



SL52045300703401TX

Date:October 30,2020

ELDO CARE CORPORATION LIMITED RM.1902, EASEY COMM. BLDG., 253-261 HENNESSY ROAD, WANCHAI, HONGKONG

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

Sample Description	:	(A)GIMA BRANDED 20700&20695&20696 3 PLY SURGICAL MASK - DISPOSABLE-EARLOOP - LIGHT BLUE
Composition Sample Color Style No. Lot No. Manufacturer	: : : :	(A)67% non-woven, 33% melt-blown (A)LIGHT BLUE ELFM3600 Not provided ELDO CARE CO., LIMITED
Test Performed	:	Selected test(s) as requested by applicant
Sample Receiving Date Testing Period Test Result(s)	:	Oct 19, 2020 Oct 21, 2020 - Oct 30, 2020 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).

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

Juan Helen

Helen 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)

Sample: A Test Side Test Area 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. ~176mm x 154mm 1889.5 CFU < 1 CFU 3.0 ±0.3µm
Mean Particle Size Test bacteria	:	3.0 ±0.3μm Staphylococcus aureus ATCC 6538

Test Item	Specimen No.	Result
	1	99.9%
Bacterial Filtration Efficiency (BFE)	2	99.9%
	3	99.9%
	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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Sara Guo (Account Executive)



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Test Result

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

Clause 5.2 Performance Requirement

Clause 5.2.3 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 tested area (Pa/cm ²)	The average value for each test specimen (Pa/cm ²)
	1-1	56.8	
	1-2	54.6	
1	1-3	46.2	53
	1-4	59.5	
	1-5	48.6	
	2-1	59.5	
	2-2	45.8	
2	2-3	41.6	50
	2-4	52.6	
	2-5	51.9	
	3-1	52.8	
	3-2	52.5	
3	3-3	48.9	52
	3-4	52.6	
	3-5	53.7	
	4-1	49.6	
	4-2	48.7	
4	4-3	52.8	49
	4-4	46.7	
	4-5	44.8	
	5-1	46.9	
	5-2	53.2	
5	5-3	53.3	52
	5-4	56.2	
	5-5	49.6	

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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Test Result

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

:

:

:

Clause 5.2 Performance Requirement

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	None Seen	Pass	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	None Seen	Pass
12	None Seen	Pass	28	None Seen	Pass
13	None Seen	Pass	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:		32			
Overall result:			Acce	ptable	

Remark:

- Performance Requirement Type I: N/A, Type II: N/A, Type IIR: ≥16.0kPa 1)
- 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.



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Dongjing Liu (Authorized Signatory)



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Test Result

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Clause 5.2 Performance Requirement

Clause 5.2.5 Microbial Cleanliness

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

Sample: A

Test Specimen#	Mask Weight(g)	Total Bioburden, (CFU/mask)	Total Bioburden, (CFU/g)
1#	3.04	15	4.93
2#	3.01	18	5.98
3#	3.03	9	2.97
4#	3.01	15	4.98
5#	2.98	6	2.01

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



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