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 connection with the successful performance of the annual surveillance audits.

EU Certificate no.: 50069-60-01

Certificate valid from: 2023-08-29
Certificate valid to: 2026-10-11

Previous certificate no. 50069-60-00, dated 2022-10-05
Change(s) to previous certificate: change of certificate structure, addition of product VITA VIONIC DENT DISC multiColor, cl. IIa (change notification no. 50069-CN23-01)

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, 2023-08-29
Notified Body ID number: 0124