General Equipment COVID-19

MEDICAL EQUIPMENT

KN95 Protective Mask

Product Features

4-Layer High-effective filter the bacteria structure.

- The First Layer--Double soft non-woven 70g, Absorbs dust, odors and harmful particles.
- The Second Layer-- ES Hot air filter cotton 50g, Pre-filtering the particulate matter and bacteria.
- The Third Layer—Melt blown non-woven 25g, Thicker than the others, that filters more than 98% of bacteria.
- The Forth Layer--Skin-friendly non-woven 50g, Not only filter particulates but also friendly to sensitive skin.

MEDICAL EQUIPMENT

Single Use Disposable Mask

Product Features

- 3- Ply
- Disinfection Treatment.
- Prevents spread of germs
- Single use

Includes CE Certification

Specifications:

- BFE (98%) ASTM F2101:2007
- Haemodialysis (ASTM F 1862, ISO22609) 120mmHg
- Anti-flaming 16CRF Pat 1610
- Material: non-woven fabrics, polypropylene fibre
- Size: 17.3*9.2
- Weight: 3g

MEDICAL EQUIPMENT

Certificate:

Certificate – Сертификат – 證明書 – Certificat – 증명서 – شهادة

Form QAT_10-M04, version 00, effective since March 6th, 2020

Certificate of Compliance

No. 1P200318.ZRMUS34

Certificate's Holder:	Zhengzhou Ruipu Medical Technology Co. Ltd. 13th Floor, Building A, Dianshang Building, Huanan 2 Road, Longhu Town, Xinzheng City, Zhengzhou City, Henan Province.
Certification ECM Mark:	
Product:	Disposable protective mask
Model(s):	Disposable Protective mask, Disposable kn95 mask
Verification to:	Standard: EN 149:2001+A1:2009 related to CE Directive(s): R 2016/425 (Personal Protective Equipment)

Remark: This document has been issued on a voluntary basis and upon request of the manufacturer. It is our opinion that the technical documentation received from the manufacturer is satisfactory for the requirements of the ECM Certification Mark. The conformity mark above can be affixed on the products accordingly to the ECM regulation about its release and its use.

Additional information and clarification about the Marking:

CE The manufacturer is responsible for the CE Marking process. This document has been issued on the basis of the regulation on ECM Voluntary Mark for the certification of products. RG01_ECM rev 3 available at: www.entecerma.it

Issuance date: 18 March 2020

Expiry date: 17 March 2025

Reviewer
Technical expert
Amanda Payne

Approver
ECM Service Director
Luca Sedponi

Ente Certificazione Macchine Srl
Via Ca' Bella, 243 – Loc. Castello di Serravalle – 40053 Valsamoggia (BO) - ITALY
☎ +39 051 6705141 🖷 +39 051 6705156. ✉ info@entecerma.it 🌐 www.entecerma.it

Corona Protection Visor

Protection Visor for full face protection against droplets and bacteria.

Covers up the face area and gives a double protection layer to additional googles and facemark which are usually base of the first layer.
The face shield decreases chances of infections enormous.

Corona Protection Visor

Certificate:

Shenzhen CTO Technology Service Co.,Ltd
9 / F, Block B, 196 Tangtou Avenue, Langxin Community, Shiyan Street, Baoan District, Shenzhen China.

CERTIFICATE OF CONFORMITY

Certificate No.	:	CTO200316099NCS
Applicant	:	Shenzhen Stardeal Industrial Co., Ltd
Address	:	No.18 Rongshuxia Industrial Zone, Tongxin Community, Baolong Street, Longgang District, Shenzhen, Guangdong, China
Manufacturer	:	Shenzhen Stardeal Industrial Co., Ltd
Address	:	No.18 Rongshuxia Industrial Zone, Tongxin Community, Baolong Street, Longgang District, Shenzhen, Guangdong, China
Product	:	Protective Face Shield
Trade Name	:	N/A
Model(s)	:	01
Test Report No.	:	CTO200316092KRS
Test Standards	:	**EN 166:2001**

The EUT described above has been tested by us with the listed standards and found in compliance with the council **PPE Directive 2016/425**. It is possible to use **CE** marking to demonstrate the compliance with this PPE Directive.

CE

For Chief Executive / Zaccheo Lee
Date: Mar. 16, 2020

This certificate of conformity is based on a single evaluation of the submitted sample(s) of the above mentioned product. It does not imply an assessment of the whole production and other relevant directives have to be observed.

Web:www.cto-cert.com E-mail: cto-service@163.com

Item Introduction:

Name：Transparent Protective FaceVisor

Material：PET Thickness 0.25mm

Weight：40g

Size ：32x22cm

Use：Direct Splash Protection

Packing：Each into Nylon bag and 200pcs to carton

Carton size: 70X36X45cm / GW 10.0kgs

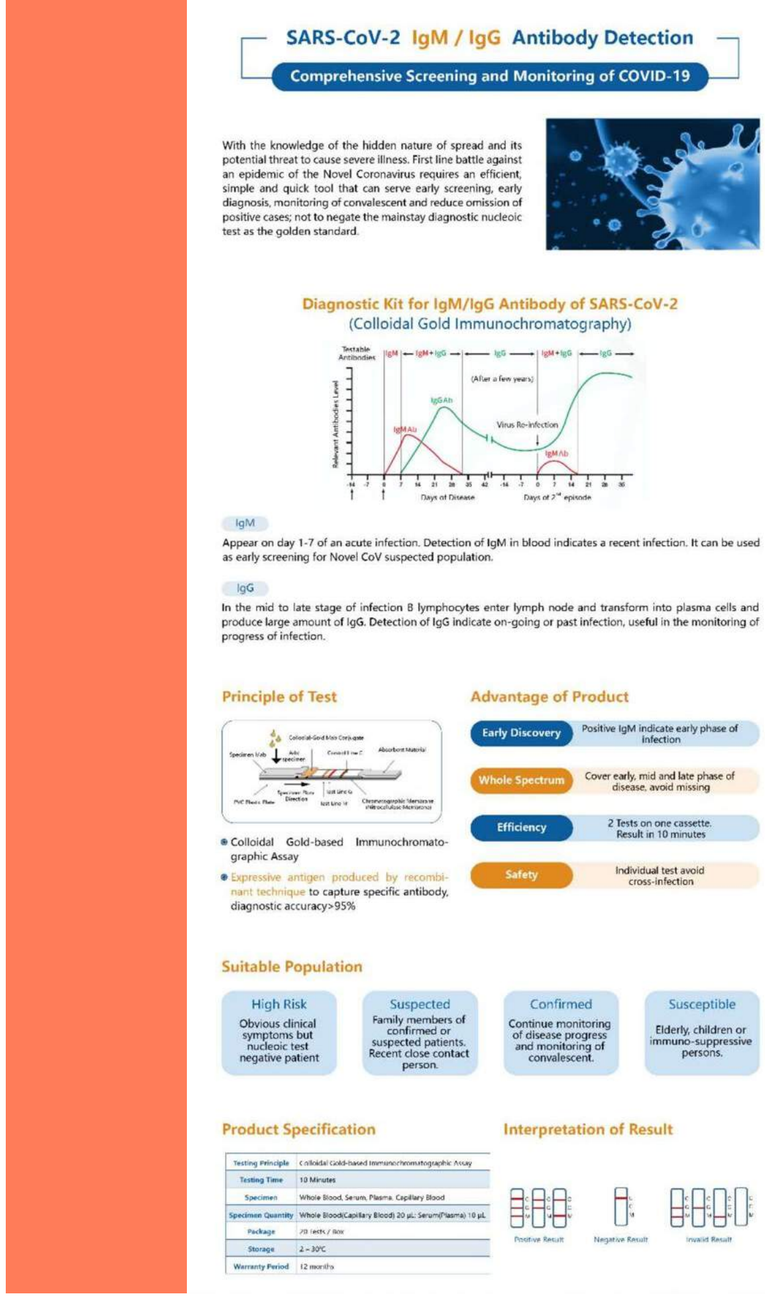
Quality Standard: ROHS EN166

Corona Virus Rapid Test Kit

- Human Use Coronavirus Disease Detection Device COVID-19 Rapid Test Kit for New Corona Virus Coronavirus AgG/AgM Test

There are three types of Novel coronavirus antibody test kit : 1,IgG/IgM three lines test kit ; 2,IgM/Total antibody three-line test kit ; 3,IgG antibody and IgM antibody two-lines test kit.

The test platform including Immunochromatography, ELISA and CLIA.



Corona Virus Rapid Test Kit

CERTIFICATE OF CONFORMITY

Certificate No.	:	ZUOCE200309610
Company Name	:	[redacted]
Company Address	:	[redacted]
Product Name	:	Diagnostic Kit for Antibody IgM/IgG of Novel Coronavirus COVID-19
Related Directives and Annex	:	98/79/EC In Vitro Diagnostic Medical Directive
Related Standards	:	EN ISO 13485:2016; EN ISO 14971:2012; EN 1041:2008; EN ISO 18113-2:2011; EN 13612: 2002; ISO 15223-1:2012; EN ISO 14155:2011; ISO 13640:2002; ISO 23640:2011 EN 980:2008
Examination Intent	:	*Examination the completeness of the Technical Documentation according to the requirements of the In Vitro Diagnostic Medical Directive 98/79/EC Annex III*
Review result	:	*During the examination of the provided Technical Documentation (No.:LR-TCF-001, Revision: A/0, dated 2020-MAR-06), no Non-compliance according to the requirements of the In Vitro Diagnostic Medical Directive 98/79/EC Annex III was detected.*
Valid From	:	03.09.2020
Valid Until	:	03.08.2025

CE

Authorized Signer :

Wallace Xu

Job Title : Certification Manager

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Date of Issue : 2020-03-09

Protective Coverall

Type 6 Protective Coverall	EN 13034
ICU Protective Coverall	EN14126 / GB19082
Hospitals Protective Coverall	CE Certified
Civil Protective Coverall	CE Certified

Protective Coverall

Certificate:

CENTROCOT
Innovation experience

EC TYPE-EXAMINATION CERTIFICATE

N° 97217151

According to Directive 89/686/EEC dated 21.12.1989 concerning the nearing of legislation in Member States, for what regards Personal Protective Equipment, and of Legislative Decree n 475 dated 4.12.92 and next amendment and supplement

Centro Tessile Cotoniero e Abbigliamento S.p.A.
P.zza Sant'Anna, 2 - 21052 Busto Arsizio (VA) - Italia

EEC Notified Body N° 0624

- in view of the firm's request submitted on the: 20th April 2017;
- in view of the positive results of the verification of the Technical File submitted by the firm together with the above mentioned request;
- taking into account the manufacturer Declaration reported in the Technical File attached to the above mentioned request
- in view of the conformity of technical specification of construction with basic requirements specified in annex II of law decree n. 475/1992 and the conformity of technical manufacturing documentation with those specifications;
- in view of the positive results of the tests carried out on the model representative for the production, in observance of what requested by paragraphs 6 and 8 of art. 7 of Legislative Decree n. 475/92 and by the Standard:

EN 13034:2005+A1:2009, EN ISO 13982-1:2004+A1:2010

Issues to:

Henan JoinKona Medical Products Stock Co., Ltd
Xinxing Road, South of Industry District, Lushan, Pingdingshan, Henan, China

the EC Type-Examination Certificate to following Personal Protective Equipment model:

OVERALL to protect against chemical hazards
Code TC0103-MIC

Category: III (third)

The model of Personal Protective Equipment is subject to the control system for the final product - Art. 11 A or B of the Directive 89/686/CEE

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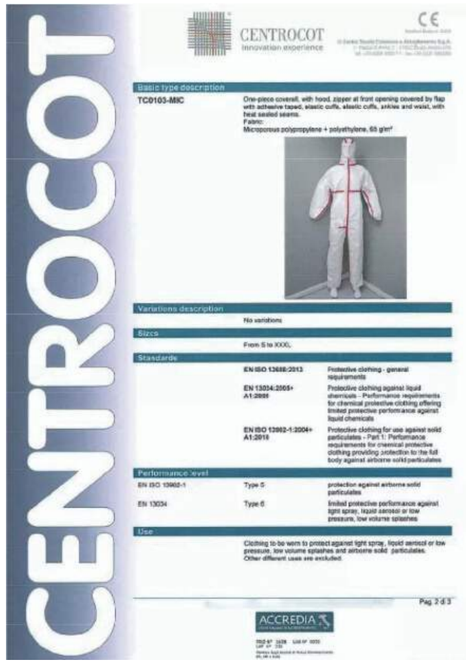
ACCREDIA



CENTROCOT
Innovation experience

Basic type description	
TC0103-MIC	One-piece coverall, with hood, zipper at front opening covered by flap with adhesive taped, elastic cuffs, elastic cuffs, ankles and waist, with heat sealed seams. Fabric: Microporous polypropylene + polyethylene, 65 g/m²

Variations description	
	No variations
Sizes	
	From S to XXXL
Standards	
EN ISO 13688:2013	Protective clothing - general requirements
EN 13034:2005+ A1:2009	Protective clothing against liquid chemicals - Performance requirements for chemical protective clothing offering limited protective performance against liquid chemicals
EN ISO 13982-1:2004+ A1:2010	Protective clothing for use against solid particulates - Part 1: Performance requirements for chemical protective clothing providing protection to the full body against airborne solid particulates
Performance level	
EN ISO 13982-1	Type 5 — protection against airborne solid particulates
EN 13034	Type 6 — limited protective performance against light spray, liquid aerosol or low pressure, low volume splashes
Use	
	Clothing to be worn to protect against light spray, liquid aerosol or low pressure, low volume splashes and airborne solid particulates. Other different uses are excluded.

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ACCREDIA

CENTROCOT
Innovation experience

Tests

The test results are contained in the report: 17RA00402, 17RA00498, 17RA00371

Marking

Following information are reported on the label inside the garment:

- EC marking
- Company name
- Article code
- Standards
- Pictograms standardized
- Maintenance symbols
- Fabric composition
- PPE category

Validity

- The present certificate has 5-year validity from the date of issue. On expiration date it will be responsibility of the Manufacturer to ask for the renewal.
- The applied standards are those in force on the date on the present Certificate. The validity of the certificate is not assured if reviewed standards are issued before its expiry date.
- Any modifications to the model and materials object of the present Certificate shall be notified and approved by Centro Tessile Cotoniero e Abbigliamento

Busto Arsizio,

Issue date:
20th April 2017

Expiry date:
20th April 2022

General Manager
Dr. Grazia Cerini

EC type-examination certificate n. 97217100 – Xiaotao TeiChan

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ACCREDIA

Instant Hand Sanitizer

- Instant Hand Sanitizer
  Kills 99.99% of germs and bateria
  NET: 100ml

INGREDIENTS
Alcohol, Aqua, Glycerin, Aloe Barbadensis Leaf Extract, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Triethanolamine.

DIRECTIONS
Squeeze gel onto your palm and massage your hands together until dry.

WARNING: FLAMMABLE
Keep away from the source of fire and flame.

FOR EXTERNAL USE ONLY
Avoid contact with the eyes.
Do not apply to damaged, inflamed or broken skin.
If skin allergies occurs, stop using.

- Keep out of the reach of children
- Use only under adult supervision
- Non staining, may discolor certain fabrics

Instant Hand Sanitizer

Certificate:

Fiscal Year 2020

CERTIFICATION OF FDA REGISTRATION

This certifies that:

SINGWONG ASIA PACIFIC (BOLUO)CO.,LTD.
Ping'an Industrial Park,Baitang Town, Boluo County Huizhou, Guangdong, 516008, CHINA

has completed the FDA Establishment Registration and Device Listing with the US Food & Drug Administration, through

Shenzhen CCT Testing Technology Co., Ltd.

Number: [redacted]

Device Listing#:

Listing No	Code	Device Name	Proprietary Name
[redacted]		Disinfectant, medical devices	INSTANT HAND SANITIZER, 50ml,60ml,100ml,250ml,300ml,500ml,1000ml

CCT 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. CCT 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. CCT assumes no liability to any person or entity in connection with the foregoing.

Pursuant to 21 CFR 807.39, "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. CCT is not affiliated with the U.S. Food and Drug Administration.

Chief engineer
Issued: 03/18/2020
Expiration Date:12/31/2020

Isolated stretcher

- Full air circulation whithin the chamber, by means of a carbon filter and blower, enables patients to remain inside long enough for a full diagnosis and intial treatment, whilst eliminating any expulsion of contaminated air to the outside
  Rechargeable batteries allow full mobility of the units for up to 10 hours of continuous operation

  A fully transparent chamber, fitted with conduit sleeves to allow diagnosisand medical treatment. Also fitted are facilities for connecting monitoring equipment cables, ventilation tubes or intravenous drips

  The units can accommodate patients of all ages, whether conscious, unconscious, or on ventilation

  Isolation chambers allow full operation of the stretcher or pediatric bed- including height adjustment, back-rest adjustment and trendelenburg function for CPR administration