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

Renfert GmbH

Single Registration Number (SRN): DE-MF-000005448
Untere Gießwiesen 2, 78247 Hilzingen, 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 CNo50910-00.

EU Certificate no.: 50910-60-00

Certificate valid from: 2023-02-08
Certificate valid to: 2027-02-07

DEKRA Certification GmbH • D-70565 Stuttgart, Handwerkstraße 15 • DEKRA

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

Karin Leicht
DEKRA Certification GmbH, Stuttgart, 2023-02-08
Notified Body ID number: 0124

Annex to the EU Certificate no. 50910-60-00

valid from 2023-02-08 to 2027-02-07

Revision status of the annex: 0 dated 2023-02-08

Following devices/device categories are included in this certificate:

Class IIa

- Strengthening net and strengthening grid for dental prothestic dentures

Karin Leicht
DEKRA Certification GmbH, Stuttgart, 2023-02-08
Notified Body ID-number: 0124