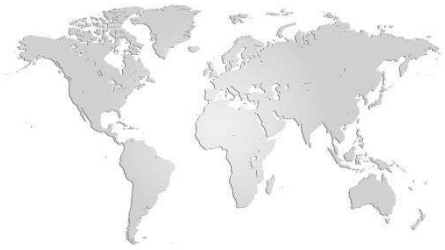
EU Certificate

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

VITA Zahnfabrik H. Rauter GmbH & Co. KG

Single Registration Number (SRN): DE-MF-000005906
Spitalgasse 3, 79713 Bad Säckingen, 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 CNo50069-00 and is only valid in conjunction with the successful completion of the annual surveillance audits.

EU Certificate no.: 50069-60-02-00

Certificate valid from: 2025-05-20
Certificate valid to: 2026-10-11

Previous certificate no. 50069-60-01, issued on 2023-08-29

Digitally signed by Markus RAINER Kopf
Date: 2025-05-20 16:21:46+02:00

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