

Test Report

SL52045311965401TX

Date: November 19, 2020

Page 1 of 4

CHONGQING COE DISPLAY TECHNOLOGY CO., LTD.
NO.3, TONGGUI AVENUE, YUFENSHAN TOWN, YUBEI DISTRICT, CHONGQING

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

Sample Description : (A) Disposable medical mask

Composition : (A) 65% non-woven, 35% Meltblown

SGS Internal Ref.No. : CQ21310390

Sample Color : (A) blue

Model No. : CCDM-NC

Product Specification : 14.5cm*9.5cm

Lot No. : C201103CN01

Product status : non-sterile

Manufacturer : CHONGQING COE DISPLAY TECHNOLOGY CO., LTD.

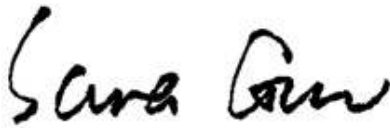
Test Performed : Selected test(s) as requested by applicant

Sample Receiving Date : Nov 05, 2020

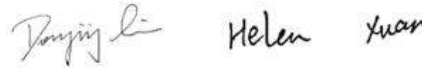
Testing Period : Nov 06, 2020 - Nov 19, 2020

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).

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



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)

Sample: A
 Test Side : Inside
 Test Area : Approximately 60 cm²
 Flow Rate : 28.3 L/min
 Pre-Conditioning : Minimum of 4 hours at 21±5°C and 85±5% R.H.
 Dimensions of test specimen : ~141mm x 154mm
 Positive Control Average : 2719.5 CFU
 Negative Monitor Count : < 1 CFU
 Mean Particle Size : 3.0 ±0.3µm
 Test bacteria : Staphylococcus aureus ATCC 6538

Test Item	Specimen No.	Result
Bacterial Filtration Efficiency (BFE)	1	99.9%
	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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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 cm²

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	1-1	35.7	37
	1-2	35.4	
	1-3	38.1	
	1-4	36.2	
	1-5	39.6	
2	2-1	35.0	38
	2-2	39.9	
	2-3	38.7	
	2-4	39.6	
	2-5	35.5	
3	3-1	35.4	36
	3-2	30.5	
	3-3	39.8	
	3-4	37.5	
	3-5	38.2	
4	4-1	37.2	36
	4-2	38.2	
	4-3	35.4	
	4-4	39.2	
	4-5	31.7	
5	5-1	34.6	36
	5-2	34.2	
	5-3	37.5	
	5-4	33.3	
	5-5	38.5	

Remark:

- 1) Performance Requirement: Type I<40 Pa/cm², Type II<40 Pa/cm², Type IIR<60 Pa/cm²
- 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 5.2.5 Microbial Cleanliness

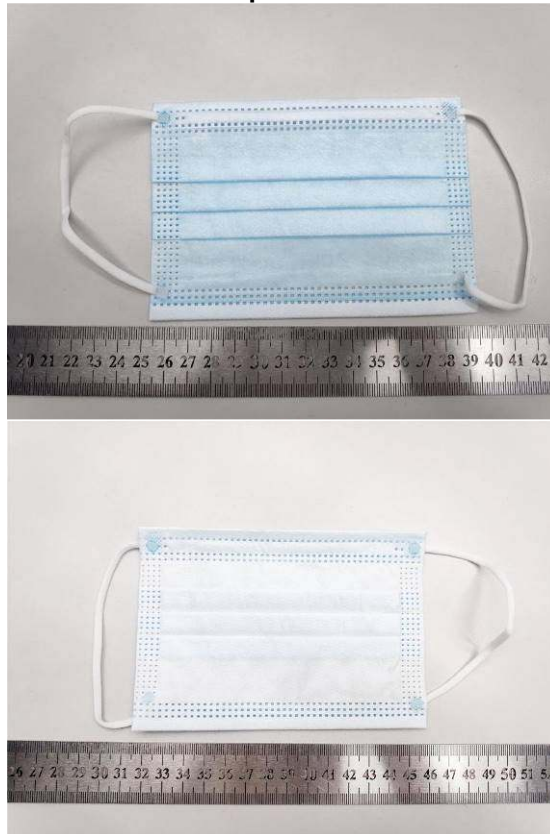
(EN 14683:2019+AC:2019 Annex D and EN ISO 11737-1:2018)

Sample: A

Test Specimen#	Mask Weight(g)	Total Bioburden, (CFU/mask)	Total Bioburden, (CFU/g)
1#	2.54	<3	<1.18
2#	2.58	3	1.16
3#	2.56	6	2.34
4#	2.62	6	2.29
5#	2.61	6	2.30

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

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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