Declaration of conformity

Manufacturer: Jiangsu Kangjin Medical Instrument Co., Ltd.
Zhenglu Town
213111 Changzhou
PEOPLE'S REPUBLIC OF CHINA
European Representative: Shanghai International Holding Corp. GmbH (Europe)
Eiffestrasse 80 20537 Hamburg GERMANY

Product Name: Biopsy Forceps for Single Use
Model Number: FB-ATS, FB-ATG, FB-AES,FB-AEG,
FB-BTS, FB-BTG, FB-BES, FB-BEG
FB-CTS, FB-CTG, FB-CES, FB-CEG
FB-DTS, FB-DTG, FB-DES, FB-DEG
UMDNS CODE: 16268
Classification (MDD, Annex IX): II a, Rule 6
Conformity Assessment Route: MDD Annex Ⅱ excluding (4)

We herewith declare that the above mentioned products meet the transposition into national law, the provisions of the following EC Council Directives and all applicable harmonized Standards. All supporting documentations are retained under the premises of the manufacturer.
Jiangsu Kangjin Medical Instrument Co., Ltd is exclusively responsible for DOC.

DIRECTIVES

General applicable directives:
Medical Device Directive: COUNCIL DIRECTIVE 93/42/EEC of 14 June 1993 concerning medical devices (MDD 93/42/EEC). Amended by DIRECTIVE 2007/47/EC of 5 September 2007

Notified Body: TÜV SÜD Product Service GmbH, Ridlerstr.65, 80339 München, Germany NB
NB Identification NO.: 0123
(EC)Certificate(s): G1 039452 0033 Rev.01
Expire date of the Certificate: 2024-05-26
S**tart of CE Marking:** Aug.20th,2006
Place, Date of Issue: Kangjin, 2021.01.06

Signature: JIANGSU KANGJIN MEDICAL INSTRUMENT CO., LTD.
Name:
Position: President