

## **Certificate of Analysis for MRA-919**

## 4a-3B, Anopheles gambiae Cell Line

Catalog No. MRA-919

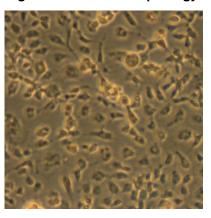
**Product Description:** The *Anopheles gambiae* cell line 4a-3B was established from neonate larvae of the 4a r/r strain.

Lot<sup>1</sup>: 70000595 Manufacturing Date: 09DEC2016

TEST	SPECIFICATIONS	RESULTS
Growth Properties	Adherent monolayer	Adherent monolayer
Cellular Morphology <sup>2</sup>	Adherent monolayer	Adherent monolayer (Figure 1)
Cell Count (pre-freeze)	≥ 1.0 × 10 <sup>6</sup> cells per vial	1.7 × 10 <sup>7</sup> cells per vial
Post-Freeze Viability	≥ 75% viable cells	78.9% viable cells
Sterility (21-day incubation) Harpo's HTYE broth³, 37°C and 26°C, aerobic Tryptic Soy broth, 37°C and 26°C, aerobic Sabouraud Dextrose broth, 37°C and 26°C, aerobic DMEM with 10% FBS, 37°C, aerobic Sheep blood agar, 37°C, aerobic Sheep blood agar, 37°C, anaerobic Thioglycollate broth, 37°C, anaerobic	No growth	No growth
Mycoplasma Contamination DNA Detection by PCR	None detected	None detected

MRA-919 was produced by cultivation of MR-MRA-919 lot 58422736 in Schneider's *Drosophila* medium supplemented with 10% fetal bovine serum (Gemini Bio-Products 100-135; insect cell qualified), penicillin (100 U/mL) and streptomycin (100 U/mL) at 25°C in sealed flasks for 9 days, after which the cells reached 80%-90% confluency and were harvested. Every 1 to 4 days, media was replaced with fresh culture medium. Cells were reseeded on day 1 (1:6 split), day 5 (1:3 split) and day 7 (1:2 split).

Figure 1: Cellular Morphology



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<sup>&</sup>lt;sup>2</sup>12 days at 25°C in Schneider's *Drosophila* medium supplemented with 10% fetal bovine serum, penicillin (100 U/mL) and streptomycin (100 U/mL) Atlas, Ronald M. <u>Handbook of Microbiological Media</u>. 3rd ed. Ed. Lawrence C. Parks. Boca Raton: CRC Press, 2004, p. 798.



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Date: 31 JAN 2017

Signature:

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