

To: 'Middelen Corona'[5.1.2e]@NFU.nl]
From: coronabeschermingsmiddelen
Sent: Tue 3/24/2020 10:27:17 AM
Subject: FW: Beschermingsmiddelen verkopen
Received: Tue 3/24/2020 10:27:17 AM
[FFP2](#)

Van: 5.1.2e

Verzonden: dinsdag 24 maart 2020 11:00

Aan: coronabeschermingsmiddelen

Onderwerp: Re: Beschermingsmiddelen verkopen

Geachte heer/mevrouw,

Na aanleiding van onderstaande email een korte toelichting op onze werkwijze. Wij werken met een vaste leverancier en hebben een lokaal team die alle leveringen controleert. Wij kunnen ook andere producten leveren waaronder overalls, wegwerp overschoenen en wegwerp bedovortrekken.

Hierbij onze prijzen voor de mondmaskers:

- KN95 FFP2 (zie bijlage) - 5.1.1c - 5.1.1c per stuk DDU

- Standaard 3 ply - QTY 100.000 - 5.1.1c per stuk DDU

Payment terms: 5.1.1c

Shipping time: 5.1.1c

Wij voeren op locatie een kwaliteitscontrole uit en kunnen hierdoor de levering ook garanderen. De mondmaskers zijn CE gecertificeerd en kunnen ook in grotere hoeveelheden dan hierboven vermeld afgenomen worden (tot 1.000.000 stuks per levering).

Ik hoor graag van u, bij verdere vragen kunt u mij ook bellen op onderstaand nummer.

Kind regards / Met vriendelijke groet,

5.1.2e

| BD | LADM Ltd. Israel-Ghana-Bulgaria-Kenya-Uganda-Tanzania-Nigeria-China-Holland | Tel : +31(0)6

5.1.2e

5.1.2e

European
Commission

Verification of Conformity

Applicant: [REDACTED]
 Address: [REDACTED]
 Product(s): Overall, Lab Coat, Shoe Cover, Pillow Case, Sleeve Cover, Apron,
 Face Mask, Cap, Surgical Gown, Patient Gown, Isolation Gown, Bed
 Sheet/Bed Cover
 Type(s): See annex
 Product Classification: Class I

The submitted technical files including test report of the above products have been reviewed against the self declaration requirements of conformity for CE marking according to Medical Device Directive (93/42/EEC).

Standard(s) used for showing compliance with the essential requirements in the specified directive(s):

Standard(s): EN ISO 14971:2012; EN ISO 15223-1:2016;
 EN 1041:2013; EN ISO 10993-1:2009/AC:2010;
 EN ISO 10993-5:2009; EN ISO 10993-10:2013

The review result of the technical files and test report support the self declaration for the devices listed above. Where the manufacturer affix's the CE marking to the product listed they must ensure that all the requirements of the appropriate EU directive(s) have and continue to be met.

SUNGO Cert GmbH



Executive Director
 Issued: Dec. 10 2019
 Cert. No.: EU158518
 Expiration Date: Dec. 9 2024



Fiscal Year 2020 CERTIFICATION OF REGISTRATION

This certifies that:

[REDACTED]
has completed the FDA Establishment Registration (as manufacturer , contract manufacturer) and Device Listing with the US Food & Drug Administration, thr

U.S. Agent for FDA
Communications:

SUNGO TECHNICAL SERVICE INC.
6050 W EASTWOOD AVE APT 201, CHICAGO,
ILLINOIS 60630, USA
Telephone: +1-855-957-7779 / E-mail: sungo.group@yahoo.c

Registration Number: [REDACTED]

Device Listing#: See annex

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

Pursuant to 21 CFR 807.39, "Registration of a device establishment or assignment of a regi number does not in any way denote approval of the establishment or its products. Any represe that creates an impression of official approval because of registration or possession of a regi



Fiscal Year 2020 CERTIFICATION OF REGISTRAT

Annex to Cert. No.: [REDACTED]

| Listing No | Code | Device Name |
|------------|------|--|
| D090469 | OEA | Non-surgical isolation gown (Non-Surgical Isolation Gown) |
| D098875 | FYF | CAP, SURGICAL (Cap, Surgical) |
| D098877 | LYU | ACCESSORY, SURGICAL APPAREL (Accessory, Surgical Apparel) |
| D119564 | FXO | SUIT, SURGICAL (Coveralls / Scrub Suits) |
| D200578 | FXP | COVER, SHOE, OPERATING-ROOM (Cover, Shoe, Operating-Room) |
| D268879 | KME | BEDDING, DISPOSABLE, MEDICAL (Bedding, Disposable, Medical) |

END OF THE ANNEX

CIC

CIC

Certificate of Registration

Certificate No.:282Q19060618121

Awarded to

[REDACTED] Co.,Ltd

Organization Code No./Unified Social Code:91330802MA2DG8140T
 address:No.1,Fengshan 1st Road, Hangbu Town, Kecheng District, Quzhou
 Zhejiang Province.

Beijing CIC Inspection & Certification CO.,LTD (CIC) certify that
 Quality Management System of the above organization has been assessed and
 in accordance with the requirements of the standard:

GB/T19001-2016/ISO9001:2015

SCOPE OF CERTIFICATION/REGISTRATION

Clothing manufacturing and sales.

(Qualifications-based operations requiring qualification permit)

This certificate is made valid when used with certification scopes and the requirements
 laws and regulations. These requirements include but are not limited to administrative
 scopes of qualifications, and CCC requirements.

Subject to operation conditions in requirements conformity with Quality Management

This Certificate is valid for a period of three years only.

Date from:Jun.18th,2019 To:Jun.17th,2022

The effectiveness of this Certificate shall be Validated by periodic surveillance audit or

**The time limit of the certificate is to Jun.11th,2020,please contact
 the surveillance or re-certification assessment before Jun.11th**

If the assessment is overdue,the certificate is invalid.

CERTIFICATE

شهادة - 증명서 - Certificat - 證明書 - Сертификат - Certificate of Compliance

Certificate of Compliance

No. 4J200310.ZBI0W99

Technical Construction File no. TCF-GGC-20200306-PPE-214

Certificate's
Holder:

[Redacted] Garmer
Co., Ltd.
No.1 Fengshan 1 road industrial park, Hang
town, KeCheng, QuZhou, ZheJiang, China

Certification ECM
Mark:



Product:
Model(s):

Single-Use Face Mask
Ear-hang Type

Verification to:

Standard:
EN 149:2001+A1:2009

related to CE Directive(s):
R 2016/425 (Personal Protective Equipment)

Remark: The product(s) has been verified on a voluntary basis. The product(s) satisfies the requirements of the Certification Mark of ECM, in reference to the above listed Standard(s). The above Compliance Mark can be affixed on the product(s) accordingly to the ECM regulation about its release and regulation can be found at www.entecerma.it. This Certificate of Compliance can be checked at www.entecerma.it.

This verification doesn't imply assessment of the production of the product(s).

Additional information, clarification about the **CE** Marking:

CE

We attest that a TCF for the **CE** Marking process is in place. The Manufacturer is Responsible to start the **CE Marking Certification Process** on an appointed Notified Body and to perform all the necessary activities, as required by the Directive and accepted by the Notified Body, before placing the product(s) on the market.

Date of issue 10 March 2020

Expiry date 09 March 2021

Chief Manager
Marta Morina

Deputy Manager
Amanda Poy

认证证书

CERTIFICATE

认证证书

POSI

CERTIFICATE

This is to certify that the Quality Management

[REDACTED]
Business license number: 91[REDACTED]6

Registered Address: No.6, HeFeng Industrial Park, [REDACTED]

Audit Address: No.6, HeFeng Industrial Park, [REDACTED]

applicable to

**Production and sale of disposable non sterilization
products(clothing,masks,sleeves,hats,shoe cover,
cover,bear sleeve,sheets,pillpwcase,aprons)(Only**

has been assessed and registered by POSI against the
ISO13485:2016

This registration is subject to the company maintaining a quality management system
to the above standard, which will be monitored by POSI

Please consult the website: www.posicert.com

The certificate information is also available on the CNCA official website



Op 23 mrt. 2020, om 11:50 heeft coronabeschermingsmiddelen <coronabeschermingsmiddelen@minvws.nl> het volgende geschreven:

Beste heer/mevrouw,

Nogmaals dank voor uw mail.

Wij hebben uw aanbieding beoordeeld en we moeten u helaas melden dat we op dit moment niet op uw aanbod ingaan.

Het centraal inkooppunt zal eerst aan de slag gaan met partijen die op korte termijn grote aantallen adequate beschermingsmiddelen kunnen leveren en direct in contact staan met leveranciers.

Niettemin wil ik u hartelijk danken voor het meedenken.

Met vriendelijke groet,

Directie Geneesmiddelen en Medische Technologie

Team Corona

Ministerie van Volksgezondheid, Welzijn en Sport

Parnassusplein 5 | 2511 VX | Den Haag

Postbus 20350 | 2500 EJ | Den Haag |

-----Oorspronkelijk bericht-----

Van: [REDACTED] <[REDACTED]@live.nl>

Verzonden: maandag 16 maart 2020 17:14

Aan: coronabeschermingsmiddelen <coronabeschermingsmiddelen@minvws.nl>

Onderwerp: Beschermingsmiddelen verkopen

Geachte heer/mevrouw,

Na aanleiding van uw oproep aan Nederlandse ondernemers voor het aanvullen van mondkapjes en andere beschermingsmiddelen voor ziekenhuizen stuur ik u deze email.

Wij leveren al geruime tijd 'medical equipment' aan landen in Oost-Europa en Afrika vanuit voornamelijk China, India en Turkije. Vanwege de onzekerheid in de wereld is het exporteren vanuit deze landen moeilijker geworden en zijn de prijzen gestegen. Vorig jaar kostte een mondkapje bij onze Chinese producent \$0,04, twee weken geleden \$0,42 en vandaag de dag \$0,60. Hierom hebben wij vandaag een order van 500.000 mondkapjes moeten annuleren.

Graag ontvangen wij een overzicht van de benodigde producten en zullen hier spoedig op reageren. Wij zullen na het ontvangen van een overzicht direct bij onze leveranciers inventariseren welke producten inclusief volumes geleverd kunnen worden tegen welke prijs.

Wat verder een idee kan zijn en waar wij graag over willen sparren is het fabriceren van bepaalde beschermingsmiddelen in eigen land. Onze partner heeft de Israëliëse overheid geadviseerd bij de aankoop van de juiste machines om mondkapjes en latex handschoenen te produceren. Door het beschikbaar stellen van een locatie kan deze faciliteit binnen 4 weken operationeel zijn en kan deze deels voorzien in de huidige en toekomstige vraag. Ik hoor graag van u.

Met vriendelijke groet,

[REDACTED]

06-25450685