

EU Quality Management System Certificate

We hereby certify the company

Mytronic GmbH
Riedelbach – Technologiepark
In den Ensterwiesen 7
61276 Weilrod
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 2024-03-22
Valid until 2028-06-22

Registration No. D1407200010
Report No. P23-01421-282437

Stuttgart, 2024-03-22



Notified Body



Devices:

MC 3FP/3FP-S Syringe

Risk class: IIa
