EC Certificate Full Quality Assurance System: Certificate BE99/50313

The management system of

Gynétics Medical Products N.V.

Rembert Dodoensstraat 51
3920 Lommel, Belgium

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

**Intra-uterine devices containing copper. Follicle aspiration needles.
Endometrial suction curette (curettage/sampling).
Embryo and oöcytes pipettes for transfer and manipulation.
Medium and medium sets for semen separation.**

For placing on the market of Class III devices covered by this certificate, an EC Design Examination Certificate according to Annex II (Section 4) is required.

This certificate is valid from 10 April 2014 until 30 January 2019 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 30 December 2016.
Issue 15. Certified since 30 January 1999.

Certification is based on reports numbered BE/AND 09307.

Authorised by

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