

Sky Xtra munnbind: Viktige ytelse egenskaper

Merk: Dette dokumentet er en høflighets oversettelse av det engelske hoved dokumentet tilgjengelig på https://www.flashbay.com/images/certificates/Sky Xtra Performance.pdf og hvis det er en forskjell i meningen mellom denne oversettelsen og hoved dokumentet skal betydningen av hoved dokumentet skal ha forrang.

Ytelsen til Sky Xtra munnbind mot funksjonell partikkel filtrering og krav til pusteevne i populære standarder for munnbind er uavhengig bestemt som følger:

	FFP2	CWA 17553:2020 - Level 90%
Filtration EN 149:2001+A1:2009, Clause 8.11 & AFNOR-SPEC-S76-001:2020, Reference to EN13274-7: 2019 Modified	PASS	PASS
Breathability EN 149:2001+A1:2009, Clause 8.9 & EN ISO 9237-1995	PASS	PASS

Testing mot FFP2 funksjonelle ytelseskrav

Sky Xtra munnbind har blitt uavhengig testet av NTEK mot de funksjonelle ytelseskravene til FFP2-standarden og det er bekreftet at den har følgende nøkkelegenskaper når den er ny:

	Requirement	Result*	
Penetration of Filter Material (EN 149:2001+A1:2009, Clause 8.11)	Maximum penetration of test aerosol: Sodium chloride @ 95 L/m ≤ 6% Paraffin oil @ 95 L/m ≤ 6%	Sodium chloride ≤ 2.07% Paraffin oil ≤ 4.39%	PASS
Breathing Resistance (EN 149:2001+A1:2009, Clause 8.9)	Maximum permitted resistance (mbar): Inhalation @ 30 L/min ≤ 0.7 Inhalation @ 95 L/min ≤ 2.4 Exhalation @ 160 L/min ≤ 3.0	Inhalation @ 30 L/min ≤ 0.4 Inhalation @ 95 L/min ≤ 1.46 Exhalation @ 160 L/min ≤ 1.27	PASS
Total Inward Leakage (EN 149:2001+A1:2009 Clause 8.5)	Total inward leakage ≤ 8%	Total inward leakage < 8%	PASS

^{*}NTEK testrapporter inkludert som vedlegg

Testing for samsvar med CWA 17553:2020

I tillegg har Sky Xtra munnbind blitt testet uavhengig av Intertek i henhold til vanlige standarder for partikkelfiltreringseffektivitet (PFE), både ny og etter 25 60 ° C maskinvaskesykluser, og det er bekreftet at den har følgende nøkkelegenskaper:

	Requirement	New*	After 25 washes*
Particulates Filtration Efficiency (PFE) (AFNOR-SPEC-S76-001:2020, Reference to EN13274-7: 2019 Modified)	Level 90%: ≥ 90% Level 70: ≥ 70%	> 99.5% (Average) PASS - Level 90%	> 90% (Average) PASS - Level 90%

^{*} Intertek testrapporter inkludert som vedlegg

I tillegg til NTEKs måling av pustebestandighet i henhold til EN 149: 2001 + A1: 2009 har Intertek målt luftpermeabilitet i henhold til EN ISO 9237-1995 og med et testtrykk på 100 Pa og et testområde på 20 cm2 ble Sky Xtra bekreftet å ha en luftpermeabilitet på 153,0 L/s/m2 når den er ny, komfortabelt i overkant av CWA 17553:2020-kravet på større enn eller lik 96 L/s/m2.

Testresultatene for Sky Xtra munnbind presenteres på de neste sidene.

Flashbay

Februar 2021



Report No.: \$21020400101E-R1 page 1 of 5

Test Report

Applicant: Flashbay Electronics

Address: Building 2, Jixun Industial Park, Xinjiao, Dong'ao Village, Shatian Town,

Huiyang District, Huizhou City, Guangdong Province, P.R.China

The following sample(s) was/were submitted and identified on behalf of the client as:

Product name: Face Mask

Model: Sky Xtra(SKX)

Manufacturer: Flashbay Electronics

Address: Building 2, Jixun Industial Park, Xinjiao, Dong'ao Village, Shatian Town,

Huiyang District, Huizhou City, Guangdong Province, P.R.China

Classification: FFP2 NR Sample quantity: 30 Pcs

Sample Received

Feb. 04, 2021

Date:

Testing Period: Feb. 04, 2021~ Feb. 22, 2021

Test Requirement:

According to the requirement of the client, the test item(s) of the sample is referring to the standard EN 149:2001+A1:2009.

Test Result(s): Please refer to the following page(s)

Test Method: Please refer to the following page(s)

Compiled by:	Vanly	Reviewed by:	May	N. C.
Approved by:	New blias	Date:	2021-02-23	A.



Report No.: \$21020400101E-R1 page 2 of 5

Test Result

Clause 7.9.2 Penetration of Filter Material

(EN 149:2001+A1:2009, Clause 8.11)

	* 3	Test Requir	ement	+		Results
V -		filter of the par of the following	ticle filtering half g table.	mask shall	4	407
Classificati	ion So	aximum penetra odium chloride est 95 L/min	tion of test aeros Paraffin o test 95 L/m	Detail re	efer to Appendix 1	
FFP1	4	20	20		*	
FFP2	14	6	6		47	
FFP3	4	1	1	*	3	4

Appendix 1: Summarization of Test Data

Penetration of filter material

4	. 3	~	Penetrat	ion (%)			
Aerosol	Condition	Sample No.	Average in 30s	Max. during			
<u> </u>	7	+ 20	after 3 min	exposure			
4 4		1#	2.07	1分 多			
Sodium chloride test	A.R.	2#	1.64	£ 2			
		3#	1.19	1			
4 5	A A	4#	4.38	* *			
Paraffin oil test	A.R.	5#	3.86	31			
1	7	6#	4.39	1			
, 4	Flow rate of test aerosol: 95.0 L/min						



Report No.: S21020400101E-R1 page 3 of 5

Clause 7.9.1 Total Inward Leakage

(EN 149:2001+A1:2009 Clause 8.5)

Test Requirement	Results
For particle filtering half masks fitted in accordance with the	
manufacturer's information, at least 46 out of the 50 individual exercise	4
results (i.e. 10 subjects x 5 exercises) for total inward leakage shall be	4 3
not greater than:	
25% for FFP1	
11% for FFP2	Detail refer to Appendix 2
5% for FFP3	Detail Telef to Appendix 2
and, in addition, at least 8 out of the 10 individual wearer arithmetic	
means for the total inward leakage shall be not greater than:	A 2
22% for FFP1	- L
8% for FFP2	A Comment of the Comm
2% for FFP3	4 5

Appendix 2: Summarization of Test Data

			Normal	Head	Head	Speak	Normal 🗸	Mean
Subject	Sample	Condition	Breathing	Side/Side	Up/Down	Loudly	Breathing	
	4		(%)	(%)	(%)	(%)	(%)	(%)
Gu	7#	A.R.	7.2	7.3	7.5	7.6	7.3	7.38
Hu	8#	A.R.	6.8	6.9	7.2	7.4	6.9	7.04
Wang	9#	A.R.	6.5	6.6	6.7	6.8	6.6	6.64
Long	10#	A.R.	7.4	7.6	7.7	7.9	7.5	7.62
Gao	11#	A.R.	6.9	7.1	7.2	7.4	7.1	7.14
Huang	15#	A.R.	6.9	7.1	7.2	7.3	7.1	7.12
Zhou	16#	A.R.	5.2	5.4	5.6	5.7	5.3	5.44
Ma	17#	A.R.	7.2	7.3	7.4	7.6	7.4	7.38
Wu	7 18#	A.R.	7.5	7.7	7.8	7.9	7.6	7.70
L Li 🍣	19#	A.R.	6.2	6.3	6.4	6.6	6.4	6.38



Report No.: S21020400101E-R1 page 4 of 5

Facial Dimension:

Subject	Length of Face	Width of Face	Depth of Face	Width of Mouth
Oubject	(mm)	(mm)	(mm)	(mm)
Gu	114	127	119	52
Hu	128	144	135	53
Wang	112	136	122	50
Long	119	134	128	51
Gao	130	154	144	52
Huang	130	140	125	53
Zhou	100	148	125	55
Ma	120	158	110	50
Wu	110	148	121	44
	112	146	112	50

Clause 7.16 Breathing Resistance

EN 149:2001+A1:2009, Clause 8.9)

4 5	Results							
The breathing res	The breathing resistances apply to valved and valveless filtering half							
masks and shall m	The second second							
L 5	Maximum pe	ermitted resista	4					
Classification	Inhala	ation	Exhalation		Detail refer to Appendix 3			
	30 L/min	95 L/min	160 L/min		4 3			
FFP1	0.6	2.1	3.0	4	5			
FFP2	0.7	2.4	3.0	4	L 3			
FFP3	1.0	3.0	3.0					

Appendix 3: Summarization of Test Data

Appendix	Appendix of Canimarization of Test Buta							
		Inhalation(mbar)		Inhalation(mbar) Exhalation resistance(mbar)			(mbar)	70
Specimen	Condition	At 30	At 95	4	At	160 L/min	4	
		L/min	L/min	Α	В	С	D	Е
12#	4	0.38	1.43	1.25	1.26	1.24	1.25	1.25
13#	A.R.	0.39	1.45	1.26	1.25	1.26	1.26	1.25
14#		0.40	1.46	1.26	1.25	1.26	1.27	1.26

A: facing directly ahead; B: facing vertically upwards; C: facing vertically downwards; D: lying on the left side; E: lying on the right side

Remark:

According to the requirement of the client, only the specimen of "A.R." has been tested.

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Report No.: \$21020400101E-R1 page 5 of 5

Sample photo(s):



Fig.1



Fig.2

This testing report displaces the original report of No. S21020400101E, and the original one No. S21020400101E was invalid since the date of this testing report released.

****End of Report****

The test report is effective only with both signature and specialized stamp, the result(s) shown in this report refer only to the sample(s) tested. Without written approval of NTEK, this report can't be reproduced except in full; The laboratory is not responsible for the authenticity of the sample information provided by the customer; The laboratory is not responsible for any deviation of results due to methods/standards provided by the customer.

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Number: GZHT02363627-S1

Report Ref:	GZHT02363627-S1	THIS IS TO SUPERSEDE REPORT NO.		
		GZHT02363627 DATED Dec 01, 2020		
Date received:	Nov 16, 2020	Date Issued: Dec 10, 2020		

Company Name: Address:	FLASHBAY ELECTRONICS BUILDING 2,JIXUN INDUSTRIAL PARK DONG'AO VILLAGE,SHATIAN TOWN HUIYANG DISTRICT,HUIZHOU CITY GUANGDONG PROVINCE,P.R.CHINA
Contact Name:	Levin

The Following Sample Was Subm	itted And Identified By/On Behalf Of The Applicant As:
End Uses :	Face Mask
Ratings :	-
Sample Name :	Knitted Face Mask
No. Of Sample :	One(53 pieces)
Size :	-
Colour :	Black
Standard :	-
Date received/ Test Started :	Nov 16, 2020
Ref :	Sky

Test was conducted on specific items, at our client's request.

Prepared And Checked By:

For Intertek Testing Services Shenzhen Ltd. Guangzhou Branch

Lin Lin

General Manager

QIN / hilaryxu



Page 1 Of 7

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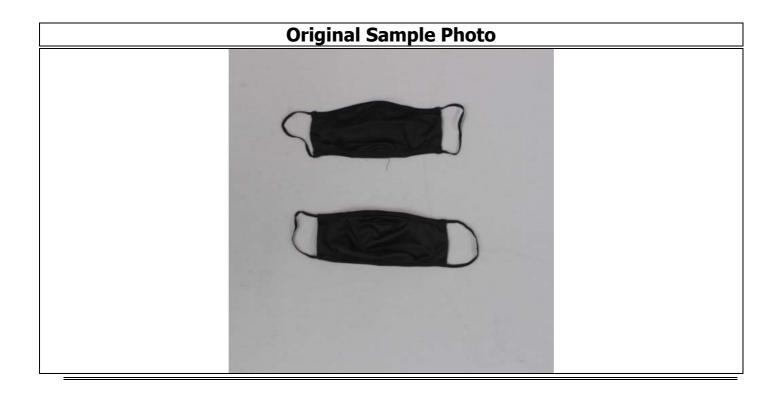
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Number: GZHT02363627-S1



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Lin Lin General Manager

QIN / hilaryxu



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Page 2 Of 7

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(10)



Number: GZHT02363627-S1

Tests Conducted (As Requested By The Applicant)

Penetration Test As Received (AFNOR-SPEC-S76-001:2020, Reference to EN13274-7: 2019 Modified) 1

TEST RESULTS:

	Efficiency of Filter M	laterial		
Aerosol	Standard terms Methods	Unit	Resu	ılt
	Aerosol particles: NaCl		#1	99.96
Sodium Chloride	Flow rate: 6cm/s		#2	99.94
	Sampling time: 1min		#3	99.98
	Temperature: 22.3℃	%	#4	99.94
	Relative humidity: 36%RH		#5	97.71
	Test area: 56.7cm ² Particle Diameter: around 3 µ m		Average	99.51
	Aerosol particles: Paraffin oil		#1	99.98
Paraffin Oil	Flow rate: 6 cm/s		#2	99.93
	Sampling time: 1min		#3	99.23
	Temperature: 22.3°C	%	#4	99.76
	Relative humidity: 36%RH		#5	99.91
	Test area: 56.7cm ² Particle Diameter: around 3 µ m		Average	99.76

Remark: The test was performed by an approved third party subcontractor laboratory.

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Page 3 Of 7

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Number: GZHT02363627-S1

Tests Conducted (As Requested By The Applicant)

2 Bacterial Filtration Efficiency (BFE)

Test Method: With reference to EN 14683: 2019+AC: 2019 Annex B

Summary of Test Method:

A specimen of the mask material is clamped between a six-stage cascade impactor and an aerosol chamber. The bacterial aerosol is introduced into the aerosol chamber using a nebulizer and a culture suspension of Staphylococcus aureus. The aerosol is drawn through the medical face mask material using a vacuum attached to the cascade impactor. The six-stage cascade impactor uses six agar plates to collect aerosol droplets which penetrate the medical face mask material. Control samples are collected with no test specimen clamped in the test apparatus to determine the upstream aerosol counts. The agar plates from the cascade impactor are incubated for (20 to 52) h and counted to determine the number of viable particles collected.

The bacterial filtration efficiency (BFE) of the mask is given by the number of colony forming units passing through the medical face mask material expressed as a percentage of the number of colony forming units present in the challenge aerosol.

Conditioning of the Specimens: 4 h at (21 ± 5) °C and (85 ± 5) % relative humidity

Test Condition:

Biological Aerosol: Staphylococcus aureus (ATCC 6538)

Testing side: Inside of the test specimen was facing towards the challenge aerosol

Test area: 78 cm² Flow rate: 28.3 L/min

The average plate count results of the positive controls: $2.5x10^3$ CFU The average plate count results of the negative controls: < 1 CFU

Mean particle size (MPS): 2.7µm

Incubation condition: (37 ± 2) °C for (20 to 52) h

Number of test specimens: 5

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Page 4 Of 7

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conomic & Technological Development District, Guangzhou, China



Number: GZHT02363627-S1

Tests Conducted (As Requested By The Applicant)

Test Procedure:

- 1. Preparation of the bacterial challenge: Dilute the culture in peptone water to achieve a concentration of approximately 5×10^5 CFU/mL.
- 2 Deliver the challenge to the nebulizer using a peristaltic or syringe pump. Connect tubing to nebulizer and peristaltic pump and into the challenge suspension; purge tubing and nebulizer of air bubbles.
- Perform a positive control run without a test specimen clamped into the test system to determine the number of viable aerosol particles being generated.
- 4. Initiate the aerosol challenge by turning on the air pressure and pump connected to the nebulizer.
- 5. Immediately begin sampling the aerosol using the cascade impactor. Adjust the flow rate through the cascade impactor to 28.3 L/m.
- 6. Time the challenge suspension to be delivered to the nebulizer for 1 min.
- 7. Time the air pressure and cascade impactor to run for 2 min.
- 8. At the conclusion of the positive control run, remove plates from the cascade impactor.
- 9. Place new agar plates into the cascade impactor and clamp the test specimen into the top of the cascade impactor, with the inside oriented toward the challenge as intended.
- 10. Repeat the challenge procedure for each test specimen and positive control sample.
- 11. Perform a negative control sample by collecting a 2 min sample of air from the aerosol chamber. No bacterial challenge should be pumped into the nebulizer during the collection of the negative control sample.
- 12. Incubate agar plates at (37 ± 2) °C for (20 to 52) h.
- 13. Count each of the six-stage plates of the cascade impactor.
- 14. Total the counts from each of the six plates for the test specimens and positive controls. Calculate the filtration efficiency percentages.

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Page 5 Of 7



GZHT02363627-S1 Number:

Tests Conducted (As Requested By The Applicant)

Calculation:

The Bacterial Filtration Efficiency (BFE), was calculated as a percentage using the following equation:

% BFE= (C-T)/C \times 100

where,

C = Average plate counts total for test controls;

T = Plate count total for the test specimen.

Test Result:

<u>Tested</u>	<u>Result</u>		
<u>Specimen</u>	The Total Plate Count (T)	Bacterial Filtration Efficiency	
-	(CFU)	(BFE) (%)	
Specimen (1)	201	91.9	
Specimen (2)	573	76.8	
Specimen (3)	233	90.6	
Specimen (4)	454	81.6	
Specimen (5)	591	76.1	

Remarks:

CFU = Colony Forming Unit

This item was conducted in Caipin Road, Guangzhou Science City, GETDD, Guangzhou, Guangdong.

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Page 6 Of 7

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Number: GZHT02363627-S1
Tests Conducted (As Requested By The Applicant)

3 Air Permeability As Received (EN ISO 9237-1995):

153.0 L/s/m²

Remark: Test Pressure = 100 Pa Test Area = 20 cm²

End of Report

This report is made solely on the basis of your instructions and/or information and materials supplied by you. It is not intended to be a recommendation for any particular course of action. Intertek does not accept a duty of care or any other responsibility to any person other than the Client in respect of this report and only accepts liability to the Client insofar as is expressly contained in the terms and conditions governing Intertek's provision of services to you. Intertek makes no warranties or representations either express or implied with respect to this report save as provided for in those terms and conditions. We have aimed to conduct the Review on a diligent and careful basis and we do not accept any liability to you for any loss arising out of or in connection with this report, in contract, tort, by statute or otherwise, except in the event of our gross negligence or wilful misconduct. No copy of the test report(except for full text copy) shall be made without the written approval by Intertek.

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To: FLASHBAY ELECTRONICS

Attention: Levin Date: Dec 10, 2020

Re: Report Revision Notification

Labtest Report Number GZHT02363627 date DEC 01, 2020

Please be informed that all the content recorded in the above captioned report will be void. This captioned report is now superseded by a revised Labtest Report, Number GZHT02363627-S1 , issued on Dec 10, 2020 .

Thank you for your attention

Prepared And Checked By:

For Intertek Testing Services Shenzhen Ltd. Guangzhou Branch

Lin Lin

General Manager

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Number: GZHT02368390

Report Ref:	GZHT02368390		
Date received/ Test	Nov 26, 2020	Date Issued:	Dec 09, 2020
Started:			

Company Name: Address:	FLASHBAY ELECTRONICS BUILDING 2, JIXUN INDUSTRIAL PARK DONG'AO VILLAGE, SHATIAN TOWN HUIYANG DISTRICT, HUIZHOU CITY GUANGDONG PROVINCE, P.R.CHINA
Contact Name:	Levin

The Following Sample Was Subm	itted And Identified By/On Behalf Of The Applicant As:
End Uses :	Face Mask
Ratings :	-
Sample Name :	Knitted Face Mask (After 25 times Washed by Client)
No. Of Sample :	One(46 pieces)
Size :	-
Colour :	Black
Standard :	-
Date received/ Test Started :	Nov 26, 2020
Ref	SKY(After 25 times Washed)

Test was conducted on specific items, at our client's request.

Prepared And Checked By:

For Intertek Testing Services Shenzhen Ltd. Guangzhou Branch

Lin Lin

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General Manager



Page 1 Of 6

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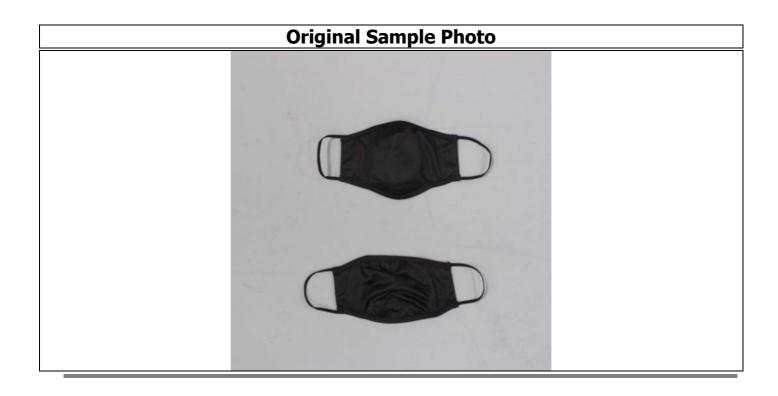
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Number: GZHT02368390



Prepared And Checked By: For Intertek Testing Services Shenzhen Ltd. Guangzhou Branch

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Page 2 Of 6

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Tests Conducted (As Requested By The Applicant)

Number: GZHT02368390

1 Penetration Test As Received (AFNOR-SPEC-S76-001:2020, Reference to EN 13274-7: 2019 Modified):

		•		_
Aerosol Particle	Test Parameters	Unit	Result	
	Flow Rate: 6 cm/s		#1	97.05
Sodium Chloride	Sampling Time: 1 min	%	#2	96.35
	Temperature: 22.1℃		#3	98.55
	Relative Humidity: 36% RH		#4	96.28
	Test Area: 56.7 cm ²		#5	98.71
	Particle Diameter: Around 3 µm		Average	97.39
	Flow Rate: 6 cm/s		#1	84.32
Paraffin Oil	Sampling Time: 1 min	%	#2	91.57
	Temperature: 22.1℃		#3	92.47
	Relative Humidity: 36% RH		#4	91.48
	Test Area: 56.7 cm ²		#5	91.93
	Particle Diameter: Around 3 µm		Average	90.63

Remark: The test was performed by an approved third party subcontractor laboratory.

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Page 3 Of 6

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2 Bacterial Filtration Efficiency (BFE)

Test Method: With reference to EN 14683: 2019+AC: 2019 Annex B

Summary of Test Method:

A specimen of the mask material is clamped between a six-stage cascade impactor and an aerosol chamber. The bacterial aerosol is introduced into the aerosol chamber using a nebulizer and a culture suspension of Staphylococcus aureus. The aerosol is drawn through the medical face mask material using a vacuum attached to the cascade impactor. The six-stage cascade impactor uses six agar plates to collect aerosol droplets which penetrate the medical face mask material. Control samples are collected with no test specimen clamped in the test apparatus to determine the upstream aerosol counts. The agar plates from the cascade impactor are incubated for (20 to 52) h and counted to determine the number of viable particles collected.

The bacterial filtration efficiency (BFE) of the mask is given by the number of colony forming units passing through the medical face mask material expressed as a percentage of the number of colony forming units present in the challenge aerosol.

Conditioning of the Specimens: 4 h at (21 ± 5) °C and (85 ± 5) % relative humidity

Test Condition:

Biological Aerosol: Staphylococcus aureus (ATCC 6538)

Testing side: Inside of the test specimen was facing towards the challenge aerosol

Test area: 78 cm² Flow rate: 28.3 L/min

The average plate count results of the positive controls: 2.4×10^3 CFU The average plate count results of the negative controls: < 1 CFU

Mean particle size (MPS): 2.7 µm

Incubation condition: (37 ± 2) °C for (20 to 52) h

Number of test specimens: 5

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Page 4 Of 6

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Test Procedure:

- Preparation of the bacterial challenge: Dilute the culture in peptone water to achieve a concentration of approximately 5×10⁵ CFU/mL.
- 2 Deliver the challenge to the nebulizer using a peristaltic or syringe pump. Connect tubing to nebulizer and peristaltic pump and into the challenge suspension; purge tubing and nebulizer of air bubbles.
- 3 Perform a positive control run without a test specimen clamped into the test system to determine the number of viable aerosol particles being generated.
- 4. Initiate the aerosol challenge by turning on the air pressure and pump connected to the nebulizer.
- 5. Immediately begin sampling the aerosol using the cascade impactor. Adjust the flow rate through the cascade impactor to 28.3 L/m.
- 6. Time the challenge suspension to be delivered to the nebulizer for 1 min.
- Time the air pressure and cascade impactor to run for 2 min. 7.
- At the conclusion of the positive control run, remove plates from the cascade impactor.
- Place new agar plates into the cascade impactor and clamp the test specimen into the top of the 9. cascade impactor, with the inside oriented toward the challenge as intended.
- 10. Repeat the challenge procedure for each test specimen and positive control sample.
- Perform a negative control sample by collecting a 2 min sample of air from the aerosol chamber. No bacterial challenge should be pumped into the nebulizer during the collection of the negative control sample.
- 12. Incubate agar plates at (37 ± 2) °C for (20 to 52) h.
- Count each of the six-stage plates of the cascade impactor. 13.
- Total the counts from each of the six plates for the test specimens and positive controls. Calculate the filtration efficiency percentages.

Calculation:

The Bacterial Filtration Efficiency (BFE), was calculated as a percentage using the following equation:

% BFE= (C-T)/C \times 100

where,

C = Average plate counts total for test controls;

T =Plate count total for the test specimen.

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Page 5 Of 6

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Tests Conducted (As Requested By The Applicant)

Test Result:

<u>Tested</u>		<u>Result</u>	
Specimen	The Total Plate Count (T)	Bacterial Filtration Efficiency	
	(CFU)	(BFE) (%)	
Specimen (1	579	75.7	
Specimen (2)	582	75.5	
Specimen (3	537	77.4	
Specimen (4	513	78.4	
Specimen (5	444	81.3	

Remarks:

CFU = Colony Forming Unit

This item was conducted in Caipin Road, Guangzhou Science City, GETDD, Guangzhou, Guangdong.

End of Report

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Page 6 Of 6

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