

## CE ATTESTATION OF CONFORMITY

Related Directives: Medical Devices Directive (93/42/EEC)

CLASS: CLASS 1

Description of Product: PROTECTIVE GOWN WITH RIB AT CUFFS

PROTECTIVE GOWN WITH ELASTICATED CUFFS PROTECTIVE GOWN WASHABLE AND REUSABLE

PROTECTIVE MEDICAL PYJAMA

PROTECTIVE POLYPROPYLENE FACE MASK

PROTECTIVE COVERALL

PROTECTIVE SHOES - PROTECTIVE CAP

FABRIC COMPOSITION: 100% POLYPROPYLENE

Manufactured by

## **CROSSING SARL**

LOT 140 HAY MLY RACHID BEN M'SICK CASABLANCA MAROC

Certificate No.: SISMAR062020103 **Issue Date (Original): 02.06.2020 Issue Date(Latest): 02.06.2020** 

**Expiry Date: 01.06.2021** 

Re-Certification date: 02.05.2023



This Certificate is issued under the following conditions:

1.It applies only to the above referenced models of the medical devices.

2.It does not imply that the SIS has performed any surveillance or control of their manufacture.

3. The manufacture is obligated to assure conformity of all in medical devices of

the respective model to type assessed by the mean of this certificate.

4. The certificate remains valid until the manufacturing condition, the quality system or relevant legislation are changed.

5. After fulfilling of the relevant EU legislation requirements, the manufacture shall affix to each medical device, of the above referenced models, the CE-marketing according to this example:

## Remark:

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

Additional information and clarification about the marking:

The manufacturer is responsible for the CE marking process. This document has been issued on the basis of the regulation on SIS Voluntary mark for the certification of products.



Note: This certificate is valid only if produced with the

continuation letter after the surveillance is carried out successfully.

documentation and Implementation has been reviewed and found to The Organization's comply with the relevant standard rules. This certificate of Registration is based on the evaluation of the mentioned scope given above. Organization is responsible for maintaining the responsibilities of the relevant standard rules. Any significant changes in the scope of the certification or standard referred above render this certificate invalid. This is an accredited certificate issued by SIS Certifications Pvt. Ltd. sanctioned for issue by International Organization for Accreditation Services, Burn point P.O.Box.293 St.George's, Greneda.

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The status of this certificate can be verified on "http://www.siscertifications.com



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