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

• Sterile FAS: sterile follicle aspiration needles
• Sterile embryo and oöcytes handling / manipulation pipettes (transfer /manipulation)

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 24 July 2018 until 30 January 2022
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 30 December 2019
Issue 19. Certified since 30 January 1999

Certification is based on reports numbered BE/AND 09307

Authorised by

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