Notified Body Number: 2841

Product TS EN ISO/IEC 17065 AB-0128-U

AB Tip İnceleme Sertifikası
EU Type-Examination Certificate

Belge No / *Certificate No*	**:** 74011335
Belgelendirme Tarihi - Bir Sonraki Belge Tarihi / *Certification Date / Certificate Validity Date*	**:** 26.01.2026-26.01.2031
Belge Geçerlilik Tarihi / *Document Validity Period*	**:** 5 yıl / *5 years*
Firma Unvanı ve Adresi / *Company Name and Address*	**:** GRAFOPLAS DEL NOROESTE, S.A. LG. A MARİSQUEİRA, S/N (POL. IND. ALMEİRAS) 15180 CULLEREDO (A CORUÑA) - SPAIN
Marka /Modeller / *Brand / Models*	**:** RADEX / MFR001G, MFVR002G
Direktifi / *Directive*	**:** 2016/425 REGULATION
Modülü/Kategori / *Module / Category*	**:** B MODÜLÜ/ KATEGORİ III *MODULE B / CATEGORY III*
Teknik Değerlendirme Rapor No/ *Technical Evaluation Report No*	**:** 74011335

Ürün Tipi / *Product Type:*

- EN 149:2001+ A1:2009 Solunumla ilgili koruyucu cihazlar - Parçacıklara karşı koruma amaçlı filtreli yarım maskeler/ *Respiratory protective devices - Filtering half masks to protect against particles*

Ürünün Malzeme Bilgisi / *Product Material Information*: RADEX / MFR001G, MFVR002G model ürünleri kumaş, elastik kayış, soluk verme valfi, burun klipsi ve filtre katmanı kullanılarak imal edilmiştir./ *RADEX / MFR001G, MFVR002G model products are manufactured using fabric, elastic strap, nose clip, exhalation valve and filter layer.*

Karar Verici / Approver ***Şirket Müdürü / General Manager***

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MNA Laboratuvarları San. Tic.Ltd .Şti
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This document has been signed electronically in accordance with the Electronic Signature Law No. 5070.
The document can be checked at https://www.mnalab.com/en/sertifika-sorgula

Notified Body Number: 2841

ATTACHMENTS (74011335)

To certify the PPE product at Category III level, C2 or D module is accompanied by applying one of the conformity assessment methods along with the EU Type Examination (Module B).

Model : RADEX / MFR001G, MFVR002G

PPE SPECIFICATION	PERFORMANCE LEVELS
Classification	FFP3
Reusable / Single Shift Use	NR
Clogging performance	D (MFVR002G with valve)

PPE produced as a single unit to fit an individual user, all the necessary instructions for manufacturing such PPE on the basis of the approved basic model:

MARKING

MANUFACTURER: GRAFOPLAS DEL NOROESTE, S.A.
LG. A MARİSQUEİRA, S/N (POL. IND. ALMEİRAS) 15180 CULLEREDO (A CORUÑA) – SPAIN

PPE TYPE:

- EN 149:2001+ A1:2009 Respiratory protective devices - Filtering half masks to protect against particles

MODEL: RADEX / MFR001G, MFVR002G
PRODUCT SIZE: Standard
PICTOGRAM AND PERFORMANCE LEVELS:

RADEX MFR001G EN 149:2001+ A1:2009 FFP3 NR for the without valve
RADEX MFR002G EN 149:2001+ A1:2009 FFP3 NR D for the with valve



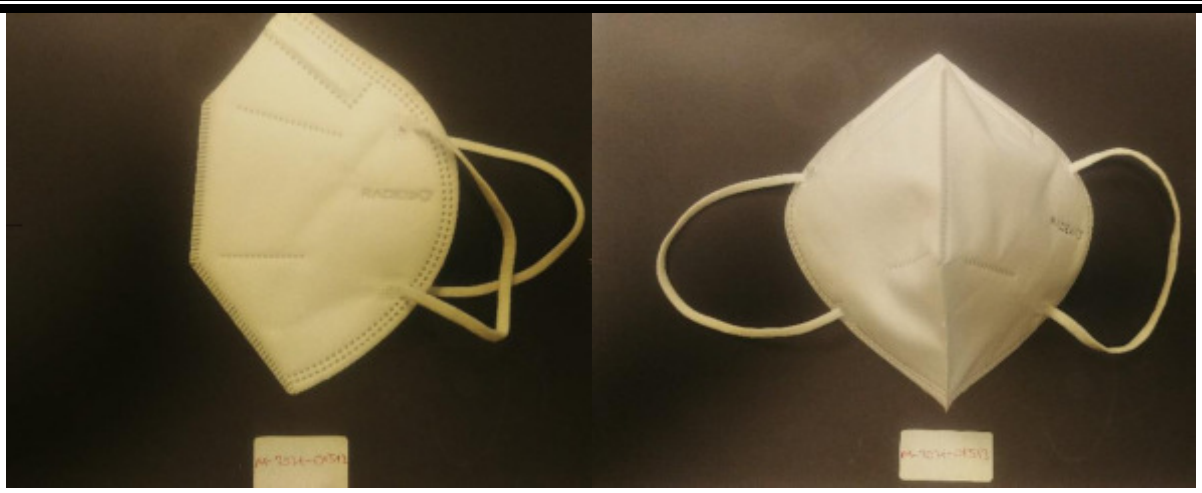
2841

Or Condition of Storage

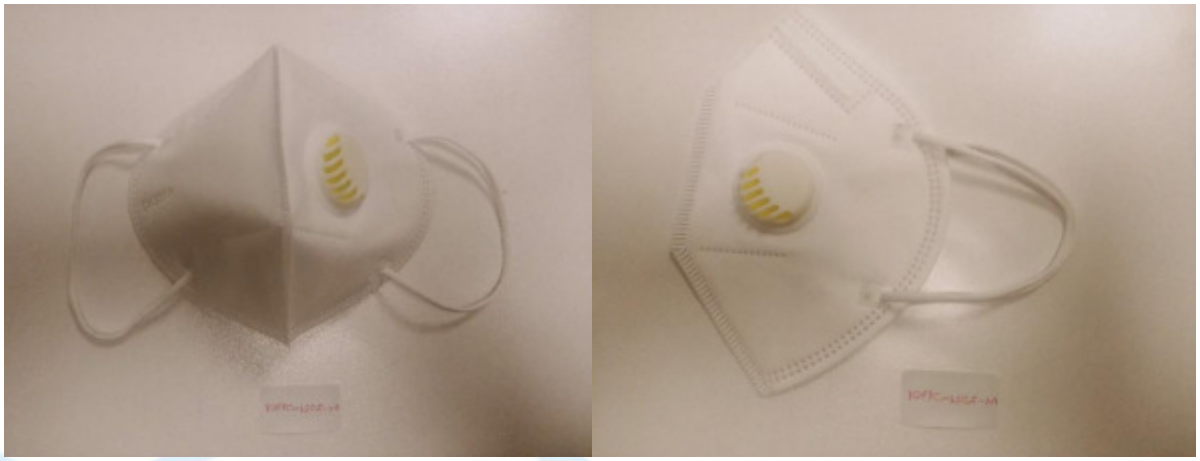
MNA LABORATORIES SAN. TIC. LTD. ŞTİ declares that the above-mentioned product meets the requirements of the directive according to the EU Directive 2016/425, the safety of the product is covered by the conditions and use specified in this certificate and in the technical file.

ATTACHMENTS (74011335)

PRODUCT PICTURES



RADEX MFR001G



RADEX MFVR002G

DOCUMENTS IN THE TECHNICAL FILE

- Test Reports
- Technical Report

Report No : 74011335

Report Date : 26.01.2026

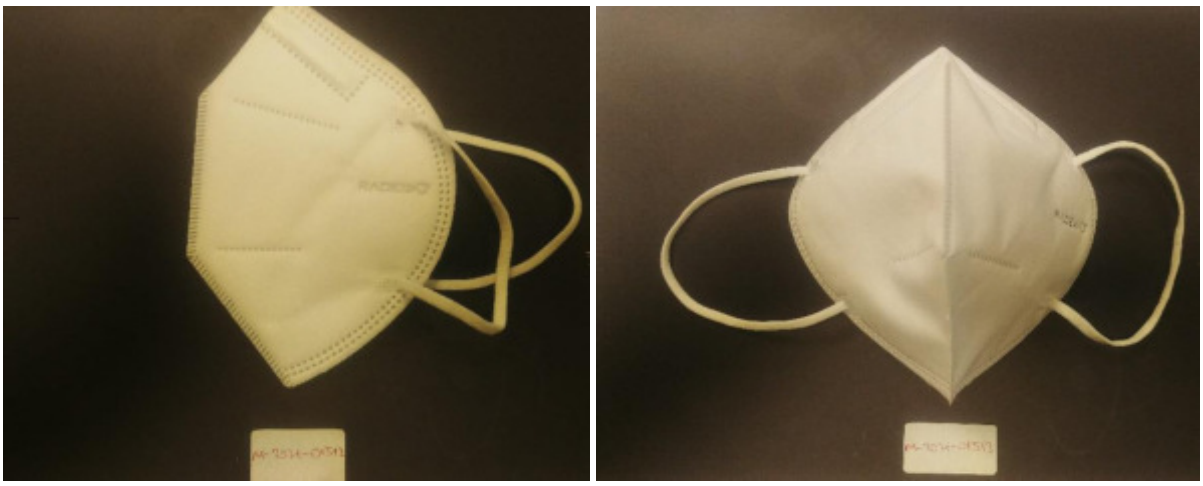
Application No : 74011335

1. **COMPANY INFORMATION:**
   GRAFOPLAS DEL NOROESTE, S.A.
   LG. A MARİSQUEİRA, S/N (POL. IND. ALMEİRAS) 15180 CULLEREDO (A CORUÑA) – SPAIN
   Tel: +34 981 677 677
   Fax: +34 981 677 020
   E-mail: admin@grafoplas.com

2. **PPE INFORMATION:**
   Disposable and non-sterile half mask made of particulate protection filter material.

3. **PPE TYPE IDENTIFICATION**
   EN 149:2001+A1:2009 Respiratory protective devices – Filtering half masks to protect against particles - Requirements, testing, marking

4. **PPE PICTURES**



RADEX MFR001G



RADEX MFVR002G

5. **PPE DIMENSIONS:**
   Produduct has been found to be produced using standard size.

6. **PPE PRODUCT MATERIAL INFORMATION:**
   The product is made of elastic strap, exhalation valve, nonwoven fabric on the outer and inner layers and filter material on the middle layer.

7. ESSENTIAL HEALTH AND SAFETY REQUIREMENTS

- A visual inspection was made according to EN 149:2001 +A1:2009 for ergonomics.
- Protection levels and degrees are defined by the manufacturer.
- Suitable construction materials were determined by visual inspection according to EN 149:2001 +A1:2009.

8. ANALYSIS EVALUATION AND MARKING:

EN 149:2001 +A1:2009

TESTS	PARAMETER	PERFORMANCE LEVELS			RESULTS	PERFORMANCE LEVELS	EVALUATION
		FFP1	FFP2	FFP3			
Part 7.3 Visual inspection	Shall also the marking and the information supplied by the manufacturer				Appropriate	-	PASS
Banned Azo Dyes	< 30 mg/kg				Not applicable	-	Not applicable
Part 7.4 Packaging	Particle filtering half mask shall be offered for sale packaged in such a way that they are protected against mechanical damage and contamination before use.				Appropriate	-	PASS
Part 7.5 Material	When conditioned in accordance 8.3.1 & 8.3.2 the particle filter half mask shall not collapse.				Appropriate	-	PASS
Part 7.6 Cleaning and disinfecting	After cleaning and disinfecting the re-usable particle filtering half mask shall satisfy the penetration requirement of the relevant class.				Not applicable	-	Not applicable
Part 7.7 Practical performance	No negative comments should be made by the test subject regarding any of the criteria evaluated.				Appropriate	-	PASS
Part 7.8 Finish of parts	Parts of the device likely to come into contact with the wearer shall have no sharp edge or burrs.				Appropriate	-	PASS

TESTS	PARAMETER	PERFORMANCE LEVELS			RESULTS	PERFORMANCE LEVELS	EVALUATION
		FFP1	FFP2	FFP3			
Part 7.9.1 Total inward leakage	At least 46 out of the 50 individual exercise result	≤25	≤11	≤5	See the table below	FFP3	PASS
	At least 8 out of the 10 individual wearer arithmetic means	≤22	≤8	≤2	See the table below	FFP3	PASS

Total Inward Leakage (%)- Without valve						
	Exercise 1	Exercise 2	Exercise 3	Exercise 4	Exercise 5	Average
Subject 1 (As received)	2,7	1,7	0,9	2,9	1,2	1,9
Subject 2 (As received)	2,4	0,7	0,5	1,2	1,1	1,2
Subject 3 (As received)	0,7	0,9	0,9	0,8	0,6	0,8
Subject 4 (As received)	0,6	0,6	0,7	0,2	0,3	0,5
Subject 5 (As received)	1,8	3,0	2,4	0,1	1,9	1,8
Subject 6 (After temperature conditioning)	0,3	0,3	0,5	0,3	0,4	0,4

Subject 7 (After temperature conditioning)	2,1	2,3	2,0	1,0	1,9	1,9
Subject 8 (After temperature conditioning)	1,2	1,8	1,3	1,1	1,6	1,4
Subject 9 (After temperature conditioning)	0,6	0,6	0,7	0,5	0,3	0,5
Subject 10 (After temperature conditioning)	1,0	0,4	0,7	0,1	0,9	0,6

Total Inward Leakage (%) -With Valve-						
	Exercise 1	Exercise 2	Exercise 3	Exercise 4	Exercise 5	Average
Subject 1 (As received)	1,7	1,5	0,7	0,7	0,7	1,1
Subject 2 (As received)	1,7	0,5	0,3	0,4	0,5	0,7
Subject 3 (As received)	0,5	0,7	0,7	2,2	2,3	1,3
Subject 4 (As received)	0,4	0,4	0,5	0,7	0,8	0,6
Subject 5 (As received)	1,6	1,6	1,2	0,9	1,0	1,3
Subject 6 (After temperature conditioning)	1,8	3,1	3,4	2,9	3,6	3,0
Subject 7 (After temperature conditioning)	1,9	2,1	0,7	0,7	0,4	1,2
Subject 8 (After temperature conditioning)	1,0	1,6	0,4	0,4	0,7	0,8
Subject 9 (After temperature conditioning)	0,4	0,4	1,6	2,2	1,8	1,3
Subject 10 (After temperature conditioning)	0,8	0,2	0,1	0,1	0,4	0,3

Subject facial dimensions

Subject	**Face Length (mm)**	**Face Width (mm)**	**Face Depth (mm)**	**Mouth Width (mm)**
1	133	132	132	65
2	125	144	116	67
3	126	135	124	75
4	123	133	134	74
5	117	135	122	73
6	122	142	133	66
7	113	132	114	75
8	135	123	123	65
9	122	135	133	74
10	135	142	125	83

TESTS	**PARAMETER**	**PERFORMANCE LEVELS**			**RESULTS**	**PERFORMANCE LEVELS**	**EVALUATION**
		FFP1	**FFP2**	**FFP3**			
Part 7.9.2 Penetration of filter material	Sodium chloride, 95 L/min %, max	% 20	% 6	% 1	See the table below	FFP3	PASS
	Paraffin oil, 95 L/min %, max	% 20	% 6	% 1	See the table below	FFP3	PASS

Penetration of filter material - without valve	**Sodium Chloride (%)**	**Paraffin Oil (%)**
As received	0,1	0,1
As received	0,2	0,1
As received	0,1	0,2
After the simulated wearing treatment	0,2	0,2
After the simulated wearing treatment	0,2	0,1
After the simulated wearing treatment	0,1	0,2
Mechanical strength and temperature conditioning (120 mg)	0,2	0,3
Mechanical strength and temperature conditioning (120 mg)	0,2	0,2
Mechanical strength and temperature conditioning (120 mg)	0,1	0,2

Penetration of filter material- with valve	**Sodium Chloride (%)**	**Paraffin Oil (%)**
As received	0,3	0,3

As received	0,4	0,3
As received	0,3	0,4
After the simulated wearing treatment	0,4	0,4
After the simulated wearing treatment	0,3	0,3
After the simulated wearing treatment	0,3	0,4
Mechanical strength and temperature conditioning (120 mg)	0,6	0,8
Mechanical strength and temperature conditioning (120 mg)	0,5	0,8
Mechanical strength and temperature conditioning (120 mg)	0,6	0,6
After clogging	0,5	0,1
After clogging	0,4	0,8
After clogging	0,5	0,7

TESTS	PARAMETER	PERFORMANCE LEVELS			RESULTS	PERFORMANCE LEVELS	EVALUATION
		FFP1	FFP2	FFP3			
Part 7.10 Compatibility with skin	Materials shall not be known to be likely to cause irritation or any other adverse effect to health				Appropriate	-	PASS
Part 7.11 Flammibility	Mask shall not burn or not to continue to burn for more than 5 s				Flame not seen	-	PASS
Part 7.12 Carbondioxide content of the inhalation air	Shall not exceed an average of 1%				Without valve 0,70 0,69 0,74 With Valve 0,77 0,71 0,72	-	PASS
Part 7.13 Head harness	It can be donned and removed easily				Appropriate	-	PASS
Part 7.14 Field of vision	The field of vision shall acceptable in practical performance test.				Appropriate	-	PASS
Part 7.15 Exhalation valve(s)	It shall withstand axially a tensile force of 10 N apply for 10 s. If fitted, shall continue to operate correctly after a continuous exhalation flow of 300 L/min over a period of 30 s.				Appropriate	-	PASS

TESTS	PARAMETER	PERFORMANCE LEVELS			RESULTS	PERFORMANCE LEVELS	EVALUATION
		FFP1	FFP2	FFP3			
Part 7.16 Breathing Resistance	Inhalation 30L/min	0,6 mbar	0,7 mbar	1,0 mbar	See the table below	FFP3	PASS
	Inhalation 95L/min	2,1 mbar	2,4 mbar	3,0 mbar	See the table below	FFP3	PASS
	Exhalation 160L/min	3,0 mbar	3,0 mbar	3,0 mbar	See the table below	FFP3	PASS

Breathing Resistance (mbar)-without valve	Inhalation 30L/min	Inhalation 95L/min
As received	0,5	1,8
As received	0,5	1,7
As received	0,4	1,7
After temperature conditioning	0,4	1,8
After temperature conditioning	0,5	1,7
After temperature conditioning	0,4	1,7
After the simulated wearing treatment	0,4	1,7
After the simulated wearing treatment	0,4	1,8
After the simulated wearing treatment	0,5	1,7
After the flow conditioning	-	-

After the flow conditioning	-	-
After the flow conditioning	-	-

Breathing Resistance (mbar) -with Valve-	Inhalation 30L/min	Inhalation 95L/min
As received	0,5	2,0
As received	0,5	1,9
As received	0,6	2,0
After temperature conditioning	0,5	1,9
After temperature conditioning	0,6	2,0
After temperature conditioning	0,5	2,0
After the simulated wearing treatment	0,5	1,9
After the simulated wearing treatment	0,6	1,9
After the simulated wearing treatment	0,6	2,0
After flow conditioning	0,6	2,0
After flow conditioning	0,5	2,0
After flow conditioning	0,5	1,9

Breathing Resistance After Clogging (mbar)- with valve	Inhalation 95 L/min
Clogging Breathing Resistance	3,5
Clogging Breathing Resistance	3,1
Clogging Breathing Resistance	3,0

Breathing Resistance 160L/min (mbar)- without valve	Facing directly ahead	Facing vertically upwards	Facing vertically downwards	Lying on the left side	Lying on the right side
As received	2,5	2,6	2,5	2,6	2,6
As received	2,6	2,6	2,5	2,6	2,6
As received	2,5	2,5	2,5	2,5	2,6
After temperature conditioning	2,6	2,6	2,6	2,6	2,5
After temperature conditioning	2,6	2,5	2,6	2,5	2,6
After temperature conditioning	2,6	2,6	2,5	2,6	2,6
After the simulated wearing treatment	2,5	2,5	2,6	2,6	2,5
After the simulated wearing treatment	2,6	2,5	2,5	2,6	2,6
After the simulated wearing treatment	2,6	2,6	2,6	2,6	2,5
After flow conditioning	-	-	-	-	-
After flow conditioning	-	-	-	-	-
After flow conditioning	-	-	-	-	-

Breathing Resistance 160L/min (mbar) -With Valve-	Facing directly ahead	Facing vertically upwards	Facing vertically downwards	Lying on the left side	Lying on the right side
As received	2,5	2,5	2,6	2,6	2,5
As received	2,6	2,5	2,5	2,5	2,6
As received	2,5	2,6	2,5	2,5	2,5
After temperature conditioning	2,5	2,5	2,5	2,6	2,6
After temperature conditioning	2,6	2,6	2,6	2,5	2,5
After temperature conditioning	2,6	2,5	2,5	2,5	2,6
After the simulated wearing treatment	2,6	2,6	2,6	2,6	2,6
After the simulated wearing treatment	2,5	2,5	2,5	2,5	2,6
After the simulated wearing treatment	2,6	2,6	2,5	2,6	2,5
After flow conditioning	2,5	2,6	2,5	2,5	2,5
After flow conditioning	2,5	2,5	2,5	2,6	2,6
After flow conditioning	2,6	2,6	2,6	2,5	2,5

Breathing Resistance After Clogging 160 L/min (mbar) Exhalation- with valve	Exhalation- with valve
Clogging Breathing Resistance	2,9
Clogging Breathing Resistance	2,8
Clogging Breathing Resistance	2,7

TESTS	PARAMETER	PERFORMANCE LEVELS			RESULTS	PERFORMANCE LEVELS	EVALUATION
		FFP 1	FFP 2	FFP3			
Part 7.17 Clogging	After clogging the inhalation resistances at 95 L/min shall not exceed. (valved)	4 mbar	5 mbar	7 mbar	3,5 3,1 3,0	FFP3	PASS
	The exhalation resistance shall not exceed 3 mbar at 160 L/ min continuous flow. (valved)				2,9 2,8 2,7	FFP3	PASS
	After clogging the inhalation and exhalation resistances shall not exceed. (valveless)	3 mbar	4 mbar	5 mbar	Not applicable	-	Not applicable
Part 7.18 Demountable part	All demountable parts (if fitted) shall be readily connected and secured were possible by hand.				Not applicable	-	Not applicable
Part 9 Marking	The packaging information shall be clearly and durably marked on the smallest commercially available packaging or legible through it if the packaging is transparent.				Appropriate	-	PASS

9. ATTACHMENTS

- Test Report (M-2021-01701, M-2021-01512, 2024CO6195)

CONTROLLER : Digitally signed by VOLKAN AKIN
Date: 2026.01.26 17:50:46 +03'00'

MNA LABORATORIES
TEST REPORT

Report No: M-2021-01701	Date: 01.12.2021	Page 1 of 4	Rev:

Purpose of Analysis	: SPECIAL REQUEST	**Brand**	:
Sample Type	: PROTECTIVE MASK	**Model**	: MASCARILLAS FFP3 V
Sample Send Org.	: GRAFOPLAS DEL NOROESTE S.A.	**Sampler**	: CUSTOMER
Manufacturer Name	: GRAFOPLAS DEL NOROESTE S.A.		
Analysis Date	: 23.11.2021		
Sample Quantity	: 60 pieces		
Other informations	:		

TESTS	LIMIT	RESULTS
EN ISO 14362-1 / EN ISO 17234-1 Banned Azo Dyes	<30 mg/kg	-
EN 149+ A1 Part 7.9.1 Total inward leakage	At least 46 out of the 50 individual exercise result: FFP1<25 FFP2<11 FFP3<5 At least 8 out of the 10 individual wearer arithmetic means: FFP1<22 FFP2<8 FFP3<2	See below table

Total Inward Leakage (%)
EN 149+ A1 Part 7.9.1

	Exercise 1	Exercise 2	Exercise 3	Exercise 4	Exercise 5	Average
Subject 1 (As received)	1,7	1,5	0,7	0,7	0,7	1,1
Subject 2 (As received)	1,7	0,5	0,3	0,4	0,5	0,7
Subject 3 (As received)	0,5	0,7	0,7	2,2	2,3	1,3
Subject 4 (As received)	0,4	0,4	0,5	0,7	0,8	0,6
Subject 5 (As received)	1,6	1,6	1,2	0,9	1,0	1,3
Subject 6 (After temperature conditioning)	1,8	3,1	3,4	2,9	3,6	3,0
Subject 7 (After temperature conditioning)	1,9	2,1	0,7	0,7	0,4	1,2
Subject 8 (After temperature conditioning)	1,0	1,6	0,4	0,4	0,7	0,8
Subject 9 (After temperature conditioning)	0,4	0,4	1,6	2,2	1,8	1,3
Subject 10 (After temperature conditioning)	0,8	0,2	0,1	0,1	0,4	0,3

Subject facial dimensions

Subject	Face Length (mm)	Face Width (mm)	Face Depth (mm)	Mouth Width (mm)
1	133	132	132	65
2	125	144	116	67
3	126	135	124	75
4	123	133	134	74
5	117	135	122	73
6	122	142	133	66
7	113	132	114	75
8	135	123	123	65
9	122	135	133	74
10	135	142	125	83

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MNA LABORATORIES
TEST REPORT

Report No: M-2021-01701	Date: 01.12.2021	Page 2 of 4	Rev:

Purpose of Analysis	: SPECIAL REQUEST	**Brand**	:
Sample Type	: PROTECTIVE MASK	**Model**	: MASCARILLAS FFP3 V
Sample Send Org.	: GRAFOPLAS DEL NOROESTE S.A.	**Sampler**	: CUSTOMER
Manufacturer Name	: GRAFOPLAS DEL NOROESTE S.A.		
Analysis Date	: 23.11.2021		
Sample Quantity	: 60 pieces		
Other informations	:		

TESTS	LIMIT	RESULTS
EN 149+ A1 Part 7.9.2 Penetration of filter material	Sodium chloride, 95 L/min% FFP1≤20 FFP2≤6 FFP3≤1 Paraffin oil, 95 L/min% FFP1≤20 FFP2≤6 FFP3≤1	See below table

Penetration of filter material EN 149+ A1 Part 7.9.2	**Sodium Chloride (%)**	**Paraffin Oil (%)**
As received	0,3	0,3
As received	0,4	0,3
As received	0,3	0,4
After the simulated wearing treatment	0,4	0,4
After the simulated wearing treatment	0,3	0,3
After the simulated wearing treatment	0,3	0,4
Mechanical strength and temperature conditioning (120 mg)	0,6	0,8
Mechanical strength and temperature conditioning (120 mg)	0,5	0,8
Mechanical strength and temperature conditioning (120 mg)	0,6	0,6

TESTS	LIMIT	RESULTS
EN 149+ A1 Part 7.11 Flammibility	Mask shall not burn or not to continue to burn for more than 5 s	Flame not seen
EN 149+ A1 Part 7.12 Carbondioxide content of the inhalation air	Shall not exceed an average of % 1	0,77 0,71 0,72
EN 149+ A1 Part 7.16 Breathing Resistance	Inhalation 30L/min FFP1≤0,6mbar FFP2≤0,7mbar FFP3≤1,0mbar Inhalation 95L/min FFP1≤2,1mbar FFP2≤2,4mbar FFP3≤3,0mbar Exhalation 160L/min FFP1≤3,0mbar FFP2≤3,0mbar FFP3≤3,0mbar	See below table

EN 149+ A1 Part 7.16 Breathing Resistance (mbar)	**Inhalation 30L/min (mbar)**	**Inhalation 95L/min (mbar)**
As received	0,5	2,0
As received	0,5	1,9
As received	0,6	2,0
After temperature conditioning	0,5	1,9
After temperature conditioning	0,6	2,0
After temperature conditioning	0,5	2,0
After the simulated wearing treatment	0,5	1,9
After the simulated wearing treatment	0,6	1,9
After the simulated wearing treatment	0,6	2,0
After the flow conditioning	0,6	2,0
After the flow conditioning	0,5	2,0

Küçükbakkalköy Mahallesi Yenidogan Caddesi No: 21 Atasehir İstanbul
info@mnalab.com tel: 02165740708 faks: 02165751331

Purpose of Analysis	: SPECIAL REQUEST	**Brand**	:
Sample Type	: PROTECTIVE MASK	**Model**	: MASCARILLAS FFP3 V
Sample Send Org.	: GRAFOPLAS DEL NOROESTE S.A.	**Sampler**	: CUSTOMER
Manufacturer Name	: GRAFOPLAS DEL NOROESTE S.A.		
Analysis Date	: 23.11.2021		
Sample Quantity	: 60 pieces		
Other informations	:		

After the flow conditioning	0,5	1,9

Breathing Resistance 160L/min (mbar) EN 149+ A1 Part 7.16	**Facing directly ahead**	**Facing vertically upwards**	**Facing vertically downwards**	**Lying on the left side**	**Lying on the right side**
As received	2,5	2,5	2,6	2,6	2,5
As received	2,6	2,5	2,5	2,5	2,6
As received	2,5	2,6	2,5	2,5	2,5
After temperature conditioning	2,5	2,5	2,5	2,6	2,6
After temperature conditioning	2,6	2,6	2,6	2,5	2,5
After temperature conditioning	2,6	2,5	2,5	2,5	2,6
After the simulated wearing treatment	2,6	2,6	2,6	2,6	2,6
After the simulated wearing treatment	2,5	2,5	2,5	2,5	2,6
After the simulated wearing treatment	2,6	2,6	2,5	2,6	2,5
After the flow conditioning	2,5	2,6	2,5	2,5	2,5
After the flow conditioning	2,5	2,5	2,5	2,6	2,6
After the flow conditioning	2,6	2,6	2,6	2,5	2,5

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MNA LABORATORIES
TEST REPORT

Report No: M-2021-01701	Date: 01.12.2021	Page 4 of 4	Rev:

Purpose of Analysis	: SPECIAL REQUEST	**Brand**	:
Sample Type	: PROTECTIVE MASK	**Model**	: MASCARILLAS FFP3 V
Sample Send Org.	: GRAFOPLAS DEL NOROESTE S.A.	**Sampler**	: CUSTOMER
Manufacturer Name	: GRAFOPLAS DEL NOROESTE S.A.		
Analysis Date	: 23.11.2021		
Sample Quantity	: 60 pieces		
Other informations	:		

Operating as an experimental laboratory, MNA Laboratories have been accredited by TURKAK with AB-1183-T and TS_EN_ISO / IEC_17025: 2017 standard. Turkish Accreditation Agency (TÜRKAK) signed a multilateral agreement with the European Accreditation Association (EA) on the recognition of test reports and a mutual recognition agreement with the International Laboratory Accreditation Association (ILAC).
*** Analysis is under accreditation.**

Note :

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3. Unsigned and unsealed reports are invalid.
4. This analysis report cannot be used in judicial-administrative procedures and for advertising purposes.
5. Results are valid for the sample as received.
6. The decision rule is the rule that determines how measurement uncertainty is taken into account when specifying the PASS density to a specified specification. According to the TLM-052 Decision Rule Implementation instruction, the Decision Rule Implementation Method selected in agreement with CUSTOMER is clearly stated in the report.
7. Limit Values are determined by taking from analysis methods.
8. The laboratory is not responsible if the information provided by the CUSTOMER affects the validity of the results.
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10. Water Repellency Determination Hydrostatic Pressure Determination TS ISO 811 (Hydrostatic Pressure Tester E / N: 53) Analysis, Seam Strength EN ISO 13965-2 (Strength Test Device E / N: 50) Analysis and resistance to liquid chemical permeation TS EN 659 - A1 Part 3.18 (Liquid Chemical Transfer Device E / N: 107) Analysis is carried out in the conditioning room and ISO 139 PART 3.2 conditions (23 ± 2 ° C temperature and 50 ± 4% relative humidity) are applied for ambient conditions.
11. List of phthalates analyzed is below.
Di-iso-nonyl phthalate (DINP), CAS number: 28553-12-0 or 68515-48-0
Di- (2-ethylhexyl) phthalate (DEHP), CAS number: 117-81-7
Di-n-octyl phthalate (DNOP), CAS number: 117-84-0
Di-iso-decyl phthalate (DIDP), CAS number: 26761-40-0 or 68515-49-1
Butyl benzyl phthalate (BBP), CAS number: 85-68-7
Di-butyl phthalate (DBP), CAS number: 84-74-2

Selin GERGİN
Sampling and Reporting Officer

Erhan ÜSTÜNEL
PPE Laboratory Responsible

Confirmed
01.12.2021
Volkan AKIN
Laboratory Manager

Küçükbakkalköy Mahallesi Yenidogan Caddesi No: 21 Atasehir İstanbul
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MNA LABORATORIES
TEST REPORT

Report No: M-2021-01512	Date: 10.11.2021	Page 1 of 4	Rev:

Purpose of Analysis	:	SPECIAL REQUEST	**Brand**	: -
Sample Type	:	PROTECTIVE MASK	**Model**	: MFR001G
Sample Send Org.	:	GRAFOPLAS DEL NOROESTE, S.A.	**Sampler**	: CUSTOMER
Manufacturer Name	:	GRAFOPLAS DEL NOROESTE, S.A.		
Analysis Date	:	06.10.2021		
Sample Quantity	:	100 pieces		
Other informations	:			

TESTS	LIMIT	RESULTS
EN 149+ A1 Part 7.9.1 **Total inward leakage**	At least 46 out of the 50 individual exercise result: FFP1<25 FFP2<11 FFP3<5 At least 8 out of the 10 individual wearer arithmetic means: FFP1<22 FFP2<8 FFP3<2	See below table

Total Inward Leakage (%)
EN 149+ A1 Part 7.9.1

	Exercise 1	Exercise 2	Exercise 3	Exercise 4	Exercise 5	Average
Subject 1 (As received)	2,7	1,7	0,9	2,9	1,2	1,9
Subject 2 (As received)	2,4	0,7	0,5	1,2	1,1	1,2
Subject 3 (As received)	0,7	0,9	0,9	0,8	0,6	0,8
Subject 4 (As received)	0,6	0,6	0,7	0,2	0,3	0,5
Subject 5 (As received)	1,8	3,0	2,4	0,1	1,9	1,8
Subject 6 (After temperature conditioning)	0,3	0,3	0,5	0,3	0,4	0,4
Subject 7 (After temperature conditioning)	2,1	2,3	2,0	1,0	1,9	1,9
Subject 8 (After temperature conditioning)	1,2	1,8	1,3	1,1	1,6	1,4
Subject 9 (After temperature conditioning)	0,6	0,6	0,7	0,5	0,3	0,5
Subject 10 (After temperature conditioning)	1,0	0,4	0,7	0,1	0,9	0,6

Subject facial dimensions

Subject	Face Length (mm)	Face Width (mm)	Face Depth (mm)	Mouth Width (mm)
1	133	132	132	65
2	125	144	116	67
3	126	135	124	75
4	123	133	134	74
5	117	135	122	73
6	122	142	133	66
7	113	132	114	75
8	135	123	123	65
9	122	135	133	74
10	135	142	125	83

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MNA LABORATORIES
TEST REPORT

Report No: M-2021-01512	Date: 10.11.2021	Page 2 of 4	Rev:

Purpose of Analysis	:	SPECIAL REQUEST	**Brand**	:	-
Sample Type	:	PROTECTIVE MASK	**Model**	:	MFR001G
Sample Send Org.	:	GRAFOPLAS DEL NOROESTE, S.A.	**Sampler**	:	CUSTOMER
Manufacturer Name	:	GRAFOPLAS DEL NOROESTE, S.A.			
Analysis Date	:	06.10.2021			
Sample Quantity	:	100 pieces			
Other informations	:				

TESTS	LIMIT	RESULTS
EN 149+ A1 Part 7.9.2 Penetration of filter material	Sodium chloride, 95 L/min% FFP1≤20 FFP2≤6 FFP3≤1 Paraffin oil, 95 L/min% FFP1≤20 FFP2≤6 FFP3≤1	See below table

Penetration of filter material EN 149+ A1 Part 7.9.2	**Sodium Chloride (%)**	**Paraffin Oil (%)**
As received	0,1	0,1
As received	0,2	0,1
As received	0,1	0,2
After the simulated wearing treatment	0,2	0,2
After the simulated wearing treatment	0,2	0,1
After the simulated wearing treatment	0,1	0,2
Mechanical strength and temperature conditioning (120 mg)	0,2	0,3
Mechanical strength and temperature conditioning (120 mg)	0,2	0,2
Mechanical strength and temperature conditioning (120 mg)	0,1	0,2

TESTS	LIMIT	RESULTS
EN 149+ A1 Part 7.11 Flammibility	Mask shall not burn or not to continue to burn for more than 5 s	Flame not seen
EN 149+ A1 Part 7.12 Carbondioxide content of the inhalation air	Shall not exceed an average of % 1	0,70 0,69 0,74
EN 149+ A1 Part 7.16 Breathing Resistance	Inhalation 30L/min FFP1≤0,6mbar FFP2≤0,7mbar FFP3≤1,0mbar Inhalation 95L/min FFP1≤2,1mbar FFP2≤2,4mbar FFP3≤3,0mbar Exhalation 160L/min FFP1≤3,0mbar FFP2≤3,0mbar FFP3≤3,0mbar	See below table

EN 149+ A1 Part 7.16 Breathing Resistance (mbar)	**Inhalation 30L/min (mbar)**	**Inhalation 95L/min (mbar)**
As received	0,5	1,8
As received	0,5	1,7
As received	0,4	1,7
After temperature conditioning	0,4	1,8
After temperature conditioning	0,5	1,7
After temperature conditioning	0,4	1,7
After the simulated wearing treatment	0,4	1,7
After the simulated wearing treatment	0,4	1,8
After the simulated wearing treatment	0,5	1,7
After the flow conditioning	-	-
After the flow conditioning	-	-
After the flow conditioning	-	-

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MNA LABORATORIES
TEST REPORT

Report No: M-2021-01512	Date: 10.11.2021	Page 3 of 4	Rev:

Purpose of Analysis	:	SPECIAL REQUEST	**Brand**	: -
Sample Type	:	PROTECTIVE MASK	**Model**	: MFR001G
Sample Send Org.	:	GRAFOPLAS DEL NOROESTE, S.A.	**Sampler**	: CUSTOMER
Manufacturer Name	:	GRAFOPLAS DEL NOROESTE, S.A.		
Analysis Date	:	06.10.2021		
Sample Quantity	:	100 pieces		
Other informations	:			

Breathing Resistance 160L/min (mbar) EN 149+ A1 Part 7.16	**Facing directly ahead**	**Facing vertically upwards**	**Facing vertically downwards**	**Lying on the left side**	**Lying on the right side**
As received	2,5	2,6	2,5	2,6	2,6
As received	2,6	2,6	2,5	2,6	2,6
As received	2,5	2,5	2,5	2,5	2,6
After temperature conditioning	2,6	2,6	2,6	2,6	2,5
After temperature conditioning	2,6	2,5	2,6	2,5	2,6
After temperature conditioning	2,6	2,6	2,5	2,6	2,6
After the simulated wearing treatment	2,5	2,5	2,6	2,6	2,5
After the simulated wearing treatment	2,6	2,5	2,5	2,6	2,6
After the simulated wearing treatment	2,6	2,6	2,6	2,6	2,5
After the flow conditioning	-	-	-	-	-
After the flow conditioning	-	-	-	-	-
After the flow conditioning	-	-	-	-	-

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MNA LABORATORIES
TEST REPORT

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Purpose of Analysis	:	SPECIAL REQUEST	**Brand**	:	-
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Sample Send Org.	:	GRAFOPLAS DEL NOROESTE, S.A.	**Sampler**	:	CUSTOMER
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Analysis Date	:	06.10.2021			
Sample Quantity	:	100 pieces			
Other informations	:				

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Erhan ÜSTÜNEL
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Confirmed
10.11.2021
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Laboratory Manager

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info@mnalab.com tel: 02165740708 faks: 02165751331

INFORME DE ENSAYO / *TEST REPORT*

2024CO6195

FECHA DE RECEPCIÓN / ***DATE OF RECEPTION***
Date Format: dd/MM/yyyy 28/10/2024

FECHA DE ENSAYOS / ***DATE TESTS***
Inicio / *Starting*: 28/10/2024
Finalización / *Ending*: 31/10/2024

SOLICITANTE / ***APPLICANT***

GRAFOPLÁS DEL NOROESTE S.A.
LG. A MARISQUEIRA S/N (POL.IND. ALMEIRAS)
ES-15180 CULLEREDO (La Coruña)
España

Att. ESTHER PEREIRA

IDENTIFICACIÓN Y DESCRIPCIÓN DE LAS MUESTRAS/ ***IDENTIFICATION AND DESCRIPTION OF SAMPLES***

Referencia AITEX ***Reference by AITEX***	**Referencia Cliente** ***Reference by customer***	**Descripción AITEX** ***AITEX sample description***
2024CO6195-S01	MFVR002G	Mascarilla blanca con válvula *White facemask with valve*

ENSAYOS REALIZADOS / ***TESTS CARRIED OUT***

- ACONDICIONAMIENTO TÉRMICO / *THERMAL CONDITIONING.*
- OBSTRUCCIÓN / *CLOGGING TEST.*

Asociación de Investigación de la Industria Textil – C.I.F.: G03182870

Los ensayos marcados con * no están amparados por la acreditación de ENAC.
*Tests marked with * are not included within the scope of the ENAC accreditation.*

RESUMEN DE INFORME / *EXECUTIVE SUMMARY*

	Referencia */ Reference*	**Ensayo/Norma** */ Test/Standard*	**Resultado** *Result*
EN 149:2001+A1:2009	2024CO6195-S01	OBSTRUCCIÓN*/ CLOGGING TEST* **EN 149:2001+A1:2009 (Ptos. 8.9,8.10, 8.11) / EN 13274-7:2008**	NIVEL FFP3 *LEVEL FFP3*

RESUMEN DE REQUISITOS / *REQUIREMENT SUMMARY*

OBSTRUCCIÓN
CLOGGING TEST

Requisitos a satisfacer según EN 149:2001+A1:2009 según punto 7.17
Requirements to be met according to EN 149:2001+A1:2009, point 7.17

La resistencia a la respiración especificada no debe excederse antes de que se haya alcanzado una carga de polvo de 833 mg·h/m^3
The specified breathing resistance must not be exceded before a dust load of 833 mg·h/m^3

Requisitos a satisfacer según EN 149:2001+A1:2009 según punto 7.17.2.1
Requirements to be met according to EN 149:2001+A1:2009, point 7.17.2.1

Con válvula/ *with valved*

Resistencia máxima a la respiración *Maximum resistance to breathing* **(mbar)**		
Clasificación *Classification*	**Inhalación 95l/min** *Inhalation 95l/min*	**Exhalación 160l/min** *Exhalation 160l/min*
FFP1	**4,0**	**3,0**
FFP2	**5,0**	
FFP3	**7,0**	

Sin válvula / *without valved*

Resistencia máxima a la respiración *Maximum resistance to breathing* **(mbar)**		
Clasificación *Classification*	**Inhalación 95l/min** *Inhalation 95l/min*	**Exhalación 95l/min** *Exhalation 95l/min*
FFP1	**3,0**	**3,0**
FFP2	**4,0**	**4,0**
FFP3	**5,0**	**5,0**

Requisitos a satisfacer según EN 149:2001+A1:2009 según punto 7.9
Requirements to be met according to EN 149:2001+A1:2009, point 7.9

Classificación *Classification*	**Penetración máxima del aerosol** *Maximum aerosol penetration* **(%)**
FFP1	20
FFP2	6
FFP3	1

DESCRIPCIÓN DE LA MUESTRA/S
SAMPLE/S DESCRIPTION

Referencia por AITEX *AITEX Reference*	**Descripción de la muestra** *Description of the sample*
2024CO6195-S01	Media máscara de partículas con válvula de exhalación que cubre nariz, boca y mentón con arnés de oreja, color blanco. *Particle filtering half mask with exhalation valve covering nose, mouth and chin with ear harness, white colour.*

2024CO6195-S01



2024CO6195-S01



2024CO6195-S01



2024CO6195-S01



///

RESULTADOS/*RESULTS*

ACONDICIONAMIENTO TÉRMICO
THERMAL CONDITIONING

Norma
Standard

EN 149:2001+A1:2009 (punto 8.3.2)
EN 149:2001+A1:2009 (point 8.3.2)

Fecha
Date

Referencia *Reference*	Inicial *Initial*	Final
2024CO6195-S01	28/10/2024	30/10/2024

Descripción del método
Method description

Atmósfera seca < 20%HR, 24 h. a (70 ± 3) ºC y 24h. a (-30 ± 3) ºC.
Dry atmosphere < 20%RH, 24 h. at (70 ± 3) ºC and 24h. at (-30 ± 3) ºC.

_____///

RESULTADOS/*RESULTS*

OBSTRUCCIÓN
CLOGGING TEST

Norma
Standard

EN 149:2001+A1:2009 (punto 8.10)
EN 149:2001+A1:2009 (point 8.10)

Aparato
Apparatus

Equipo generador de aceite de parafina y cloruro sódico y equipo de obstrucción de polvo de dolomita.
Paraffin oil and sodium chloride generator equipment's and dolomite dust clogging equipment.

Condiciones de ensayo
Testing conditioning

Fecha de ensayo *Test date*	Inicial *Initial*	Final
30/10/2024	23,0 ºC / 50 % R.H.	23,3 ºC / 48,3 % R.H.
31/10/2024	22,3 ºC / 50 % R.H.	22,3 ºC / 54,3 % R.H.

Pre-acondicionamiento de muestras
Sample Conditioning

- Muestra 1: En original.
Sample 1: As received.
- Muestra 2 y 3: Acondicionamiento térmico.
Samples 2 and 3: Thermal conditioning.

Incertidumbre del ensayo
Test uncertainty

La incertidumbre expandida para el ensayo de Resistencia a la respiración es ± 1,8% del valor del mesurando, para una probabilidad de cobertura del 95%.
The expanded uncertainty for the Breathing resistance test is ± 1,8% of the value of the measured for a probability of coverage of 95%.
La incertidumbre expandida es para FFP1 ± 14%, para FFP2 ± 18% y para FFP3 ± 23% del valor del mesurando, para una probabilidad de cobertura del 95%.
The expanded uncertainty is for FFP1 ± 14%, for FFP2 ± 18% and for FFP3 ± 23% of the value of the measured for a probability of coverage of 95%.
La incertidumbre expandida del ensayo de penetración de NaCl es ± 15% del valor del mesurando, para una probabilidad de cobertura del 95%.
The expanded uncertainty of NaCl penetration test is ± 15% of the value of the measured for a probability of coverage of 95%.

>>>

Referencia
Reference

2024CO6195-S01

Fecha *Date*	Concentración (mg/m³) *Concentration (mg/m³)*	Tiempo de exposición para 833 mg·h/m³ de polvo de dolomita (min) *Exposure time for 833 mg-h/m³ of dolomite dust (min)*
30/10/2024	440	114
31/10/2024	469	107

Muestra nº. *Sample no.*	Completa el tiempo de exposición para 833 mg·h/m³ de polvo de dolomita. *Complete exposure time for 833 mg-h/m³ of dolomite dust.*	Duración de exposición al polvo de dolomita (min) *Duration of exposure time to dolomite dust (min)*
1	SI *YES*	114
2	SI *YES*	107
3	SI *YES*	107

Resistencia a la respiración durante la exposición al polvo de dolomita
Resistance to breathing during exposure to dolomite dust

Muestra nº. *Sample no.*	Inhalación mbar *Inhalation mbar*		Exhalación mbar *Exhalation mbar*	
	Inicial *Initial*	Final	Inicial *Initial*	Final
1	1,63	3,45	1,36	2,75
2	1,34	3,17	1,08	2,17
3	1,21	2,86	0,96	2,47

Resistencia a la respiración después de la exposición al polvo de dolomita
Breathing resistance after exposure to dolomite dust

Muestra nº. *Sample no.*	Resistencia la inhalación 95L/min mbar *Resistance to breathing 95L/min mbar*
1	3,5
2	3,1
3	3,0

Muestra nº. *Sample no.*	Resistencia a la exhalación 160L/min mbar *Resistance to exhalation 160L/min mbar*
1	2,9
2	2,8
3	2,7

>>>

Penetración después de la exposición al polvo de dolomita
Penetration after exposure to dolomite dust

Ensayo de penetración de aceite de parafina *Paraffin oil penetration test*	
Muestra nº. ***Sample no.***	**Valor medio de penetración %** ***Average value of penetration %***
1	0,10
2	0,80
3	0,71

Ensayo de penetración de cloruro sódico *Sodium chloride penetration test*	
Muestra nº. ***Sample no.***	**Valor medio de penetración %** ***Average value of penetration %***
1	0,50
2	0,35
3	0,53

_____///

Begoña Picó
Responsable Unidad de Gestión Licitaciones Públicas
Head of Public Tenders Division

Date: 31/10/2024 17:57:52
Digitally Signed by:BEGOÑA PICO MONLLOR - NIF:21647435L

CLAUSULAS DE RESPONSABILIDAD

1.- AITEX responde únicamente de los resultados sobre los métodos de análisis empleados, consignados en el informe y referidos exclusivamente a los materiales o muestras que se indican en el mismo y que queden en su poder, limitando a éstos la responsabilidad profesional y jurídica del Centro. Salvo mención expresa, las muestras han sido libremente elegidas y enviadas por el solicitante.
2.- AITEX no se hace responsable en ningún caso del mal uso de los materiales ensayados ni de la interpretación o uso indebido que pueda hacerse de este documento. Los laboratorios de AITEX no realizan muestreos.
3.- La Oferta o Pedido a la que da conformidad el solicitante a través de firma y sello, constituye el Acuerdo Legalmente ejecutable en el que AITEX es responsable de salvaguardar y garantizar, la confidencialidad absoluta, de la gestión de toda la información obtenida o creada durante el desempeño de las actividades contratadas.
4.- Ante posibles discrepancias entre informes, se procederá a una comprobación dirimente en la sede central de AITEX. Asimismo, el solicitante se obliga a notificar a AITEX cualquier reclamación que reciba con causa en el informe, eximiendo a este Centro de toda responsabilidad en caso de no hacerlo así, y considerando los plazos de conservación de las muestras.
5.- AITEX proporcionará a solicitud del interesado, el procedimiento de tratamiento de quejas. En el caso de que se desee poner una, diríjanla a: calidad@aitex.es.
6.- AITEX no se hace responsable de la información proporcionada por los clientes, que se refleja en el Informe, y pueda afectar a la validez de los resultados. La información suministrada por el cliente, no se encuentra cubierta por el alcance de acreditación de ENAC.
7.- AITEX no se hace responsable de un estado inadecuado de la muestra recibida que pudiera comprometer la validez de los resultados, expresando tal circunstancia, en los informes de ensayo.
8.- AITEX podrá incluir en sus informes, análisis, resultados, etc., cualquier otra valoración que juzgue necesaria, aún cuando ésta no hubiere sido expresamente solicitada.
9.- Cuando se solicite Declaración de Conformidad, de no indicarse lo contrario, se aplicará la regla de decisión según ILAC-G8: 2009 con una zona de seguridad de 1U y una Probabilidad de Aceptación Falsa <2,5%.
10.- Las incertidumbres de ensayos, que se explicitan en el Informe de resultados, se han estimado para una k=2 (95% de probabilidad de cobertura). En caso de no informarse, éstas se encuentran a disposición del cliente en AITEX.
11.- Los materiales originales, o muestras sobrantes no sometidas a ensayo, se conservarán en AITEX durante los DOCE MESES posteriores a la emisión del informe, por lo que toda comprobación o reclamación que, en su caso, deseará efectuar el solicitante, se deberá ejercer en el plazo indicado.
12.- Este informe sólo puede enviarse o entregarse en mano al solicitante o a la persona debidamente autorizada por él.
13.- Los resultados de los ensayos y la declaración de cumplimiento con la especificación en este informe se refieren solamente a la muestra de ensayo tal como ha sido analizada/ensayada y no a la muestra/ítem del cual se ha sacado la muestra de ensayo.
14.- El cliente debe prestar atención, en todo momento, las fechas de la realización de los ensayos.
15.- De acuerdo a la Resolución EA (33) 31, los informes de ensayo deben incluir la identificación única de la muestra pudiendo añadirse además cualquier marca o etiquetado del fabricante. No está permitido reemitir informes de ensayo de denominaciones de muestras (referencias) no ensayadas, sólo se pueden volver a reemitir para la corrección de errores o la inclusión de datos omitidos que ya estaban disponibles en el momento del ensayo. El laboratorio no puede asumir la responsabilidad por la que se declara que el producto con el nuevo nombre comercial / marca comercial es estrictamente idéntico al ensayado originalmente; esta responsabilidad es del cliente.
16.- Este informe no podrá ser reproducido parcialmente sin la aprobación por escrito del laboratorio que lo emite.

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. AITEX laboratories do not carry out sampling.
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. The information provided by the client is not covered by the scope of ENAC accreditation.
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 according to ILAC-G8: 2009 will be applied with a security zone of 1U and a Probability of False Acceptance <2.5%.
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.