Assessment of dispersal and population structure of Norway rats (*Rattus norvegicus*) in a seaport setting

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Abstract
Seaports are introduction hotspots for invasive alien species (IAS). This is especially true for rodents, which have accompanied humans around the globe since the earliest days of ocean-going voyages. The rapid spread of IAS soon after arrival in a new environment is facilitated by further human-mediated transport or landscape features, like roads. By measuring genetic diversity and structure to investigate dispersal pathways, we gained insight into the transport, spread and establishment stages of a biological invasion, leveraging the most common rodent species (*R. norvegicus*) in this setting. We characterized the genetic structure of three Norway rat populations along a busy industrial road used by trucks to access the Port area in Paranaguá city (Brazil). A total of 71 rats were genotyped using 11 microsatellite markers. The results revealed a pattern of gene flow contrary to the expected stepping-stone model along the linear transect, with the two furthest apart populations being clustered together. We hypothesize that the observed outcome is explained by natural dispersal along the corridor being lower than human-mediated transport. The sampled area furthest from the port is a gas station frequented by trucks which are considered the most likely mode of transportation. In terms of management strategies, we suggest more emphasis should be put on cargo surveillance to lower the risk of Norway rat dispersal, not only for biosecurity, but also for sanitary reasons, as this port is a major grain trading point.

Keywords Human-mediated dispersal · Microsatellite · Road · Invasive · Urban areas · Zoonotic disease

Introduction

Biological invasions are a serious threat to biodiversity around the world, causing not only disturbances to ecosystems, but also threats to human health and negative impacts on agriculture (Myers et al. 2000; Miller et al. 2016). By definition, invasions occur when species are intentionally or accidentally introduced outside of their native or historic range, and successfully spread in the new environment (Levine 2008).

Rodents, especially from the genus *Rattus*, are commensal species considered to be the most widely introduced vertebrates, accompanying humans around the globe (Drake and Hunt 2009). Their occurrence often has a dramatic impact on native biodiversity, affecting individual demographic rates, genetic (through hybridization), trophic structure and/or resources availability (Parker et al. 1999), and they are causing massive economic loss because of destruction of grains and stocked materials (Pimentel et al. 2000). Rodents are also known for their role in facilitation of the transmission of numerous diseases to humans (Himsworth et al. 2013). These animals are particularly problematic because of their high reproductive rate, the capability of exploiting a wide range of food sources, survival in vastly different ecological conditions and ability to travel long distances unnoticed, utilising existing transportation infrastructure and shipping routes (Varnham 2010).
Seaports are considered introduction epicentres for invasive alien species (IAS) (Drake and Lodge 2004; Miller et al. 2016). It is therefore crucial that ports implement surveillance measures, such as containment facilities and the use of biocontrol agents, chemicals (toxins) or mechanical devices (traps), to control, and prevent the establishment and spread of alien species in a new environment (Lodge et al. 2006; Gren 2008).

Paranaguá is a seaport city located in Paraná state and is home to the main port of Southern Brazil. The port of Paranaguá is the second largest port in export of grains, fruits, and seeds in Brazil handling 13% of the national cargo and it is the third-largest port in cereal shipment (i.e. considering the volume imported and exported) (11.3% of the national cargo) (ANTAQ 2020). Because of the high volumes of food being handled and the resulting abundance of food sources for pest species, Paranaguá has long dealt with pest infestations. The port has implemented a vector proliferation control program that aims to reduce and control pigeon, rat, and mice populations by using poison bait and bird repellent. In addition, to reduce the food availability for pest species, cleaning procedures are in place to remove spilled grains in the primary area of loading/unloading and surroundings (APPA 2016). There is no information about how effective these measures are in terms of avoiding spreading of these pests to the city.

Previous studies made in urban environment show that Rattus norvegicus present strong site fidelity and individual movements are usually limited to 30-150 m. Also, dispersal from their natal site seems to occur over short distances (Gardner-Santana et al. 2009; Byers et al. 2019). If that applies to the rat population in Paranaguá, the control measures implemented in the Port should be enough to avoid spread of new immigrants that arrive in the port. If an IAS manages to pass through the implemented control measures after its arrival in the new environment, linear features in the landscape, like roads, railways, and canals can facilitate its spread and colonization (Brown et al. 2006; Cameron and Bayne 2009). In Paranaguá a sampling collection through the city revealed a positive association of Norway rats (Rattus norvegicus) with the main roads used by trucks to access the Port area (Gatto-Almeida et al. 2020a). In this study we aimed to evaluate how effective are the control measures implemented by Paranaguá’s port to prevent rats spread from the port. For this purpose, we genetically characterize three R. norvegicus putative populations along the main road used by trucks in Paranaguá i.e. a linear feature that may be facilitating their spread, to understand possible migrations and gene flow that might be occuring. This road crosses the entire city on its longest axis connecting its main entrance to the port. If the control measures were effective we expected individuals from the two areas outside the port would be genetically similar between them, and port’s population would present higher genetic differentiation.

Material and methods

Study area

The study was conducted in Paranaguá municipality (25°30’31"S 48°30’39"W), a port city with a population of 153,600, and 827 km² area, located in Paraná state, South of Brazil (Fig. 1). The port’s history dates back to 1872, when it was initially used as a berth, until its official foundation in 1935. It has an organized area of 425 km², and is the most important port in south Brazil, moving mainly solid bulk (34 million tons in 2019), but also liquid bulk, and containers (ANTAQ 2020; APPA 2019).

Sample collection

Considering some limiting factors we had to carry out the study (e.g. stolen traps, safety and financial resources), we selected three locations to survey in Paranaguá. Norway rat tissue samples were collected from these three locations along the main road used by trucks to transport goods, which we a priori treated as population units. Samples were obtained from rat carcasses found in the respective areas, or from individuals captured by trapping. We performed nine field campaigns between January 2017 and July 2018, and applied a total effort of 535 trap-nights. Rats were captured in live traps and euthanized for tissue collection following the procedures recommended by UFPR ethics committee (CEUA Nº 1211).

We were interested in characterizing gene flow along the road leading away from the port, and for that purpose we selected three sampling sites in Avenida Ayrton Senna da Silva (Fig. 1, black line). The first sampling location was the Port complex (Fig. 1, cyan circles). Through the establishment of a partnership with the port administration (Administração dos Portos de Paranaguá e Antonina – APPA), we were able to access rat carcasses collected inside the Port area by the zoonosis control team and installed live traps in areas where we had access. For comparison of genetic diversity along the road, rats were sampled in an area ~2 km away from the Port (Area 2) chosen because of the presence of wasteland, residences and a big storage courtyard (Fig. 1, red squares). The third area (Area 3) was ~6 km away from the Port and was chosen because of a gas station, where truckers regularly stay overnight, and surrounded by wasteland and residences (Fig. 1, purple triangles). Both Area 2 and 3 had grassy berms along the road where traps were set and the carcasses were found.
Samples were obtained under License Nº 23,102–1 SISBIO from the Brazilian environmental agency.

**DNA extraction, genotyping, and sequencing**

Genomic DNA extractions were done from muscle tissue samples preserved in 70% ethanol. We used 11 microsatellite markers developed by Jacob et al. (1995) from Rattus norvegicus: D12Rat4, D5Rat83, D7Rat13, D9Rat13, D10Rat20, D16Rat81, D11Mgh5, D15Rat77, D2Rat234, D18Rat96, and D20Rat46. Markers were chosen from different chromosomes to avoid physical linkage. The M13 fluorescent labeling protocol (Schuelke 2000) was used to amplify the loci and incorporate the fluorescent dyes 6-FAM, and NED (Integrated DNA Technologies®).

Polymerase chain reactions (PCR) were performed in 15 µl volumes containing 2 µl of extracted DNA (> 60 ng DNA/ µl), 1 X Buffer, 1.5 mM MgCl2, 0.2 mM dNTPs, 0.1 µM of forward primer, 0.2 µM of each reverse and M13-labelled primer, and 1 U Taq. The thermocycler conditions were: initial denaturation of 94 ºC for 5 min, 25 cycles of 94º for 45 s, 55 ºC for 50 s and 72 ºC for 50 s, followed by 8 cycles of 94º for 45 s, 53 ºC for 40 s and 72 ºC for 50 s, and final extension of 72 ºC for 15 min.

PCR products were mixed with 0.4 µL of GeneScan™ 600 LIZ® (Applied Biosystems) and 7.6 µL of Hi-Di Formamide (Applied Biosystems), followed by heat shock treatment, 95 ºC for 5 min, 4 ºC for 5 min. Genotyping runs were performed on an ABI3500 (Applied Biosystems) automated sequencer and analysed using GENEIOUS prime version 2019.0.4 (www.geneious.com) including the Microsatellite Analysis External Plugin version 1.4.6 (Biomatters Ltd.) Only individuals with at least six genotyped loci were kept in the dataset for analysis.

**Table 1** Measures of genetic diversity per sampling location. Sampling location, number of sampled individuals (N_s), number of alleles reported for 11 loci (N_A), allelic richness (Ar) (Ar; based on min. sample size of 13 individuals), private alleles (PA), within population gene diversity (H_s), and observed heterozygosity (H_O) derived from H_s and F_IS values as H_O = H_s - (F_IS × H_s). F-IS values (none were significant, based on 21,000 randomisations)

|       | N_s | N_A | Ar  | PA | H_s  | H_O  | F_IS |
|-------|-----|-----|-----|----|------|------|------|
| Port  | 25  | 79  | 6.528| 18 | 0.782| 0.784| -0.003|
| Area 2| 24  | 66  | 5.430| 4  | 0.708| 0.672| 0.051 |
| Area 3| 22  | 76  | 6.158| 13 | 0.730| 0.702| 0.038 |
The software Micro-Checker version 2.2.3 (Van Oosterhout et al. 2004) was used to test for the presence of null alleles, false alleles and allelic dropout across the sampling sites data, using Bonferroni correction, and 3,000 randomisations. The number of alleles (\(N_A\)), allelic richness (\(Ar\)), observed heterozygosity (\(H_O\)) and intrapopulation gene diversity (\(H_S\)) according to Nei (1987), mean pairwise \(F_{ST}\) and \(F_{IS}\) for each locus following Weir and Cockerham (1984) (** indicates significance \(p<0.001\), *** \(p<0.01\) and * \(p<0.05\), based on 21,000 randomisations)

| Locus Name | \(N_A\) | \(Ar\) | \(H_O\) | \(H_S\) | \(F_{ST}\) | \(F_{IS}\) |
|------------|--------|--------|--------|--------|----------|----------|
| D5Rat83    | 12     | 7.743  | 0.685  | 0.757  | 0.12***  | 0.105    |
| D10Rat20   | 10     | 6.935  | 0.706  | 0.747  | 0.028*** | 0.046    |
| D2Rat234   | 8      | 6.540  | 0.768  | 0.78   | 0.073*** | 0.019    |
| D20Rat46   | 10     | 6.187  | 0.723  | 0.731  | 0.057*** | 0.028    |
| D11Mgh5    | 9      | 4.396  | 0.692  | 0.588  | 0.117*** | -0.181   |
| D7Rat13    | 13     | 9.516  | 0.773  | 0.829  | 0.060*** | 0.065    |
| D16Rat81   | 14     | 9.174  | 0.782  | 0.839  | 0.053*** | 0.07     |
| D18Rat96   | 11     | 8.289  | 0.783  | 0.782  | 0.083*** | -0.004   |
| D9Rat13    | 7      | 5.351  | 0.675  | 0.709  | 0.027**  | 0.043    |
| D15Rat77   | 7      | 5.280  | 0.774  | 0.726  | 0.01     | -0.063   |
| D12Rat4    | 7      | 4.301  | 0.548  | 0.649  | 0.032**  | 0.158*   |
| Mean       | 9.82   | 6.701  | 0.719  | 0.740  |          |          |

**Results**

Our study included a total of 71 individuals: 25 from the Port area (20 carcasses collected by the zoonosis control team and 5 captured in live traps), 24 from Area 2 (11 carcasses found in the area and 13 captured in live traps) and 22 from the Area 3 (11 carcasses found in the area and 11 captured in live traps). Overall we were able to amplify and genotype 656 of the 781 aimed loci (genotyping success of 83.9%). As expected, the Port sampling site was more run with 200,000 Markov chain Monte Carlo interactions for 10 runs, after a burn-in period of 1,000,000 and number of clusters ranging \(K=1\) to 4. The output file from STRUCTURE was used in STRUCTURE HARVESTER v0.6.94 (Earl and vonHoldt 2012) to calculate \(\Delta K\) values (the rate of change in the log probability of data between successive \(K\) values) as suggested by Evanno et al. (2005).

Individual membership assignments estimated in STRUCTURE were aligned by CLUMPP 1.1.2 (Jakobsson and Rosenberg 2007) and DISTRUCT 1.1 (Rosenberg 2004) was used to generate the bar plot (in Fig. 2).

We used the saddlepoint approximation method to genetically assign individuals using the visualization approach described by McMillan and Fewster (2017) (Fig. 3) and also performed a principal components analysis (PCA) (Fig. 4A) on the raw genetic data using R 3.5.2 with the package geneplot v0.1.0 (McMillan and Fewster 2017). Discriminant analysis of principal component (DAPC) (Fig. 4B) was also used to assess genetic clustering with the package adegenet (Jombart 2008) in R 3.5.2. DAPC relies on data transformation using PCA as a prior step to discriminant analysis and is particularly well suited to assess between-groups variability minimizing variation within the groups, while PCA aims to summarize the overall variability among individuals (Jombart et al. 2010).
Fig. 2 STRUCTURE plot using the admixture model. Estimated proportion of cluster membership for each rat. The most likely structure of $K=2$ across 10 replicates was indicated by STRUCTURE HARVESTER.

Fig. 3 GenePlots for Norway rats in Paranaguá. (A) Genetic assignment for individuals from Port, Area 2 and Area 3 to reference populations Port and Area 2. (B) Genetic assignment for individuals from Port and Area 3 to reference populations Port and Area 3.
diverse, as evident by the number of alleles (n = 79), compared to 66 and 76 alleles for Area 2 and Area 3 respectively. Other indexes showed a similar pattern: allelic richness (Ar = 6.528), gene diversity (Hs = 0.782) and observed heterozygosity (Ho = 0.784) were highest for the Port (Table 1). Port also showed a very low FIS value (-0.003), suggesting that this is the least inbred site among the three analysed. Interestingly, samples from Area 2, which is 2 km away from the Port, showed less genetic diversity than individuals sampled in Area 3, located 6 km away from the Port. Samples from Area 3 also showed similar values compared to the Port for number of alleles, allelic richness and private alleles.

Micro-Checker detected an excess of homozygotes for D12 locus, which can indicate presence of null alleles. Locus D12 was the only marker showing a significant FIS value (FIS = 0.158*), which indicates departure from Hardy–Weinberg equilibrium (HWE) (Table 2). We performed downstream analysis with and without locus D12 (data not shown) and since inclusion of D12 did not have significant effects on the results, the final analysis is based on all 11 markers. No evidence of false alleles or allelic dropout was detected.

Mantel’s test revealed no significant pattern of IBD (r = -0.71, p-value = 0.833) and according to Wright’s (1978) criteria, the pairwise FST values suggested moderate genetic differentiation between Port and Area 2 (FST = 0.0793, p < 0.01) as well as between Area 2 and Area 3 (FST = 0.0642, p < 0.01), but only little differentiation between Port and Area 3 (FST = 0.0456, p < 0.05), even though Port and Area 3 were the locations that were farthest apart. The Jost’s D values also produced a similar result, with Port and Area 3 having the least differentiation. These results are shown in Table 3.

The Bayesian clustering analysis performed in STRUCTURE indicated the most likely number of clusters (K) was two, and further confirmed the FST results (i.e. clustering of Area 3 and Port), and the differentiation of these two sampling sites from Area 2 (Fig. 2). The results from the saddlepoint approximation method used in GenePlots (McMillan and Fewster 2017) corroborated with the STRUCTURE results and FST values, indicating greater genetic similarity between Port and Area 3 than between Port and Area 2. The genetic assignment of individuals using Area 2 and Port as reference populations (Fig. 3A) revealed that more than half of the individuals from Area 3, lie below the tick diagonal line (Port assignment area) indicating greater posterior probability of originating from Port than Area 2. The remaining Area 3 individuals were mostly assigned over the tick diagonal line suggesting an equal probability of the individual originating from Port or Area 2. Lastly, four individuals from Area 3 were plotted above the diagonal lines implying higher probability of originating from Area 2.

Most of individuals sampled in Port were plotted below the 1% line for Area 2 (red dashed line) and therefore have good fit only for the Port population (see also Online Resource 1). By contrast, almost half of the individuals from Area 2 were nested above the Port’s 1% line (cyan dashed line) and between the 1% and 100% line for its own population, indicating that those individuals from Area 2 represents a subset of the Port population.

The genetic assignment using Area 3 and Port as reference (Fig. 3B) highlighted the relation between those two sites. Half of the represented individuals from Area 3 were allocated above the 1% line for Port line (cyan dashed line) and between the 1% and 100% line for its own population, indicating that those Area 3 individuals most likely represent a subset of the Port area.

PCA (Fig. 4A) and DAPC (K = 3) (Fig. 4B) also showed higher associations between Port and Area 3 than between Port and Area 2. PCA detected two genetically divergent individuals from the Port location that did not fit with any of the sampled sites, which might indicate new migrants from an area outside of the sampled region.
Discussion

Many studies of invasive rat species focus on island settings and serve often as eradication assessment (i.e. investigating the eradication viability or its success by verifying the survivor versus reinvader scenario) (e.g. Savidge et al. 2012; Pichlmueller and Russell 2018; Gatto-Almeida et al. 2020b). As pest eradication (i.e. complete removal of all rats) is not feasible on a continental scale, we aimed at gaining a better understanding of rat dispersal in urban areas, which can help to develop and refine management strategies.

In this study we genetically characterized *R. norvegicus* populations in a Brazilian port city, considering a busy road as a dispersal axis. The aim was to evaluate how effective the control measures were in the port area. The samples were collected from three sites along one of the main roads used by trucks to access the port in Paranaguá and results indicated that both areas outside the port presented individuals with good fit to Port population. Additionally, our samples did not show significant IBD pattern (r = -0.71, p-value = 0.833), and even suggested greater genetic similarity between the furthest apart areas, Port and Area 3, while the central location (i.e. Area 2) expected to be more similar to Area 3, if control measures in Port were effective, showed moderate genetic differentiation from the other two.

Roads are known key pathways for the spread of IAS (Brown et al. 2006; Cameron and Bayne 2009) and this spread can occur via self-dispersion (e.g. displacement by walking or swimming) (Innes et al. 2010; Russell et al. 2010; Brown et al. 2006) or human-mediated dispersion, usually associated with long distance dispersal (e.g. transport by vehicles) (Von Der Lippe and Kowarik 2007). Berthier et al. (2016), for example, found that a road network heavily used by trucks for commercial operations facilitates passive dispersal of black rats over long distances through unfavourable landscapes in south-western Niger.

Our target species, *R. norvegicus*, is known to exhibit strong site fidelity and low frequency of long-distance dispersal (> 500 m) in urban environments, especially if food and harbourage are available (Gardner-Santana et al. 2009; Combs et al. 2018; Byers et al. 2019). Hence on the spatial scale of our sampling (6 km), assuming only self-dispersion of Norway rats, we expected to observe an IBD pattern if the control measures in port were not being effective. Individuals from the two closest sampling areas (Port and Area 2) were expected to be more genetically similar and the furthest apart populations (Port and Area 3) would show greater differentiation, as reported by Gardner-Santana et al. (2009) for Norway rats population in Baltimore (East coast, USA).

In Baltimore, the sampled populations were 0.07–13.4 km apart and their pairwise FST values increased as the pairwise distance between areas increased (Gardner-Santana et al. 2009). In France, Desvars-Larrive et al. (2019) also found IBD pattern for two out of three sampled municipalities, but in contrast with those findings, our analysis suggests that the genetic patterns of rats from the two furthest apart sampling areas (Port-Area 3) are more similar, than between either Port-Area 2 or Area 2-Area 3, indicating that rats, in Paranaguá, disperse not only by self-dispersion, but also by human assistance.

Genetic assignment using GenePlots supports this idea as more than 50% of Area 3 individuals showed a good fit for the Port population, while only a few were assigned with confidence to Area 2. Vice versa, four individuals from Port had a good fit for Area 3, suggesting that gene flow occurs in both directions, although it is probably more likely to have individuals from the Port arriving in Area 3, considering GenePlot also suggests Area 3 as a subset of Port.

When individuals from just one of two populations display a good fit to both populations, it suggests that this population is a subset of the other (McMillan and Fewster 2017), like we registered between Port and Area 3. This pattern was also observed in comparison between Port and Area 2. The most likely explanation for these subsetting results is that individuals from the Port location were introduced to Area 2 and Area 3 (i.e. individuals coming from Port established in Area 2 and 3), but a different frequency and/or number of new migrants arriving in the respective areas could explain the difference in genetic differentiation observed.

The Area 2 population possibly received fewer immigrants and hence has undergone a bottleneck, resulting in the loss of alleles. This was reflected in lower diversity indices, such as the lower number of private alleles, number of alleles and allelic richness. In contrast, Area 3 appears to have received new immigrants more frequently, resulting in higher genetic diversity and more genetic similarity to the founder population (Port).

Overall, all three populations showed reasonably genetic distinctions highlighted by the differentiation indexes (Table 3) and by PCA and DAPC plots (Fig. 4). These are congruent with previous studies that evaluated *R. norvegicus* populations in urban environments and found long-distance dispersal occurring at low frequency (Gardner-Santana et al. 2009; Byers et al. 2019). Despite of that, we were also able to register human-mediated transport between two populations of rats 6 km apart. These results suggest that besides self-dispersion, human-mediated dispersion could play an important role in the dispersal dynamics of *R. norvegicus* in Paranaguá. Most of Area 3 samples were collected near a gas station where truck drivers stay overnight with loaded trucks (either coming out or going to the Port). Trucks can accidentally transport rats,
just like vessels (Sanu and Newport 2010) and any area constantly being accessed by these modes of transportation (i.e. gas station) is susceptible to the arrival of new immigrants, facilitating gene flow between populations.

Human-mediated dispersal was also proposed to cause the absence of patterns of isolation by distance in brown rats in eastern France (Desvars-Larrive et al. 2019). The study sampled one urban and two rural municipalities and found a weak IBD pattern between the urban habitat and one of the rural municipalities, while Saint-Romain-de-Popey, the other rural municipality, showed an absence of correlation between genetic diversity and geographical distances. The authors argue that this departure is expected when dispersal between sites is rare or/and when passive dispersal (human-mediated) occurs. They however did not evaluate which human routes could have been involved in dispersal.

On a larger scale, Berthier et al. (2016) evaluated Rattus rattus dispersal between cities apart in Niger (103 to 285 km distance). The authors did not find patterns of IB:IB between the cities, but DAPC analysis and FST values revealed greater genetic similarity between cities connected by roads, even between those furthest apart.

To summarise, our data indicates that even with control measures implemented by the Port authorities, it was not sufficient to prevent dispersal of rats from the Port to surrounding areas and vice versa. The road that we evaluate seems to be an important pathway to R. norvegicus dispersal in Paranaguá and therefore should be kept under pest surveillance, to prevent rats from boarding when loaded trucks park on the road, as much as trucks’ cargo to prevent contaminated cargo to be loaded.

Conclusions

This study is, to our knowledge, the first to explore the role of roads in human-mediated dispersal of Norway rats inside an urban environment in America. Trucks serve as the most probable vector of rat dispersal in the studied area and vehicles can be carrying high number of individuals out of the port and facilitating the establishment of new colonies and invading new areas along the roads in Paranaguá city. Traffic out of the Port area is increasing and hence rat dispersal is expected to occur at increased rates. Therefore, we suggest stronger cargo surveillance, especially when transporting grains and other perishable goods. This would not only lower the risk of unintended rodent dispersal out of the Paranaguá Port, but also avoid cargo contamination by rat droppings and subsequently the potential spread of zoonotic disease. Members of the genus Rattus, notoriously known as reservoir for plague, leptospirosis and hantavirus, have been crucial in the circulation of such diseases in densely populated areas. Often emergence or re-emergence of these diseases is linked to rat dispersal and human activities overlapping in shared habitats. Such contaminations not only pose a risk to human and animal health, they are also a financial burden for cargo companies if the goods are spoiled.

Supplementary information The online version contains supplementary material available at https://doi.org/10.1007/s11252-021-01171-x.

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Author contribution FGA, LMT and IH conceived and designed the research, FGA and TMD collected the data, FGA, AAS and TMD conducted experiments. FGA and FP analysed data. FGA, LMT, FP and IH wrote the manuscript. All authors read and approved the manuscript.

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Data availability Additional information can be found in the electronic supplementary material. The microsatellite data file is available in the Figshare digital repository and can be accessed at: https://doi.org/10.17608/k6.auckland.12363914.

Declarations

Ethical approval Animal handling followed the procedures recommended by UFPR ethics committee (CEUA Nº 1211).

Consent to participate Not applicable.

Consent for publication Not applicable.

Conflicts of interest The authors declare that there is no conflict of interest.

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