EU Certificate

for the assessment of the quality management system

according to Medical Device Regulation (EU) 2017/745, Annex IX Chapter 1

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 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.

EU Certificate no.: 50069-60-00

Certificate valid from: 2022-10-07 Certificate valid to: 2026-10-11

DEKRA

Natascha Jezyschek DEKRA Certification GmbH Stuttgart; 2022-10-07 Notified Body ID number: 0124

