

EC CERTIFICATE

Number: 2153004CE02

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)

Sterile catheterization accessory sets, sterile single use forceps, clamps and scissors, sterile and non sterile dressing sets, sterile dialysis sets, sterile delivery sets, sterile surgical sets, sterile washing sets, sterile and non sterile gauze and non woven swabs and balls and sterile syringes with 0,9% sodium chloride

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 2153004CN, initially dated 8 September 2012

Addendum, initially dated 8 September 2012

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.

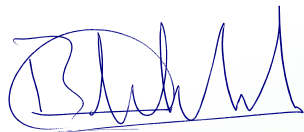
This certificate is valid until: 1 September 2023

Issued for the first time: 8 September 2012

Revised: 4 June 2019

Reissued: 2 November 2018

DEKRA Certification B.V.



B.T.M. Holtus
Managing Director



J.A. van Vugt
Certification Manager

© Integral publication of this certificate and adjoining reports is allowed

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
T +31 88 96 83000 F +31 88 96 83100 www.dekra-certification.com Company registration 09085396

ADDENDUM

Belonging to certificate: 2153004CE02

1/2

CE MARKING OF CONFORMITY MEDICAL DEVICES

Sterile catheterization accessory sets, sterile single use forceps, clamps and scissors, sterile and non sterile dressing sets, sterile dialysis sets, sterile delivery sets, sterile surgical sets, sterile washing sets, sterile and non sterile gauze and non woven swabs and balls and sterile syringes with 0,9% sodium chloride

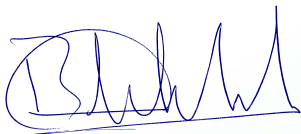
Issued to:

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

This certificate covers the following product(s):

- Sterile catheterization accessory sets
- Sterile and non sterile dressing sets
- Sterile dialysis sets
- Sterile delivery sets
- Sterile surgical sets
 - o Sterile anaesthesia sets
 - o Sterile puncture sets
 - o Sterile suture accessory sets
- Sterile washing sets
- Sterile and non sterile gauze and non woven swabs and balls
 - o Sterile and non sterile gauze balls
 - o Sterile and non sterile gauze balls on a pin
 - o Sterile and non sterile gauze balls star shape
 - o Sterile and non sterile gauze balls tampon
 - o Sterile and non sterile gauze balls triangular shape
 - o Sterile and non sterile gauze balls with X-ray
 - o Sterile and non sterile gauze swabs with X-ray
 - o Sterile and non sterile abdominal swabs with X-ray
 - o Gauze swabs flat
 - o Preparation balls with X-ray

DEKRA Certification B.V.



B.T.M. Holtus
 Managing Director



J.A. van Vugt
 Certification Manager

© Integral publication of this certificate and adjoining reports is allowed

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
 T +31 88 96 83000 F +31 88 96 83100 www.dekra-certification.com Company registration 09085396

ADDENDUM

Belonging to certificate: 2153004CE02

2/2

CE MARKING OF CONFORMITY MEDICAL DEVICES

Sterile catheterization accessory sets, sterile single use forceps, clamps and scissors, sterile and non sterile dressing sets, sterile dialysis sets, sterile delivery sets, sterile surgical sets, sterile washing sets, sterile and non sterile gauze and non woven swabs and balls and sterile syringes with 0,9% sodium chloride

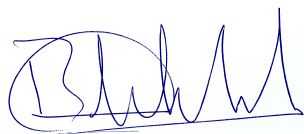
Issued to:

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

- o Sterile and non sterile non woven balls
- o Sterile and non sterile non woven balls triangular shape
- o Non woven balls on a pin
- o Non woven balls with X-ray
- o Non woven swabs with X-ray
- o Sterile syringes with 0,9% sodium chloride.
- Sterile forceps, clamps and scissors

Initial date: 8 September 2012
 Revision date: 4 June 2019

DEKRA Certification B.V.



B.T.M. Holtus
 Managing Director



J.A. van Vugt
 Certification Manager

© Integral publication of this certificate and adjoining reports is allowed

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
 T +31 88 96 83000 F +31 88 96 83100 www.dekra-certification.com Company registration 09085396