Detection of *Ehrlichia canis* in bone marrow aspirates of experimentally infected dogs

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ABSTRACT

The present work describes the detection of infected cells in the bone marrow aspirates of dogs experimentally infected with a Brazilian isolate of *Ehrlichia canis*. Dogs were monitored twice a day by clinical evaluation and peripheral blood smear examination. Every three days, blood samples were collected for cell counts. Weekly, aspirates from the bone marrow were examined and serum samples were tested by IFAT. The clinical signs observed were fever, pallid membranes, lymphadenopathy, serous nasal secretions, and pronounced weight loss. Hematological alterations included normocytic normochromic anemia, decrease of neutrophils and lymphocytes, and thrombocytopenia. Few *E. canis* infected cells were seen in blood smears. However, stages of *E. canis* were visualized in bone marrow aspirates 15 days post infection.

Key words: *Ehrlichia canis*, diagnosis, bone marrow puncture, dogs.

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Canine ehrlichiosis, also known as Tropical pancytopenia, is a fatal hemorrhagic syndrome, characterized by thrombocytopenia, epistaxe and progressive pancytopenia (HUXSOLL et al., 1970). The causal agent, *Ehrlichia canis*, is a rickettsia transmitted by the tick *Rhipicephalus sanguineus*, that multiplies inside mononuclear leukocytes, affecting several organs, such as the spleen, lymphonodes and liver (WOLDEHIWET & RISTIC, 1993).

During the acute phase, the parasite can be detected in the cytoplasm of circulating monocytes for a short period of time. The
surviving animals remain infected, but the parasite cannot be seen in blood smears, which is still the routine diagnostic method for detection of hemoparasites.

Cases of ehrlichiosis in dogs have increased considerably over the last years (MOREIRA et al., 2003) and there is a need for more sensitive and specific tests to enable clinicians implementing early treatment.

In the present study, we evaluated clinically and by laboratory tests two mongrel dogs that had been experimentally inoculated with 5 ml of cryopreserved blood infected with a Brazilian isolate of *Ehrlichia canis* (Jaboticabal strain). The *E. canis* strain was first isolated from a naturally infected dog and was frozen in liquid nitrogen at the Veterinary Teaching Hospital, UNESP, Jaboticabal, SP, Brazil (unpublished data).

The dogs were monitored daily by corporal temperature and peripheral blood smears, and every two days by hematological parameters and blood cell counts (erythrocytes, leukocytes and platelets). One of the dogs was also monitored weekly by direct examination of bone marrow aspirates. Blood smears and smears made from bone marrow aspirates were stained with Giemsa and examined by optical microscopy. Serum sample were obtained from the two animals before the experimental inoculation and 15 days after infection to evaluate humoral responses. These samples were tested for specific IgG response to *E. canis* by IFAT (Indirect Fluorescent Antibody test) using a commercial kit (MegaScreen FLOREHLICHIA c®). The two dogs were seronegative prior to infection.

The first clinical signs were detected 15 days after infection (PI), when the animals presented hyperthermia (mean temperature 40°C), oculus-nasal discharge and weight loss. The hematological alterations included normocytic normochromic anemia during the first week PI. In the third week PI, the total number neutrophils decreased, with a decrease of segmented neutrophils and lymphocytes, in addition to a thrombocytopenia.

Single morula were detected in peripheral blood smears on day 11 (animal 1) and on day 13 (animal 2) (Figure 1), while several *E. canis* inclusions were seen in the cytoplasm of monoblasts and mature monocytes in bone marrow aspirates on day 15 PI (Figure 2).

Inclusions of *E. canis* infecting bone marrow of dogs have been previously reported associated to a super acute disease with non-regenerative anemia, in which infected leukocytes were rare (Meinkoth et al., 1989). At that time, the infected dogs showed positive antibody titers for *E. canis* detected at 1:64 dilutions.

One of the problems associated with treatment of clinical cases of canine erlichiosis is related to difficulties in detecting morula in peripheral blood smears. Therefore, detection of parasites in other sites and in other biological fluids would represent improvements for the diagnosis and evaluation of recovery of clinically affected dogs.

It was concluded that *Ehrlichia canis* develops a parasitism in the bone marrow that allows...
Figure 2 - Microphotography of a bone marrow aspirate of a dog infected with *Ehrlichia canis* showing inclusions in the cell cytoplasm (arrow) (Giemsa staining 1,000 x). Bar 2.5mm.

detection of inclusions during the acute phase, suggesting that the direct examination of bone marrow aspirates can be useful for easier identification of *E. canis* infections in dogs even when no peripheral intracellular morulae are detectable.

This study (protocol n. 035/03) is in agreement with the Ethical Principles in Animal Experimentation, adopted by the **Ethics Committee on Animal Experimentation** (CETEA/UFMG) and was approved in December 2003.

**REFERENCES**


