Population dynamics of *Megalobulimus paranaguensis* (Gastropoda: Pulmonata) in the southeast coast of Brazil

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ABSTRACT. The population dynamics of *Megalobulimus paranaguensis* (Pilsbry & Ihering, 1900), a large and long-lived land gastropod from Brazil’s Southeast (Atlantic Forest) was studied between 2006 and 2009, at an urban park in the city of Santos, state of São Paulo. The study included biometry, weighing, and marking and recapture of adult individuals. The variables obtained from specimens were correlated with the environmental variables of the study period. The survival rate of the adult snail population was 96.7%. Recruitment showed several peaks during the year, and was concentrated between April and August, except in 2009, when there was only one peak in August. Specimen abundance progressed from 2006 on, with a trend towards relative stability during that period. The condition factor remained relatively stable as well, decreasing from December to February. The potential evapotranspiration and precipitation influenced the number of captures, and the mean temperature influenced the condition factor. *Megalobulimus paranaguensis* goes through a period of dormancy in the tropical summer, which is between November and February, but remains active in the winter. The survival rate of *M. paranaguensis* was high, and may be the result of having its annual cycle synchronized with the tropical climate of the Atlantic forest.

KEY WORDS. Activity; abundance; dormancy; land snail.

*Megalobulimus* Miller, 1878, which includes species popularly known as aruá-do-mato, is a group of Neotropical giant snails of Megalobuliminae. Not much is known about the biology, behavior and ecology of species of *Megalobulimus* and other Brazilian terrestrial gastropods. *Megalobulimus* spp. populations tend to occur in low densities (Eston et al. 2006) and, like other native species of terrestrial mollusks, they have low reproductive potential, spawning only two to five eggs per batch (Sobreira & Molina 2002). They are herbivorous, go through a dormancy period, and bury themselves in the rainforest soil or leaf litter during the day (Bequaert 1948). Individuals of *Megalobulimus* spp. also have a long lifespan (Fontenelle & Miranda 2012). Since species of this genus are not commonly found, their population dynamics remains poorly understood. Morphologically, species of *Megalobulimus* are distinguished by having large thick and round shells with vertical nepionic riblets, thickened outer lip in adults (Bequaert 1948) and buccal fringe (Leme 1973).

*Megalobulimus paranaguensis* (Pilsbry & Ihering, 1900), the subject of this study, has a mean shell length of 98 mm. This species is distributed in the North of the state of Santa Catarina, Paraná and in the south of the state of São Paulo (Morretes 1954, Simão 2006, Agudo-Padron 2014). Currently, the northernmost site from its original distribution is the Island of São Vicente. Little is known about the possible reasons that led to this range expansion. Recently, Fontenelle & Miranda (2012) analyzed the age and growth of *M. paranaguensis* and Miranda et al. (2015) analyzed the population structure of the species in the presence of the invasive *Achatina fulica* Bowdich, 1822, and pointed out that long-term studies need to be conducted to better understand the population cycles of the species. The aim of this study was to analyze the population dynamics of *M. paranaguensis* from Santos during four years (2006-2009).

MATERIAL AND METHODS

The study took place from 2006 to 2009, in the Parque Zoobotânico Orquidário Municipal de Santos (PZOMS) (UTM: 23K E 362672.57, 7348904.53 N), an urban park located in the municipality of Santos, Island of São Vicente, state of São Paulo. The park is surrounded by buildings. Specimens of *M. paranaguensis* were collected within the boundaries of the PZOMS, Monday through Friday, from 7:00 am to 8:00 am, totaling 800 collecting hours in four years. The specimens were marked using Fontenelle & Miranda (2012) code tags. The total length and weight of specimens were recorded and then specimens were released. Only adult individuals identified by thicker
shell were marked. Voucher specimens of the population studied were deposited in mollusk collection of Museu de Zoologia da Universidade de São Paulo (lot MZSP 54629). The following environmental variables were obtained from the website http://www.ciagro.sp.gov.br/ciagroonline/#Monitoramento (maintained by EMBRAPA – Agronomic Institute of Campinas) for the municipality of Santos, SP: mean temperature, potential evapotranspiration, number of rainy days, precipitation and temperature range.

Captures and recaptures took place between March and October, since M. paranaguensis individuals aestivate from November to March (Miranda et al. 2015). They were quantified to estimate the population parameters according to the model of Schwarz & Arnason (1996). The fundamental parameters of this model are apparent survivorship (\( \Phi \)), probability that an individual will be captured (\( p \)) and probability of an individual entering the population (\( \beta \)). The model also has derived parameters, which are recruitment (\( B \)) and the estimated abundance (Population size) in each sample (\( N_i \)). To assess any violations of the assumptions of mark-recapture analysis, the \( \chi^2 \) test was used.

The basic parameters are either constant or they can vary over time, and combinations thereof generate different models. To find out which model best explained the data, the Akaike information criterion with correction for small samples (AICc) (Burnham & Anderson 2001) was used. Models with smaller AICc values have greater statistical support. Furthermore, to determine which model is most supported by the data the AIC Delta (\( \Delta \) AIC) was used, and Akaike weight to ascertain if the probability of the model in question is the most parsimonious (Burnham & Anderson 2001, 2004, Johnson & Omland 2004).

The relationship between length and weight was determined by a regression analysis, adjusting the power function using shell length as an independent variable. The resulting allometric coefficient was tested using the t-test to determine whether the growth was isometric (\( b = 3.00 \)), positive allometric (\( b > 3.00 \)) or negative allometric (\( b < 3.00 \)). Then, the condition factor determined the adult individuals. Stepwise regressions using AICc as variables selection criterion were conducted between environmental variables and the number of captures and the condition factor.

**RESULTS**

During this study there were two peaks of increased snail activity, which were represented by an increase in the number of captures. A major peak occurred from March to May, and a minor one from August to September. The number of monthly captures (Fig. 1) was lower than from December to February, when the species is dormant, and there was a decline in the number of captures during June and July. The average length of the shell of M. paranaguensis was 96.60 ± 4.83 (mean ± SD) mm (minimum shell length = 70.00 mm and maximum shell length = 105.00). The mean weight of specimens was 159.21 ± 35.89 g (minimum = 40g and maximum = 226 g). The resulting biometric ratio (weight = 0.0007.\( \text{Length}^{2.687} \); \( r^2 = 0.8467 \)) showed negative allometry (\( t = 4.97, \text{DF} = 331, p <0.0001 \)). The condition factor remained relatively stable as well, with a decrease from December to February (Fig. 2).

A total of 82 adult individuals were captured. Of these, 56 were recaptured at least once (recapture rate = 0.68). There were no violations of the assumptions of the mark-recapture method (\( \chi^2 = 47.33, p = 0.78, \text{DF} = 56 \)). The model that best explained the data was the \( \Phi(t)p(t)\beta(t) \), which assumes constant survival and capture probabilities and variables entry over time (Table 1). The apparent survivorship of the population was 96.7% (CI: 94.0-97.9%) during the four years of analysis. The capture probability showed two peaks per year: a higher peak from April to May and a shorter peak from August to September (Fig. 3), probably due to the variation of temperature and potential evapotranspiration in the area. Recruitment showed several peaks during the year, concentrated between April and August, except for the months of November and December when the species aestivates.
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Potential evapotranspiration and mean temperature. Potential evapotranspiration and mean temperature fluctuated significantly, while fluctuations in precipitation were more irregular, generally declining in February and peaking in July. The stepwise regression kept the variables potential evapotranspiration and precipitation. The relationship between the abundance of *M. paranaguensis* and potential evapotranspiration was negative ($t = 3.99$, $p < 0.0001$, $b = -0.02$), whereas abundance and precipitation were positively correlated ($t = 1.72$, $p = 0.08$, $b = 0.004$). For the condition factor, the relationship between mean temperature and abundance was negative ($t = 2.06$, $p = 0.04$, $b = 2.06 \times 10^{-5}$).

**DISCUSSION**

*Megalobulimus paranaguensis* has a well-defined seasonal pattern, with aestivation in the warmer months, followed by increased post-dormancy activity. This pattern is influenced by evapotranspiration, precipitation and environmental temperature. Our data show that the population of *M. paranaguensis* is highly resistant and has a high survival rate. The life cycle of the species is synchronized with the tropical environment, and individuals cope with heat stress by becoming numb during the hot summer months.

The extremely high survival rate of *M. paranaguensis* coupled with the recruitment peaks has resulted in an increase in the numbers of individuals over the years. Miranda et al. (2015) estimated the survival rate of *M. paranaguensis* at 73% in Porchat Island, São Vicente. Three factors seem to have played a role in the higher survival rate found in our study: 1) there are no predators to control the population in Parque Zoobotânico Orquidário Municipal de Santos; 2) the Orquidário has a deeper layer of plant litter, is more densely covered with vegetation and is less impacted than the Porchat Island. The latter has several areas with lower plant density and household waste in its interior; and 3) young individuals were not marked in this study. In general, the newer cohort studies have the highest mortality rate (Hellier 2001) and the exclusion of juveniles tends to overestimate the calculated survival. This has been confirmed by Romero (2004), who documented a mortality rate higher than 94% in some years in a cohort of *Megalobulimus mogianensis* Simone & Leme, 1998. In his study, most individuals in captivity died before their first dormancy.

The capture probability estimated by the model followed the same tendency as the mean number of catches throughout the year. The capture probability of terrestrial snails is associated with activity (Cook 2001), which in turn is influenced by moisture, precipitation and evapotranspiration in the environment.

Accroding to the number of recruitment peaks, *M. paranaguensis* is an iteroparous species, since there is more than one posture peak during the year. This same pattern has been found in *Megalobulimus abbreviatus* (Bequaert, 1948) (Horn et al. 2005). Iteroparity (more than one breeding season) is relatively common among pulmonate mollusks (Calow 1978, 1979).
Table 1. Results of mark – recapture model selection of *Megalobulimus paranaguensis* in the Parque Zoobotânico Orquidário Municipal de Santos (PZOMS). The values of the most parsimonious model are in bold.

| Model | Description | $\text{AIC}_c$ | $\Delta\text{AIC}_c$ | $\text{AIC}_c$ weight | Parameters |
|-------|-------------|----------------|----------------------|------------------------|------------|
| $\Phi(.|p(t)|\beta(t)$ | Constant survivorship; variable Entry and Capture probabilities | 1,140.120 | 0.000 | 1.00 | 61 |
| $\Phi(.|p(.)|\beta(t)$ | Constant Survivorship and Capture probability; variable Entry probability | 1,196.200 | 56.070 | 0.00 | 32 |
| $\Phi(t|p(t)|\beta(t)$ | Saturated model | 1,246.690 | 106.560 | 0.00 | 90 |
| $\Phi(t|p(.|\beta(t)$ | Variable Survivorship and Entry probability; constant Capture probability | 1,263.120 | 122.990 | 0.00 | 61 |
| $\Phi(t|p(t)|\beta(.)$ | Variable survivorship and Capture probability; constant Entry probability | 14,844.500 | 13,704.370 | 0.00 | 62 |
| $\Phi(t|p(.|\beta(.)$ | Variable survivorship and Capture probability; constant Entry probability | 38,162.693 | 32,024.739 | 0.00 | 33 |
| $\Phi(.|p(.|\beta(.)$ | Constant Survivorship and Entry probability; constant Capture probability | 39,012.140 | 37,872.010 | 0.00 | 32 |
| $\Phi(.|p(.)|\beta(.)$ | Reduced model | 39,417.610 | 38,277.490 | 0.00 | 4 |

Figure 6-8. Mean values of temperature in °C (6), potential evapotranspiration in% (7) and precipitation in mm (8) during the year in the municipality of Santos.
Lazaridou-Dimitriadou (1995), and seems to be the reproductive pattern in Megalobuliminae. Recruitment also occurs in the same period as found by Miranda et al. (2015), but in their study only one recruitment peak during a period of six months was found.

Aestivation in this study was found from November to February, consistent with Miranda et al. (2015). Aestivation is a state of dormancy where there is a lot of heat or dryness (Booth 1974), is relatively common among terrestrial and freshwater mollusks (Cáceres 1997). According to Iglesias et al. (1996), precipitation controls the beginning and end of dormancy in *Helix aspersa* Muller, 1774, whereas periods of rainless days trigger dormancy and the return of the rain marks the end of it. Lazaridou-Dimitriadou & Saunders (1986) suggested that the onset of dormancy in *H. lucorum* Linnaeus, 1758 is controlled by low humidity. Another factor that may also be involved in the activity cycles of *M. paranaguensis* is the presence of a possible pattern of endogenous activity. This pattern was also speculated by Lazaridou-Dimitriadou & Saunders (1986). Therefore, aestivation in *M. paranaguensis* is caused by temperature rise and evapotranspiration between December and March, which remove moisture from the environment and cause a slowdown in snail activity. This was reflected in the condition factor of *M. paranaguensis*, which dropped from December to February. This drop reflects dehydration and weight loss during aestivation. Conversely, the progressive increase of this index during the months of activity show that the snails accumulated energy reserves to prepare for the state of dormancy.

The effects of environmental variables on the abundance of terrestrial mollusks are well documented in the literature. In the study of Choi et al. (2004), mathematical modeling using derived ordinary equations found that mortality in young *Deroceras reticulatum* (Muller, 1774) is more influenced by temperature, while recruitment rates in young individuals and the mortality rate of adults are more influenced by precipitation. Recent work by Nunes & Santos (2012) showed that the variables moisture, air temperature, soil temperature and depth of plant litter influence the distribution and abundance of terrestrial gastropods in the Atlantic Forest. Snails depend on high humidity to have an active life (Baker 1958), being sensitive to desiccation, which tends to restrict their movements (Getz 1974). D’Avala & Bessa (2005) and Dias et al. (2007) showed that substrate moisture is important for growth, reproduction, survival and resistance to desiccation in *Subulina octona* (Bruguère, 1792). As the evaporation removes moisture from the substrate, it decreases the number of captures, while the high precipitation tends to increase the number of catches by increasing humidity, which influences positively the activity of *M. paranaguensis*. Furthermore, according to Rizzatti & Romero (2001), body hydration increases heart rate in *M. mogianensis*, and since heart rate indicates activity, precipitation is also an important factor modulating activity in *M. paranaguensis*. Miranda et al. (2015), in a nearby location of this study, showed that the abundance of *M. paranaguensis* increases with increasing environmental humidity, and that humidity decreased in the period when evapotranspiration and temperature increased in this study.

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