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

DeltaMed GmbH

Single Registration Number (SRN): DE-MF-000007882
Raiffeisenstraße 8a, 61169 Friedberg, 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 CNo50245-01 and is only valid in conjunction with the successful completion of the annual surveillance audits.

EU Certificate no.: 50245-60-02-00

Certificate valid from: 2025-03-04
Certificate valid to: 2030-03-03

Previous certificate no. 50245-60-01, issued on 2023-02-23

Digitally signed by Markus RAINER Kopf
Date: 2025-02-20 18:31:01+01:00

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

DEKRA Certification GmbH, Stuttgart
Notified Body ID number: 0124