DECLARATION OF CONFORMITY

EC COUNCIL DIRECTIVE 93/42/EEC FOR MEDICAL DEVICES

Manufacturer: **LM-Instruments Oy**
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Herewith we declare that the periodontal, cavity and root canal preparation instruments that we are producing as well as LM Implant Misura instruments, LM-EndoMax endodontic hand files, LM-ErgoHold3 and extraction instruments, comply with the relevant regulations of the guideline 93/42/EEC for medical devices as they have been modified with the directive 2007/47/EC. The products are classified in Medical Devices, Class I.

Date, place: 25.5.2021, Parainen

Kari Lehtonen
Quality Manager
LM-Instruments Oy