Note on otolith growth in elvers, *Anguilla anguilla* (L.), and the relative otolith size during somatic growth

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Abstract. Newly pigmented glass-eels were kept in a cold water quarantine, before being exposed to increased water temperature and weaned with cod roe and commercial dry feed. The otoliths started to grow before fish growth was observed. As a consequence, the relative otolith size increased. This increase was higher at higher temperature. During the first period with significant eel growth, no further change in the otolith/fish size ratio was observed. Initially, the otolith size was not correlated to fish size, but a positive correlation occurred in growing elvers. The results are discussed with reference to variation in relative otolith size of larger eels. As otoliths have frequently been used in back-calculation of previous eel size, further studies on factors determining otolith growth are suggested.

Keywords: eel elvers, otolith growth, relative otolith size

Introduction

The otoliths (sagittae) are the calcified structures most commonly used for age determination of eels (Liew 1974, Vollestad et al. 1988). Additionally, a linear relationship between otolith length (or the length of a particular radius) and total fish length has been taken as a criterion justifying back-calculation of previous eel length (e.g. Penaz and Tesch 1970, Rossi and Villani 1980, Moriarty 1983, Vollestad and Jonsson 1986, Paulovitz and Biro 1986, Fernandez-Delgado et al. 1989, Nagiçê and Bahnsawy 1990).

The research on otolith growth patterns of larvae and juvenile fish was intensified by the discovery of daily growth increments (Pannella 1971). Campana and Nielson (1985) critically reviewed how well otolith growth reflects fish age and growth at the daily level of precision. An uncoupling of the otolith-fish size relationship may occur under suboptimal conditions. At low temperature the otolith growth appears to cease, although body growth continues at a reduced rate (Marshall and Parker 1982). On the other hand, the otoliths continue to grow through periods of starvation and cessation of somatic growth (Brothers 1981, Campana 1983). Mosegaard et al. (1988) suggested that some metabolic activity, and thus an increasing function of fish size and temperature, is responsible for the otolith growth rate. Modified models have later been proposed (Secor and Dean 1989, Wright et al. 1990).

A daily formation of otolith increments is validated in Japanese elvers, *Anguilla japonica* T. and S. (Tsukamoto 1989), but the increment deposition is disrupted by low temperature as well as food deprivation (Umezawa and Tsukamoto 1991).

This study is based on European elvers, *Anguilla anguilla* (L.), reared at different temperatures in a long-term experiment on the sex differentiation of
eels. The eels were not size graded during their first eight months of culture due to a long period with high mortality and low growth rates. Otolith samples were collected on several occasions in order to get indications whether the relative size of otoliths could reflect metabolic state and/or recent growth history. The results are discussed with reference to observed residuals in otolith length versus fish length regressions from earlier studies on larger eels.

Material and Methods

Experimental fish and rearing conditions

Glass eels, caught in the English River Severn (Bristol Channel Fisheries Ltd.), were imported by a Swedish eel farm (Scandinavian Silver Eel AB, Helsingborg) in April 1991. About 38 800 specimens were kept in a cold water quarantine (3-5°C) for eight weeks, before being transported to the Institute of Freshwater Research, Drottningholm. During transport the water temperature accidently increased to 10-13°C.

At arrival all surviving eels were moved to one of the tanks in a recirculating freshwater system (Holmgren et al. 1992). During the first five days the water temperature in the tank was gradually raised from 9.3 to 17.0°C. Initially, the elvers were given frozen and subsequently thawed cod roe. Later on, the feed was served as a paste of cod roe and an increasing proportion of commercial dry feed (as powder and in granulated form). Manual feeding was performed 4-11 times per 24 hours. Finally, dry feed was given as hourly rations from a feeder.

After five weeks, the eels were randomly grouped and distributed into six tanks (about 1300 eels/tank). In two tanks the temperature was retained at 17°C. In another two tanks it was gradually raised to 20°C, and the final temperature in the last two tanks was 26°C. At daily basis, the feed rations were given in excess at all temperatures according to Seymour (1989). This particular study is based on samples of small eels, taken during the first eight months of culture, i.e. before the first grading.

General sampling of surviving and dying eels

The abundance and size distributions of surviving and dying eels were monitored by the sampling scheme listed in Table 1. Alive eels were anaesthetised using 0.12 g benzocaine/L of culturing water. Total length was measured to the nearest mm. Excess water was shaken off, and total weight recorded with the accuracy of 0.01 g. The pigmentation stage was defined according to Strubberg (1913) and Elie et al. (1982).

Special sampling of otoliths

On six occasions (Table 1), the sagittal otoliths were collected in random samples of eels, with individual data on total length, total weight and pigmentation stage. The right sagitta was mounted in clear nail polish on an objective slide. By use of an ocular micrometer, otolith length and width was measured with the accuracy of 0.01 mm. Otolith length was defined as the maximum length parallel to the sulcus, and otolith width as the maximum length perpendicular to the defined otolith length. Relative otolith length was computed as the otolith length in percent of total fish length.

Statistical analyses

The program package „SPSS/PC+”, V4.0 (Norusis 1990) was used in the computations. Parametric t-tests, analyses of variance (ONEWAY) and correlation tests were used, as Kolmogorov-Smirnov Goodness of Fit tests and plotted histograms indicated no or only minor deviations from assumed normal distributions and equal variances.
Results

Total length and pigmentation stage of surviving elvers

The total length of the imported elvers varied between 62-80 mm (mean 71.0 mm). There was no difference in total length between samples taken on 4 April, 28 May, 9 June and 10-11 July (Fig. 1), i.e. the elvers did not grow during eight weeks in the cold water quarantine or during six weeks of weaning at 17°C. Mean total weight decreased from 0.33 to 0.23 g (t-test, P=0.001). As only fresh weights were taken, these data were considered to be of minor value in the further discussions. In August some growing individuals (up to 100 mm) were observed at 20 and 26°C, but this did hardly affect the mean value. Ten weeks later, the majority of surviving elvers had total length above the initial mean value and a considerable number were larger than the initial maximum.

When the elvers arrived from England in April, they were already more or less pigmented (Table 2), with a mode in stage VIA2. After eight weeks in quarantine, 52.7% of initial numbers arrived to the laboratory in Drottningholm. At this point, 95% of surviving eels had reached the last elver stage, VIA4. During the following weeks, some eels became fully pigmented, stage VIB, but
eels in earlier stages, VIA₃ and VIA₄, still occurred in October.

**Otolith length of surviving elvers**

The smaller subsamples taken for otolith measurements (Table 1) were considered representative, as estimates of mean total length and standard deviation were comparable to that of the larger samples (Fig. 2a and Fig. 1, t-tests, P-values 0.05). There was no difference in otolith length between samples taken 0, 8 or 9 weeks after arrival to Sweden (Fig. 2b, ONeway, P=0.251). The pooled mean value was 0.349 mm, with a coefficient of variation of 6.5%. In these initial samples there was no difference in otolith length between stages of pigmentation (ONEway, P=0.314). In July, after another five weeks at 17°C, the otolith length had increased compared to the pooled material of earlier samples (t-test, p.001), and in August virtually all otoliths were larger than the initial maximum. Additionally, the otoliths were significantly larger at 26°C than at the lower temperatures (ONEway, P=0.0003).

**Otolith length in relation to Total fish length**

The small subsamples of living eels taken on 4 April, 28 May and 9 June, and the subsample of dying eels on 9 June, were all homogeneous with respect to total length and otolith length. In these initial samples, there was no correlation between otolith length and total length (Fig. 3). The relative otolith length varied between 0.42-0.60%, with a mean value of 0.50%. From June to August, when the otoliths increased in length in spite of no change in

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Table 2

Pigmentation stage in samples of living elvers, from arrival to Sweden (Helsingborg) 4 April 1991 and during the following months until October (see specifications and sample sizes in Table 1). Listed figures are the relative frequencies (%) per class

<table>
<thead>
<tr>
<th>Number of weeks in Sweden</th>
<th>VB</th>
<th>VIA₀</th>
<th>VIA₁</th>
<th>VIA₂</th>
<th>VIA₃</th>
<th>VIA₄</th>
<th>VIB</th>
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<td>36</td>
<td>22</td>
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<td>0</td>
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<td>0</td>
<td>1</td>
<td>1</td>
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<td>0</td>
<td>0</td>
<td>15</td>
<td>45</td>
<td>40</td>
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<tr>
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<td>0</td>
<td>0</td>
<td>10</td>
<td>60</td>
<td>30</td>
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<tr>
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<td>0</td>
<td>0</td>
<td>5</td>
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<td>11</td>
<td>57</td>
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</table>
total length, there was a linear increase in relative otolith length (Fig. 2c). During the following weeks until October, there was no further change in relative otolith length at 20 or at 26°C (t-tests, P-values 0.756 and 0.430, respectively), indicating an isometric relation between the length increases of otoliths and somatic tissue. This was also expressed as positive correlations between otolith length and total length, at both temperature regimes (Fig. 4).

Discussion

With a set of data indicating no increase in total length of glass eels/elvers during at least four months, one might be lost in efforts to explain the growth failure. Such a discussion could include aspects on keeping already pigmented elvers in cold water, on stress related to transport and on infections with fungi and bacteria during the weaning period. That is definitely not the object of this study.

In this experiment, not more than 12.4% of initial numbers survived from April to October. Neither total length nor otolith length was repeatedly measured on individual elvers. Nevertheless, the insignificant change in total length distributions until August, and the fact that most otoliths in August were larger than the largest otolith before and after eight weeks in the cold water quarantine, together indicate uncoupling between otolith and somatic growth. Umezawa and

Figure 2. Total length, otolith length and relative otolith length of elvers in the smaller subsamples for the otolith study (see Table 1). Symbols for mean values, standard deviations and total ranges are used as in Figure 1.
Tsukamoto (1991) studied otolith increment deposition of Japanese elvers during 50 days. In unfed groups and/or at low temperature (12°C) total length remained unchanged and otolith growth was sharply reduced compared to the other groups. In the present study, feed was provided in excess throughout the period of weaning and culturing in heated water. Surviving elvers were probably feeding, although initially at insufficient rates for somatic growth. The growth of their otoliths might be seen as an index of metabolic activity (Mosegaard et al. 1988, Wright 1991). The observation of largest relative otolith length at the highest temperature (26°C) in August is an accordance with the highest expected metabolic rate.

In the Loire estuary, mean total length decreased and otolith radius increased from the glass eel stages (VA/VB) to the late elver stages (VIA₃/VIA₄).
Within each pigmentation stage the coefficient of variation (CV) was about 5% in total length and about 10% in maximum otolith radius. In the initial samples from the present study, the mean otolith length was 0.349 mm with a CV of 6.5%, which was not explained by variation in total length or pigmentation stage. The hypothesis that otolith growth reflects metabolic scope has been adopted in studies of salmonids, where early fry size was poorly predicted by otolith size. Selection for fish larger otoliths at emergence was observed in brown trouts which later established territories (Titus and Mosegaard 1991).

Furthermore, otolith size and dominance status at first feeding were good predictors of subsequent growth of Atlantic salmon (Metcalfe et al. 1992). The otolith size of glass eels can not as simply as in salmonid alevins be explained by standard metabolic rate. The glass eels have previously been feeding and growing as leptocephali.

When the cultured elvers began to increase in total length, there was at least initially no further increase in relative otolith length. In October this isometric relationship between otolith and somatic growth was expressed as positive correlations between otolith and total length at both 26°C ($r=0.900$) and 20°C ($r=0.817$). Such levels of correlations have earlier been taken as acceptable criteria for back-calculations of previous eel size (Rossi and Villani 1980, Vollestad and Jonsson 1986, Nagiec and Bahnsawy 1990). In other fish species, it has been shown that slow growing fish have larger otoliths than fast growing individuals of the same size (Secor and Dean 1989, Reznick et al. 1989). Growth back-calculations usually assume that the relationship between fish and otolith length is linear through time, at least within individuals. Campana (1990) reduced back-calculation bias by defining individual fish trajectories and a biological determined intercept.

The present study on elvers is a part of a long term experiment. As preliminary results it can be mentioned that the eels which first attained total weights above 150 g had relative otolith lengths within the same range as glass eels (Holmgren, unpublished results). This might indicate that the otolith growth in relation to fish growth can switch repeatedly within an individuals life time. Further research on the subject is suggested. For the time being, it is recommended to consider uncoupling between otolith and somatic growth as a possible bias when interpreting data from back-calculations.

**References**


Norusis M.J. 1990 – SPSS/PC+ V.4.0. Base manual for the IBM PC/XT/AT and PS/2. SPSS inc., Chicago, USA.


