

Certificate

We hereby certify the company

Robert Hofmann GmbH
An der Zeil 6
96215 Lichtenfels
Germany

the introduction and application of a

Quality management system according to EN ISO 13485

in the scope

Project management and production of individual and series parts for medical devices by means of 3D plastic and metal printing as well as machining processes and plastic injection moulding processes

An audit by mdc has proven that this quality management system meets the requirements of the following standard:

EN ISO 13485:2016 + AC:2018 + A11:2021 - ISO 13485:2016
Medical devices – Quality management systems – Requirements for regulatory purposes

Valid from 2025-05-25
Valid until 2028-05-24

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Berlin, 2025-05-16


Certification Body