

Hemosporidian parasites of free-living birds in the São Paulo Zoo, Brazil

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Abstract Numerous studies addressed the diversity of bird *Plasmodium* and *Haemoproteus* parasites. However, a few have been carried out in continental avian hotspot regions such as Brazil, a country with markedly different biomes, including Amazon, Brazilian Savanna, Atlantic Forest, Caatinga, Pantanal, and Pampas. We present the first study on hemosporidian (Haemosporida) parasites in free-living birds from an Atlantic Forest fragment where more than 80 avian species have been reported. Within this area, the São Paulo Zoo locates, and it is the fourth largest zoo in the world and the largest in Latin America. A total of 133 free-living bird samples representing 12 species were collected in the zoo, with the overall hemosporidian prevalence of 18 % by PCR-based diagnostics. Twenty-four positive PCR signals were reported from four different bird species, including migratory ones. *Columba livia*, an urban species, considered nowadays a pest in big cities, showed 100 % prevalence of *Haemoproteus* spp., mainly *Haemoproteus columbae*. We discuss the

epidemiological importance of new parasites introduced by migratory birds in the São Paulo Zoo area and the risk it poses to the captive species, which are natives or exotics. We also warn about the influence these parasites can have on the biodiversity and the structure of host populations by altering the competitive interaction between the free-living and the captive birds.

Keywords *Plasmodium* · *Haemoproteus* · Free-living birds · Atlantic Forest · São Paulo Zoo

Introduction

Avian hemosporidians (Sporozoa, Haemosporida) are protozoan parasites with over 250 species of the genera *Plasmodium*, *Haemoproteus*, *Leucocytozoon*, and *Fallisia*, which are described in birds. They are transmitted exclusively by dipteran blood-sucking insects (Garnham 1966; Atkinson et al. 2008; Valkiūnas 2005). Although these parasites are cosmopolitan in distribution, a review of global diversity of *Plasmodium* and *Haemoproteus* genetic lineages has shown a substantial geographical bias in studies on continental avian non-hotspot regions in Europe and North America whose focus has been mainly on the Passeriformes species (Clark et al. 2014).

Brazil hosts one of the greatest diversities of fauna and flora in the world, consisting of markedly different biomes (Amazon, Cerrado, Atlantic Forest, Caatinga, Pantanal, and Pampas) and is also considered a hotspot for avian richness (Orme et al. 2005). However, more specifically for the Atlantic Forest, a few studies on the prevalence of blood parasites in birds have been conducted (Ribeiro et al. 2005; Belo et al. 2009; Sebaio et al. 2012; Lacorte et al. 2013; Motta et al. 2013). São Paulo

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is the second Brazilian state to hold large remaining areas of Atlantic forest (<http://mapas.sosma.org.br/dados/>) that is one of world's biodiversity hotspots (Ribeiro et al. 2009). In the county of São Paulo, the state's capital, there is an area of 26,664 km² of forest that includes a state park named “Fontes do Ipiranga” (Parque Estadual das Fontes do Ipiranga—PEFI), where over 80 avian species have been reported. Within the PEFI, the São Paulo Zoo locates. This is the fourth largest zoo in the world and the largest in Latin America. This area has a high density and diversity of mosquitoes that can transmit avian malaria (Bueno et al. 2010), and *Plasmodium (Novyella) nucleophilum* has been identified and characterized molecularly from a captive goose born at the Zoo (Chagas et al. 2013). Biting midges (Ceratopogonidae) and hippoboscid flies (Hippoboscidae), which are vectors of avian *Haemoproteus* species, have also been reported in the São Paulo State (Felippe-Bauer and Oliveira 2001; Arzua and Valim 2010). In this paper, we present results of the first study on hemosporidian parasites in free-living birds from this area. We also discuss the effects of new parasites introduced by migratory birds to the São Paulo Zoo area and the possible risk of such introduction to the exotic captive bird species.

Materials and methods

Study area

Established in 1958, the São Paulo Zoo is located in an area of 824,529 m² within one of the last Atlantic Forest remnants within the City of São Paulo, the largest city in Brazil (Fig. 1a). The waters of the Ipiranga stream form a lake that receives several wild species of birds including migratory species (Fig. 1b). Around the lake, the forest shelters wild native animals, forming a diverse wildlife along with the zoo animal collection where more than 3000 animals are on display, representing species of mammals, birds, reptiles, amphibians, and invertebrates. During migration and the wintering of birds from various American countries, as well as from other Brazilian regions, the space destined for zoo animals is shared by migratory species, native and opportunistic species. The latter are behaviorally flexible birds living in variable environmental conditions and sustaining from different food sources.

Collection of blood samples

Samples of different bird species were collected between September 2012 and June 2015. Healthy individual samples were collected primarily for other projects: blood from *Dendrocygna viduata* was collected for a research project

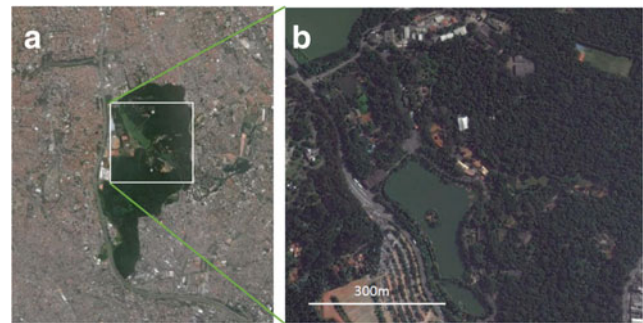


Fig. 1 Satellite images of the Atlantic forest fragment within the city of São Paulo (a) and the São Paulo Zoo within this fragment (b). Google Earth

aiming isolation of microorganisms and bacterial and fungal resistance genes in migrating birds (SISBIO 35166-2); blood from *Penelope superciliaris* was collected for the study of the biology of *Penelope* sp. (Galliformes: Cracidae) and its ecological importance in the PEFI (SISBIO 41010-1). All other samples were collected for routine activities of the veterinary staff in the park since the birds presented clinical signs of diseases (SISBIO 34605-2).

For capturing *Penelope* sp., a trap of wooden structure measuring 80.0 × 80.0 × 6.0 cm as a base and covered with mesh braided line fishing net of 1.0 × 1.0 cm was installed on the ground in the most suitable locations in order to minimize stress and injuries to the birds. To attract birds to the trap, cracked corn was used as bait. For capturing the Anatidae species, three traps of 3.0 × 2.4 × 0.3 m were placed on the lake border. These traps were built of wood and fence with doors in the upper side to facilitate the birds' removal. To attract the birds, we use the same methodology as for catching *Penelope* sp., but rice and Anatidae species food were added to bait, as established by CEMAVE/IBAMA (1994). The sampled birds were ringed after identification and blood sampling; CEMAVE's model rings were used.

Blood sampling and microscopic examination

After being caught, the birds were physically restrained and the venous blood was collected. A thin blood smear was prepared without anticoagulant and the remaining blood was placed in a lithium heparin tube and frozen for DNA extraction. The thin blood smear was fixed by 100 % methanol, stained by Giemsa in a work solution prepared with buffered water (10 % stain stock solution was used), and then examined microscopically for 20–25 min: 100 fields were examined at low magnification (×400) and 100 fields were viewed at high magnification (×1000) (Valkiūnas 2005) using an Olympus BX41 light microscope. The intensity of parasitemia was determined by actual counting of the number of parasites per 1000 erythrocytes, as recommended by Godfrey et al. (1987).

Genomic DNA (gDNA) extraction, PCR amplification of *Haemoproteus* sp. and *Plasmodium* sp. *cytb* fragments, sequencing, and the sequence data analysis

DNA from blood samples was extracted with the Wizard® SV 96 Genomic DNA Purification System (Promega) with the following modifications. First, 10 µl of red blood cell pellets was completed to 200 µl with ultrapure water and an initial lysis was performed with Proteinase K (0.8 mg/200 µl sample) for 1 h at 37 °C. Next, 400 µl of Whole Blood Lysis Buffer (Wizard® SV Lysis Buffer with 10 % Triton X-100) was added and incubated overnight at room temperature. The lysates were transferred into the columns and washed according to the manufacturer's instructions. The gDNA was eluted in 100 µl of Nuclease-Free Water and stored at -20 °C. The PCR reactions were conducted using a nested polymerase chain reaction targeting the mitochondrial cytochrome *b* (*cytb*) gene of *Haemoproteus* and *Plasmodium* (Hellgren et al. 2004). Primers HaemNFI and HaemNR3 and 50 ng gDNA were used in the first reaction and 1-µl aliquot of this product was used as a template for a nested reaction with primers HaemF and HaemR2. Two blood samples with different parasitemias (<0.01 and 6.49 %) of *Plasmodium* sp. were used as positive control, and ultrapure water was used as a negative control. The PCR product was sequenced by Big Dye Terminator v3.0 Cycle Sequencing Kit in ABI Genetic Analyzer (ABI, USA), using PCR oligonucleotides (HaemF and HaemR2). The sequences of ~480 bp were obtained and used in this study. Sequences possessing at least one different nucleotide were considered unique lineages. In order to determine already available lineage names, the obtained sequences were aligned with sequences deposited in MalAvi database (available at <http://mbio-serv2.mbioekol.lu.se/Malavi/blast.html>, see Bensch et al. 2009) using BLASTN (Basic Local Alignment Search Tool, see Altschul et al. 1997). Those absent from the MalAvi database were considered as new and were named according to the MalAvi nomenclature (Bensch et al. 2009) and deposited in GenBank.

Phylogenetic analysis

The phylogenetic relationship among reported parasites was inferred using *cytb* gene sequences. The phylogenetic reconstruction was performed separately for *Haemoproteus* and *Plasmodium* lineages, with 24 sequences used in each analysis. The phylogenetic tree was constructed using the Bayesian inference method implemented in MrBayes v3.2.0 (Huelsenbeck and Ronquist 2001). Bayesian inference was performed with two Markov Chain Monte Carlo searches of 3 million generations, each with sampling of 1 in 300 trees. After a burn-in of 25 %, the remaining 15,002 trees were used to calculate the 50 % majority-rule consensus tree. The

phylogeny was visualized using FigTree version 1.4.0 available at <http://tree.bio.ed.ac.uk/software/figtree/> (Rambaut 2006).

Results

In total, 133 samples belonging to 124 birds from 12 species were collected and screened for *Plasmodium* or *Haemoproteus* infections (Table 1). We detected an overall prevalence of 18 % (24 positive samples) by PCR-based diagnostics. These positive results were found in four different bird species (Table 1), but the prevalence varied significantly among these hosts. *Columba livia* had highest prevalence (100 %), while other eight sampled bird species were not infected. Of these PCR positive samples, 21 samples were also positive by microscopic examination: one was positive for *Plasmodium* sp., with a light parasitemia, and 20 samples positive for *Haemoproteus*, with parasitemias ranging between 0.3 and 15.5 %. No co-infection with hemosporidians belonging to the same or different genera was detected by microscopy. *Haemoproteus* (*Haemoproteus*) *columbae* (Fig. 2) was the only species of hemosporidian parasites morphologically identified. For the first time, we provided morphological support for molecular characterization of several lineages of this parasite (Figs. 2 and 3). Although *Plasmodium* sp. was seen, the parasites were not identified to species level because of light parasitemia.

The *cytb* sequences recovered from the PCR-positive samples showed four birds positive for *Plasmodium* spp. (3 %) and 20 positive for *Haemoproteus* spp. (15 %), with seven different *cytb* lineages (three of *Plasmodium* and four of *Haemoproteus* lineages), of which five were obtained from resident birds and two from migratory birds (Table 1). We detected four new lineages: (1) *C. livia* (sample identification number, ID 481) presented a sequence with 98 % of identity with the lineage AFR119 (*Haemoproteus* sp.); (2) *D. viduata* (ID 263) presented a sequence with 98 % of identity with STUSUP02 and VIOLI07 (*Plasmodium* sp.); (3) *D. viduata* (ID 249) presented a sequence with 99 % of identity with GRW06 (*Plasmodium elongatum*); and (4) *P. superciliaris* (ID 885) and *N. nycticorax* (ID 854) presented a sequence with 99 % of identity with PESA01 (*Plasmodium* sp.). Three *Haemoproteus* lineages had been already deposited in MalAvi: HAECOL1, COQUI05, and COLIV03. However, only the first one has been attributed to certain morphospecies (*H. columbae*).

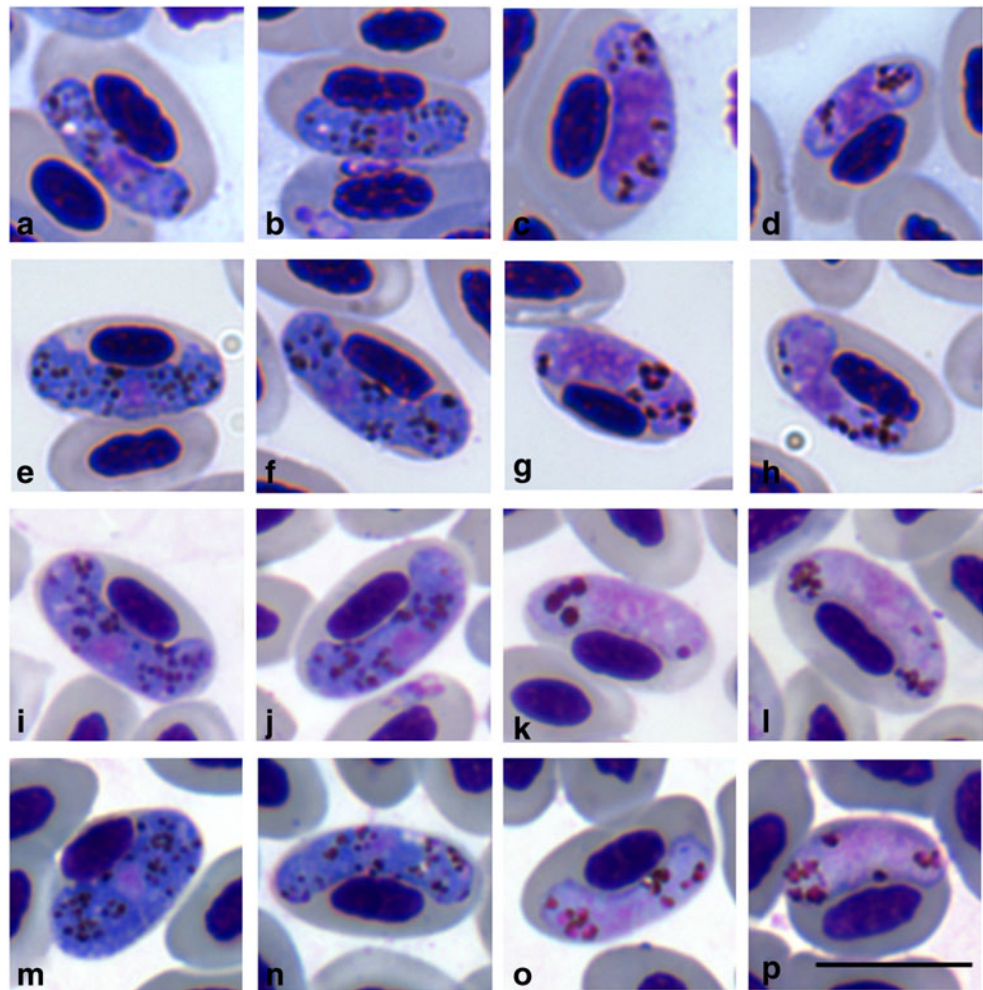
The phylogenetic reconstruction of *Haemoproteus* spp. lineages showed two well-supported clusters (posterior probability is of 100 %), corresponding to each subgenus of this genus (Fig. 3, clades a and b). One included species of the subgenus *Haemoproteus* (*Haemoproteus*), with *H. columbae* lineage HAECOL1, *Haemoproteus multipigmentatus* and

Table 1 Birds sampled and the results of PCR-based and microscopic diagnostics of *Plasmodium* and *Haemoproteus* parasites and their lineages

Order (Family)	Host species (Common name)	Number of samples	Number of PCR (positive)	Smear (positive)	(Sample ID) parasitemia, %	Hemosporidian lineages identified (parasite morphospecies)	GenBank number
Anseriformes (Anatidae)	<i>Dendrocygna bicolor</i> ^a (fulvous whistling duck)	4	3 (0)	4 (0)	–	–	–
	<i>Dendrocygna viduata</i> ^a (white-faced whistling duck)	84	81 (2)	81 (1)	(263) <0.01 %	DENV1D01 (263) (<i>Plasmodium</i> sp.) DENV1D02 (249) (<i>P. elongatum</i>) ^c	KU057966 KU057965
Columbiformes (Columbidae)	<i>Columba livia</i> (rock dove)	20	20 (20)	20 (20)	(425) 12.0 (430) 1.8 (478) 15.5 (481) 0.3	COLIV03 ^b (<i>H. columbae</i>) COQU105 ^b (<i>H. columbae</i>) COLIV06 (<i>Haemoproteus</i> sp.) ^d + <i>H. columbae</i> ^e HAECOL1 ^b (<i>H. columbae</i>) ^e	KU131584 KU131585 KU057964 KU131583
Falconiformes (Falconidae)	<i>Caracara plancus</i> (southern crested caracara)	3	3 (0)	3 (0)	–	–	–
Galliformes (Cracidae)	<i>Penelope superciliosus</i> (rusty-margined guan)	6	6 (1)	3 (0)	–	–	–
Piciformes (Ramphastidae)	<i>Ramphastos dicolorus</i> (red-breasted toucan)	1	1 (0)	1 (0)	–	–	–
Pelecaniformes (Ardeidae)	<i>Egretta thula</i> (Snowy Egret)	1	1 (0)	1 (0)	–	–	–
	<i>Nycticorax nycticorax</i> (black-crowned night heron)	3	3 (1)	3 (0)	–	–	–
Strigiformes (Strigidae)	<i>Asio clamator</i> (striped owl)	6	6 (0)	6 (0)	–	–	–
	<i>Megascops choliba</i> (tropical screech owl)	1	1 (0)	1 (0)	–	–	–
	<i>Athene cunicularia</i> (burrowing owl)	2	2 (0)	2 (0)	–	–	–
Suliformes (Phalacrocoracidae)	<i>Phalacrocorax brasilianus</i> (Neotropical cormorant)	2	2 (0)	2 (0)	–	–	–
					(424) 1.4, (426) 0.4, (427) 3.6, (428) 2.2, (429) 1.0, (431) 2.3, (432) 3.0, (433) 0.5, (434) 0.3, (435) 3.5, (436) 3.6, (437) 0.8, (476) 0.5, (477) 0.6, (479) 0.5, (480) 2.4	(885) NYCNYC01 (<i>Plasmodium</i> sp.)	KU057967
						(854) NYCNYC01 (<i>Plasmodium</i> sp.)	KU057967

^a Migratory species^b Lineage previously reported^c Parasite morphospecies previously reported^d Parasite not detected by microscopic examination^e Parasite seen in blood films, but not detected by PCR

Fig. 2 *Haemoproteus columbae* gametocytes from the blood of *Columba livia* (rock pigeon) in São Paulo Zoo. Bird ID 425 (lineage COLIV03): macrogametocytes, **a–b** and microgametocytes, **c–d**; Bird ID 430 (lineage COQUI05): macrogametocytes, **e–f** and microgametocytes, **g–h**; Bird ID 478 (lineage COQUI05): macrogametocytes, **i–j** and microgametocytes, **k–l**; Bird ID 481 (this parasite was readily seen in blood films, but was not detected by PCR): macrogametocytes, **m–n** and microgametocytes, **o–p**. Note large aggregations of volutin and pigment granules readily visible in microgametocytes of all lineages of this parasite. Giemsa-stained thin blood films. Bar = 10 μ m



Haemoproteus iwa. The other included species belonging to the subgenus *Haemoproteus* (*Parahaemoproteus*). All *Haemoproteus* sequences obtained from doves in this study clustered with subgenus *Haemoproteus* (*Haemoproteus*). The majority of them appeared closely related with *H. columbae* lineage HAECOL1, while the sequence obtained in *C. livia* (ID 481) clustered with *H. multipigmentatus* and *H. iwa*. No *Parahaemoproteus* parasite lineages were reported in this study.

The phylogenetic tree of *Plasmodium* spp. (Fig. 4) showed low phylogenetic resolution, and morphospecies belonging to different subgenera (*Haemamoeba*, *Giovannolaia*, *Novyella*, *Bennettinia*, and *Huffia*) were included in more than one clade. However, the sequences recovered from *P. superciliares* (ID 885) and *N. nycticorax* (ID 854) grouped with a well-supported cluster with the majority of *P. (Haemamoeba)* sequences (94 % of posterior probability); the sequence recovered from *D. viduata* (ID 263) (DENVID01) grouped with a well-supported cluster containing exclusively *P. (Novyella)* sequences (96 % of posterior probability); and the sequence recovered from *D. viduata* (ID 249)

(DENVID02) grouped with the only *P. (Huffia)* sp. sequence available in the tree, with a high posterior probability (100 %).

Discussion

This study adds to the better understanding of the occurrence and genetic diversity of *Plasmodium* and *Haemoproteus* parasites in different groups of birds living freely in the São Paulo Zoo. The majority of examined birds belonged to the Anatidae species, among which two migrant species were investigated and one was found positive. This was *Dendrocygna viduata* (white-faced whistling duck), a species found in Africa, including Madagascar and Comores Islands, Central America Islands, and South America, including Brazil. *Dendrocygna javanica*, *D. viduata*, and *D. autumnalis* had been previously observed infected with *Plasmodium* sp. (Beadell et al. 2004; Valkiunas 2005). In this study, we found two white-faced whistling ducks positive for *Plasmodium* lineages, with two different new lineages

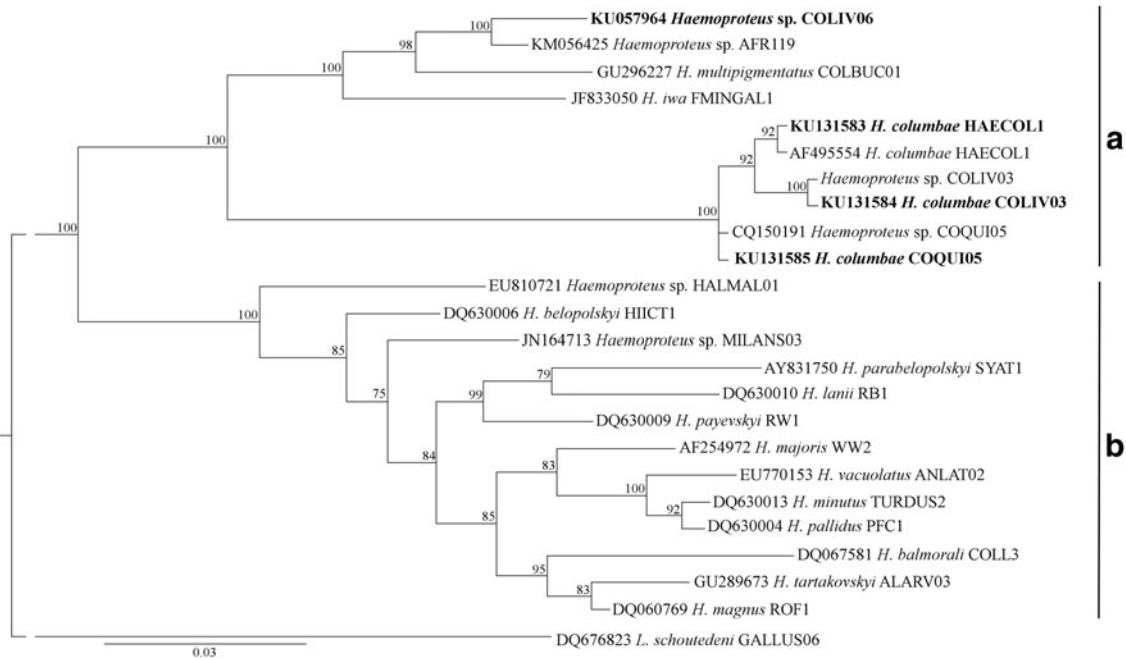


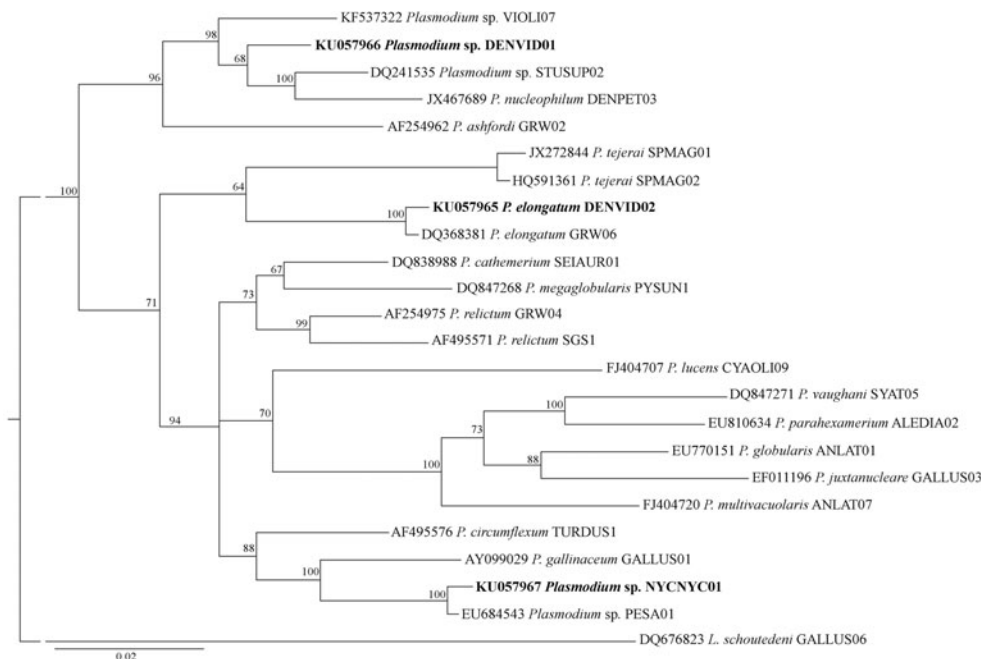
Fig. 3 Bayesian phylogeny of cytochrome *b* gene lineages of *Haemoproteus* species of avian hemosporidian parasites. Lineages recorded in this work are given **bolded**. Names of the lineages are given after the species names of parasites. GenBank accession numbers of the

lineages are provided before the parasite species names. Nodal support values (in percentage) indicate posterior clade probabilities. **a** Species of the subgenus *Haemoproteus* (*Haemoproteus*). **b** Species belonging to the subgenus *Haemoproteus* (*Parahaemoproteus*)

(DENVID01 and DENVID02). The lineage DENVID01 is similar (98 % of identity) to two different lineages (STUSUP02 and VIOLI07), which, in turn, have 97 % of identity with each other. The lineage DENVID02 shows high identity (99 % of similarity due to a C-T nucleotide substitution at position 186) to a sequence that has already been characterized as *P. elongatum* (GRW06), the cosmopolitan and

virulent parasite in many bird species (Garnham 1966; Valkiūnas et al. 2008). The lineage DENVID01 has been reported before in other birds (mainly belonging to the Anatidae) in the São Paulo Zoo (K. Kirchgatter, unpublished observation). Inversely, the lineage DENVID02 has not been found in other infected bird species in the Zoo (K. Kirchgatter, unpublished observations). This lineage was confirmed in

Fig. 4 Bayesian phylogeny of cytochrome *b* gene lineages of *Plasmodium* species of avian hemosporidian parasites. Lineages recorded in this work are given **bolded**. Names of the lineages are given after the species names of parasites. GenBank accession numbers of the lineages are provided before the parasite species names. Nodal support values (in percentage) indicate posterior clade probabilities



different assays of PCR and sequencing, indicating a possibility that this bird came to the Zoo being already infected. In response to seasonal fluctuations in food and water availability, the white-faced whistling duck may undertake short migrations (usually less than 500 km) in search of favorable foraging grounds (Madge and Burn 1988; del Hoyo et al. 1992). Their migration is conditioned to the availability of water bodies (Carboneras and Kirwan 2014) and they may use even the polluted ones (Sick 1997). Considering the wide extension of Brazil, the migratory distance of 500 km by *D. viduata* populations allows us to determine the origin of São Paulo State's populations only to this country, and probably ranging from South to Southeast, as these birds usually look for sites with warm temperatures. *Plasmodium elongatum* (here the lineage DENVID02) has a wide range of distribution, and it is of broad specificity having been reported in birds belonging to ten different orders (see MalAvi and Garnham 1966; Valkiūnas 2005), including magellanic penguins from southern Brazil (Vanstreels et al. 2014a).

Birds of the Columbidae family were the second most largely sampled group in this study, with 20 individual rock doves (*Columba livia*) tested. This bird was introduced to Brazil in XVI century from Europe, and it became well adapted to the urban environment due to its ecological plasticity in the ability to inhabit different environments (Sigrist 2014), particularly those located close to human settlements (Sick 1997; Develey and Endrigo 2011). Despite some studies showing its ability to migrate over hundreds of kilometers, the native populations have been considered sedentary (Baptista et al. 1997). In São Paulo Zoo, the birds usually find food and shelter easily, and that contributes to the spread of the populations. Four different lineages of hemosporidians were found in *C. livia* from São Paulo Zoo: (1) the lineage HAECOL1 has been formerly found in *C. livia* in several countries: Botswana, Nigeria, Colombia, and Italy (Waldenstrom et al. 2002; Karamba et al. 2012; Gonzalez et al. 2015; Scaglione et al. 2015); (2) COQUI05 has been found in *Coquillettidia* (Culicidae, Diptera) mosquitoes from Cameroon (Njabo et al. 2009); (3) COLIV03 has already been found in *C. livia* from Nigeria (Karamba et al. 2012); (4) COLIV06 is a new sequence, and possibly it belongs to a *Haemoproteus* species since it showed 10 % of genetic distance with other sequences from HAECOL1 group. However, the bird with COLIV06 was probably co-infected whereas parasites that could present this sequence were not visible during microscopic examination of blood films. Absence of gametocytes in birds PCR positive for hemosporidian parasites has been described and can be explained by light gametocyte parasitemia, amplification of DNA of circulating sporozoites (Valkiūnas et al. 2009), or even the presence of remnants of parasites, which abort development during tissue stage of development (Ferrell et al. 2007; Donovan et al. 2008; Olias et al. 2011; Chagas et al. 2013). COQUI05 and COLIV03 sequences showed 1–

2 % of genetic distances with HAECOL1 group sequences and were confirmed by microscopy in this study as *H. columbae* (Fig. 2), a parasite widespread in at least 14 species of the Columbiformes (Valkiūnas 2005).

Haemoproteus spp. is the most common blood parasite in birds, followed by *Leucocytozoon* spp. and *Plasmodium* spp. (Garnham 1966; Valkiūnas 2005). However, in South America, *Haemoproteus* lineage diversity does not follow the high diversity found among avian and mosquito species (Clark et al. 2014). It is possible that a bias to preferential amplification of *Plasmodium cytb* sequences exists (Valkiūnas et al. 2006). During this study, we found 83 and 17 % of the positive free-living birds infected with *Haemoproteus* spp. and *Plasmodium* spp., respectively, while *Leucocytozoon* spp. has not been observed in the lowlands of Brazil (Valkiūnas 2005). The high prevalence of *Haemoproteus* spp. reported in this study is somewhat striking since only 2 % of the positive captive birds of the same region have been found infected with this parasite (K. Kirchgatter, unpublished observation). However, this difference can be attributed to a bias due to high rate of infection found in rock doves. High prevalence (100 %) of *Haemoproteus* infection is consistent with that found in a study, which examined rock doves in the State of São Paulo (Adriano and Cordeiro 2001).

Columba livia, in its natural habitat, plays an important ecological role, controlling the population of insects and spreading some seeds. As invasive species in the study area and big cities such São Paulo, it is known as a sinantropic animal, competing with local species. However, the parasites that rock doves harbor can be transmitted to other columbiform birds, compromising their fitness. *C. livia* has a status of *least concern* (IUCN RED LIST 2015), but in São Paulo county, where PEFI locates, not only rock doves are present but also other species of Columbiformes, such as *Columbina talpacoti*, *Patagioenas picazuro*, *Patagioenas cayennensis*, *Patagioenas plumbea*, *Patagioenas speciose*, *Zenaida auriculata*, *Geotrygon montana*, *Geotrygon violacea*, *Leptotila verreauxi*, and *L. rufaxilla* (http://www.prefeitura.sp.gov.br/cidade/secretarias/upload/meio_ambiente/arquivos/TRE_EIA.pdf). Although *Haemoproteus* sp. usually does not cause clinical disease in adapted avian hosts (Valkiūnas 2005), infections with this parasite causing sudden death in non-adapted passerine birds have been reported in San Diego Zoo, USA (Donovan et al. 2008). *Haemoproteus* (*Parahaemoproteus*) parasites cause severe disease in some bird species in Europe and Australia (Olias et al. 2011; Cannell et al. 2013) and can be lethal for blood-sucking insects (Valkiūnas et al. 2014). Because rock doves have expanded their range into environmental conservation areas, there is a greater risk of spreading its infections to the native fauna; therefore, population control activities with these birds can be required.

We also found infected *Penelope superciliaris* and *Nycticorax nycticorax*. The former bird is the most well-known species of the Cracidae family, which includes 50 species and 20 of them are present in Brazil (Del Hoyo and Kirwan 1994; CBRO 2014). *Penelope superciliaris* is found from north to south of Brazil, Paraguay, and north of Argentina (Guix 1997). This species eats basically insects, flowers, and fallen fruits, which makes it an important seed disperser in the forest ecosystems (Del Hoyo and Kirwan 1994; Mikich 2002; Sick 1997). *Nycticorax nycticorax* (Ardeidae) has nightly and twilight habits (CBRO 2014) and can be found living close to water from Canada to south Argentina, including Old World where it has migration habits (Martinez-Vilalta et al. 2014). Its diet consists of fish, crustaceans, amphibians, mollusks, crabs, aquatic insects, polychaetes, small lizards, snakes, rodents, bats, eggs and chicks, and occasionally carcass (Sick 1997). In São Paulo Zoo, it is an opportunistic bird that does not belong to the park, but lives in the neighborhood and feeds using remnants of food of captive animals. Thus, living freely in the area of the zoo, such birds can rapidly take advantage of favorable conditions when they arise. In a report of *Plasmodium* sp. causing mortality in penguins from São Paulo Zoo, we detected a mosquito engorged with *N. nycticorax* blood (Bueno et al. 2010). As these birds were frequently found inside the penguin enclosures feeding on the leftover of fish, they might be a source of *Plasmodium* infections for other bird species in São Paulo Zoo. According to MalAvi database, the lineage found in *N. nycticorax* (PESA01) has been found in three different hosts of three different orders: *Phaeomyias murina*, a passeriform species from Brazil (Lacorte et al. 2013); *Calidris melanotos*, a migratory wader from Alaska (Yohannes et al. 2009); and *Leptotila verreauxi*, a Columbidae species from Uruguay (Durrant et al. 2006), showing a generalist parasite that may infect a broad range of avian hosts. However, it is impossible to proceed species identification using only the sequence information since a small genetic difference (96 % or less of identity) was reported in readily distinguishable hemosporidian morphospecies, all of *Haemamoeba* subgenus. Additional morphologic studies are needed to determine *Plasmodium* species identity of the PESA01 lineage.

We also tested owls that live freely in the Zoo. Although none was found infected in our study, hemosporidian parasites, mainly *Haemoproteus* parasites, are widespread in owls worldwide and have been found in the USA (Tavernier et al. 2005; Ishak et al. 2008), Europe (Krone et al. 2008; Bukauskaitė et al. 2015), Africa (Ishak et al. 2008), Asia (McClure et al. 1978), and South America (White et al. 1978). The prevalence of infections exceeds 50 % in many owl populations worldwide (Valkiūnas 2005). Owl haemoproteids belong to the subgenus *Parahaemoproteus* and are transmitted by biting midges of the genus *Culicoides*

(Bukauskaitė et al. 2015). In São Paulo State, one individual of *Megascops choliba* sampled in 2009 at Carapicuíba was found positive for *Haemoproteus syrni* (Vanstreels et al. 2014b). The absence of positives among the owl samples in our study could reflect the low number of birds tested here (nine samples of three birds). However, all the 35 samples tested from captive birds of Strigiformes order from the same area were negative by PCR (K. Kirchgatter, unpublished observation). This result might indicate weak transmission of *Parahaemoproteus* parasites due to the absence of susceptible species of biting midges at our study site warranting further investigation.

The cost of the infections to the birds depends on many factors, including the age, host immunity, breeding season, migration, and food availability (Sorci and Cornet 2010). In captivity, the animals do not usually need to compete for food, territory, or females, decreasing these stress levels that would affect animals under natural conditions. Wild animals sampled in this study have access to food from the captive animals. Since parasites can influence the biodiversity and the structure of host populations by altering the competitive interaction between the free-living and the captive birds, biodiversity preservation studies should be considered in areas with these two animal communities.

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