EC DESIGN-EXAMINATION CERTIFICATE

Number: 2154875DE04

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

Manufacturer:

Gynemed GmbH & Co. KG

Lübecker Straße 9
23738 Lensahn
Germany

For the product

Cell culture media with specific supplement for use in IVF, ICSI or similar procedures for ART

Documents, that form the basis of this certificate:

Certification Notice 2154875CNCN, initially dated 27 November 2012
CE Marking of Conformity 2154875CE01
Addendum, initially dated 13 October 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 2018
Issued for the first time: 13 October 2016

DEKRA Certification B.V.

drs. G.J. Zoetbrood
Managing Director

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

ADDENDUM

Belonging to certificate: 2154875DE04

EC DESIGN-EXAMINATION
MEDICAL DEVICES

Cell culture media with specific supplement for use in IVF, ICSI or similar procedures for ART

Issued to:

Gynemed GmbH & Co. KG
Lübecker Straße 9
23738 Lensahn
Germany

This certificate covers the following product(s):

Product name	**GM501 Flush**
Product code	4 GM 501F-50

Initial date: 13 October 2016

DEKRA Certification B.V.

drs. G.J. Zoetbrood
Managing Director

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