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1 **Age estimation obtained from analysis of octopus (*Octopus vulgaris***

2 **Cuvier, 1797) beaks: improvements and comparisons**

3

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14

14 **ABSTRACT**

15

16 Two methods are currently available for age estimation in octopus beaks. They have  
17 been applied to the same specimen from a sample of 30 individuals of *Octopus vulgaris*  
18 caught in central-eastern Atlantic waters. These techniques aim at revealing growth  
19 increments in the Rostrum Sagittal Sections (RSS) and Lateral Wall Surfaces (LWS) of  
20 octopus upper and lower beaks. Both methods were improved to reduce the time of  
21 sample preparation and to enhance the appearance of the increments. For each  
22 individual, two independent readings were done for upper and lower beak sections, as  
23 well as for the lateral wall surfaces. Vertical reflected light (epifluorescence) and Image  
24 Analysis System were shown to be useful in the observation and analysis of the  
25 sequence of increments. Precision of the ageing, increment counts obtained by both  
26 techniques, and increment widths were discussed. Using upper beak RSS led to more  
27 precise age estimates, whereas preparing LWS was quicker and simpler, and revealed a  
28 higher number of increments. Therefore, our study recommends counting growth  
29 increments in LWS of beaks to age adult common octopus.

30

31 *Keywords*

32

33 *Octopus vulgaris*, age, growth, beaks, techniques

## 34 1. Introduction

35

36 Determination of age and growth is critical to understand the life history of  
37 harvested species and to model the dynamics of their populations. Sound knowledge on  
38 life history and population dynamics is essential for assessment and management  
39 purposes. Identifying and interpreting growth increments in calcified structures (otoliths  
40 and scales of fish, statoliths of cephalopods, among other structures) produce reliable  
41 estimations of the absolute age of wild marine animals (Boyle and Rodhouse, 2005). In  
42 spite of the difficulties raised by the age determination in cephalopods, those ageing  
43 methods based on the study of incremental growth structures (Bettencourt and Guerra,  
44 2000; Lipinski and Durholtz, 1994) are considered the most appropriate for exploited  
45 species of this group. Other available methods (Caddy, 1991) such as length frequencies  
46 are not suitable for cephalopods, since this group has high and variable growth rates,  
47 short life cycles and massive mortalities after spawning (Jereb et al., 1991; Perales-  
48 Raya, 2001; Semmens et al., 2004).

49 The common octopus *Octopus vulgaris* Cuvier, 1797 is one of the most important  
50 target species in the world, with catches of about 42 420 t/year for the period 2003-2007  
51 (FAO, 2009). However, there is still not a validated and standardized age determination  
52 method for using on *O. vulgaris*, mainly due to the uselessness of statoliths for ageing  
53 species from the Octopodidae family. Recently, Doubleday et al. (2006) and Leporati et  
54 al. (2008) validated the daily deposition of increments in stylets of adults *Octopus*  
55 *pallidus* Hoyle, 1885 of known age. The high mortality of the paralarvae in captivity of  
56 *O. vulgaris* has not yet allowed the obtaining of known-age adults for validation  
57 purposes. However, preliminary results using chemical marking in stylets (Hermosilla  
58 et al., 2010) and beaks (Oostuizen, 2003; Perales-Raya, unpublished results) have

59 shown a daily deposition of increments in adults of this species, although definitive  
60 validation is still necessary for ageing common octopus.

61 As beaks are present in all cephalopod species (Mangold and Bidder, 1989), any  
62 improvement in their preparation technique for ageing purposes should be useful to  
63 many commercially exploited species of this group. Beaks are composed of a chitin-  
64 protein complex (Hunt and Nixon, 1981) and secreted by a single layer of tall columnar  
65 cells, known as beccublasts that are responsible for their growth (Dilly and Nixon,  
66 1976). The chitinization and hence growth process related to lateral walls and rostrum  
67 takes place from the rostrum tip to the wing edges (Cherel and Hobson, 2005; Miserez  
68 et al., 2008).

69 The beaks are structures easy to extract and manipulate. The previous freezing of the  
70 animal (samples from industrial fisheries are obtained frozen) has no effect on the  
71 visualization of the growth increments. Another advantage is that microstructures are  
72 preserved in the beak sections after being prepared according to our method. However,  
73 the possible erosion of the rostral tip during the life of the animal may bias age  
74 determination and has to be taken into account. Sections of other hard structures, such  
75 as stylets, have been recently used for octopus ageing with good results. Nevertheless,  
76 microstructure disintegration has been reported within several minutes after preparation  
77 (Doubleday et al., 2006) and the sections showed significant cracks when the animal  
78 had previously been frozen (Sousa Reis and Fernandes, 2002).

79 Octopus beaks have been used for ageing by Raya and Hernández-González (1998)  
80 who developed a method using sagittal sections of the rostral area. Later, Hernández-  
81 López et al. (2001) proposed a technique using the inner surfaces of lateral walls, as  
82 previously done by Clarke (1965) for *Moroteuthis ingens*.

83 The aims of this study were: (1) to improve and simplify the present techniques for  
84 revealing growth increments in the beaks of the common octopus; (2) to estimate the  
85 precision of the increment counts in upper and lower beak sections and lateral wall inner  
86 surfaces; (3) to compare, for each sampled animal, the number of increments counted in  
87 the upper and lower beak sections, and in the lateral wall inner surfaces; and (4) to  
88 establish the best method for counting growth increments in the beaks of the common  
89 octopus.

90

91

## 92 **2. Material and Methods**

93

94 The study was carried out with a sample of 30 frozen animals from both sexes,  
95 ranging in total body weight from 90 to 5361 g (Table 1). These individuals were  
96 caught during 2007 in central east Atlantic waters (off Mauritania) by the Spanish  
97 industrial freezer trawler fleet. Once thawed, specimens were weighed and their beaks  
98 removed, cleaned and preserved in 70% ethanol. Before preparation, the beaks were  
99 rehydrated in distilled water for several days. The upper and lower beaks were weighed  
100 (mg) and the main lengths (as defined by Clarke, 1986) were obtained (mm): Hood  
101 Length (HL), Height (H), Crest Length (CL) and Rostral Length (RL).

102 Rostrum sagittal sections (RSS) were prepared following an improved technique  
103 based on the method developed by Raya and Hernández-González (1998) for upper and  
104 lower beaks. The rostrum area was cut with scissors and mounted in polyester resin with  
105 the lateral side facing up. After hardening of the applied resin cover, the piece was  
106 ground down with 1200 grit carborundum sandpaper. After reaching the central plane  
107 we polished with 1  $\mu\text{m}$  diamond paste to obtain a smooth surface of the sagittal section.

108 This section revealed a banding pattern from the rostral tip to the joining point of the  
109 hood and the crest (Fig. 1). Since the increments were visible under vertical reflected  
110 light (ultraviolet epi-illumination, if possible), it was not necessary to sand down both  
111 sides like other cephalopod hard structures such as statoliths and stylets.

112 Lateral wall surfaces (LWS) were prepared based on the method described by  
113 Hernández-López et al. (2001) for the upper beaks. We sagittally sectioned them with  
114 scissors to obtain two symmetrical half beaks which were cleaned by hand with water to  
115 remove any mucus attached to the inner surfaces of lateral walls. The LWS were also  
116 epi-illuminated, but here the violet light led to better results than ultraviolet one, due to  
117 the darkness of this beak zone.

118 The magnification chosen for RSS ranged between 200X and 400X, and we used  
119 50X for viewing the LWS. Increments were identified and marked under the *live* camera  
120 mode (which allows for multi focal imagery), and several photos were taken to cover  
121 the whole studied area. We measured the distances between growth marks (increment  
122 width) and performed the increment count with an image analysis system (IAS,  
123 software Age&Shape). When extrapolation was necessary because increment visibility  
124 was poor (i.e. first and last portions of the anterior and posterior borders of the LWS),  
125 the IAS carried it out by using the average width of the nearest and most visible  
126 increments. To avoid tip erosion effects, the first increments located at the rostral tip of  
127 the RSS were counted in the dorsal area.

128 Precision is defined as the reproducibility of repeated measurements (age readings)  
129 on a given structure, whether or not those measurements are accurate (Kalish et al.,  
130 1995). The same trained reader made two repeated counts. Coefficients of Variation  
131 (CV) of the age estimates were calculated to assess precision. This method is favoured  
132 for microstructure studies as it is statistically more rigorous and thus more flexible than

133 the use of average percent error (APE) because of the absence of an assumed  
 134 proportionality between the standard deviation and the mean (Campana, 2001). For each  
 135 sampled individual, we calculated the CV for the six readings: two for the upper beak,  
 136 two for the lower beak, and two for the lateral walls. We obtained a total of 180  
 137 readings. For this study, CV was calculated as the ratio of the standard deviation over  
 138 the mean:

139

$$140 \quad CV = 100\% \times \sqrt{\frac{(R1 - R)^2 + (R2 - R)^2}{R}}$$

141

142 where  $R1$  and  $R2$  were the number of increments from the first and the second reading  
 143 respectively;  $R$  was the mean number of increments for both readings.

144 The normal distribution of the data was checked with the one-sample Kolmogorov-  
 145 Smirnov test. Homogeneity of the variances was assessed with the Levene's test.  
 146 Differences in both readings ( $R1$  and  $R2$ ) for each preparation (upper and lower RSS,  
 147 LWS) were compared by performing a one-way analysis of variances (ANOVA) [Zar,  
 148 1984], a Tukey's honestly significant difference (HSD) test and a Bonferroni's multiple  
 149 range post hoc test. When a normal distribution and/or homogeneity of the variances  
 150 were not achieved, data were subjected to a non-parametric Kruskal-Wallis test and a  
 151 Games-Howell post hoc test. For all the statistical tests performed, significance level  
 152 (statistically different readings) was chosen to be  $P < 0.05$ . The statistical analysis was  
 153 performed using the SPSS package (version 9.0) from SPSS Inc.

154 The relationships between the number of increments and the beak measurements  
 155 (HL, H, CL and RL) were calculated, as well as the relationships between the increment  
 156 counts and the total body weight. Relationships calculated using the second readings  
 157 ( $R2$ ) showed the highest regression values when plotted against beak measurements.



158 Besides, the second reading is supposed more reliable because of greater experience and  
159 practice.

160

161

### 162 **3. Results**

163

#### 164 *3.1. Methodological improvements*

165

166 Although 70% ethanol was used for the preservation of the beaks during the  
167 biological sampling, our laboratory observations recommend preserving them in  
168 distilled water at a cold temperature (around 5 °C) (Perales-Raya, unpublished results).

169 The beaks preserved in ethanol for long time periods showed the poorest visibility of  
170 the increments, probably because ethanol dehydrates the beaks. Instead of using  
171 sections of beaks, as described by Raya and Hernández-González (1998), our cutting  
172 technique allowed the embedding of only the rostrum area, thus reducing time for  
173 grinding and polishing. Etching the section surfaces was not necessary as the ultraviolet  
174 light allowed the obtaining of more information from the deeper planes.

175 Vertical reflected light (ultraviolet for the sections and violet for the lateral walls)  
176 gave good results for observation of increments. Fig. 2 shows the sequence of  
177 increments in the inner surface of the lateral walls, from the anterior to the posterior  
178 edge of these structures.

179 In the upper and lower RSS, patterns of increments were observed from the rostrum  
180 tip to the joining point of the hood and the crest (Fig. 3A). The increments located at the  
181 rostrum tip were lost, probably due to the erosion of the rostrum during the feeding  
182 process. To avoid the tip erosion effects we usually counted the first increments in the

183 dorsal area of the rostral sections, where defining a transect for counting a sequence of  
184 thin increments until the dorsal border of the hood was possible (Fig. 3B).  
185 Unfortunately, the lateral walls had no alternative reading zones, but it appeared that  
186 feeding erosion (if it exists) did not affect in the same way the readings performed in the  
187 anterior region of the lateral wall area as it did in the rostral tip of the sections.

188

### 189 3.2. Ageing precision, reading comparisons and growth curves

190

191 Table 1 shows the second reading values ( $R2$ ) and Table 2 shows the results of mean  
192 CV for the three preparations of each sampled individual (upper beak RSS, lower beak  
193 RSS and LWS). RSS of the upper beak showed to be the most precise technique.  
194 Although the CV obtained were quite similar, the results showed that the less precise  
195 readings were performed in the lateral walls.

196 Significant differences were found in the number of increments between readings of  
197 LWS and upper beak RSS both in repeated readings  $R1$  ( $df = 89$ ,  $F = 7.37$ ,  $P = 0.001$ )  
198 and  $R2$  ( $df = 89$ ,  $F = 6.91$ ,  $P = 0.002$ ), according to ANOVA and HSD Tukey post-hoc  
199 test, with a mean difference of 38 increments more in the LWS with respect to upper  
200 RSS. However, HSD Tukey test did not show significant differences between lower and  
201 upper RSS ( $P = 0.055$  for  $R1$ , and  $P = 0.123$  for  $R2$ ). Even if HSD Tukey test did not  
202 find significant differences between lower RSS and LWS ( $P = 0.315$  for  $R1$ , and  $P =$   
203  $0.198$  for  $R2$ ), a mean difference of 16 increments more was observed in the LWS.

204 Fig. 4 shows the relationship between the total body weight of the sampled  
205 individual and the number of increments counted in RSS and LWS. For the same  
206 weight, a higher number of increments was counted in LWS (formula in the figure).  
207 Upper beak RSS produced the lowest counts. Regression values were:  $Y = 27.978X^{0.249}$

208 ( $r^2 = 0.75$ ) for the LWS;  $Y = 26.277X^{0.241}$  ( $r^2 = 0.49$ ) for the lower beak RSS;  $Y =$   
209  $31.395X^{0.200}$  ( $r^2 = 0.54$ ) for the upper beak RSS. Poor relationships were observed  
210 between the number of increments in lower beak RSS and the total body weight for  
211 animals over 2 000 g, and between upper beak RSS and total body weight for animals  
212 over 3 000 g.

213 Fig. 5A shows the results of the beak growth. The best regression (power model;  $R^2$   
214  $= 0.76$ ) was obtained plotting the weight of the upper beak versus the number of  
215 increments ( $R^2$ ) in the LWS. Concerning beak measurements (Fig. 5B), the best  
216 regression fit (power model;  $R^2 = 0.75$ ) was obtained for the hood length (HL) of upper  
217 beak versus the number of increments ( $R^2$ ) in the LWS.

218 Mean widths were calculated for each increment counted in the second reading ( $R^2$ )  
219 of the upper beak RSS, where the highest reading precision was achieved (Fig. 6A).  
220 Mean widths were also calculated for each increment counted in the second reading  
221 ( $R^2$ ) of the LWS, where the highest number of rings were counted (Fig. 6B). Figure 6A  
222 shows that the approximately first 50 increments (counted in the dorsal area of the RSS)  
223 were much thinner than rest of the growth marks (counted along the main axis of the  
224 RSS). This figure also shows a constant decreasing trend until approximately increment  
225 number 180, being highly scattered afterwards. Figure 6B showed a more constant trend  
226 in the mean distances of each increment in the LWS, the values being mostly comprised  
227 between 75 and 100 microns. Also here, dispersion increased from increment 180  
228 onwards.

229

230

#### 231 4. Discussion and conclusions

232

233 Upper and lower beak RSS produced similar readings in terms of increment  
234 numbers, although the upper beak showed to give more precise age estimates. Readings  
235 performed in the LWS produced higher increment numbers than the readings in RSS  
236 (average of 38 increments more). In spite of the lower precision of the age readings in  
237 the LWS, this technique showed to be the simplest and quickest one. Those differences  
238 could be due to the fact that there were more increments to count in the LWS than in the  
239 upper and lower RSS.

240 Preliminary laboratory results of validation obtained so far indicate that increments  
241 seem to be laid down on a daily basis (Oosthuizen, 2003; Perales-Raya, unpublished  
242 results) in both of the studied octopus beak zones. For octopus paralarvae, increments  
243 have been shown to deposit daily on the lateral walls (Hernández-López et al., 2001).

244 Two hypothesis are suggested to explain the viewing of more increments in the  
245 LWS of the beak: (i) feeding erosion of the rostral tip, and even in the dorsal-posterior  
246 area of the hood (where first increments were counted), could have biased increment  
247 count toward underestimation; or (ii) increment number is underestimated in the RSS  
248 because growth marks start depositing in the rostrum several weeks after hatching. As  
249 the feeding erosion is greater in the anterior region of the beak and we performed the  
250 increment counts in the dorsal edge of the hood (where growth marks were identifiable  
251 until the posterior end), the underestimation would be negligible. At hatching, the  
252 buccal mass is fully formed and functional (Nixon and Mangold, 1996), but maybe at  
253 this stage, when the beaks are transparent and oral denticles are present in both upper  
254 and lower jaws of the paralarvae (Villanueva and Norman, 2008), the formation of  
255 internal increments inside the rostrum has not yet started.

256 When looking at the average widths of the increments, upper beak RSS showed a  
257 general decreasing trend for the increments counted along the central axis of the RSS

258 starting at approximately increment 90. As for this value the increment width is the  
259 widest, we can think that the fastest growth corresponds to the age of about 90 days.  
260 The thin increments counted in the dorsal area of the RSS showed an increasing trend  
261 from the edge to approximately increment 50, even if this increasing trend was not  
262 comparable to those of the increments counted along the central axis. From about  
263 increment 180, the points were highly scattered. This fact could be due to the lower  
264 number of available samples with more than 180 increments for calculating the average  
265 widths, and to the higher variability of widths observed in the posterior edge of the  
266 counting area. The trend of the average increment width observed in the LWS  
267 preparations seems to reflect the probable more constant growth of those beak surfaces.  
268 Values were also more scattered from increment 180 onwards.

269 Considering all the facts presented and discussed in this study, we recommend using  
270 the LWS to perform growth increment counts in the beaks of common octopus. Even if  
271 the readings were less precise than those performed in the RSS, the method is simpler  
272 and quicker. In addition, LWS are less eroded during the life of the octopus, thus  
273 avoiding the eventual underestimation problems. When daily deposition of those  
274 increments will definitively be validated for common octopus beaks, counting the  
275 growth marks of the lateral walls appears as the most suitable ageing technique for  
276 *Octopus vulgaris*.

277

278

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280

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284

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- 364



365 **Fig. 1.** Drawing of upper beak sagittal section. Reading area inside the left circle, where it is  
 366 shown the rostral section and the increments.

367  
 368 **Fig. 2.** Increments in the inner surface of lateral walls (50X): (A) anterior region with the first  
 369 increments showing with an arrow the extrapolated area; (B) medium region with increments;  
 370 (C) posterior region with last increments where arrow shows the extrapolated area of the edge.

371  
 372 **Fig. 3.** (A) Appearance of increments in the central area of the beak sections (200X). (B) Dorsal  
 373 region of the beak sections, where it was possible to count thin increments until the dorsal  
 374 border of the hood (at the top of the image, magnification 300X)

375  
 376 **Fig. 4.** Relationship between total weight (g) and number of increments of the octopus beaks  
 377 (*Octopus vulgaris*). Square: lower section, circle: lateral wall, cone: upper section. Black curve:  
 378 regression for lateral walls, equation above.

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 380 **Fig. 5.** (A) Relationship between number of increments in the lateral wall and upper beak  
 381 weight (mg) of the octopus beaks (*Octopus vulgaris*). (B) Relationship between number of  
 382 increments in lateral wall and main beak measurements of upper beak. x: height, square: rostral  
 383 length, cone: hood length, circle: crest length. The best regression values were obtained for the  
 384 hood length and its regression line is displayed in the graph.

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 386 **Fig. 6.** (A) Trend of increment width in the upper sections, and (B) trend of increment width in  
 387 the lateral walls of the octopus beaks (*Octopus vulgaris*).

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**Table 1**

393 Sampling details for the octopus *Octopus vulgaris*  
 394 beaks used in the present study. R2: Number of  
 395 increments of the second reading.

Total weight (g)	R2 Upper Beak	R2 Lower Beak	R2 Lateral Wall
91	78	78	74
532	125	120	131
537	101	92	138
605	85	107	135
647	130	101	155
708	139	167	165
724	114	143	147
900	105	108	127
925	129	160	221
1 074	159	177	168
1 106	107	117	138
1 277	112	175	176
1 315	112	127	162
1 416	128	127	202
1 526	137	172	153
1 569	149	157	173
1 741	148	205	171
1 868	140	162	196
1 879	157	173	163

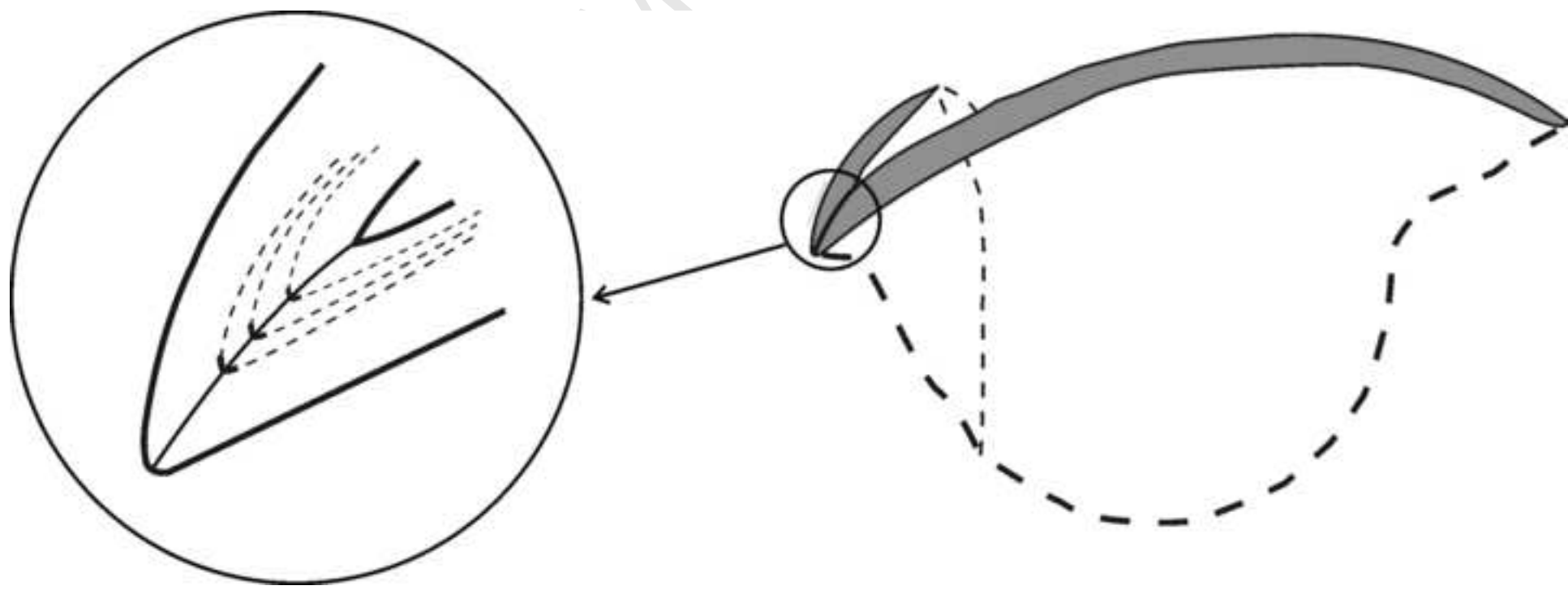
2 000	151	158	187
2 176	128	167	194
2 211	156	193	198
2 485	146	143	194
3 065	167	218	156
3 217	243	298	235
3 431	192	227	207
	Mean	Confidence interval	N
	CV	(+/- 95%)	
Rostrum Sagittal Section (Upper Beak)	3.93	1.29	30
Rostrum Sagittal Section (Lower Beak)	4.49	1.46	30
Latera Wall (Upper Beak)	4.84	1.47	30
3 765	160	229	237
4 522	126	131	211
5 156	136	166	227
5 361	181	152	243

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**Table 2**

Precision of the two counts for Section Upper Beak, Section Lower Beak and Lateral Wall in the common octopus (*Octopus vulgaris*). CV (Coefficient of variation), N (number of samples).

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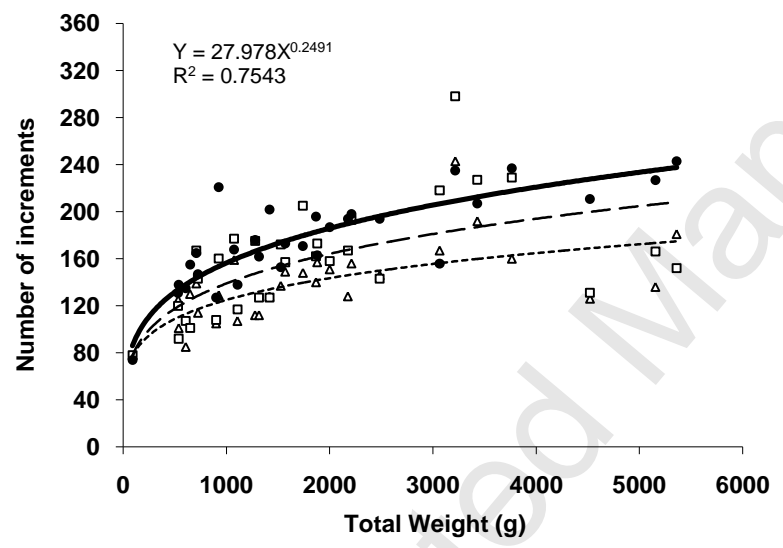




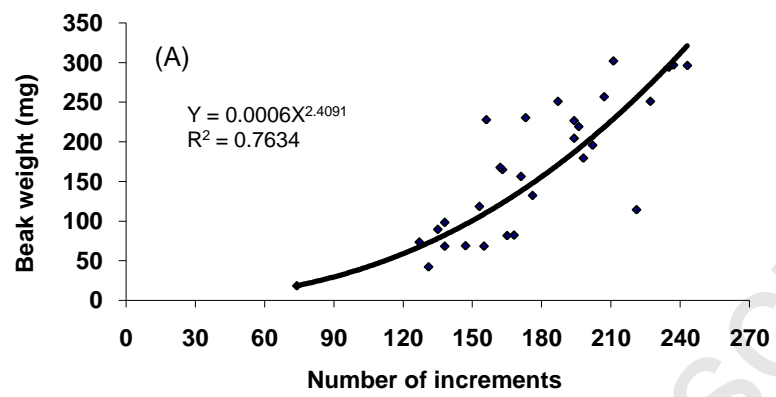




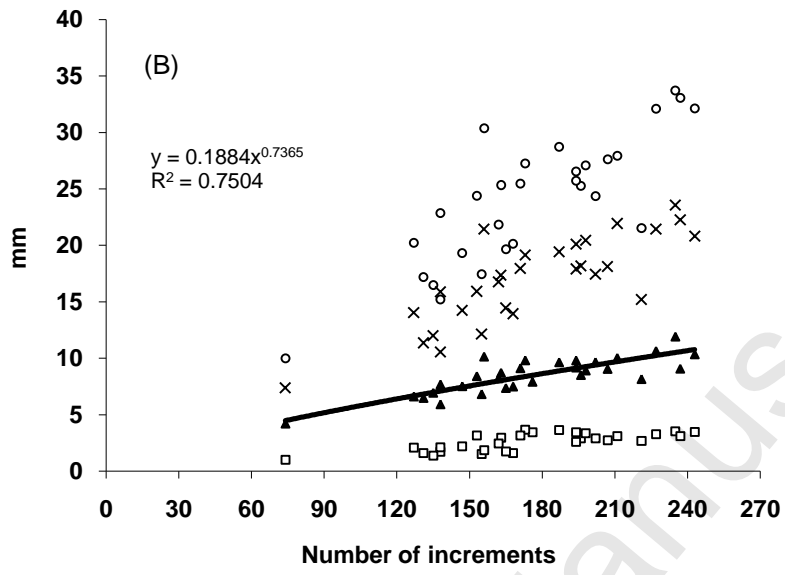




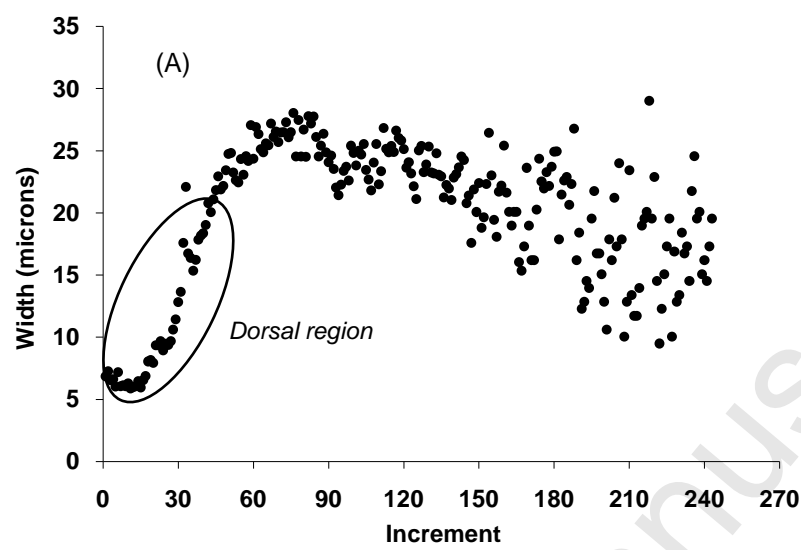




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