Extrait de la copie du rapport de test SGS - performances selon EN14683:2019+AC:2019 : filtration bactérienne (EFB), respirabilité, résistance aux projections et absence de charge microbienne (propreté microbienne). Le certificat original émis par SGS est fourni sur demande à tout client ayant commandé ce produit. **Robé Médical**

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 cm^2
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	~176mm x 158mm
Positive Control Average	2473 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	99.8%
	3	99.9%
	4	99.8%
	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%.

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±5ºC 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	48.4	
	1-2	49.5	
1	1-3	43.6	46
	1-4	43.3	
	1-5	43.6	
	2-1	45.9	
	2-2	43.4	
2	2-3	42.3	45
	2-4	46.7	
	2-5	44.8	
	3-1	46.7	
	3-2	42.6	
3	3-3	43.1	47
	3-4	51.9	
	3-5	48.8	
	4-1	55.1	
	4-2	45.3	
4	4-3	41.0	49
	4-4	54.7	
	4-5	46.8	
	5-1	46.9	
	5-2	48.1	
5	5-3	41.9	46
	5-4	43.5	
	5-5	47.8	

Remark:
1) Performance Requirement: Type I<40 Pa/cm^2, Type II<40 Pa/cm^2, Type IIR<60 Pa/cm^2
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%.

Clause 5.2.4 Splash Resistance
(ISO 22609 :2004)

Sample: A
Test Blood Pressure 16.0kPa
Pre-Conditioning Minimum of 4 hours at 21±5°C and 85±5% R.H.
Distance of the mask to the tip of cannula 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:			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)

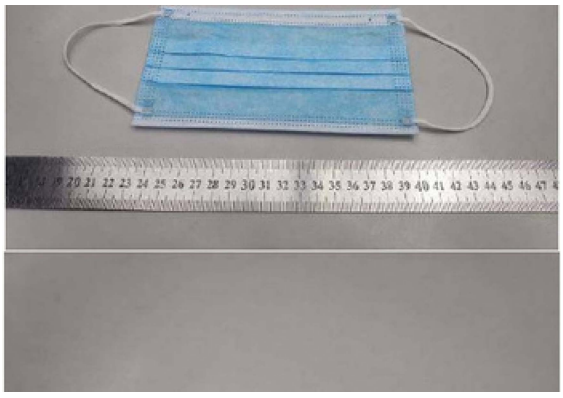
Sample: A

Test Specimen#	Mask Weight(g)	Total Bioburden, (CFU/mask)	Total Bioburden, (CFU/g)
1	2.91	18.60	6.39
2	2.88	22.32	7.75
3	2.92	29.76	10.19
4	3.00	22.32	7.44
5	2.98	26.04	8.74

Recovery Efficiency 80.4 %
Correction Factor 1.2

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