





Date Received: Apr. 08, 2020

Name of Sample	Air Purifier	Source of Sample	Delivery
Applicant	Air Doctor LLC	Client	Sherry Huang
Manufacturer	Killian Salah Salah Amerikan	Brand	AirDoctor
Type and Specification	AD3000	Quantity of Sample	1Set (Two machine and a set of filters
Date of Production	- 6	State of Sample	Machine
Batch Number	- 12 M	Packing of Sample	In box
		AIRDoctor	
Sample Picture			
Sample Picture Standard and Methods	GB 21551.3-2010 Antibacter electrical appliances-Particular	rial and cleaning function for	household and simila
	electrical appliances-Particular 1. Eliminating Bacterial Rat ATCC 6538, Escherichia	rial and cleaning function for requirements of air cleaner e (Staphylococcus albus 8032, Stacoli 8099, Klebsiella pneumoniae, P.Aeruginosa ATCC 15442, CanaATCC 16404)	phylococcus aureus ATCC 4352, Serratia







Date Received: Apr. 08, 2020 Date Analyzed: Apr. 09, 2020

Test Method for Air Purifier Disinfection Performance:

- 1. Test Equipment
 - 1) Strain: Staphylococcus albus, Staphylococcus aureus, Escherichia coli, Klebsiella pneumoniae, Serratia marcescens, P.Aeruginosa, Candida albicans
 - 2) Microbial aerosol generator: TK-3
 - 3) Culture media: NA, SDA
 - 4) Sampling equipment: six-stage sieve sampler
- 2. Test Conditions
 - 1) The volume of the test chamber: 30 m³
 - 2) Environment temperature: (20~25) ℃
 - 3) Environment humidity: (50~70) %RH
- 3. Operation Conditions of the Air Purifier

Set the switch to position "The highest wind speed", "negative ions".

- 4. Test Procedure
 - Get a slant culture (4~5 generation) which is incubated at 37 °C for 24 h, wash the culture from this slant with 10 mL broth, filter the liquid culture by aseptic cotton buds, and dilute this inoculums with broth as appropriate.
 - 2) The equipments are placed in the test chambers, close the door, and turn on the HEPA filter system. Simultaneously operate the environmental control devices until the temperature reaches (20~25)°C, relative humidity reaches (50~70)%. Turn off the chamber environmental control system.
 - 3) Release microbial aerosol: turn on the microbial aerosol generator, then turn on the ceiling fan, turn off the fan after 10 min, and let stand for 15 min.
 - 4) Original bacteria aerosols collected by six-stage sieve sampler.
 - 5) The air purifier is adjusted to the highest air cleaning mode setting for test (test group). Bacteria aerosols (control group and test group) are collected at 60 min.
 - 6) Choose 2 NA plates (the same batch) as the negative control, and culture them on the same condition with the samples.
 - 7) Run the test three times and take the mean as the final result.
- 5. Computational Formula

Natural decay rate
$$N_t(\%) = \frac{V_0 - V_t}{V_0} \times 100$$

Where: V_0 = original bacteria count of control group; V_t = bacteria count after treatment of control group.

Eliminating Bacterial Rate
$$K_t$$
 (%) = $\frac{V_1 \times (1 - N_t) - V_2}{V_1 \times (1 - N_t)} \times 100$

Where: V_1 = original bacteria count of test group; V_2 = bacteria count after treatment of test group. ***To be continued***







Date Received: Apr. 08, 2020 Date Analyzed: Apr. 09, 2020

Test results

				C	ontrol Group		Test	Group	Eliminating
Number of T	Test Time (min)	Test Bacteria	Test Number	Original Bacteria Count V_0 (cfu/m 3)	Bacteria Count after Treatment V_t (cfu/m 3)	Natural Decay Rate N_t (%)	Original Bacteria Count V1 (cfu/m³)	Bacteria Count after Treatment V ₂ (cfu/m ³)	Bacterial Rate K_t (%)
			1	1.33×10 ⁵	1.03×10 ⁵	22.56	1.34×10 ⁵	7	99.99
		Staphylococcus	2	1.28×10 ⁵	1.01×10 ⁵	21.09	1.25×10 ⁵	7	99.99
		albus	3	1.36×10 ⁵	1.10×10 ⁵	19.12	1.39×10 ⁵	7	99.99
			Mean			(A) 40			99.99
. (13)			1	1.11×10 ⁵	8.93×10 ⁴	19.55	1.26×10 ⁵	7	99.99
J20201603-1		Staphylococcus	2	1.21×10 ⁵	9.79×10 ⁴	19.09	1.18×10 ⁵	7	99.99
.J20201003-1	60	aureus	3	1.15×10 ⁵	9.15×10 ⁴	20.43	1.05×10 ⁵	7	99.99
			Mean						99.99
			1	1.44×10 ⁵	9.52×10 ⁴	33.89	1.37×10 ⁵	7	99.99
		Escherichia	2	1.39×10 ⁵	9.35×10 ⁴	32.73	1.28×10 ⁵	7	99.99
		coli	3	1.47×10 ⁵	1.02×10 ⁵	30.61	1.41×10 ⁵	7	99.99
			Mean	1. 16				0	99.99

Note: The negative control group was sterile growth.

To be continued







Date Received: Apr. 08, 2020 Date Analyzed: Apr. 09, 2020

Test results

1est results									
	5	-		Co	ontrol Group)	Test	Group	
Number of T	Test Time (min)	Test Bacteria	Test Number	Original Bacteria Count V_0 (cfu/m 3)	Bacteria Count after Treatment V_t (cfu/m 3)	Natural Decay Rate N _t (%)	Original Bacteria Count V1 (cfu/m³)	Bacteria Count after Treatment V ₂ (cfu/m ³)	Eliminating Bacterial Rate K_t (%)
			1	1.27×10 ⁵	9.45×10 ⁴	25.59	1.12×10 ⁵	7	99.99
		Klebsiella	2	1.22×10 ⁵	9.31×10 ⁴	23.69	1.23×10 ⁵	7	99.99
		pneumoniae	3	1.20×10 ⁵	8.99×10 ⁴	25.08	1.04×10 ⁵	7	99.99
		(0)	Mean		1000	N. Service			99.99
		Serratia	1	1.17×10 ⁵	8.37×10 ⁴	28.46	1.24×10 ⁵	7	99.99
			2	1.13×10 ⁵	7.75×10 ⁴	31.42	1.10×10 ⁵	7	99.99
		marcescens	3	1.21×10 ⁵	8.54×10 ⁴	29.42	1.33×10 ⁵	7	99.99
KJ20201603-1	60 –	Day.	Mean					10.	99.99
1820201003 1	- 00	, O	1	1.38×10 ⁵	1.07×10 ⁵	22.46	1.30×10 ⁵	7	99.99
		P. 4	2	1.33×10 ⁵	1.04×10 ⁵	21.80	1.42×10 ⁵	7	99.99
		P.Aeruginosa	3	1.40×10 ⁵	1.13×10 ⁵	19.29	1.39×10 ⁵	7	99.99
	_		Mean				1/19		99.99
			1	1.05×10 ⁵	7.37×10 ⁴	29.81	1.13×10 ⁵	7	99.99
		Candida	2	9.07×10 ⁴	6.61×10 ⁴	27.12	1.06×10 ⁵	7	99.99
		albicans	3	1.08×10 ⁵	7.51×10 ⁴	30.46	1.19×10 ⁵	7	99.99
			Mean			- (A)	V G		99.99

Note: The negative control group was sterile growth.







GUANG ZHOU INSTITUTE OF MICROBIOLOGY

TEST REPORT

Date Received: Apr. 08, 2020 Date Analyzed: Apr. 09, 2020

Air Disinfection Test Method:

1. Test Equipment

1) Strain: Aspergillus niger

Microbial aerosol generator: TK-3

3) Culture media: PDA

4) Sampling equipment: six-stage sieve sampler

2. Test Conditions

1) The volume of the test chamber: 30 m³

2) Environment temperature: (20~25) ℃

3) Environment humidity: (50~70) %RH

3. Operational Conditions of the Machine

Set the switch to position "The highest wind speed", "negative ions".

4. Test Procedure

- 1) To the 4th to 5th generation of Aspergillus niger roxell culture, add 5.0 ml to 10.0 ml of 0.05% (v / v) Tween 80 aqueous PBS solution, scrap the Aspergillus niger conidia in solution and transfer the spore suspension with glass beads in the flask, lightly shaking 1 min and filter removed hypha. Centrifuge 20min in the range of 5000r / min ~ 6000r / min . Then observe under the microscope (400 times), if there are still hypha in the suspension, to be centrifuged. Diluted with physiological saline solution to the appropriate concentration before use.
- 2) The equipments are placed in the test chambers respectively, close the door, and open the HEPA filter. Simultaneously operate the environmental control devices until the experimental cabin temperature to be (20~25) °C, relative humidity to be (50~70) %, Turn off the chamber environmental control system.
- 3) Release microbial aerosol: turn on the microbial aerosol generator, then turn on the ceiling fan, turn off the fan after 5 min, and let stand for 5 min.

4) Original Bacteria aerosols collected by six-stage sieve sampler.

- 5) The air purifier is adjusted to the highest air cleaning mode setting for test (test group), Bacteria aerosols (control group and test group) are collected at 60 min respectively.
- 6) Choose 2 PDA plates (the same batch) as the negative control, and culture them on the same condition with the samples.
- 7) Run the test three times and take the mean as the final result.

Computational Formula

Natural decay rate
$$N_t(\%) = \frac{V_0 - V_t}{V_0} \times 100$$

Where: V_0 = original bacteria count of control group; V_t = bacteria count after treatment of control group.

Eliminating Bacterial Rate
$$K_{t}(\%) = \frac{V_{1} \times (1 - N_{t}) - V_{2}}{V_{1} \times (1 - N_{t})} \times 100$$

Where: V_1 = original bacteria count of test group; V_2 = bacteria count after treatment of test group. ***To be continued***







Date Received: Apr. 08, 2020 Date Analyzed: Apr. 09, 2020

Test Results

				(Control Group		Test	Group	Eliminating
Number of Sample	Time		Test Number	Original Bacteria Count V_0 (cfu/m ³)	Bacteria Count after Treatment V_t (cfu/m 3)	Natural Decay Rate N, (%)	Original Bacteria Count V ₁ (cfu/m ³)	Bacteria Count after Treatment V_2 (cfu/m 3)	Bacterial Rate K_t (%)
			1	7.59×10 ⁴	5.27×10 ⁴	30.57	7.92×10 ⁴	7	99.99
W.100001.000.1	60	Aspergillus	2	7.76×10 ⁴	5.61×10 ⁴	27.71	8.35×10 ⁴	7	99.99
KJ20201603-1	60	niger	3	7.83×10 ⁴	5.79×10 ⁴	26.05	8.13×10 ⁴	7	99.99
			Mean			/ X /	9-	3	99.99

Note: The negative control group was sterile growth.

To be continued







GUANG ZHOU INSTITUTE OF MICROBIOLOGY

TEST REPORT

Date Received: Apr. 08, 2020 Date Analyzed: Apr. 09, 2020

Removal of Escherichia coli Phage Performance Test Evaluation Method:

1. Test Equipment

1) Strain: Escherichia coli Phage

2) Microbial aerosol generator: TK-3

3) Culture media: NA

2. Test Conditions

1) The volume of the test chamber: 30 m³

2) Environment temperature: (20~25) °C

3) Environment humidity: (50~70) %RH

3. Operational Conditions of the Machine

Set the switch to position "The highest wind speed", "negative ions".

4. Test Procedure

- Thawed the phage solution before test, diluted with sterile deionized water until liquid surface tension was 65 to 69×10^{-3} N/m.
- The equipments are placed in the test chambers respectively, close the door and open the HEPA filter. Simultaneously operate the environmental control devices, After reaching requirements and turn off the chamber environmental control system.
- 3) Open the aerosol generator, while stirring side of the bacteria, spray dyeing bacteria after the end of the fan to continue stirring 2 min, and then set aside for 2 min.
- 4) Open the sampling pump to collect the initial suspended phage from the air in the test chamber. Immediately after sampling, the prototype is to be tested and the timepiece is started.
- After 60 min of action, turn off the sample to be tested while opening the sampling pump and collecting the suspended phage from the air in the test compartment.
- 6) Run the test three times and take the mean as the final result.
- 7) Choose 2 NA plates (the same batch) as the negative control, and culture them on the same condition with the samples.

5. Computational Formula

Natural decay rate $N_i(\%) = \frac{V_0 - V_i}{V_0} \times 100$

Where: V_0 = original bacteria count of control group; V_t = bacteria count after treatment of control group.

Killing Rate $K_t(\%) = \frac{V_1 \times (1 - N_t) - V_2}{V_1 \times (1 - N_t)} \times 100$

Where: V_1 = original bacteria count of test group; V_2 = bacteria count after treatment of test group.







Date Received: Apr. 08, 2020 Date Analyzed: Apr. 09, 2020

Test Results

Number of Ti				C	Control Group		Test	Group	Virus Removal rate K _t (%)
	Test Time (min)	Test Strain	Test Number	Original Virus Count V_0 (cfu/m ³)	Virus Count after Treatment V_t (cfu/m 3)	Natural Decay Rate N, (%)	Original Virus Count V1 (cfu/m³)	Virus Count after Treatment V ₂ (cfu/m ³)	
			1	8.79×10 ⁴	5.83×10 ⁴	33.67	7.87×10 ⁴	7	99.99
KJ20201603-1	60	Escherichia coli Phage	2	7.23×10 ⁴	5.18×10 ⁴	28.35	7.55×10 ⁴	O 7	99.99
11020201005 1	00	1 nage	3	8.54×10 ⁴	5.96×10 ⁴	30.21	8.76×10 ⁴	7	99.99
		6	Mean						99.99

Note: The negative control group was sterile growth.

End of report

Editor A Checker

Issuer 4

Date Reported









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Contact Address, NO.1 Jiantashan Road, Huangpu District, Guangzhou City, Guangdong Province
Test Address, (only fill in when it's different from the contact address)

Postal Code, 510663

Tel., (8620)61302671

URL, http://www.ggtest.com.cn

Date Received: Jun. 17, 2020 Date Analyzed: Jun. 21, 2020

Name of Sample	Air purifier	Source of Sample	Delivery Michael pedersen	
Applicant	Air Doctor LLC	Client		
Manufacturer		Brand	AirDoctor	
Type and Specification	AD3000	Quantity of Sample	1PC	
Date of Production		State of Sample	Machine	
Batch Number		Packing of Sample	In box	
	and the second s			
Sample Picture	1 Referring to GR/T 18801-2015 Ai	AlROctor	30	
Sample Picture Standard and Methods	Referring to GB/T 18801-2015 Ai Referring to <technical effect="" evaluation="" standard="" td="" test<=""><td>r cleaner</td><td>2.1.3 Air disinfection</td></technical>	r cleaner	2.1.3 Air disinfection	
	2. Referring to <technical standard<="" td=""><td>r cleaner d For Disinfection> 2002-2</td><td>2.1.3 Air disinfection</td></technical>	r cleaner d For Disinfection> 2002-2	2.1.3 Air disinfection	

To be continued

Date Received: Jun. 17, 2020 Date Analyzed: Jun. 21, 2020

Test Method for Purification Effect of Airborne Virus Aerosols

- Test Equipment
 - Strain: Influenza A virus A/PR8/34 H1N1
 - Cells: MDCK
- **Test Conditions**
 - Environment temperature: (23~25) ℃
 - Environment relative humidity: (50~60) %
 - 3) Test time: 60 min
 - The volume of the test chamber: 30 m³ 4)
 - Machine setting: "The highest wind speed", "Negative ions". 5)

Test Results

			Virus Tite	er of Control Gro	oup	Virus Titer o	f Test Group	
Virus	Test Time (min)	Test Number	Original Concentration (TCID ₅₀ /m³)	Final Concentration (TCID ₅₀ /m³)	Natural Decay Rate (%)	Original Concentration (TCID ₅₀ /m³)	Final Concentration (TCID ₅₀ /m³)	Removal Rate (%)
a		1	3.69×10 ⁶	5.85×10 ⁵	84.15	2.49×10 ⁶	/	≥99.99
A/PR8/34 (H1N1)	60	2	1.17×10 ⁶	1.73×10 ⁵	85.21	3.69×10 ⁶	1	≥99.99
		3	1.98×10 ⁶	3.69×10 ⁵	81.36	5.46×10 ⁶	1	≥99.99

Note: "/" means not detected.

*** End of report***

Issuer '

Date Reported

Statements

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