



Product Service

EC Certificate

Full Quality Assurance System

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

No. G1 18 04 70354 009

Manufacturer: Labotect Labor-Technik-Göttingen GmbH

Kampweg 12
37124 Rosdorf
GERMANY



Facility(ies):

Labotect Labor-Technik-Göttingen GmbH
Kampweg 12, 37124 Rosdorf, GERMANY

BeLoTec GmbH
Kampweg 12, 37124 Rosdorf, GERMANY

Product

Category(ies):

Sterile medical products and medical products for assisted reproduction, consisting of biopsy needles and sets for follicle aspiration, suction pumps, cell and tissue culture systems

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

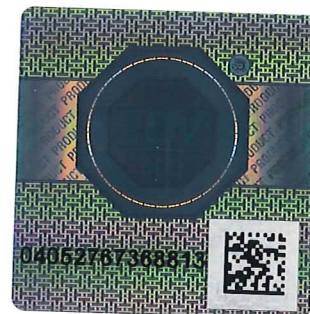
Report No.: 713132064

Valid from: 2018-07-11

Valid until: 2023-07-10

Date, 2018-07-11

Stefan Preis



TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

Page 1 of 1