Reproduction and nutritional values of the edible limpet *Nacella magellanica* (Gastropoda: Patellogastropoda)

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Summary: *Nacella magellanica* is an edible limpet that has been consumed since pre-Hispanic times by human populations along the coasts of Patagonia, but studies of its nutritional value and reproduction are not yet available. We investigated the reproductive cycle and the seasonal variation in the nutritional composition (proteins, lipids and carbohydrates) of the whole body of this limpet in order to analyse some aspects of its importance as a formal fishery resource. Throughout a single year, the spawning period extended over all months except June in males, while females spawned from late winter to spring, with an increase from August to November. The nutritional data obtained for *N. magellanica* are within the ranges of widely consumed species of molluscs, with annual average values of 29.8% proteins, 2.7% lipids and 1.8% carbohydrates. The best nutritional values for human consumption (highest concentration of proteins, body weight), avoiding the reproductive period, were found in April but taking into account the minimum size of capture. Our results are useful for increasing the policies aimed at managing this abundant edible limpet as a formal resource, since it is widely consumed in southern South America.

Keywords: marine gastropods; fishery resources; biochemical composition; reproduction; limpets.

Reproducción y valores nutricionales de la lapa comestible *Nacella magellanica* (Gastropoda: Patellogastropoda)

Resumen: *Nacella magellanica* es un caracol comestible, consumido desde tiempos prehispánicos por poblaciones a lo largo de las costas patagónicas. Sin embargo, aún no existen estudios sobre sus valores nutricionales ni su ciclo reproductivo. En este trabajo se estudió el ciclo reproductivo y la variación estacional en su composición nutricional (proteínas, lípidos y carbohidratos) en todo el cuerpo de la lapa, para analizar algunos aspectos de su importancia como recurso pesquero formal. El periodo de liberación de gametas en los machos se extendió a lo largo de todo el año, mientras que en las hembras fue desde el invierno tardío hasta la primavera, con un incremento desde agosto a noviembre. La información nutricional obtenida para *N. magellanica* se encuentra dentro de los rangos de otras especies de moluscos ampliamente consumidos, con promedios anuales de proteínas de 29.8%, 2.7% de lípidos y 1.8% de carbohidratos. Los mejores valores nutricionales para el consumo humano (mayor concentración de proteínas y peso corporal), evitando el período reproductivo, se encontraron durante abril, aunque habría que tener en cuenta el tamaño mínimo de captura. Nuestros resultados son útiles para incrementar las políticas existentes para el manejo de esta abundante lapa comestible como un recurso formal, ya que es consumida ampliamente en el sur de Sudamérica.

Palabras clave: gasterópodos marinos; recursos pesqueros; composición bioquímica; reproducción; lapas.


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INTRODUCTION

Marine molluscs are abundant and easy to collect, and possess healthful properties such as a high proportion of proteins and a low level of saturated fatty acids (Usai and Busarova 1984, Manzano and Aranda 1998, D’Armas et al. 2010), making them a desirable seafood. Several studies have highlighted the dietary value of marine resources, and particularly of sea snails, for human consumption (Leiva and Castilla 2002, Vasconcelos et al. 2008, Bigatti et al. 2015). Therefore, knowledge of the nutritional composition of potential marine food can increase the possibilities for alternative foods worldwide. Studies on marine invertebrates have demonstrated that the reproductive strategies provide valuable information for fisheries regarding marine population dynamics (Underwood 1979, Collin 2003, Bigatti et al. 2008).

Coastal gastropod species such as limpets are consumed worldwide (Branch 1975, Pombo and Escofet 1996, Castilla 1999). In southern South America, limpets of the genus *Nacella* are commercially exploited in Chile and captured manually in Argentina from the provinces of Tierra del Fuego (Conti et al. 2012) to Chubut. The species *Nacella magellanica* was a resource widely used by the native populations of the Patagonian and Fueguian coasts in pre-Hispanic times (Gómez Otero et al. 1998, Orquera 1999, De Aranzamendi et al. 2009). At present, the harvesting of gastropods in Atlantic Patagonia is restricted to artisanal fisheries (Elías and Pereiro 2003). Therefore, owing to its high abundance, the current consumption and the simplicity of collection during low tides, *N. magellanica* could become a formal artisanal fishery resource on the Patagonian Atlantic coasts.

*N. magellanica* has a wide distribution around the Magellanic region in southern South America, from Puerto Montt in the Pacific Ocean (42°S) to the Buenos Aires province in the Atlantic Ocean (35°S), including the Strait of Magellan, Cape Horn, Tierra del Fuego and the Malvinas Islands (Powell 1973, Pastorino 1995, González-Wevar et al. 2012). The genus *Nacella* is a broadcast spawner (it releases its gametes into the water) (Moriconi 1999, González-Wevar et al. 2012) and, like many other marine gastropods, its reproduction is correlated with the photoperiod and the water temperature (Moriconi 1999, Bigatti et al. 2008, Cumplido et al. 2010).

In recent years, *N. magellanica* has been extended as an incipient resource to the internal markets in Argentina, without any form of official control. This study is accordingly aimed at adding to a series of efforts made to establish regulatory policies for marine gastropod fisheries in northern Patagonia (Bigatti and Ciocco 2008, Penchasazdeh et al. 2009, Bigatti et al. 2015, among others).

The study reported here investigated for the first time the reproductive cycle and seasonal variation in total body mass and biochemical composition of *N. magellanica* in order to determine its nutritional values. The results obtained for this coastal resource can be useful for strengthening the management policies of several marine gastropod species.

MATERIALS AND METHODS

Sampling

From January to December 2012, 50 specimens of *Nacella magellanica* were collected each month on the rocky intertidal shore of Punta Ninfas, Chubut (42°58’42”S, 64°18’33”W). The analysis were made only with limpets above the maturity size [1.73 cm high in females and 1.5 cm in males, *sensu* Nieto Vilela (2014)]. Unfortunately, no individuals could be collected during the month of May because the extreme weather conditions made it impossible to access to the sampling site. Punta Ninfas is located in the open sea, about 100 km from the city of Puerto Madryn, an area lacking fishing activity and inhabited by a stable community of marine elephants (*Mirounga leonina*) and particularly frequented by outdoor tourists.

All the specimens collected were taken directly to the laboratory and 30 were fixed in Bouin’s fluid for 48 h and stored in 70% (v/v) aqueous ethanol for histology. Because physical procedures enhance spawning in other molluscs (Velasco et al. 2007, Aji 2011), no sand depuration was performed and the biofilm over the shell was not removed. For each sampling, morphometric measurements were recorded on 30 individuals with callipers (precision 0.1 mm) to establish the height, length and width. To determine the total weight, each shell was removed and the shell and body weights were recorded on a digital balance (precision ±0.001 g). Sex was determined on the basis of gonadal *frotis* under an optical microscope and related to gonadal colour: females were green and males yellow (see Results). From the total of sampled individuals, 5 specimens were used for nutritional values and 5 were used for ash determination. The 10 remaining specimens were used as a back-up, because in the process of shell extraction the tissue was often broken, making the specimens useless for nutritional composition determination or histological gonadal analysis.

Gonadal cycle

Each month, only around 10 individuals were suitable for gonadal determination (Table 1). Gonads were processed following standard histological procedures to study the annual cycle, as described by several

| Table 1. – Number of individuals per gametogenic cycle of the 30 analysed per month. |
|-----------------------------------------|--------|----------|--------|
|                                       | Analyzable | Parasited | Sand/Indet |
| January                                | 10      | 4        | 16      |
| February                               | 10      | 3        | 17      |
| March                                  | 10      | 1        | 19      |
| April                                  | 10      | 2        | 18      |
| June                                   | 12      | 0        | 18      |
| July                                   | 12      | 0        | 18      |
| August                                 | 11      | 1        | 18      |
| September                              | 10      | 0        | 20      |
| October                                | 11      | 2        | 17      |
| November                               | 10      | 2        | 18      |
| December                               | 12      | 2        | 16      |
Between 10 and 100 oocytes with a visible nucleolus using the ZEN 2.3.0.13 program. Measurements (length and width mean) in all cells with the gonadal cycle was estimated by oocyte diameter digital camera (Sound Vision 2.0). For each female, (Zeiss Axiostar, Germany) and photographed with a eosin (Gabe 1968).

Mazurkiewicz and Pokryszko 2005). The gonad was sectioned in anteroposterior slices (toward both front and back), embedded in paraffin and sectioned at 5 to 7 µm with a digital microtome (Leica), then heated for 24 h at 60°C, and finally stained with haematoxylin and minium film and heated at 60°C to constant weight. Dried tissues were triturated before sample processing to obtain a composite sample and an average monthly value for protein, lipid and carbohydrate content.

These determinations were performed by colorimetry according to the respective protocols of Lowry et al. (1951) using albumin as standard, Zöllner and Kirsch (1962) using cholesterol as standard and Fraga (1956) using glucose as standard. The measurements were made in a Hewlett Packard Model 8452A spectrophotometer, and the results were expressed as percent dry weight of tissues.

For ash determination, a single aliquot of wet body mass of five individuals was first heated at 60°C until the dried weight (DW) became constant. Thereafter, the sample was calcinated in a muffle oven at 550°C for 12 h and the weight of the resulting ash (CW) was recorded. The percent ash content was calculated as (CW/DW)×100. The estimation of moisture content was performed with the same aliquot of the monthly wet sample by recording the initial weight (IW) and the value obtained after heating at 80°C to constant weight (CW). The percent moisture content was then expressed as ((IW-CW)/IW)×100.

Statistical analyses

Because the parametric assumption was not met, differences between weight and length in males and females were evaluated with Wilcoxon and Mann-Whitney tests. Differences between average oocyte size and seasonal biochemical comparisons were assessed with a Kruskal-Wallis test. For significant results (P<0.05) a pairwise comparison test of subgroups was applied. The sex ratio was determined and compared with a 1:1 proportion by a chi-squared test. All the testes and modal determination were performed with the Statistica 7.0 software package (StatSoft, Inc., Tulsa).

To infer whether the sea surface temperature (SST) in °C and the gametogenic cycle were associated, we made in situ measurements in each sampling event and estimated the data from standard mapped satellite images (SMIs, NOAA, http://oceancolor.gsfc.nasa.gov/) using those of seasonal satellite-derived SSTs from MODIS Aqua for the period January to December 2012. The images have a spatial resolution of approximately 9.2 km. The global SMI data were subsampled from the region bounded by latitude from 33° to 43°S and longitude from 49° to 65°W. The satellite data were extracted from the sea 2 km off the coast. To evaluate seasonal variation in oocyte diameter, we performed a Kruskal-Wallis test and a pairwise comparison test of ranks.

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### Table 2. – Stages of gametogenic cycle in males and females.

<table>
<thead>
<tr>
<th>Stages</th>
<th>Female</th>
<th>Male</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Developing</td>
<td>Gonads formed by tubules supported by a germinal epithelium lying alongside (Fig. 1). The oocytes proliferate inside the tubule attached to the wall. The basophilic oocytes (40-80 µm) were usually of pyriform shape. The oocytes also had a conspicuous external membrane and were found next to companion cells.</td>
<td>The tubules were associated with multiple layers of spermatogonia (Fig. 1). In this stage, the lumen of the tubules contained some empty space along with some spermatocytes and spermatids.</td>
</tr>
<tr>
<td>2. Ripe</td>
<td>Increase in oocyte diameter. Short distance between the mature acidophilic oocytes, producing right edges in the mature oocytes (100-140 µm, Figs 1 and 2).</td>
<td>A minimum of empty space was observed in the tubules, while spermatocytes (SC) and spermatooza (S) were present (Fig. 1).</td>
</tr>
<tr>
<td>3. Spawned</td>
<td>Empty tubule lumen. Mature oocytes attached to the lumen were very uncommon, while the basophilic oocytes were more abundant (Figs 1 and 3). The lumen contained some athresic oocytes, and the trabeculae became more enlarged.</td>
<td>At this stage, the spermatogonia were formed by two or three layers, whereas the spermatooza became concentrated in the centre of the tubule (Fig. 1). Epididymis full of mature spermatocytes.</td>
</tr>
<tr>
<td>4. Resorption</td>
<td>The oocytes exhibited an irregular shape and size, and the follicular walls became more conspicuous (Figs 1 and 4). Presence of phagocytic-nutritive cells.</td>
<td>The spermatooza began to withdraw from the tubules and became separated from each other while phagocytes were present in the centre of the tubules (Fig. 1).</td>
</tr>
</tbody>
</table>

Authors (Ramón and Amor 2002, Horn et al. 2005, Mazurkiewicz and Pokryszko 2005). The gonad was sectioned in anteroposterior slices (toward both front and back), embedded in paraffin and sectioned at 5 to 7 µm with a digital microtome (Leica), then heated for 24 h at 60°C, and finally stained with haematoxylin and eosin (Gabe 1968).

Each sample was observed by light microscopy (Zeiss Axiostar, Germany) and photographed with a digital camera (Sound Vision 2.0). For each female, the gonadal cycle was estimated by oocyte diameter measurements (length and width mean) in all cells with a visible nucleolus using the ZEN 2.3.0.13 program. Between 10 and 100 oocytes with a visible nucleolus were measured in every female. The gametogenic cycle of both sexes was divided into four stages (Table 2) according to the stage of the reproductive cells; the classification was based on previous studies on Patagonian gastropods (Morriconi 1999, Bigatti et al. 2008, Averbuj et al. 2010) (Fig. 1).

### Nutritional composition

For nutritional value determination, five randomly selected individuals with a size range of 0.9 to 2.7 cm height were analysed monthly. The determination was performed in males and females together, because both sexes are collected for consumption at the same time due to the absence of external sexual dimorphism. Total body masses were placed on aluminium film and heated at 60°C to constant weight. Dried tissues were triturated before sample processing to obtain a composite sample and an average monthly value for protein, lipid and carbohydrate content.

These determinations were performed by colorimetry according to the respective protocols of Lowry et al. (1951) using albumin as standard, Zöllner and Kirsch (1962) using cholesterol as standard and Fraga (1956) using glucose as standard. The measurements were made in a Hewlett Packard Model 8452A spectrophotometer, and the results were expressed as percent dry weight of tissues.

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Fig. 1. – Gonadal stages for females (left) and males (right), 1, Developing; 2, Ripe; 3, Spawning; 4, Resorption. Abbreviations: oocytes (Oo), companion cell (CC), nucleus (N), follicular wall (FW), spermatocytes (SC), spermatozoa (S), phagocytes (Ph). Scale bar females: 100 µm.
Scale bar males 1-4: 10 µm; 2-3: 100 µm.
RESULTS

Of the 358 Nacella magellanica sampled, 176 were females, 170 were males and 12 were indeterminate. The sex ratio in the N. magellanica population from Punta Ninfas was not significantly different from parity (1:1, n=346, d.f.=1, P>0.05) throughout the study period. The female mean total weight (12.05±3.75 g) and body weight (6.23±2.0 g) were significantly higher (P<0.05) than those of males (10.22±3.34 g and 5.23±1.8 g, respectively). The average female height (1.98±0.31 cm) and width (3.82±0.37 cm) were significantly higher (P<0.05), than those of males (1.89±0.28 cm and 3.68±0.34 cm, respectively). The average length was similar in both sexes: 3.07±0.32 cm in females and 3.01±0.29 cm in males (P>0.05).

Gonadal cycle

We found a gonad coloration pattern across the reproductive cycle; in the case of females, green gonads recorded during maturation become darker in totally spawned specimens. In males, the gonad was yellow upon maturing and in ripe (fully mature) specimens, but brown in individuals after gamete release. Of 30 fixed specimens, only a mean of 10 were suitable for gonadal determination, because of rupture of several slides caused by sand content in the gut or because they were full of parasites Gymnophalloides (Table 1). Castration due to digenean parasites (Gymnophalloides), a phenomenon well described previously for this species (Cremonte et al. 2013, Bagnato et al. 2015), was detected in 16% of the analysed individuals (Table 3).

The monthly value of the oocyte modal diameter, from January to December 2012, ranged from 12 µm in January and February to 145 µm in October. We observed two modal diameters in several months due to the presence of growing and ready-to-spawn oocytes (Table 3). The low oocyte modal diameter observed in January, February and March (12-16 µm) may correspond to a phase of recovering after spawning, with resorption and gamete developing, and the gonad shows a developing stage in March, with oocytes with a modal diameter of 129 µm. This stage continued until September, with an increase in the oocyte size until spawning; in October a second developing period started and continued with the increase and spawning (Table 3, Fig. 1). Oocytes with mode diameters bigger than 140 µm seemed to be mature and ready to spawn when we recorded them free in the gonadal lumen.

We observed that female gamete liberation is low in January, March and April and increases in August, continuing until the end of the year (Fig. 2). The male gametogenic cycle was different to the female one (Fig. 3). The first individuals in the ripe stage appeared

<table>
<thead>
<tr>
<th>Protein</th>
<th>Lipid</th>
<th>Carbohydrate</th>
<th>Ashes</th>
<th>Humidity</th>
<th>Oocyte mean</th>
<th>Oocyte 1st mode</th>
<th>Oocyte 2nd mode</th>
<th>Prevalence</th>
</tr>
</thead>
<tbody>
<tr>
<td>January</td>
<td>19.9±3.8</td>
<td>3.3±0.3</td>
<td>2.4±0.4</td>
<td>14.42</td>
<td>78.03</td>
<td>14</td>
<td>12</td>
<td>13</td>
</tr>
<tr>
<td>February</td>
<td>22.8±4.1</td>
<td>2.9±0.3</td>
<td>0.4±0.3</td>
<td>41.15</td>
<td>80.87</td>
<td>12</td>
<td>11</td>
<td>10</td>
</tr>
<tr>
<td>March</td>
<td>25.8±4.9</td>
<td>2.7±0.08</td>
<td>1.31±0.1</td>
<td>21.60</td>
<td>84.12</td>
<td>58</td>
<td>16</td>
<td>129</td>
</tr>
<tr>
<td>April</td>
<td>63.4±12.2</td>
<td>3.6±0.2</td>
<td>1.1±0.07</td>
<td>20.68</td>
<td>84.27</td>
<td>213</td>
<td>129</td>
<td></td>
</tr>
<tr>
<td>May</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
</tr>
<tr>
<td>June</td>
<td>59.2±18.4</td>
<td>2.4±0.2</td>
<td>0.9±0.2</td>
<td>30.48</td>
<td>82.95</td>
<td>125</td>
<td>129</td>
<td></td>
</tr>
<tr>
<td>July</td>
<td>10.3±4.5</td>
<td>2.2±0.1</td>
<td>0.4±0.2</td>
<td>29.70</td>
<td>80.55</td>
<td>113</td>
<td>123</td>
<td></td>
</tr>
<tr>
<td>August</td>
<td>27.3±2.1</td>
<td>2.1±0.09</td>
<td>1.2±0.3</td>
<td>27.46</td>
<td>82.89</td>
<td>122</td>
<td>129</td>
<td></td>
</tr>
<tr>
<td>September</td>
<td>33.2±3.6</td>
<td>2.1±0.3</td>
<td>1.4±0.6</td>
<td>24.85</td>
<td>80.10</td>
<td>118</td>
<td>111</td>
<td></td>
</tr>
<tr>
<td>October</td>
<td>18.7±12</td>
<td>2.6±0.05</td>
<td>5.4±0.8</td>
<td>18.45</td>
<td>81.23</td>
<td>124</td>
<td>145</td>
<td></td>
</tr>
<tr>
<td>November</td>
<td>16.9±7.2</td>
<td>2.8±0.2</td>
<td>2.5±0.2</td>
<td>25.91</td>
<td>82.44</td>
<td>121</td>
<td>116</td>
<td></td>
</tr>
<tr>
<td>December</td>
<td>30.3±0.3</td>
<td>2.9±0.4</td>
<td>2.7±0.9</td>
<td>20.49</td>
<td>82.44</td>
<td>48</td>
<td>24</td>
<td>113</td>
</tr>
</tbody>
</table>

Fig. 2. – Female gonadal stages as determined from histological analysis. Key to bar textures: black, developing; grey, ripe; crosshatched, spawned; diagonally hatched, resorption.

Fig. 3. – Male gonadal stages as determined from histological analysis. Key to bar textures: Black, developing; grey, ripe; crosshatched, spawned; diagonally hatched, resorption.
from April to December, with a maximum in November. Gamete release in males occurred throughout most of the year, except in June. Thereafter, resorption was found throughout the year except in June and October.

Although histological sections of *N. magellanica* showed that individuals were in more than one gonadal stage (Figs 2 and 3) at the same time, we were able to determine a relationship between the oocyte sizes and the seasons. With a rank comparison we found three groups (selected by oocyte size) that were significantly different (P<0.05): A, summer oocytes with a low size; B, autumn oocytes with a medium size; and C, spring and winter oocytes grouped together.

Sea temperature values recorded in situ were similar to those recorded by satellite (Fig. 4). Throughout the sampling period the measurements made by the MODIS Aqua satellite recorded an average water surface temperature of 12.8°C (max. 17.13°C, min. 8.72°C) and a mean photoperiod of 13.0 h (max. 16.3 h, min. 10.06 h). The increase in photoperiod started after 21 June and reached a maximum of 16.3 h of light in December, while the SST began to increase in August and reached its maximum value of 17°C in January (satellite information, Fig. 4). The spawning started in August and continued until December, with the concomitant increase in the water temperature (Fig. 4). During the autumn and winter (March to August), the oocytes became mature, with diameters greater than 140 µm before gamete release in spring (September, October and November). Between the seasons, the average size of the oocytes was significantly higher in winter (120.72 µm) and spring (122.54 µm) (P<0.05). The smallest mean value was recorded during the summer (32.4 µm).

Finally, the female spawning period for *N. magellanica* was associated with a decrease in body weight (Fig. 5). During the month of July, when the gonads were mature and ready to spawn, the body weight underwent a slight increase, and decreased later during the spring spawning peak.

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**Nutritional composition**

Results of biochemical analysis are shown in Table 3. The average nutritional values (dry weight) were 29.8±3.25% proteins, 2.71±0.12% lipids, and 1.8±0.26% carbohydrates. The average ash content was 25.0±2.2% and the average moisture 81.8±0.6%. The abnormally high percentages of ashes are related to the abundant content of inorganic sediment particles within the organism’s digestive system. Average seasonal concentrations showed no significant variations for proteins, but for lipids the winter average was significantly lower than that of the other seasons (P<0.05), while for carbohydrates the winter and spring averages were significantly lower and higher, respectively, than those of the other seasons (P<0.05).

We found a body weight variation in males and females across the sample period, though female body weight remained higher than male body weight during most of the year. Proteins were high in April and June, lipids remained low throughout the year with a slight decrease during the spawning period (Table 3), and carbohydrates increased during the spawning period with a maximum in October (Fig. 5).

**DISCUSSION**

The nutritional values of *Nacella magellanica* are within the ranges of those of widely consumed species of molluscs. The spawning period is related to changes in environmental conditions, such as the photoperiod and temperature of southern latitudes in the southwestern Atlantic.

The spawning period recorded in *N. magellanica* differed from that of previous studies on the southern species *Nacella deaurata* from the Beagle Channel.
The high concentration of protein found in April and June has no clear correlation with the abundance of available food to be expected in spring and summer (Giese 1967, Morais et al. 2003, Najmudeen 2007), so it could be attributed to reserves after the ingestion of biofilms prior to this month that are energetically favourable for the gamete-developing period. High protein proportions were also found in the Antarctic Nacella concinna (Congjie et al. 2005), which would indicate that the genus Nacella can store high protein concentrations naturally. The highest concentration of lipids in April is probably related to the food intake during the summer and the occurrence of gametogenesis at that time.

In general, marine invertebrates with external fecundity require reserves of carbohydrates to survive within the environment (Spikes 1949, Anderson and Personne 1970). Because carbohydrates are the primary energy source of molluscs (Barber and Blake 1981), the decline observed in the percentages of total body carbohydrates in N. magellanica in July could be attributable to a drop in glycogen levels as a result of the usual shortage of food in the winter (Morais et al. 2003, Ren et al. 2003). Furthermore, since carbohydrates are transformed into lipids during gamete formation (Ren et al. 2003), a decrease in the percentage of carbohydrates during gametogenesis might also be expected.

The annual nutritional values and the reproductive cycle reported here can enhance and expand current efforts to protect and conscientiously exploit this commonly consumed resource. According to our results, the optimum values for human consumption (i.e. the highest concentration of proteins, the highest body weight and the gonadal stage) are in April, corresponding to the austral autumn. The capture of the species in this season is recommended only if the minimum size of capture, 2 cm, is respected, as stated in fishery regulation 199/18 (Secretaría de Pesca del Chubut) based on preliminary studies of Nieto Vilela (2014), allowing at least two spawning periods to ensure the sustainability of the resource.

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