

**EU Declaration of Conformity**  
**according to Annex IV**  
Medical Device Regulation 2017/745/EU

**KAVO**

**Manufacturer**

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

**EUDAMED SRN**

DE-MF-000006471

**Product / REF**

EXPERTsurg LUX / 1.008.3500  
MASTERsurg LUX Wireless / 1.009.1200  
SM5 / 1.011.4900

**Basic UDI-DI**

++EKAVG512Z8

**Classification**

Class IIa, Rule 9

**Intended use of the product (s)**

This KaVo product is intended only for use in the field of dentistry, for surgery to expose and dissect oral tissue structures or endodontic treatments (e.g. periodontal gap, gingiva, bone, jaw, extractions, implantations) and must be used by expert medical staff only. Any other type of use is not permitted.  
*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. 51512-60-00-01 according to the conformity assessment procedure of regulation 2017/745/EU, Annex IX.

**Notified Body**

2017/745/EU

**DEKRA Certification GmbH**

Handwerkstraße 15  
70565 Stuttgart

**0124**

**Validity**

Issued on

2025-03-31

Valid until

2026-09-28

  
Klaus Reisenauer  
Senior Director Regulatory Affairs  
Quality Assurance