

A Case Report of *Babesia* sp. in Feline Host: Diagnosis and Successful Outcome

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Abstract

Babesiosis is a disease of mammals including cats caused by a hemoparasitic protozoan *Babesia* sp. affecting erythrocytes and vascular endothelium. A two-year-old, intact male, Persian cat with an indoor-outdoor lifestyle was presented to the District Veterinary Centre, Palakkad, Kerala, with a history of persistent vomiting, progressive lethargy, and anorexia. Hematological analysis demonstrated marked anemia, thrombocytopenia, and leukopenia. Microscopic examination of Giemsa-stained peripheral blood smear prepared from ear tip revealed the presence of *Babesia* organisms within erythrocytes. Based on morphological characteristics, a presumptive diagnosis of feline babesiosis was made; however, definitive species-level identification remained inconclusive due to the absence of molecular characterization. The cat was treated with a combination therapy consisting of diminazene aceturate and doxycycline. The animal showed rapid clinical improvement, with resolution of clinical signs and had an uneventful recovery. This case highlights the importance of hematological and blood smear evaluation for early detection of feline babesiosis, alongside low-cost molecular methods for species confirmation and the potential efficacy of combination therapy.

Keywords: *Babesia* sp.; Cat, Kerala; Diminazene aceturate; Anemia; Palakkad.

1. Introduction

Babesiosis is a worldwide disease affecting mammals caused by protozoan parasites of the genus *Babesia*, which invade erythrocytes by rapid multiplication and impact the vascular endothelium. In domestic cats, babesiosis is caused by several *Babesia* species. These include small-forms- *B. felis* and *B. cati*, as well as large-forms- *B. herpailuri* and *B. pantherae* with a varying size of 1.0–2.5 µm and 2.5–5.0 µm, respectively.[1] Multiple reports have been documented in cats affected with *B. canis* and *B. gibsoni* which is a common parasite found in dogs rather than cats.[2,3] *Babesia felis* is the species most frequently responsible for causing babesiosis in cats. Feline babesiosis usually presents with anorexia, lethargy, and weakness as the most frequent signs, whereas pyrexia, an enlarged spleen, icterus, vomiting, and respiratory symptoms are seen occasionally. In extreme cases, cats may experience multiorgan dysfunction complications like kidney failure, lung edema, liver and neurological affections.[4] *Babesia* species have merozoites that appear very similar in form, relying on morphology alone is insufficient for accurate classification and hence molecular studies play a crucial role in distinguishing and properly identifying these organisms.[1] The present study reveals a case report of feline babesiosis diagnosis and successful management.

2. Case Presentation

A two-year-old, intact male, Persian cat weighing 3.1 kg was presented dull and depressed to District Veterinary Centre, Palakkad, Kerala, India with history of vomiting, lethargy and progressive reduction in appetite for 4 days; the cat had a mixed indoor–outdoor lifestyle. While on examination clinically, an elevated body temperature of 103.9°F, pale conjunctival and buccal mucous membranes, dyspnea and salivation were noticed. No ticks were detected on the body of the cat during clinical examination. Haematological evaluation was performed using an automated haematology analyser (Exigo H400, Boule Diagnostics, Sweden) according to the manufacturer’s instructions. The analysis revealed leukopenia, characterized by decreased neutrophil and eosinophil counts, along with an increased monocyte count. The lymphocyte count remained in the normal range. A noticeable reduction was found in total erythrocyte counts, haemoglobin and haematocrit. The red cell indices—mean corpuscular volume, mean corpuscular haemoglobin, and mean corpuscular haemoglobin concentration—remained within the established normal reference ranges, indicating a normocytic, normochromic anaemia. A severe reduction in platelet count ($12 \times 10^9/L$) was also noticed, suggestive of thrombocytopenia. The hematological parameters observed in the present case are summarized in Table 1. Thin peripheral blood smears prepared from the ear tip were air-dried, fixed in absolute methanol for 30 seconds, and stained with Giemsa working solution for 30 minutes at room temperature. The smears were gently rinsed with buffered water, air-dried, and examined under a $\times 100$ oil-immersion objective. Single and paired piroplasms arranged at an acute angle were observed within

erythrocytes, suggestive of *Babesia* sp. (Fig. 1).

Table 1: Hematological parameters observed in the present case

Blood Parameters	Reference range	Results
Total leukocyte Count ($10^9/L$)	5.5-19.5	4.1
Neutrophils ($10^9/L$)	2.8-13	1.5
Eosinophils ($10^9/L$)	0.1-1	<0.1
Monocytes ($10^9/L$)	0.2-1	1.2
Lymphocytes ($10^9/L$)	1-7	1.3
Total Erythrocyte Count ($10^{12}/L$)	5-11	3.86
Haemoglobin (g/dL)	8-15	6
Haematocrit (%)	25-45	16.8
Mean Corpuscular Volume (fL)	39-50	43.6
Mean Corpuscular Hemoglobin (pg)	12.5-17.5	15.5
Mean Corpuscular Haemoglobin Concentration (g/dL)	31-38.5	35.6
Platelet count ($10^9/L$)	200-500	12



Fig. 1: Babesia sp. piroplasm in RBC on Giemsa-stained peripheral blood smear (100x).

Treatment and Evaluation

The animal was initially treated with an intramuscular injection of diminazene aceturate (Berenil®, MSD) at a dose of 3.5 mg/kg. Additionally, doxycycline was administered orally at 10 mg/kg (Doxypet, Savavet) for a duration of 21 days. Supportive care, including fluid administration, multivitamin injections, and oral haematinics, was also provided as per the requirement. The animal was handled humanely throughout the diagnostic and therapeutic procedures. All clinical interventions were performed in accordance with standard veterinary practices, with appropriate measures taken to minimize pain, stress, and suffering. Care was taken to provide appropriate supportive and therapeutic management in view of the cat’s dyspnea and severe anemia. The animal’s appetite began to improve from the second day of treatment and showed rapid clinical improvement. After 21 days, hematological parameters returned to normal and the cat had an uneventful recovery.

3. Results and discussion

Feline babesiosis has been reported worldwide, with documented cases from several regions including Spain,

Portugal, Israel, Poland, and Thailand.^[4] Reports of babesiosis in India have been documented in regions such as Tamil Nadu, Gujarat and Himachal Pradesh.^[5-7] In Kerala, multiple studies have reported the occurrence of feline babesiosis. Investigations by Sabu *et al.* [2013], Rejimon *et al.* [2022] and Vincy and Tressamol [2023] provide evidence of its prevalence across different hosts.^[8-10] Furthermore, the identification of a novel feline species, *Babesia panickeri*, by Panicker *et al.* [2020] underscores the endemic nature of babesiosis in Kerala and highlights its continued presence in the region.^[4]

The cat in the present case had a regular access to both indoor and outdoor environments, potentially increasing its opportunity for exposure to arthropod vectors. Clinical manifestations observed in this study, including pale mucous membranes, fever, and anorexia, are consistent with those reported in previous cases of babesiosis.^[5,11,12] Similarly, the haematological alterations like anaemia, thrombocytopenia, and leukopenia align with the findings of Aziz *et al.* [2025] and Rejimon *et al.* [2022].^[5,9] These changes are characteristic of babesiosis and reflect the underlying pathological processes, which commonly involve reductions in red blood cell count, haemoglobin concentration, haematocrit, and platelet count.^[13] The concurrent absolute monocytosis observed in this study may reflect an increased demand for erythrophagocytic activity, resulting in increased circulating monocytes. Although leukopenia accompanied by monocytosis is rarely reported in babesiosis, this hematological finding may have been associated with immunosuppression secondary to a concurrent infection, such as *Mycoplasma haemofelis* and may have resolved following doxycycline administration.

Although *Babesia felis* is the most commonly discussed *Babesia* sp. in cats, several reports have documented the occurrence of large *Babesia* species in felines, including those associated with both cats and dogs. Since the tick was not detected from cat, the transmission of the infection remains uncertain. Although primaquine phosphate is considered efficacious and the drug of choice in feline babesiosis, its temporary unavailability necessitated the use of a combination therapy consisting of diminazene aceturate (Berenil®) and doxycycline.^[5,14,15] Several species of *Babesia* have been reported in cats including *B. felis*, *B. canis* and *B. lengau*.^[3,16,17] In the present case, microscopic examination revealed intraerythrocytic piroplasms morphologically suggestive of a large *Babesia* species. However, morphological assessment alone is insufficient for definitive species-level identification of *Babesia*. Therefore, molecular characterization is essential to confirm the species identity and distinguish closely related *Babesia* spp. The absence of molecular confirmation represents an important limitation of the present study and precludes definitive species-level identification of the organism based solely on its morphological characteristics. Thus, the present study highlights the need for low-cost molecular diagnostics in modern feline veterinary practice and explores research opportunities in endemic regions such as Kerala, where a novel *Babesia* species has been identified in cats.

4. Conclusions

Feline babesiosis should be considered in cats presenting with anemia, thrombocytopenia, and nonspecific clinical signs. This case emphasizes the importance of careful hematological evaluation and blood smear examination for the early detection of feline babesiosis, relevance of low-cost molecular techniques for species confirmation at veterinary practice and demonstrates the potential effectiveness of combination therapy in managing the infection.

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CRedit Author Contribution Statement

Farhan Eravaramkunnath: Conceptualization; Formal Analysis; Methodology. **Fathima Dhina:** Conceptualization; Data Curation. **Arya Ramachandran Nair Sujatha Devi:** Formal Analysis; Investigation; Supervision. **Bindhu Lakshmanan:** Supervision, Validation, Writing – Review & editing. All authors have read and approved the final version of the manuscript for publication and agree to be accountable for all aspects of the work, ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

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Institutional Review Board Statement

All procedures were conducted in accordance with institutional guidelines and applicable national animal welfare regulations. As this was a retrospective case report of routine clinical care, IACUC approval was not required. The report was prepared in accordance with the CARE guidelines.

Informed Consent Statement

Informed consent was obtained from the owner for the treatment of the animal.

Consent to Publish Statement

Informed consent was obtained from the owner for the publication of the clinical findings and associated information.

Data Availability Statement

The datasets generated and/or analyzed during the current study that support the findings are available from the corresponding author upon reasonable request.

Conflict of Interest

There is no conflict of interest.

Artificial Intelligence (AI) Use Disclosure

The authors declare that artificial intelligence (AI)-

assisted tools were used only for language refinement, grammar improvement, and manuscript structuring purposes during the preparation of this work. All technical content, experimental implementation, results, and interpretations were independently developed and verified by the authors.

Supporting Information

Not applicable.

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