## EC DESIGN-EXAMINATION CERTIFICATE

Number: 2154875DE02

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

(Devices in Class III)

Manufacturer:

Gynemed GmbH & Co. KG

Lubecker Straße 9 23738 Lensahn Germany

For the product

GM501 CULT - Cell culture media for the in vitro culture of human embryos and embryo transfer

Documents, that form the basis of this certificate

Certification Notice 2154875CN, initially dated 27 November 2012 CE Marking of Conformity 2154875CE01 Addendum, initially dated 2 March 2016

DEKRA hereby declares that the design of the product(s) falling within the product category mentioned above, 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, based on an examination in accordance with Annex II (4) of this Directive. The manufacturer has implemented a quality assurance system for the above mentioned product category in accordance to the provisions of Annex II (4) of Council Directive 93/42/EEC of June 14, 1993 and is subject to periodical surveillance.

The necessary information and the reference to the relevant documentation, of the products concerned and the examinations and 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: 2 March 2016
Reissued: 12 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: 2154875DE02

## EC DESIGN-EXAMINATION MEDICAL DEVICES

GM501 CULT - Cell culture media for the in vitro culture of human embryos and embryo transfer

Issued to:

Gynemed GmbH & Co. KG Lubecker Straße 9

Lubecker Straße 9 23738 Lensahn Germany

This certificate covers the following product(s):

Devices:

GM501 Cult GM501 Cult with Gentamicin GM501 Cult with Gentamicin and Phenolred II

Initial date: 2 March 2016

DEKRA Certification B.V.

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

Helligh

© 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