



C/ Sofía, 3-5 Pol. Ind. Cabezo Beaza C.P. 30353 CARTAGENA (Spain) CIF. B30668420

DOCUMENTATION

REF. CV-41 / YYC1028

MASK ULTRA PROTECTION FFP2

MASCARILLA ULTRA PROTECCIÓN FFP2 MASQUE FFP2 ULTRA PROTECTION FFP2 MASCHERINA PROTEZIONE ULTRA FFP2



MASTER BOX: 1000 pcs



ITEM: YYC1028 DESCRIPTION: NAAMIO

MATERIAL:

5 PLY (40% non woven, 38% Meltblown, 22% algodón).

QUANTITY: 1.000

G.W N.W

CNT SIZE

BATCH NUMBER: PRODUCTION DATE:

VALIDITY:

MADE IN P.R.C.









BAG: 1 pc





NOTA IMPORTANTE:

En todos los procesos de fabricación de nuestras mascarillas, no se utiliza grafeno o derivados del mismo.

Colaboramos con

































Manufacturer: JIANGMEN YANYANG TRADING CO.,LTD

NO.1,4TH FLOOR,BUILDING 2,NO.18 XINYI ROAD,JIANGHAI DISTRICT,JIANGMEN CITY,GUANGDONG PROVINCE, CHINA

EU Declaration of Conformity

We, the manufacturer, herewith declare that the products

Filtering half mask

Brand: CRDLIGHT Item No.: YYC1028

Meet the essential requirements and relevant provisions of EU Directive:

Personal Protective Equipment Regulation(EU) 2016/425 Annex 5

This Declaration of conformity is valid in connection with the following test report:

Certificate No:2163-PPE-1732

Report Date: November 23, 2020

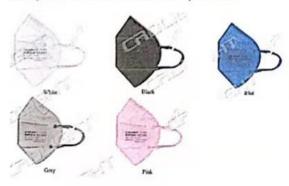
Test Reported by: Jiangsu Guojian Testing Technology Co., Ltd

Application Standard: EN149:2001+A1:2009 FFP2 NR

Classifications: FFP2 NR

The above mentioned declaration of conformity is exclusively under the responsibility of

JIANGMEN YANYANG TRADING CO.,LTD NO.1,4TH FLOOR,BUILDING 2,NO.18 XINYI ROAD,JIANGHAI DISTRICT,JIANGMEN CITY,GUANGDONG PROVINCE, CHINA



Jiangmen China 2020-December-04

Place,

Date



EU Declaration of Conformity Document No: Page 1 of 1



NB 2163

EU TYPE EXAMINATION CERTIFICATE

Certificate No: 2163-PPE-1732

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

Single shift use particle filtering half mask for protection against solid and liquid aerosols, is a folding type, 5 layers cotton and polypropylene fabrics, without valve, fitted with ear loops, with internal nose clip and inside sponge strip.

Brand Name: CRDLIGHT Model: YYC1028 Classification: FFP2 NR Model have white, black, grey, pink, and blue versions

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 04/12/2020 and will be valid for 5 years, if there is no change in the relevant harmonised standard affecting the essential health and safety requirements.



Director



TECHNICAL ASSESSMENT REPORT

REPORT DATE / NO: 04.12.2020 / 2163-KKD-1732

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 Jiangsu Guojian Testing Technology Co. Ltd. accredited by CNAS (Chinese Accreditation Body), signatory to ILAC MRA, with number CNAS L10118 for the product identified below, dated 23.11.2020 with Serial Id 2020-WSZ FHL No.8631 based on EN 149: 2001 + A1: 2009 standard and test reports on the material safety by means of toxic, carcinogen, irritationg and sensitivity evaluation.

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 Personel 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: Single shift use particle filtering half mask for protection against solid and liquid aerosols, is a folding type, 5 layers cotton and polypropylene fabrics, without valve, fitted with ear loops, with inside nose bridge.

Component and Materials:

Component	Material	Grade / Size	
1st layer (Outer)	Non-Wowen Fabric	50 g/m ² (±5 g/m ²)	
2nd layer	Melt-blown - non-wowen fabric	25 g/m² (±5 g/m²)	
3rd layer	Melt-blown - non-wowen fabric	25 g/m² (±5 g/m²)	
4th layer	Hot Air cotton	25 g/m ² (±5 g/m ²)	
5th layer (Inner)	Non-Wowen Fabric	25 g/m ² (±5 g/m ²)	
Internal Nose Clip	Polypropylene	52 mm (±5 mm)	
Ear Loop	Polyester + Cotton	23 cm (±1 cm)	
Sponge Strip	Polyurethane	120 mm (±1 mm)	
Plastic Clip	Polypropylene	50 mm (±1 cm)	

Classification: FFP2 NR

Brand Name: CRDLIGHT Model: YYC1028

Colored samples of the mask





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 prossible 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 fore seeable 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 permessible user impediment

Any inpediment 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:

- a) In addition to the name and addressof the manufacturer and/or his authorized representative established in the Community
- b) 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;
- c) Performance as recorded during technical tests to check the levels or classes of protection provided by the PPE in guestion;
- Suitable PPE accessories and the characteristics of appropriate spare parts;
- e) The classes of protection appropriate to different levels of risk and the corresponding limits of use;
- The obsolescence deadlineor period of obsolescence of PPEor certain of its components; f)
- g) The type of packaging suitable for transport;
- h) The significance of any markings(see 2.12)
- Where appropriate the references of the Directives applied inaccordance with Article5(6) (b); i)
- j) 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



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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 rem ain perfectly legible throughout the foreseeableuseful 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

3.10.2. Protection against cutaneous and ocular contact

PPE intended to prevent the surface contact of all or part of the body with substances and mixtures which are hazardous to health or with harmful biological agents must be capable of preventing the penetration or permeation of such substances and mixtures and agents through the protective integument under the foreseeable conditions of use for which the PPE is intended.

To this end, the constituent materials and other components of those types of PPE must be chosen or designed and incorporated so as to ensure, as far as possible, complete leak-tightness, which will allow where necessary prolonged daily use or, failing this, limited leak-tightness necessitating a restriction of the period of wear.

Where, by virtue of their nature and the foreseeable conditions of their use, certain substances and mixtures which are hazardous to health or harmful biological agents possess high penetrative power which limits the duration of the protection provided by the PPE in question, the latter must be subjected to standard tests with a view to their classification on the basis of their performance. PPE which is considered to be in conformity with the test specifications must bear a marking indicating, in particular, the names or, in the absence of the names, the codes of the substances used in the tests and the corresponding standard period of protection. The manufacturer's instructions must also contain, in particular, an explanation of the codes (if necessary), a detailed description of the standard tests and all appropriate information for the determination of the maximum permissible period of wear under the different foreseeable conditions of use.

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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 1		and the state of t					
Anti-la	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;							
5		Filtering Efficiency and Maximum Total Inward Leakage: Classified as FFP2						
	Mask is classified for single shift use, NR							
Article	Packing: Particle filtering half masks are packaged to protect them from contamination before use and with cardboard boxes to protect them from contamination before use and with cardboard boxes to protect them from contamination before use and with cardboard boxes to protect them from contamination before use and with cardboard boxes to protect them from contamination before use and with cardboard boxes to protect them from contamination before use and with cardboard boxes to protect them from contamination before use and with cardboard boxes to protect them from contamination before use and with cardboard boxes to protect them from contamination before use and with cardboard boxes to protect them from contamination before use and with cardboard boxes to protect them from contamination before use and with cardboard boxes to protect them from contamination before use and with cardboard boxes are protected by the protect them from contamination before use and with cardboard boxes are protected by the protect them from contamination before use and with cardboard boxes are protected by the protect them from contamination before use and with cardboard boxes are protected by the protect them from contamination before use and the protected by the prot							
7.4		mechanical damage. The packaging design and the product is considered to withstand the foreseeable conditions of use based on the						
	inspection results given in the test report. Details given in Annex 5 of Technical File Material: Materials used in particle filtering half masks, according to the simulated wearing treatment and temperature conditioning res							
				earing treatment and temperature conditioning range half mask is designed to be used, it suffered in				
				air flow through the filter has not constitute a				
				cturing of the mask does not have an adverse a				
				adverse effect for the wearers health in Section				
	Technical File.	colares that the mater	iai do not nave any	adverse effect for the weaters health in Section				
Article		t collapse when subje	ect to simulated wear	ing and temarature conditioning. No nuisance				
7.5	reported during the practical performance te			ing and temarature conditioning. No nuisance s				
				e most outer layer of the mask, with the earloo				
				DC20110368 (Blue), DC20110370 (Grey), DC				
	The second secon			Q2588-6EN (white) prepared by TST Testing T				
				spunbound fabric) used in the most outer layer of				
	is considered to be safe for use on the mask.							
Article				ble. No cleaning or disinfection procedure provi				
7.6	manufacturer.	6 mail mask is not do.	igned to be as re-asa	ole. No cleaning of disinfection procedure provi				
	Practical Performance :							
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	security of fastenings and field of vision. Al issues.	so no imperfactions re	eported during total in	ure by means of head harness / straps/ earloop award tests about the comfort, field of vision and				
	security of fastenings and field of vision. Al	so no imperfactions re		ure by means of head harness / straps/ earloop				
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	Condition	No. o Samp	-	Sodium Chloride Testi 95 L/min max (%)		equirements in accordance EN 149:2001 + A1:200		Result	
	(A.R.)	- Samp	10	0,1		211712001 1711.200			
Article 7.9.2	(A.R.)			0,1					
	(A.R.)			0,1		FFD1 < 20.0/	Filtering half masks fulfill trequirements of the standar		
	(S.W.)			0,2		FFP1 \leq 20 %			
	(S.W.)			0,1		FFP2 ≤ 6 %		EN EN 149:2001 + A1:200 given in 7.9.2 in range of the first, and second protection	
	(S.W.)	the state of the s		0,2					
	(M.S. T.C.)	(M.S. T.C.) -		0,6 0,5		FFP3 ≤ 1 %		classes.	
	(M.S. T.C.)							FFP1, FFP2, FFP3	
	(M.S. T.C.)	-		0,6				,,	
	Conditioning: (M	Conditioning: (M.S.) Mechanical Strength				95 1	$L/min = 1.6 \text{ dm}^3.\text{sn}^{-1}$		
	(A	.R.) As Receiv	re Conditioning red, original wearing treatm						
	Penetration of filte						***************************************		
	Con	dition	No. of	Paraffin Oil Test 95 L/min max (quirements in accordance EN 149:2001 + A1:2009		Result	
		L D \	Sample		(70) With	EN 149.2001 + A1.2009			
		1.R.)	-	0,2					
		A.R.)	-	0,3				half masks fulfill the	
		1.R.)		0,2		FFP1 ≤ 20 %	requirem	nents of the standard	
Article		s.W.)	-	0,3				49:2001 + A1:2009	
7.9.2				0,2		FFP2 ≤ 6 %		7.9.2 in range of the	
		S.W.) S. T.C.)	-	0,3		EED2 < 1.04	first and	d second protection	
	***************************************	5. T.C.)	-	0,7		FFP3 ≤ 1 %	DED	classes.	
		S. T.C.)		0,6 0.8			FFF	21, FFP2, FFP3	
	Conditioning: (M.			0,8					
			e Conditioning						
		R.) As Receive							
		Care Commission Commission	wearing treatme						
Autiala									
Article	Compatibility with	elcine In Proof							
		SKIII. III FIACI	ical Performance	ce report, the likelihoo	od of mask ma	terials in contact with the s	kin causi	ng irritation or other	
7.10	adverse effect on he	alth was not re	ical Performant ported.	ce report, the likelihoo	od of mask ma	terials in contact with the s	kin causi	ng irritation or other	
7.10	adverse effect on he	ealth was not re	ported.	ce report, the likelihoo	od of mask ma	terials in contact with the s	skin causi	ng irritation or other	
7.10	adverse effect on he	alth was not re	ported.	ce report, the likelihoo				ng irritation or other	
7.10	adverse effect on he	No. of	ported.	sual inspection	Requiren	nents in accordance with E		ng irritation or other	
7.10	Flammability: Condition	No. of Sample	ported.	sual inspection	Requiren	nents in accordance with E 149:2001 + A1:2009		Result	
4rticle	adverse effect on he Flammability: Condition (A.R.)	No. of Sample	ported.	sual inspection	Requiren	nents in accordance with E 49:2001 + A1:2009 Filtering half mask			
	adverse effect on he Flammability: Condition (A.R.) (A.R.)	No. of Sample	ported.	sual inspection 0,4s 0,4s	Requiren	nents in accordance with E 149:2001 + A1:2009 Filtering half mask shall not burn or not	N	Result Passed	
Article	Condition (A.R.) (A.R.) (T.C.)	No. of Sample	ported.	sual inspection 0,4s 0,4s 0,5s	Requiren	nents in accordance with E 49:2001 + A1:2009 Filtering half mask	N Filte	Result Passed ering half masks fulfill	
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Article 7.17.2	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 7.17.3	Penetration of filter material: This test is not applied to Particle Filtering Half Mask which is not reusable.
Article 7.18	Demountable Parts: There are no demountable parts on the product.
Article	Marking – Packaging: Necessary markings are available on the product package (box). The name and trademark of the manufacturer is stated to exist on the carton boxes. The type of the mask and the classification including the status of re-usability, the reference to EN 149:2001+A1:2009 standard, the year of end 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 and annex 6 of the technical file.
9	The technical documentation for mask design (drawing) also evaluated for marking requirements, drawing Annex 6. The mask template (drawing) indicates that the mask will carry information about the brandname 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 samples by the laboratory do not carry necessary marking information as stated in the technical documentation, the manufacturer shall follow marking instructions for serial production. CRD-F-001 drawing which 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 commertially available package.

PREPARED BY	APPROVED BY
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