EC CERTIFICATE

Number: 2113863CE01

Full Quality Assurance System

Directive 93/42/EEC on Medical devices, Annex II excluding (4)

(Devices in Class IIa, IIb or III)

Manufacturer:

Oté Pharma Sol B.V.

Vluchtoord 38 5406 XP Uden The Netherlands

For the product category(ies)

Multipurpose Solutions and Cleaning Solutions for Contact Lens Care

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 2113863CN, initially dated 10 July 2008 Addendum, initially dated 10 July 2008

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 July 2023 Issued for the first time: 10 July 2008 Reissued: 2 July 2018

DEKRA Certification B.V.

B.T.M. Holtus Managing Director J.A. van Vugt Certification Manager

Aulugh

© 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-product-safety.com Company registration 09085396

ADDENDUM

Belonging to certificate: 2113863CE01

CE MARKING OF CONFORMITY MEDICAL DEVICES

Multipurpose Solutions and Cleaning Solutions for Contact Lens Care

Issued to:

Oté Pharma Sol B.V.

Vluchtoord 38 5406 XP Uden The Netherlands

This certificate covers the following product(s):

Variants:

- 001.xx All-in-one-Solution
- 002.xx Neutralisation Solution
- 003.xx Hydrogen Peroxide Solution
- 004.xx Preserved Buffered Isotonic Sodium Chloride Solution
- 005.xx Buffered Isotonic Sodium Chloride Solution
- 006.xx Storage, Wetting and disinfecting Solution
- 008.xx Isotonic Sodium Chloride 0,9% Solution
- 009.xx All-in-one-Solution for hard gas permeable lenses
- 010.xx Cleaner
- 011.xx Cleaner with polishing agent
- 014.xx Multifunctional solution
- 016.xx Multifunctional solution HydroShield
- 017.xx Neutralisation Tablets
- 020.xx All-in-one solution UG
- 025.xx All-in-one solution soft B5
- 028.xx Multifunctional solution US
- 029.xx Cleaner intensive
- 031.xx Monosept
- 043.xx All-in-one-Solution for hard gas permeable lenses B5
- 045.xx Multi Sol-comfort

Initial date: 10 July 2008 Revision date: 16 May 2021 DEKRA Certification B.V.

B.T.M. Holtus Managing Director J.A. van Vugt Certification Manager

Aulligh

© 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-product-safety.com Company registration 09085396