EC CERTIFICATE

Number: 2154875CE01

Full Quality Assurance System

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

Manufacturer:

Gynemed GmbH & Co. KG

Lubecker Straße 9
23738 Lensahn
Germany

For the product category(ies)

Media for the washing, cryopreservation and in vitro culture of human embryos, ova and spermatozoa for use in In Vitro Fertilization (IVF) / IntraCytoplasmatic Sperm Injection (ICSI), Assisted Reproductive Techniques (ART) and Embryo Transfer (ET).

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 2154875CN, initially dated 27 November 2012
Addendum, initially dated 27 November 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 October 2023
Issued for the first time: 27 November 2012
Reissued: 12 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: 2154875CE01

CE MARKING OF CONFORMITY
MEDICAL DEVICES

Media for the washing, cryopreservation and in vitro culture of human embryos, ova and spermatozoa for use in In Vitro Fertilization (IVF) / IntraCytoplasmatic Sperm Injection (ICSI), Assisted Reproductive Techniques (ART) and Embryo Transfer (ET).

Issued to:

Gynemed GmbH & Co. KG
Lubecker Stra?e 9
23738 Lensahn
Germany

This certificate covers the following product(s):

Media with gentamicin (Class III)
Media with human albumin (Class III)
Media with human albumin and gentamicin (Class III)
Media with specific medium supplement (Class III)

Initial date: 27 November 2012
Revision date: 13 October 2016

DEKRA Certification B.V.

B.T.M. Holtus
Managing Director

J.A. van Vugt
Certification Manager

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