CE

EG- Konformitätserklärung

Wir **FIT PHARM TECHNOLOGIES GMBH**
48619 Heek,
Benzstr. 13

bescheinigen hiermit in alleiniger Verantwortung, dass das

Produkt : Partikelfiltrierende Halbmaske
Typ : FIT F261
Schutzklasse : FFP2 NR

auf das sich diese Erklärung bezieht, mit den grundlegenden Gesundheitsschutz- und Sicherheitsanforderungen der PSA-Verordnung

- **(EU) 2016/425**

Anhang II unter Anwendung der harmonisierten Norm

- **EN 149:2001+A1:2009**

übereinstimmt. Die Risikoeinstufung erfolgte gem. PSA- Verordnung (EU) 2016/425, Anhang I in Kategorie III.

Durchgeführt wurde die Baumusterprüfung gem. PSA- Verordnung (EU) 2016/425, Anhang V (Modul B) durch die notifizierte Stelle

- **MNA LABORATORIES IND.TRADE.CO.LTD. Türkei, NB-Nr.2841**

die auch die EU- Baumusterprüfungsbescheinigung (Zertifikat-Nr. 77-20-02) ausgestellt hat und die Konformität mit der Bauart auf Grundlage einer internen Fertigungskontrolle mit überwachter Produktprüfung in unregelmäßigen Abständen gem. PSA- Verordnung (EU) 2016/425, Anhang VII, Modul C2 überwacht.

Das Produkt wird durch die FIT PHARM TECHNOLOGIES GMBH 48619 Heek, Benzstr. 13 hergestellt.

Die Erklärung verliert ihre Gültigkeit, wenn das bezeichnete Produkt wesentlich verändert wird.

Stimfort, 19.11.2020 SBoh

Ort, Datum Geschäftsführer

PHARM TECHNOLOGIES
FIT Pharm Technologies GmbH
Benzstraße 13
48619 Heek, Germany

MNA LABORATUVARLARI

Notified Body Number: 2841

AB Tip İnceleme Sertifikası
EU Type-Examination Certificate

Belge No / *Certificate No* : 77-20-02
Belgelendirme Tarihi - Bir Sonraki Belge Tarihi /
Certification Date / Certificate Validity Date : 18.11.2020-18.11.2025
Belge Geçerlilik Tarihi / *Document Validity Period* : 5 yıl / *5 years*
Firma Unvanı ve Adresi /
Company Name and Address : Fit Pharm Technologies GmbH Benzstraße 13 48619 Heek Nordrhein-Westfalen Deutschland

Ürün Adı /Modeller / *Product Name / Models* : FIT F261
Direktifi / *Directive* : 2016/425 REGULATION
Modülü/Kategori / *Module / Category* : B MODÜLÜ/ KATEGORİ III
MODULE B / CATEGORY III
Test Rapor No/ları / *Test Report No* : MNA M-2020-00445
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***: FIT F261 model ürünleri kumaş, kulak kayışı, burun klipsi ve filtre katmanı kullanılarak imal edilmiştir./ *FIT F261 model products are manufactured using fabric, ear strap, nose clip, filter layer.*

Volkan AKIN	***Okan AKEL***
18.11.2020	***18.11.2020***
Karar Verici / Approver	***Şirket Müdürü / General manager***

Notified Body Number: 2841

ATTACHMENTS (77-20-02)

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 : FIT F261

PPE SPECIFICATION	PERFORMANCE LEVELS
Classification	FFP2
Reusable / Single Shift Use	NR

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: Fit Pharm Technologies GmbH
PPE TYPE :

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

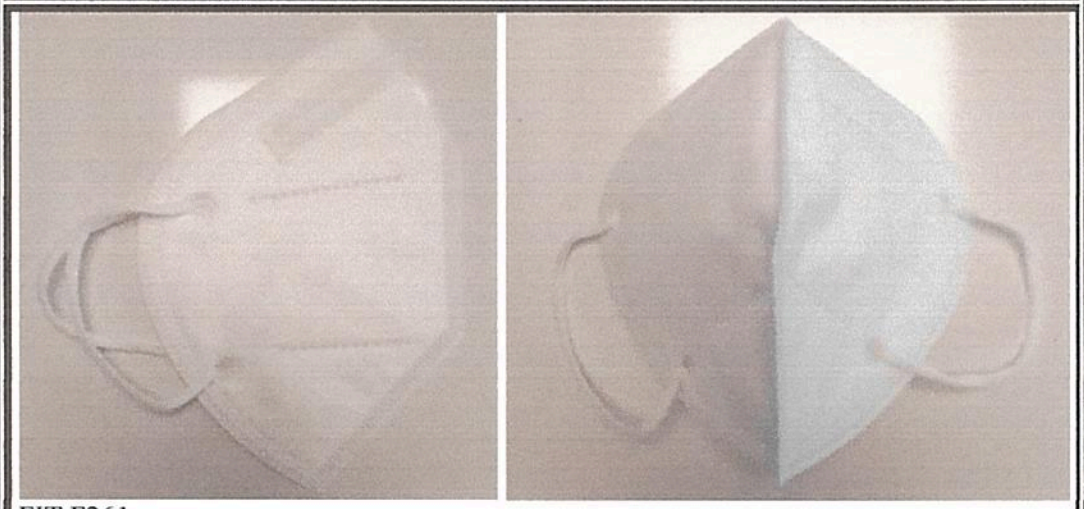
MODEL: FIT F261
PICTOGRAM AND PERFORMANCE LEVELS:

EN 149:2001+ A1:2009 FFP2 NR

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 (77-20-02)

PRODUCT PICTURES

FIT F261

DOCUMENTS IN THE TECHNICAL
- Basic Health Safety Requirements - Risk Assessment - Test Reports - Technical Report

MNA LABORATUVARLARI

Notified Body Number: 2841

CONFORMITY TO TYPE BASED ON INTERNAL PRODUCTION CONTROL PLUS SUPERVISED PRODUCT CHECKS AT RANDOM INTERVALS (MODULE C2)

MODÜL C2 - ÜRETİMİN DÂHİLÎ KONTROLÜ VE ÜRÜNÜN RASTGELE ARALIKLARLA DENETİMLİ MUAYENESİNE DAYALI TİPE UYGUNLUK

Belge No / *Certificate No*	: 59071770
Belgelendirme Tarihi - Bir Sonraki Belge Tarihi / *Certification Date / Certificate Validity Date*	: 18.09.2023-18.09.2024
Belge Geçerlilik Tarihi / *Document Validity Period*	: 1 yıl / *1 year*
Firma Unvanı ve Adresi / *Company Name and Address*	: Fit Pharm Technologies GmbH Industriestraße 45, 48629 Metelen Deutschland
Marka / Model / *Brand / Model*	: FIT F261
Direktifi / *Directive*	: 2016/425 REGULATION
Modülü/Kategori / *Module / Category*	: C2 MODÜLÜ/ KATEGORİ III *MODULE C2 / CATEGORY III*
Teknik Değerlendirme Rapor No/ *Technical Evaluation Report No*	: MNA 59071770

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***: FIT F261 model ürünleri kumaş, elastik kayış, burun klipsi ve filtre katmanı kullanılarak imal edilmiştir./ FIT F261 *model products are manufactured using fabric, elastic strap, nose clip, filter layer.*

Volkan AKIN
18.09.2023
Karar Verici / Approver

Okan AKEL
18.09.2023
Şirket Müdürü / General Manager

Notified Body Number: 2841

CONFORMITY TO TYPE BASED ON INTERNAL PRODUCTION CONTROL PLUS SUPERVISED PRODUCT CHECK AT RANDOM INTERVALS (MODULE C2, ANNEX VII) (59071770)

Report No : 59071770

Report Date : 18.09.2023

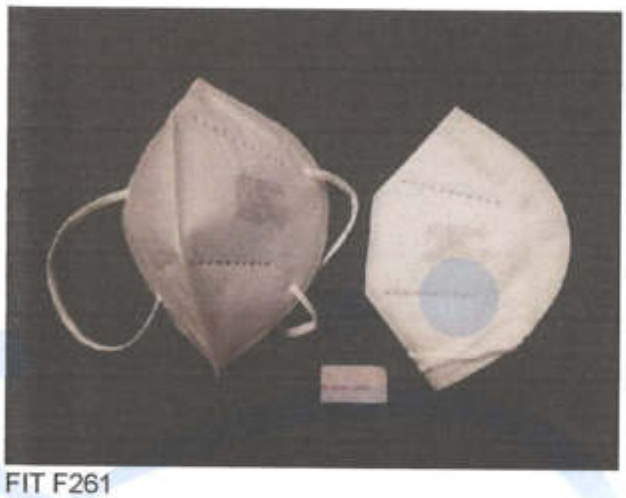
Application No : 59071770

1. **COMPANY INFORMATION:**
   Fit Pharm Technologies GmbH
   Industriestraße 45, 48629 Metelen Deutschland

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**



FIT F261

5. **PPE DIMENSIONS:**
   FIT F261 model has been found to be produced using standard size.

6. **PPE PRODUCT MATERIAL INFORMATION:**
   The mask is made of elastic strap, nonwoven fabric on the outer and inner layers and fitler 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	FFP 2	FFP3			
Part 7.3	Shall also the marking and the information supplied by the manufacturer				Appropriate	-	PASS

Visual inspection				
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	FFP2	PASS
	At least 8 out of the 10 individual wearer arithmetic means	≤22	≤8	≤2	See the table below	FFP2	PASS

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

Subject facial dimensions

Subject	Face Length (mm)	Face Width (mm)	Face Depth (mm)	Mouth Width (mm)
1	120	145	105	61
2	128	155	112	68
3	110	128	105	55

4	123	140	133	57
5	116	128	99	58
6	120	130	91	56
7	138	151	119	65
8	110	130	96	55
9	120	131	85	58
10	135	142	125	83

TESTS	PARAMETER	PERFORMANCE LEVELS			RESULTS	PERFORMANCE LEVELS	EVALUATION
		FFP 1	FFP 2	FFP3			
Part 7.9.2 Penetration of filter material	Sodium chloride, 95 L/min %, max	% 20	% 6	% 1	See the table below	FFP2	PASS
	Paraffin oil, 95 L/min %, max	% 20	% 6	% 1	See the table below	FFP2	PASS

Penetration of filter material	Sodium Chloride (%)	Paraffin Oil (%)
As received	1,5	3,5
As received	1,2	3,0
As received	1,6	3,1
After the simulated wearing treatment	1,6	2,5
After the simulated wearing treatment	1,8	2,9
After the simulated wearing treatment	1,7	2,8
Mechanical strength and temperature conditioning (120mg)	3,3	4,9
Mechanical strength and temperature conditioning (120mg)	2,8	5,1
Mechanical strength and temperature conditioning (120mg)	3,7	4,3

TESTS	PARAMETER	PERFORMANCE LEVELS			RESULTS	PERFORMANCE LEVELS	EVALUATION
		FFP 1	FFP 2	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				0,52 0,58 0,50	-	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.				Not applicable	-	Not applicable

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

Breathing Resistance (mbar)	Inhalation 30L/min	Inhalation 95L/min
As received	0,4	1,2
As received	0,4	1,5
As received	0,5	1,5
After temperature conditioning	0,4	1,2
After temperature conditioning	0,4	1,3
After temperature conditioning	0,4	1,2
After the simulated wearing treatment	0,5	1,2
After the simulated wearing treatment	0,5	1,2
After the simulated wearing treatment	0,4	1,2

Breathing Resistance 160L/min (mbar)	Facing directly ahead	Facing vertically upwards	Facing vertically downwards	Lying on the left side	Lying on the right side
As received	1,9	1,9	1,9	1,8	1,9
As received	1,8	1,8	1,8	1,9	1,9
As received	1,8	1,9	1,9	1,8	1,9
After temperature conditioning	1,6	1,6	1,7	1,7	1,7
After temperature conditioning	1,7	1,7	1,7	1,6	1,7
After temperature conditioning	1,6	1,7	1,6	1,6	1,7
After the simulated wearing treatment	1,8	1,9	1,9	1,8	1,9
After the simulated wearing treatment	1,9	1,9	1,8	1,8	1,8
After the simulated wearing treatment	1,9	1,8	1,8	1,8	1,9

TESTS	PARAMETER	PERFORMANCE LEVELS			RESULTS	PERFORMANCE LEVELS	EVALUATION
		FFP 1	FFP 2	FFP3			
Part 7.17 Clogging	After clogging the inhalation resistances shall not exceed. (valved)	4 mbar	5 mbar	7 mbar	Not applicable	-	Not applicable
	The exhalation resistance shall not exceed 3 mbar at 160 L/ min continuous flow. (valved)				Not applicable	-	Not applicable
	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 Reports (M-2023-0510)

CONTROLLER : **VOLKAN AKIN**

SIGNATURE :

DATE : **18.09.2023**

MNA LABORATORY
ANALYSIS REPORT

Report Nu. : M-2023-0510	Date : 2023-09-13 11:28:00	Page : 1 / 5	Rev:

Purpose of Analysis	: Special request
Sample Send Org.	: FIT Pharm Technologies GmbH
Address	: Industriestr. 45, 48629 Metelen
Sample Acceptance Date	: 2023-08-24 15:54:35
Analysis Date	: 2023-08-25 09:44:33
Sample Quantity	: 120 Pieces
Sample Description	: FIT F261
Other informations	:

Flammability

Device: Flammability tester

Measurement uncertainty:-

Tests	Analysis result	Limit Value	Method	Evaluation	Physical Condition
Flammability	**No flame seen**	**Shall not burn for more than 5 sec after removal from the flame**	**EN 13274-4**	**PASS**	**-**

Penetration Of Filter Material

Device:Filter Test System

Measurement uncertainty:±0,080

Tests	Analysis result	Limit Value	Method	Evaluation	Physical Condition
Penetration Of Filter Material	**Check the table**	**FFP1≤20 FFP2≤6 FFP3≤1**	**EN 149+A1 Part 8.11, EN 13274-7**	**PASS (FFP2)**	**-**

	Sodium Chloride (%)	Paraffin Oil (%)
As received 1	**1,5**	**3,5**
As received 2	**1,2**	**3,0**
As received 3	**1,6**	**3,1**
After the simulated wearing treatment 1	**1,6**	**2,5**
After the simulated wearing treatment 2	**1,8**	**2,9**
After the simulated wearing treatment 3	**1,7**	**2,8**
Mechanical strength and temperature conditioning (120 mg) 1	**3,3**	**4,9**

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

Mechanical strength and temperature conditioning (120 mg) 2	**2,8**	**5,1**
Mechanical strength and temperature conditioning (120 mg) 3	**3,7**	**4,3**

Carbon Dioxide Content Of The Inhalation Air

Device:Carbon DioxideTester

Measurement uncertainty:±0,072

Tests	Analysis result	Limit Value	Method	Evaluation	Physical Condition
Carbon Dioxide Content Of The Inhalation Air	**Check the table**	**Maximum %1**	**EN 149+A1 Part 8.7**	**PASS**	**-**

	CO2 (%)
Sample 1	**0,52**
Sample 2	**0,58**
Sample 3	**0,50**

Total Inward Leakage

Device: Total Inward Leakage Tester

Measurement uncertainty:±0,090

Tests	Analysis result	Limit Value	Method	Evaluation	Physical Condition
Total Inward Leakage	**Check the table**	**See the limits table.**	**EN 149+A1 Part 8.5**	**PASS (FFP2)**	**-**

	At least 46 out of the 50 individual exercise result shall be not greater than	At least 8 out of the 10 individual wearer arithmetic means shall be not greater than
FFP1	**≤25**	**≤22**
FFP2	**≤11**	**≤8**
FFP3	**≤5**	**≤2**

	Exercise 1	Exercise 2	Exercise 3	Exercise 4	Exercise 5	Average
Subject 1 (As received)	**2,1**	**2,6**	**3,5**	**5,2**	**5,1**	**3,7**

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

Breathing Resistance

Device:Breathing Resistance Tester

Measurement uncertainty: Inhalation 30L/min:±0,160,Inhalation30 L/min:±0,026 Exhalation 160 L/min:0,046

Tests	Analysis result	Limit Value	Method	Evaluation	Physical Condition
Breathing Resistance	**Check the table**	**See the limits table.**	**EN 149+A1 Part 8.9**	**PASS (FFP2)**	**-**

Classification	30 L/min max basınç (mbar)	95 L/min max basınç (mbar)	160 L/min max basınç (mbar)
FFP1	**0,6**	**2,1**	**3,0**
FFP2	**0,7**	**2,4**	**3,0**
FFP3	**1,0**	**3,0**	**3,0**

Inhalation	30 L/min	95 L/min
As received 1	**0,4**	**1,2**
As received 2	**0,4**	**1,5**
As received 3	**0,5**	**1,5**

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After temperature conditioning 1	**0,4**	**1,2**
After temperature conditioning 2	**0,4**	**1,3**
After temperature conditioning 3	**0,4**	**1,2**
After the simulated wearing treatment 1	**0,5**	**1,2**
After the simulated wearing treatment 2	**0,5**	**1,2**
After the simulated wearing treatment 3	**0,4**	**1,2**
After the flow conditioning 1	**-**	**-**
After the flow conditioning 2	**-**	**-**
After the flow conditioning 3		

Exhalation 160L/min	**Facing directly ahead**	**Facing vertically upwards**	**Facing vertically downwards**	**Lying on the left side**	**Lying on the right side**
As received 1	**1,9**	**1,9**	**1,9**	**1,8**	**1,9**
As received 2	**1,8**	**1,8**	**1,8**	**1,9**	**1,9**
As received 3	**1,8**	**1,9**	**1,9**	**1,8**	**1,9**
After temperature conditioning 1	**1,6**	**1,6**	**1,7**	**1,7**	**1,7**
After temperature conditioning 2	**1,7**	**1,7**	**1,7**	**1,6**	**1,7**
After temperature conditioning 3	**1,6**	**1,7**	**1,6**	**1,6**	**1,7**
After the simulated wearing treatment 1	**1,8**	**1,9**	**1,9**	**1,8**	**1,9**
After the simulated wearing treatment 2	**1,9**	**1,9**	**1,8**	**1,8**	**1,8**
After the simulated wearing treatment 3	**1,9**	**1,8**	**1,8**	**1,8**	**1,9**
After the flow conditioning 1	**-**	**-**	**-**	**-**	**-**
After the flow conditioning 2	**-**	**-**	**-**	**-**	**-**
After the flow conditioning 3					

Küçükbakkalköy Mahallesi Yenidogan Caddesi No: 21 Ataşehir İstanbul
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MNA LABORATORY
ANALYSIS REPORT

Report Nu. : M-2023-0510	Date : 2023-09-13 11:28:00	Page : 5 / 5	Rev:

Operating as a test laboratory, MNA Laboratories is accredited by TÜRKAK according to AB-1183-T and TS_EN_ISO/IEC_17025:2017 standards has been done. A multilateral agreement with the European Accreditation Association (EA) on the recognition of the Turkish Accreditation Agency (TÜRKAK) test reports and It has signed a mutual recognition agreement with the International Laboratory Accreditation Association (ILAC).

***The analysis is within the scope of accreditation.**

Note :

1. No part of this analysis report may be used alone or separately and may be partially copied or reproduced without the written permission of the laboratory. It cannot be reproduced, used by third parties or as a means of advertising.
2.Analysis results are valid for the sample sent and analyzed by the company/institution/individual to MNA Laboratories. represent the whole may not.
3. Unsigned and Unsealed reports are invalid.
4. This analysis report cannot be used in judicial-administrative proceedings and for advertising purposes.
5. Results are valid for the sample received.
6. A decision rule is a rule that determines how measurement uncertainty is to be taken into account when specifying compliance with a specified specification.TLM-052 Decision Rule According to the implementation instruction, the decision rule chosen in agreement with the customer will be applied if necessary.
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.
9. Test and / or measurement results, expanded measurement uncertainties (if any) and test methods are given in the following pa ges, which are the supplementary part of this certificate.
10. Water Repellency Determination Hydrostatic Pressure Determination T S 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.

Selin Gergin

Sample Acceptance and Reporting Officer

2023-09-13 11:26:30

Erhan Üstünel

Laboratory Responsible

2023-09-13 11:16:59

VOLKAN AKIN

Laboratory Manager

2023-09-13 11:20:35

Küçükbakkalköy Mahallesi Yenidogan Caddesi No: 21 Ataşehir İstanbul
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