fishes (e.g. Tomo and Barrera Oro, 1986). These “difficult” sections were jointly re-examined. Afterwards, if readings still differed by more than two units, these were discarded.

Since no systematic differences were observed between readers, we assumed that both the higher/lower value of each pair had about

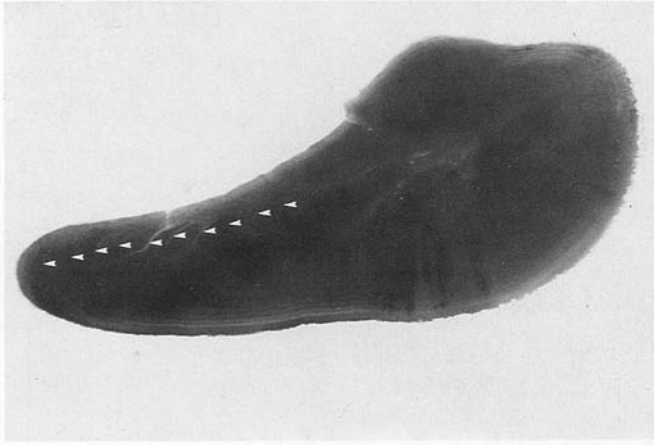


Fig. 1. Otolith section from a ten-year-old female of *Chionodraco hamatus*

equal probabilities of being correct. A "good" estimate of the mean lengths at age could be obtained by entering approximately the same number of "high" and "low" individual ages. However, when readings differed by two units, we adopted the intermediate integer. On the whole, 32 out of 39 females and 70 out of 79 males were aged.

Results

Forty specimens (ten of which were excluded) of *C. hamatus* were captured during 13 fishing operations by trammel-net and 140 during 10 fishing operations by gill-net. The gill-net's higher efficiency was probably linked to the fact that it stood higher on the sea bottom (the Channichthyidae are indeed semipelagic) rather than to different mesh size and/or selectivity. Most *C. hamatus* specimens were caught by entanglement, which is relatively independent of the body size and mesh size.

Even if no proper chi-square test for heterogeneity between the nets could be done because of the low number of fishes captured by the trammel-net, the two length frequencies appeared to be relatively similar. Consequently, they were lumped together for further analysis. Figure 2a, b shows the resulting length frequency of females and males, respectively. The size difference between sexes can be clearly appreciated (median TL of females = 398.5 mm, median TL of males = 358.5 mm).

Figure 3a, b depicts the log-transformed regression lines calculated on our length and weight data for females and males, respectively. In detail, the two regression lines are:

$$\begin{aligned} \text{LnEWt} &= 3.0064 \text{ LnTL} - 12.1911 \text{ (females, } R = 0.84) \\ \text{LnEWt} &= 3.5858 \text{ LnTL} - 15.5125 \text{ (males, } R = 0.83) \end{aligned}$$

where LnTL are the logarithms of total lengths and LnEWt the logarithms of weights drawn from the same specimens once they had been eviscerated. Original measures were expressed in millimetres and grams.

As total weights vary throughout the year in connection with the maturation cycle of the gonads, we believed that eviscerated weights could be better compared with future data from other periods and/or areas.

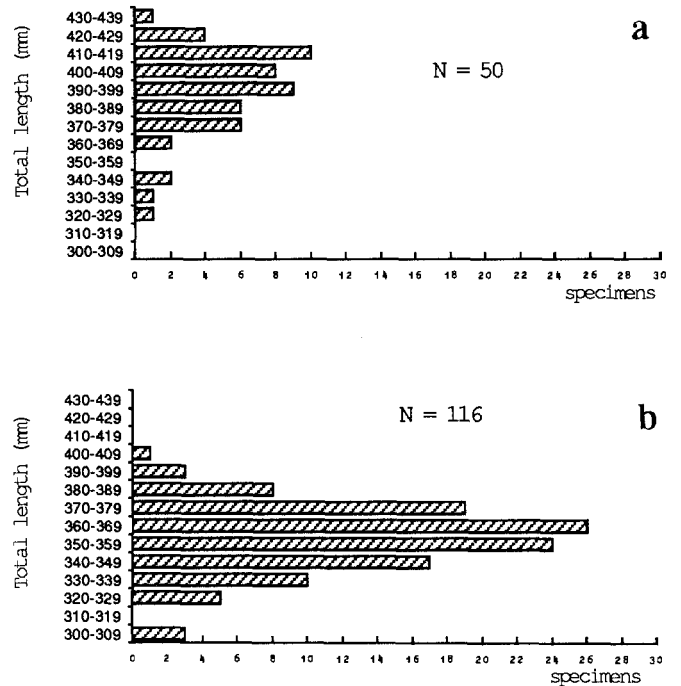


Fig. 2a,b. Length frequency of *Chionodraco hamatus* females (above) and males (below) caught in Terra Nova Bay in 1988

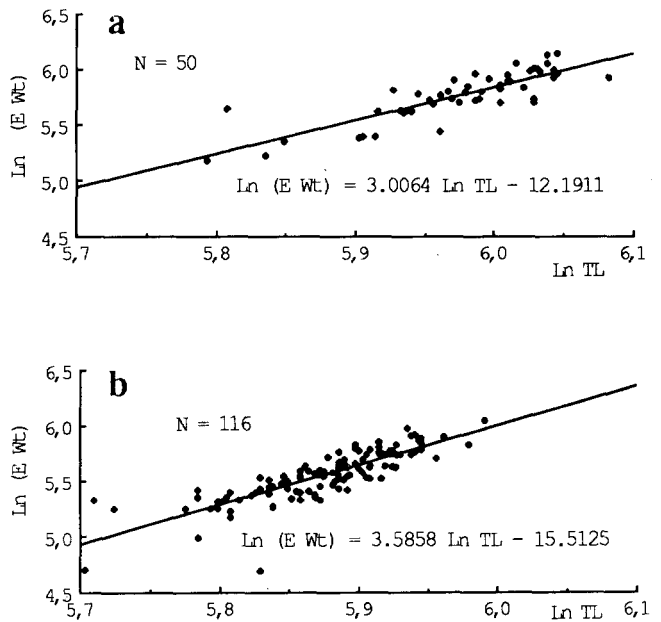


Fig. 3a,b. Regression line, logarithm of weights vs logarithm of total lengths (Ln EWt vs Ln TL), of *Chionodraco hamatus* females (above) and males (below) caught in Terra Nova Bay in 1988

Table 1 shows the age at length values estimated from our specimens by counting the annuli of single otolith sections. The classic Von Bertalanffy growth function was not fitted to the experimental data because we were dealing only with relatively old animals; such a fitting would probably have "smoothed" the extrapolation curve.

Table 1. Age composition of *Chionodraco hamatus* samples caught in Terra Nova Bay in 1988

AGE	FEMALES			MALES					Average TL ± SE			
	Total Length (mm)	Average TL ± SE		Total Length (mm)								
5 years	333	340.7 ± 4.1		300					325.0 ± 25.0			
	342											
	347											
6 years	328	353.7 ± 12.8		302	325	346	351	336.5 ± 4.7				
	366			322	330	347	355					
	367			325	332	348	355					
7 years	378	386.6 ± 3.4		331	340	350	358	367	351.9 ± 2.5			
	379			333	342	350	360	368				
	390			335	343	354	360	370				
	391			339	345	355	364	370				
	395			340	349	356	366					
8 years	382	399	399.8 ± 3.3		325	362	368	374	380	388	369.6 ± 2.5	
	392	400			356	362	371	374	381			
	393	407			358	363	372	376	382			
	396	414			360	364	372	379	382			
	398	417			361	365	373	379	382			
9 years	388	415	421	411.7 ± 4.4		362	390	380.8 ± 6.4				
	405	419	367			395						
	414	420	372			399						
10 years	385	422		409.5 ± 8.3								
	415											
	416											

Our *C. hamatus* samples were comprised mainly of breeders (Everson's stages 4 and 5) which consequently attained large TLs. Lack of young individuals was probably due more to bathymetric/geographic distribution than to our fishing gears, as the same fishing gear also caught some small specimens (less than 250 mm TL) from other Channichthyidae species very similar to *C. hamatus* in their body shape and maybe swimming behaviour (e.g. *Pageopsis macropterus* and *Chaenodraco wilsoni*).

Discussion

The sagittae sections showed a clear sequence of opaque/hyaline bands which complied well with Everson's description of annuli, so we assumed they expressed regular switches in the growth rate of fishes and were deposited a year apart. The annuli appeared in the sections with a very regular shape as well as with decreasing width from the nucleus to the external border as described by Everson (1980) and expected on the grounds of a positive correlation usually existing in fishes between TL and otolith size.

However, the individual ages obtained by examination of the otolith sections must be regarded simply as "apparent ages" since we could not really know which was the first true annulus among those located in the proximity of

the nucleus. Furthermore, we do not have proof that annuli are laid down yearly, as we had no means of comparison with other age estimates, e.g. correlating them with prominent peaks in the length frequencies of large samples composed of young year classes too.

The age data listed in Table 1 match well the report by Gubsch (1982) in which samples of *Chionodraco rastrospinosus* collected near both the South Orkney Islands and King George Islands were composed of year classes IV/VII in the length range 340–470 mm.

Our data show, if correct, that both sexes grow very slowly in the sampled length range, but females are on average larger than males of the same age and increase their TL at a greater rate. Moreover, the females seem to grow less as they get older (the growth rate passes from 33.2 mm/year between age classes VII/VIII to, paradoxically, negative growth for classes IX/X), whereas males follow a linear growth curve (approx. 11/18 mm per year).

In the end, the observation that most specimens present in our samples were breeders (Everson's stages 4 and 5) and males were somewhat younger (and smaller) than females (mean ages of males = 7.31 ± 0.96 years, mean age of females = 7.84 ± 1.46 years, $p < 0.001$, Student's test) could imply (even if other explanations are possible) that, in the area sampled, males come to maturity earlier than females.

Acknowledgements. We are much indebted to the personnel of the MAFF Laboratory of Lowestoft (East Anglia, UK) for preparing the otolith slides used in this study.

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