

AB Tip İnceleme Sertifikası EU Type-Examination Certificate

Belge No / Certificate No

Belgelendirme Tarihi - Bir Sonraki Belge Tarihi /

Certification Date / Certificate Validity Date

Belge Geçerlilik Tarihi / Document Validity Period

Firma Unvanı ve Adresi /

Company Name and Address

: JEDX MEDCARE

: 15.03.2024-27.09.2026

: 72031125

: 5 yıl / 5 years

Köysikuja 1, 01640 Vantaa, FINLAND

Marka /Modeller / Brand / Models

: JedX 3805 W FT HLV(White), JedX MIL 5482 FT HLV(Khaki) JedX 3805 W FT HLPV (White) : 2016/425 REGULATION

Direktifi / Directive

Modülü/Kategori / Module / Category

: B MODÜLÜ/ KATEGORİ III MODULE B / CATEGORY III

Teknik Değerlendirme Rapor No/ Technical Evaluation Report No

: MNA 72031125

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: JedX 3805 W FT HLV(White), JedX MIL 5482 FT HLV(Khaki), JedX 3805 W FT HLPV (White) model ürünleri kumaş, elastik kayış, soluk verme valfı, burun klipsi ve filtre katmanı kullanılarak imal edilmiştir./ JedX 3805 W FT HLV(White), JedX MIL 5482 FT HLV(Khaki), JedX 3805 W FT HLPV (White) model products are manufactured using fabric, elastic strap, nose clip, exhalation valve and filter layer.

Revizyon nedeni/ Reason for revision: Sertifikaya farklı valf görseli eklemesi yapılmıştır. / Different valve image has been added to the certificate..

Karar Verici / Approver

Şirket Müdürü / General Manager





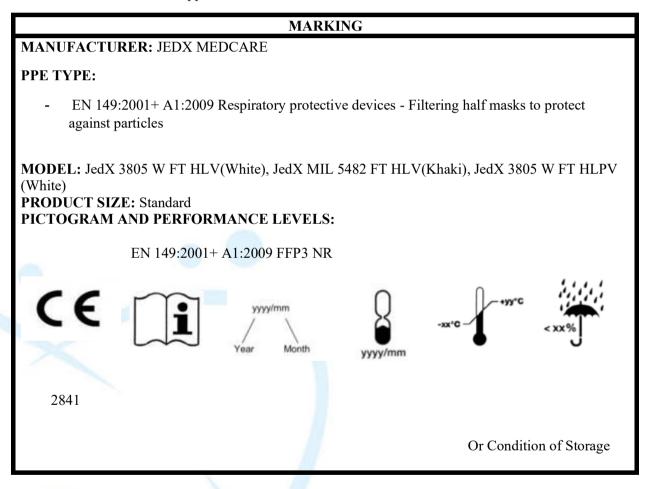
ATTACHMENTS (72031125)

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: JedX 3805 W FT HLV(White), JedX MIL 5482 FT HLV(Khaki), JedX 3805 W FT HLPV (White)

PPE SPECIFICATION	PERFORMANCE LEVELS
Classification	FFP3
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:



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 (72031125)

PRODUCT PICTURES JedX 3805 W FT HLV(White) JedX 3805 W FT HLPV (White)



JedX MIL 5482 FT HLV(Khaki)

DOCUMENTS IN THE TECHNICAL FILE

- Test Reports
- Technical Report

MNA Laboratuvarları San. Tic.Ltd .Şti Adres: Küçükbakkalköy Mahallesi Yenidoğan Cad.No:21 Ataşehir/ İstanbul Tel: 0216 574 07 08 Faks: 0216 575 13 31 www.mnalab.com



TECHNICAL EVALUATION REPORT (72031125)

Report No :72031125

Report Date :15.03.2024

Application No :72031125

1. COMPANY INFORMATION:

JEDX MEDCARE

Köysikuja 1, 01640 Vantaa, FINLAND

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





JedX 3805 W FT HLV(White)

JedX 3805 W FT HLPV (White)



JedX MIL 5482 FT HLV(Khaki)



TECHNICAL EVALUATION REPORT (72031125)

5. PPE DIMENSIONS:

JedX 3805 W FT HLV(White), JedX MIL 5482 FT HLV(Khaki), JedX 3805 W FT HLPV (White) model 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	PERFORMAN CE LEVELS	EVALUATIO N	
		FFP1		FFP3			
Part 7.3 Visual inspection	Shall also the markin supplied by the manuf		ne infor	mation	Appropriate	-	PASS
Banned Azo Dyes	< 30 mg/kg				<5 mg/kg	-	PASS
Part 7.4 Packaging	sale packaged in suc	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 8.3.2 the particle filte collapse.				Appropriate	-	PASS
Part 7.6 Cleaning and disinfecting	particle filtering half i	eaning and disinfecting the re-usable filtering half mask shall satisfy the ation requirement of the relevant			Not applicable	-	Not applicable
Part 7.7 Practical performance		its should be made by ding any of the criteria			Appropriate	-	PASS
Part 7.8 Finish of parts	Parts of the device like with the wearer shall I burrs.				Appropriate	-	PASS

TESTS	PARAMETER	PERFORMANCE LEVELS		RESULTS	PERFORMAN CE LEVELS		
		FFP1	FFP 2	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



TECHNICAL EVALUATION REPORT (72031125)

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

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	Sodium chloride, 95	% 20	% 6	% 1	See the table	FFP3	PASS
Penetration	L/min				below		
of filter	%, max						
material	Paraffin oil, 95 L/min	% 20	% 6	% 1	See the table	FFP3	PASS
	%, max				below		

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



TECHNICAL EVALUATION REPORT (72031125)

TESTS	PARAMETER	PERFORMANCE		PARAMETER PERFORMANCE RESULTS LEVELS		RESULTS	PERFORMANCE LEVELS	EVALUATION
		FFP1	FFP2	FFP3		227220		
Part 7.10	Materials shall not b	e know	n to be	likely to	Appropriate	-	PASS	
Compatibility	cause irritation or an	y other a	adverse	effect to				
with skin	health							
Part 7.11	Mask shall not burn o	or not to	continue	e to burn	Flame not	-	PASS	
Flammibility	for more than 5 s				seen			
Part 7.12	Shall not exceed an	average	of % 1		0,77	-	PASS	
Carbondioxide					0,70			
content of the					0,68			
inhalation air					, , , ,			
Part 7.13	It can be donned and	d remov	ed easily	/	Appropriate	-	PASS	
Head harness								
Part 7.14	The field of vision sha	all accep	table in	practical	Appropriate	-	PASS	
Field of vision	performance test.							
Part 7.15	It shall withstand axi	ally a te	nsile for	ce of 10	Appropriate	-	PASS	
Exhalation	N apply for 10 s.							
valve(s)	If fitted, shall contin							
	after a continuous		on flow	of 300				
	L/min over a period of	ot 30 s.						

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

Breathing Resistance (mbar)	Inhalation 30L/min	Inhalation 95L/min
As received	0,3	1,1
As received	0,3	1,1
As received	0,2	1,0
After temperature conditioning	0,2	1,1
After temperature conditioning	0,2	1,0
After temperature conditioning	0,3	1,0
After the simulated wearing treatment	0,3	1,1
After the simulated wearing treatment	0,3	1,1
After the simulated wearing treatment	0,3	1,0
After the flow conditioning	0,2	1,0
After the flow conditioning	0,2	1,1
After the flow conditioning	0.2	1.1

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	1,5	1,4	1,4	1,4	1,5
As received	1,4	1,5	1,5	1,4	1,4
As received	1,4	1,5	1,4	1,4	1,4
After temperature conditioning	1,5	1,4	1,4	1,4	1,5
After temperature conditioning	1,4	1,5	1,5	1,4	1,4
After temperature conditioning	1,5	1,4	1,4	1,4	1,5

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After the simulated wearing treatment	1,4	1,5	1,5	1,4	1,4
After the simulated wearing treatment	1,4	1,5	1,4	1,4	1,4
After the simulated wearing treatment	1,5	1,4	1,4	1,4	1,5
After the flow conditioning	1,5	1,4	1,4	1,4	1,5
After the flow conditioning	1,4	1,5	1,5	1,4	1,4
After the flow conditioning	1,4	1,5	1,4	1,4	1,4

TESTS	PARAMETER	PERF	ORMAN LS	ICE	RESULTS	PERFORMANCE LEVELS	EVALUATION
		FFP	FFP	FFP3			
		1	2				
Part 7.17	After clogging the	4	5	7	Not applicable	-	Not applicable
Clogging	inhalation	mbar	mbar	mbar			
	resistances shall						
	not exceed.						
	(valved)						
	The exhalation resist				Not applicable	-	Not applicable
	3 mbar at 160 L/ min continuous flow.			s flow.			
	(valved)						
	After clogging the	3	4	5	Not applicable	-	Not applicable
	inhalation and	mbar	mbar	mbar			
	exhalation						
	resistances shall						
	not exceed.						
	(valveless)						
Part 7.18	All demountable par				Not applicable	-	Not applicable
Demountable		connected and secured were		were			
part	· · · · · · · · · · · · · · · · · · ·	ossible by hand.					
Part 9		the packaging information shall be clearly nd durably marked on the smallest ommercially available packaging or legible			Appropriate	-	PASS
Marking							
	1						
	through it if the packaging is transparent.						

9. ATTACHMENTS

• Test Report (M-2021-01464, M-2022-00013)

Reason for Revision: Different valve image has been added to the certificate..

CONTROLLER :

SIGNATURE :

DATE :



Brand

Model

Sampler: CUSTOMER

: JedX 3805 W FT HLV

Report No: M-2021-01464 Date: 24.09.2021 Page 1 of 4 Rev:

Purpose of Analysis : SPECIAL REQUEST

Sample Type : PROTECTIVE MASK

Sample Send Org. : JEDX MEDCARE

Manufacturer Name : JEDX MEDCARE

Analysis Date : 16.09.2021 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

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

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



Report No: M-2021-01464 Date: 24.09.2021 Page 2 of 4 Rev:

Purpose of Analysis : SPECIAL REQUEST Brand

Sample Send Org. : JEDX MEDCARE Sampler : CUSTOMER

Manufacturer Name : JEDX MEDCARE

Analysis Date : 16.09.2021 Sample Quantity : 100 pieces

Other informations :

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

Penetration of filter material EN 149+ A1 Part 7.9.2	Sodium Chloride (%)	Paraffin Oil (%)
As received	0,5	0,4
As received	0,5	0,5
As received	0,4	0,6
After the simulated wearing treatment	0,5	0,5
After the simulated wearing treatment	0,6	0,4
After the simulated wearing treatment	0,5	0,6
Mechanical strength and temperature conditioning (120 mg)	0,7	0,7
Mechanical strength and temperature conditioning (120 mg)	0,7	0,8
Mechanical strength and temperature conditioning (120 mg)	0,8	0,8

TESTS	LIMIT	RESULTS
EN 149+ A1 Part 7.11	Mask shall not burn or not to continue to burn for more than 5 s	Flame not seen
Flammibility		
EN 149+ A1 Part 7.12	Shall not exceed an average of % 1	0,77
Carbondioxide content of the inhalation air		0,70
		0,68
EN 149+ A1 Part 7.16	Inhalation 30L/min FFP1≤0,6mbar FFP2≤0,7mbar FFP3≤1,0mbar	See below table
Breathing Resistance	Inhalation 95L/min FFP1≤2,1mbar FFP2≤2,4mbar FFP3≤3,0mbar	
	Exhalation 160L/min FFP1≤3,0mbar FFP2≤3,0mbar FFP3≤3,0mbar	

EN 149+ A1 Part 7.16	Inhalation 30L/min (mbar)	Inhalation 95L/min (mbar)	
Breathing Resistance (mbar)			
As received	0,3	1,1	
As received	0,3	1,1	
As received	0,2	1,0	
After temperature conditioning	0,2	1,1	
After temperature conditioning	0,2	1,0	
After temperature conditioning	0,3	1,0	
After the simulated wearing treatment	0,3	1,1	
After the simulated wearing treatment	0,3	1,1	
After the simulated wearing treatment	0,3	1,0	
After the flow conditioning	-	-	
After the flow conditioning	-	-	
After the flow conditioning	-	-	



Report No: M-2021-01464 Date: 24.09.2021 Page 3 of 4 Rev:

Purpose of Analysis : SPECIAL REQUEST

Brand

Sample Send Org. : JEDX MEDCARE Sampler : CUSTOMER

Manufacturer Name : JEDX MEDCARE

Analysis Date : 16.09.2021 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 downward s	Lying on the left side	Lying on the right side
As received	1,5	1,4	1,4	1,4	1,5
As received	1,4	1,5	1,5	1,4	1,4
As received	1,4	1,5	1,4	1,4	1,4
After temperature conditioning	1,5	1,4	1,4	1,4	1,5
After temperature conditioning	1,4	1,5	1,5	1,4	1,4
After temperature conditioning	1,5	1,4	1,4	1,4	1,5
After the simulated wearing treatment	1,4	1,5	1,5	1,4	1,4
After the simulated wearing treatment	1,4	1,5	1,4	1,4	1,4
After the simulated wearing treatment	1,5	1,4	1,4	1,4	1,5
After the flow conditioning	-	-	-	-	-
After the flow conditioning	-	-	-	-	-
After the flow conditioning	-	-	-	-	-



Report No: M-2021-01464 Date: 24.09.2021 Page 4 of 4 Rev:

Purpose of Analysis : SPECIAL REQUEST

Sample Type : PROTECTIVE MASK Model : JedX 3805 W FT HLV

Sample Send Org. : JEDX MEDCARE Sampler: CUSTOMER

Manufacturer Name : JEDX MEDCARE

Analysis Date : 16.09.2021 Sample Quantity 100 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:

- 1. No part of this analysis report can be used alone or separately, and may not be partially copied or reproduced, used to third parties and as a means of advertising without the written permission of the laboratory.
- 2. Analysis results are valid for the above mentioned sample sent by MNA Laboratory company / institution / person. It may not represent the whole.
- 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.
- 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 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 \pm 2 $^{\circ}$ C temperature and 50 \pm 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 (DDP), 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**

Brand

Confirmed / 29.04.2021 Volkan AKIN **Laboratory Manager**



MNA LABORATORIES TEST REPORT

Report No: M-2022-00013 Date: 18.01.2022 Page: 1 / 2 Rev:

Purpose of Analysis : SPECIAL REQUEST Brand : JEDX

Sample Type : MASK Model : 3805 MFT HLV FFP3 VALVE

Sample Send Org. : JEDX MEDCARE Sampler : COSTUMER

Manufacturer Name : JEDX MEDCARE

Analysis Date : 05.01.2022 Sample Quantity : 100 pieces

Other informations :

No	Tests	Results	Limit Value	Method	Evaluation	Physical Condition
1	DETERMINATION OF BANNED AZO DYES	< 5 (mg/kg)	P P	TS EN ISO 14362-1+ TS EN ISO 17234-1	PASS	

SAMPLE PLACE

1. Line Sample :Brown fabric



MNA LABORATORIES TEST REPORT

Page:

18.01.2022

Purpose of Analysis : SPECIAL REQUEST **Brand** : JEDX : MASK Model

Date:

Sample Type : 3805 MFT HLV FFP3 VALVE

Sample Send Org. : JEDX MEDCARE Sampler: COSTUMER

Manufacturer Name : JEDX MEDCARE

: 05.01.2022 **Analysis Date** Sample Quantity : 100 pieces

Other informations

Report No: M-2022-00013

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:

- 1. No part of this analysis report can be used alone or separately, and may not be partially copied or reproduced, used to third parties and as a means of advertising without the written permission of the laboratory.
- 2. Analysis results are valid for the above mentioned sample sent by MNA Laboratory company / institution / person. It may not represent the whole.
- 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.
- 9. Test and / or measurement results, expanded measurement uncertainties (if any) and test methods are given in the following pa which are the supplementary part of this certificate.
- 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 \pm 2 $^{\circ}$ C temperature and 50 \pm 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 GERGIN Sampling and Reporting Officer

Erhan ÜSTÜNEL **PPE Laboratory Responsible**

2 / 2

Rev:

Confirmed / 18.01.2022 Volkan AKIN **Laboratory Manager**