

STUDY REPORT

Study Title

Antibacterial Activity and Aerosol Efficacy of Scientific Air Management's Device

Test Method Custom Aerosol Study

Study Identification Number NG6997

<u>Test Facility</u> MicrochemLaboratory 1304 W.Industrial Blvd Round Rock, TX 78681 (512) 310-8378



Test Device Information

The test device was received on 10 MAR 2016.



(note: photos depict the test device analyzed in this study) Test device received: Scientific Air Management SAM 400



Test MicroorganismInformation

The following test microorganisms were selected for this test:



MS2 Bacteriophage (MS2), 15597-B1

This virus is a non-enveloped positive-stranded RNA virus of the bacteriophage family Leviviridae. Bacterial cells are the hosts for bacteriophages, and *E. coli* 15597 serves this purpose for MS2 bacteriophage. Its small size, icosohedral structure, and environmental resistance has made MS2 ideal for use as a surrogate virus (particularly in place of picornaviruses such as poliovirus and human norovirus) in water quality and disinfectant studies.

Permissive Host Cell System for MS2: Escherichia coli, 15597





Summary of the Procedure

- Bacterial cultures, fungal culture, and a virus stock are pooled to target concentrations as appropriate.
- The test inoculum is split into two equal parts and added to two nebulizers. Liquid culture should not exceed 18ml per nebulizer.
- The chamber is setup and the safety check list is completed prior to test initiation.
- The nebulizers are activated for 60 minutes to ensure target microbial concentrations are achieved prior to activation of the device.
- AnSKC bio-samplerisused to take a time zero sample to determine starting chamber concentration for baseline comparison.
- Device is activated for the study sponsor determined contact time. After each contact time, the SKC bio-samplers are activated for 10 minutes to determine microbial concentrations.
- Samples are enumerated using standard dilution and plating techniques.
- Microbial concentrations are determined after 24-48 hours of incubation for bacteria and viruses. Fungal plates are incubated at room temperature for 5-7 days.
- Reductions of microorganisms are calculated relative to concentration at Time Zero.

Study Timeline



Amended report delivered 31 AUG 2016



Criteria for Scientific Defensibility of a Custom Device Study

For Microchem Laboratory to consider a DeviceStudy to be scientifically defensible, the following criteria must be met:

- 1. The average number of viable bacteria recovered from the time zero samples must be approximately 1×10^5 cells/m³ or greater.
- 2. Positive/Growth controls must demonstrate growth of the appropriate test microorganism.
- 3. Negative/Purity controls must demonstrate no growth of test microorganism.

Passing Criteria

Because of the nature of the study, passing criteria may be determined by the Study Sponsor.

Testing Parameters used in this Study

Volume of Inoculum added to Nebulize <u>r</u>	15 ml per nebulizer (30 ml total)		
SKC Biosampler Media (Vol.)	Phosphate Buffered Saline (20 ml)		
Nebulization Time	60 minutes		
SKC Biosampler Time	10 minutes		
Sampling Time Points	Time zero, 5 minutes 15 minutes, 30 minutes		
SKC Biosampler Sampling Rate	12.5 L/minute		
SKC Biosampler Liters Sampled	125 L sampled per sampling time point		

TestMicroorganism_	MS2 Bacteriophage		
CultureGrowthMedia	N/A Stock Solution		
Culture Growth Time	N/A Stock Solution		
CultureDilutionMedia	Phosphate buffered saline		
TargetConcentration	≥1.0 x 10 ⁶ CFU/m ³		
EnumerationMedia	50% Tryptic SoyAgar		
EnumerationType	Poured with <i>E. coli</i> 15597		
EnumerationIncubationTime	18-24 hours		



Study Notes

No additional observations or notations were made for this study.

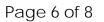
Study Photographs





Photographs are the device received by the laboratory on 10 MAR 2016. The device is photographed operating within the chamber and the SKC bio-sampler in the foreground.







Control Results

Neutralization Method:Notapplicable Growth Confirmation: Confirmed

Media Sterility: Sterile

Calculations

Percent Reduction = $(\frac{B-A}{B}) \times 100$

Where:

B = Number of viable test microorganisms on the control carriers immediately after inoculation A = Number of viable test microorganisms on the test carriers after the contact time

$$Log_{10}Reduction = Log(\frac{B}{A})$$

Where:

B = Number of viable test microorganisms on the control carriers immediately after inoculation A = Number of viable test microorganisms on the test carriers after the contact time

$$CFU/m^{3}=1000 \times \left(\frac{\frac{CFU}{ml}x(V_{s})}{T_{s}(12.5)}\right)$$

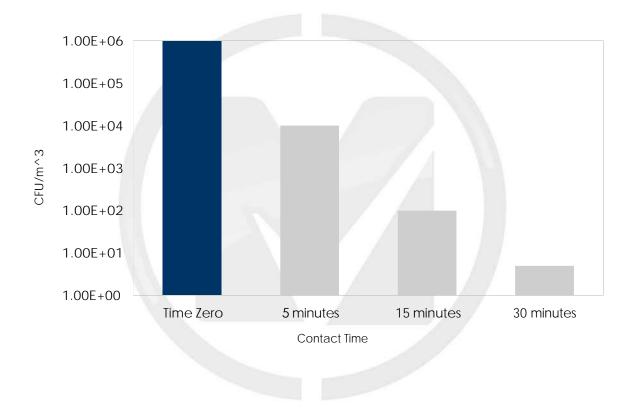
Where: V_s = Bio-sampler volume (ml) T_s = Time sampled (min)

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Results of the Study

Test Device	Microorganism	Inoculum Concentration (CFU/ml)	Treatment Time Point	Recovery (CFU/m³)	Percent Reduction vs. Time Zero	Log ₁₀ Reduction vs. Time Zero
Scientific Air Manage- ment \$400 MS2 Bacteriophage ATCC 15597-B1	1.00E+07	Time Zero	1.00E+06	N/A		
		5 minutes	1.00E+04	99.0%	2.00	
		15 minutes	1.00E+02	99.990%	4.00	
			30 minutes	5.00E+00	99.9995%	5.30



The results of this study apply to the tested substances (s) only. Extrapolation of findings to related materials is the responsibility of the Sponsor.

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