Genetic variability of two populations of *Pseudoplatystoma reticulatum* from the Upper Paraguay River Basin

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Abstract

Catfishes of the genus *Pseudoplatystoma* are very important species due to both their high commercial value and their ecological role as voracious predators. They undertake lengthy migratory movements during their life-cycle, this including reproductive migration which occurs from October to December in the rainy season. In the present study, seven microsatellite loci were analyzed to assess genetic variability in two samples of *P. reticulatum* from the Upper Paraguay Basin. The loci were highly polymorphic (mean = 7.28). According to all analysis, the two samples of *P. reticulatum* revealed pronounced genetic differentiation. *Fₚ* value was 0.2290, *Rₚ* value 0.1067 and AMOVA 22.90% (*Fₛ*) and 10.67% (*Rₛ*), all being highly significant (*p* < 0.001). The division of the fishes into two groups was confirmed by microsatellite multi-locus Bayesian assignment testing. The results obtained present evidence of genetic structuring in a *P. reticulatum* population.

Key words: fish, population genetics, microsatellite, homing, population structure.

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Catfishes of the genus *Pseudoplatystoma* belong to the family Pimelodidae, the 93 species of which occurring in the major river basins of South America (Ferraris, 2007). They are migratory species of high commercial value, besides playing an important ecological role in the basins where they occur, due to their predatory behavior (Sato et al., 1988; Miranda, 1997). The upriver migratory movement for reproduction usually occurs from October to December (Resende, 2003). Almost all the stocks of these giant catfishes had declined in the last years (Barthem and Goulding, 2007). The loss of biodiversity in aquatic environments is one among the most serious problems faced by countries all over the world (Moyle and Leidy, 1992), compromising the ecosystem functioning as a whole. The conservation of these ecosystems is vital for various economical sectors in many countries (Ehrlich and Ehrlich, 1992). According to Allan and Flecker (1993), several factors have been identified as causing the decline of fish diversity in several aquatic ecosystems, such as the introduction of exotic species, industrialization, urbanization, destruction of forests and riparian vegetation, pollution by pesticides and gold mining camps, and the construction of physical barriers for the generation of electricity.

The Pantanal area is located at the Upper Paraguay River Basin, with a drainage basin extending from the border between Brazil and Paraguay up to the limits of the Amazon River Basin, and covering about 140,000 km² (Vila da Silva, 1995). In the Pantanal, fisheries constitute the second activity in economic importance. Furthermore, and apart from their ecological importance, fish resources are fundamental for subsistence, amateur, professional and sports fisheries, and the transaction of native crafts (Catella, 2003).

Microsatellite markers have been extensively used in studies on the genetics of fish populations. These markers have a co-dominant inheritance pattern, a high degree of polymorphism, and allow for easy analysis through experiments involving the polymerase chain reaction (PCR) (Wright and Bentzen, 1994; Triantafyllidis et al., 2002; Salgueiro et al., 2003; Barroso et al., 2005; Mäkinen et al., 2006). Studies using microsatellites have revealed pronounced genetic differences even among populations isolated by short geographic distances (Koskinen et al., 2002). In this study, we analyzed two natural populations of *Pseudoplatystoma reticulatum* (“cachara”) collected in the Upper Paraguay River Basin, by using seven microsatellite loci.
loci originally developed for *P. corruscans* to access the genetic diversity of these samples. These data can be used as subsidies for setting up management policies for the development and conservation of these species.

A total of 52 adult specimens of *P. reticulatum* were caught by gill nets at two points on the Paraguay River Basin. 31 specimens were collected from the Paraguay River (PR) (16°04′ 00″ S 57°41′ 00″ W) and 21 from the Jauru River (PJ) (15°51′ 00″ S 27°27′ 00″ W). Fin clips were the source of nuclear DNA. These were collected from freshly caught fish and immediately preserved in 95% ethanol. After collection, the fishes were sold. For the extraction of genomic DNA, about 0.1 mg of tissue was incubated in 200 μL of 5% Chelex (Sigma®) at 65 °C overnight.

The *P. reticulatum* samples were screened for variation at each of the seven microsatellite loci. Five of these loci have already been described by Revaldaves et al. (2005) (Pcor01, Pcor05, Pcor08, Pcor10, and Pcor21), and two by Pereira et al. (2009) (Pcor23 and Pcor28). PCR amplification reactions were conducted in a thermocycler PTC-100 (MJ Research) with a final volume of 12.5 μL, consisting of about 10 ng of DNA, 0.25 μM of each primer, 0.2 mM of dNTP, 1.2 mM of MgCl₂, 0.2 U of Taq-Pht DNA polymerase, 1X PCR buffer (50 mM KCl, 10 mM Tris-HCl, 0.1% Triton X-100, and 1.5 mM MgCl₂) and water. We used the following PCR profile for the loci Pcor01, Pcor02, Pcor05, Pcor08, Pcor21, Pcor23 and Pcor28: initial denaturation at 95 °C for 5 min, 30 cycles of 10 s at 95 °C, 15 s at an annealing temperature of 55 °C, 15 s at 72 °C and a final extension at 72 °C for 10 min. For the loci Pcor10, the PCR profile consisted of an initial denaturation at 95 °C for 5 min, 30 cycles of 30 s at 95 °C, 30 s at an annealing temperature of 48 °C, 30 s at 72 °C and a final extension at 72 °C for 10 min. Amplified products were resolved on 6% polyacrylamide gels stained with silver nitrate. Microsatellite alleles were identified by their size in base-pairs. Allele lengths were estimated by comparison with a 10 bp ladder (10 pb DNA Ladder - Invitrogen), using Kodak Digital Science 1D software.

Allelic count, expected and observed heterozygosity (*H₀*, *Hₑ*), inbreeding coefficient (*Fₛ*) and gene flow (*Nm = 0.25(1-Fₛ)/Fₛ*) were obtained with POPGEN 1.32 software (Yeh and Boyle, 1997). Allelic richness and Nei gene diversity were obtained with Fstat v2.9.3 software (Goudet, 2001). Deviation from Hardy-Weinberg equilibrium (HWE) was tested with the GENEPOP 3.3 package (Raymond and Rousset, 1995). MICRO-CHECKER 2.2.1 (van Oosterhout et al., 2004) software was used to infer the most probable cause of HWE departures.

In order to investigate the genetic structure in samples of *P. reticulatum*, the number of alleles per polymorphic locus ranged from 5 to 11 (Pcor08), and allelic richness from 1.722 (Pcor28) to 9.503 (Pcor08) (Table 1). Nei gene diversity ranged from 0.056 (Pcor28) to 0.870 (Pcor01). A total of 51 alleles were detected, of which 26 were private (Table 2). The private alleles showed frequencies ranging from 0.0167 (six alleles) to 0.3333 (one allele) (Table 2). Expected and observed heterozygosities ranged from 0.0556 (Pcor28 to PRJ) to 0.8615 (Pcor01 - PRJ) and from 0.0556 (Pcor28 - PRP) to 0.5556 (Pcor08 - PRJ and PRP), respectively (Table 1). Significant departures from HWE (p < 0.025 adjusted according to Bonferroni correction) were detected in 8 loci (Table 1). The occurrence of genotyping errors due to null alleles, stuttering or large allele drop-out were checked with the MICRO-CHECKER program. Significant values were, however, not found due to stuttering or large allele drop-out. Estimates of the occurrence of null alleles revealed positive values for all cases in which departure from HWE was identified. The *Fₛ* index suggested the existence of heterozygote deficiency in eight out of 14 comparisons in the populations analyzed (Table 1).

Genetic differentiation between the two populations of *P. reticulatum*, estimated through the *Fₛ* index for all the loci, was 0.2290 and was statistically highly significant (p < 0.025, after Bonferroni correction), thereby showing the existence of strong genetic differentiation among the analyzed samples. The *Rₛ* index estimated for all the loci was 0.1067, which was also statistically highly significant (p < 0.025, after Bonferroni correction).
Individual multi-locus genotypes were used to assign individuals to their respective population of origin. On considering the correct assignment of all individuals, 98.3% were correctly assigned to the PRP population and 99.1% to the PRJ population.

Hierarchical AMOVA revealed that most total genetic variance was to be found within populations $F_{ST} = 77.10\%$ and $R_{ST} = 89.33\%$. The values for variability between populations were $F_{ST} = 22.90\%$ and $R_{ST} = 10.67\%$. These were highly significant ($p < 0.0001$), thus revealing the strong structure of *P. reticulatum* populations.

Structure analysis without admixture inferred that the two populations were genetically distinct, with $K = 2$ populations maximizing the estimated log likelihood in the model (Figures 1 and 2).

The value of gene flow parameter $N_{m}$ was calculated from the mean $F_{ST}$ value. The mean value obtained was $N_{m} = 0.8417$, indicating that some gene exchange had occurred among the sampled populations.

The microsatellites displayed a high degree of polymorphism (mean $= 7.28$), consistent with the mean number observed in other fish species (DeWoody and Avise, 2000).

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**Table 1** - Summary of microsatellite data on each analyzed population of *Pseudoplatystoma reticulatum* analyzed. $N$, Number of individuals; $A$, number of alleles; AR, Allelic Richness; ND, Nei gene diversity; $H_{o}$, observed heterozygosity; $H_{e}$, expected heterozygosity; $F_{IS}$, inbreeding coefficient; HWE result of Hardy-Weinberg probability test on deviation from expected Hardy-Weinberg proportions with $p$-value = 0.05 (adjustment Bonferroni correction $p = 0.025$; $k = 2$). *, significant; ns, not significant and $r$, null allele frequency per loci.

<table>
<thead>
<tr>
<th>Loci</th>
<th>Pcor01</th>
<th>Pcor05</th>
<th>Pcor08</th>
<th>Pcor10</th>
<th>Pcor21</th>
<th>Pcor23</th>
<th>Pcor28</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>P. reticulatum</em> Jauru (PRJ)</td>
<td></td>
<td></td>
<td></td>
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<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>$N$</td>
<td>20</td>
<td>21</td>
<td>18</td>
<td>13</td>
<td>21</td>
<td>21</td>
<td>18</td>
</tr>
<tr>
<td>$A$</td>
<td>9</td>
<td>8</td>
<td>11</td>
<td>2</td>
<td>4</td>
<td>2</td>
<td>2</td>
</tr>
<tr>
<td>AR</td>
<td>8.132</td>
<td>7.085</td>
<td>9.503</td>
<td>2.000</td>
<td>3.475</td>
<td>2.000</td>
<td>1.722</td>
</tr>
<tr>
<td>ND</td>
<td>0.870</td>
<td>0.813</td>
<td>0.856</td>
<td>0.269</td>
<td>0.337</td>
<td>0.343</td>
<td>0.056</td>
</tr>
<tr>
<td>$H_{o}$</td>
<td>0.5500</td>
<td>0.3333</td>
<td>0.5556</td>
<td>0.3077</td>
<td>0.2857</td>
<td>0.3333</td>
<td>0.0556</td>
</tr>
<tr>
<td>$H_{e}$</td>
<td>0.8615</td>
<td>0.8014</td>
<td>0.8508</td>
<td>0.2708</td>
<td>0.3357</td>
<td>0.2846</td>
<td>0.0556</td>
</tr>
<tr>
<td>$F_{IS}$</td>
<td>0.3452</td>
<td>0.5739</td>
<td>0.3284</td>
<td>-0.1818</td>
<td>0.1280</td>
<td>-0.2000</td>
<td>-0.0286</td>
</tr>
<tr>
<td>HWE</td>
<td>(0.0010)*</td>
<td>(0.0000)*</td>
<td>(0.0000)*</td>
<td>(1.0000)ns</td>
<td>(0.0738)ns</td>
<td>(0.5324)ns</td>
<td>(-) ns</td>
</tr>
<tr>
<td>$r$</td>
<td>0.1576</td>
<td>0.2519</td>
<td>0.1472</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
</tbody>
</table>

| *P. reticulatum* Paraguai (PRP) |
| $N$   | 27     | 31     | 18     | 31     | 25     | 30     | 30     |
| $A$   | 5      | 10     | 8      | 3      | 2      | 7      | 5      |
| AR    | 4.888  | 7.430  | 8.824  | 2.706  | 1.775  | 5.597  | 3.641  |
| ND    | 0.661  | 0.756  | 0.819  | 0.210  | 0.078  | 0.767  | 0.301  |
| $H_{o}$ | 0.3704 | 0.5312 | 0.5556 | 0.2258 | 0.0800 | 0.2667 | 0.3000 |
| $H_{e}$ | 0.6401 | 0.7361 | 0.7921 | 0.2089 | 0.0784 | 0.7605 | 0.306  |
| $F_{IS}$ | 0.4105 | 0.2668 | 0.2786 | -0.0987 | -0.0417 | 0.6434 | -0.0150 |
| HWE   | (0.0038)* | (0.0000)* | (0.0024)* | (1.0000)ns | (1.0000)ns | (0.0000)* | (0.2382)ns |
| $r$   | 0.1439 | 0.1185 | 0.1183 | -      | -      | 0.2557 | -      |

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**Table 2** - Private allele counts. Allele number and relative frequency (in parentheses) are listed for each locus analyzed.

<table>
<thead>
<tr>
<th>Loci</th>
<th>Pcor01</th>
<th>Pcor05</th>
<th>Pcor08</th>
<th>Pcor10</th>
<th>Pcor21</th>
<th>Pcor23</th>
<th>Pcor28</th>
</tr>
</thead>
<tbody>
<tr>
<td>Jauru (PRJ)</td>
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<tr>
<td>107 (0.1500)</td>
<td>133 (0.0476)</td>
<td>165 (0.0278)</td>
<td>104 (0.0238)</td>
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<tr>
<td>109 (0.1250)</td>
<td>179 (0.0278)</td>
<td>114 (0.1190)</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>119 (0.0250)</td>
<td>120 (0.0476)</td>
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<td></td>
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<td></td>
<td></td>
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<tr>
<td>121 (0.0500)</td>
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<tr>
<td>Paraguai (PRP)</td>
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<tr>
<td>151 (0.0185)</td>
<td>137 (0.0167)</td>
<td>147 (0.0263)</td>
<td>139 (0.0484)</td>
<td>116 (0.0400)</td>
<td>091 (0.3333)</td>
<td>100 (0.0167)</td>
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<tr>
<td>145 (0.0167)</td>
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<tr>
<td>153 (0.0333)</td>
<td></td>
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<tr>
<td>159 (0.0167)</td>
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</table>
There was significant populational structuring between the two samples of _P. reticulatum_ analyzed, as demonstrated through the different tests employed in this study. The _F_{ST}\text{ and } R_{ST}\text{ index observed between populations of } P. reticulatum\text{ showed high (0.2290) and moderate (0.1067) values, respectively (p < 0.025), thus suggesting there is a strong genetic structure. The differences of values between indices were probably due to differences in the mutation models on which they were based. While } F_{ST}\text{ is based on IAM, } R_{ST}\text{ is based on SMM. The } R_{ST}\text{ index may be the best for microsatellite analysis, and it is expected that } R_{ST}\text{ values under a strict SMM pattern would be higher than those of } F_{ST}\text{ (Slatkin, 1995). However, as can be seen from our results, } F_{ST}\text{ values were higher than } R_{ST}.\text{ This may be explained by the fact that probably not all microsatellite loci evolve strictly in accordance to a SMM model (Balloux and Lugen-Moulin, 2002). This presupposes that mutations occur by the addition or subtraction of a single repetition unit in the microsatellite immediately anterior or posterior to a known and highly related mutation. Departure from this strict SMM pattern would result in the inferior performance of } R_{ST}\text{ in relation to } F_{ST}\text{ (Slatkin, 1995, Balloux et al., 2000). Departure from the SMM model observed in the present study may be related to the fact that in three of our eight loci repetitions were imperfect ( } P_{corr05}^{-(TC)9GC(TC)9-}; P_{corr05}^{-(TC)9CC(TC)15-}\text{ and } P_{corr10}^{-(GTCG)15(GT)9(CC-)}\text{).}

The results obtained through AMOVA analysis revealed the occurrence of significant differentiation between populations of _P. reticulatum_ on using both indices ( } F_{ST}\text{ and } R_{ST},\text{ thereby depicting the occurrence of strong genetic structuring. The values of molecular variation among populations were 22.90% ( } F_{ST}\text{) and 10.67% ( } R_{ST}\text{), both with highly significant p values (p < 0.0001). Through structure analysis without admixture, it was shown that both populations were genetically distinct, with } K = 2 \text{ populations maximizing the estimated log-likelihood in the model (Figures 1 and 2).}

The genetic flow of all loci in the two populations was estimated as } Nm = 0.8417 \text{ migrants per generation. According to Nei (1987), } Nm \text{ values above 1 suggest that genetic flow constitutes a positive factor against genetic differentiation among populations (Spieth, 1974). Thus, our data showed that genetic flow between the two populations analyzed did not exist or was very low, thereby reinforcing the hypothesis of genetic structuring. The results obtained in the assignment tests were extremely positive, as described in the literature for other fish groups (Triantafyllidis et al., 2002). In the present study involving seven microsatellites ( } F_{ST} = 0.2290), 98.3% and 99.1% of
the individuals were correctly assigned to the location from which they were sampled (PRP and PRJ, respectively). These results are consistent with the values found by Pereira et al. (2009) when analyzing six P. corruscans populations (values ranging from 93.6% to 98.2%). Cornuet et al. (1999) showed by simulations that 100% correct assignments can be achieved through the Bayesian method with as few as 10 microsatellite loci and 10 individuals sampled per population, when populations are sufficiently diverged ($F_{ST} \sim 0.1$).

Twenty-six private alleles were found in the two populations of P. reticulatum analyzed. In five of these, frequency was higher than 10% (Table 2). The existence of private alleles places in evidence the absence of gene flow, or at least that it is at a minimum and frequently present in structured populations. Thus, these data reinforce the existence of firm structuring in the populations. Thus, our work renders preliminary evidence of the genetic structure in P. reticulatum. As the results presented are consistent with those obtained for P. corruscans (Pereira et al., 2009) and, on considering the similar behavior between these two species, we suggest that P. reticulatum also presents homing behavior.

The ability to identify and define biological populations is crucial for taking informed decisions concerning conservation and management (Waples and Gaggiotti, 2006). Considering the ecological and economic importance of P. reticulatum, the present data constitute a particularly important element when contemplating their management and conservation. It also places in evidence, the importance of preserving each population to further a positive outcome in the genetic conservation of these species.

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References


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