

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

KaVo Dental GmbH

Single Registration Number (SRN): DE-MF-000006471

Bismarckring 39, 88400 Biberach, 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 CNo50512-00 and is only valid in conjunction with the successful completion of the annual surveillance audits.

EU Certificate no.: 51512-60-00-00

Certificate valid from:

2024-05-24

Certificate valid to:

2026-09-28



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

BS-MDR-092

DEKRA Certification GmbH, Stuttgart 2024-05-24
Notified Body ID number: 0124

Annex to the EU Certificate no. 51512-60-00-00

Following device categories are included in this certificate:

Class IIa

MDA 0311

Basic UDI-DI

Dental Treatment Units	++EKAVG001YA
Dental Intraoral Cameras and Diagnostic Devices	++EKAVG513ZA
Dental Multifunctional Handpieces	++EKAVG101YF
Dental straight and contra-angle Handpieces Straight and contra-angle Handpieces Air motor instruments	++EKAVG502Z5 ++EKAVG507ZF
Dental Turbines	++EKAVG501Z3
Dental Scaler Air-powered	++EKAVG503Z7
Dental Powder Jet Handpieces	++EKAVG505ZB
Dental Motors Air motors Low-voltage motors	++EKAVG508ZH ++EKAVG510Z4
Dental treatment and surgical Devices Treatment Devices Surgical Devices	++EKAVG511Z6 ++EKAVG512Z8