

# Efficacy of ecoVIOX Air Purifier in Removing Microbes

**PACS ID#:** 07204

Work Order#: 026808

Customer: Lucia Bonino

**Dates of Testing:** 03/20/2021

**Date Completed: 03/24/2021** 

**Date of Report: 03/25/2021** 

# **Environmental Diagnostics Laboratory**

A Division of Pure Air Control Services, Inc.

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March 25, 2021

To

Lucia Bonino ecoVIOX LLC 1994 NE 147<sup>th</sup> Terrace North Miami, FL 33181

Reference: PACS ID 07204 Work Order 026808

Efficacy Testing for ecoVIOX LLC

Dear Lucia Bonino,

We appreciate the opportunity to provide you with our professional environmental microbiology services. EDLab is pleased to submit this report that describes the efficacy testing for ecoVIOX.

This report summarizes the findings and other relevant data based on experiments set-up for efficacy evaluation of ecoVIOX.

Please call me at 1-800-422-7873, ext. 301 should you have any questions. We look forward in assisting you in future projects.

Respectfully submitted,

Kasandra Cuellar

Staff Microbiologist

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Laboratory Director













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PACS ID: 07204 Work Order: - 026808

Sample Numbers/Lab Numbers:001/193344

**Product description:** This system is equipped with a charcoal activated HEPA filter, UVC lamp, ionizer, along with an ozone generator.

# **Test Challenge Organisms:**

Bacteria: Staphylococcus aureus

Fungi: Aspergillus niger

**TEST:** Microbial Removal Efficacy

**Methodology:** To determine the microbial removal efficacy of an air purification device. A known concentration of the selected challenge organisms Staphylococcus aureus and Aspergillus niger were aerosolized through the device. Air samples were collected both when the device was switched off (device not running) and switched-on (all components of the device were running) utilizing the settling plate method at two different time intervals (0 and 60 minutes) with Typic Soy Agar (TSA) and Malt Agar Extract (MEA) microbial culture media for S. aureus and A. niger, respectively. Samples for pretest sterility verification of the test chamber was collected in addition to the negative and positive control samples. The viability and concentration of the suspension utilized for the aerosolization of the test organisms (S. aureus and A niger) was determined by serial dilution technique.

#### **Controls:**

# **Negative Controls:**

Reagent Water Lot:476425 Exp:06-26-2021

#### **Positive Controls:**

Staphylococcus aureus ATTC: 25923 Exp:06-30-2021 Aspergillus niger ATTC:6275 Exp:01-31-2022





# Microbiological media used:

TSA= Lot #: 135372 Exp:06-07-2021 MEA= Lot #:47409 Exp:05-17-2021

## **Challenge concentration:**

Staphylococcus aureus 50.8 x 10<sup>4</sup>

Aspergillus niger  $25.0 \times 10^2$ 

#### **Observations:**

All the experiments were conducted under standard laboratory conditions. *Table - 1* records the sterility verification of the environmental test chamber. *Table - 2* contains the results of the efficacy of the device tested.

Table – 1: Sterility Verification

Trial	Bacteria	Fungi
Sterility Verification	BDL	BDL

<sup>\*</sup>BDL=Below Detection Limit

Table – 2: Microbiological Efficacy Results

Challenge Organisms	Pre-treatment		Post-treatment	
	Colonies (CFU/m²) 0 min	Colonies (CFU/m²) 60 min	Colonies (CFU/m²) 0 min	Colonies (CFU/m²) 60 min
S. aureus	899,422	373	2,986	BDL
A. niger	8,584	1,866	187	BDL





Figure – 1 Efficacy on Microbes

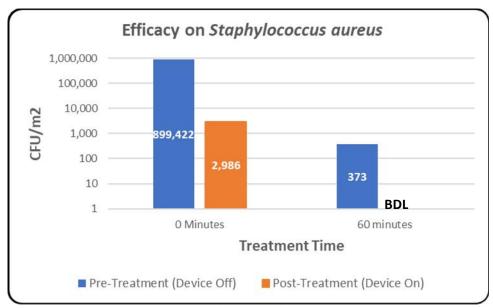


Figure – 1A

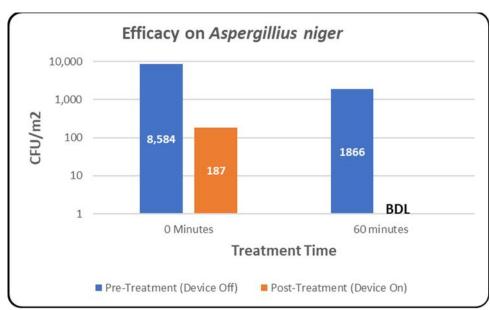


Figure – 1B
\*BDL= Below Detection Limit





### Conclusion

This device is rated as effective against the microbes (Staphylococcus aureus and Aspergillus niger). A more comprehensive study is encouraged to understand the details on the reduction efficacy of this air purifying device.

Note: No other testing or evaluation has been performed, except for those mentioned above

Analyst:

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