



Florina
Iustina
Ionescu

Digitally signed by Florina Iustina Ionescu
DN: cn=Florina Iustina Ionescu, sn=Ionescu, givenName=Florina Iustina, serialNumber=201104209F811, email=ionescuscmyahoo.ro, ou=Comitet director, o=Spitalul Clinic Judetean de Urgenta Arad, I=ARAD, st=ARAD, c=RO, title=Director financiar contabil
Date: 2020.05.21 12:12:59 +0300

spital <scjuarad.bap@gmail.com>

Solicitare clarificare negociere 13 loturi

1 mesaj

spital <scjuarad.bap@gmail.com>

22 aprilie 2020, 18:33

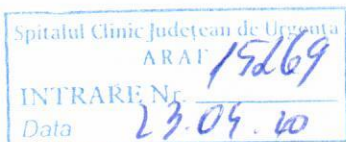
Către: suport@neramo.ro

Buna ziua,

va rugam sa ne transmiteti certificatul de conformitate pentru loturile ofertate in cadrul procedurii asumat cu semnastura si stampila ofertantului.

Termen de raspuns 23.04.2020 ora 10:00.

Multumim.



spital <scjuarad.bap@gmail.com>

Solicitare clarificare negociere 13 loturi

2 mesaje

spital <scjuarad.bap@gmail.com>

22 aprilie 2020, 18:33

Către: suport@neramo.ro

Buna ziua,

va rugam sa ne transmiteti certificatul de conformitate pentru loturile ofertate in cadrul procedurii asumat cu semnastura si stampila ofertantului.

Termen de raspuns 23.04.2020 ora 10:00.

Multumim.

suport <suport@neramo.ro>

23 aprilie 2020, 08:57

Către: spital <scjuarad.bap@gmail.com>, Vanzari Neramo <vanzari@neramo.ro>

Buna ziua

Atasat veti gasi certificate de conformitate pentru loturile ofertate.

Daca mai aveti nevoie de alte informatii nu ezitati sa ne contactati.

Cu stima

Valery Enache

0773 366258

[Textul citat a fost ascuns]

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Cu stima

Valery Enache

NERAMO Distribution srl

Tel: (004)0773 366 258

 **certificate.pdf**
1090K

CELAB®

Via Maira snc
04100 Latina
Italy
celab@celab.com

celab ✓

CERTIFICATE

Certificate Number UCN : 802695723197
Job : J28921
Date of Issue : 2020-03-10
Certificate valid up to : 2024-03-09

Brand Name : HONSUN
Type : Goggles
Model N : Y-01, C-01, X-01, T-01, S-01, G-01

Manufacturer : Jiangsu Hongxu Desheng Technology Co., Ltd.
Address : Xinghu Road, Development Zone, Danyang City,
Jiangsu Province China

Standard Used : EN 166:2001

Conclusion :

After inspection of the technical documentation issued by the customer, and in his request, we express our opinion that the product meets the technical requirement of the following directives and standards:
(EU)2016/425 Personal protective equipment (PPE)

This opinion is only valid for the directive, the equipment and configuration described, in conjunction with the test data detailed above and with compliance with all applicable legal requirement for the product .
The following manufacturer documents was inspected:

Presence of Declaration of conformity template	✓ OK
Presence of test report using standards as indicated in the declaration of conformity Test report reference : (2020)wei song zi spectacles 0292	✓ OK
Presence of CE symbol in the product label.	✓ OK
Presence of instruction manual	✓ OK
Use of valid Harmonized standard in the declaration of conformity	✓ OK
Presence of product description in the technical construction file	✓ OK

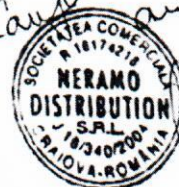
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Massimiliano Bertoldi
General Manager - CELAB
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Doc 121 Voluntary Certificate rev 3.32



EC Certificate
Directive 93/42/EEC Annex V
Production Quality Assurance
Medical Devices



Registration No.: DD 60111758 0001

Report No.: 15095978 001

Manufacturer: Sunsmmed Protective Products Ltd.
No. 18, Industrial Park, Maozui Town
Xiantao City
433000 Hubei
China

Products:

Aspects of manufacture concerned with securing and maintaining sterile conditions:
Sterile Surgical Caps, Sterile Surgical Face Masks,
Sterile Surgical Gowns, Sterile Bed Protections,
Sterile Surgical Drapes, Sterile Surgical Packs

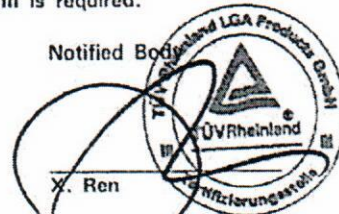
Expiry Date: 2021-08-11

The Notified Body hereby declares that the requirements of Annex V of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Effective Date: 2016-08-12

Date: 2016-08-12

Notified Body



TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.

