



青岛威立防护用品有限公司

Qingdao Weili Protective Articles Co., Ltd.

口罩系列产品介绍

Introduction of mask series products



QMC 威立

始于2020

专注口罩研发行业领军者

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01



威立口罩简介

COMPANY PROFILE OF WEILI

取得国外标准认证或注册的医疗物资生产企业清单

Name List of Medical Devices and Supplies Companies with Certification/Authorization from other Countries

中国医药保健品进口商会

CHINA CHAMBER OF COMMERCE FOR IMPORT & EXPORT OF MEDICINES & HEALTH PRODUCTS



中国医药保健品进出口商会

服务产业链 | 助力国际化

首页

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商会会刊

取得国外标准认证或注册的医疗物资生产企业清单

动态更新：2020年8月4日下载

2058	青岛威立防护用品有限公司 Qingdao Weili Protective Articles Co., Ltd.	91370281MA3RJRH101	欧盟CE
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青岛威立防护用品有限公司简介

青岛威立防护用品有限公司，专注于医疗防护用品的研发与制造，秉承“用品质捍卫生命”的企业发展原则，服务于全球消费者。“品质及格是尽社会的责任，品质卓越是对社会的奉献。”质量第一，威立专注于把检产品质量，追求精益求精的制作工艺。

产品种类 KN95/KP95/KP100/N95/N99/FFP2/FFP3, 一次性医用非灭菌等各类防护口罩。共有生产流水线十多条。可日产平面一次性口罩150万，折叠口罩50万个左右。现有无菌医用车间和无尘防护用品车间工厂。SGS ISO CE 各类检验检测齐全。威立会继续挑战更高端产品技术。

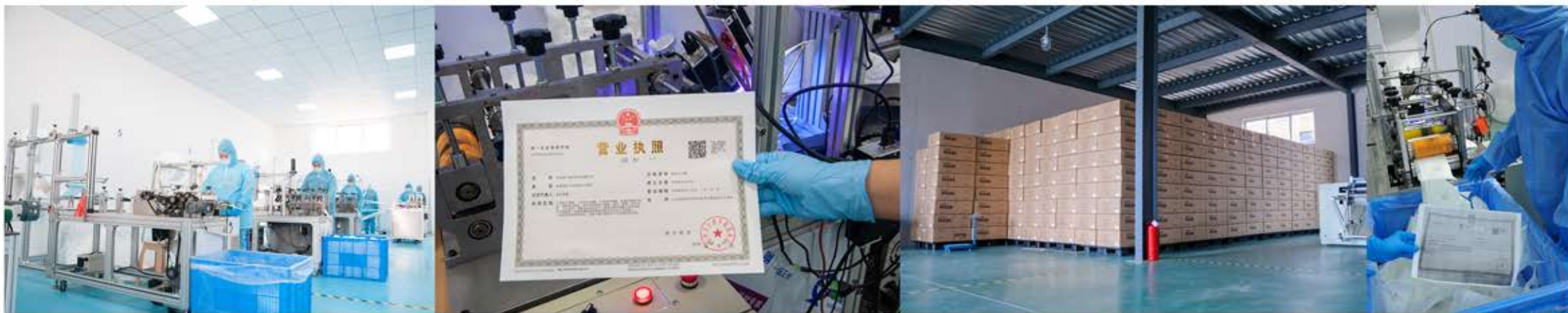
Brief introduction of Qingdao Weili Protective Articles Co., Ltd.

Qingdao Weili Protective Articles Co., Ltd. focuses on the R&D and manufacture of medical protective articles. We serve the customers worldwide by inheriting the principle of company development "Defending lives by quality."

"Ordinary quality is the responsibility to the society, extraordinary quality is the devotion to the society." Quality comes first. Qingdao Weili focuses on the quality inspection and pursues excelsior manufacture craft.

The products range include KN95/KP95/KP100/N95/N99/FFP2/FFP3 and disposable medical masks (non-sterile) etc. There are over ten production lines with daily output of 1.5million disposable masks and 0.5million folding masks. We are also equipped with sterile medical workshop and dustless workshop for protective articles. A complete range of inspections, such as SGS, ISO, CE, is available.

Qingdao Weili will go on with the challenge of higher product technology.



02



企业资质

ENTERPISE CERTIFIACTES

	
营 业 执 照	
(副 本) 1-1	
统一社会信用代码 91370281MA3RJRH101	 扫描二维码 “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 青岛威立防护用品有限公司	注册 资 本 壹佰万元整
类 型 有限责任公司(自然人独资)	成 立 日 期 2020年03月17日
法 定 代 表 人 姜许雯婧	营 业 期 限 2020年03月17日至 年 月 日
经 营 范 围 1类医疗器械、II类医疗器械、III类医疗器械、普通劳保防护用品、美容美发用品、洗护用品的生产销售、批发、零售；医疗用品、卫生用品、货物及技术进出口（国家法律法规禁止经营的项目不得经营，法律法规限制经营的取得许可证后方可经营）。。 (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)	住 所 山东省青岛市胶州市胶西办事处杜村工业园
登 记 机 关 	
2020 年 04 月 01 日	
国家企业信用信息公示系统网址: http://www.gsxt.gov.cn	市场主体应当于每年1月1日至6月30日通过 国家企业信用信息公示系统报送公示年度报告
国家市场监督管理总局监制	



对外贸易经营者备案登记表

统一社会信用代码: 91370281MA3RJRH101
进出口企业代码: 03572777

经营者中文名称	青岛威立防护用品有限公司		
经营者英文名称	Qingdao Weili Protective Articles Co., Ltd		
组织机构代码		经营者类型 (由备案登记机关填写)	有限责任公司
住 所	山东省青岛市胶州市胶西办事处杜村工业园		
经营场所 (中文)	山东青岛胶州市胶西办事处杜村工业园		
经营场所 (英文)	Du Cun Industrial Park, Jiaozhou Office, Jiaozhou City, Qingdao City, Shandong Province		
联系电话	17864183832	联系传真	
邮政编码	266300	电子邮箱	310386525@qq.com
工商登记注册日期	2020-3-17	工商登记注册号	

依法办理工商登记的企业还须填写以下内容

企业法定代表人姓名	姜许雯婧	有效证件号	370281199604010060
注册资金	壹佰万元		(折美元)

依法办理工商登记的外国（地区）企业或个体工商户（独资经营者）还须填写以下内容

企业法定代表人/ 个体工商户负责人姓名		有效证件号	
企业资产/个人财产			(折美元)

备注

填表前请认真阅读背面的条款，并由企业法定代表人或个体工商户负责人签字、盖章。

备案登记机关
签 章
2020 年 05 月 12 日

第二类医疗器械经营备案凭证

备案编号：鲁青食药监械经营备 20202096 号

企业名称	青岛威立防护用品有限公司
法定代表人	姜许雯婧
企业负责人	杜文杰
经营方式	批零兼售
住 所	山东省青岛市胶州市胶西办事处杜村工业园
经营场所	山东省青岛市胶州市胶西办事处杜村工业园
库房地址	山东省青岛市胶州市胶西办事处杜村工业园
经营范围	目类：6840 诊断试剂（诊断试剂不用低温冷藏运输贮存） 目类：06 医用成像器械，08 呼吸、麻醉和急救器械，09 物理治疗器械，11 医疗器械消毒灭菌器械，14 运输、护理和防护器械，15 患者承载器械

备案部门（公章）
备案日期：2020 年 05 月 29 日



第 28371399 号

商 标 注 册 证

OMC

核定使用商品/服务项目 (国际分类: 26)

第 26 类: 假胡子; 假发; 发髻; 发束; 假髻; 假发套; 遮秃假发; 接发片; 人类的头发 (截止)

注 册 人 青岛鼎发祥工艺品有限公司

注册人地址 山东省青岛市胶州市里岔镇工业园河北村

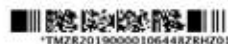
注册日期 2018年12月07日 有效期至 2028年12月06日

局 长



申长雨

发证机关



商标转让证明

兹核准第 28371399 号商标转让注册。

受让人名称 青岛华扬联合国际贸易有限公司

地址 山东省青岛市胶州市胶州经济技术开发区长江路1号



注: 转让注册商标的, 本证明与《商标注册证》一并使用。受让人自标注之日起享有商标专用权。受让人为多个的, 第一人为代表人。

青岛威立防护用品有限公司
对公付款账户

开户名称：青岛威立防护用品有限公司
地 址：山东省青岛市胶州市胶西办事处
杜村工业园
电 话：82209111
开户银行：青岛农村商业银行股份有限公司
胶州阜安支行新城分理处
银行账户：2060006724205000012616



山东农村商业银行
RURAL COMMERCIAL BANK OF SHANDONG

业务凭证

基本存款账户信息

账户名称：青岛威立防护用品有限公司

账户号码：2060006724205000012616

开户银行：青岛农村商业银行股份有限公司胶州阜安支行新城分理处

法定代表人：
(单位负责人) 姜许雯婧

基本存款账户编号：J4525006494402

2020 年 04 月 02 日

03



“威立” 口罩产品信息

‘WEILI’ MASK SERIES



FFP2 NR

MEETS EN149:2001+A1:2009 STAND

CE CERTIFICATE NO.

符合 EN149:2001+A1:2009 标准要求。



FFP2 NR

MEETS EN149:2001+A1:2009 STANDARD REQUIREMENTS.

CE CERTIFICATE NO.

符合 EN149:2001+A1:2009 标准要求。





Soft sponge nose clip

FFP2 NR

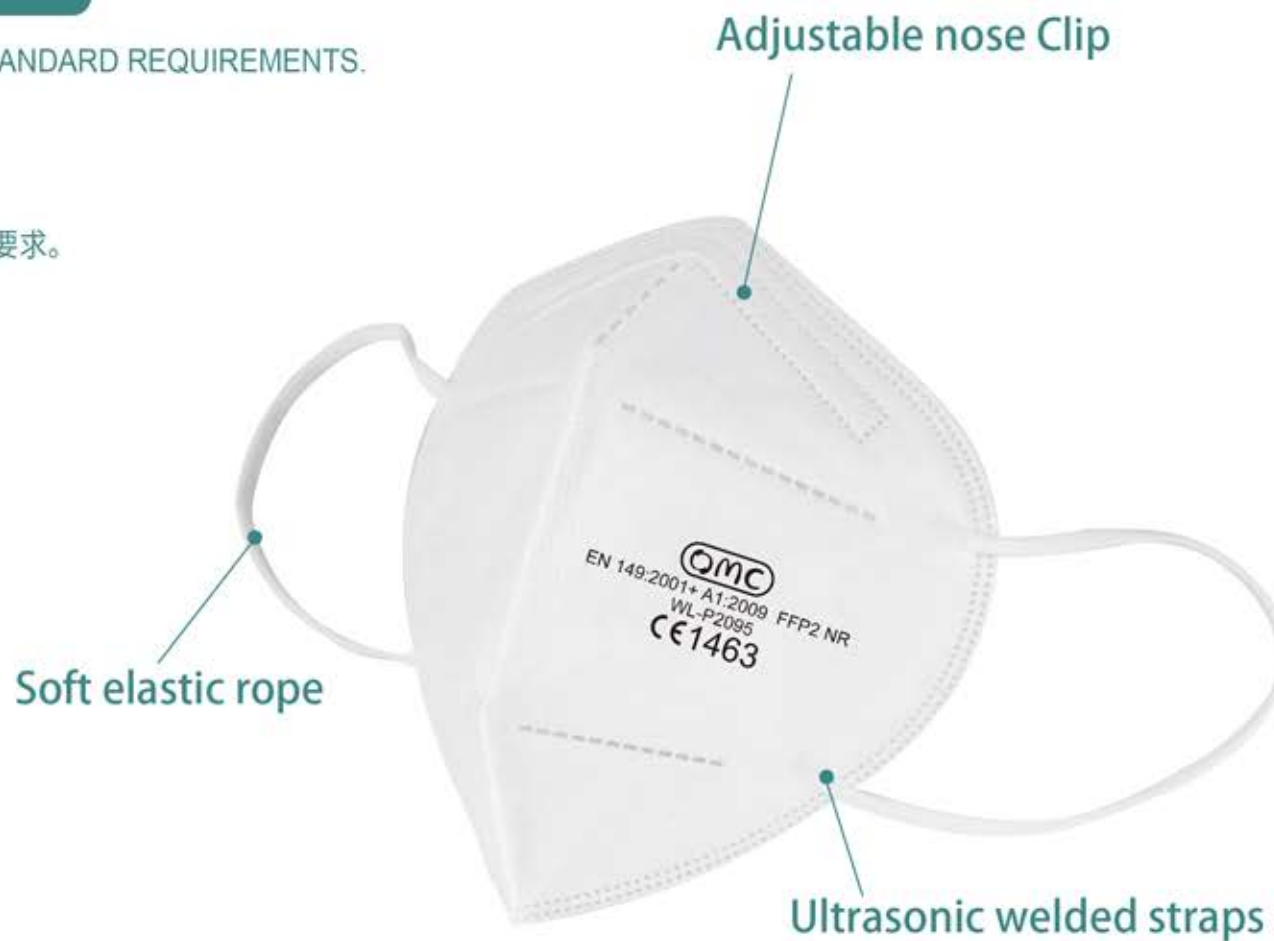
MODEL: OM-P2195

MEETS EN149:2001+A1:2009 STANDARD REQUIREMENTS.

CE CERTIFICATE NO.

型号: WL-P2095

符合 EN149:2001+A1:2009 标准要求。





FFP2 NR

MEETS EN149:2001+A1:2009 STANDARD REQUIREMENTS.

CE CERTIFICATE NO.

符合 EN149:2001+A1:2009 标准要求。



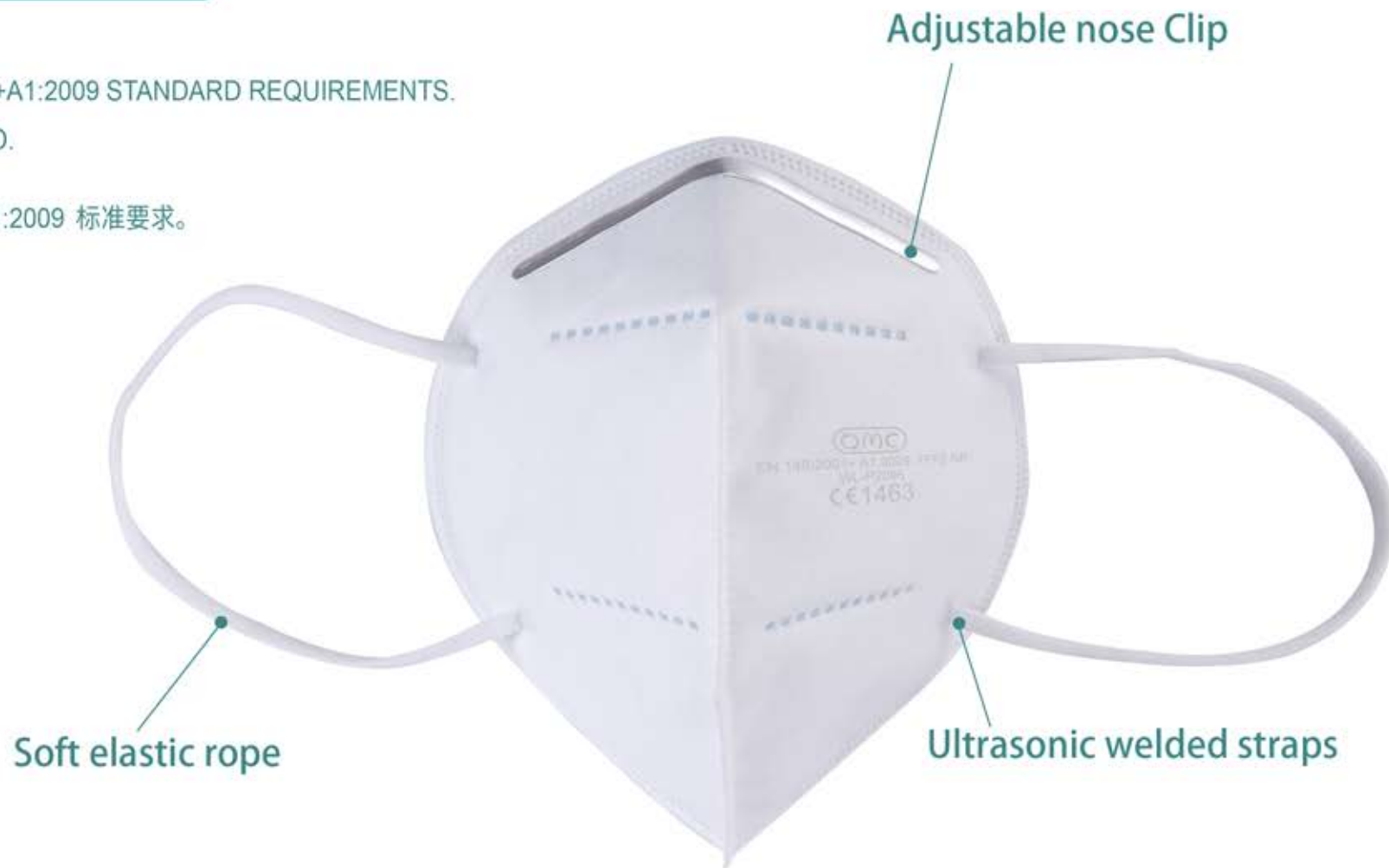


FFP2 NR

MEETS EN149:2001+A1:2009 STANDARD REQUIREMENTS.

CE CERTIFICATE NO.

符合 EN149:2001+A1:2009 标准要求。





FFP2 NR

MEETS EN149:2001+A1:2009 STANDARD REQUIREMENTS.

CE CERTIFICATE NO.

符合 EN149:2001+A1:2009 标准要求。





DISPOSABLE MEDICAL MASK TYPE IIR

MEETS EN:14683:2019+AC:2019 STANDARD REQUIREMENTS.

SGS (EU)

符合 EN:14683:2019+AC:2019 标准要求。



- THREE LAYERS OF NON-WOVEN FABRIC
- ADJUSTABLE NOSE CLIP

- ULTRASONIC WELDED STRAPS
- SOFT ELASTIC ROPE

MODEL: OM-P145-M

MEETS T/GDMDMA 0005-2020 STANDARD REQUIREMENTS.
GTTC (EU)

符合 T/GDMDMA 0005-2020 标准要求。



- THREE LAYERS OF NON-WOVEN FABRIC
- ADJUSTABLE NOSE CLIP

- ULTRASONIC WELDED STRAPS
- SOFT ELASTIC ROPE

04



特殊资质
OTHER CERTIFICATE

FDA REGISTRATION NUMBER

FDA注册证书及FDA官网型号认证

FDA registration certificate

FDA 注册证明

Fiscal Year 2020

Qingdao Weili Protective Articles Co.,Ltd.

Industrial Park,Ducun,Jiaoxi office,Jiaozhou,Qingdao, Shandong

Owner/Operator Number: 10063841

Registration Number:3016733892

Device Listing#:

Listing No	Code	Device Name	Proprietary Name
D400766	QKR	FACE MASK (EXCEPT N95 RESPIRATOR) FOR GENERAL PUBLIC/HEALTHCARE PERSONNEL PER IIE GUIDANCE	Disposable Medical Mask(Non-Sterile), 17.5*9.5cm
D400767	ORW	N95 respirator with antimicrobial/antiviral agent for use by the general public in public health medical emergencies	N95 Medical Mask(Non-Sterile),N95, KN95 Medical Mask(Non-Sterile),KN95, N95 Protective Mask,N95
D377227	IWP	Radiographic protective Glove	Medical Gloves (Non Sterile) S/M/L/XL
D382040	HQY	Sunglasses (non-prescription including photosensitive)	Medical Goggles
D382042	OEA	Non-Surgical Isolation Gown	Disposable Protective Clothing S/M/L/XL/XXL/XXXL

CCTC will confirm that such registration remains effective upon request and presentation of this certificate until the end of the calendar year stated above, unless said registration is terminated after issuance of this certificate. CCTC makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. This certificate does not denote endorsement or approval of the certificate-holder's device or establishment by the U.S. Food and Drug Administration. CCTC assumes no liability to any person or entity in connection with the foregoing.

Pursuant to 21 CFR 807.30, "Registration of a device establishment or assignment of a registration number does not in any way denote approval of the establishment or its products. Any representation that creates an impression of official approval because of registration or possession of a registration number is misleading and constitutes misbranding." The U.S. Food and Drug Administration does not issue a certificate of registration, nor does the U.S. Food and Drug Administration recognize a certificate of registration. CCTC is not affiliated with the U.S. Food and Drug Administration.

咨询网址: <https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfRL/rl.cfm>

Issued Date: 03/14/2020

Expiration Date/有效期至: 12/31/2020

Web: <http://www.fda.gov>

Tel: 1-888-INFO-FDA(1-888-463-6332)

e-mail: webmail@oc.fda.gov

U.S. Department of Health & Human Services

FDA

U.S. FOOD & DRUG ADMINISTRATION

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Cosmetics

Tobacco Products

Establishment Registration & Device Listing

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Medical Devices

Databases

1 result found for Establishment Registration or FEI Number : 3016733892

New Search

Establishment Name	Registration Number	Current Registration Yr
QINGDAO WEILI PROTECTIVE ARTICLES CO., LTD. CHINA	3016733892	2020
Disinfectant, Medical Devices - Hands-Washing-Free Sanitizer Gel		Foreign Exporter; Manufacturer
Radiographic Protective Glove - Disposable Vinyl Glove; Medical Gloves Non Sterile S/M/L/XL		Foreign Exporter; Manufacturer
Mask, Scavenging - KN95 Protective Mask		Foreign Exporter; Manufacturer
Non-Surgical Isolation Gown - Disposable Protective Clothing S/M/L/XL/XXL/XXXL		Foreign Exporter; Manufacturer
Accessory, Surgical Apparel - Disposable Medical Mask(Non-Sterile) N95 Medical Mask(Non-Sterile) KN95 Medical Mask(Non-Sterile) N95 Protective Mask 17.5*9.5cm N95 KN95		Foreign Exporter; Manufacturer
Show all 8 Listings For QINGDAO WEILI PROTECTIVE ARTICLES CO., LTD, CHINA		

TEST REPORT OF FFP2 CE



AC 114

CERTYFIKAT BADANIA TYPU UE (MODUŁ B) EU TYPE-EXAMINATION CERTIFICATE (MODULE B)

Nr
No. CW/PPER/3/08/2020

ZASWIADCZA SIĘ,

że Polski Rejestr Statków S.A. (PRS) przeprowadził procedurę badania typu wymienionego niżej wyrobu i stwierdził jego zgodność z wymaganiami określonymi w załączniku V do Rozporządzenia Parlamentu Europejskiego i Rady (UE) 2016/425 (PPE) w sprawie środków ochrony indywidualnej oraz uchylenia dyrektywy Rady 89/686/EEG, ze zmianami.

THIS IS TO CERTIFY

that Polski Rejestr Statków S.A. (PRS) did undertake the EU type-examination procedure for the product identified below which was found to be in compliance with the requirements of Annex V to the Regulation (EU) 2016/425 of the European Parliament and of the Council of 9 March 2016 on personal protective equipment and repealing Council Directive 89/686/EEC, as amended.

Wnioskodawca
Applicant

Qingdao Weili Protective Articles Co., Ltd.
Industrial Park, Ducun, Jiaoxi Office
Jiaozhou, Qingdao, Shandong, China 266327.

Producent
Manufacturer

Qingdao Weili Protective Articles Co., Ltd.
Industrial Park, Ducun, Jiaoxi Office
Jiaozhou, Qingdao, Shandong, China 266327.

Typ wyrobu
Product type

Sprzęt ochrony układu oddechowego. Półmaski filtrujące do ochrony przed cząstkami.
Respiratory protective devices. Filtering half masks to protect against particles.

Opis wyrobu
Product description

Półmaska filtracyjna, model: WL-P2095 (klasa FFP2 NR).
Filtering Half Mask, Model: WL-P2095 (FFP2 class NR).

Zastosowane normy
Specified standards

PN-EN 149+A1:2010 [EN 149:2001+A1:2009]

Niniejszy certyfikat pozostaje ważny do czasu uchylenia przy zachowaniu warunków uznania (patrz str. 2).
This certificate remains valid unless cancelled or revoked, provided the approval conditions (see page 2) are complied with.

Data ważności
Expiry date

2025-08-03



Dyrektor Pionu Certyfikacji
Certification Division Director

Michał Chudziński

Gdańsk, 2020-08-04



Nr jednostki notyfikowanej
No. of notified body
1463

NOTIFIKA (Polski Rejestr Statków S.A.)
NOTIFIED BODY (Polski Rejestr Statków S.A.)
80-416 Gdańsk, Poland

tel. (+48) (58) 346 17 00
fax (+48) (58) 341 77 69
e-mail: do@prs.pl
www: http://www.prs.pl/

CW/PPER/3/08/2020

Wykaz dokumentacji
List of documents

1. Instrukcja użytkownika - zatwierdzona przez PRS S.A. dnia 2020-07-31.
2. Ocena ryzyka - zatwierdzona przez PRS S.A. dnia 2020-07-31.
3. Dokumentacja techniczna „Półmaska filtracyjna, model: WL-P2095” - zatwierdzona przez PRS S.A. dnia 2020-07-31.
4. Raport z badań nr [2020] WSZ FHL NO.6952 wydany przez Jiangsu Guojian Testing Technology Co., Ltd. z akredytacją CNAS L10118 z dnia 2020-07-08.
5. Sprawozdanie z przeglądu PRS S.A. nr CW/PPER/MAP/18/2020 z dnia 2020-08-03.

1. Instruction of use - approved by PRS S.A. on 2020-07-31.
2. Risk analysis - approved by PRS S.A. on 2020-07-31.
3. Technical documentation "Filtering Half Mask, Model: WL-P2095" - approved by PRS S.A. on 2020-07-31.
4. Test report No. [2020] WSZ FHL NO.6952 issued by Jiangsu Guojian Testing Technology Co., Ltd. with CNAS accreditation no. L10118 dated on 2020-07-08.
5. PRS S.A. Survey Report No. CW/PPER/MAP/18/2020 dated on 2020-08-03.

Miejsca produkcji
(inne niż podane na stronie 1)
Places of production
(different than given on page 1)

Ograniczenia uznania
Approval limitations

1. Dane techniczne:
- półmaska filtrująca z regulowanym klipsem na nos,
- półmaska filtrująca wykonana z 5 warstwowej włókniny z filtrem z tkaniny,
- wymiary: 105x155 mm,
- docelowa grupa użytkowa: dorośli dla obu płci,
- kolor: biały.
2. Półmaska filtrująca przeznaczona do jednorazowego użytku.
3. Dokumentacja techniczna zatwierdzona w języku angielskim.
4. Produkt ten nie może być stosowany jako maska przeciwgazowa w środowisku toksycznym.
5. Półmaska filtrująca nie jest przeznaczona do użytkowania medycznego i chirurgicznego.

1. Specifications:

- particle filtering half mask with adjustable nose clip,
- particle filtering half mask made with 5 layers non-woven fabric with melt-blown fabric filter,
- size: 105x155mm,
- target group: unisex,
- color: white.

2. Respirator Mask shall not be used for more than one shift.

3. Technical documentation approved in English.

4. This product can not be used as a gas mask in a toxic environment.

5. Respirator Mask can not be used for medical and surgical purposes.

Warunki uznania
Approval conditions

1. Niniejszy certyfikat straci ważność po wprowadzeniu zmian lub modyfikacji w wyrobie bez uprzedniego uzgodnienia z PRS.
This certificate becomes invalid after changes or modifications to the product without prior agreement with PRS.

2. Znak zgodności może być umieszczony na uznanym wyrobie oraz może być wystawiona deklaracja zgodności tylko pod warunkiem, że łączna z badaniem typu UE zostanie przeprowadzona ocena zgodności produkcji pod nadzorem jednostki notyfikowanej, według załącznika VII lub VIII wymienionego wyżej rozporządzenia.

The Mark of Conformity may only be affixed to the above type approved product and a manufacturer's Declaration of Conformity issued provided the production is assessed under surveillance of a notified body according to Annex VII or VIII of the a/m Regulation.

证书报告 CERTIFICATES



CIBG
Ministerie van Volksgezondheid,
Welzijn en Sport

> Retouradres Postbus 16114 2500 BC Den Haag

Lotus NL B.V.
T.a.v. de heer X. Wei
Koningin Julianaplein 10
2595 AA 's-Gravenhage

Datum: 21 juli 2020
Betreft: notificatie medisch hulpmiddel klasse I

Geachte heer Wei,

Hierbij bevestig ik de ontvangst op 13 juli 2020 van de notificatie van het medische hulpmiddel klasse I, dat bedrijf QINGDAO WEILI PROTECTIVE ARTICLES CO., LTD., met Europees gemachtigde Lotus NL B.V., als fabrikant overeenkomstig Verordening (EU) 2017/745 (MDR) op de markt wenst te gaan brengen.

Het product is onder volgend kenmerk geregistreerd. Ik verzoek u om in alle verdere correspondentie over dit product het bijbehorende kenmerk te vermelden en het bij telefoongesprekken bij de hand te houden.

Disposable medical mask (non-sterile)
(geen merknaam) (NL-CA002-2020-52592)

Ik wijs u erop dat medische hulpmiddelen die op de markt gebracht worden volgens de MDR over een systeem voor hulpmiddelindicatie (UDI) moeten beschikken¹ en dat fabrikanten, gemachtigden en importeurs in de Europese databank voor Europese hulpmiddelen (Eudamed) moeten worden geregistreerd². Bijlage VI van de MDR bevat de bij de registratie te verstrekken gegevens.

Op dit moment is Eudamed nog niet in gebruik, zodat het wat betreft het bovenstaande voldoende is dat u uw product overeenkomstig de huidige wet- en regelgeving hebt genotificeerd.

Zodra Eudamed volledig in gebruik is, wordt de fabrikant of diens gemachtigde geacht binnen achttien maanden bovenstaand hulpmiddel te registreren in Eudamed.³

Farmatec

Bezoekadres:
Hoffen
Rijnstraat 50
2515 XP Den Haag
T 070 340 6661

<http://hulpmiddelen.farmatec.nl>

Inflichtingen bij:
M.P. Meijer - Michiels

medische_hulpmiddelen@
minvws.nl

Ons kenmerk:
CIBG-20203488

Bijlagen

Uw aanvraag
13 juli 2020

Correspondentie uitsluitend
richten aan het retouradres met
vermelding van de datum en het
kenmerk van deze brief.

Tevens wijs ik u er voor de goede orde nog op dat de registratie van uw mededeling betreffende de aflevering van het bovengenoemde product slechts een administratieve handeling betreft. Deze ontvangstbevestiging bevestigt dan ook geen besluit betreffende de kwalificatie van het desbetreffende product als medisch hulpmiddel in de zin van art. 1 WMH, noch betreffende de indeling in risicoklasse I.

De Minister voor Medische Zorg en Sport,
namens deze,

Afdelingshoofd
Farmatec

Dr. M.J. van de Velde

¹ O.g.v. art. 29 MDR.

² O.g.v. art. 31 MDR.

³ www.casct-europe.eu/wp-content/uploads/2018/05/FAQ_MDR_180117_V1.0-1.pdf. Zie vraag en antwoord nummer 20.

THE TEST REPORT FOR EN149:2001+A1: 2009 FROM CHINA COMPONENTS TEST



Test Report

Report No.: [2020] WSZ FHL NO.6952

Product Name Filtering half mask

Applicant Qingdao Weili Protective Articles Co.,Ltd.

Manufacturer Qingdao Weili Protective Articles Co.,Ltd.

Test Type Entrusted inspection

Jiangsu Guojian Testing Technology Co., Ltd.
3/F., Unit D, Xingye Building, Taihu International Tech-Park, Wuxi, Jiangsu, China

中国国检
检测通过
EN 149:2001
+A1: 2009
测试报告

Report No.: [2020] WSZ FHL NO.6952

Test Report

Product name	Filtering half mask	Model name	WL-P2095
		Brand	OMC
Laboratory/ Add	Jiangsu Guojian Testing Technology Co., Ltd./ 3/F., Unit D, Xingye Building, Taihu International Tech-Park, Wuxi, Jiangsu, China		
Applicant/ Add/Tel	Qingdao Weili Protective Articles Co.,Ltd./Dacun Industrial Park, Jiaoxi Office, Jiaozhou City, Qingdao City, Shandong Province, China/13963921920		
Manufacturer/ Add/Tel	Qingdao Weili Protective Articles Co.,Ltd./Dacun Industrial Park, Jiaoxi Office, Jiaozhou City, Qingdao City, Shandong Province, China/—		
Sample classification	FFP2	Sample number	GW6952-2020
Sample quantity	200 pcs	Date of receipt of sample	01/07/2020
Test type	Entrusted inspection	Article/Batch/Style number	—
Date (s) of performance of tests	01/07/2020-08/07/2020	Testing location	Same as the Laboratory
Sample state	Meeting the requirements of testing	Sample description	Refer to page 3
Test standard(s)	EN 149:2001+A1:2009 Respiratory protective devices - Filtering half masks to protect against particles - Requirements, testing, marking		
Test items	Packaging, material, practical performance, finish of parts, compatibility with skin, flammability, carbon dioxide content of the inhalation air, head harness, field of vision, penetration of filter material, breathing resistance, total inward leakage		
Test conclusion	The samples upon testing comply with FFP2 classification requirements according to the standard EN 149:2001+A1:2009. The details of test results see on Page 3-11		
Note	Date of issue: 23/07/2020		
	The test results presented in this report relate only to the submitted sample as received.		

Lu Bing 卢兵
Approver (name, signature)

Wan Heng 万恒
Reviewer (name, signature)

Yang Ying 杨莹
Chief Tester (name, signature)

THE TEST REPORT FOR GB2626-2006



TEST REPORT

(Electronic version)



VERIFICATION WEBSITE: www.gtgc.net.cn

VERIFICATION CODE: ROB-5495-34



No: 200151480

ISSUE DATE: 2020-06-09

APPLICANT: QINGDAO WEILI PROTECTIVE ARTICLES CO., LTD.
ADDRESS: DUCUN INDUSTRIAL PARK, JIAOXI OFFICE, JIAOZHOU CITY, QINGDAO CITY, SHANDONG PROVINCE.

INFORMATION CONFIRMED BY APPLICANT:

K95 MASK

QUANTITY: 70 PIECES

MANUFACTURE'S NAME: QINGDAO WEILI PROTECTIVE ARTICLES CO., LTD.

DATE RECEIVED/DATE TEST STARTED: 2020-06-01

CONCLUSION:

FILTRATION EFFICIENCY TO NaCl PARTICULATE MATTER M
INSPIRATORY RESISTANCE M
EXPIRATORY RESISTANCE M
VISUAL FIELD M
HEAD BAND M
FLAMMABILITY M
DEAD SPACE M
APPEARANCE INSPECTION[2 PIECES] M

NOTE: "M" -MEET THE STANDARD'S REQUIREMENT "F" -FAIL TO MEET THE STANDARD'S REQUIREMENT
"---" -NO COMMENT

REMARK:
THE AUTHORIZATION OF APPEARANCE INSPECTION AND DEAD SPACE IS NOT RECEIVED FROM CNAS.
THIS REPORT IS THE ENGLISH TRANSLATION VERSION OF THE REPORT 200130029.
ALL THE TESTED ITEMS ARE TESTED UNDER THE STANDARD CONDITION (EXCEPT FOR INDICATION).
COPIES OF THE REPORT ARE VALID ONLY RE-STAMPED.
THE EXPERIMENT WAS CARRIED OUT AT No.1, ZHJIANG ROAD, PANYU DISTRICT, GUANGZHOU, GUANGDONG, P.R.CHINA.

APPROVED BY:
Yuan Liu ENGINEER

刘圆



总部: 广州市番禺区江湾1号
花都实验室: 广州市花都区新街镇河滨西路1号

电话: 020-51994596/51994599
电话: 020-37721161

THE TEST REPORT FOR DISPOSABLE PROTECTIVE MASKS

检验检测报告

Test Report

STFWT202014028

共 2 页 第 1 页

Page 1 of 2

产品名称 Product Name	一次性防护口罩 (非医用)	规格型号 Specification Type	17.5cm*9.5cm
		商 标 Trademark	OMC
委托单位 Trust Unit	青岛威立防护用品有限公司	电 话 Tel	13963921920
生产单位 Manufacturer	青岛威立防护用品有限公司	样品等级 Sample Grade	—
样品数量 Sample Quantity	50 只	送样日期 Sample Receiving Date	2020-05-19
检验检测类别 Test Category	委托送样检验	批号/货号 Serial Number	—
样品状态 Samples Conditions	符合检测要求		
检验检测及判定依据 Document and Decide Accordance	GB/T 32610-2016 《日常防护型口罩技术规范》		
检验检测结论 Test Conclusion	检验检测结果见第二页		
备 注 Remarks	产品未明示性能等级, 综合判定按标准最低性能等级进行判定。 本报告检验结论仅对所检项目得出, 不代表未经检验的项目或功能符合要求。 本报告仅对来样负责。		



批准:
Approver

陈敏

审核:
Examiner

吴亮亮

主 检:
Major tester

凌青梅

SGS



中国认可
国际互认
检测
TESTING
CNAS L0599

Page 1 of 6

Test Result(s) : Unless otherwise stated the results shown in this test report refer only to the sample(s) tested, for further details, please refer to the following page(s).

Sample No.	Recommendation Level
(A)	FFP2 NR

Sara Ann

Sara Guo (Account Executive)

[illegible]

Member of the SGS Group (SGS SA)

PACKING PARAMETERS



50PCS/BOX



50pcs/bag(contains 10pcs/bag*5)
1000pcs/carton
The Carton Size: 685*595*26mm
Weight:7.5kgs/Carton

PACKING PARAMETERS



50PCS/BOX



50pcs/bag
2000pcs/carton
The Carton Size: 430*430*480mm
Weight:8.16kgs/Carton