

***Babesia microti* IFA Kits**

**(Catalog BMG-120 and BMM-120)**

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**Performance Characteristics**

**SENSITIVITY**

The indirect immunofluorescence antibody assay (IFA) for *Babesia microti* was described in the literature in 1978<sup>1</sup> and has served thereafter as the most common method for serodiagnosis. The Fuller Laboratories test uses *Babesia microti*-(GI strain)-infected hamster or mouse erythrocytes as the antigen substrate.

Specific IgM antibody is often detectable at the onset of parasitemia, with IgG detection following within 1-2 weeks. Due to the wide variety of antigens present on the whole organism by the IFA technique, sensitivity is approximately equal to Western immunoblot assay using whole cell lysates<sup>4</sup>. Sensitivity of the igM test is 91% and IgG 39% in the acute phase. In convalescent phase sera the sensitivity rises to a range of 88-96%, depending upon the testing center performing the assay<sup>1-4</sup>..

**SPECIFICITY**

There have been no reports of cross-reactivity in the IFA procedure, with the exception of related *Babesia* spp. Specificity has been reported as 90-100% in a comparison of test centers<sup>2</sup>, with IgM reported as 99% specific<sup>3</sup>. Sera from a non-endemic region were tested in-house, 94 from Southern California. There were no positives (100% specific).

**References**

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2. Krause, P.J., S.R. Telford III, R. Ryan, P.A. Conrad, M. Wilson, J.W. Thomford, and A. Spielman. 1994. Diagnosis of babesiosis: evaluation of a serologic test for the detection of *Babesia microti* antibody. *J. Infect. Dis.* 169:923-926.
3. Krause, P.J., R. Ryan, S.R. Telford III, D.H. Persing, and A. Spielman. 1996. Efficacy of an immunoglobulin M serodiagnostic test for the rapid diagnosis of acute babesiosis. *J. Clin. Microbiol.* 34:2014-2016.
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