

Sponsor: Amy Orrell Mazza Healthcare LLC 2101 Waukegan Rd. Bannockburn, IL 60015

Viral Filtration Efficiency (VFE) Final Report

Test Article: Purchase Order: Study Number:		
Study Received Date:		
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:InsideTest Area:~40 cm²VFE Flow Rate:28.3 Liters per minute (L/min)Conditioning Parameters:85 ± 5% relative humidity (RH) and 21 ± 5°C for a minimum of 4 hoursPositive Control Average:2.5 x 10³ PFUNegative Monitor Count:<1 PFU</td>MPS:3.0 μm



Sarah Guzman electronically approved for

Study Director

James Luskin

11 Jun 2020 21:33 (+00:00) Study Completion Date and Time

kxh



Results:

Test Article Number	Percent VFE (%)	
1	99.5	
2	99.3	
3	98.9	
4	99.3	
5	99.3	

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