‘Candidatus Mycoplasma haemominutum’, a low-virulence eperythrozcytic parasite of cats

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The phylogenetic position and some taxonomically relevant characteristics of a small, low-virulence bacterial parasite of cats are described. A 16S rDNA analysis revealed that the organism was in the Mycoplasma clade and was most closely related to a parasite of pigs previously designated Eperythrozoon suis. As the organism has not been cultured in vitro and is maintained in serial passage in cats in vivo, Candidatus status is proposed for this novel taxon as ‘Candidatus Mycoplasma haemominutum’.

Keywords: ‘Candidatus Mycoplasma haemominutum’, feline infectious anaemia, Haemobartonella felis

Morphological characteristics

Cells of ‘Candidatus Mycoplasma haemominutum’ are approximately 0·3 μm in diameter, approximately half the size of ‘Candidatus Mycoplasma haemofelis’ (Fig. 1). Cells of both species are Gram-positive, epicellular coccoid bodies. Ultrastructurally, ‘Candidatus Mycoplasma haemominutum’ is similar to other mollicutes, with a plasma membrane, no cell wall and no internal membranous structures.

Clinical disease

Clinical signs in cats infected with ‘Candidatus Mycoplasma haemominutum’ may be minor or absent and are not associated with fatality (Foley et al., 1998). Likewise, haematological abnormalities are minor, if present. In one study, the mean packed cell volume (PCV) declined during the course of infection from 48·6 to 31·4%, but did not go below reference ranges (Foley et al., 1998). Leukocyte counts also declined but stayed within reference ranges. ‘Candidatus Mycoplasma haemominutum’ organisms were visible for 7–21 d, beginning on day 14 post-inoculation (p.i.), and were detected singly or occasionally multiply on each erythrocyte. In most cases, the date of maximal parasitism was the date of minimum PCV. Tetracycline abolished patent infections associated with ‘Candidatus Mycoplasma haemominutum’ within 24 h and PCR tests became negative by 12 h after initiation of therapy (Foley et al., 1998).

Following the acute stage of parasitaemia, cats may enter a series of infection cycles consisting of periods of latency and recrudescence (Foley et al., 1998; Small & Ristic, 1987). Evidence of cycling was detected in four cats with ‘Candidatus Mycoplasma haemominutum’.

Abbreviations: FeLV, feline leukaemia virus; FIV, feline immunodeficiency virus; PCV, packed cell volume.
infections (Foley et al., 1998; Small & Ristic, 1987). Cat #88506 had a reoccurrence of anaemia and depression 7 months after acute anaemia, with no clinical signs or haematological abnormalities during the intervening period. During the reactivated attack, the PCV declined from 44.9 to 21.3%. Approximately 5% of erythrocytes appeared to be parasitized by ‘Candidatus Mycoplasma haemominutum’ in the later attack.

Co-infection of ‘Candidatus Mycoplasma haemominutum’ with feline retroviruses has been evaluated (J. George and N. C. Pedersen, unpublished data). The cats with the most severe erythrocyte reductions were those infected with the ‘Candidatus Mycoplasma haemominutum’ and feline leukaemia virus (FeLV), with or without feline immunodeficiency virus (FIV) infection. There was no mortality as a direct result of ‘Candidatus Mycoplasma haemominutum’ infection, but two cats in the FeLV-only group and one cat with FeLV and FIV developed myeloproliferative disorders characteristic of some end-stage FeLV infections during weeks 6 and 7.

Cats infected with ‘Candidatus Mycoplasma haemominutum’ develop IgG class antibodies that are somewhat cross-reactive with antibodies to ‘Candidatus Mycoplasma haemofelis’. Experimentally infected cats were evaluated with an indirect immunofluorescent antibody assay using acetone-fixed whole-blood smears from acutely infected cats as substrate (Foley et al., 1998). Smears were coated with a 1:25 dilution of patient serum in PBS and incubated for 30 min at 37 °C with moisture. After washing, smears were coated with a 1:25 dilution of fluorescein-conjugated rabbit anti-cat IgG (Antibodies Inc.) and lightly counter-stained with Evans blue. Characteristic bright-green fluorescence was observed when ‘Candidatus Mycoplasma haemominutum’ substrate smears were reacted with ‘Candidatus Mycoplasma haemofelis’ sera but not ‘Candidatus Mycoplasma haemofelis’ sera. In contrast, ‘Candidatus Mycoplasma haemofelis’ smears reacted with both ‘Candidatus Mycoplasma haemofelis’ and ‘Candidatus Mycoplasma haemominutum’ sera. Seroconversion in cats infected with ‘Candidatus Mycoplasma haemominutum’ occurred in most cats by day 21 p.i., with a maximum titre of 400 by day 28 p.i. Infection with ‘Candidatus Mycoplasma haemominutum’ protected cats against challenge with the more virulent ‘Candidatus Mycoplasma haemofelis’ (Foley, 2001).

‘Candidatus Mycoplasma haemominutum’ has not been cultivated in artificial media; its entire existence is intimately associated with erythrocyte surfaces. Mycoplasmas in general require highly enriched media for growth. Unpublished unsuccessful culture protocols have included culture on mycoplasma media and on fresh cat blood agar. However, the type specimen is maintained frozen at the University of California, Davis, and the organism is propagated by in vivo passage in cats.

The epidemiology of ‘Candidatus Mycoplasma haemominutum’ infection is not known. Based on other feline haemobartonella-like organisms, the routes of infection may include in utero or lactogenic, iatrogenic, possibly oral and haematophagous arthropod transmission (Cretillat, 1984; Fisher, 1983; Harbut, 1963; Harvey & Gaskin, 1977; Nash & Bobade, 1986; Splitter et al., 1956).

**PCR and genetic analysis**

A PCR for ‘Candidatus Mycoplasma haemominutum’ and *M. haemofelis* was designed to amplify DNA extracted from feline whole blood using the QIAamp tissue extraction kit (Qiagen) following the manufacturer’s instructions (Foley et al., 1998). Specific PCR primers are at position 1183 forward (5’-GCAT-AAGTGTGCAATC-3’) and 1290 reverse (5’-GT- TCCAATAGTACTTCTCC-3’) of the 16S rRNA gene. ‘Candidatus Mycoplasma haemominutum’/‘Candidatus Mycoplasma haemofelis’ PCR was performed using 20 pmol of each of these primers in 100 µl containing final concentrations of 50 mM KCl, 10 mM Tris/HCl, 0.1% Triton X-100, 2.5 mM MgCl2 and 0.2 mM each dNTP, 2 U Taq DNA polymerase and 60-6 µl water. The cycling conditions were 94 °C for 4 min, followed by 35 cycles of 94 °C for 30 s, 53 °C for 1 min and 70 °C for 45 s, followed by a 5 min elongation step at 70 °C and cooling to 4 °C. The
positive band on ethidium bromide-stained, 1·5 % agarose gels corresponded to a fragment of 130 bp. The 130 bp products were confirmed as ‘Candidatus Mycoplasma haemominutum’ amplicons by DNA sequencing. In ten cats with experimental ‘Candidatus Mycoplasma haemominutum’ infection, PCR tests were initially negative and became positive 8·8 d after inoculation on average. A published PCR protocol for 16S amplification of ‘Candidatus Mycoplasma haemofelis’ (Messick et al., 1998) does not amplify ‘Candidatus Mycoplasma haemominutum’ DNA.

The 16S rRNA sequence of ‘Candidatus Mycoplasma haemominutum’ has been reported by Rikihisa et al. (1997) as the California strain of H. felis. The similarity between ‘Candidatus Mycoplasma haemominutum’ and ‘Candidatus Mycoplasma haemofelis’ is 83%. Analysis of the DNA sequence of the relatively conserved 16S rRNA gene revealed that ‘Candidatus Mycoplasma haemominutum’ is only distantly related to the rickettsias and should instead be placed in the Mycoplastae (Foley et al., 2001). The phylogenetic data suggest that two distinct strains of Haemobartonella felis in domestic cats. Am J Vet Res 59, 1581–1588.


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