EU Declaration of Conformity
according to Article 19
Medical Device Regulation 2017/745/EU

Manufacturer

KaVo Dental GmbH
Bismarckring 39
88400 Biberach
Germany
www.kavo.com

EUDAMED SRN DE-MF-000006471

Product / REF KaVo Lumina 2.021.5000

Basic UDI-DI ++EKAVG200YJ

Classification Class I, Rule 13

Intended use of the product (s)
KaVo treatment lights are used to illuminate a patient's oral cavity during dental treatment. The lights are mounted on a KaVo device system (dental treatment unit) or on the KaVo support system (Centro column).
For detailed description of product and accessories see instructions for use.

EU Marking in accordance with
Regulation on medical devices (MDR) 2017/745/EU

Common Specifications
Currently not available

Statement
We declare under our sole responsibility that the products manufactured by us to which this declaration relates conform to the essential safety and performance requirements in accordance with the provisions of the regulation above and their applicable annexes.

This declaration is supported by the certificate with registration no. 51512-21-01 for the implementation and application of a QM system according to EN ISO 13485:2016+AC:2018+A11:2021.

Validity
Issued on 2025-04-02
Valid until 2027-09-28

Klaus Reisenauer
Senior Director Regulatory Affairs
Quality Assurance