



## Molecular discrimination of pouched four-eyed opossums from the Mamirauá Reserve in the Brazilian Amazon

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### Abstract

Previous cytochrome B (CytB) mtDNA studies have suggested four species for the opossum genus *Philander* (four-eyed opossums), three (*P. mcilhennyi*, *P. andersoni* and *P. opossum*) from the Amazon and one (*P. frenata*) from the Brazilian Atlantic forest. During a faunal survey nine specimens of *Philander sp.* and four of *Didelphis marsupialis* were collected in the Mamirauá Sustainable Reserve, Amazonas State, Brazil. Preliminary analyses based on morphology and geographical distributions were not conclusive, suggesting that *Philander* specimens could belong to either *P. andersoni* or *P. opossum*. In order to elucidate the relationship of this taxon to the remaining Amazonian taxa, seven *Philander* and two *Didelphis* specimens animals were sequenced for the cytB mtDNA gene and compared to other previously studied taxa. The maximum likelihood (ML), neighbor-joining (NJ) and maximum parsimony (MP) consensus bootstrap trees depicted six groups: *Didelphis*., *P. frenata*, *P. andersoni*, *P. mcilhennyi*, *P. o. opossum* and *Philander sp.* and *Philander canus* in a common assemblage supported by significant bootstrap values, suggesting that the *Philander sp.* from Mamirauá in fact belongs to the species *Philander canus*.

*Key words:* cytochrome B, Mamirauá, molecular phylogeny, mtDNA, *Philander*, pouched four-eyed opossum.

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According to Nowak (1999) the order Marsupialia presents 15 families, with the Didelphidae being the larger extant New World representative of this order. The Didelphidae encompasses 11 recent genera and 77 species, members of which occur from southeastern Canada through the eastern United States and Mexico and into South America to about 47 degrees South in Argentina. Kirsch (1977) proposed the recognition of two living subfamilies: Didelphinae (*Marmosa*, *Monodelphis*, *Lestodelphis*, *Metachirus*, *Didelphis*, *Philander*, *Lutreolina*, *Chironectes*) and Caluromyinae (*Caluromys*, *Caluromysiops*, and *Glironia*). Recently, nuclear gene sequence investigations by Jansa and Voss (2000) have found strong support for the monophyly of both the family Didelphidae (American opossums) and the subfamily Didelphinae. On the other hand, Gardner and Creighton (1989) suggested that the genus *Marmosa* is not monophyletic but is comprised of a series of separate taxa of unknown phylogenetic relationships, these authors creating the genus *Gracilinanus* to include some species while also suggesting that *Marmosops*, *Micoureus*, and *Thylamys*

should be raised to the status of genera. The lack of monophyly in the *Marmosa* has been corroborated by all recent molecular analyses (Jansa and Voss 2000; Kirsch and Palma 1995; Patton *et al.* 1996). In relation to the genus *Philander* (four-eyed opossums), Tiedemann 1808, which is widely distributed and eurytopic in the tropical Amazon forest, the taxonomic status of many of species and subspecies is still highly controversial. According to Voss and Emmons (1996), congeneric species of *Philander* may be sympatric in several parts of the Amazon forest. Based on morphological data, Gardner (1993) recognized two *Philander* species, *P. andersoni* distributed from southern Venezuela to Peru and *P. opossum* occurring from Mexico to Paraguay and northeastern Argentina. Mitochondrial cytochrome B (CytB) data prompted Patton and Silva (1997) to propose four *Philander* species, three (*P. mcilhennyi*, *P. andersoni* and *P. opossum*) from the Amazon region and one (*P. frenata*) from the Brazilian Atlantic forest.

In the study described in this paper, we collected a sample of *Philander* specimens from the Mamirauá Sustainable Developmental Reserve, which is situated at the confluence of the Solimões and Japura rivers between 3°10' S and 64°41' W in the Brazilian state of Amazonas. Preliminary analyses based on morphology and geographical distributions were not conclusive, suggesting that the

specimens could belong either to *P. andersoni* or *P. opossum*. Considering the large amount of data available on didelphid mitochondrial CytB, we decided to test whether or not this marker could be helpful in solving this question. During a faunal survey at the Mamirauá Reserve we used *Tomahawk* (17 x 17 x 61 cm) traps to capture nine *Philander* (two of which were released) and four *Didelphis* specimens (Table 1) which were subsequently sacrificed. The specimens were measured and their skin and skulls prepared and deposited at the Mastozoology collection of the Emilio Goeldi Paraense Museum, Belém (PA), Brazil. Fresh tissues (blood or liver) were collected and preserved in 95% ethanol for the DNA procedures. Table 1 show details of all the opossum specimens used in this study.

The DNA was extracted according to Sambrook *et al.* (1989) and about 440 base pairs of the CytB region of the mitochondrial DNA (mtDNA) was sequenced for seven *Philander* and two *Didelphis* specimens using the MVZ05 (5'CGAAGCTTGATATGAAAAA CCATCGTTG3') and

MVZ14 (5'GGTCTTCATCTYHGGYTTACAAGAC3') primers (Smith and Patton, 1991). The polymerase chain reaction (PCR) was carried out in a final volume of 50 µL containing 10 ng of genomic DNA, 10 mM Tris-HCl (pH 8.85), 25 mM KCl, 5 mM (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, 0.2 mM dNTP, 50 pM of each primer and 1 unit of Taq DNA polymerase (Qiagen, USA). Amplification was performed in a MJ Research thermocycler with a cycling profile of 94 °C for 3 min followed by 30 cycles of 94 °C for 1 min, 40 °C for 45 s, 72 °C for 1 min and an additional extension period of 72 °C for 5 min in the last cycle. The PCR products were purified using ExoSAP-IT (USB, USA) and then sequenced using a dye-terminator cycle sequencing kit in a 377ABI automatic sequencer, according to protocols supplied by the manufacturers (Applied Biosystems, Foster, CA, USA).

The *Philander* and *Didelphis* Mamirauá specimens sequenced in the present work (deposited in GenBank under accession numbers DQ236271-DQ236279) together

**Table 1** - Specimens and cytochrome B (CytB) sequences used in our study.

Taxon <sup>1</sup>	Code <sup>2</sup>	Origin	Opossum specimen register number <sup>3</sup>	Sequence origin
<i>Didelphis</i> species				
<i>D. albiventris</i>	-	Venezuela	-	GenBank U34667
<i>D. marsupialis</i>	1	Rio Juruá, Amazonas (AM), Brazil	-	GenBank U34665
<i>D. marsupialis</i>	2	Mamirauá-AM, Brazil	MPEG24569	Present work
<i>D. marsupialis</i>	3	Mamirauá-AM, Brazil	MPEG24570	Present work
<i>D. virginiana</i>	-	-	-	GenBank <u>Z29573</u>
<i>Philander</i> species				
<i>Philander</i> sp.	1-7	Mamirauá-AM, Brazil	MPEG24571-24573, 26340, 26342, 26343, 26346	Present work
<i>P. andersoni</i>	1	Rio Jaú-AM, Brazil	YL1139	JP <sup>4</sup>
<i>P. andersoni</i>	2	Parque yasuri, Ecuador	ROM104030	JP
<i>P. andersoni</i>	3	Peru	6893	JP
<i>P. andersoni</i>	4	Peru	KU144120	JP
<i>P. frenata</i>	1	Espirito Santo, Brazil	MAM189	JP
<i>P. frenata</i>	2	Rio de Janeiro, Brazil	MNTJ-ORG1	JP
<i>P. mcilhennyi</i>	1	Rio Juruá, Acre state, Brazil	MNFS1103	JP
<i>P. mcilhennyi</i>	2	Rio Juruá-AM, Brazil	JLP15702	JP
<i>P. mcilhennyi</i>	3	Rio Juruá-AM, Brazil	MNFS383	JP
<i>P. mcilhennyi</i>	4	Rio Juruá-AM, Brazil	JLP16069	JP
<i>P. mcilhennyi</i>	5	Rio Urucu-AM, Brazil	MNFS146	JP
<i>P. o canus</i>	1-2	Rio Juruá-AM, Brazil	JLP15395	JP
<i>P. opossum opossum</i>	1	Rio Negro-AM, Brazil	16785	JP
<i>P. o. opossum</i>	2	Guiana	31732	JP
<i>P. o. opossum</i>	3	Guiana	31047	JP
<i>P. o. opossum</i>	4	Rio Xingu, Pará, Brazil	542907	JP
<i>P. o. fuscogriseus</i>	-	Boca del Toro, Panama	USNM464248	JP

<sup>1</sup>Common names: *D. albiventris* = White-eared opossum; *D. marsupialis* = Southern opossum; *P. andersoni* = Black four-eyed opossum; *P. frenata* = A four-eyed opossum; *Philander mcilhennyi* = Mcilhenny's four-eyed opossum; and *P. o. canus*, *P. o. opossum*, *P. o. fuscogriseus* and *Philander* sp. = Gray four-eyed opossum; <sup>2</sup>Code = numbers used in Fig. 1; <sup>3</sup>Emilio Goeldi Paraense Museum, Belem, Brazil; <sup>4</sup>CytB sequences provided by James Patton.

with sequences from the *Philander* species *P. opossum opossum*, *P. opossum fuscogriseus*, *P. opossum canus*, *P. andersoni*, *P. mcilhennyi*, *P. frenata* and the *Didelphis* species *D. albiventris* and *D. virginiana* (kindly provided by James Patton) and the *Didelphis virginiana* GenBank Z29573) were automatically aligned using the Clustal W program (Thompson *et al.*, 1994) and subsequently slightly modified using the BIOEDIT sequence editor (Hall, 1999).

A divergence matrix was constructed using evolutionary model parameters selected by the Modeltest program version 3.06 (Posada and Crandall, 1998) and phylogenetic analysis was carried out using maximum parsimony (MP) and neighbor joining (NJ) analysis and the PAUP\* program version 4b10 (Swofford, 2003). The Treefinder program (Jobb *et al.*, 2004) was used in the maximum likelihood (ML) analyses including bootstraps. Tree topology confidence was assessed by bootstrapping 1000 pseudo-replicates for the MP and NJ analyses and 500 pseudo-replicates for ML analyses.

The CytB sequences were tested for saturation by plotting TN93 distances against transition and transversion values using the DAMBE program (Xia and Xie, 2001) and no indication of saturation was observed even at the highest divergencies. The evolutionary model that best fitted the data set was TN93 (Tamura and Nei, 1993) with rates following a gamma distribution. The parameters of the evolutionary model chosen by the ModelTest used in the PAUP\* program were as follows: Base = 0.3205, 0.2335 and 0.1409; Nst = 6; Rmat = 1.0000, 6.5252, 1.0000, 1.0000 and 19.0258; Rates = gamma; Shape = 0.2352; and Pinvar = 0. Distances were estimated using both the PAUP4b10 and MEGA 2.1 (Kumar *et al.*, 2001) programs, which generate similar values except that Mega 2.1 has the option of comparing average genetic divergencies between groups (Table 2) as well as group standard errors.

We found that the TN93 genetic distances (d) varied from 0.04 to 0.36, with our *Philander sp.* specimens from Mamirauá clearly being most closely related to *P. canus* ( $d = 0.004 \pm 0.02$ ) followed by *P. mcilhennyi* ( $d = 0.047 \pm 0.02$ ). The next grouping most strongly related to our *Philander sp.* specimens was that of *P. andersoni* and *P. opossum* whose distance from the *Philander sp.* ( $d = 0.07$  to

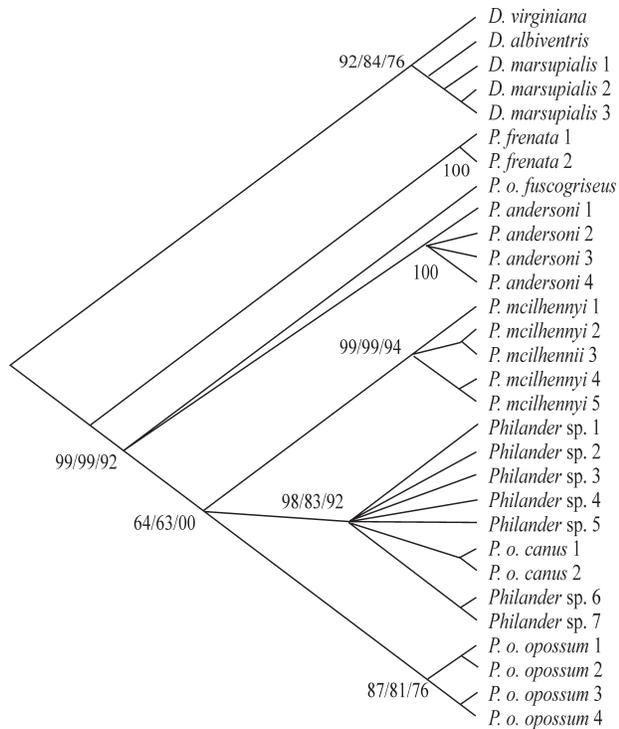
0.08) were quite similar. Our data shows that *P. frenata* and *Didelphis* are undoubtedly the most divergent taxa, being this degree of divergence compatible to differences between distinct genera and similar results having been described previously (Patton and Silva, 1997).

Our MP analyses showed that among the 435 characters, 328 were constant, 24 were parsimony-uninformative and 83 parsimony-informative. The score of the best 7410 equally parsimonious trees was 187 and the consistency index was 0.615. All methods used yielded the same tree topology. The consensus tree is presented in the Figure 1 with bootstrap values at the nodes (ML, NJ and MP). The topology clearly shows six groups: *Didelphis* (ML = 92%, NJ = 84% and MP = 76%); *P. frenata* (ML, NJ and MP = 100%); *P. a. andersoni* (ML, NJ and MP 100%); *P. a. mcilhennyi* (ML = 99%, NJ = 99%, MP = 94%) and *P. o. opossum* (ML = 87%, NJ = 81% and MP = 76%), with *Philander sp.* and *Philander o. canus* belonging to a common assemblage (ML = 98, NJ = 83 and MP = 92%).

Our ML, NJ and MP data suggests that the *Philander sp.* specimens from Mamirauá belong to the species *Philander o. canus*. However, the relationships of *P. o. canus*, *P. andersoni* and *P. mcilhennyi*, *P. o. opossum*, *P. o. fuscogriseus*, were not resolved. Indeed, the bootstrap values supporting *P. o. canus*, *P. mcilhennyi* and *P. o. opossum* (ML = 64%, NJ = 63 and MP = 0%) also suggests an unresolved phylogenetic relationship, as previously observed by Patton and Costa (2003). Clearly, the hypothesis of *P. frenata* from the Brazilian Atlantic forest being a different taxonomic entity (Patton and Costa, 2003) was also confirmed by our work. Indeed, our genetic divergence estimates between *P. frenata* and other *Philander* species shown in Table 2 strongly suggest that they may belong to different genus instead of different species of the same genus. Our genetic divergence estimates using the evolutionary model that best fitted the data showed divergence values of *P. frenata* in relation to the other opossums of the same scale of those from *Didelphis* in relation to the other opossums. Considering our results, the mitochondrial cytochrome B sequences proved to be useful as a complementary tool for taxonomic inference. As pointed out by Patton and Costa (2003) the true diversity of rainforest mar-

**Table 2** - Genetic distances estimated according to Tamura and Nei (1993) modeled by a gamma distribution with alpha parameter equal to 0.23 (lower diagonal). Standard errors estimated using bootstrap (upper diagonal).

	<i>Philander sp.</i>	<i>P. canus</i>	<i>P. andersoni</i>	<i>P. opossum</i>	<i>P. mcilhennyi</i>	<i>P. frenata</i>	<i>Didelphis</i>
<i>Philander sp.</i>		0.002	0.031	0.026	0.018	0.126	0.120
<i>P. canus</i>	0.004		0.029	0.025	0.017	0.133	0.114
<i>P. andersoni</i>	0.087	0.079		0.027	0.032	0.281	0.217
<i>P. opossum</i>	0.072	0.065	0.088		0.031	0.252	0.198
<i>P. mcilhennyi</i>	0.047	0.041	0.100	0.092		0.143	0.184
<i>P. frenata</i>	0.242	0.246	0.345	0.343	0.270		0.133
<i>Didelphis</i>	0.244	0.232	0.347	0.363	0.327	0.286	



**Figure 1** - Consensus tree of 433 base pairs of the mitochondrial Cytochrome B (CytB) gene with bootstrap values on the right corresponding to maximum likelihood (ML), Neighbor-Joining (NJ) and Maximum parsimony (MP) analyses, respectively.

supials will require a concentrated effort of multiple individuals working in collaboration, and we believe that the work reported in our present paper was an initial step in this direction.

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