

TEST REPORT

2021TM2065

DATE OF RECEPTION

05/08/2021

APPLICANT

UAB "PDSA"
Lakūnų g. 3A
LT-09108 Vilnius

DATE TESTS

Starting: 05/08/2021
Ending: 23/08/2021

Att. Arūnas Mirinas

IDENTIFICATION AND DESCRIPTION OF SAMPLES

REFERENCES

MASK REF. MM-2R

TESTS CARRIED OUT

- IN VITRO DETERMINATION OF BACTERIAL FILTRATION EFFICIENCY (BFE).
- DETERMINATION OF BREATHABILITY (DIFFERENTIAL PRESSURE).
- DETERMINATION OF PRESSURE OF SPLASH RESISTANCE.
- DETERMINATION OF A POPULATION OF MICROORGANISMS ON PRODUCTS.



SAMPLE DESCRIPTION

PHOTOGRAPHY



Reference ⁽¹⁾
MASK REF. MM-2R

LOT number ⁽¹⁾

⁽¹⁾ Data provided for the customer

///



RESUMEN / SUMMARY

Of the tests carried out on the following reference:

MASK REF. MM-2R

ORIGINAL. No pretreatment has been performed.

Tests according to the standard EN 14683:2019+AC: 2019.

Having obtained the following results:

TESTS	RESULTS
Pto 5.2.2 Bacterial Filtration Efficiency (BFE) (%)	99,35
Pto 5.2.3 Breathability: Differential pressure (Pa/cm ²)	16,1
Pto 5.2.4 Splash resistance pressure (kPa)	Failure 1 of 32 at 16 kPa

Notes

- The rest of the standard tests not indicated in this report, have not been evaluated.

-----///



RESULTS

IN VITRO DETERMINATION OF BACTERIAL FILTRATION EFFICIENCY (BFE)

Standard

EN 14683:2019+AC:2019

Test date

05/08/2021 - 06/08/2021

Batch n^o[1]

Reference

MASK REF. MM-2R

Number of test specimen

5

Size of test specimen

10 cm x 10 cm

Tested area of the test specimen

50 cm²

Sample side was oriented toward the challenge aerosol

Inner side

Equipment

Six stage Andersen Sampler (03285E12)

Flow of air

28.3 l/min

Test germ

Staphylococcus aureus ATCC 6538

Incubation conditions

24 h at 37 ± 2 °C

Uncertainty of the test

The relative expanded uncertainty of the test is ± 5 % assay value of the measured.

----->>>



RESULTS

Test sample values							
	Level1 (cfu/plate)	Level2 (cfu/plate)	Level3 (cfu/plate)	Level4 (cfu/plate)	Level5 (cfu/plate)	Level6 (cfu/plate)	Total count (ufc)
1	0	0	0	3	4	0	7
2	0	0	0	5	13	0	18
3	0	0	0	1	5	0	6
4	0	0	0	8	8	0	16
5	0	0	3	3	3	0	9

Legend meaning: cfu: colony forming units

Pre-treatment Original. No pretreatment has been performed.

Calculation of bacterial filtration efficiency:

Test	Filtration efficiency (%)
1	99,59
2	98,95
3	99,65
4	99,07
5	99,47
Mean	99,35

Notes

- The "positive hole" conversion factor described by A. Andersen has been applied to the number of CFU colony forming units collected by the cascade impactor for the sample and positive control.
- Tested samples were supplied by the customer.
- Mean of the plate counts of the negative controls: 0 ufc.
- Mean of the total plate counts of the two positive controls: 1713 cfu.

- ^[1] Data provided by the customer.

///



RESULTS

DETERMINATION OF BREATHABILITY (DIFFERENTIAL PRESSURE)

Standard

EN 14683:2019+AC:2019

Principle

It is measure the differential pressure required to move air through a measured surface area at a constant flow of air, with the aim of measuring the pressure of air exchange of the material of the mask.

Test date

20/08/2021 - 23/08/2021

Batch n^{o(1)}

Reference

MASK REF. MM-2R

Number of test specimen

5

Size of test specimen

4.9 cm²

Tested area of the test specimen

Circular, diameter 2.5 cm

Sample conditioning

T^a 21 ± 5 °C Hr 85 ± 5 %

Flow of air

(8 ± 0,3) l/min

Pre-treatment

Original. No pretreatment has been performed.

Uncertainty of the test

The relative expanded uncertainty of the test is ± 6 % assay value of the measured

----->>>



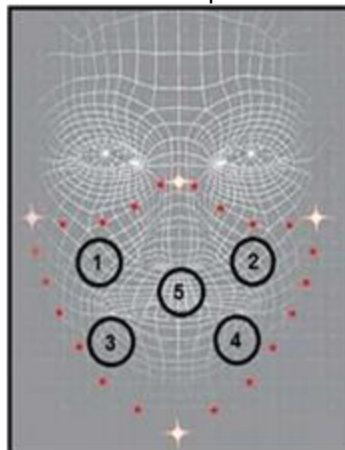
RESULTS

Results

Test specimen	Pos1 Pa	Pos2 Pa	Pos3 Pa	Pos4 Pa	Pos5 Pa	Average Pa	ΔP (Pa/cm ²)
1	73,2	70,2	77,0	91,2	73,5	77,0	15,7
2	73,2	66,7	75,3	88,7	72,2	75,2	15,4
3	80,6	82,2	80,9	83,2	80,3	81,4	16,6
4	74,0	78,3	77,2	85,8	74,2	77,9	15,9
5	95,8	76,2	83,2	72,2	90,2	83,5	17,0
					Average	79,0	16,1

Notes

- Tested samples were supplied by the customer.
- The specimens of each mask have been taken from the positions according to the image:



- ⁽¹⁾Data provided by the customer.

_____///



RESULTS

DETERMINATION OF PRESSURE OF SPLASH RESISTANCE

Standard EN 14683:2019+AC:2019 **Test method** ISO 22609:2004

Principle:

A defined volume of synthetic blood is shot with defined speeds of a pneumatically checked valve at the test specimen, in order to simulate a squirting of blood and other body fluids for the sample material. The back of the mask is examined by means of visual inspection and swab on penetrating liquid. The more the resistance against liquid splashes, the more merrier is the liquid resistance.

Test date

12/08/2021 - 13/08/2021

Batch n^o(1)

Reference

MASK REF. MM-2R

Material of test sample

Fabric

Tested area of the test specimen

19.6 cm²

Sample Conditioning

T^a 21 ± 5 °C

Hr 85 ± 5 %

Test environmental test conditions

T^a 21 ± 5 °C

Hr 36 ± 5 %

Test parameters 16 KPa (120 mm Hg) **Volume of synthetical blood** 2.0 mL

Pre-treatment

Original. No pretreatment has been performed.

----->>>



RESULTS

Results Pressure 16 KPa (120 mm Hg)		
Replica	Passed	Failed
1	X	
2	X	
3	X	
4	X	
5	X	
6		X
7	X	
8	X	
9	X	
10	X	
11	X	
12	X	
13	X	
14	X	
15	X	
16	X	
17	X	
18	X	
19	X	
20	X	
21	X	
22	X	
23	X	
24	X	
25	X	
26	X	
27	X	
28	X	
29	X	
30	X	
31	X	
32	X	

>>>



RESULTS

Remarks

- To pass the test no more than 3 of 32 samples may fail.

- ⁽¹⁾Data provided by the customer.

///



RESULTS

DETERMINATION OF A POPULATION OF MICROORGANISMS ON PRODUCTS

Standard

EN 14683:2019+AC:2019; EN ISO 11737-1:2018

Reference

MASK REF. MM-2R

Batch number ⁽¹⁾

Sample size (SIP)

4,14 g

Replica number

5

Test date

18/08/2021 – 23/08/2021

Test equipments

Incubator (03068E05) and Incubator (03202E05)

Results

Parameter	Replica 1 (ufc/g)	Replica 2 (ufc/g)	Replica 3 (ufc/g)	Replica 4 (ufc/g)	Replica 5 (ufc/g)	Average (ufc/g)
Aerobic bacteria to 33 ± 2°C	1	1	4	2	6	3
Moulds and yeasts to 22 ± 2°C	<1	<1	<1	<1	<1	<1



RESULTS

Notes

⁽¹⁾Data provided from customer

The total count of microorganisms in the sample is 3 cfu/g

In accordance with the standard EN 14683:2019+AC:2019, the results must be in the values of the following table:

Parameter		
Cleanliness microbial	ufc/g	≤ 30

-----///



Judit Sisternes
Head of Health & Hygiene Products Division

LIABILITY CLAUSES

- 1.- AITEX is liable only for the results of the methods of analysis used, as expressed in the report and referring exclusively to the materials or samples indicated in the same which are in its possession, the professional and legal liability of the Centre being limited to these. Unless otherwise stated, the samples were freely chosen and sent by the applicant.
- 2.- AITEX shall not be liable in any case of misuse of the test materials nor for undue interpretation or use of this document
- 3.- The Offer and / or Order to which the applicant gives approval through signature and seal, constitutes the Legally Executable Agreement in which AITEX is responsible for safeguarding and guaranteeing the absolute confidentiality of the management of all the information obtained or created during the performance of the contracted activities.
- 4.- In the eventuality of discrepancies between reports, a check to settle the same will be carried out in the head offices of AITEX. Also, the applicants undertake to notify AITEX of any complaint received by them as a result of the report, exempting this Centre from all liability if such is not done, the periods of conservation of the samples being taken into account.
- 5.- AITEX will provide at the request of the person concerned, the treatment of complaints procedure. In the event that you want to make it, direct it to: calidad@aitex.es.
- 6.- AITEX is not responsible for the information provided by customers, which is reflected in the Report, and may affect the validity of the results.
- 7.- AITEX is not responsible for an inadequate state of the sample received that could compromise the validity of the results, expressing such circumstance, in the test reports.
- 8.- AITEX may include in its reports, analyses, results, etc., any other evaluation which it considers necessary, even when it has not been specifically requested.
- 9.- When a Declaration of Conformity is requested, if not indicated otherwise, the decision rule will be applied according to ILAC-G8 & ISO 10576-1, in case of ambiguity, or indeterminacy
- 10.- The uncertainties of tests, which are made explicit in the Results Report, have been estimated for a $k = 2$ (95% probability of coverage). If not informed, they are available to the client in AITEX.
- 11.- The original materials and rests of samples, not subject to test, will be retained in AITEX during the twelve months following the issuance of the report, so that any check or claim which, in his case, wanted to make the applicant, should be exercised within the period indicated.
- 12.- This report may only be sent or delivered by hand to the applicant or to a person duly authorised by the same.
- 13.- The results of the tests and the statement of compliance with the specification in this report refer only to the test sample as it has been analyzed / tested and not the sample / item which has taken the test sample.
- 14.- The client must attend at all times, to the dates of the realization of the tests.
- 15.- According to Resolution EA (33) 31, the test reports must include the unique identification of the sample, and any brand or label of the manufacturer may be added. It is not allowed to re-issue test reports of untested sample names (references), they can only be re-issued for error correction or inclusion of omitted data that were already available at the time of the test. The laboratory can not assume responsibility for declaring that the product with the new trade name / trademark is strictly identical to the one originally tested; This responsibility belongs to the client.
- 16.- This report may not be partially reproduced without the written approval of the issuing laboratory.