

Certificate of Analysis for NR-44229

Genomic RNA from Human Respiratory Syncytial Virus, A1998/3-2

Catalog No. NR-44229

Product Description: Genomic RNA was isolated from a preparation of cell lysate and supernatant from HEp-2 cells¹ infected with human respiratory syncytial virus (RSV), A1998/3-2.

Lot²: 61968413 Manufacturing Date: 27AUG2013

TEST	SPECIFICATIONS	RESULTS
Sequencing of Species-Specific Region (820 nucleotides)	Consistent with human RSV, A1998/3-2 virus Identical to NR-28529	99% identity with human RSV, A1998/3-2 virus (GenBank: JX069801) Identical to NR-28529
Functional Activity by RT-PCR Amplification ³	~ 900 bp amplicon	~ 900 bp amplicon (See Figure 1)
Total RNA Content by RiboGreen [®] Measurement (Viral, Cellular and Carrier)	Report results	153 ng per 100 μL
Virus Inactivation 10% of total yield inoculated on HEp-2 cells ¹ and evaluated for cytopathic effect and expression of viral antigens ^{4,5}	No virus detected	No virus detected

¹HEp-2 cells: ATCC[®] CCL-23™

Figure 1 Base Pairs 1500 --- 900 bp amplicon

Date: 04 NOV 2013 Signature: Minhael Q. Gyyrla

Title: Technical Manager, BEI Authentication or designee

ATCC[®], on behalf of BEI Resources, hereby represents and warrants that the material provided under this certificate has been subjected to the tests and procedures specified and that the results described, along with any other data provided in this certificate, are true and accurate to the best of ATCC[®]'s knowledge.

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²Nucleic acid was extracted from a preparation of Human Respiratory Syncytial Virus, A1998/3-2 (BEI Resources NR-28529, Lot 59927457), using a QIAamp[®] Viral RNA Mini Kit (Qiagen 52906).

³Amplified using iScript™ One-Step RT-PCR Kit with SYBR® Green (Bio-Rad 170-8892) with 2 µL of NR-44229 in a 25 µL reaction

⁴Using LIGHT DIAGNOSTICS™ Respiratory Syncytial Virus FITC Reagent (Millipore 5022)

⁵Use of the QIAamp[®] Viral RNA Mini Kit has been demonstrated to consistently inactivate 100% of human respiratory syncytial virus, as shown by the absence of cytopathic effect (CPE) and expression of viral antigens after plating the entire extract on virus-susceptible cells.