

# EU Quality Management System Certificate

We hereby certify the company

**MEIKO Maschinenbau GmbH & Co. KG**  
**Englerstraße 3**  
**77652 Offenburg**  
**Germany**

the introduction and application of a quality management system in accordance with Annex IX, Chapter I and III of Regulation (EU) 2017/745 for conformity assessment.

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

## **Annex IX – Chapter I (Quality Management System)**

of Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices.

Surveillance is carried out in accordance with Annex IX, Section 3 of Regulation (EU) 2017/745.

This certificate from mdc medical device certification GmbH (Notified Body 0483) consists of 2 pages. Details about the devices covered as well as further information and conditions are contained on the following pages.

Valid from 2024-02-16  
Valid until 2028-04-12

Registration No. D1182200009  
Report No. P23-01230-290972

Stuttgart, 2024-02-16



Notified Body



## Devices:

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MEIKO TOPIC 10  
MEIKO TOPIC 20  
MEIKO TOPIC 40

Risk class: IIa

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MEIKO TopLine 10  
MEIKO TopLine 20  
MEIKO TopLine 30  
MEIKO TopLine 40

Risk class: IIa

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MEIKO TopLine 10 Type M2  
MEIKO TopLine 20 Type M2  
MEIKO TopLine 30 Type M2  
MEIKO TopLine 40 Type M2

Risk class: IIa

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## The certificate is based on the previous certificate

D1182200008 dated 2023-08-21 with the following changes:

Restricted by "MEIKO TopLine 40E"