EU Quality Management System Certificate PK25/00000133

The management system of

Seemann Technologies

500M DASKA ROAD, GHUINKI STOP 51310, Sialkot, Pakistan

SRN Number: PK-MF-000022513

has been assessed and certified as meeting the requirements of

MDR (EU) 2017/745 Quality Management System certificate (Annex IX Chapter I and III)

For the following products

The Scope of Registration appears on page 2 of this certificate

This certificate is valid from 12 August 2025 until 12 August 2030 and remains valid subject to satisfactory surveillance audits.
Recertification audit due before 12 February 2030

Issue 1. Certified since 12 August 2025

Authorised by
Virginie Siloret
Global Medical Device
Certification Manager
SGS Belgium NV NB1639
SGS House Noorderlaan 87 2030 Antwerp Belgium
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Seemann Technologies

MDR (EU) 2017/745 Quality Management System certificate (Annex IX Chapter I and III)

Class Is device
MDA0312, MDS1005
Sterile Single Use Electrosurgical Monopolar and Bipolar Connecting Cables
Basic UDI: 896110280TF07-0169

Class IIb device
MDA0312, MDS1005, EMDN: K02010106

Sterile/Non-sterile Single Use Bipolar Forceps Hand-piece with/without Cable, with/without NONSTICK, Irrigation and Suction Feature
Intended Use: Electrosurgical bipolar forceps, with accessories, used in Electrosurgery for cutting and coagulating.
Basic UDI: 896110280TF01-014X

Sterile/Non-sterile Single Use Monopolar Electrosurgical Electrodes
Active/Neutral for open, minimal and laparoscopic invasive surgery with/without Handpiece Cautery Pencils
Intended use: Electrosurgical Single Use Electrodes used in Electrosurgery for cutting and coagulating, with accessories, and intended to remove tissue and control bleeding.
Basic UDI: 896110280TF05-015T

Conditions for & limitation to the validity of the certificate:
For placing on the market of Class III or class IIb implantable devices (except sutures, staples, dental fillings, dental braces, tooth crowns, screws, wedges, plates, wires, pins, clips and connectors and Annex VIII rule 12 devices) covered by this certificate, a Technical Documentation Assessment Certificate according to Annex IX section 4 and 5 is required.

For Class I devices, audit done by SGS Belgium N.V. is limited to the specific aspect described in Article 52 section 7 of MDR (EU) 2017/745 (sterility, reusability or measurement function).

List of examinations and tests performed, which may include reference to relevant CS and harmonised standards, as per Annex XII, Chapter II, section 10 is available “on request" per email to NB1639@sgs.com.

Limitation: N/A

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Certification is based on following reports: - PK/SKT/22415 - S2A 1.0 + TFR 1.3 + TFR 1.4
Authorized representative name and address (if relevant): European Device Solutions (UK) Limited & European Healthcare and device solutions (Ireland) Ltd.,
Stratton House, Bishopstown Road, Cork, T12 Y9TC, Ireland

Previous certificate number: N/A

Change in between this certificate and previous one: N/A

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