





广州市微生物研究所有限公司

GUANG ZHOU INSTITUTE OF MICROBIOLOGY CO., LTD.

检测报告

TEST REPORT

Report Number

KJ20210566

Name of Sample

Air Purifier

Applicant

Healthy Air Technology Ltd.







GUANG ZHOU INSTITUTE OF MICROBIOLOGY CO., LTD. TEST REPORT

Date Received: Feb. 22, 2021

	- A.	Date Analyzed: Mar 01, 2021			
Name of Sample	Air Purifier	Source of Sample	Delivery		
Applicant	Healthy Air Technology Ltd.	Client	Yu Huang		
Manufacturer	Healthy Air Technology Ltd.	Brand	Healthy Air Technology		
Type and Specification	HA800	Quantity of Sample	1PC		
Date of Production		State of Sample	Machine		
Batch Number	202006	Packing of Sample	In box		
AN	V A A A A A A A A A A A A A A A A A A A				





3	1. GB/T 18801-2015 Air cleaner
Standard and Methods	2. GB 21551.3-2010 Antibacterial and cleaning function for household and similar
	electrical appliances-Particular requirements of air cleaner
Items of Analysis	Eliminating Bacterial Rate (Staphylococcus albus 8032)
Remarks	

To be continued







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Test Method for Air Purifier Eliminating Bacterial Performance:

- 1. Test Equipment
 - 1) Strain: Staphylococcus albus
 - 2) Microbial aerosol generator: TK-3
 - 3) Culture media: NA
 - 4) Sampling equipment: six-stage sieve sampler
- 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. Operation Conditions of the Machine Set the switch to position "Turbo".
- 4. Test Procedure
 - 1) Get a bacteria slant culture (4~5 generation) which is incubated at 37 °C for 24 h, wash the culture from this slant with 10 mL NB, filter the liquid culture by aseptic cotton buds, and dilute this inoculum with NB to suitable concentration. Then make atomized bacterial suspension.
 - 2) The equipment is placed in the two 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.
 - 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) At the same time, the test group and the control group were sampled with six-stage sieve sampler.
 - The test group started the sample and sampled after 60 min of action, and the control group also sampled in the corresponding time period.
 - 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***







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TEST REPORT

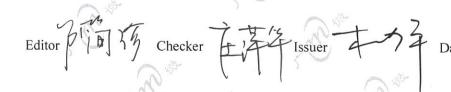
Date Received: Feb. 22, 2021 Date Analyzed: Mar 01, 2021

Test Results

9 % ·	30			Control Group		Test Group Eliminatin		Eliminating	
Number of	Test	Test Time	Toot	Original Bacteria	Bacteria Count after	Natural Decay	Original	Bacteria	Bacterial Rate
Sample	Strain	(min)	Mumbar	Count	Treatment V.	Rate	Bacteria Count	Count after Treatment	K_{ℓ}
- 0		-	w.	(cfu/m^3)	(cfu/m^3)	N_t (%)	V_1 (cfu/m ³)	V_2 (cfu/m ³)	(%)
	Se de la constant de		1	1.21×10 ⁵	1.14×10 ⁵	5.79	1.15×10 ⁵	<7	>99.99
KJ20210469-1	Staphylococcus albus	60	2	1.14×10 ⁵	1.09×10 ⁵	4.38	1.17×10 ⁵	7>1	>99.99
		, 000	3	1.12×10 ⁵	1.05×10 ⁵	6.25	1.05×10 ⁵	<7	>99.99
24			Mean		30		××-	50	>99.99

Note: The negative control group was sterile growth.

End of report





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Statements

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- 2. For the received sample, the sample information in the report is claimed by the applicant, the inspection unit is not responsible for its authenticity. The report is responsibility for the received sample only.
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- 6. The test data and results of items which are not accredited by CMA, only used associentific research, teaching or internal quality control.
- 7. Any ambiguity by the language which used in the report, the Chinese shall prevail

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