SUPPORTING INFECTIOUS DISEASE RESEARCH

Human Respiratory Syncytial Virus, B1

Catalog No. NR-56243

Product Description:

Human respiratory syncytial virus (RSV), B1 was developed by multiple passages in Vero cells from an original human isolate in 1985, in West Virginia, USA. NR-56243 lot 70062310 was produced by infecting *Chlorocebus* (formerly *Cercopithecus*) *aethiops* kidney epithelial cells (Vero; ATCC[®] CCL-81[™]) and incubating in Eagle's Minimum Essential Medium (ATCC[®] 30-2003[™]) supplemented with 2% fetal bovine serum (ATCC[®] 30-2020[™]) for 6 days at 37°C with 5% CO₂.

Passage History:

V(Unk)/V(2) (Prior to BEI Resources/BEI Resources); V= Cercopithecus aethiops kidney cells; Unk = Unknown

Lot: 70062310

Manufacturing Date: 14AUG2023

TEST	SPECIFICATIONS	RESULTS
Identification by Infectivity in Vero Cells	Cell rounding and detachment	Cell rounding and detachment
Identification by Fluorescent Antibody (FA) Assay ¹	Fluorescence observed	Fluorescence observed
Sequencing of Species-Specific Region (~ 810 nucleotides)	≥ 98% identity with RSV, B1 (GenBank: AF013254)	99.9% identity with RSV, B1 (GenBank: AF013254)
Titer by TCID₅0 Assay in Vero Cells by Fluorescent Antibody Assay ^{1,2} (8 days at 37°C with 5% CO ₂)	Report results	1.6 × 10 ⁶ TCID ₅₀ /mL
Sterility (21-day incubation)		
Harpo's HTYE broth, 37°C and 26°C, aerobic ³	No growth	No growth
Trypticase Soy broth, 37°C and 26°C, aerobic	No growth	No growth
Sabouraud broth, 37°C and 26°C, aerobic	No growth	No growth
Sheep blood agar, 37°C, aerobic	No growth	No growth
Sheep blood agar, 37°C, anaerobic	No growth	No growth
Thioglycollate broth, 37°C, anaerobic	No growth	No growth
DMEM with 10% FBS, 37°C, aerobic	No growth	No growth
Mycoplasma Contamination		
Agar and broth culture (14-day incubation at 37°C)	None detected	None detected
DNA detection by PCR of extracted Test Article nucleic acid	None detected	None detected

²The Tissue Culture Infectious Dose 50% (TCID₅₀) endpoint is the 50% infectious endpoint in cell culture. The TCID₅₀ is the dilution of virus that under the conditions of the assay can be expected to infect 50% of the culture vessels inoculated, just as a Lethal Dose 50% (LD₅₀) is expected to kill half

the conditions of the assay can be expected to infect 50% of the culture vessels inoculated, just as a Lethal Dose 50% (LD₅₀) is expected to kill half of the animals exposed. A reciprocal of the dilution required to yield the TCID₅₀ provides a measure of the titer (or infectivity) of a virus preparation. ³Atlas, 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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