EU Certificate

for the assessment of the quality management system

according to Medical Device Regulation (EU) 2017/745 Annex IX Chapter I+III

As a Notified Body of the European Union DEKRA Certification GmbH certifies, that the manufacturer

Stihler Electronic GmbH

Single Registration Number (SRN): DE-MF-000006188
Gaussstrasse 4, 70771 Leinfelden-Echterdingen, Germany

applies a quality management system according to Annex IX Chapter I+III of the Medical Device Regulation (EU) 2017/745 for the medical devices listed in the annex. This certificate is based on the assessments listed in CNo50192-00 and is only valid in conjunction with the successful completion of the annual surveillance audits.

EU Certificate no.: 50192-60-00-01

Certificate valid from: 2025-06-20
Certificate valid to: 2028-07-23

Digitally signed by Markus RAINER Kopf
Date: 2025-06-20 12:34:44+02:00

DEKRA Certification GmbH, Stuttgart
Notified Body ID number: 0124

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
BS-MDR-092

Annex to the EU Certificate no. 50192-60-00-01

Following devices/device categories are included in this certificate:

Class IIa

ASTOTUBE®

- Article number: M77432002
  M77449004
  M77449003
  M77455502
  M77460002
- Basic-UDI-DI: 426011414TUBE5P
- EMDN Code: A030201
- MDN Code 1202

Change(s) to previous certificate:
Correction of MDN Code