

An Online mtDNA Tool for Identification of Neotropical Psittacid Species and Taxonomic Issues: A Study Case of the *Amazona ochrocephala* Complex

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Abstract

Parrots are among the most popular pets in the world and they are also some of the most illegally traded, particularly in Brazil. Some computational tools were recently developed by researchers based on molecular databases for taxonomy support, forensic identification and conservation purposes. In this study, the DNA Surveillance platform was used to build an online database tool for molecular identification of Brazilian Psittacids using DNA sequences of six mitochondrial genes. To illustrate possible taxonomic issues of the online tool due to interspecific hybridization or unresolved taxonomy, we focused on *Amazona aestiva* that is considered as one of the most common parrots in Brazil, commonly bred as pets, and considered to be part of a species complex with *Amazona ochrocephala* from South America. We provide three curated sequence databases, which allow the species identification of individuals or tissue samples of birds of the Psittacidae family using mitochondrial DNA markers, and a comprehensive description of a taxonomic issue involving the *A. ochrocephala* complex. The results obtained corroborate previous studies suggesting that these species are not reciprocally monophyletic, due to either an ancient hybridization in central Brazil, or, they maybe just are morpho-varieties of the same species. Alternatively, if *A. aestiva* and *A. ochrocephala* were considered as sister species, the data could be interpreted either as a result of secondary contact or incipient speciation. Beyond the use of mtDNA for spe-

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cies identification, the high mtDNA haplotype diversity observed in *A. aestiva* indicates its potential use in discrimination of lineages that could be an important auxiliary tool to certify the captive origin of legally commercialized parrots.

Keywords

Amazona aestiva, Illegal Trade of Birds, Psittaciformes, Phylogeography, DNA-Surveillance

1. Introduction

Psittacidae is the avian family with the highest percentage of endangered species, with 27% of its species listed as vulnerable, endangered or critically endangered [1]. Psittacids of genus *Amazona* are popularly known as parrots and are among the most threatened of the order Psittaciformes [2]. This genus is comprised of 33 species; 16 of them are threatened and two (*A. martinicana* and *A. violacea*) are extinct, according to the IUCN Red List [3]. All species are listed in Appendix I or II of CITES [4]. The evolutionary history of genus *Amazona* was investigated in a few studies [5]-[7]. The study by Russello and Amato [5] indicated that *Amazona* is not a monophyletic group. *Amazona aestiva*, which is considered as one of the most common parrots in Brazil, forms a species complex with *A. ochrocephala* from South America [5] [7].

Parrots, by virtue of their high ability to imitate the human voice, besides their beauty, gentleness and intelligence, are among the most sought birds as pets in the world and they are common in the illegal traded [8]. It is estimated that only about 5% of psittacids traded in Brazil are captive bred, the remainder being withdrawn from nature [9] [10]. Seizures between 1999 and 2000 by the environment agency from Brazil (IBAMA), indicated that parrots were the 3rd most common animal group illegally traded [9].

Some global efforts aim to expedite and streamline molecular approaches to help taxonomy, forensic identification and conservation. One initiative is the *Barcoding of Life Data Systems*, which links curated specimens and their corresponding molecular data (DNA barcodes), and is publicly accessible on the web [11]. Since 1990's, genetic tracking techniques were introduced in the surveillance of protected animals, in the attempt to inhibit the illegal trade and trafficking. Important studies detected meat from the illegal hunting of whales in popular markets in Japan and Korea, triggering a revision of international agreements controlling whaling, and promoting new resolutions of the International Whaling Commission (IWC) [12]-[14]. Besides, genetic monitoring was adopted for these animal products' control, and it was also used on the surveillance of countries under the jurisdiction of CITES [15], which reported recently that illegal trade routes of whale meat between Japan, Korea and the United States remain active [14].

Custom databases for molecular identification of particular taxa in the DNA Surveillance online tool [16] can be used by forensic genetics against biopiracy and illegal trade. This surveillance started with a tool for identification of cetaceans that includes molecular data organized in a database and identification protocols [13]. The applicability of this tool has been expanded to other taxa, such as for the identification of morphologically similar rat species (*Rattus* spp.) from Southeast Asia and Pacific region [17], and the identification of rodent preys in the stomach content of Carnivores based on the technique of mini-barcodes obtained from DNA extracted from carnivore faeces [18].

In this study, we added in the DNA Surveillance online tool three databases for molecular identification of Brazilian psittacids based on mitochondrial sequences. We also discuss possible limitations of this tool due to taxonomic issues, using as example the species complex *Amazona aestiva/A. ochrocephala*. We sequenced three regions of two mitochondrial genes of various captive specimens from these taxa and analyzed them in the online tool to identify their species. The results showed that phylogenetic and taxonomic issues are essential information to allow a correct species identification.

2. Material and Methods

2.1. Samples, Amplifications and Sequencing

The analyzed blood samples were taken from 46 specimens of *Amazona aestiva* (two *A. aestiva xanthopteryx*,

and the remaining, *A. aestiva aestiva*) kept in captivity in the Vale Verde Ecological Park, a legal commercial breeding facility that is located in the city of Betim, Minas Gerais state, Brazil. All birds are banded, which allows for maintenance of individual records of the birds. The majority of the animals used in this study are founders or matrices from nature, which were originally apprehended by environmental monitoring agents of the Brazilian Government.

Segments of two mitochondrial genes (ND2 and COI) were amplified and sequenced. Early studies with psittacids used the 3' end of COI gene, but with the DNA-Barcode initiative, a growing number of sequences of the 5' end of COI are being produced. Thus, we generated and analyzed these two sections of this gene. The nomenclature and abbreviations used for phylogenetic groups and clades found in our results, tables and figures follow the same ones used in previous studies [7] [19] [20].

For DNA extraction, we used a modified phenol-chloroform-isoamyl alcohol protocol [21]. DNA samples were stored in the collection of the Universidade Federal de Minas Gerais (BD-LBEM), licensed by the Brazilian Ministry of Environment (MMA/CGEN).

Two mitochondrial genes were entirely amplified to avoid amplification of NuMts [22]. The ND2 gene (1041 bp) was amplified with specific primers: H6313 and L5216 [23], and the COI gene (1540pb) was amplified with primers L6615 and H8121 [24]. PCR reactions of ND2 were performed under the following conditions: 94°C for 2 min, 35 cycles of 94°C for 30 s, 63°C for 40 s, 72°C for 2 min, 94°C for 40 s and a final extension of 10 min at 72°C. For COI, we used the same cycle but with annealing temperatures ranging from 58°C to 61°C for 40 seconds. The amplifications were carried out in 12.5 µL reactions containing 0.5 U of *Taq* polymerase (Phonectria), 1 × buffer with 1.5 mM MgCl₂ (Phonectria), 200 µM dNTPs 0.5 µM each primer, and 2 µL of genomic DNA (~40 ng). The amplification products were purified by precipitation in PEG 8000 (20% polyethyleneglycol, 2.5 M NaCl) and finally dissolved in ultrapure water [21]. The sequencing reactions consisted of 35 cycles of 95°C for 25 s, 50°C for 15 s, 60°C for 3 min in a total volume of 10 µL, which contained 4 µL of the sequencing Kit (ET DYE Terminator Kit for Mega BACE, Amersham Biosciences), 3 µL of ultrapure water, 2 µL of purified PCR amplicons, and 1 µL of each primer (0.5 µM final concentration). The following primers were used for sequencing reactions: H6313 and L5216 for ND2 gene, and socoiF1 and H6035COI_Tyr [21], and LCO1490, HCO2198 [24] for COI gene. Sequencing products were purified using ammonium acetate and ethanol, then dissolved with formamide-EDTA buffer and run in the automatic sequencer MegaBACE 1000 (Amersham Biosciences).

2.2. Molecular Analyses

Contigs for each sample were obtained from a total of four forward and reverse sequences, derived from at least two different PCR products, using programs Phred v 0.20425 [25], Phrap v0.990319 and Consed v19.0 [26]. High quality consensus ND2 and COI sequences presented at least a Phred 20 score (99% confidence) for every nucleotide position. Final consensus sequences for each individual are deposited in GenBank (accession numbers JX476306 to JX476426). ND2 and/or COI sequences from 86 individuals of *A. aestiva* and 53 of *A. ochrocephala* (30 of South American subspecies and 23 of Central American subspecies) were downloaded from GenBank [5] [7] [20]. Sequence alignments were obtained using MUSCLE [27] implemented in the program MEGA v5.01 [28].

The programs Arlequin v3.15 [29] and DnaSP v5 [30] were used to estimate the hierarchical distribution of genetic diversity through the Analysis of Molecular Variance (AMOVA) [31], and genetic differences between species (*A. aestiva*, *A. ochrocephala*), subspecies (*A. a. aestiva*, *A. a. xanthopteryx*, *A. o. ochrocephala*, *A. o. xantholaema*, *A. o. nattereri*, *A. o. panamensis*, *A. o. tresmariae*, *A. o. oratrix*, *A. o. auropalliata*, *A. o. belizensis* and “*A. ochrocephala* CO-VE”) and groups (South America [SA], Central America [CA] and Northern South America [NSA]). Arlequin also calculated the haplotype diversity (h), average number of nucleotide differences (k), nucleotide diversity (π), the number of sites with substitutions (S), the composition of nucleotides, frequency haplotypes, Tajima's D and Fu's Fs neutrality tests, which may reveal signs of population expansion (when negative, and P value is less than 0.05). In Arlequin, analyses were performed with the model of pairwise distance (p). We built a haplotype network with the median-joining algorithm (MJ) [32] available in the software NETWORK 4.6 for inferring phylogenetic relationships among haplotypes and their possible geographical correlation.

2.3. Molecular Identification Databases and Tools

Mitochondrial sequences from GenBank (**Supplementary Material SM2**) were used in the three databases. The “Brazilian Parrots” database holds all mitochondrial sequences from all Psittacidae species, and some *Amazona aestiva* and *A. ochrocephala* sequences generated by our approaches. The database “Amazona Identification” presents ND2 and COI sequences of individuals from known geographic origin of the *Amazona aestiva*/*A. ochrocephala* complex. The “All *Amazona*” database has ND2 and COI sequences from genus *Amazona*. These databases were added to the DNA Surveillance platform (<http://www.dna-surveillance.auckland.ac.nz/>). ClustalX 2.012 [33] was used to align and generate these datasets in PHYLIP format.

The query results were extracted in FASTA format and TinySeqXML flatfiles. FASTA files were concatenated with the “Amazona Identification” dataset, aligned with ClustalX 2.012, manually corrected with MEGA v5.01 [28], generated the “All *Amazona*” database haplotypes on DnaSP v5 [30] and realigned at ClustalX to generate DNA Surveillance datasets. With MS Excel 2007, the TinySeqXML flatfiles were converted as spreadsheets and used as database for bulk annotation of DNA Surveillance datasets, exported as Tabbed Separated Values format (TSV).

The online identification protocol aligns the user input sequence by a simple profile alignment against the prealigned dataset of reference sequences, using the penalty values: transitions = 1, transversions = 2, gap creation = 3, gap extension = 1, with the F84 model of substitution to calculate genetic distances and a neighbor-joining (NJ) tree, which is built from the table of distances, graphically and in Newick text format [34].

To test the species identification performance of the domain “Brazilian Psittacidae v1.0” database we used non-curated GenBank sequences obtained with the Entrez queries: “txid9224[Organism:exp] COI”, “txid9224 [Organism:exp] ND2”, “txid9224 [Organism:exp] CytB”, “txid9224 [Organism:exp] 16S” and “txid9224 [Organism:exp] Control Region”. Functionality and confidence tests were performed using the “fvv039” *Amazona* sp. individual as input.

3. Results

3.1. Brazilian Parrots Platform

The three databases and the identification tool are available in the DNA-Surveillance website in the following link: <http://dna-surveillance.fos.auckland.ac.nz:23060/page/parrots/title>.

The “All *Amazona* species” database had 40 COI sequences from 23 species and 16 ND2 sequences from seven species. The “*Amazona* Identification” database had 11 COI sequences and 11 ND2 haplotypes from the *A. aestiva*/*A. ochrocephala* species complex. The “Brazilian Psittacidae v1.0” database had 55 COI sequences from 19 species, 107 ND2 sequences from 25 species, 229 CytB sequences from 68 species, 62 16SrDNA sequences from 25 species and 149 Control Region sequences from 44 species.

Figure 1 shows the result of the identification test of the COI sequence from sample fvv039 using the “*Amazona* Identification” database. This input sequence (query) grouped with those from *A. aestiva* and *A. ochrocephala*, confirming its identification (**Supplementary Material SM1**). The COI sequence from this same sample (fvv039) was tested using the “Brazilian Psittacidae v1.0” database and it correctly grouped with sequences of other samples of *Amazona* (**Supplementary Material SM2**).

3.2. Genetic Diversity and Phylogenetic Relationships

Sequence alignment matrices of genus *Amazona* contained 502 characters of the 5’ end of ND2, and 401 characters of the 3’ end and 474 of 5’ end of COI. ND2 showed the greatest number of variable sites, as well as the highest values of haplotype and nucleotide diversities (0.842 \pm 0.032 and 0.012 \pm 0.006 respectively). The 5’ end COI segment (DNA Barcode) showed the lowest genetic variation, presenting only two parsimony informative sites among eight variable ones, but it was informative for phylogenetic reconstructions.

Diversity indices, numbers of transitions and transversions, and neutrality test results are shown in **Table 1**.

Based on 903 characters of the concatenated sequence of ND2 and the 3’ end of COI of genus *Amazona*, we found 34 haplotypes with 74 substitutions (69 synonymous and five non-synonymous) in 73 polymorphic sites, of which 46 were parsimony informative sites, and 27 corresponded to singleton sites. Genetic diversity was estimated to be 0.016 \pm 0.006 and haplotype diversity, 0.884 \pm 0.028 (**Table 1**). The neutrality tests showed no significant values.

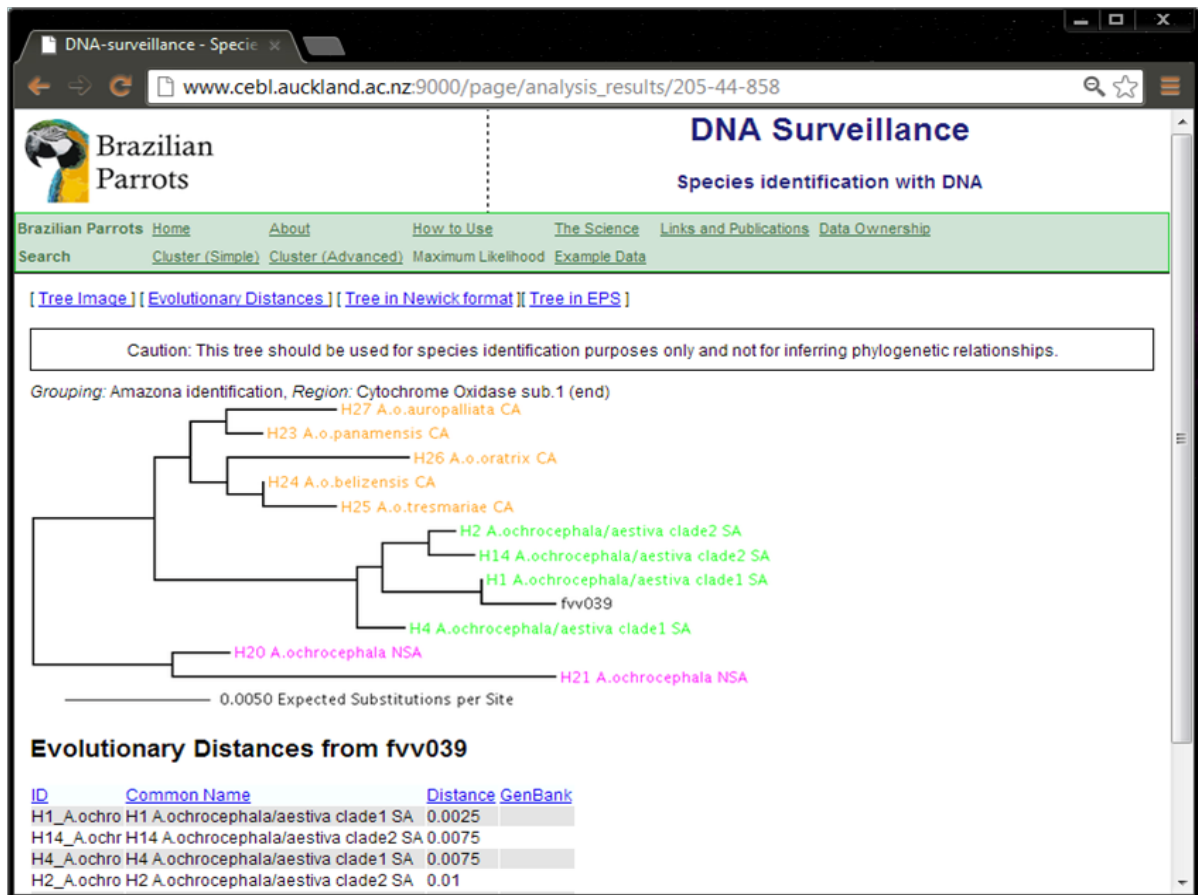


Figure 1. Identification result using sample fvv039 as input data and the “Amazona Identification” database.

Table 1. Indices of genetic diversity, numbers of transitions and transversions, and results of neutrality test indices. H: number of haplotypes, h: haplotype diversity, K: mean number of pairwise differences, π : nucleotide diversity (average over loci), S: no. of sites with substitutions, ts: no. transitions, tv: no. transversions, D: Tajima’s D, and Fu’s FS test. Statistically significant values are highlighted in gray.

	N	Indices of Diversity					ts	tv	Neutrality Tests	
		H	h	k	π	S			D	Fu’s Fs
Concatenated (903 characters)										
Genus <i>Amazona</i>	87	34	0.884	10.407	0.016	73	69	5	-0.92877 (P: 0.18800)	-5.17525 (P: 0.11900)
<i>A. aestiva</i>	45	15	0.715	3.493	0.004	23	23	1	-1.10675 (P: 0.10800)	-3.55190 (P: 0.08900)
<i>A. ochocephala</i> (BR)	19	9	0.901	5.263	0.006	17	17	0	0.31095 (P: 0.67300)	-0.06540 (P: 0.48400)
<i>A. a. aestiva</i>	41	13	0.666	3.027	0.003	21	20	1	-1.26950 (P: 0.08600)	-2.96931 (P: 0.10100)
<i>A. a. xanthopteryx</i>	4	3	0.833	3.167	0.003	5	5	0	1.54056 (P: 0.89300)	0.81143 (P: 0.58400)
<i>A. o. xantholaema</i>	6	2	0.533	0.533	0.001	1	1	0	0.85057 (P: 0.87900)	0.62543 (P: 0.46300)
<i>A. o. nattereri</i>	9	4	0.750	4.000	0.004	11	11	0	-0.05493 (P: 0.48500)	2.00500 (P: 0.85600)
<i>A. o. ochrocephala</i>	4	3	0.833	5.167	0.006	8	8	0	1.81983 (P: 0.93400)	1.55690 (P: 0.71800)
<i>A. ochocephala</i> CO-VE	1	1	1.000	0.000	0.000	0	0	0	-	-
<i>A. o. panamensis</i>	4	1	0.000	0.000	0.000	0	0	0	-	-
<i>A. o. tresmariae</i>	5	5	1.000	3.400	0.004	8	5	3	-0.80734 (P: 0.31500)	-2.00426 (P: 0.03900)

Continued

<i>A. o. oratrix</i>	7	3	0.762	1.047	0.001	2	2	0	1.16843 (P: 0.87800)	0.10980 (P: 0.42600)
<i>A. o. auropalliata</i>	2	2	1.000	2.000	0.002	2	2	0	0.00000 (P: 1.00000)	0.69315 (P: 0.37000)
<i>A. o. belizensis</i>	4	2	0.500	3.500	0.004	7	7	0	-0.81734 (P: 0.12800)	3.25086 (P: 0.90900)
5' ND2 (502 characters)										
<i>Genus Amazona</i>	100	32	0.842	6.033	0.012	53	49	5	-1.30989 (P: 0.06500)	-9.29404 (P: 0.01500)
<i>A. aestiva</i>	57	15	0.659	2.000	0.004	17	16	1	-1.39925 (P: 0.07000)	-6.54483 (P: 0.00200)
<i>A. o. ochocephala (BR)</i>	20	9	0.890	3.074	0.006	10	10	0	0.31634 (P: 0.69700)	-1.66643 (P: 0.19900)
<i>A. a. aestiva</i>	53	13	0.613	1.746	0.003	15	14	1	-1.43307 (P: 0.06300)	-5.44982 (P: 0.00600)
<i>A. a. xanthopteryx</i>	4	3	0.833	2.500	0.005	4	4	0	1.36522 (P: 0.86800)	0.46110 (P: 0.50900)
<i>A. o. xantholaema</i>	6	2	0.533	0.533	0.001	1	1	0	0.85057 (P: 0.86600)	0.62543 (P: 0.46700)
<i>A. o. nattereri</i>	9	4	0.750	2.222	0.004	6	6	0	0.02885 (P: 0.53800)	0.61728 (P: 0.61100)
<i>A. o. ochrocephala</i>	5	4	0.900	3.400	0.007	6	6	0	1.24100 (P: 0.87800)	-0.12801 (P: 0.36400)
<i>A. o. ochocephala CO-VE</i>	1	1	1.000	0.000	0.000	0	0	0	0.00000 (P: 1.00000)	-
<i>A. o. panamensis</i>	4	1	0.000	0.000	0.000	0	0	0	0.00000 (P: 1.00000)	-
<i>A. o. tresmariae</i>	5	5	1.000	3.000	0.006	7	4	3	-0.74682 (P: 0.33500)	-2.23755 (P: 0.02800)
<i>A. o. oratrix</i>	7	3	0.762	1.048	0.002	2	2	0	1.16843 (P: 0.86300)	0.10980 (P: 0.44500)
<i>A. o. auropalliata</i>	2	1	0.000	0.000	0.000	0	0	0	0.00000 (P: 1.00000)	-
<i>A. o. belizensis</i>	4	2	0.500	2.000	0.004	4	4	0	-0.78012 (P: 0.21100)	2.19722 (P: 0.83000)
3' COI (401 characters)										
<i>Genus Amazona</i>	173	27	0.747	3.353	0.008	31	31	0	-1.09586 (P: 0.11900)	-8.87367 (P: 0.01400)
<i>A. aestiva</i>	121	14	0.589	1.639	0.004	14	14	0	-0.99048 (P: 0.18500)	-4.61238 (P: 0.04800)
<i>A. o. ochocephala (BR)</i>	24	7	0.841	1.986	0.005	7	7	0	0.18506 (P: 0.62000)	-0.86489 (P: 0.34600)
<i>A. a. aestiva</i>	97	10	0.456	1.207	0.003	11	11	0	-1.14046 (P: 0.12400)	-3.08310 (P: 0.09900)
<i>A. a. xanthopteryx</i>	24	6	0.500	0.569	0.001	5	5	0	-1.66676 (P: 0.02200)	-3.82216 (P: 0.00200)
<i>A. o. xantholaema</i>	7	1	0.000	0.000	0.000	0	0	0	0.00000 (P: 1.00000)	-
<i>A. o. nattereri</i>	11	5	0.818	1.491	0.004	5	5	0	-0.48520 (P: 0.33500)	-1.08441 (P: 0.16100)
<i>A. o. ochrocephala</i>	6	2	0.533	1.067	0.003	2	2	0	1.03194 (P: 0.86900)	1.72310 (P: 0.77600)
<i>A. o. ochocephala CO-VE</i>	5	4	0.900	3.400	0.009	6	6	0	1.24100 (P: 0.86500)	-0.12801 (P: 0.36300)
<i>A. o. panamensis</i>	5	1	0.000	0.000	0.000	0	0	0	0.00000 (P: 1.00000)	-
<i>A. o. tresmariae</i>	5	2	0.400	0.400	0.001	1	1	0	-0.81650 (P: 0.28800)	0.09021 (P: 0.30100)
<i>A. o. oratrix</i>	7	1	0.000	0.000	0.000	0	0	0	0.00000 (P: 1.00000)	-
<i>A. o. auropalliata</i>	2	2	1.000	2.000	0.005	2	2	0	0.00000 (P: 1.00000)	0.69315 (P: 0.37200)
<i>A. o. belizensis</i>	4	2	0.500	1.500	0.004	3	3	0	-0.75445 (P: 0.23800)	1.71605 (P: 0.75600)
5' COI-DNA Barcode (474 characters)										
<i>A. aestiva</i>	42	10	0.456	0.821	0.002	8	7	1	-1.57664 (P: 0.04100)	-7.08857 (P: 0.00000)
<i>A. a. aestiva</i>	38	8	0.380	0.553	0.001	6	5	1	-1.65321 (P: 0.02400)	-6.30674 (P: 0.00000)
<i>A. a. xanthopteryx</i>	4	4	1.000	2.000	0.004	4	4	0	-0.78012 (P: 0.19700)	-1.87180 (P: 0.02600)

AMOVA based on the concatenated ND2 and 3' end COI matrix showed moderate F_{ST} and Φ_{ST} values (6.9% and 13.5% respectively) between *A. aestiva* and *A. ochrocephala*, which suggested that the greatest genetic diversity is found among individuals and populations within the complex, and not between these taxa (Table 2).

Among the 502 characters of ND2 from the 45 captive *A. aestiva* individuals, 13 haplotypes were found, 10 of which were not yet described for this species. The remaining three haplotypes are shared with *A. ochrocephala* from South America (haplotypes H1, H5 and H10). Genetic diversity was estimated to be 0.003 ± 0.001 and haplotype diversity was 0.656 ± 0.093 . For the COI DNA Barcode segment (474 characters), 10 haplotypes were found, seven of them were only found in the captive specimens, sequenced here and two haplotypes (H1 and H4) were shared with two of three specimens previously sequenced [19].

The haplotype network based on the concatenated sequence matrix of the *A. aestiva/A. ochrocephala* complex showed three main divergent groups (South America [SA], Central America [CA] and northern South America [NSA]) (Figure 2). SA showed many highly divergent haplotypes, with two subgroups (Clade 1 and Clade 2). The star-shaped topology observed in Clade 1 suggests recent population expansion. The most common haplotypes were H1 ($n = 27$) and H5 ($N = 11$), which altogether account for the majority of the *A. aestiva* specimens.

NSA corresponds to a highly divergent *A. ochrocephala* individual from Colombia, which is frequently observed in close relationship with other specimens from Venezuela using other molecular markers in previous

Table 2. AMOVA results.

	ND2 (502 bp)		COI (401 bp)		ND2-COI (903 bp)	
	Φ_{ST}	F_{ST}	Φ_{ST}	F_{ST}	Φ_{ST}	F_{ST}
Between SA, CA, NSA groups	0.59734	0.18371	0.73730	0.30803	0.63712	0.14841
South America (between SA \times NSA)	0.43994	0.22504	0.65709	0.41608	0.47586	0.22855
Between SA (Clade 1 \times Clade 2)	0.38830	0.21707	0.55670	0.42132	0.41253	0.22287
Between <i>A. aestiva</i> \times <i>A. ochrocephala</i>	0.15029	0.06961	0.12170	0.08798	0.13511	0.06899

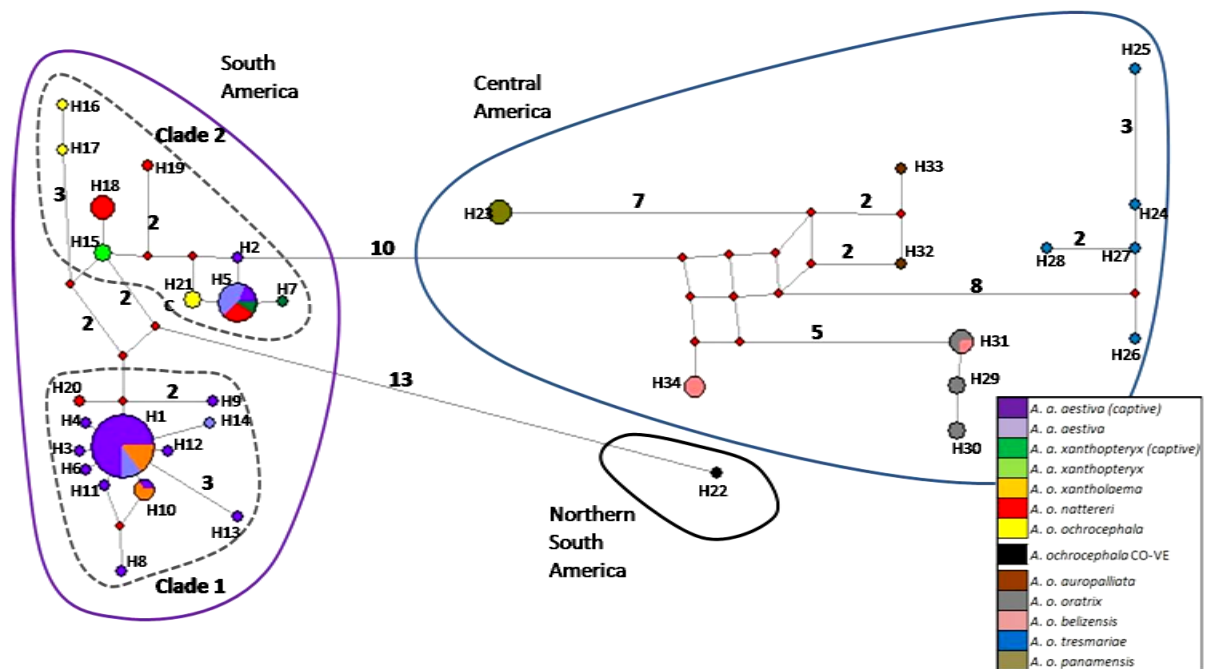


Figure 2. Haplotype network based on 902 bp of ND2 and COI of captive *A. aestiva* and other specimens of the *A. aestiva/A. ochrocephala* complex. The number of substitutions is proportional to the length of the line connecting haplotypes and is also shown. The size of the circles is proportional to the frequency of the haplotype obtained in this sample. Red diamonds represent median vectors.

studies (**Supplementary Material SM3** and **Supplementary Material SM4**). CA group corresponds to *A. ochrocephala auropalliata*, *A. o. oratrix*, *A. o. belizensis*, *A. o. tresmariae*, and *A. o. panamensis* and it forms a clearly separated group from the *A. aestiva/A. ochrocephala* complex from South America. All haplotypes found in captive specimens sequenced here were grouped in Clades 1 and 2 from South America.

The haplotype network based on the 3' end COI segment showed the same topology as previously obtained by Caparroz *et al.* [20], but included more branches and new haplotypes (**Supplementary Material SM3**). The same general group arrangement was also found based only on ND2 (**Supplementary Material SM4**). The haplotype network based on COI-DNA Barcode segment showed no genetic structure (**Supplementary Material SM5**), but its star-like shape is congruent with the results of tests of neutrality tests, thus indicating recent population expansion (**Table 1** and **Figure 1**).

Based on the sampling localities of previously sequenced samples of the *A. aestiva/A. ochrocephala* complex [6] [7] [20] we tried to assign the possible geographical origin of the captive parrots sequenced here (**Table 3**). However, as the majority of the haplotypes presents wide distribution, this analysis not conclusive.

4. Discussion

4.1. Phylogenetic Relationships and the Limits of Species in the *Amazona ochrocephala* Complex

Our data endorse previous studies [6] [7] [20] as members of the *A. aestiva/A. ochrocephala* complex are more closely related amongst themselves than to other *A. ochrocephala* subspecies from Central America. We confirmed previous results [6] [7] [20] [35] that indicated that within this species complex there are three lineages (South America—SA, Central America—CA and North South America—NSA). The current taxonomy of these (once) subspecies from Central America recognizes three species (*A. tresmariae*, *A. oratrix*, and *A. auropalliata*) [36]. In addition, *Amazona barbadensis* [35] (from several isolated populations in dry forest in the coastal areas of Southern Venezuela and Caribbean region) has a strong and basal affiliation with the NSA lineage of the *A. aestiva/A. ochrocephala* complex from South America.

Eberhard and Bermingham [6] recovered a polytomy involving these three lineages. In the study of Ribas *et al.* [9] the SA lineage is sister to the CA lineage, and NSA appears in a basal position, suggesting an ancestral distribution of the group in northern South America with subsequent diversification to the north and south. Urantówka *et al.* [35] tested combinations of genetic markers and exclusion of rapidly evolving sites, and provided support for the close relationship between NSA and SA.

Our results indicate that none of the subspecies of the *A. aestiva/A. ochrocephala* complex from South America (*A. o. ochrocephala*, *A. o. natereri*, *A. o. xantholaema*, *A. a. aestiva*, and *A. a. xanthopteryx*) formed reciprocally monophyletic groups. Instead, *A. aestiva* was paraphyletic in relation to *A. ochrocephala*, as observed in previous studies [6] [7] [20] [35].

Sampling four places representing a lineage that is distributed across the whole Amazon Basin, Eberhard and

Table 3. Possible geographic origin of captive parrots.

Haplotype	Possible geographic area	Conclusive?
3' COI		
H1	TO/BA/MG/DF/GO/Ilha de Marajó, PA	no
H2	MS/Provincia del Chaco, Provincia del Jujuy/Altamira, PA/Beni, Bolivia	no
H4	BA	maybe
ND2		
H1	MG/Ilha de Marajó, PA/Santa Cruz Bolívia/Beni Bolívia, AC/MS	no
H11	Ilha do Marajó, PA	maybe

Brazilian States (TO: Tocantins, BA: Bahia, MG: Minas Gerais, DF: Distrito Federal, PA: Pará, MS: Mato Grosso do Sul, AC: Acre).

Birmingham [6] considered that there is a strong phylogenetic structure in the CA lineage, contrasting with a lack of structure in the SA lineage. Ribas *et al.* [7] showed in a larger sampling (three out of 12 localities represented *A. aestiva*) that there is genetic structure in the SA lineage, with two well-supported and closely related clades. Caparroz *et al.* [20] sampled over eight locations for *A. aestiva*, including six in central Brazil and two in northern Argentina, but the increase of sampling did not show a more structured phylogeographic pattern. However, Caparroz *et al.* [20] observed a major divergence between two groups: “north-eastern” (corresponding to central Brazil states of Bahia, Tocantins, Minas Gerais, Distrito Federal, and Goias), and “southwestern” (Mato Grosso do Sul in Brazil, and northern Argentina). These groups are respectively related to clades 1 and 2 observed here and by Ribas *et al.* [7].

Urantówka *et al.* [35] suggest that NSA represent the descendants of the most ancient lineage of *A. aestiva/A. ochrocephala* complex, that about 1 Mya colonized Venezuela. Thereafter, it colonized Central America originating the CA lineage and, southern regions of South America originating the SA clade. Within the SA lineage, Caparroz *et al.* [20] suggested that Mato Grosso do Sul could represent the ancestral distribution of the two *A. aestiva* clades identified in their study and the current distribution of lineages could be the result of a recent population expansion towards the northeast and the southwest. Considering that the divergence between these two clades occurred in the end of Middle Pleistocene (about 300,000 years ago [20]), this expansion could be related to climatic changes and the cycles of expansion and contraction of the savannah areas at that time [20].

As in [7] [20] [35], we also consider the process of incomplete lineage sorting unlikely to explain the current distribution of the genetic variability of the *A. aestiva/A. ochrocephala* complex from South America, because if this was the case, *A. aestiva* groups should have maintained the same pattern of haplotype distribution found in *A. ochrocephala* during the population expansion events, which is not observed in our data. However, it is possible that hybridization followed by recurrent introgression after isolated diversification could be a better hypothesis. Gene flow due to migration and introgression may have prevented genetic differentiation within the SA clade [7] [35]. The presence of haplotypes shared between *A. aestiva* and South American *A. ochrocephala* (Figure 2) support the hypothesis of historical hybridization/introgression. Alternatively, incipient speciation accompanied by morphological differentiation during population expansion could be an alternative hypothesis. In any case, our results do not support that *A. aestiva* and South American *A. ochrocephala* are two distinct species.

4.2. Species Identification and Traceability of the mtDNA Lineages in Captivity

The illegal removal of animals from their natural environments is one of the main problems to be solved by wildlife protection agencies [37]. Most of the animals that are seized or voluntarily delivered in Brazil are sent to screening centers of wild animals (CETAS) [37]-[39]. Recently, a study using the mitochondrial marker Cyt-b, reported the success of identification of species of 17 specimens (seven carcasses, and ten eggs) of birds seized in an operation of the Brazilian Federal Police [40]. Some specimens were identified as psittacids: *Alipiopsitta xanthops*, *Ara chloropterus*, *Aratinga aurea*, and also *A. aestiva/A. ochrocephala* complex. In a study of all animals sent to CETAS/BH (IBAMA) in the state of Minas Gerais (MG) during the period of 1992-2012 it was shown that several endangered and endemic bird species seized in MG came from other regions of the country [36].

The main objective of the “Brazilian Parrots” tool is to allow species identification of individuals or tissue samples of birds of the Psittacidae family. Unfortunately, the identification of the origin of the sample is a more complex issue as the taxa under investigation should have a well characterized population genetic structure.

The high mtDNA haplotype diversity (H) observed in the captive *A. aestiva* specimens (Table 1) could be used to identify specific maternal lineages. In the case of a legal commercial breeding facility this data could help to certify the captive origin of the commercialized progeny. According to the Brazilian law, only the F1 or subsequent generations of captive bred *A. aestiva* and *A. ochrocephala* can be legally traded. Therefore, the mtDNA could be used as a marker of the maternal lineage of the breeding females used as matrices. This matrilineal genetic marker can be used as a lineage certificate for the progeny that is born in captivity and is legally sold. This molecular certificate can be used to identify, for example, illicit trafficking due to commercialization of wild animals that were illegally implanted with chips or rings, to legalize animals as captive-born. The combination of mitochondrial DNA analyses with other tools and devices such as microchip (properly controlled by

IBAMA-Brazil) could help to reduce illegal trafficking, or at least could facilitate the detection of illegal specimens. In this scenario, a mtDNA database of captive-bred females and their descendants could be created for this purpose, which could be publicly consulted by breeders, animal trade organizations, importers, and institutions worldwide.

Acknowledgements

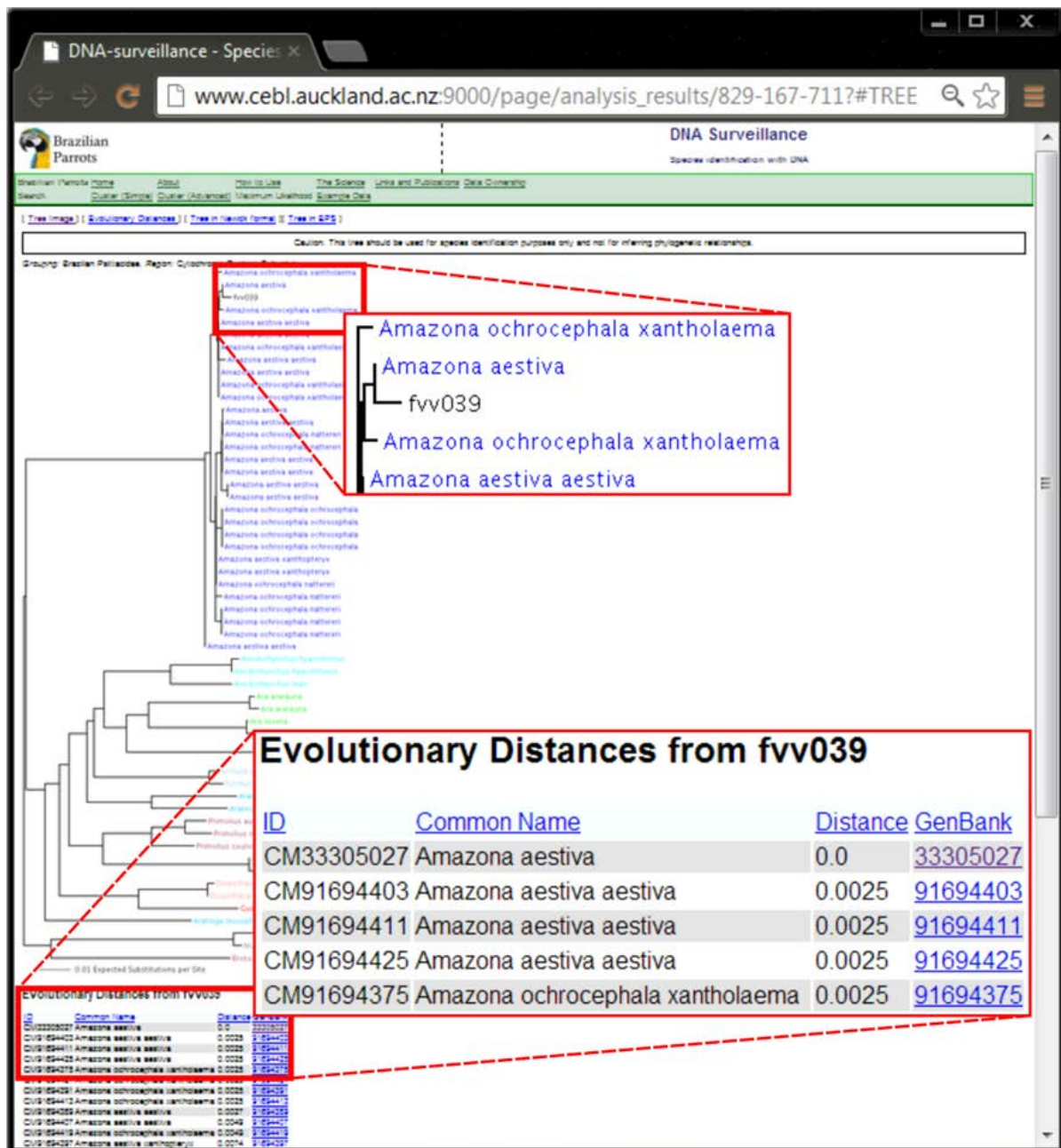
We thank Paulo Machado and Fazenda Vale Verde, who provided blood samples of captive animals used in this study. Thanks to Camila Clozato Lara and Ricardo França who helped in the execution of some laboratory experiments. This study was supported by FAPEMIG (PRONEX), FAPEMIG-Vale SA (RDP-00043-10/2010-1), CNPq (473428/2004-0; 303558/2009-1; 473809/2008-7 and Pro-Reitoria de Pesquisa da UFMG.). A.V.C. received studentship from FAPEMIG and CNPq; F.A.A.S. was supported by FAPEMIG and CNPq; C.Y.M. was supported by FAPESP, CNPq, and CAPES. F.R.S was supported by CNPq.

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Supplementary Material



SM1. Result of the analysis of the ND2 sequence from sample fvv039 in the DNA Surveillance Brazilian parrots database.

SM2. Samples, origin and Genbank accession numbers of *Amazona aestiva* used in this study.

Taxon	Provenance	ID sample	GenBank		
			ND2	COI-Barcode	COI
<i>A. a. aestiva</i>	Criad. Bico Torto	fvv MG 074	JX476391	JX476315	
		fvvmg95153	JX476419	JX476335	JX476370
		CBT31100002	JX476384	JX476308	JX476347
		fvv066	JX476390	JX476314	JX476351
		fvv078	JX476392	JX476316	JX476352
		fvvmg95154	JX476420	JX476343	
		fvvmg95161	JX476393	JX476317	JX476353
		fvv385	JX476407	JX476325	JX476362
	Foz do Iguaçu IBAMA	fvv326	JX476402		JX476360
	IBAMA	fvv MG 95192	JX476394	JX476319	JX476355
		fvv573	JX476411	JX476329	JX476366
		fvv588b	JX476414	JX476342	JX476379
		fvvmg032	JX476417	JX476333	
		fvv049	JX476389	JX476313	
		fvv267	JX476397	JX476320	JX476356
		fvv583	JX476413		
		fvv597	JX476416	JX476332	JX476368
		fvvmg95168	JX476421	JX476336	JX476373
		fvvmg95198	JX476424	JX476338	JX476372
		fvv589	JX476415	JX476331	
		fvvmg059	JX476418	JX476334	JX476369
		fvvmg95169	JX476422	JX476337	JX476371
		fvvmg95189	JX476423		JX476380
		fvvmg039	JX476426	JX476312	JX476350
	Nilsea L. Santos	fvv279	JX476399	JX476340	JX476377
	Zoo Americana	fvv265	JX476395		
		fvv276	JX476398	JX476321	JX476357
		fvv320	JX476401		JX476359
		fvv557	JX476409	JX476327	JX476364
		fvv558	JX476410	JX476328	JX476365
		fvv266	JX476396		
		fvv332	JX476417	JX476333	
		fvv339	JX476405	JX476323	
fvv553		JX476408	JX476326	JX476363	

Continued

		fvv576	JX476412	JX476341	JX476378
	Zoo Brasília	4045	JX476381	JX476339	JX476375
		4056	JX476383	JX476307	JX476346
	Zoo Curitiba	4162	JX476382	JX476306	JX476345
<i>A. aestiva?</i>	?	CONSER004	JX476387	JX476310	JX476348
	?	CONSER005	JX476388	JX476311	JX476349
	?	CONMG172	JX476385		JX476376
	?	CONMG427	JX476386	JX476309	
	?	fvvr97	JX476425	JX476344	JX476374
<i>A. a. xanthopteryx</i>	Foz do Iguaçu IBAMA	fvv343	JX476406	JX476324	JX476361
		fvv310	JX476400		JX476358
<i>A. a. Aestiva</i>	Gurupi, Tocantins, Brazil (TO)	UCB 155			EU340675
		UCB 157			EU340676
		UCB 158			EU340677
		UCB 159			EU340678
		UCB 160			EU340679
		UCB 161			EU340680
		UCB 162			EU340681
		UCB 163			EU340682
		UCB 164			EU340683
		UCB 165			EU340684
		UCB 166			EU340685
		UCB 167			EU340686
		UCB 168			EU340687
		UCB 169			EU340688
	Feira de Santana, Bahia, Brazil (BA)	USP 742			EU340651
		USP 743			EU340652
		USP 744			EU340653
		USP 745			EU340654
		USP 746			EU340655
		USP 747			EU340656
		USP 748			EU340657
	Chapada Gaúcha, Minas Gerais, Brazil (MG)	USP 4051			EU340641
		USP 4052	DQ453647		EU340642
		USP 4053	DQ453657		EU340643
		USP 4054			EU340644
		USP 4055	DQ453648		EU340645
		USP 4056	DQ453650		EU340646
		USP 4057	DQ453646		EU340647
		USP 4058			EU340648
		USP 4059			EU340649
		USP 4060			EU340650

Continued

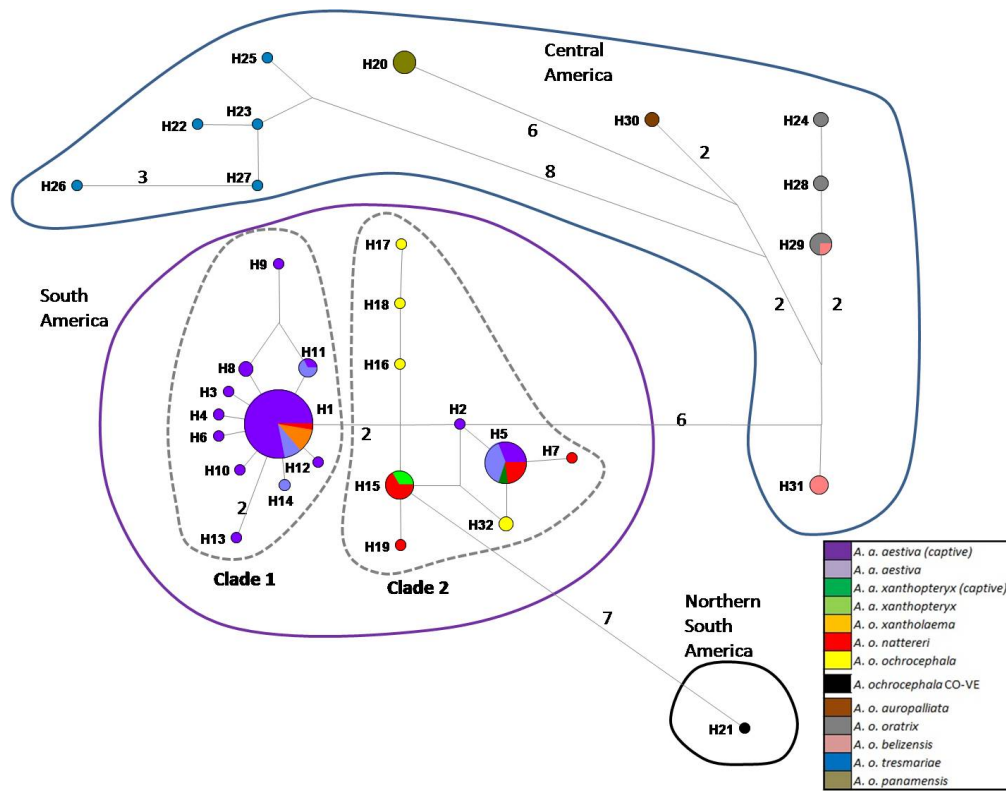
	Distrito Federal, Brazil (DF)	UCB 02		EU340658
		UCB 03		EU340659
		UCB 04		EU340660
		UCB 05		EU340661
		UCB 06		EU340662
		UCB 07		EU340663
		UCB 09		EU340664
		UCB 11		EU340670
		UCB 109		EU340669
		UCB 115		EU340671
		UCB 141		EU340672
		UCB 143		EU340673
	Chapadão do Céu, Goiás, Brazil (GO)	UCB 104		EU340667
		UCB 107		EU340668
		UCB 152		EU340674
	Miranda, Mato Grosso do Sul, Brazil (MS)	USP 2180	DQ453639	EU340628
		USP 2181		EU340629
<i>A. a. aestiva</i>	Miranda, Mato Grosso do Sul, Brazil (MS)	USP 2182		EU340630
		USP 2183	DQ453636	
		USP 2184		EU340631
		USP 2186	DQ453638	EU340632
		USP 2188		EU340633
		USP 2189	DQ453645	
		USP 2190		EU340634
		USP 2192	DQ453656	EU340635
		USP 2193		EU340636
		USP 2195		EU340637
		USP 2196		EU340638
		USP 2199		EU340639
		USP 2201		EU340640
	Unknown (captive bird)	aest1	AY194434	

Continued

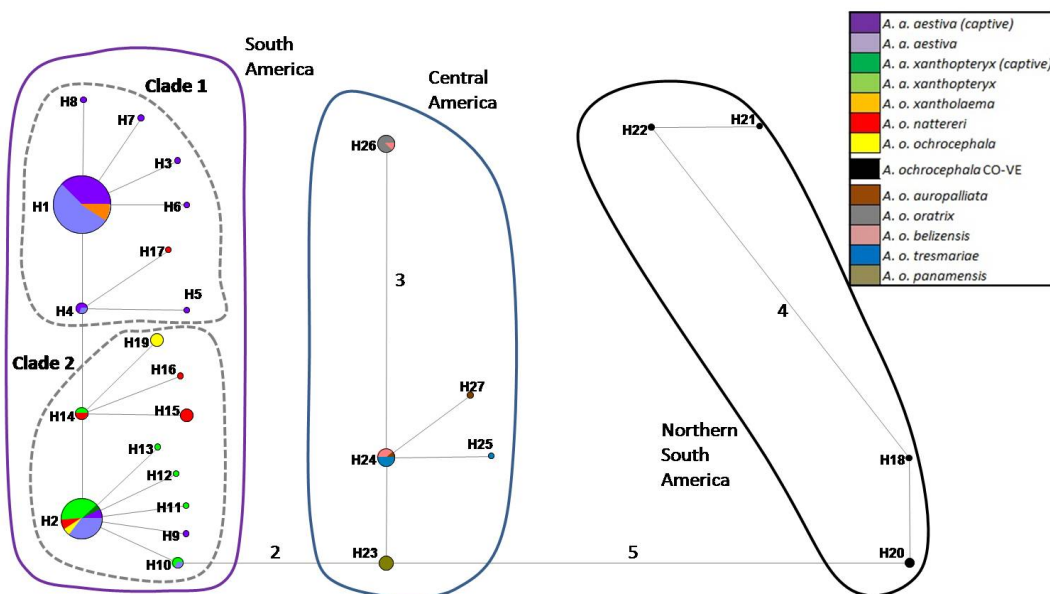
<i>A. a. xanthopteryx</i>	Provincia del Chaco, Argentina (AR1)	UCB 47		EU340689
		UCB 48		EU340690
		UCB 49		EU340691
		UCB 52		EU340692
		UCB 55		EU340693
		UCB 59		EU340694
		UCB 60		EU340695
		UCB 61		EU340696
		UCB 64		EU340697
		UCB 67		EU340698
		UCB 70		EU340699
		UCB 74		EU340700
		UCB 77		EU340701
		UCB 86		EU340702
Provincia del Jujuy, Argentina (AR2)	UCB 97		EU340703	
	UCB 98		EU340704	
	UCB 99		EU340705	
	UCB 100		EU340665	
	UCB 101		EU340666	
Vila Bela da Stsma.Trindade, MT	1320	DQ453643		
	1319	DQ453652		
	MACN-Or-ct_2370		FJ027055	
	MACN-Or-ct_3075		FJ027054	
	MACN-Or-ct_3060		FJ027053	
<i>A. o. xantholaema</i>	Ilha do Marajó, Pará, Brazil (IM)	USP 1587	DQ453640	DQ453612
		USP 1589	DQ453655	DQ453627
		USP 1590	DQ453632	DQ453604
		USP 1031	DQ453651	DQ453623
		USP 1042	DQ453654	DQ453626
	Loro Parque, band# NB-91-4ESY3	xanth1	AY194445	
<i>A. o. ochrocephala</i>	Macapá, Amapá, Brazil (MA)	USP 1556	DQ453659	DQ453631
		USP 1563	DQ453649	DQ453621
		USP 1565	DQ453653	DQ453625
		USP 1572	DQ453635	DQ453607

Continued

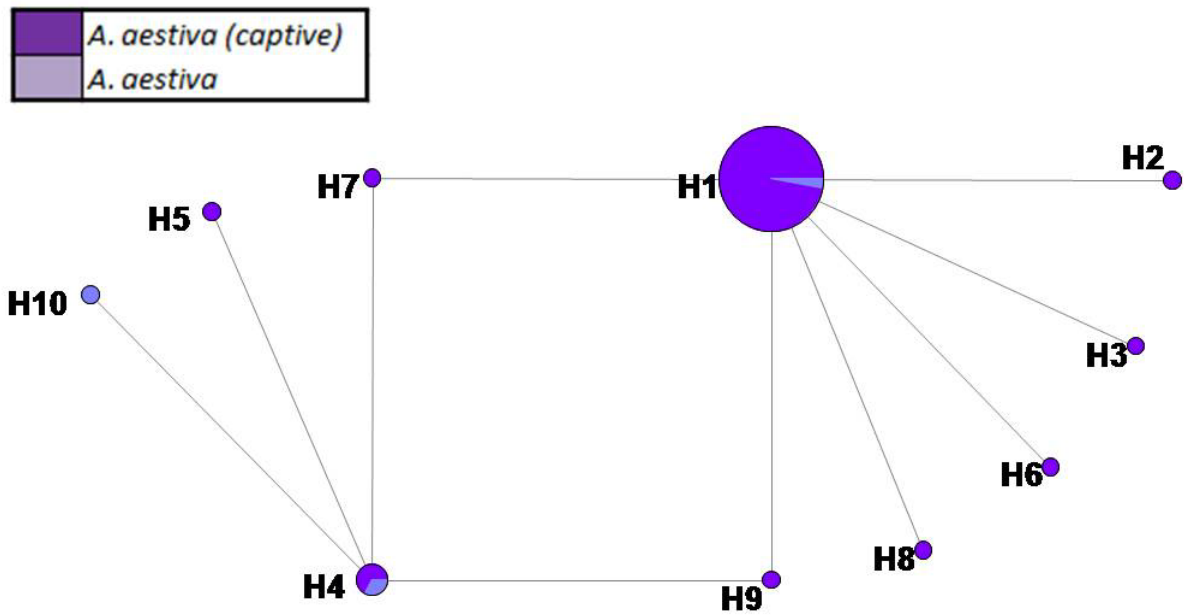
<i>A. o. ochrocephala</i>	Altamira, Pará, Brazil (PA)	NMNH B06867	AY194435	AY194368
		NMNH B07034	AY194436	AY194369
	Carimaguá, Colombia (CO)	STRI-x-61	AY194460	AY194393
	Caicara, Venezuela (VE)	AMNH 177109		AY194400
	Maipures, Venezuela (VE)	AMNH 437237		AY194401
<i>A. o. nattereri</i>	Jacareacanga, Pará, Brazil (JA)	USP 2951	DQ453642	DQ453614
	Xapurí, Acre, Brazil (AC)	USP 2078	DQ453637	DQ453609
	Rio Iaco, Acre, Brazil (AC)	USP 2068	DQ453644	DQ453616
	Basiléia, Acre, Brazil (AC)	USP 2074	DQ453641	DQ453613
	Assis, Acre, Brazil (AC)	USP 2076	DQ453634	DQ453606
	Rio Itimarí, Acre, Brazil (AC)	USP 2084	DQ453633	DQ453605
	Beni, Bolivia (BE)	LSU B-25220	AY194438	AY194371
	Pando Department, Bolivia (BO)	USP 2075	DQ453658	DQ453630
		LSU B9409	AY194439	AY194372
Santa Cruz Department, Bolivia (SC)	LSU B12973	AY194437	AY194370	
<i>A. o. panamensis</i>	Coclé, Panama	STRI-x-26	AY194462	AY194395
	Chiriqui, Panama	STRI-x-27	AY194463	AY194396
		STRI-x-30	AY194464	AY194397
		STRI-x-34	AY194465	AY194398
<i>A. o. auropalliata</i>	Guanacaste, Costa Rica	STRI-x-98	AY194444	AY194377
	Chiapas; Marqués de Comillas, Mexico	auro2	AY194449	
<i>A. o. oratrix</i>	Tamaulipas, Los Colorados, México	STRI-x-47		AY194390
	Mexico: Tamaulipas; Los Colorados	ort1	AY194451	
		ort2	AY194452	
		ort3	AY194453	
		ort4	AY194457	
		ort5	AY194450	
	Mexico: Veracruz; Tempoal	ort6	AY194447	
ort7		AY194448		
<i>A. o. tresmariae</i>	Nayarit, Isla Maria Madre, Mexico	STRI-x-50	AY194454	AY194388
		STRI-x-51	AY194455	AY194389
		tres3	AY194456	
		tres4	AY194458	
		tres5	AY194459	
<i>A. o. belizensis</i>	Belize Zoo, band # 90185	beliz1	AY194440	
		beliz2	AY194441	
		beliz3	AY194442	
		beliz4	AY194443	



SM3. Haplotype network based on 502 bp of ND2 of captive *A. aestiva* and other specimens of the *A. aestiva/A. ochocephala* complex. The number of substitutions is proportional to the length of the line connecting haplotypes and is also shown. The size of the circles is proportional to the frequency of the haplotype obtained in this sample. Red diamonds represent median vectors.



SM4. Haplotype network based on 474 bp of COI of captive *A. aestiva* and other specimens of the *A. aestiva/A. ochocephala* complex. The number of substitutions is proportional to the length of the line connecting haplotypes and is also shown. The size of the circles is proportional to the frequency of the haplotype obtained in this sample.



SM5. Haplotype network based on 474 bp of COI-DNA Barcode of captive *A. aestiva* and other specimens of the *A. aestiva*/*A. ochocephala* complex. The size of the circles is proportional to the frequency of the haplotype obtained in this sample.

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