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Viral Filtration Efficiency (VFE) Final Report

Test Article: RESPILON RFT
Purchase Order: 20200401001
Study Number: 1307230-S01
Study Received Date: 05 Jun 2020

Testing Facility: Nelson Laboratories, LLC

6280 S. Redwood Rd.

Salt Lake City, UT 84123 U.S.A.

Test Procedure(s): Standard Test Protocol (STP) Number: STP0007 Rev 16

Deviation(s): None

Summary: The VFE test is performed to determine the filtration efficiency of test articles by comparing the viral control counts upstream of the test article to the counts downstream. A suspension of bacteriophage Φ X174 was aerosolized using a nebulizer and delivered to the test article at a constant flow rate and fixed air pressure. The challenge delivery was maintained at 1.1 - 3.3 x 10³ plaque forming units (PFU) with a mean particle size (MPS) of 3.0 µm ± 0.3 µm. The aerosol droplets were drawn through a six-stage, viable particle, Andersen sampler for collection. The VFE test procedure was adapted from ASTM F2101.

All test method acceptance criteria were met. Testing was performed in compliance with US FDA good manufacturing practice (GMP) regulations 21 CFR Parts 210, 211 and 820.

Test Side: Inside
Test Area: ~40 cm²

VFE Flow Rate: 28.3 Liters per minute (L/min)

Conditioning Parameters: $85 \pm 5\%$ relative humidity (RH) and 21 ± 5 °C for a minimum of 4 hours

Positive Control Average: 2.5 x 10³ PFU Negative Monitor Count: <1 PFU

MPS: 3.0 µm





Leah Tiberius electronically approved for

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22 Jul 2020 23:47 (+00:00)

Study Completion Date and Time

Study Director

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Results:

Test Article Number	Percent VFE (%)
1	>99.9
2	99.8
3	99.7
4	>99.9 ^a
5	>99.9 ^a

^a There were no detected plaques on any of the Andersen sampler plates for this test article.

The filtration efficiency percentages were calculated using the following equation:

$$\% VFE = \frac{C - T}{C} x \ 100$$

C = Positive control average

T = Plate count total recovered downstream of the test article Note: The plate count total is available upon request

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