EU REP CERTIFICATE

Authorized Representative
Article 11 MDR 2017/745
No. AMS/CE/2022/3072016DE
SRN DE-AR-000008078

Manufacturer: SUMA MEDICAL DEVICES
24/617 Potters Street, Dharowal
Sialkot-51310, Pakistan

HAS GIVEN A MANDATE TO

EU-Representative: ANTEX MEDI SOLUTION
HANS BUNTE STRASSE 6
69123 HEIDELBERG GERMANY

TO ACT AS
EU REPRECENTATIVE FOR

Medical Device Products CLASS 1 LISTED IN ATTACHED ANNEX I

The certificate remains valid until the expiration agreement of EU REP, manufacturing conditions, the quality system or relevant legislation are changed. The validity is conditioned by positive results of periodic surveillance audits.

The product liability rests with the manufacturer in accordance with applicable directive and standard, after fulfilling of the relevant EU legislation requirements, the manufacturer shall affix relevant CE marking to all below mentioned models of the medical device. The product classes covered are in ATTACHED ANNEX I.

Date of Initial Registration: 17-07-2020
Certificate Renewal: 17-07-2022
Certificate Valid Until: 16-07-2023

EC REP ASSOCIATION ERA

Authorized Signatory
ANTEX MEDI SOLUTION

CE

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ANNEX I PAGE 1/1:

List of Medical devices under the scope of the agreement

SURGICAL & DENTAL INSTRUMENTS

Scissors
Forceps
Mouth Mirrors
Needle Holders
Ear Speculas
Excavators
Curettes
Nose Speculas
Scalpels
Scalpel Handles
Scalers
Bone Rasps
Excavators
Buck Hammers

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