



中国认可 国际互认 检测 TESTING CNAS L0599

Test Report

SL52035297686301TX

Date:September 21,2020

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to the

NANJING N.E. ORIENTAL IMPORT.& EXPORT. CO., LTD 702 FLOOR, B4 BUILDING, NO.118, RUANJIAN ROAD, YUHUATAI DISTRICT, NANJING

The following sample(s) was/were submitted and identified on behalf of the client as: Sample Description : (A) Surgical mask (Claimed Type IIR)

Buyer Composition Sample Color Style No.	::	MD FONSCARE (A)55%Non-woven 25%melt-blown 10%bridge 10%earloop (A)Blue MD FONSCARE MD-319R
Manufacturer Supplier Country of Destination	: : :	Jiangsu jinshaxiang technology Co., Ltd NANJING N.E. ORIENTAL IMPORT.& EXPORT. CO.,LTD France
Test Performed	:	Selected test(s) as requested by applicant
Sample Receiving Date Testing Period	:	Sep 08, 2020 Sep 09, 2020 - Sep 21, 2020
Test Result(s)	:	Unless otherwise stated the results shown in this test report refer only to the sample(s) tested, for further details, please refer to the following page(s).

Comment:

EN 14683:2019+AC:2019 Medical Face Masks-Requirements and Test	(A)
<u>Methods</u>	
Clause 5.2 Performance Requirement	
Clause 5.2.2 Bacterial filtration efficiency (BFE)	М
Clause 5.2.3 Breathability	М
Clause 5.2.4 Splash Resistance	М
Clause 5.2.5 Microbial Cleanliness	М
Clause 5.2.6 Biocompatibility	EXCLUDED

Remark: M=Meet EN 14683:2019+AC:2019 Performance Requirement (Type IIR)

Signed for and on behalf of SGS-CSTC Standards Technical Services (Shanghai) Co., Ltd Testing Center

Ponjig li Helen

Sara Guo (Account Executive)

Dongjing Liu / Hailian Xuan (Authorized Signatory)



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Test Result

EN 14683:2019+AC:2019 Medical Face Masks-Requirements and Test Methods

Clause 5.2 Performance Requirement

Clause 5.2.2 Bacterial Filtration Efficiency (BFE)

(EN 14683:2019+AC:2019 Annex B)

Flow Rate:Pre-Conditioning:Dimensions of test specimen:Positive Control Average:Negative Monitor Count:Mean Particle Size:	Inside Approximately 60 cm ² 28.3 L/min Minimum of 4 hours at 21±5°C and 85±5% R.H. ~174mm x 157mm 2156 CFU < 1 CFU 3.0 ±0.3µm Staphylococcus aureus ATCC 6538
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Test Item	Specimen No.	Result
	1	99.6
Postarial Filtration Efficiency (PEE)	2	99.9
Bacterial Filtration Efficiency (BFE), %	3	99.7
70	4	99.9
	5	99.9

Remark:

- 1) Performance Requirement: Type I≥95%, Type II≥98%, Type IIR ≥98%
- 2) The number of specimens that shall be tested is minimum 5, but can be greater and shall be increased if necessary to allow for an AQL(Acceptable Quality Level) of 4%.



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Clause 5.2.3 Breathability

(EN 14683 :2019+AC:2019 Annex C)

Sample: A		
Test Side	:	Randomly test in different location (1 around and 4 away from the centric point) on each of the 5 masks
Pre-Conditioning	:	Minimum of 4 hours at 21±5°C and 85±5% R.H.
Test Area	:	4.9 cm ²
Flow Rate	:	8 l/min

Specimen No.	Test Area No.	Different Pressure for each	The average value for each test
		tested area (Pa/cm ²)	specimen (Pa/cm ²)
	1-1	38.2	
	1-2	35.2	
1	1-3	38.0	38
	1-4	36.9	
	1-5	39.2	
	2-1	39.5	
	2-2	38.5	
2	2-3	38.8	38
	2-4	38.3	
	2-5	36.6	
	3-1	34.2	
	3-2	35.8	
3	3-3	36.3 37	
	3-4	39.7	
	3-5	38.5	
	4-1	29.7	
4	4-2	32.1	
	4-3	31.6	33
	4-4	36.1	
	4-5	36.3	
	5-1	30.4	
F	5-2	27.4	7
5	5-3	26.3	30
	5-4	35.7	7
	5-5	30.4	1

Remark:

- 1) Performance Requirement: Type I<40 Pa/cm², Type II<40 Pa/cm², Type IIR<60 Pa/cm²
- The number of specimens that shall be tested is minimum 5, but can be greater and shall be increased if 2) necessary to allow for an AQL(Acceptable Quality Level) of 4%.



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Clause 5.2.4 Splash Resistance

(ISO 22609 :2004)

Sample: A Test Blood Pressure Pre-Conditioning Distance of the mask to the tip of cannula

16.0kPa

Minimum of 4 hours at 21±5°C and 85±5% R.H.

: 300±10mm

Test Specimen#	Penetration on inside surface	Conclusion	Test Specimen#	Penetration on inside surface	Conclusion
1	None Seen	Pass	17	None Seen	Pass
2	None Seen	Pass	18	None Seen	Pass
3	None Seen	Pass	19	None Seen	Pass
4	None Seen	Pass	20	None Seen	Pass
5	None Seen	Pass	21	None Seen	Pass
6	None Seen	Pass	22	None Seen	Pass
7	Seen	Fail	23	None Seen	Pass
8	None Seen	Pass	24	None Seen	Pass
9	None Seen	Pass	25	None Seen	Pass
10	None Seen	Pass	26	None Seen	Pass
11	None Seen	Pass	27	Seen	Fail
12	None Seen	Pass	28	None Seen	Pass
13	Seen	Fail	29	None Seen	Pass
14	None Seen	Pass	30	None Seen	Pass
15	None Seen	Pass	31	None Seen	Pass
16	None Seen	Pass	32	None Seen	Pass
Number of Pass:		29			
Overall result:		Acceptable			

Remark:

- 1) Performance Requirement Type I: N/A, Type II: N/A, Type IIR: ≥16.0kPa
- 2) Test was conducted within 60s after removal from conditioning chamber.
- 3) An acceptable quality limit of 4.0% is met for a single sampling plan when 29 or more of the 32 tested specimens show pass results.

Clause 5.2.5 Microbial Cleanliness

(EN 14683:2019+AC:2019 Annex D and EN ISO 11737-1:2018)

	Mask Weight(g)	Total Bioburden, (cfu/mask)	Total Bioburden, (cfu/g)
Sample Number			
1#	3.61	3	0.83
2#	3.64	6	1.65
3#	3.63	3	0.83
4#	3.66	6	1.64
5#	3.67	3	0.82

Remark: Performance Requirement: Type I≤30 CFU/g, Type II≤30 CFU/g, Type IIR≤30 CFU/g

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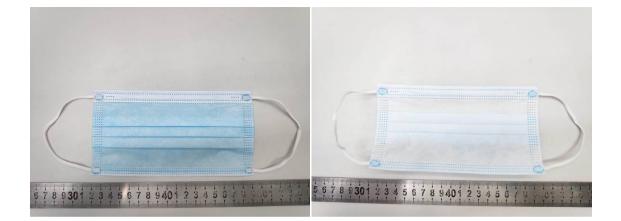
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Sample Photo



The statement of conformity in this test report is only based on measured values by the laboratory and does not take their uncertainties into consideration.

End of Report



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