



EC Certificate

Full Quality Assurance System

Certificate No.:
239534-2017-CE-IND-NA-PS Rev. 1.0

Project No.:
PRJC-535567-2015-MSL-IND

Valid Until:
06 September 2022

This is to certify that the quality system of:

MRK HEALTHCARE PRIVATE LIMITED

S. No. 153/P& 310, Panch Pippal, Hansapur, Runi, Unjha-Patan Road, Patan-384265, Gujarat, India

For design, production and final product inspection/testing of:

Disposable Medical Devices

Has been assessed with respect to:

The conformity assessment procedure described in Article 11.3.a and Annex II excluding section 4 (Module H2) of Council Directive 93/42/EEC on Medical Devices, as amended

and found to comply.

Further details of the product(s) and conditions for certification are given overleaf.

Place and Date:
Høvik, 16 May 2018



For:
DNV GL NEMKO PRESAFE AS

Mariann Jeremiassen

The Certificate has been digitally signed.
See www.presafe.com/digital_signatures for more info

Notice: The Certificate is subject to terms and conditions as set out in the Certification Agreement. Failure to comply may render this Certificate invalid.

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Jurisdiction

Application of Council Directive 93/42/EEC of 14 June 1993, adopted as “Forskrift for Medisinsk Utstyr” by the Norwegian Ministry of Health and Care Services.

Certificate history:

Revision	Description	Issue Date
	Original Certificate	2017-09-06
1.0	Addition of Foley Catheter	2018-05-16

Products covered by this Certificate:

Product Description	Product Name	Class
Surgical Rubber Gloves and Speciality Gloves	Surgical latex gloves - sterile & non-sterile (Powdered Gloves, Powder Free Chlorinated Gloves, Powder Free Polymer Coating Gloves) Elbowlength gynaecology procedure latex gloves - sterile & non-sterile (Powdered Gloves, Powder Free Chlorinated Gloves, Powder Free Polymer Coating Gloves) Double pair speciality gloves (high risk gloves) - sterile (Powdered Gloves, Powder Free Chlorinated Gloves, Powder Free Polymer Coating Gloves) Orthopaedic speciality gloves - sterile (Powdered Gloves, Powder Free Chlorinated Gloves, Powder Free Polymer Coating Gloves) Micro surgery speciality gloves - sterile (Powdered Gloves, Powder Free Chlorinated Gloves, Powder Free Polymer Coating Gloves) Ultra Nulife sterile surgical gloves (beadless & beaded gloves) (Powdered Gloves, Powder Free Chlorinated Gloves, Powder Free Polymer Coating Gloves)	Ila
Examination Latex Gloves	Sterile & non-sterile (Powdered Gloves, Powder Free Chlorinated Gloves, Powder Free Polymer Coating Gloves)	Is
Examination Latex-free	Sterile & non-sterile (Powdered Gloves, Powder	Is

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Gloves	Free Chlorinated Gloves, Powder Free Polymer Coating Gloves)	
Non-active devices for emergency & intensive care	Penrose Drainage Tubing	Ila
	Male Incontinence device (U-Drain)	Is
	Foley Balloon Catheter - Three Way and Two Way (for urine drainage from bladder)	Ila

The complete list of devices is filed with the Notified Body

Sites covered by this certificate

MRK HEALTHCARE PRIVATE LIMITED, S. No. 153/P& 310, Panch Pippal, Hansapur, Runi, Unjha-Patan Road, Patan-384265, Gujarat, India

EU Representative

Obelis s.a, 34, Av de Tervuren, Bte 44, B-1040 Brussels, BELGIUM

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Terms and conditions

The certificate is subject to the following terms and conditions:

- Any producer (see 2001/95/EC for a precise definition) is liable for damage caused by a defect in his product(s), in accordance with directive 85/374/EEC, as amended, concerning liability of defective products.
- The certificate is only valid for the products and/or manufacturing premises listed above.
- The Manufacturer shall fulfil the obligations arising out of the quality system as approved and uphold it so that it remains adequate and efficient.
- The Manufacturer shall inform Presafe of any intended updating of the quality system and Presafe will assess the changes and decide if the certificate remains valid.
- Periodical audits will be held, in order to verify that the Manufacturer maintains and applies the quality system. Presafe reserves the right, on a spot basis or based on suspicion, to pay unannounced visits.

The following may render this Certificate invalid:

- Changes in the quality system affecting production.
- Periodical audits not held within the allowed time window.

Conformity declaration and marking of product

When meeting with the terms and conditions above, the producer may draw up an EC declaration of conformity and legally affix the CE mark followed by the Notified Body identification number of Presafe.

End of Certificate