



3EC International a.s., Hraničná 18, 821 05 Bratislava, Slovak Republic
Notified body No. 2265

EU QUALITY MANAGEMENT SYSTEM CERTIFICATE No. 2022-MDR/QS-010

Healthium Medtech Limited

Registered Place of Business: No. 472 D, 13th Cross, 4th Phase, Peenya Industrial Area,
Bangalore, Karnataka – 560 058, India
Manufacturing Site 01: No. 472 D, 13th Cross, 4th Phase, Peenya Industrial Area, Bangalore,
Karnataka – 560 058, India
Manufacturing Site 02: Plot No: 1605, Portia Road, Sri City, Tirupati District,
Andhra Pradesh – 517 646, India
SRN No.: IN-MF-000008421

Name and address of the Authorized representative:

MED DEVICES LIFESCIENCES B.V., Keizersgracht 482, 1017 EG Amsterdam, The Netherlands

This EU Quality Management System Certificate issued in accordance with the Regulation (EU) 2017/745 of the European Parliament and of the Council on medical devices as amended confirms, that quality management system of medical device:

**MONOFILAMENT POLYAMIDE (NYLON)
STERILE NON-ABSORBABLE SURGICAL SUTURE**

(for detailed list refer to Annex I)

Intended purpose: Annex II

MD class III

meets the requirements on quality management system according to the Chapter I Annex IX of the Regulation (EU) 2017/745 of the European Parliament and of the Council on medical devices as amended.

Conditions for or limitations to the validity of the certificate: N/A

Validity of the certificate is conditional upon positive results of regular surveillance audits.

Notified body No. 2265 has performed assessment of the quality management system of the abovementioned medical device and found that it meets the requirements stated above. The outcome of the assessment of the quality management system of the abovementioned medical device is stated in the MD Technical Documentation Assessment Report No. MDR025_2021 from August 4, 2022, MD Clinical Evaluation Report No. MDR025_2021 from August 4, 2022 and MD Audit Report No. SK-0643-22 from August 4, 2022. Information on all examinations and tests performed is stated in the abovementioned reports and is available on request.

This **EU Quality Management System Certificate** applies only to the quality management system of the abovementioned medical device. For the placing on the market of the MDs which this certificate covers, the EU Technical Documentation Assessment Certificate issued in accordance with the Regulation (EU) 2017/745 on medical devices as amended is required. The certificate validity is conditional upon fulfilment of relevant legal requirements by the manufacturer.



Valid from: **October 18, 2022**
Valid until: **August 15, 2027**
First issue: **August 15, 2022**
Revision: **01**
History: **Annex III**



3EC International a.s.
Katarína Tomin Srdošová, PhD.
Director of NB2265

In Bratislava, Slovakia, October 18, 2022