A new lizard malaria parasite *Plasmodium intabazwe* n. sp. (Apicomplexa: Haemospororida: Plasmodiidae) in the Afromontane *Pseudocordylus melanotus* (Sauria: Cordylidae) with a review of African saurian malaria parasites

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**Abstract**

**Background:** Saurian malaria parasites are diverse apicomplexan blood parasites including the family Plasmodiidae Mesnil, 1903, and have been studied since the early 1900s. Currently, at least 27 species of *Plasmodium* are recorded in African lizards, and to date only two species, *P. zonuriae* (Pienaar, 1962) and *P. cordyli* Telford, 1987, have been reported from the African endemic family Cordylidae. This paper presents a description of a new malaria parasite in a cordylid lizard and provides a phylogenetic hypothesis for saurian *Plasmodium* species from South Africa. Furthermore, it provides a tabular review of the *Plasmodium* species that to date have been formally described infecting species of African lizards.

**Methods:** Blood samples were collected from 77 specimens of *P. melanotus* (A. Smith, 1838) from Platberg reserve in the Eastern Free State, and two specimens of *Cordylus vittifer* (Reichenow, 1887) from the Roodewalshoek conservancy in Mpumalanga (South Africa). Blood smears were Giemsa-stained, screened for haematozoa, specifically saurian malaria parasites, parasite stages were photographed and measured. A small volume was also preserved for TEM studies. *Plasmodium* and *Haemoproteus* primer sets, with a nested-polymerase chain reaction (PCR) protocol, were employed to target a fragment of the cytochrome-β (*cyt-b*) gene region. Resulting sequences of the saurian *Plasmodium* species’ isolates were compared with each other and to other known *Plasmodium* spp. sequences in the GenBank database.

**Results:** The presence of *P. zonuriae* in both specimens of the type lizard host *C. vittifer* was confirmed using morphological characteristics, which subsequently allowed for the species’ molecular characterisation. Of the 77 *P. melanotus*, 44 were parasitised by a *Plasmodium* species, which when compared morphologically to other African saurian *Plasmodium* spp. and molecularly to *P. zonuriae*, supported its description as a new species *Plasmodium intabazwe* n. sp. (Continued on next page)
Background
Saurian malaria parasites are diverse apicomplexan blood parasites including the family Plasmodiidae Mesnil, 1903, and have been reported and described from a range of vertebrates since the early 1900s, see [1]. Reptilian malaria parasites have in the past been represented by four genera: Plasmodium Marchiafava & Celli, 1885; Haemoproteus Kruse, 1890; Saurocytozoon Lainson & Shaw, 1969; and Haemocystidium Castellani & Willey, 1904. However, species of Haemoproteus infecting reptiles (lizard, snake and chelonian) have recently been proposed to belong, on both a morphological and molecular basis, to the resurrected genus Haemocystidium during the revision of the haemoproteid genera by Pineda-Catalan et al. [2]. These genera, as well as the subgenera Sauramoeba Garnham, 1966; Carinamoeba Garnham, 1966; Lacertamoeba Telford, 1988; Paraplasmidium Telford, 1988; Asiamoeba Telford, 1988; Garnia Lainson, Landau & Shaw, 1971; and Ophidiella Garnham, 1966, were in the past solely differentiated by the morphology of the erythrocytic stages and development in the tissues of the vertebrate host, as well as by the development in the haematophagous vector [1]. The first saurian Plasmodium species, Plasmodium agamae Wenyon, 1909 and Plasmodium mabuiae Wenyon, 1909, were described by Wenyon (1909) from an agamid lizard (Agama agama) and from a scincid lizard (Trachylepis quinquetaeniata) in Africa, respectively. On a global scale, Plasmodium species descriptions remained rare until the 1960s when research on these organisms greatly increased [3, 4]. Today, at least 27 species of Plasmodium are recorded from African lizards, but only two species have been described from African cordylid lizards (Table 1).

The first saurian malaria parasite from South Africa, Plasmodium zonuriae (Pienaar, 1962), was described from a cordylid lizard (Cordylus vittifer) collected at Elandsfontein (North-West province) [5]. Following the original description by Pienaar [5], an additional host, Pseudocordylus microlepidotus (Cuvier, 1829), collected in 1972, and locality, the vicinity of Cape Town (Western-Cape province), was recorded for this malaria parasite [6]. Later, in 1986, specimens of the type-host (C. vittifer) from roughly the same region as the type-locality as well as further north within South Africa (Pretoria region), were confirmed positive for P. zonuriae, thereby again increasing the distribution range of this parasite [6]. In his review on haemoparasites of the reptilia, Telford [6] reported a second species of Plasmodium, collected from South African C. vittifer, Plasmodium cordyli Telford, 1987; a parasite which he had originally described from Cordylus tropidosternum in Tanzania [6, 7]. Thus, by 2009, two species of Plasmodium had been described and reported parasitising cordylid lizards of South Africa, namely P. zonuriae from both C. vittifer and P. microlepidotus and P. cordyli from C. vittifer [6].

The majority of malaria parasite species descriptions in African lizards were based on the morphology of the intraerythrocytic parasites, and new species were often named based on new host infection records. This is equally true for other apicomplexan blood parasites such as the haemogregarines. However, recently there has been a concerted effort to describe known and new species of haemogregarines from African lizards using both morphological and molecular methods (see [8]). In this study we intended to do the same for three species of African saurian malaria parasites and as such the aim of this study was to morphologically confirm and molecularly characterise the two Plasmodium species, P. zonuriae and P. cordyli, reporting infected South African cordylids, review all existent saurian malaria parasites based on their morphometrics and describe, name and molecularly diagnose an unknown species of Plasmodium parasitising the South African Aframontane cordylid species Pseudocordylus melanotus.

Methods
Lizard collection, blood smear preparation and screening
Specimens of Pseudocordylus melanotus (n = 77) were collected by hand during the summer months (September to April), over a period of 5 years (2008–2013), at Platberg Reserve (28°14′36.71″S, 29°09′45.45″E) in the Eastern Free State Province, South Africa. Additionally, specimens of Cordylus vittifer (n = 2) were collected by hand in the Roodewalshoek Conservancy (25°00′51.20″S, 30°19′23.17″E) (1441 m), Mpumalanga, South Africa.

Blood was taken with an insulin syringe from the femoral arteries or veins and lizards were immediately released thereafter at the site of capture. Thin blood smears were prepared, air-dried, fixed and stained using Giemsa-
| Lizard host species by family | Plasmodid species | Original host localities | Intracellular meront dimensions (L × W in μm); [LW in μm²]; Number of merozoites | Intracellular gametocyte dimensions (L × W in μm); [LW in μm²] | Gametocyte morphology | Effects of gametocytes on host cell | Reference |
|-------------------------------|-------------------|-------------------------|----------------------------------------|----------------------------------------|------------------------|-----------------------------|----------|
| **Agamidae**                 |                   |                         |                                        |                                        |                        |                             |          |
| *Agama agama* (syn. *Agama* | Plasmodium (Sauramoeba) | Gbanga, Liberia (9–18 × 4–11) | (9–22 × 4–10) | Round to elongate or bulky | [3, 27–29] |
| *Agama cyanogaster* (Southgate, 1970); | giganteum Theiler, 1930 | [52–165] 28–74 | [45–145] | – | |
| *Agama mossambica* (Peters, 1854) | Plasmodium (Lacertamoeba) | Bahr El Ghazal, Sudan (4–11 × 3–6) | (6–19 × 3–8) | Hypotrophy | [3, S, 27–32] |
| *Agama agama* other hosts: Acanthocercus atricollis (Smith, 1849); | agamae (Wennyon, 1909) | Province, Sudan (12–55) 4–15 | [33–105] | Dislocation and occasional distortion of nuclei | |
| *Agama mossambica* | Plasmodium (Lacertamoeba) | Morogoro Region, Tanzania (5–15 × 3–7) | (6–17 × 3–8) | Elongate | [6] |
| *Agama mossambica* | giganteum Theiler, 2009 | [20–75] 6–34 | [36–84] | – | |
| **Chamaeleonidae**           |                   |                         |                                        |                                        |                        |                             |          |
| *Chamaeleo breviceps* | Plasmodium (Sauramoeba) | Moramanga, Subprefecture, Madagascar (11–23 × 7–11) | (9–20 × 5–13) | Oval to elongate or bulky | [33] |
| Günther, 1879; | robinsoni (Brygoo, 1962) | [90–184] 40–74 | [72–221] | – | |
| other host: Calumma parsoni crucifer (Cuvier, 1824) | Plasmodium (Lacertamoeba) | Périnet, Madagascar (6–9 × 5–8) | (9–15 × 5–10) | Hypotrophy | [33] |
| *Chamaeleo bivittatus* | giganteum Theiler, 1930 | [36–64] 10–16 | [66–126] | Dislocation and occasional distortion of nuclei | |
| *Kinyongia fischeri* (Reichenow, 1887) | Plasmodium (Sauramoeba) | Tanga Region, Tanzania | – | – | [34] |
| *Kinyongia fischeri* | giganteum Theiler, 1930 | Tanga Region, Tanzania (9 × 6) | (8–11 × 5–8) | Oblong to elongate | [35] |
| *Kinyongia oxyrhina* (Naver & Böhmke, 1988) | Plasmodium (Sauramoeba) | Kilombero district, Tanzania (6–15 × 4–8) | (6–14 × 4–8) | Hypotrophy | [36] |
| *Kinyongia oxyrhina* | giganteum Theiler, 1930 | Morogoro Region, Tanzania (5–7 × 4–6) | (5–11 × 4–6) | Dislocation of nuclei | |

**Table 1**: African saurian malaria species across the families Agamidae, Chamaeleonidae, Cordylidae, Gekkonidae, Lacertidae, Opluridae and Scincidae.
### Table 1: African saurian malaria species across the families Agamidae, Chamaeleonidae, Cordylidae, Gekkonidae, Lacertidae, Opluridae and Scincidae (Continued)

| Lizard host species by family | Plasmodium species | Original host localities | Intracellular meront dimensions (L x W in μm); [LW in μm²]; Number of merozoites | Intracellular gametocyte dimensions (L x W in μm); [LW in μm²] | Gametocyte morphology | Effects of gametocytes on host cell | Reference |
|-------------------------------|-------------------|-------------------------|-----------------------------------------------------------------------------------|----------------------------------------------------------|---------------------|----------------------------------|-----------|
| **Trioceros werneri**         | *Plasmodium*      |                         |                                                                                   |                                                          |                     |                                  |           |
| (Torner, 1899)                | *Lacertamoeba*    |                         |                                                                                   |                                                          |                     |                                  |           |
| **Trioceros werneri**         | *Plasmodium*      |                         |                                                                                   |                                                          |                     |                                  |           |
| (Tornier, 1899)               | *arachniformis*   |                         |                                                                                   |                                                          |                     |                                  |           |
| **Trioceros werneri**         | *Plasmodium*      |                         |                                                                                   |                                                          |                     |                                  |           |
| (Reichenow, 1887)             | *Lacertamoeba*    |                         |                                                                                   |                                                          |                     |                                  |           |
| **Cordylus t. tropidosternum**| *Plasmodium*      |                         |                                                                                   |                                                          |                     |                                  |           |
| (Cope, 1869)                  | *Carinamoeba*     |                         |                                                                                   |                                                          |                     |                                  |           |
| **Cordylus vittifer**         | *Plasmodium*      |                         |                                                                                   |                                                          |                     |                                  |           |
| (Reichenow, 1887)             | *Lacertamoeba*    |                         |                                                                                   |                                                          |                     |                                  |           |
| **Cordylus vittifer**         | *Plasmodium*      |                         |                                                                                   |                                                          |                     |                                  |           |
| *(Pienaar, 1962)*             | *Lacertamoeba*    |                         |                                                                                   |                                                          |                     |                                  |           |
| **Pseudocordylus melanotus**  | *Plasmodium*      |                         |                                                                                   |                                                          |                     |                                  |           |
| *(A. Smith, 1838)*            | *Carinamoeba*     |                         |                                                                                   |                                                          |                     |                                  |           |
| **Cnemaspis barbouri Perret, 1986** | *Plasmodium* | Morogoro Region, Tanzania | (6–13 × 3–7) [24–91] | (7–14 × 3–9) [32–108] | Elongate (active) | Ovoid or rounded (chronic) | [25] |
| **Cnemaspis cremaspi**         | *Plasmodium*      | Morogoro Region, Tanzania | (6–13 × 3–7) [24–91] | (7–14 × 3–9) [32–108] | Elongate (active) | Ovoid or rounded (chronic) | [25] |
| **Hemidactylus platyccephalus** | *Plasmodium*     | Morogoro Region, Tanzania | (4–10 × 2–6) [12–54] | (5–10 × 4–7) [20–63] | Ovoid          | Hypertrophy                  | [25] |
| Peters, 1854                  | *Lacertamoeba*    | Morogoro Region, Tanzania | (4–10 × 2–6) [12–54] | (5–10 × 4–7) [20–63] | Ovoid          | Hypertrophy                  | [25] |
| **Lygodactylus capensis grateri** | *Haemocystidium lygodactyi* | Morogoro region, Tanzania | (5–10 × 9–16) [20–70] | (8–25 × 5–11) | Elongate to oval | Distortion of host cell | [37] |
| Sternfeld, 1911               | *lygodactyi*      | Morogoro region, Tanzania | (5–10 × 9–16) [20–70] | (8–25 × 5–11) | Elongate to oval | Distortion of host cell | [37] |
| **Lygodactylus capensis grateri** | *Haemocystidium lygodactyi* | Morogoro region, Tanzania | (5–10 × 9–16) [20–70] | (8–25 × 5–11) | Elongate to oval | Distortion of host cell | [37] |
Table 1  African saurian malaria species across the families Agamidae, Chamaeleonidae, Cordylidae, Gekkonidae, Lacertidae, Opluridae and Scincidae  (Continued)

| Lizard host species by family | Plasmodiid species | Original host localities | Intracellular meront dimensions (L × W in μm); [LW in μm²]; Number of merozoites | Intracellular gametocyte dimensions (L × W in μm); [LW in μm²] | Gametocyte morphology | Effects of gametocytes on host cell | Reference |
|-------------------------------|--------------------|--------------------------|-----------------------------------------------------------------|-----------------------------------------------------------------|----------------------|-----------------------------------|-----------|
| *Lygodactylus l. luteopicturatus* Pasteur, 1964 Other host: *Lygodactylus capensis grotei* | *Plasmodium* (Lacertamoeba) lovendi Telford, 1984 | Morogoro Region, Tanzania | (5–15 × 3–7) [20–91] 6–26 | (8–23 × 3–11) [48–176] | Elongate, rarely rounded | Hypotrophy Distortion of host cell Displacement of nuclei | [25] |
| *Tarentola mauritanica deserti* (Linnaeus, 1758) Other host: *Tarentola annularis* (Geoffroy De St-Hilaire, 1827) | *Haemocystidium tarentolae* (Parrot, 1927) Paperna & Landau, 1991 | El Kantara, Algeria | – | (8–18 × 4–12) | Elongate | Slight hypertrophy and distortion of host cell Lateral displacement of nuclei | [38, 39] |
| *Holaspis guentheri* Gray, 1863 | *Plasmodium* (Lacertamoeba) holaspi Telford, 1986 | Morogoro Region, Tanzania | (5–13 × 4–7) [25–66] 8–18 | (6–18 × 3–8) [28–98] | Elongate | Distortion of host cell Displacement and distortion of nuclei | [40] |
| *Oplurus cuvieri* Gray, 1831 | *Haemocystidium opluri* Paperna & Landau, 1991 | Baie de Loukaio, Madagascar | – | (12–19 × 3–12) | Oblong, oval | Lateral hypertrophy Displacement of nuclei | [39] |
| *Trachylepis maculilabris* (Gray, 1845) | *Plasmodium* (Lacertamoeba) maculilabre Schwetz, 1931 | Kisangani, Congo | (10.0 × 6.9) [69] 15–20 | (7–13 × 5–8) [42–91] | Ovoid to elongate | Hypertrophy Distortion of host cell Displacement and occasional distortion of nuclei | [41] |
| *Trachylepis quinquetaeniata* (Lichtenstein, 1823) Other hosts: *Trachylepis maculilabris, Trachylepis striata* (Peters, 1844) | *Plasmodium* (Carinamoeba) mabuiae (Wenyon, 1909), Telford, 1983 | Bahr-El-Ghazal Province, Sudan | (4–9 × 2–5) [10–30] 4–12 | (5–11 × 3–5) [18–44] | Elongate, rarely ovoid or round | Hypertrophy Occasional distortion of host cell Displacement and occasional distortion of nuclei | [31, 42] |
| *Trachylepis striata* | *Plasmodium* (Sauramoeba) heischi Gamham & Telford, 1984 | Nairobi, Kenya | (8–18 × 6–11) [48–144] 20–65 | (8–12 × 4–9) [60–120] | Large, spindle-shaped | Distortion of host cell Lateral displacement of nuclei | [43] |
| Other hosts: *Trachylepis maculilabris, Trachylepis quinquetaeniata, Trachylepis varia* (Peters, 1867) | *Plasmodium* (Lacertamoeba) pitmani Hoare, 1932 | Lake Victoria, Uganda | (4–11 × 3–7) [12–66] 4–25 | (5–16 × 4–9) [25–91] 2.07 | Ovoid | Distortion of nuclei | [44, 45] |

Abbreviations: L, length; W, width; LW = L × W; – Indicates dimensions not recorded
stain (FLUKA, Sigma-Aldrich, Steinheim, Germany). Subsequently, smears were screened with the aid of a 100× oil immersion objective, images captured and parasites measured as described previously [9–12]. Morphometric measurements [length (L) and width (W) in micrometres] of the developmental stages in the lizard erythrocytes were taken using Deltapix (Nikon, The Netherlands) software. Following the method of Telford [6], the area or size was calculated as (length × maximum width) (LW) (in μm²) for all parasite stages examined, and the relative size of meront and gametocyte stages to the host cell nucleus (LW/HNLW) and to the nuclei of uninfected host cells (LW/NNLW) determined. A few drops of blood were fixed in 2.5% glutaraldehyde for TEM, and the remaining blood, that was not used in blood smear preparation, was placed in sterile PCR microtubes (Eppendorf, Germany) with an equal volume of 70% ethanol to be processed molecularly at a future date.

Transmission electron microscopy (TEM)

Four to five drops of fresh blood were fixed in 2.5% glutaraldehyde (0.2 M, pH 7.2) for 1 h at 4 °C. Blood was pelleted by centrifugation (10,000 rpm) and rinsed in 0.2 M Sorensen’s phosphate buffer and post fixed in 2% solution of osmium tetroxide in the same buffer. Samples were then dehydrated in a graded ethanol series, dried over a 4 molecular sieve and embedded in Agar 100 medium (Agar Scientific, Ltd., Stansted, UK). Thin sections, cut with a glass knife on an Ultracut III Ultra-microtome (Leica, Wetzlar, Germany) were collected on copper, 300 hexagonal-mesh grids and stained for 20 min in 10% uranyl acetate in Analar grade methanol, washed with Analar methanol and allowed to dry. Sections were then stained for 20 min in Reynolds’ lead citrate solution, washed with 0.02 M sodium hydroxide solution followed by distilled water, before examination with a Philips CM100 transmission electron microscope (Philips Electron Optics, Eindhoven, The Netherlands) operated at 80–100 kV. Digital images were captured with a MegaView II side mounted digital Olympus camera with accompanying iTEM software.

DNA extraction and sequence analysis

Ethanol-preserved blood from the nine parasitised individuals of the 77 collected of *P. melanotus* and both *C. vittifer* were used for molecular work. Genomic DNA of *Plasmodium* spp. was extracted from the samples following the standard protocol for the Kapa Express Extract kit (Kapa Biosystems, Cape Town, South Africa). Amplification of *Plasmodium* spp. DNA was initially completed using primer sets, with *Leucocytozoon* spp. DNA was initially completed using and both Thermal Cycler PCR and Leucocytozoon. Comparative sequences of *Plasmodium* species were downloaded from GenBank and aligned to the sequences generated within this study. Two species of *Leucocytozoon*, *Leucocytozoon gentili* and *Leucocytozoon majoris* (GenBank: DQ451435, DQ451439) were used as the outgroup, as species of *Leucocytozoon* were shown to be a sufficient outgroup of the focal taxa by [2]. All phylogenetic analyses were further undertaken in the bioinformatics software program Geneious Ver. 7.1. Sequences were identified using the Basic Local Alignment Search Tool (BLAST) [14], and deposited in the NCBI GenBank database under the accession numbers KX121601–KX121609. Comparative sequences of *Plasmodium* species were downloaded from GenBank and aligned to the sequences generated within this study. Two species of *Leucocytozoon*, *Leucocytozoon gentili* and *Leucocytozoon majoris* (GenBank: DQ451435, DQ451439) were used as the outgroup, as species of *Leucocytozoon* were shown to be a sufficient outgroup of the focal taxa by [2]. All phylogenetic analyses were further undertaken in the bioinformatics software program Geneious Ver. 7.1. Sequences were aligned using the MUSCLE alignment tool [15]. To infer phylogenetic relationships a Bayesian inference (BI) method was used. A comprehensive model test was preformed to determine the most suitable nucleotide substitution model, according
to the Akaike information criterion using jModelTest 2.1.7 [16, 17]. The best-fit model selected was the General Time Reversible with estimates of invariable sites and a discrete Gamma distribution (GTR + I + Γ). The dataset comprised 31 cytochrome b (cyt-b) mitochondrial sequences, with an alignment length of 497 nt. The BI analysis was implemented from within Geneious 7.1 using MrBayes 3.2.2 [18]. The Markov Chain Monte Carlo (MCMC) algorithm was run for 10 million generations. The Markov chain was sampled every 100 cycles, and the MCMC variant contained 4 chains with a temperature of 0.2. The log-likelihood values of the sample point were plotted against the generation time and the first 25 % of the trees were discarded as ‘burn-in’ with no ‘burn-in’ samples being retained. Results were visualised in Trace (implemented from within Geneious) to assess convergence and the ‘burn-in’ period.

Results

General observations

Parasites of what appeared to be a Plasmodium species were discovered in the peripheral blood of 44/77 (prevalence 57 %) specimens of *P. melanotus* captured on various rocky outcrops on the summit of Platberg Reserve in the Eastern Free State. The peripheral blood of some lizards occasionally revealed mixed infections, comprising most likely Hepatozoon species, filarial nematode species and so-called Sauroplasma infections (unpublished data).

Co-infections between the Plasmodium species and Hepatozoon species (of which there appeared to be possibly two species) were found in 27/77 (35 %) of the total lizards sampled. One of these Hepatozoon species was recently described as *Hepatozoon affluomaloti* Van As, Davies & Smit 2015, occurring in 14/77 (18 %) of the lizards [19].

*Plasmodium zonuriae* (Pienaar, 1962) infections were confirmed, based on morphological characteristics (see Table 1, Fig. 1 and Discussion), in both *C. vittifer* specimens (100 % prevalence) from the Roodewalshoek Conservancy, Mpumalanga. These infections from both lizard hosts were present in the peripheral blood erythrocytes (Figs. 1 and 2). No Plasmodium species, morphologically comparable to *P. cordyli*, were observed in either of the *C. vittifer* specimens or in any of the *P. melanotus* specimens.

*Plasmodium zonuriae* (Pienaar, 1962) Telford, 1987

**Type-host:** Cordylus vittifer (Reichenow, 1887) (Sauria: Cordylidae).

**Vector:** Unknown.

**Type-locality:** Elandsfontein (North-West province) [5].

**Other localities:** Pretoria (Gauteng) and Middelburg (Mpumalanga) [6]; Roodewalshoek (Mpumalanga) (present study).

**Voucher material:** Voucher material, 2× blood smears from Cordylus vittifer NMB P 414 and C. vittifer NMB P 415, respectively, deposited in the Protozoan Collection of the National Museum, Bloemfontein (NMB), South Africa.

**Representative DNA sequences:** The cyt-b gene sequences have been submitted in the GenBank database under the accession numbers KX121608 and KX121609.

Redescription

**Trophozoite.** Amoeboid forms (Fig. 1a) with abundant pseudopodia, measuring 1.8–2.8 × 1.6–3.1 (2 ±
0.9 × 2.7 ± 1.4) μm (n = 55), containing discrete brown or black pigment granules. Oval to pyriform forms with purple-stained periphery and pinkish central vacuole (Fig. 1e arrowhead) measuring 1.1–1.8 × 1.0–1.7 (1.4 ± 0.3 × 1.2 ± 0.35) μm (n = 60); LW: 3.9–21.8 (8.4 ± 5) μm² (n = 60). Mature trophozoites elongated (Fig. 1b), with distinct purple staining peripheral areas, 3.0–4.6 μm in diameter. Dark golden brown pigment granules dispersed throughout cytoplasm present.

Meront. Ovoid to pear-shaped (Fig. 1c), with slightly basophilic cytoplasm, measuring 4.6–10.2 × 4.2–7.2 (6.9 ± 1.3 × 6.2 ± 0.9) μm (n = 40), mostly polar in position within host cell cytoplasm. Dark golden brown pigment granules dispersed throughout meronts present (Fig. 1c), aggregating peripherally.

Mature meront. Polymorphic in shape (Fig. 1d-e), often slightly kidney shaped (Fig. 1d) to broadly oval (Fig. 1e),
measuring 4.8–9.9 × 4.1–6.9 (5.9 ± 1.1 × 5.8 ± 0.9) μm (n = 33); LW: 28–49.4 (38.4 ± 7) μm², with golden brown pigment granules aggregating either peripherally (Fig. 1d) or centrally within faintly eosinophilic cytoplasm (Fig. 1e). Meronts produced 12–25 (18.2 ± 3.5) (n = 33) pink-stained merozoites (Fig. 1d arrow), lying at periphery. Ruptured meronts dispersed free-purple-stained merozoites within host cell cytoplasm (Fig. 1e arrowhead). Meront size relative to host cell nucleus (LW/HNLW): 0.7–1.4 (1.0 ± 0.2) and to normal erythrocyte nuclei (LW/NNLW): 0.9–1.6 (1.2 ± 0.2) (n = 50).

**Gametocyte.** Variable in shape, often oval to kidney-shaped, positioned in polar regions of host erythrocyte (Fig. 1f–g). Mature gametocytes, elongated or kidney-shaped, pale pink-stained, with fine malarial pigment (Fig. 1f arrow), or rounded with predominantly basophilic cytoplasm (Fig. 1f–g), producing strands of malarial pigment centrally (Fig. 1g) and peripherally (Fig. 1h).

**Microgametocyte.** Kidney-shaped, pale-staining (Fig. 1f), measuring 8.4–10.4 × 3.9–4.6 (9.9 × 4.1) μm; LW: 45.6–55.5 (52.0) μm² (n = 18); tending to curve around host cell nucleus, sometimes slightly displacing it. Cytoplasm pale-pink, contains fine irregular brown to black pigment granules, loosely aggregating peripherally. Distinct pink staining nuclei aggregate peripherally (Fig. 1f arrow). Gametocyte size relative to host cell nucleus (LW/HNLW): 1.3–2.3 (1.7 ± 0.4) and to normal erythrocyte nuclei (LW/NNLW): 1.7–2.9 (2 ± 0.5) (n = 50).

**Macrogametocyte.** Deep basophilic staining, kidney-shaped, curving around nucleus (Fig. 1g) measuring 6.1–10.5 × 6.2–7.9 (10.1 ± 1.1 × 7.3 ± 0.9) μm (n = 44); LW: 52.7–91.4 (62.5 ± 16) μm²; with course, rounded dark-brown pigment often loosely clumped (Fig. 1g) or evenly dispersed centrally (Fig. 1h). Dark chromatin strands arranged peripherally (Fig. 1g–h). Macrogametocyte size relative to host cell nucleus (LW/HNLW): 1.6–2.0 (1.8 ± 0.2) and to normal erythrocyte nuclei (LW/NNLW): 1.4–21.8 (1.6 ± 0.2) (n = 50).

**Remarks**

Prior to this study, only two species of *Plasmodium* had been described parasitising cordyloid lizards from South Africa, *Plasmodium cordyli* and *Plasmodium zonuriae*, originally described from cordylids of Tanzania and South Africa, respectively [6]. Infections of *Plasmodium zonuriae* (Pienaar, 1962) Telford, 1987 were confirmed in the type-host lizard in a new locality Roodewalshoek in the Mpumalanga Province. Telford [6] stated that he found *P. cordyli* infecting *C. vittifer* from the then Transvaal Province (incorporating the present Mpumalanga Province), the same host and region from which *P. zonuriae* was originally described by Pienaar [5] and reported in this study. Thus it is possible that in future with the addition of more *C. vittifer* samples, we may be able to isolate *P. cordyli* for molecular analysis. Interestingly, Telford [6] described two morphological variants of *P. cordyli*, accounting for these differences as a result of parasitism of different host species.

**Plasmodium intabazwe n. sp.**

**Type-host:** *Pseudocordylus melanotus* (A. Smith, 1838) (Sauria: Cordylidae).

**Vector:** Unknown.

**Type-locality:** Donkey Pass (28°16’28.99"S, 29°12’21.39”E) (2,312 m), Platberg Reserve (Free State Province).

**Other localities:** Gibson Dam (28°16’33.12”S, 29°12’40.13”E) (2,292 m) and Platberg Pan (28°14’35.28”S, 29°09’47.02”E) (2,258 m) across the summit of Platberg, Platberg Reserve (Free State Province).

**Type-material:** Hapantotype, 1× blood smear from *Pseudocordylus melanotus* deposited in the Protozoan Collection of the National Museum, Bloemfontein, South Africa under accession number NMB P 412; parahapantotype, 1× blood smear from *Pseudocordylus melanotus*; deposited in the Protozoan Collection of the National Museum, Bloemfontein (NMB), South Africa, under accession number NMB P 413.

**Representative DNA sequences:** The cyt-b gene sequences have been submitted in the GenBank database under the accession numbers KX121601–KX121607.

**ZooBank registration:** To comply with the regulations set out in article 8.5 of the amended 2012 version of the International Code of Zoological Nomenclature (ICZN) [20], details of the new species have been submitted to ZooBank. The Life Science Identifier (LSID) of the article is urn:lsid:zoobank.org:pub:5BD63E9C-CA4C-4292-9AE2-E0D79169259F. The LSID for the new name *Plasmodium intabazwe* is urn:lsid:zoobank.org:act:797829AA-9E9A-4C98-B5A3-E4080DF9DBE2.

**Etymology:** The species epithet is derived from the Zulu word for ‘flat mountain’ since the first records of this malarial parasite were found in the blood of *Pseudocordylus melanotus* collected on Platberg (Afrikaans for ‘flat mountain’) in the eastern Free State, South Africa.

**Description**

**Trophozoite.** Rounded (Fig. 2a, d arrow) to oval (Fig. 2b arrow), with purple-stained periphery (Fig. 2a-b) and pinkish central vacuole, measuring 1.6–2.2 × 1.4–2.3 (1.9 ± 0.1 × 1.6 ± 0.25) μm (n = 30); LW: 1.4–10.4 (3.6 ± 3) μm² (n = 50).
**Immature meront.** Broadly rounded (Fig. 2c-d), sometimes amoeboid, stained deeper pink, measuring 4.2–5.5 × 2.4–5.2 (4.9 ± 0.3 × 4.2 ± 0.7) μm (n = 30); LW: 6.7–20.8 (15.6 ± 4.2) μm² (n = 50); mostly polar in position within host cells. Dark golden-brown pigment granules dispersed throughout meronts present (Fig. 2c), often aggregating at one end (Fig. 2d arrowhead). Immature meront size relative to host cell nucleus (LW/HNLW): 0.2–0.6 (0.5 ± 0.1) and to normal erythrocyte nuclei (LW/NNLW): 0.2–0.6 (0.5 ± 0.1) (n = 50).

**Mature meront.** Polymorphic in shape (Fig. 2e-f), often oval (Fig. 2e) to rounded (Fig. 2f arrowhead), or kidney-shaped (Fig. 2g), measuring 3.8–6.5 × 3.7–6.7 (5.2 × 5.2) μm (n = 18); LW: 25–44.4 (34.5) μm² (n = 18); with irregular dark-brown or black pigment granules often loosely aggregating at periphery (Fig. 2e–g arrowhead). Meronts produced 8–14 pink-staining nuclei (merozoites) (Fig. 2e arrow), lying at periphery. TEM micrographs of erythrocytes containing *P. intabazwe* n. sp. revealed a meront lying in a polar position in an erythrocyte (Fig. 2j). The meront was sectioned so that two nuclei (merozoites) were exposed. Also observable was a surface micropore (mp in Fig. 2j), various intracytoplasmic vacuoles (Fig. 2j arrows), and a structure resembling a dense body (db in Fig. 2j). The meront showed no evidence of a parasitophorous vacuole, the parasite surface membrane being in direct contact with the host erythrocyte cytoplasm (cy in Fig. 2j). Mature meront size relative to host cell nucleus (LW/HNLW): 0.7–1.4 (1.1 ± 0.3) and to normal erythrocyte nuclei (LW/NNLW): 0.7–1.3 (1 ± 0.2) (n = 50).

**Gametocyte.** Variable in shape, often kidney-shaped, positioned in polar regions of host erythrocyte (Fig. 2h–i). Later forms, elongated or kidney-shaped, pale pink-stained, with fine malarial pigment (Fig. 2h), or deeply purple-stained, containing coarse evenly distributed dark granules of malarial pigment (Fig. 2i).

**Microgametocyte.** Kidney-shaped to loosely oval (Fig. 2h), pale-staining, tending to curve slightly around host cell nucleus, measuring 6.5–7.7 × 3.6–4.1 (6.9 ± 5.4) μm (n = 8); LW: 47–57.5 (51.5) μm² (n = 8); seldom causing displacement of host cell nucleus. Fine irregularly shaped, dark-brown malarial pigment tends to aggregate peripherally (Fig. 2h). Diffuse pale-pink nuclei (Fig. 2h) often clumping at extremities of cytoplasm. Microgametocyte size relative to host cell nucleus (LW/HNLW): 1.5–2 (1.7 ± 0.2) and to normal erythrocyte nuclei (LW/NNLW): 1.4–1.7 (1.5 ± 0.1) (n = 50).

**Macrogametocyte.** Deep staining, elongate kidney-shaped, curving around the nucleus (Fig. 2i), with evenly dispersed dark-brown pigments, measuring 5.5–7.1 × 5.4–6.6 (6.4 × 5.8) μm (n = 8); LW: 52.7–91.5 (62.5) μm² (n = 8). Macrogametocyte size relative to host cell nucleus (LW/HNLW): 0.01–1.5 (1.1 ± 0.4) and to normal erythrocyte nuclei (LW/NNLW): 0.01–1.4 (1.0 ± 0.4) (n = 50).

**Remarks**

Observation of division stages within the peripheral blood, such as meronts, indicated that this parasite belongs to the genus *Plasmodium*, as compared to the closely related *Haemocystidium*, a genus of reptile plasmodiid parasite that shows no divisional stages within the peripheral blood (see [2, 6, 9]). *Plasmodium intabazwe* n. sp. has unique morphological and morphometric features when compared to the other two *Plasmodium* spp. from cordylids, *P. cordyli* and *P. zonuriae* (see Table 1). Trophozoites and young meront stages of *P. cordyli* could not be compared with those of *P. intabazwe* n. sp. as they were not described for *P. cordyli*. Meronts of *P. intabazwe* n. sp. are similar in appearance and size to those of *P. cordyli* infecting the type-host *C. tropidosternum* in that both species demonstrate polymorphic meronts ([7], measuring (mean 12 vs 7.5 × 4.5 μm), with merozoite numbers averaging eight and seven, respectively. However, meronts of *P. intabazwe* n. sp. do not possess the characteristic fan-shaped and are not strongly nucleophilic as was reported to be evident in *P. cordyli* meronts. Notably, meronts of *P. cordyli* in the South African *C. vittifer* also lack the tendency to be nucleophilic and furthermore, the characteristic fan-shaped meronts were described as rare. Also, meronts of *P. cordyli* infecting *C. vittifer* were even larger than meronts of *P. cordyli* infecting the type-host *C. tropidosternum*, as well as being larger than the meronts of *P. intabazwe* n. sp. at (mean 7.0 × 5.6 vs 5.0 × 4.1 vs 5.2 × 5.2 μm, respectively) and contained more merozoites (mean 12 vs 7.5 vs 11, respectively). Gametocyte morphotypes of *Plasmodium cordyli* in both hosts, *C. tropidosternum* and *C. vittifer*, were reported by Telford [6] to be almost identical (dimorphic gametocytes were not reported), demonstrating rounded or ovoid gametocytes measuring on average 6.5 × 5.5 in *C. tropidosternum* vs 7.0 × 5.5 μm in *C. vittifer*, [6, 7] as compared to the kidney-shaped gametocytes of *P. intabazwe* n. sp. measuring on average 6.3 × 6 μm, which are sexually dimorphic and easily distinguishable based on pigment distribution, size and staining properties.

Microscopical examinations of the meront and gametocyte morphology led to the identification of *P. zonuriae*. Even though there was a degree of variation in the sizes of these stages in *P. zonuriae* between descriptions (see Table 1) ([5, 6], this study), there were common characteristic features. These include merozoite numbers (mean 18) and features such as the pigment granules being dispersed within both the meronts and gametocytes,
and these stages curving around the host cell nucleus. Morphologically, trophozoite stages of *P. intabazwe* n. sp. and *P. zonuriae* bore a close resemblance in size, trophozoites of both species measuring on average ~1.9 × 1.6 μm. Trophozoites of *Plasmodium intabazwe* n. sp. are polymorphic as are trophozoites of *P. zonuriae*. Furthermore, early meront stages of *P. intabazwe* n. sp. also resembled those of *P. zonuriae* in size and appearance, forming shapes from oval to elongated, or even on occasion amoeboid shapes. However, the division stages (meronts) in the peripheral blood of the two species differ. The meronts of *P. intabazwe* n. sp. contain eosinophilic cytoplasm with fine, yet abundant pigment that aggregates peripherally or centrally as compared to the division stages of *P. zonuriae*, which contain evenly distributed pigment at the periphery. Furthermore, meronts of *P. intabazwe* n. sp. are overall smaller than the meronts of *P. zonuriae* (mean 5.2 × 5.2 vs 12.0 × 6.5 μm) (unfortunately these dimensions were not provided in the original description of *P. zonuriae* [5]) see [6]. In addition, meronts of *P. intabazwe* n. sp. contain eight merozoites on average (vs 18 in *P. zonuriae*) [5, 6]. The gametocyte stages of *P. intabazwe* n. sp. and *P. zonuriae* also differed substantially. Gametocytes of *P. intabazwe* n. sp. are sexually dimorphic and usually kidney-shaped, measuring on average 6.3 × 6.0 μm, and differ from *P. zonuriae*, which are usually elongated or rounded, measuring on average 10.9 × 5.7 μm ([5, 6], this study). Although microgametocyte stages of *P. intabazwe* n. sp. may at times resemble those of *P. zonuriae* in appearance and size, the pigment granules were in general much finer and evenly dispersed ([5, 6], this study).

At a cytopathological level, erythrocytes infected with *P. intabazwe* n. sp. were neither hypertrophied nor dehaemoglobinized and rarely stained paler than non-infected cells. On the other hand, infections with *P. zonuriae* in *C. vittifer* are reported to have caused severe anaemic changes in the peripheral blood, causing hypertrophy, and displacing and distorting host cell nuclei [5, 6]. However, these effects may not always be present, as reported for the meronts of *P. zonuriae* infecting its other host *P. micropelidotus* from the Western Cape Province, which caused little distortion of either the host cell or its nucleus. Similarly, *P. intabazwe* n. sp. also showed little effect on host cells, but in some cases meronts and gametocytes tended to curve around the host cell nucleus, though they never displaced or deformed it.

**Molecular analysis**

Amplicons of 500 nt from *Plasmodium zonuriae* and *Plasmodium intabazwe* n. sp., were derived from the blood of two *C. vittifer* and seven *P. melanotus*, respectively. Sequences isolated in the current study from *P. zonuriae* and *P. intabazwe* n. sp. confirmed the morphological observations by representing two distinct morphotypes, with an interspecific divergence of 3.4 % based on 497 nt sequence comparisons (percentage of bases/identical residues). For the phylogenetic analysis, only a single nucleotide sequence from both *P. zonuriae* and *P. intabazwe* n. sp. was used, as all the nucleotide sequences obtained in the current study from both *P. zonuriae* (*n* = 2) and *P. intabazwe* n. sp. (*n* = 7) were identical. These sequences were analysed together with sequences for 18 *Plasmodium* spp., four *Haemoproteus* spp., five *Haemocystidium* spp. and two *Leucocytozoon* spp. (used as the outgroup), all downloaded from the GenBank database (see Table 2).

The BI tree (Fig. 3) showed *Plasmodium* species isolated from lizard and bird hosts to be polyphyletic. *Plasmodium* species isolated from primate hosts were shown to form a well-supported monophyletic clade, forming a sister clade to *Plasmodium chrichtonae* (AY099061) and *Plasmodium mexicanum* (AY099060) isolated from North American lizard hosts. *Haemoproteus* species isolated from bird hosts formed a well-supported monophyletic clade separate from *Plasmodium* and *Haemocystidium* species. At the base of the phylogeny *Haemocystidium* species isolated from reptilian hosts were shown to be paraphyletic. The phylogenetic analysis confirmed the generic placement of the two species, *P. zonuriae* and *P. intabazwe* n. sp., within a large monophyletic clade consisting of only *Plasmodium* species. The two species from the current study were also shown to be closely related, forming a monophyletic clade with strong nodal support. Furthermore, these sequences clustered with *Plasmodium* species isolated from other lizard hosts.

**Discussion**

*Plasmodium intabazwe* n. sp. can be distinguished from other African cordyliid malarial parasites, both morphologically in terms of not only its morphometric characteristics, but also its staining properties, distribution of pigment and cytopathology.

Even though the morphological findings strongly suggested that *P. intabazwe* n. sp. is a different species to both *P. cordyli* and *P. zonuriae*, *P. intabazwe* and *P. zonuriae* did share similar morphological characteristics for a number of their peripheral blood life stages. This highlights the importance, when possible, of using both morphology and molecular characteristics in species descriptions. Even in cases where the morphology of two potential species is different, it is important to use molecular methods to differentiate them. As emphasized by Perkins [21], morphology of parasites can alter depending on the processing conditions of the slides, never mind the potential plasticity of the traditional characteristics used to differentiate these organisms. Unfortunately, the morphological findings could not be supported molecularly...
for *P. cordyli*. It would have been preferable to have molecularly compared *P. cordyli* to both the other species. However, if found in future through more extensive sampling, we would still be cautious in characterising this species as *P. cordyli* given that *C. vittifer* and South Africa are not the type-host or locality and given the morphological differences between the two variants.

The morphological findings regarding *P. zonuriae* and *P. intabazwe* n. sp. were further supported by the molecular data obtained for both species with an interspecific divergence of 3.4 %. Although [22] and [23] have suggested that the mean ‘cut-off’ for distinguishing between different *Plasmodium* and *Haemoproteus* species is an interspecies divergence of 5 %, a number of species such as *Haemoproteus pallidus* Valkiūnas & Iezhova, 1991 and *Haemoproteus minutus* Valkiūnas & Iezhova, 1992, with an interspecific divergence of approximately 1 % in sequences of the cyt-b gene, have been shown to be separate species when combining both morphological and molecular data (see [22, 24]).

In the phylogenetic analysis, *P. zonuriae* and *P. intabazwe* n. sp. were well nested within a larger monophyletic clade comprising only *Plasmodium* species. Although this clade contained moderate posterior probability support, the overall tree topology was similar to other studies such as [24], and particularly with a review on the closely

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**Table 2** List of organisms used in the phylogenetic analyses of this study according to associated host group, with associated GenBank accession numbers, host and references

| Group                        | Accession number | Parasite       | Host                  | Reference                      |
|------------------------------|------------------|----------------|-----------------------|--------------------------------|
| *Plasmodium* spp. of lizards |                  | *P. chiricahuae* | *Sceloporus jarrovi*  | [46]                           |
|                              |                  | *P. fairchidi*  | *Norops cupreus*      | [47]                           |
|                              |                  | *P. floridense* | *Anolis acutatus*     | [46]                           |
|                              |                  | *P. gemini*     | *Hysilurus modestus*  | [48]                           |
|                              |                  | *P. intabazwe* n. sp. | *Pseudocordylus melanotus* | Current study                  |
|                              |                  | *P. koreafense* | *Sphenomorphus johiensis* | [48]                           |
|                              |                  | *P. lacertiliae* | *Ernnea longicauda*   | [48]                           |
|                              |                  | *P. megalotrypa* | *Sphenomorphus simus* | [48]                           |
|                              |                  | *P. mexicanum*  | *Sceloporus occidentalis* | [46]                           |
|                              |                  | *P. minuvirkide* | *Prasinohaema prehensicauda* | [48]                           |
|                              |                  | *P. zonuriae*   | *Cordylus vittifer*   | Current study                  |
| *Plasmodium* spp. of birds   |                  | *P. circumflexum* | *Sylvia atricapilla*  | Unpublished                     |
|                              |                  | *P. elongatum*  | *Passer domesticus*   | [49]                           |
|                              |                  | *P. gallinaceum* | *Gallus gallus*       | [46]                           |
|                              |                  | *P. relictum*   | *Hemignathus viens*   | [50]                           |
| *Plasmodium* spp. of primates|                  | *P. falciplarium* | *Homo sapiens*       | [49]                           |
|                              |                  | *P. knowlesi*   | *Old World monkeys*   | [49]                           |
|                              |                  | *P. malariae*   | *Homo sapiens*        | [49]                           |
|                              |                  | *P. reichenowi* | *Pan troglodytes*     | [49]                           |
|                              |                  | *P. vivax*      | *Homo sapiens*        | [49]                           |
| *Haemoproteus* spp.          |                  | *H. balmorali*  | *Muscicapa striata*   | [22]                           |
|                              |                  | *H. lanii*      | *Lanius collurio*     | [22]                           |
|                              |                  | *H. pallidus*   | *Ficedula hypoleuca*  | [22]                           |
|                              |                  | *H. parabelopolskyi* | *Sylvia atricapilla*  | Unpublished                     |
| *Haemocystidium* spp.        |                  | *H. kopki*      | *Teratoscincus scincus* | [46]                           |
|                              |                  | *H. mesnili*    | *Naja annulifera*     | [2]                            |
|                              |                  | *H. pacayae*    | *Podocnemis unifilis* | [2]                            |
|                              |                  | *H. pelletopahli* | *Podocnemis expansa* | [2]                            |
|                              |                  | *H. ptyodactyllo* | *Pyodactylus hasselquistii* | [46]                           |
| *Leucocytozoon* spp. (outgroup) |                  | *L. gentili*    | *Passer domesticus*   | [51]                           |
|                              |                  | *L. majoris*    | *Fringilla coelebs*   | [51]                           |
related haemoprotid genera (see [2]). In the latter study, although the *Haemoproteus* spp. clade was shown to be the basal clade, the nodal support values separating the major *Plasmodium* species, *Haemocystidium* species and *Haemoproteus* species clades were all considerably higher. This was most likely attributable to the multigene and concatenated approach that was used for the phylogenetic analysis by [2], as compared to the current study’s cyt-*b* gene analysis, which is expected to provide a less robust analysis due to its variability.

Outside South Africa, and other than *P. cordyli* in Tanzania [7], *P. intabazwe* n. sp. is perhaps morphologically closest to *Plasmodium ulugurum* Telford, 1984 described from the gecko *Hemidactylus platycephalus* in the Uluguru Mountains, Tanzania [25]. Meronts of *P. ulugurum*, however, were more variable in size and measured on average $7 \times 4 \mu m$, while meronts of *P. intabazwe* n. sp. were more constrained in their dimensions (mean $5.2 \times 5.2 \mu m$). The gametocytes of the two species also differ, with those of *P. ulugurum* being larger and ovoid (mean...
7.5 × 5.5 μm), the pigment also not dispersed within the gametocytes, but rather forming an aggregate of dark greenish-yellow granules near the cell margin of both sexes of gametocytes. The smaller, microgametocytes of *P. intabazwe* n. sp. had distinct pigment distributed around the periphery of the parasites, measuring (6.0–7.9 × 3.5–5.6 μm). Microgametocytes of *P. uluguruense* were reported to be longer and larger than the macrogametocytes, measuring on average 8 × 5 μm [6].

Interestingly, as reported for *P. microlepidotus* during Pienaar’s [5] study, in this study *P. melanotus* were also found to be parasitised by the prositigmatic mites *Zonurobia semilunaris* [26], although they occur in disjunctive localities. These mites were the only haematophagous invertebrates observed feeding on *P. melanotus*, warranting further study in future for developmental stages of *P. intabazwe* n. sp.

**Conclusions**
The molecular characterisation of *P. intabazwe* n. sp. along with that of *P. zonuriae* represents the first combined morphological description and molecular characterisation of South African saurian malarial parasites, additionally confirming their placement within *Plasmodium*. The detection and description of the sporogonic stages in the natural vector should also be a key consideration in future species descriptions. Future research should include the identification of possible definitive hosts or vectors such as mites and mosquitoes as well as experimental transmission studies. Most importantly however, this work increases biodiversity knowledge of saurian malarial parasites, which by adding to the known taxa, contributes to further studies into these organisms’ taxonomy and phylogeny.

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**Availability of data and material**
The datasets supporting the conclusions of this article are available in the NCBI (National Center for Biotechnology Information) repository under the GenBank accession numbers KX121601–KX121609, and the ZooBank repository under the life science identifier (LSID): urn:lsid:zoobank.org:pub:58D63E9C-CA4C-4292-9AE2-E0D79169259F.

**Authors’ contributions**
All authors conceived and designed the project, participated in general data analysis and in drafting the manuscript. JVA carried out the fieldwork, prepared and examined blood smears, prepared light micrographs and compiled all measurement data. ECN participated in the molecular studies and in the sequence alignment and CAC provided significant support to the preparation of the manuscript. All authors read and approved the final manuscript.

**Competing interests**
The authors declare that they have no competing interests.

**Consent for publication**
Not applicable.

**Ethics approval and consent to participate**
This study received the relevant ethical approval from the North-West University Ethics Department (North-West University ethics approval no: NWU-00005-14-53).

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