

# 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)



*K. Leicht*

Karin Leicht

DEKRA Certification GmbH, Stuttgart, 2023-08-29

Notified Body ID number: 0124



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