

Certificate

We hereby certify the company

Mascia Brunelli S.p.A.
Viale Monza 272
20128 Milano
Italy

the introduction and application of a

Quality management system according to EN ISO 13485

in the scope

Design, manufacturing and distribution of haemostatic medical devices, in-vitro diagnostic microbiological culture media, platelet aggregation reagents and rapid tests for the detection of infectious diseases. Distribution of medical devices and in-vitro diagnostic devices.

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 2024-03-07
Valid until 2027-03-06

Registration No. D1016000055
Report No. P22-01687-252240

Stuttgart, 2024-03-07



Certification Body

