

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

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

Belge No / Certificate No : 40091178
**Belgelendirme Tarihi - Bir Sonraki Belge Tarihi /
Certification Date / Certificate Validity Date** : 27.11.2023-27.11.2024
Belge Geçerlilik Tarihi / Document Validity Period : 1 yıl / 1 year
**Firma Unvanı ve Adresi /
Company Name and Address** : PHARMAPLAST S.A.E
Address-1: Amria free zone 23512, Alexandria Egypt.
Address-2: Part number 2, block 7, third industrial zone, Borg el Arab, Alexandria, Egypt.
Address-3: KRE LTD, "Pharmaplast group", Industrial Area, 2830Katunsi, Sandanski, Bulgaria.
Marka / Model / Brand / Model : FC-F2E
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** : 40091178
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: FC-F2E model ürünleri kumaş, elastik kayış, burun klipsi ve filtre katmanı kullanılarak imal edilmiştir./ *FC-F2E model products are manufactured using fabric, elastic strap, nose clip, filter layer.*

Karar Verici / Approver

Şirket Müdürü / General manager



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**CONFORMITY TO TYPE BASED ON INTERNAL
PRODUCTION CONTROL PLUS SUPERVISED PRODUCT
CHECK AT RANDOM INTERVALS
(MODULE C2, ANNEX VII) (40091178)**

Notified Body Number: 2841

Report No : 40091178

Report Date : 27.11.2023

Application No : 40091178

1. COMPANY INFORMATION:

PHARMAPLAST S.A.E

Address-1: Amria free zone 23512, Alexandria Egypt.

Address-2: Part number 2, block 7, third industrial zone, Borg el Arab, Alexandria, Egypt.

Address-3: KRE LTD, "Pharmaplast group", Industrial Area, 2830Katunsi, Sandanski, Bulgaria.

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



FC-F2E (Red, Pink, Blue, Green, Grey, Black, White)

5. PPE DIMENSIONS:

FC-F2E 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 filter material on the middle layer.

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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				<5	-	PASS
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 (%)



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

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
		FFP1	FFP2	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	0,5	0,6
As received	0,5	0,7
As received	0,3	0,4
After the simulated wearing treatment	0,4	0,7
After the simulated wearing treatment	0,5	0,6
After the simulated wearing treatment	0,4	0,8



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Mechanical strength and temperature conditioning (120mg)	2,9	3,8
Mechanical strength and temperature conditioning (120mg)	2,8	3,6
Mechanical strength and temperature conditioning (120mg)	2,9	3,9

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				0,49 0,52 0,53	-	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
		FFP1	FFP2	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		0,5	1,7
As received		0,5	1,7
As received		0,4	1,7
After temperature conditioning		0,5	1,6
After temperature conditioning		0,5	1,6
After temperature conditioning		0,4	1,6
After the simulated wearing treatment		0,4	1,6
After the simulated wearing treatment		0,5	1,7
After the simulated wearing treatment		0,5	1,6



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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	2,8	2,7	2,8	2,7	2,8
As received	2,7	2,7	2,7	2,7	2,7
As received	2,8	2,8	2,8	2,8	2,7
After temperature conditioning	2,8	2,8	2,7	2,8	2,8
After temperature conditioning	2,8	2,8	2,7	2,7	2,7
After temperature conditioning	2,8	2,8	2,8	2,8	2,8
After the simulated wearing treatment	2,8	2,8	2,8	2,8	2,8
After the simulated wearing treatment	2,7	2,7	2,7	2,7	2,7
After the simulated wearing treatment	2,8	2,7	2,8	2,7	2,8

TESTS	PARAMETER	PERFORMANCE LEVELS			RESULTS	PERFORMANCE LEVELS	EVALUATION
		FFP1	FFP2	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-2023-0577,M-2023-0573)

CONTROLLER :

SIGNATURE :

DATE :

AB-1183-T

M-2023-0573

11-23

Report Nu. : M-2023-0573	Date : 2023-11-10 10:54:55	Page : 1 / 3	Rev:
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Purpose of Analysis	: Special request
Sample Send Org.	: Pharmaplast SAE
Address	: Amria Free Zone, 23512, Akexandria, Egypt
Sample Acceptance Date	: 2023-09-20 08:14:03
Analysis Date	: 2023-09-20 13:41:17
Sample Quantity	: 24 Pieces
Sample Description	: Pharmaplast FC-F2E
Other informations	:

Tests	Method	Expected performance level	Evaluation
Banned Azo Dyes	EN ISO 14362-1 / EN ISO 17234-1		PASS

Banned Azo Dyes *

Device:GC-MS

Measurement uncertainty: Textile:±0,350 Leather:±0,390

Compounds/Cas No: biphenyl-4-ylamine, 4-aminobiphenyl xenylamine/92-67-1, benzidine/92-87-5, 4-chloro-o-toluidine/95-69-2, 2-naphthylamine/91-59-8, o-aminoazotoluene, 4-amino-2',3'-dimethylazobenzene, 4-o-tolylazo-o-toluidine/97-56-3, 5-nitro-o-toluidine/99-55-8, 4-chloroaniline/106-47-8, 4-methoxy-m-phenylenediamine/615-05-4, 4,4'-methylenedianiline, 4,4'-diaminodiphenylmethane/101-77-9, 3,3'-dichlorobenzidine, 3,3'-dichlorobiphenyl-4,4'-ylenediamine/91-94-1, 3,3'-dimethoxybenzidine, o-dianisidine/119-90-4, 3,3'-dimethylbenzidine, 4,4'-bi-o-toluidine/119-93-7, 4,4'-methylenedi-o-toluidine/838-88-0, 6-methoxy-m-toluidine p-cresidine/120-71-8, 4,4'-methylene-bis-(2-chloro-aniline), 2,2'-dichloro-4,4'-methylene-dianiline/101-14-4, 4,4'-oxydianiline/101-80-4, 4,4'-thiodianiline/139-65-1, o-toluidine, 2-aminotoluene/95-53-4, 4-methyl-m-phenylenediamine/95-80-7, 2,4,5-trimethylaniline/137-17-7, o-anisidine, 2-methoxyaniline/90-04-0, 4-amino azobenzene/60-09-3

Tests	Analysis result	Limit Value	Method	Evaluation	Physical Condition
Banned Azo Dyes	Check the table for results.	≤30 mg/kg	EN ISO 14362-1 / EN ISO 17234-1	PASS	-

Part of Sample	Results(mg/kg)
Grey Elastic Fabric+Red Fabric+Red Elastic Fabric,Green Fabric+Green Elastic Fabric+Blue Fabric,Blue Elastic Fabric+Pink Elastic Fabric+Pink Fabric,Black Fabric+Black Elastic Fabric+Grey Fabric	<5

CAS No	Substances
92-67-1	4-aminobiphenyl
92-87-5	Benzidine
95-69-2	4-chloro-o-toluidine

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TS EN ISO/IEC 17025
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91-59-8	2-naphthylamine		
97-56-3	o-aminoazotoluene		
99-55-8	5-nitro-o-toluidine		
106-47-8	4-chloroaniline		
615-05-4	2,4-diaminoanisoole		
101-77-9	4,4-methylenedianiline		
91-94-1	3,3-dichlorobenzidine		
119-90-4	3,3-dimethoxybenzidine		
119-93-7	3,3-dimethylbenzidine		
838-88-0	4,4-methylenediotoluidine		
120-71-8	p-cresidine		
101-14-4	2,2-dichloro-4,4-methylene-dianiline		
101-80-4	4,4-oxydianiline		
139-65-1	4,4-thiodianiline		
95-53-4	o-toluidine		
95-80-7	2,4-diaminotoluene		
137-17-7	2,4,5-trimethylaniline		
90-04-0	o-anisidine		
60-09-3	4-aminoazobenzene		

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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.
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3. Unsigned and Unsealed reports are invalid.
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5. Results are valid for the sample received.
6. The decision rule is the rule that determines how measurement uncertainty is taken into account when specifying compliance with an established specification. The customer may choose to apply and/or not apply the decision rule (except in cases where legislation/standards are mandatory). If the customer prefers to apply the decision rule; According to the TLM-052 Decision Rule Application instruction published on the www.mnalab.com website, the decision rule selected in agreement is applied and reported by stating the relevant analysis and decision rule method in the "Note" section. If the customer leaves the decision rule application to the laboratory's evaluation, MNA LABORATORIES applies the simple decision rule.
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 pages, 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 13935-2 (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 $50 \pm 4\%$ relative humidity) are applied for ambient conditions.

Selin Gergin

Sample Acceptance and Reporting Officer

2023-11-10 10:54:07

VOLKAN AKIN

Laboratory Manager

2023-11-10 10:54:36

Erhan Üstünel

Laboratory Responsible

2023-09-28 18:28:52

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ANALYSIS REPORT

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Purpose of Analysis	: Special request
Sample Send Org.	: Pharmaplast SAE
Address	: Amria Free Zone, 23512, Akexandria, Egypt
Sample Acceptance Date	: 2023-09-20 08:14:34
Analysis Date	: 2023-09-20 13:52:16
Sample Quantity	: 120 Pieces
Sample Description	: Pharmaplast FC-F2E
Other informations	:

Tests	Method	Expected performance level	Evaluation
Flammability	EN 13274-4	-	PASS
Breathing Resistance	EN 149+A1 Part 8.9	-	PASS (FFP2)
Total Inward Leakage	EN 149+A1 Part 8.5	-	PASS (FFP2)
Carbon Dioxide Content Of The Inhalation Air	EN 149+A1 Part 8.7	-	PASS
Penetration Of Filter Material	EN 149+A1 Part 8.11, EN 13274-7	-	PASS (FFP2)

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	-

Breathing Resistance

Device: Breathing Resistance Tester

Measurement uncertainty: Inhalation 30L/min: $\pm 0,160$, Inhalation 30 L/min: $\pm 0,026$ 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)	-

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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	0,5	1,7
As received 2	0,5	1,7
As received 3	0,4	1,7
After temperature conditioning 1	0,5	1,6
After temperature conditioning 2	0,5	1,6
After temperature conditioning 3	0,4	1,6
After the simulated wearing treatment 1	0,4	1,6
After the simulated wearing treatment 2	0,5	1,7
After the simulated wearing treatment 3	0,5	1,6
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	2,8	2,7	2,8	2,7	2,8
As received 2	2,7	2,7	2,7	2,7	2,7
As received 3	2,8	2,8	2,8	2,8	2,7
After temperature conditioning 1	2,8	2,8	2,7	2,8	2,8
After temperature conditioning 2	2,8	2,8	2,7	2,7	2,7
After temperature conditioning 3	2,8	2,8	2,8	2,8	2,8
After the simulated wearing treatment 1	2,8	2,8	2,8	2,8	2,8
After the simulated wearing treatment 2	2,7	2,7	2,7	2,7	2,7
After the simulated wearing treatment 3	2,8	2,7	2,8	2,7	2,8
After the flow conditioning 1	-	-	-	-	-

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After the flow conditioning 2	-	-	-
After the flow conditioning 3			

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

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conditioning)			

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	0,49
Sample 2	0,52
Sample 3	0,53

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



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Mechanical strength and temperature conditioning (120 mg) 3	2,9	3,9	

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ANALYSIS REPORT

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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 or for advertising purposes.
5. Results are valid for the sample received.
6. The decision rule is the rule that determines how measurement uncertainty is taken into account when specifying compliance with an established specification. The customer may choose to apply and/or not apply the decision rule (except in cases where legislation/standards are mandatory). If the customer prefers to apply the decision rule; According to the TLM-052 Decision Rule Application instruction published on the www.mnalab.com website, the decision rule selected in agreement is applied and reported by stating the relevant analysis and decision rule method in the "Note" section. If the customer leaves the decision rule application to the laboratory's evaluation, MNA LABORATORIES applies the simple decision rule.
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 pages, 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 13935-2 (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 $50 \pm 4\%$ relative humidity) are applied for ambient conditions.

Selin Gergin

Sample Acceptance and Reporting Officer

2023-11-07 13:43:43

Erhan Üstünel

Laboratory Responsible

2023-11-07 13:16:07

VOLKAN AKIN

Laboratory Manager

2023-11-07 13:17:16