TÜVUSA

Certificate

acc. to ISO 13485:2016

Medical Devices - Quality Management Systems - Requirements for Regulatory Purposes

Certificate Registration No.: 19-1620-Q

The Certification Body TUV USA, Inc. hereby confirms as a result of the audit, assessment, and certification decision according to ISO/IEC 17021-1:2015, that the organization's quality management system is in conformance with ISO 13485:2016 for Medical Devices-Quality Management Systems-Requirements for Regulatory Purposes.

MTD Micro Molding
15 Trolley Crossing Road
Charlton, Massachusetts 01507, USA

Additional sites covered by QM System: N/A

Scope:

Contract Manufacturer of Plastic Injection Molded Medical Device Components that have Microscopic Features or are Microscopic in Size.
Assembly, packaging and labelling of Non-Active Tubing Instrument and IVD Instrument for Blood Coagulation Diagnostics.
Design and Development of Injection Mold for Customer Specific Medical Products.

The validity of this certification document can be obtained by contacting the TUV USA, Inc. Office.
TUV USA, Inc. (a Member of the TÜV NORD Group)
215 Main Street, Salem, New Hampshire 03079, USA
Tel: 001-603-870-8023, Fax: 001-603-870-8026, Email: medical-usa@tuv-nord.com

ANAB
ANSI National Accreditation Board
ACCREDITED
ISO/IEC 17021-1
MANAGEMENT SYSTEMS CERTIFICATION BODY

MEMBER OF MULTILATERAL RECOGNITION ARRANGEMENT
IAF

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Valid Until:
2025-02-27

Bradley Chen
Vice President- Medical, Americas
Medical Products Division
TUV USA, Inc.