EC CERTIFICATE

Number: 2153004CE03

Full Quality Assurance System

Directive 93/42/EEC on Medical devices, Annex II excluding (4)
(Devices in Class IIa, IIb or III)

Manufacturer:

Steripack S.A.
Zona Industrial 1 - Lote 11 a 14
4560-164 Guilhufe
Penafiel
Portugal

For the product category(ies)
Anticoagulant for use as a catheter lock

DEKRA grants the right to use the EC Notified Body Identification Number illustrated below to accompany the CE Marking of Conformity on the products concerned conforming to the required Technical Documentation and meeting the provisions of the EC-Directive which apply to them:

0344

Documents, that form the basis of this certificate:

Certification Notice 2153006CN, initially dated 8 September 2012
Addendum, initially dated 26 January 2015

DEKRA hereby declares that the above mentioned manufacturer fulfils the relevant provisions of 'Besluit Medische Hulpmiddelen', the Dutch transposition of the Council Directive 93/42/EEC of June 14, 1993 concerning Medical devices, including all subsequent amendments. The manufacturer has implemented a quality assurance system for design, manufacture and final inspection for the above mentioned product category in accordance to the provisions of Annex II of Council Directive 93/42/EEC of June 14, 1993 and is subject to periodical surveillance. For placing on the market of Class III devices an additional EC design examination certificate according to Annex II (4) is mandatory. The necessary information related to the quality management system of the manufacturer, including facilities and the reference to the relevant documentation, of the products concerned and the assessments performed, are stated in the Certification Notice which forms an integrative part of this certificate.

This certificate is valid until: 1 September 2023
Issued for the first time: 26 January 2015
Revised: 4 June 2019
Reissued: 2 November 2018

DEKRA Certification B.V.

B.T.M. Holtus
Managing Director

J.A. van Vugt
Certification Manager

DEKRA Certification B.V. is Notified Body with ID no 0344

DEKRA Certification B.V. Meander 1051, 6825 MJ Arnhem P.O. Box 5185, 6802 ED Arnhem, The Netherlands
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ADDENDUM

Belonging to certificate: 2153004CE03

CE MARKING OF CONFORMITY
MEDICAL DEVICES

Anticoagulant for use as a catheter lock

Issued to:

Steripack S.A.
Zona Industrial 1 - Lote 11 a 14
4560-164 Guilhufe
Penafiel
Portugal

This certificate covers the following product(s):

- Citra-Lock S 4%, sterile syringe containing 2,5 ml 4% Trisodium Citrate solution

Initial date: 26 January 2015
Revision date: 4 June 2019

DEKRA Certification B.V.

B.T.M. Holtus
Managing Director

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