

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:

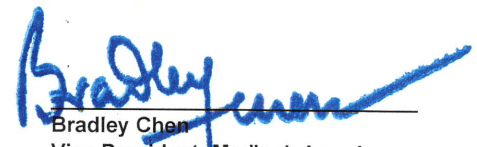
**Contract Manufacturer of Plastic Injection Molded Medical Device Components
that have Microscopic Features or are Microscopic in Size.
Assembly, packaging and labelling of Non-Active Tubing Instrument and IVD
Instrument for Blood Coagulation Diagnostics.
Design and Development of Injection Mold for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

Audit Report Reference No.: **21-3955 SA2 SPA**
Initial Certification Date: **2019-02-28**
Current Cycle Start Date: **2022-02-28**

Effective Date:
2024-03-27 / ed. 6

Valid Until:
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