SUPPORTING INFECTIOUS DISEASE RESEARCH

## Babesia duncani, WA1 (in vitro)

## Catalog No. NR-50440

**Product Description:** Babesia duncani (B. duncani), WA1 was isolated from human blood from the first reported case of babesiosis acquired in Washington, USA, and adapted to continuous *in vitro* culture in human erythrocytes.

## Lot<sup>1</sup>: 70004225

## Manufacturing Date: 28APR2017

TEST	SPECIFICATIONS	RESULTS
Cellular Morphology <sup>2</sup>	Report results	Single, double and tetrad forms
Genotyping <sup>3</sup> Sequencing of 18S ribosomal RNA (rRNA) gene (~ 680 base pairs)	Consistent with <i>B. duncani</i>	Consistent with <i>B. duncani</i>
Functional Activity by PCR Amplification <sup>3,4</sup> 18S rRNA gene	~ 930 base pair amplicon	~ 930 base pair amplicon
Level of Parasitemia <sup>3,5</sup>	Report results	13%
Viability <sup>2,6</sup>	Growth	Growth
Mycoplasma Contamination <sup>3</sup> DNA Detection by PCR	None detected	None detected

<sup>1</sup>NR-50440 was produced by cultivation of the deposited material in human erythrocytes with *Babesia* Growth Medium (HL-1<sup>™</sup> Chemically Defined, Serum-Free Medium (Lonza 77201), adjusted to contain 20% Human Serum Type A Positive, 1% HB 101<sup>®</sup> supplement (Irvine Scientific<sup>®</sup> T151), 2 mM L-glutamine, 200 µM hypoxanthine, 32 µM thymidine, 100 IU/mL penicillin, 100 µg/mL streptomycin and 0.25 µg/mL amphotericin B and 100 µg/mL gentamicin). After a series of passages, the culture was propagated in human erythrocytes with *Babesia* Growth Medium for 1 day at 37°C in a humidified atmosphere of 93% N<sub>2</sub>, 5% CO<sub>2</sub>, 2% O<sub>2</sub> until the first peak of parasitemia was reached.

<sup>2</sup>Testing completed on vialed, post-freeze material.

<sup>3</sup>Testing completed on bulk material prior to vialing and freezing.

<sup>4</sup>Primer sequences and conditions for PCR are available upon request.

<sup>5</sup>Parasitemia was determined after 1 day of infection by microscopic counts of Giemsa-stained blood smears.

<sup>6</sup>Viability of the material following cryopreservation was determined by cultivation in human erythrocytes with *Babesia* Growth Medium at 37°C in a humidified atmosphere of 93% N<sub>2</sub>, 5% CO<sub>2</sub>, 2% O<sub>2</sub> and examination of parasitemia every day for 4 days post-infection (19% parasitemia).

Date: 29 JAN 2018

Signature:

BEI Authentication or designee

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