

Sponsor: Healthy Breath Ltd 4D Pacific Rise Mt Wellington Auckland 1060 New Zealand

Bacterial Filtration Efficiency (BFE) at an Increased Challenge Level Final Report

Test Article: MEO Kids

Study Number: 1067190-S01A.1 Amended

Study Received Date: 28 Jun 2018 Study Completion Date: 18 Jul 2018

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: STP0009 Rev 10

Deviation(s): None

Summary: This test procedure was performed to evaluate the BFE of test articles at an increased challenge level. A suspension of *Staphylococcus aureus*, ATCC #6538, was delivered to the test article at a challenge level of greater than 10⁶ colony forming units (CFU). The challenge was aerosolized using a nebulizer and delivered to the test article at a fixed air pressure and flow rate of 30 liters per minute (LPM). The aerosol droplets were generated in a glass aerosol chamber and drawn through the test article into all glass impingers (AGIs) for collection. The challenge was delivered for a one minute interval and sampling through the AGIs was conducted for two minutes to clear the aerosol chamber. The mean particle size (MPS) control was performed at a flow rate of 28.3 LPM using a six-stage, viable particle, Andersen sampler for collection.

This test procedure was modified from Nelson Laboratories, LLC (NL), standard BFE procedure in order to employ a more severe challenge than would be experienced in normal use. This method 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.

Challenge Flow Rate: 30 LPM

Area Tested: ~40 cm²

Side Tested: Side Facing Up in Bag

Challenge Level: 3.3 x 106 CFU

MPS: ~2.9 µm

Test Monitor Results: Acceptable

Study Director

Janelle R. Bentz, M.S.

Amended Report Date

1067190-S01

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bsm

FRT0009-0001 Rev 11

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Results: MEO Kids:

Test Article Number	Total CFU Recovered	Filtration Efficiency (%)
1	1.2×10^2	99.9964
2	4.7×10^{1}	99.9986
3	7.2×10^{1}	99.9978

The filtration efficiency percentages were calculated using the following equation:

$$\% \ BFE = \frac{C-T}{C} \ x \ 100$$
 C = Challenge Level T = Total CFU recovered downstream of the test article

Amendment Justification: At the request of the sponsor, results were separated in to reports organized per sample ID.

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Sponsor: Healthy Breath Ltd 4D Pacific Rise Mt Wellington Auckland 1060 New Zealand

Bacterial Filtration Efficiency (BFE) at an Increased Challenge Level Final Report

Test Article: MEO Lite

Study Number: 1067190-S01B.1 Amended

Study Received Date: 28 Jun 2018 Study Completion Date: 18 Jul 2018

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: STP0009 Rev 10

Deviation(s): None

Summary: This test procedure was performed to evaluate the BFE of test articles at an increased challenge level. A suspension of *Staphylococcus aureus*, ATCC #6538, was delivered to the test article at a challenge level of greater than 10⁶ colony forming units (CFU). The challenge was aerosolized using a nebulizer and delivered to the test article at a fixed air pressure and flow rate of 30 liters per minute (LPM). The aerosol droplets were generated in a glass aerosol chamber and drawn through the test article into all glass impingers (AGIs) for collection. The challenge was delivered for a one minute interval and sampling through the AGIs was conducted for two minutes to clear the aerosol chamber. The mean particle size (MPS) control was performed at a flow rate of 28.3 LPM using a six-stage, viable particle, Andersen sampler for collection.

This test procedure was modified from Nelson Laboratories, LLC (NL), standard BFE procedure in order to employ a more severe challenge than would be experienced in normal use. This method 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.

Challenge Flow Rate: 30 LPM

Area Tested: ~40 cm²

Side Tested: Yellow Stickered Side

Challenge Level: 3.3 x 10⁶ CFU

MPS: ~2.9 µm

Test Monitor Results: Acceptable

Study Director

Janelle R. Bentz, M.S.

Amended Report Date

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Results: MEO Lite:

Test Article Number	Total CFU Recovered	Filtration Efficiency (%)
1	1.1×10^{1}	99.99967
2	1	99.999970
3	1.2×10^{1}	99.99963

The filtration efficiency percentages were calculated using the following equation:

%
$$BFE = \frac{C - T}{C} \times 100$$
 C = Challenge Level T = Total CFU recovered downstream of the test article

Amendment Justification: At the request of the sponsor, results were separated in to reports organized per sample ID.

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Sponsor: Healthy Breath Ltd 4D Pacific Rise Mt Wellington Auckland 1060 New Zealand

Bacterial Filtration Efficiency (BFE) at an Increased Challenge Level Final Report

Test Article: MEO X

Study Number: 1067190-S01C.1 Amended

Study Received Date: 28 Jun 2018 Study Completion Date: 18 Jul 2018

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: STP0009 Rev 10

Deviation(s): None

Summary: This test procedure was performed to evaluate the BFE of test articles at an increased challenge level. A suspension of *Staphylococcus aureus*, ATCC #6538, was delivered to the test article at a challenge level of greater than 10⁶ colony forming units (CFU). The challenge was aerosolized using a nebulizer and delivered to the test article at a fixed air pressure and flow rate of 30 liters per minute (LPM). The aerosol droplets were generated in a glass aerosol chamber and drawn through the test article into all glass impingers (AGIs) for collection. The challenge was delivered for a one minute interval and sampling through the AGIs was conducted for two minutes to clear the aerosol chamber. The mean particle size (MPS) control was performed at a flow rate of 28.3 LPM using a six-stage, viable particle, Andersen sampler for collection.

This test procedure was modified from Nelson Laboratories, LLC (NL), standard BFE procedure in order to employ a more severe challenge than would be experienced in normal use. This method 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.

Challenge Flow Rate: 30 LPM

Area Tested: ~40 cm²

Side Tested: Side Facing Up in Bag

Challenge Level: 3.3 x 10⁶ CFU

MPS: ~2.9 μm

Test Monitor Results: Acceptable

Study Director

Janelle R. Bentz, M.S.

Amended Report Date

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Results:

MEO X:

Test Article Number	Total CFU Recovered	Filtration Efficiency (%)
1	7.6×10^{2}	99.977
2	5.0×10^2	99.985
3	5.9×10^{2}	99.982

The filtration efficiency percentages were calculated using the following equation:

Amendment Justification: At the request of the sponsor, results were separated in to reports organized per sample ID.

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