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The Testing Center is accredited for compliance with ISO/IEC 17025:2017.

The results of tests, calibrations and/or measurements included in this document are traceable to Chinese/national standards.

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TEST REPORT

Selected test items from EN 149:2001+A1:2009 Respiratory protective devices—Filtering half masks to protect against particles—Requirements, testing, marking
According to supervised product checks procedure at random intervals (C2)

The following samples were submitted and identified on behalf of the client as:

Product : Particle Filtering Half Mask
Report No. : KZ2022018
Client : Universal Certification and Surveillance Service Trade Ltd.Co.
Model(s) : YY0525
Number of samples : 30
Received date : 2020.08.17
Date(s) of tests : 2020.08.17-2020.08.17

DESCRIPTION OF SAMPLES

General information

Manufacturer
Manufacturer address

Classification
FFP2

JIANGMEN YANYANG TRADING CO.,LTD.
NO.1,4th Floor, Building 2, NO.18 Xinyi Road, Jianghai District,
Jiangmen City, Guangdong Province, China

Main components
White folding mask

Chief Tester: 冯云
Feng Yun

Reviewer: 傅丹华
Fu Danhua

Approve: 傅科杰
Fu Kejie
Authorized Signatory,
Lab Director



Issued: 2020.08.17



Page 1 of 5

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Test Report No.KZ2022018**Conditions:**

The test results presented in this report relate to the samples tested only.

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The authenticity of this test report and its contents can be verified by contacting the laboratory.

Conclusion**Test Items**

Clause 7.9.2	Penetration of filter material	Pass
Clause 7.16	Breathing resistance	Pass

Remarks: Pass = Meet EN 149:2001+A1:2009 FFP2 Requirement
Fail = Below EN 149:2001+A1:2009 FFP2 Requirement
N/A = Not Applicable

Disclaimer Measurement Uncertainty:

Unless otherwise agreed upon, Pass or Fail verdicts are given based on the measured values without any considerations of measurement uncertainties. Please note, every test method has a measurement uncertainty which has been evaluated by the laboratory according to ISO/IEC 17025 requirements.

By taking measurement uncertainties into account it might happen that measured values can neither be assessed as Pass nor as Fail.

Test Report No.KZ2022018

Test Results

7.9.2 Penetration of filter material

Pass¹

The penetration of the filter of the particle filtering half mask shall meet the following requirements.

Sodium chloride test 95 L/min

Paraffin-oil test 95 L/min

FFP1	≤20%	≤20%
FFP2	≤6%	≤6%
FFP3	≤1%	≤1%

Note 1: FFP2 respirator. Test results are shown in Annex A Table 7.9.2.

7.16 Breathing resistance

Pass²

Classification	Maximum permitted resistance (mbar)		
	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

Note 2: FFP2 respirator. Test results are shown in Annex A Table 7.16.

End of Test Results

Test Report No.KZ2022018

Annex A: Summarization of Test Data

Table 7.9.2 Penetration of filter material

Test specification: EN 149:2001+A1:2009 Clause 8.11

Test specification: EN 14725-1:2007 Clause 6.11				
Aerosol	Condition	Sample No.	Penetration (%)	Assessment
Sodium chloride test	As received	01	0.60	Pass
		02	0.23	
		03	0.17	
Paraffin oil test	As received	04	0.72	
		05	0.64	
		06	0.54	
Flow conditioning: single filter; 95.0 L/min				

Table 7.16 Breathing resistance (mbar)

Test specification: EN 149:2001+A1:2009 Clause 8.9

As received	Flow rate		07					08					09				
			A	B	C	D	E	A	B	C	D	E	A	B	C	D	E
	Inhalation	30 L/min	0.55	0.58	0.59	0.50	0.59	0.52	0.54	0.62	0.56	0.57	0.51	0.52	0.58	0.55	0.57
		95 L/min	1.75	1.80	1.82	1.55	1.30	1.59	1.75	1.85	1.72	1.84	1.62	1.71	1.82	1.61	1.85
	Exhalation	160 L/min	2.41	2.81	2.67	2.17	2.86	2.61	2.63	2.71	2.59	2.78	2.45	2.72	2.69	2.37	2.81
Assessment			Pass														

A: facing directly ahead; B: facing vertically upwards; C: facing vertically downwards; D: lying on the left side; E: lying on the right side

End of Annex A

Test Report No.KZ2022018

Annex B: Photos of sample



End of Annex B

UNIVERSAL



EU TYPE EXAMINATION CERTIFICATE

Certificate No: 2163-PPE-834

Respiratory protective devices, filtering half masks to protect against particles manufactured by
JIANGMEN YANYANG TRADING CO., LTD.
No.1, 4th Floor, Building 2, No. 18 Xinyi Road, Jianghai District, Jiangmen City, Guangdong
Province, 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: CRDLIGHT **Model:** YY0525
Filtering half mask
Classification: FFP2 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 **25/06/2020** 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



Verify the validity with the QR code

TECHNICAL ASSESSMENT REPORT

REPORT DATE / NO: 25.06.2020 / 2163-KKD-834

Manufacturer: JIANGMEN YANYANG TRADING CO., LTD.

Address: No.1, 4th Floor, Building 2, No. 18 Xinyi Road, Jianghai District, Jiangmen City, Guangdong Province, China

This report is for the, given above, manufacturer prepared according to the test results obtained from Trust Right Testing and Certification Service (Zhongshan) Ltd, accredited by IAS (International Accreditation Service), signatory to ILAC MRA, with number TL-861 for the product identified below, dated 15.06.2020 with Serial Id R20200062 based on EN 149: 2001 + A1: 2009 standard and the technical file dated 19 June, 2020 Version 01 provided by the manufacturer.

The technical file of the manufacturer, and risk evaluation against the essential health safety requirements and the test report evaluated for their relation with Essential Requirements of Personal Protective Equipment Regulation and found to be appropriate.

This report is an annex and an integral part of the EU Type Examination Certificate issued to the manufacturer. The test results and issued certificate belongs only to the tested model. The technical report consists of a total of 6 pages.

Product Description: Particle Filtering Half Mask

Classification: FFP2 NR

Brand Name: CRDLIGHT **Model:** YY0525



**THE CLAUSES OF EN 149: 2001 + A1: 2009 STANDARD RELATED TO EUROPEAN UNION DIRECTIVE
EU 2016/425 REQUIREMENTS**

1.1. Design principles

1.1.1. Ergonomics

PPE must be so designed and manufactured that in the foreseeable conditions of use for which it is intended the user can perform the risk related activity normally whilst enjoying appropriate protection of the highest possible level.

1.1.2. Levels and classes of protection

1.1.2.1. Highest level of protection possible

The optimum level of protection to be taken into account in the design is that beyond which the constraints by the wearing of the PPE would prevent its effective use during the period of exposure to the risk or normal performance of the activity.

1.1.2.2. Classes of protection appropriate to different levels of risk

Where differing foreseeable conditions of use are such that several levels of the same risk can be distinguished, appropriate classes of protection must be taken into account in the design of the PPE.

1.2. Innocuousness of PPE

1.2.1. Absence of risks and other inherent nuisance factors

PPE must be so designed and manufactured as to preclude risks and other nuisance factors under foreseeable conditions of use.

1.2.1.1. Suitable constituent materials

The materials of which the PPE is made, including any of their possible decomposition products, must not adversely affect the health or safety of users.

1.2.1.2. Satisfactory surface condition of all PPE parts in contact with the user

Any part of the PPE that is in contact or is liable to come into contact with the user when the PPE is worn must be free of rough surfaces, sharp edges, sharp points and the like which could cause excessive irritation or injuries

1.2.1.3. Maximum permissible user impediment

Any impediment caused by PPE to movements to be made, postures to be adopted and sensory perception must be minimized; nor must PPE cause movements which endanger the user or other persons.

1.3. Comfort and effectiveness

1.3.1. Adaptation of PPE to user morphology

PPE must be designed and manufactured in such a way as to facilitate its correct positioning on the user and to remain in place for the foreseeable period of use, bearing in mind ambient factors, the actions to be carried out and the postures to be adopted. For this purpose, it must be possible to adapt the PPE to fit the morphology of the user by all appropriate means, such as adequate adjustment and attachment systems or the provision of an adequate range of sizes.

1.3.2. Lightness and design strength

PPE must be as light as possible without prejudicing design strength and efficiency.

Apart from the specific additional requirements which they must satisfy in order to provide adequate protection against the risks in question (see 3), PPE must be capable of withstanding the effects of ambient phenomena inherent under the foreseeable conditions of use

1.4. Information supplied by the manufacturer

The notes that must be drawn up by the former and supplied when PPE is placed on the market must contain all relevant information on:

- In addition to the name and address of the manufacturer and/or his authorized representative established in the Community
- Storage, use, cleaning, maintenance, servicing and disinfection, cleaning, maintenance or disinfectant protection recommended by manufacturers must have no adverse effect on PPE or users when applied in accordance with the relevant instructions;
- Performance as recorded during technical tests to check the levels or classes of protection provided by the PPE in question;
- Suitable PPE accessories and the characteristics of appropriate spare parts;
- The classes of protection appropriate to different levels of risk and the corresponding limits of use;
- The obsolescence deadline or period of obsolescence of PPE or certain of its components;
- The type of packaging suitable for transport;
- The significance of any markings (see 2.12)
- Where appropriate the references of the Directives applied in accordance with Article 5(6) (b);
- The name, address and identification number of the notified body involved in the design stage of the PPE

These notes, which must be precise and comprehensible, must be provided at least in the official language(s) of the member state of destination

2. ADDITIONAL REQUIREMENTS COMMON TO SEVERAL CLASSES OR TYPES OF PPE

2.1. PPE incorporating adjustment systems

If PPE incorporates adjustment systems, the latter must be designed and manufactured so that, after adjustment, they do not become undone unintentionally in the foreseeable conditions of use.

2.3. PPE for the face, eyes and respiratory system

Any restriction of the user's face, eyes, field of vision or respiratory system by the PPE shall be minimised.

The screens for those types of PPE must have a degree of optical neutrality that is compatible with the degree of precision and the duration of the activities of the user.

If necessary, such PPE must be treated or provided with means to prevent misting-up.

Models of PPE intended for users requiring sight correction must be compatible with the wearing of spectacles or contact lenses.

2.4. PPE subject to ageing

If it is known that the design performance of new PPE may be significantly affected by ageing, the month and year of manufacture and/or, if possible, the month and year of obsolescence must be indelibly and unambiguously marked on each item of PPE placed on the market and on its packaging.

If the manufacturer is unable to give an undertaking with regard to the useful life of the PPE, his instructions must provide all the information necessary to enable the purchaser or user to establish a reasonable obsolescence month and year, taking into account the quality level of the model and the effective conditions of storage, use, cleaning, servicing and maintenance.

Where appreciable and rapid deterioration in PPE performance is likely to be caused by ageing resulting from the periodic use of a cleaning process recommended by the manufacturer, the latter must, if possible, affix a marking to each item of PPE placed on the market indicating the maximum number of cleaning operations that may be carried out before the equipment needs to be inspected or discarded. Where such a marking is not affixed, the manufacturer must give that information in his instructions.

2.6. PPE for use in potentially explosive atmospheres

PPE intended for use in potentially explosive atmospheres must be designed and manufactured in such a way that it cannot be the source of an electric, electrostatic or impact-induced arc or spark likely to cause an explosive mixture to ignite.

2.8. PPE for intervention in very dangerous situations

The instructions supplied by the manufacturer with PPE for intervention in very dangerous situations must include, in particular, data intended for competent, trained persons who are qualified to interpret them and ensure their application by the user.

The instructions must also describe the procedure to be adopted in order to verify that PPE is correctly adjusted and functional when worn by the user.

Where PPE incorporates an alarm which is activated in the absence of the level of protection normally provided, the alarm must be designed and placed so that it can be perceived by the user in the foreseeable conditions of use.

2.9. PPE incorporating components which can be adjusted or removed by the user

Where PPE incorporates components which can be attached, adjusted or removed by the user for replacement purposes, such components must be designed and manufactured so that they can be easily attached, adjusted and removed without tools.

2.12. PPE bearing one or more identification or recognition marks directly or indirectly relating to health and safety

The identification or recognition marks directly or indirectly relating to health and safety affixed to these types or classes of must preferably take the form of harmonized pictograms or ideograms and must remain perfectly legible throughout the foreseeable useful life of the PPE. In addition, these marks must be complete, precise and comprehensible so as to prevent any misinterpretation; in particular, where such marks incorporate words or sentences, the latter must appear in the official language(s) of the Member State where the equipment is to be used.

If PPE (or a PPE component) is too small to allow all or part of the necessary marking to be affixed, the relevant information must be mentioned on the packing and in the manufacturer's notes.

3. ADDITIONAL REQUIREMENTS SPECIFIC TO PARTICULAR RISKS

3.10.1. Respiratory protection

PPE intended for the protection of the respiratory system must make it possible to supply the user with breathable air when exposed to a polluted atmosphere and/or an atmosphere having an inadequate oxygen concentration.

The breathable air supplied to the user by PPE must be obtained by appropriate means, for example after filtration of the polluted air through PPE or by supply from an external unpolluted source.

The constituent materials and other components of those types of PPE must be chosen or designed and incorporated so as to ensure appropriate user respiration and respiratory hygiene for the period of wear concerned under the foreseeable conditions of use.

The leak-tightness of the facepiece and the pressure drop on inspiration and, in the case of the filtering devices, purification capacity must keep contaminant penetration from a polluted atmosphere low enough not to be prejudicial to the health or hygiene of the user.

The PPE must bear details of the specific characteristics of the equipment which, in conjunction with the instructions, enable a trained and qualified user to employ the PPE correctly.

In the case of filtering equipment, the manufacturer's instructions must also indicate the time limit for the storage of new filters kept in their original packaging.



Technical Assessment of EN 149: 2001 + A1: 2009 Standard and other Standards it refers to, Clauses Corresponding to the (EU) 2016/425 Directive

Conforming to EN 149:2001 + A1:2009 Standard Requirements

Article 5	Classification: Particle Filtering Half Mask The mask subject to evaluation based on the test results and technical file provided by the manufacturer is classified as: Filtering Efficiency and maximum Total Inward Leakage: Classified as FFP2 Mask is classified for single shift use, NR																																						
Article 7.4	Packing: Particle filtering half masks are packaged to protect them from contamination before use and with cardboard boxes to prevent mechanical damage. The packaging design and the product is considered to withstand the foreseeable conditions of use based on the visual inspection results given in the test report.																																						
Article 7.5	Material: Materials used in particle filtering half masks, according to the simulated wearing treatment and temperature conditioning results; It is understood it withstands handling and wear over the period for which the particle filtering half mask is designed to be used, it suffered mechanical failure of the facepiece or straps, any material from the filter media released by the air flow through the filter has not constitute a hazard or nuisance for the wearer. The manufacturer declares that the materials used in manufacturing of the mask does not have an adverse affect to the health and safety of users. Based on the test results, the masks did not collapse when subject to simulated wearing and temarature conditioning. No nuisance situation is reported during the practical performance tests by human subjects.																																						
Article 7.6	Cleaning and Disinfection: Particle filtering half mask is not designed to be as re-usable. No cleaning or disinfection procedure provided by the manufacturer.																																						
Article 7.7	Practical Performance : The test report indicates that the human subjects did not face any difficulty in performing the excercises while they were wearers by the sample masks, in walking test or work simulation tests. The wearers did not report any failure by means of head harness / straps/ earloops comfort, security of fastenings and field of vision. Also no imperfections reported during total inward tests about the comfort, field of vision and fastening issues. <table><tr><th>Assessed Elements</th><th>Positive</th><th>Negative</th><th>Requirements in accordance with EN 149:2001 + A1:2009 and Result</th></tr><tr><td>2.Head harness comfort</td><td>2</td><td>0</td><td rowspan="3">Positive results are obtained from the test subjects No imperfections</td></tr><tr><td>3.Security of fastenings</td><td>2</td><td>0</td></tr><tr><td>5.Field of vision</td><td>2</td><td>0</td></tr></table> Conditioning : (A.R.) As Received, original	Assessed Elements	Positive	Negative	Requirements in accordance with EN 149:2001 + A1:2009 and Result	2.Head harness comfort	2	0	Positive results are obtained from the test subjects No imperfections	3.Security of fastenings	2	0	5.Field of vision	2	0																								
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3.Security of fastenings	2	0																																					
5.Field of vision	2	0																																					
Article 7.8	Finish of Parts: Particle filtering half masks, which are likely to come into contact with the user, do not have sharp edges and do not contain burrs.																																						
Article 7.9.1	Total Inward Leakage: The Total Inward Leakage test is conducted by 10 individual in an aerosol chamber with a walking band, and samples are taken during the conduction of the excercises defined in the standard. The samples used in the test are subjected to the conditioning required in the standard as Temperature conditioning and as received. The face dimensions of the subjects are also reported. The measurement details for each subject and for each excercise are available in the test report. It was reported that: All 50 excercise measurement results are smaller or equal to 11%, the values varies between 6,7 % and 8,9 % 9 out of 10 individual's arithmetic mean is smaller or equal to 8%, the values varies between 7,1 % and 8,3 % According to the reported results, the product meets the limits for FFP1 and FFP2 classification.																																						
Article 7.9.2	Penetration of filter material: Sodium Chloride Testing <table><tr><th>Condition</th><th>No. of Sample</th><th>Sodium Chloride Testing 95 L/min max (%)</th><th>Requirements in accordance with EN 149:2001 + A1:2009</th><th>Result</th></tr><tr><td>(A.R.)</td><td>-</td><td>1,4</td><td rowspan="3">FFP1 ≤ 20 %</td><td rowspan="9">Filtering half masks fulfill the requirements of the standard EN EN 149:2001 + A1:2009 given in 7.9.2 in range of the FFP1, FFP2 classes.</td></tr><tr><td>(A.R.)</td><td>-</td><td>1,3</td></tr><tr><td>(A.R.)</td><td>-</td><td>1,2</td></tr><tr><td>(S.W.)</td><td>-</td><td>1,3</td><td colspan="2" rowspan="3">FFP2 ≤ 6 %</td></tr><tr><td>(S.W.)</td><td>-</td><td>1,3</td></tr><tr><td>(S.W.)</td><td>-</td><td>1,4</td></tr><tr><td>(M.S. T.C.)</td><td>-</td><td>1,6</td><td colspan="2" rowspan="3">FFP3 ≤ 1 %</td></tr><tr><td>(M.S. T.C.)</td><td>-</td><td>1,6</td></tr><tr><td>(M.S. T.C.)</td><td>-</td><td>1,5</td></tr></table> Conditioning : (M.S.) Mechanical Strength (T.C.) Temperature Conditioning (A.R.) As Received, original (S.W.) Simulated wearing treatment 95 L/min = 1,6 dm³.sn⁻¹	Condition	No. of Sample	Sodium Chloride Testing 95 L/min max (%)	Requirements in accordance with EN 149:2001 + A1:2009	Result	(A.R.)	-	1,4	FFP1 ≤ 20 %	Filtering half masks fulfill the requirements of the standard EN EN 149:2001 + A1:2009 given in 7.9.2 in range of the FFP1, FFP2 classes.	(A.R.)	-	1,3	(A.R.)	-	1,2	(S.W.)	-	1,3	FFP2 ≤ 6 %		(S.W.)	-	1,3	(S.W.)	-	1,4	(M.S. T.C.)	-	1,6	FFP3 ≤ 1 %		(M.S. T.C.)	-	1,6	(M.S. T.C.)	-	1,5
Condition	No. of Sample	Sodium Chloride Testing 95 L/min max (%)	Requirements in accordance with EN 149:2001 + A1:2009	Result																																			
(A.R.)	-	1,4	FFP1 ≤ 20 %	Filtering half masks fulfill the requirements of the standard EN EN 149:2001 + A1:2009 given in 7.9.2 in range of the FFP1, FFP2 classes.																																			
(A.R.)	-	1,3																																					
(A.R.)	-	1,2																																					
(S.W.)	-	1,3	FFP2 ≤ 6 %																																				
(S.W.)	-	1,3																																					
(S.W.)	-	1,4																																					
(M.S. T.C.)	-	1,6	FFP3 ≤ 1 %																																				
(M.S. T.C.)	-	1,6																																					
(M.S. T.C.)	-	1,5																																					



Article 7.17	Clogging: This test is not applied to Particle Filtering Half Mask which is not reusable. (For single shift use devices, the clogging test is optional test. For re-usable devices test is mandatory.)
Article 7.18	Demountable Parts: There are no demountable parts on the product.
Article 8	Testing: All tests conducted according to Clause 8 of this standard is available in the test report and are evaluated in this report for qualification and classification of the mask.
Article 9	Marking – Packaging: Necessary markings are available on the product package (box). The manufacturer and its trademark is clearly visible. The type of the mask and the classification including the status of re-usability, the reference to EN 149:2001+A1:2009 standard, the end date of shelf life, using and storage instructions and pictograms and CE mark are available on the product package. The above evaluation is based on the technical document for packaging and marking, for box design. Verified on the Annex 9.1 of the technical file. The technical documentation for mask design (drawing) also evaluated for marking requirements, drawing YY0525. Even the mask template (drawing) not indicates the necessary markings, the image of the mask in the technical file carries information about the manufacturer / trademark (CRDLIGHT) of the manufacturer. Type of mask, the reference to EN 149+A1:2009 standard and classification including the re-usability of the mask. The manufacturer also printed CE mark with our Notified Body number. The mask do not have sub-assemblies. Even the tested sample by the laboratory do not carry necessary marking information as stated in the technical documentation, the manufacturer shall follow marking instructions for serial production. Model YY0525 drawing exists in the technical file of the manufacturer. Annex 6 of technical file.
Article 10	Information to be supplied by the manufacturer: In each of the smallest commercially available packaging of the product, implementation (installation instructions) pre-use controls, warning and usage limitations, storage and meanings of symbols / pictograms are defined. User instruction document in the technical file found to be appropriate, Annex 8. The manufacturer shall include this documented user information text in every smallest commercially available package.

PREPARED BY

Osman CAMCI
PPE Expert



APPROVED BY

Suat KAÇMAZ
Director




CERTIFICATE OF CONFORMANCE

Certificate No: 2163-PPE-834/01

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

JIANGMEN YANYANG TRADING CO., LTD.

NO.1, 4th Floor, Building 2, NO.18 Xinyi Road, Jianghai District,
Jiangmen City, Guangdong Province, 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
CRDLIGHT / YY0525	FFP2 NR	2163-PPE-834	25.06.2020	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 **19/08/2020** and will be valid for one year, until **18/08/2021** 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

Verify the validity with the QR code



EU Declaration of Conformity

Annex V PPE Regulation (EU) 2016/425

1. This EU Declaration of conformity refers to the following products:

Product Name	Model	Classification/Type	Batch No./Serial No./Identifier
Filtering Half Mask	YY0525	FFP2 NR	2163-PPE-834

2. Manufacturer's name and address is as below:

Name	Jiangmen Yanyang Trading Co., Ltd.
Address	No. 1, 4th Floor, Building 2, No. 18 Xinyi Road, Jianghai District, Jiangmen City, Guangdong Province, China

**3. The conformity assessment according to the essential requirements has been done by
Easepal according to**

EU Community
Legislation

Personal Protective Equipment (PPE) Regulation (EU) 2016/425
Annex VII in connection with
Annex V (Module B EU Type Examination Certification)
Certificate No.: 2163-PPE-834 (issued date: 25/06/2020)
Annex VIII (Module C2 Certificate of Conformance)
Certificate No.: 2163-PPE-834/01 (issued date: 19/08/2020)

Harmonised
standard

Personal Protective Equipment
EN 149:2001+A1:2009
Trust Right Testing and Certification Service (Zhongshan) Ltd.
Test Serial ID: R20200062 (Dated 15.06.2020)

Other specifications: **Particle filtering mask, Category III, FFP2 NR**

In cooperation with:

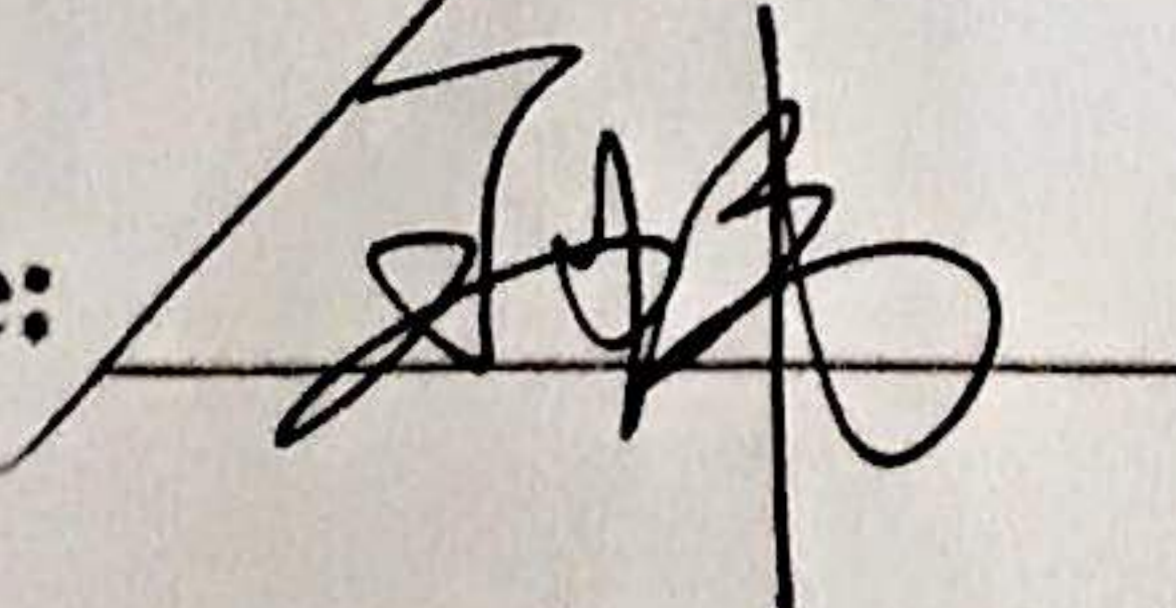
Notified Body:

UNIVERSAL SERTİFİKASYON VE GÖZETİM HİZM. TİC. LTD. ŞTİ.
(CE 2163)
Necip Fazil Bulvari Keyap Sitesi
E2 Blok No: 44/84
Yukari Dudullu Ümraniye
Istanbul, Turkey

This declaration is valid until: 14th Jan. 2022

Place and date of issue: Jiangmen, 15th Jan. 2021

Signature:



Date: 15th Jan. 2021

Company Sign:





统一社会信用代码
91440704MA52768M32

营业执照

(副本) (副本号:1-1)



扫描二维码登录“
国家企业信用信息公示系统”了解更
多登记、备案、许
可、监管信息。

名称 江门岩羊工贸有限公司
类型 有限责任公司(自然人投资或控股)

法定代表人 胡杰

经营范围 研发、生产、加工、销售: LED、驱动电源、开关电源、电子整流器、智能化设备系列产品及配件、电器产品附件、低压电器产品、照明电器附件、电子产品、电气元器件、家用电器、五金、塑料制品、医疗器械、消毒产品(不含危险化学品); 医疗器械技术研发; 生产、销售: 机械设备、橡胶制品、日用品、口罩; 货物或技术进出口(国家禁止或涉及行政审批的货物和技术进出口除外)。(依法须经批准的项目, 经相关部门批准后方可开展经营活动。) 〰

注册资本 人民币伍拾万元

成立日期 2018年08月30日

营业期限 长期

住

所 江门市江海区信义路18号2幢4层自编1号
(信息申报制、一址多照) (经营场所
: 江门市江海区信义路18号5幢5层自编A
04(信息申报制)) (一照多址)

登记机关



2020 年 5 月 12 日

CRDLIGHT

Certificate of conformity

product Name:Filtering half mask(non-medical)

Model:YY0525

Size:155*105mm

Quantity:25pcs/box

Color:White

Main Material:50% non-woven, 33.33% melt-blown,
16.67% hot air cotton

exquive Standard:EN149:2001+A1:2009FFP2NH

Pronction Date: Feb. 2021

Lot Number: 0212

Expiry Date: 12.Feb.2024

Manufacturer:jiangmen Yanyang Trading Co., Ltd.

Address: No. 18 Kt Road Jiangtai District,
jiangmen, Guangdong, Carra.

MADE IN CHINA