

Shenzhen HJR Electronics Technology Co., LTD

EU DECLARATION OF CONFORMITY

PPE: Particulate Respirators (filtering half masks)

This declaration of conformity is issued under the sole responsibility of the manufacturer:

Shenzhen HJR Electronics Technology Co., LTD, 5 / F, building A3, Xinjian Xing science and Technology Industrial Park, No. 3333, GuangQiao Avenue, Gong Ming street, Guang Ming New District, Shenzhen City, Guangdong Province.

Product Model/ Description:

HJR-CN99-06-FFP3 NR Particulate Respirator

The above model is:

A. in conformity with the provisions of Personal Protective Equipment (PPE) Regulation (EU)2016/425 and, where such is the case, with the national standard transposing harmonized standard No.EN149:2001-A1:2009

B. identical to the PPE which is the subject of EU type-examination certificate of conformity No: CE 2163-PPE-772.

This PPE is subject to the conformity assessment procedure based on internal production control plus supervised product checks at random intervals (Module C2)

under surveillance of the notified body

HERE THE NOTIFIED BODY NAME: UNIVERSAL NUMBER: NB2163

Signed for and on behalf of Shenzhen HJR Electronics Technology Co., LTD:

Date: 27 Jul 2020



EUTYPE EXAMINATION CERTIFICATE

Certificate No: 2163-PPE-772

Respiratory protective devices, filtering half masks to protect against particles manufactured by

Shenzhen HJR Electronics Technology Co., Ltd.

5th Floor, A3 Building, Xinjianxing Industrail Park, 3333#, Guanqqiao Avenue, Guangming New District, Shenzhen, Guangdong, China

are tested and evaluated according to

EN 149:2001 + A1:2009 Respiratory Protective Devices -Filtering Half Masks to Protect Against Particles -Requirements, Testing, Marking

Based on the type examination conducted with the evaluation of test reports, technical file according to Personal Protective Equipment Regulation (EU) 2016/425 Annex 5, it is approved that the product meets the requirements of the regulation.

Product Definition

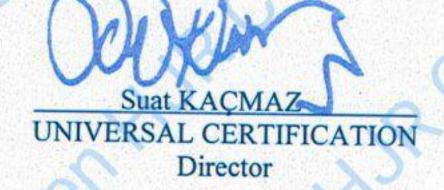
Brand Name: HJR Model: HJR-CN99-06
Filtering half mask
Classification: FFP3 NR

Here by the manufacturer is allowed to use notified body number (2163) and can fix CE mark, as shown below, on the Category III product models given above, with;

- Issuing an appropriate EU Declaration of Conformity according to Personal Protective Equipment Regulation (EU) 2016/425 Annex 9.
- Ongoing successful performance in fulfilment of the requirements set out in Personal Protective Equipment Regulation (EU) 2016/425 and harmonised standards, ensured by assessments based on Annex 7 (Module C2) or Annex 8 (Module D) of the regulation no later than 1 year from the beginning of serial production

This certificate is initially issued on 19/06/2020 and will be valid for 5 years, if there is no change in the relevant harmonised standard affecting the essential health and safety requirements.







erify the validity with the QR code



CERTIFICATE OF CONFORMANCE

Certificate No: 2163-PPE-772/01

Respiratory protective devices, filtering half masks to protect against particles manufactured by

Shenzhen HJR Electronics Technology Co., Ltd.

5th Floor, A3 Building, Xinjianxing Industrail Park, 3333#, Guanqqiao Avenue, Guangming New District, Shenzhen, Guangdong, China

continues to fulfil the requirements of

EN 149:2001 + A1:2009 Respiratory Protective Devices -Filtering Half Masks to Protect Against Particles -Requirements, Testing, Marking

Based on the evaluation of test reports and internal quality control audit reports according to EN 149+A1:2009 and Personal Protective Equipment Regulation (EU) 2016/425 Annex VII (Module C2). This certificate implies that the manufactured products show below are in conformance with the approved EU Type Examination model and meets the requirements of the regulation.

Product Definition

Model	Class	EU Type Examination Certificate			
Model	Class	Serial No	Date	Issuing NB No	
HJR-CN99-06	FFP3 NR	2163-PPE-772	19.06.2020	2163	

Here by the manufacturer is allowed to use notified body number (2163) and can fix CE mark, as shown below, on the Category III product models given above, with;

- Issuing an appropriate EU Declaration of Conformity according to Personal Protective Equipment Regulation (EU) 2016/425 Annex 9.
- Taking all measures necessary so that the manufacturing process and its monitoring ensure the homogeneity of production and conformity of the manufactured PPE with the type described in the EU type examination certificate.

This certificate is issued on 19/06/2020 and will be valid for one year, until 18/06/2021 if the manufacturer makes no major change in the product designs and manufacturing processes affecting the product performance on the essential health and safety requirement.

(E 2163

Suat KAÇMAZ
UNIVERSAL CERTIFICATION
Director

Certificate CN20/42424

The management system of

Shenzhen HJR Electronics Technology Co., Ltd.

5 / F Building A3 Xinjianxing Science and Technology Industrial Park, No. 3333 Guangqiao Avenue, Gongming Street, Guangming New District, Shenzhen City, Guangdong Province, 518106, P.R. China

has been assessed and certified as meeting the requirements of

Regulation (EU) 2016/425

Module D

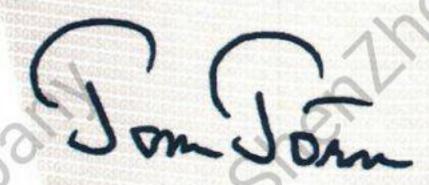
For the following activities

Manufacture of FFP2/FFP3 Protective Respirators (Note: all products marked CE0598 must have a valid EU Type Examination Certificates issued under Module B or a valid EC type examination certificate issued under Article 10 of the PPE Directive 89/686/EEC.)

This certificate is valid from 22 October 2020 until 21 October 2023 and remains valid subject to satisfactory surveillance audits.

Re certification audit due before 6 October 2023 Issue 1. Certified since 22 October 2020

Authorised by



SGS FIMKO OY, Notified Body 0598

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UNIVERSAL CERTIFICATION and SURVEILLANCE SERVICES TRADE CO.

Necip Fazil Bulvari Keyap Sitesi E2 Blok No:44/84 Yukari Dudullu Umraniye, Istanbul / TURKEY

TEST REPORT

Report Date: 18.06.2020

Report Number: 06-2020-T0139

CLIENT and SAMPLE INFORMATION

Appen own up							
TEST OWNER	Shenzhen HJR Electronics	Technology Co., Ltd.					
ADDRESS	5th Floor, A3 Building, Xinjianxing Industrail Park, 3333#, Guanqqiao Avenue, Guangming New District, Shenzhen, Guangdong, China						
SAMPLE DESCRIPTION	Folded Type Filtering Mask						
BRAND NAME - MODEL	HJR-CN99-06 EN 149+A1:2009						
TESTING STANDARD							
CASE NUMBER	CE-PPE-2713						
SAMPLE RECEIVE DATE	24.05.2020 TE	ESTING START DATE 25.05.2020					
DISINFECTION INSTRUCTION If applicable	Not given, single use only						
NUMBER OF SAMPLES	50 SAMPLE I	Ds: 1-46					
AS RECEIVED SAMPLE NO	26-46						
60, 60,	Simulated wearing treatment	1-2-3-4-5-6-7-8-9 (As Received)					
CONDITIONING SAMPLE NO	Temperature conditioning	10-11-12-13-14-15 (Sample after test of Mechanical Strength) 16-17-18-19-20-21-22-23-24-25 (As Received)					
	Mechanical strength	10-17-18-19-20-21-22-23-24-23 (As Received)					

The results given in this test report belongs to the samples tested. The report content cannot be recreated partially without the written consent of UNIVERSAL CERTIFICATION.

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CERTIFICATI
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Telefon: 0216 455 80 80 Faks: 0216 455 80 08
Sarıgazi V.S.Batok & SIVAZ

Director



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500	1. REPORT SUMM	My Kerdy		COLL
SP	TEST STANDARD	TEST NAME	RESULT	EVALUATION
	EN 149:2001 + A1:2009 clause 8.5 EN 13274-1:2001	Total Inward Leakage Testing	Pass	FFP3
1/3	EN 149:2001 + A1:2009 clause 8.11 EN 13274-7:2019	Penetration of Filter Material	Pass	FFP3
	EN 149:2001 + A1:2009 clause 8.6 EN 13274-4:2001 EN 149:2001 + A1:2009	Flammability Testing	Pass	See result
5	clause 8.7 EN 13274-6:2001 EN 149:2001 + A1:2009	Carbon Dioxide Content of The Inhalation Air Testing	Pass	See result
3/10	clause 8.9 EN 13274-3:2001	Breathing Inhalation Resistance-30 l/min Breathing Inhalation Resistance-95 l/min	Pass	See results See results
	EN 149:2001 + A1:2009 clause 8.9 EN 13274-3:2001	Exhalation Resistance, flow rate 160 l/min	Pass	See result
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2. TEST RESULTS and EVALUATION

7.4 PACKAGING (EN 149:2001 + A1:2009 clause 8.2)

Test Method: Clause 8.2-Visual inspection

REQUIREMENT	RESULTS	COMMENT
Particle filtering half masks shall be offered for sale packaged in such a way that they are protected against mechanical damage and contamination before use.	Pass	The masks were packaged in sealed plastic bags, in larger plastic bags inside a large cardboard box that gave some protection against mechanical damage or contamination before use

Lab A

7.5 MATERIAL (EN 149:2001 + A1:2009 clause 8.2, 8.3.1, 8.3.2)

Test Method: Clause 8.2-Visual inspection

Clause 8.3.1-Simulated wearing treatment

A breathing machine is adjusted to 25 cycles/min and 2,0 l/stroke. The particle filtering half mask was mounted on a Sheffield dummy head.

For testing, a saturator is incorporated in the exhalation line between the breathing machine and the dummy head, the saturator being set at a temperature in excess of 37 °C to allow for the cooling of the air before it reaches the mouth of the dummy head.

The air has been saturated at (37 ± 2) °C at the mouth of the dummy head

Clause 8.3.2-Temperature conditioning

The ambient temperature for testing has been between 16 °C and 32 °C and the temperature limits has been subject to an accuracy of ± 1 °C.

- a) for 24 h to a dry atmosphere of (70 ± 3) °C;
- b) for 24 h to a temperature of (-30 ± 3) °C; and allow to return to room temperature for at least 4 h between exposures and prior to subsequent testing. The conditioning has been carried out in a manner which ensures that no thermal shock occurs.

REQUIREMENT	RESULTS	COMMENT
Materials used shall be suitable to withstand handling and wear over the period for which the particle filtering half mask is designed to be used.	Pass	The materials used were able to withstand handling and wear during the limited laboratory testing carried out.
Any material from the filter media released by the air flow through the filter shall not constitute a hazard or nuisance for the wearer.	Pass	It was not constitute a hazard or nuisance for the wearer.
After undergoing the conditioning described in 8.3.1 none of the particle filtering half masks shall have suffered mechanical failure of the facepiece or straps.	Pass	None of the specimens conditioned suffered mechanical failure.
When conditioned in accordance with 8.3.1 and 8.3.2 the particle filtering half mask shall not collapse.	Pass	None of the specimens had not collapse after conditioning.

Lab B

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7.6 CLEANING AND DISINFECTING (EN 149:2001 + A1:2009 clause 8.4, 8.5, 8.11)

Test Method: Described in Clause 8.4, 8.5 and 8.11

REQUIREMENT	RESULTS	COMMENT
If the particle filtering half mask is designed to be re-usable, the materials used shall withstand the cleaning and disinfecting agents and procedures to be specified by the manufacturer. With reference to 7.9.2, after cleaning and disinfecting the re-usable particle filtering half mask shall satisfy the penetration requirement of the relevant class.	N/A	This article is not applicable for tested protective mask which is single use disposable mask.

7.7 PRACTICAL PERFORMANCE (EN 149:2001 + A1:2009 clause 8.4)

Test Method: Described in Clause 8.4

REQUIREMENT	RESULTS	COMMENT
The particle filtering half mask shall undergo practical performance tests under realistic conditions. These general tests serve the purpose of checking the equipment for imperfections that can not be determined by the tests described elsewhere in this standard.	No imperfections	Detail refer to Annex I
Two as received mask samples are used by two subject for the walking (10 mins walking with a speed of 6km/h) and work simulation (bended walking, crawling and basket filling excercises) tests.	(H)	Noth

Annex I-Test Result:

Number of sample: 29 (A.R), 30 (A.R)

Assessed elements	Positive Assessment	Negative Assessment	Requirements in accordance with EN 149:2001+A1:2009	Assessment of Test Result Conformity / Nonconformity
The face piece fitting Head harness comfort Security of fastenings Field of vision	2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	0 0 0 0	Filtering half masks should not have imperfections related to wearer's acceptance	Filtering half masks fulfil requirements of the standard EN 149:2001 + A1:2009 given in 7.7 No imperfections

The subjects (MEG and MA) were able to complete the exercises and did not report any nuisance or problem with the mask.

Lab B

7.8 FINISH OF PARTS (EN 149:2001 + A1:2009 clause 8.2)

Test Method: Described in Clause 8.2

REQUIREMENT	RESULTS	COMMENT
Parts of the device likely to come into contact with the wearer shall have no sharp edges or burrs.	Pass	None of the specimens used in laboratory testing showed evidence of sharp edges or burrs while visual inspection and performance tests.

Lab A

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7.9.1 TOTAL INWARD LEAKAGE (EN 149:2001 + A1:2009 clause 8.5)

Test Method: Described in Clause 8.5

REQUIREMENT	RESULTS	COMMENT
The total inward leakage consists of three components: face seal leakage, exhalation value leakage (if exhalation value fitted) and filter penetration. For particle filtering half masks fitted in accordance with the manufacturer's information, at least 46 out of the 50 individual results shall be not greater than: 25 % for FFP1, 11 % for FFP2, 5 % for FFP3 and in addition at least 8 out of the 10 individual wearer arithmetic means for the total inward leakage shall not be greater than: 22 % for FFP1, 8 % for FFP2, 2 % for FFP3	Pass	Classified as FFP3 Detail refer to Annex II

Annex II-Test Result:

The test results obtained are given in the tables as follows

Test Subject	No of sample	Cond.	1. Walk (%)	Head side/ side (%)	Head up/down (%)	Talk (%)	2. Walk (%)	Average (%)
Ol "	31	A.R.	1,49	1,54	1,57	1,45	1,87	1,58
2	32	A.R.	1,54	1,62	1,68	1,74	1,8	1.68
3	33	A.R.	1,43	1,82	1,28	1,37	1,47	1,47
4	34	A.R.	1,41	1,45	1,43	1,51	1,48	1,46
5	35	A.R.	1,69	1,82	1.7	1,36	1,65	1,64
6	16	T.C.	1,49	1,53	1,71	1,47	1,63	1,57
7 0	17	T.C.	1,77	1,83	1,75	1,79	1,58	1,74
8	18	T.C.	1,68	1,76	1,83	1,87	1,85	1,8
9	19	T.C.	2,3	2,02	2,12	1,83	1,89	2,03
10	20	T.C.	2,45	1,87	1,79	2,1	1,84	2,01
			not greater tha	n 5 %			2	Pass (FFP3)

At least 8 of 10 individual wearer arithmetic means were not greater than 2 %.

		A V			
Test Subject	Face Length (mm)	Face Width (mm)	Face Depth (mm)	Mouth Width (mm)	
	117	1.55	130	60	
2	113	148	128	62	
3	112	160	134	59	
4	115	148	125	61	
5	120	158	132	57	
6	118	150	134	59	
7	115	152	130	57	
8	117	155	134	59	
9	114	149	128	57	
10	110	150	131	55	

For Information Only

Lab B





7.9.2 PENETRATION OF FILTER MATERIAL (EN 149:2001 + A1:2009 clause 8.11)

Test Method: Described in Clause 8.11

	REQUIREMENT	12	RESULTS	COMMENT
Classification	Max penetration	of test aerosol	-10	
46,	NaCl test 95 l/min %max	Paraffin oil test 95 l/min %max	Pass	Detail refer to Annex IIIA and IIIB
FFP1	20	20	110	Detail Telef to Affilex IIIA and IIIB
FFP2	6	6	4	
FFP3		1 1		M. Ila
	C	1		n. ilo

Annex IIIA-Test Result:

The test results obtained are given in the tables as follows;

No. of Sample	Condition	Penetration of Sodium Chloride in accordance with EN 13274-7:2019 [%] Flow rate 95 1/min	Requirements in accordance with EN 149:2001+A1:2009	Assessment of Test Result Conformity / Nonconformity
36 37 38	As received	0,85 0,89 0,82	PED1 < 20.0/	Passed Filtering half masks fulfil
1 2 3	Simulated wearing treatment	0,80 0,74 0,81	FFP1 ≤ 20 % FFP2 ≤ 6 %	the requirements of the standard EN 149:2001+A1:2009 given
10 11 12	Mechanical strength + Temperature conditioned	0,93	FFP3 ≤ 1 %	in 7.9.2 in range of the first, second and third protection class (FFP1, FFP2, FFP3)

Annex IIIB-Test Result:

The test results obtained are given in the tables as follows:

THE LEST I	esuits obtained are given in	the tables as follows,		
No. of Sample	Condition	Penetration of Paraffin Oil Mist in accordance with EN 13274-7:2019	Requirements in accordance with EN	Assessment of Test Result Conformity/
	OV.	[%]	149:2001+A1:2009	Nonconformity
O,	ve,	Flow rate 95 l/min	2	-01
39	6	0.79		Passed
40	As received	0,83	2	1
41		0,82	FFP1 ≤ 20 %	Filtering half masks fulfil
4	Cimulated magning	0,97		the requirements of the
5	Simulated wearing treatment	0,81	FFP2 ≤ 6 %	standard EN
6	treatment	0,82		149:2001+A1:2009 given
13	Mechanical strength +	0.92	FFP3 ≤ 1 %	in 7.9.2 in range of the first,
14	Temperature	0,88	AMO	second and third protection
15	conditioned	0,79		class (FFP1, FFP2, FFP3

Lab A + B

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7.10 COMPATIBILITY WITH SKIN (EN 149:2001 + A1:2009 clause 8.4, 8.5)

Test Method: Described in Clause 8.4 and 8.5.

REQUIREMENT	RESULTS	COMMENT
Materials that may come into contact with the wearer's skin shall not be known to be likely to cause irritation or any other adverse effect to health.	Pass	No irritation or any other adverse effect to health or sensitivity reported by the subjects during the practical performance and TIL tests.

Lab B

7.11 FLAMMABILITY (EN 149:2001 + A1:2009 clause 8.6)

Test Method: Described in Clause 8.6

REQUIREMENT	RESULTS	COMMENT
The material used shall not present a danger for the wearer and shall not be of	4.	(0)
highly flammable nature. When tested, the particle filtering half mask shall not burn or not to continue to burn 5s after removal from the flame.	Part Part	Day 1 - A - W
built of not to continue to built 38 after removal from the flame.	Pass	Detail refer to Annex IV

Annex IV-Test Result: The test results obtained are given in the tables as follows:

No. of Sample	Condition	Visual inspection	Requirements in accordance with EN 149:2001+A1:2009	Assessment of Test Result Conformity / Nonconformity
45	Qd manifold	0,6 s	Filtering half mask	Passed
46	As received	1,1 s	shall not burn or not	Filtering half masks fulfil
21	Temperature	1,1 s	more than 5 s after	requirements of the standard EN 149:2001 +
22	conditioned	1,2 s	removal from the flame	A1:2009 given in 7.11

Lab B

7.12 CARBON DIOXIDE CONTENT OF THE INHALATION AIR (EN 149:2001 + A1:2009 clause 8.7)

Test Method: Described in Clause 8.7

REQUIREMENT	RESULTS	COMMENT
The carbon dioxide content of the inhalation air (dead space) shall not exceed an average of 1,0 % (by volume)	Pass	Detail refer to Annex V

Annex V-Test Result: The test results obtained are given in the tables as follows:

No. of Sample	Condition	CO ₂ content of the inhalation air [%] by volume	An average CO ₂ content of the inhalation air [%] by volume	Requirements in accordance with EN 149:2001+A1:2009	Assessment of Test Result Conformity / Nonconformity
26	111.	0,57	5	CO ₂ content of the	Passed
27	As received	0,63	0,59	inhalation air shall not exceed an average of 1,0% by volume	Filtering half masks fulfil requirements of the standard EN 149:2001 + A1:2009 given in 7.12
28		0,59			

Lab B

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7.13 HEAD HARNESS (EN 149:2001 + A1:2009 clause 8.4, 8.5)

Test Method: Described in Clause 8.4, 8.5

REQUIREMENT	RESULTS	COMMENT
The head harness shall be designed so that the particle filtering half-mask can be donned and removed easily.	Pass	No problem with the head harness reported by the wearers during the practical performance test.
The head harness shall be adjustable or self-adjusting and shall be sufficiently robust to hold the particle filtering half mask firmly in position and capable of maintaining total inward leakage requirements for the device.	Pass	No problem with the head harness reported by the wearers during the practical performance test.

Lab B

7.14 FIELD OF VISION (EN 149:2001 + A1:2009 clause 8.4)

Test Method: Described in Clause 8.4

REQUIREMENT	RESULTS	COMMENT
The field of vision is acceptable if determined so in practical performance tests.	Pass	There were no adverse comments following practical performance tests.

Lab B

7.15 EXHALATION VALVE (EN 149:2001 + A1:2009 clause 8.2, 8.3.4, 8.8, 8.9.1)

Test Method: Clause 8.2, 8.3.4, 8.8, 8.9.1

REQUIREMENT	RESULTS	COMMENT
A particle filtering half mask may have one or more exhalation valve(s), which shall function correctly in all orientations.	N/A	No exhalation valve in tested samples.
If an exhalation valve is provided it shall be protected against or be resistant to dirt and mechanical damage and may be shrouded or may include any other device that may be necessary for the particle filtering half mask to comply with 7.9	N/A	No exhalation valve in tested samples.
Exhalation valve(s), if fitted, shall continue to operate correctly after a continuous exhalation flow of 300 l/min over a period of 30s.	N/A	No exhalation valve in tested samples.
When the exhalation valve housing is attached to the face blank, it shall withstand axially a tensile force of 10N applied for 10s.	N/A	No exhalation valve in tested samples.

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7.16 BREATHING RESISTANCE (EN 149:2001 + A1:2009 clause 8.9)

Test Method: Described in Clause 8.9

	REQU	REMENT	3		RESULTS	COMMENT
		12		40),	· iuli
Classification	Max per	mitted resistance	e (mbar)	0		Classified as FFP3
AL.	Inhal	ation	Exhalation			(a)
(1)	30 l/min	95 l/min	160 l/min		Pass	Detail refer to Annex VIA-VIB
FFP1	0.6	2.1	3.0			- The state of the
FFP2	0.7	2.4	3.0		1 1	
FFP3	1.0	3.0	3.0		M.	
		1			M.	

Annex VIA-Test Result:

The test results obtained are given in the tables as follows;

Inhalation Resistance

mnatation	Resistance					
No. of	Condition	Inhalation Resistance (mbar)				
Sample	478	Flow rate 30 l/min [mbar]	Requirements in accordance with EN 149:2001+A1:2009	Flow rate 95 l/min [mbar]	Requirements in accordance with EN 149:2001+A1:2009	Assessment of Test Result Conformity Nonconformity
42		0,79		2,60	27	
43	As received	0,72	W.A.	2,55		
44		0,75	FFP1 ≤ 0,60	2,66	FFP1 ≤ 2,10	4.
70	Simulated	0,77	60	2,82	2	Passed
8	wearing	0,81	FFP2 ≤ 0,70	2,98	FFP2 ≤ 2,40	Qualifies FFP3
9	treatment	0,79	1/2	2,85		Qualifies 1113
23	Tamanandana	0,81	FFP3 ≤ 1,0	2,82	FFP3 ≤ 3,00	12
24	Temperature conditioned	0,88		2,80		n n
25	conditioned	0,83		2,85	~O.	

Exhalation Resistance

Condition	Flow	Facing	Facing	Facing	Lying	Lying	Requirements in	Assessment
	rate	directly						of Test
		1	upwards	downwards			THE RESERVE ASSESSMENT	Result
				1	15.6 9.5		149:2001+A1:2009	Conformity
		20			side	side	Q.	
		CV.		~				Nonconfor
							2	mity
		2,47	2,53	2,61	2,75	2,75	-	OF
As received		2,45	2,39	2,48	2,65	2,59		
	1	2,55	2,58	2,65	2,70	2,77	FFP1 < 3.0	Passed
Simulated	0,	2,52	2,61	2,62	2,85	2,62		Qualifies
wearing	160l/min	2,50	2,53	2,54	2,79	2,65	FFP2 ≤ 3,0	FFP1,
treatment		2,56	2,57	2,65	2,89	2,93	1),	FFP2, FFP3
T		2,88	2,78	2,72	2,78	2,81	FFP3 ≤ 3,0	.0_
		2,76	2,80	2,89	2,75	2,82		1 11
conditioned		2,82	2,83	2,87	2,79	2,83		
	As received Simulated wearing	As received Simulated wearing treatment Temperature	Conditioned Conditioned	Conditioned Condition Co	As received 2,47 2,53 2,61 2,45 2,39 2,48 2,55 2,58 2,65 2,52 2,61 2,62 2,50 2,53 2,54 2,56 2,57 2,65 2,88 2,78 2,72 2,76 2,80 2,89	Conditioned Condition Co	As received Continue	As received $ \begin{array}{c ccccccccccccccccccccccccccccccccccc$

Lab A



7.17 CLOGGING (EN 149:2001 + A1:2009 clause 8.9, 8.10)

Test Method: Described in Clause 8.8, 8.10

REQUIREMENT	RESULTS	COMMENT
Valved particle filtering half masks: After clogging the inhalation resistances shall not exceed: FFP1:4mbar, FFP2:5mbar, FFP3:7mbar at 95L/min continuous flow. The exhalation resistance shall not exceed 3mbar at 160L/min continuous flow. Valveless particle filtering half masks: After clogging the inhalation resistances shall not exceed: FFP1:3mbar, FFP2:4mbar, FFP3:5mbar at 95L/min continuous flow	NAS (This is optional test and not desired by client.

Lab -

7.18 DEMOUNTABLE PARTS (EN 149:2001 + A1:2009 clause 8.2)

Test Method: Described in Clause 8.2

REQUIREMENT	RESULTS	COMMENT	
All demountable parts (if fitted) shall be readily connected and secured, where possible by hand	N/A	No demountable part.	Noil

Lab .

Pass	Requirement satisfied.	1		
NCR	Requirement not satisfied. Refer to the "Result details" section for more information.			
NAs	Assessment not carried out.	0		
N/A	Requirement not applicable.	2		

LABORATORY INFORMATION

Code	Laboratory Name	Competency Explanations
Lab A	UNIVERSAL SERTIFIKASYON VE GOZETIM HIZMETLERI TIC. LTD. STI.	Internal Laboratory Services of Notified Body
Lab B	GCNTR ULUSLARARASI BELGELENDIRME, GOZETIM, EGITIM VE DIS TICARET LIMITED SIRKETI KOCAELI DILOVA SUBESI	Laboratory holds an accreditation by Turkish Accreditation Agency with number AB-1252-T according to EN ISO/IEC 17025:2017.
•	the laboratories is also under supervision / ass	NIVERSAL CERTIFICATION and the technical competence of essment of UNIVERSAL CERTIFICATION based on the ts for bodies certifying products, processes and services standard. with the issuing laboratory code.

Gr)

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