

IVROU®
IRYS-03 FFP3 NR
EN149:2001+A1:2009
CE 2163

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EN149:2001+A1:2009
CE 2163



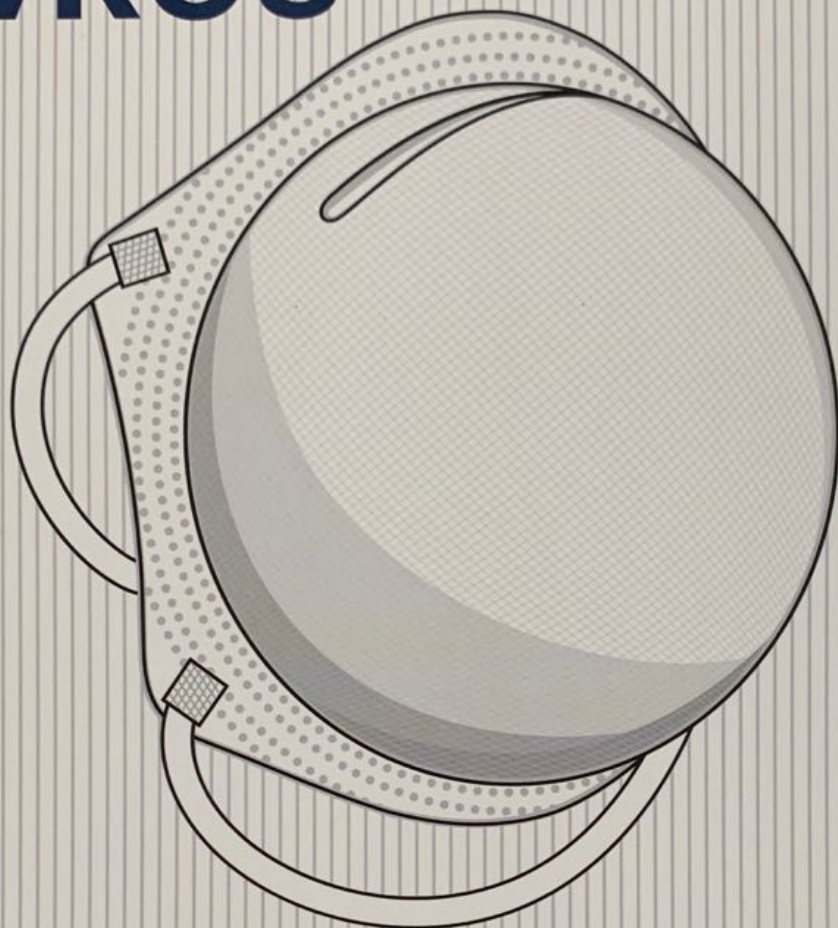




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IVROU®



Automatic absorbing filter with anti-fog

Enhanced respirator

SELF-PRIMING FILTER RESPIRATOR

FFP3 EN 149:2001+A1:2009

MASK / MASQUE / MASKE / MASKER
MASCHERA / MÁSCARA / MASCARAR



SINGLE USE

CE 2163

20_{PCS}

- High filtration efficiency
- Low respiratory resistance
- More comfortable to wear

SELF-PRIMING FILTER RESPIRATOR **IRYS-03**



Face the face of the mask without the nose clip, with the nose clip facing up. Pull the upper headband over the top of the head and the back of the ear.



Pull the lower headband over the top of the head and under the ears.



Place your finger in the middle of the nose clip and press with your fingers towards the sides of the nose.



Cover the mask with both hands and make fine adjustments to ensure that there are no significant air leaks.

NOTICE FOR USE :

- Please do not use this product near a fire source.
- Due to personal differences, fog may appear in the eyes, so please pay more attention when driving.
- As this product is a disposable mask, it can not be re-used through washing.
- Keep out of reach of children.
- Keep away from high temperature's and humidity, and keep in a clean place.
- Use individually packaged products as soon as they are unpacked.

APPLICATION :

It is used in the protection industry for dust generation during construction space, dust prevention, closer metal casting, stone mining, electronics, pharmaceutical, physical processing and grinding, which has good protection against sandstorms, haze and PM2.5. Can effectively protect against pollen, water based and oil based aerosols.

STORAGE CONDITIONS AND METHODS :

Store room temperature 10-30°C at less than 70% relative humidity and avoid direct sunlight exposure. Well ventilated and clean environment without corrosive gas, please do not use if the product gets mildewed or deteriorated due to improper storage.

EU TYPE EXAMINATION CERTIFICATE**Certificate No: 2163-PPE-902**

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

Guangzhou Carrot Mall Network Technologies Co., Ltd.Room 1406, No.4 Zhudian Road (Nansha Street), Nansha District, Guangzhou, Guangdong
Province, China

by the following manufacturer

NANYANG JIUKANG MEDICAL DEVICE CO., LTD.

No.88 Jiulong Road, Nanyang City, Henan, China

are tested and evaluated according to

**EN 149:2001 + A1:2009 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:** IVROU **Model:** IRYS-03

Filtering half mask

Classification: FFP3 NRHere by the manufacturer is allowed to use notified body number 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 **30/06/2020** and will be valid for 5 years, if there is no
change in the relevant harmonised standard affecting the essential health and safety
requirements.**Suat KACMAZ**
UNIVERSAL CERTIFICATION
Director

CERTIFICATE OF CONFORMANCE**Certificate No: 2163-PPE-902/01**

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

Guangzhou Carrot Mall Network Technologies Co., Ltd.Room 1406, No.4 Zhudian Road (Nansha Street), Nansha District, Guangzhou, Guangdong
Province, China

by the following manufacturer

NANYANG JIUKANG MEDICAL DEVICE CO., LTD.

No.88 Jiulong Road, Nanyang City, Henan, 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.

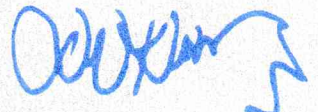
Product Definition

Model	Class	EU Type Examination Certificate		
		Serial No	Date	Issuing NB No
IVROU / IRYS-03	FFP3 NR	2163-PPE-902	30.06.2020	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 **30/06/2020** and will be valid for one year, until **29/06/2021** 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.


Suat KAÇMAZ
UNIVERSAL CERTIFICATION
Director

EU DECLARATION OF CONFORMITY

This Declaration of Conformity, issued under the sole responsibility of the manufacturer

Guangzhou Carrot Mall Network Technologies Co., Ltd.

Room 1406, No.4 Zhudian Road (Nansha Street), Nansha District,

Guangzhou, Guangdong Province, China

EC Representative: Unimama B.V., Lutkemeerweg 298, 1067 TH Amsterdam, Netherlands

hereby declaring the following Personal Protective Equipment (PPE)

Product Description: IVROU Filtering Half Mask

Product Model/s: IRYS-03 FFP3 NR

is/are in conformity with the provisions of the following European Regulation

PPE (Personal Protective Equipment) Regulation

The model is/are in conformity with the provisions of Regulation (EU) 2016/425, and with the National Standard transposing the harmonised European Standard Number(s):

EN 149:2001+A1:2009

and is/are identical to the PPE which is/are the subject of EU type-examination (Module B of Regulation (EU) 2016/425) referenced on the certificate number:

Certificate No.: 2163-PPE-902 (Issue Date: 30/06/2020)

issued by

Universal Certification and Surveillance Service Trade Ltd. Co.

Necip Fazıl Bulvarı Keyap Sitesi E2 Blok No: 44/84 Yukarı Dudullu

Ümraniye-Istanbul, Turkey (Notified Body No. 2163)

and is/are in conformance with procedures set out in Module C2 of Regulation (EU) 2016/425 under the evaluation of Universal Certification and Surveillance Service Trade Ltd. Co. (Notified Body No. 2163), referenced on the Certificate No: 2163-PPE-902/01 (Issue Date: 30/06/2020).

Guangzhou, China, 01/07/2020

Chen, Xiaoying

Quality Manager

Guangzhou Carrot Mall Network Technologies Co., Ltd.



UNIVERSAL CERTIFICATION and SURVEILLANCE SERVICES TRADE CO.
Necip Fazil Bulvarı Keyap Sitesi E2 Blok No:44/84 Yukarı Dudullu Ümraniye, İstanbul / TURKEY

TEST REPORT

Report Date: 30.06.2020

Report Number: 06-2020-T-0177

CLIENT and SAMPLE INFORMATION

TEST OWNER	Guangzhou Carrot Mall Network Technologies Co., Ltd		
ADDRESS	Room 1406, No.4 Zhudian Road (Nansha Street), Nansha District, Guangzhou, Guangdong Province, China		
SAMPLE DESCRIPTION	Cone type filtering half mask (See end of test report for sample photo)		
BRAND NAME – MODEL	IVROU / IRYS-03		
TESTING STANDARD	EN 149+A1:2009		
CASE NUMBER	CE-PPE-2876		
SAMPLE RECEIVE DATE	01.06.2020	TESTING START DATE	01.06.2020
DISINFECTION INSTRUCTION <i>If applicable</i>	Not given, single use only		
NUMBER OF SAMPLES	50	SAMPLE IDs:	1 – 46
AS RECEIVED SAMPLE NO	26-46		
CONDITIONING SAMPLE NO	Simulated wearing treatment	1-2-3-4-5-6-7-8-9 (As Received)	
	Temperature conditioning	10-11-12-13-14-15 (Sample after test of Mechanical Strength)	
	Mechanical strength	16-17-18-19-20-21-22-23-24-25 (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.

UNIVERSAL
CERTIFICATION
Necip Fazil Bulvarı, Keyap Sitesi, E2 Blok, No:44/84
Yukarı Dudullu Ümraniye/İSTANBUL
Telefon: 0216 455 80 80 Faks: 0216 455 80 08
Sergazî V.D. 892 025 8722

Suat KAÇMAZ
Director

1. REPORT SUMMARY

TEST STANDARD	TEST NAME	RESULT	EVALUATION
EN 149:2001 + A1:2009 clause 8.5 EN 13274-1:2001	Total Inward Leakage Testing	Pass	FFP3
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	Flammability Testing	Pass	See result
EN 149:2001 + A1:2009 clause 8.7 EN 13274-6:2001	Carbon Dioxide Content of The Inhalation Air Testing	Pass	See result
EN 149:2001 + A1:2009 clause 8.9 EN 13274-3:2001	Breathing Inhalation Resistance-30 l/min	Pass	See results
	Breathing Inhalation Resistance-95 l/min	Pass	See results
EN 149:2001 + A1:2009 clause 8.9 EN 13274-3:2001	Exhalation Resistance, flow rate 160 l/min	Pass	See results

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 $(37 \pm 2) ^\circ\text{C}$ 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 $\pm 1 ^\circ\text{C}$.

- a) for 24 h to a dry atmosphere of $(70 \pm 3) ^\circ\text{C}$;
- b) for 24 h to a temperature of $(-30 \pm 3) ^\circ\text{C}$; 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

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. 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 exercises) tests.	No imperfections	Detail refer to Annex I

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	2	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
Head harness comfort	2	0		
Security of fastenings	2	0		
Field of vision	2	0		

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

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 valve leakage (if exhalation valve 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 (%)
1	31	A.R.	1.48	1.53	1.45	1.38	1.59	1.49
2	32	A.R.	1.46	1.55	1.51	1.42	1.47	1.48
3	33	A.R.	1.43	1.49	1.56	1.47	1.50	1.49
4	34	A.R.	1.45	1.61	1.57	1.48	1.56	1.53
5	35	A.R.	1.49	1.52	1.57	1.50	1.53	1.52
6	16	T.C.	1.52	1.57	1.65	1.62	1.58	1.59
7	17	T.C.	1.62	1.69	1.71	1.56	1.57	1.63
8	18	T.C.	1.65	1.69	1.76	1.73	1.77	1.72
9	19	T.C.	1.69	1.75	1.86	1.79	1.83	1.78
10	20	T.C.	1.75	1.79	1.86	1.92	1.95	1.85

All 50 individual exercise results were not greater than 5 %.

At least 9 of 10 individual wearer arithmetic means were not greater than 2 %.

Pass (FFP3)

Test Subject	Face Length (mm)	Face Width (mm)	Face Depth (mm)	Mouth Width (mm)
1	117	155	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

7.10 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 TH 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 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.	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	As received	0 s	Filtering half mask shall not burn or not continue to burn for more than 5 s after removal from the flame	Passed
46		0 s		Filtering half masks fulfil requirements of the standard EN 149:2001 + A1:2009 given in 7.11
21	Temperature conditioned	0 s		
22		0 s		

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	As received	0.65	0.65	CO ₂ content of the inhalation air shall not exceed an average of 1.0% by volume	Passed Filtering half masks fulfil requirements of the standard EN 149:2001 + A1:2009 given in 7.12
27		0.67			
28		0.63			

Lab B

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.

Lab -

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

Test Method: Described in Clause 8.9

REQUIREMENT				RESULTS	COMMENT	
Classification	Max permitted resistance (mbar)			Pass	Classified as FFP3 Detail refer to Annex VIA-VIB	
	Inhalation		Exhalation			
	30 l/min	95 l/min	160 l/min			
	FFP1	0.6	2.1			3.0
	FFP2	0.7	2.4			3.0
FFP3	1.0	3.0	3.0			

Annex VIA-Test Result:

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

Inhalation Resistance

No. of Sample	Condition	Inhalation Resistance (mbar)				Assessment of Test Result Conformity / Nonconformity	
		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		
42	As received	0.65	FFP1 ≤ 0.60	2.25	FFP1 ≤ 2.10	Passed Qualifies FFP3	
43		0.68		2.31			
44		0.71		2.33			
7	Simulated wearing treatment	0.73		2.34			
8		FFP2 ≤ 0.70		2.36			FFP2 ≤ 2.40
9				0.76			
23	0.78			2.38			
24	Temperature conditioned	0.82		2.36			
25		0.84		2.37			
		0.87					

Exhalation Resistance

No. of Sample	Condition	Flow rate	Facing directly	Facing vertically upwards	Facing vertically downwards	Lying on the left side	Lying on the right side	Requirements in accordance with EN 149:2001+A1:2009	Assessment of Test Result Conformity / Nonconformity
42	As received	160l/min	2.60	2.65	2.67	2.69	2.71	FFP1 ≤ 3.0	Passed Qualifies FFP1, FFP2, FFP3
43			2.51	2.53	2.55	2.57	2.56		
44			2.53	2.58	2.63	2.65	2.60		
7	Simulated wearing treatment		2.60	2.64	2.67	2.75	2.77	FFP2 ≤ 3.0	
8			2.56	2.59	2.67	2.72	2.75		
9			2.61	2.65	2.68	2.71	2.70		
23	Temperature conditioned		2.65	2.69	2.72	2.76	2.73	FFP3 ≤ 3.0	
24			2.67	2.69	2.73	2.78	2.75		
25			2.69	2.71	2.75	2.79	2.82		

Lab A

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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.

Lab -

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

LABORATORY INFORMATION

Code	Laboratory Name	Competency Explanations
Lab A	UNIVERSAL SERTİFİKASYON VE GÖZETİM HİZMETLERİ TİC. LTD. ŞTİ.	Internal Laboratory Services of Notified Body
Lab B	GCNTR ULUSLARARASI BELGELENDİRME, GÖZETİM, EĞİTİM VE DİŞ TİCARET LIMITED SİRKETİ KOCAELİ DİLOVA SUBESİ	Laboratory holds an accreditation by Turkish Accreditation Agency with number AB-1252-T according to EN ISO/IEC 17025:2017.
<ul style="list-style-type: none"> The laboratories are contracted bodies with UNIVERSAL CERTIFICATION and the technical competence of the laboratories is also under supervision / assessment of UNIVERSAL CERTIFICATION based on the provisions of EN ISO/IEC 17065 Requirements for bodies certifying products, processes and services standard. Each test result given in this test report shown with the issuing laboratory code. 		



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VE GÖZETİM HİZM.
TİC. LTD. ŞTİ.
Necip Fazıl Bulvarı, Keleş Sitesi, E2 Blok, No:44/4
Yükarı Dudulu-Ömerliye/ŞANLIURFA
Tic. Sic. No: 0216 455 80 80 Faks: 0216 455 80 08
Sarıgül V.D. 892 025 8722

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Sample Photos of Packaging and Mask itself



• End of Report •

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SERTİFİKASYON
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TİC. LTD. ŞTİ.
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