

3EC International a.s., Hraničná 18, 821 05 Bratislava, Slovakia Notified Body No. 2265

EC CERTIFICATE

No. 2021-MDD/QS-067

issued in compliance with the Council Directive 93/42/EEC as amended, certifies that the medical device of Class I sterile,

Micromanipulation tools for assisted reproduction HOLDING, ICSI, HATCHING, BIOPSY, DENUDING

manufactured by company

Microtech IVF s.r.o. Seifertova 801/64, Lesná, 638 00 Brno, Czech Republic

is manufactured under conditions fulfilling the quality system requirements of Annex V of the Directive 93/42/EEC as amended.

The Notified Body No. 2265 has performed an audit of the above device quality system – restricted to aspects of manufacture concerned with securing and maintaining sterile conditions. The quality system has been assessed, approved and is subject to continuous surveillance according to Annex V, Section 4, of the Directive 93/42/EEC as amended. The detailed description of the system, requirements and measures applied by the manufacturer are presented in the Audit Report No. ICT_217 and the Final protocol No. 310549/2021.

This certificate is issued under the following conditions:

It applies only to the quality system maintained in the manufacture of the above referenced model of medical device and it does not substitute the design or type-examination procedures, if requested. The certificate remains valid until the manufacturing conditions or the quality system are changed but until May 26th, 2024 at the latest. The certificate validity is conditional upon positive results of regular surveillance audits and fulfillment of relevant legal and other requirements by manufacturer.



Dr. Katarina Tomin Srdošová Responsible to act on behalf of NB 2265



At Bratislava, on May 25th, 2021
This certificate supersedes the EC Certificate No. 2016-MDD/QS-025 issued on October 26th, 2016

Certificate history:

Revision	Date of issue	Application for Conformity Assessment of MD number	Description
0	May 25 th , 2021	310549	Recertification