Certificate

acc. to ISO 13485:2016

Medical Devices - Quality Management Systems - Requirements for Regulatory Purposes

Certificate Registration No.: 19-1626-Q

TUV USA, Inc. hereby certifies that the quality management system of the company mentioned below is in conformance with ISO 13485:2016 for Medical Devices-Quality Management Systems-Requirements for Regulatory Purposes.

Wise Plastics Technologies
3810 Stern Avenue
St. Charles, IL 60174
USA

Additional sites covered by QM System: See Annex I

Scope:

The Manufacture and Value Added Process Design of Engineered Injection Molded Products and Assemblies

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, Suite 1, Salem, NH 03079, USA
Tel: 001-603-870-8023, Fax: 001-603-870-8026, Email: medical-usa@tuv-nord.com

Audit Report Reference No.:19-8008 RC
Certificate Initial Issue Date: 13-MAY-2019
Current Cycle Start Date: 13-MAY-2019
Certificate Revised Date: 13-DEC-2019

Effective Date:
13-DEC-2019 / ed. 2

Valid Until:
12-MAY-2022

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

Certificate Registration No.: **19-1626-Q / ed. 2**
Company Name: ***Wise Plastics Technologies***
Central Office Address: ***3810 Stern Ave, St. Charles, IL 60174, USA***

Additional Site(s). covered by the QM System:

Location	Scope of Certification
Wise Plastics Technologies **1601 West Hawthorne Ln** **West Chicago, IL 60185** **USA**	The Manufacture and Value Added Process Design of Engineered Injection Molded Products and Assemblies

---End of list---

The validity of this certification document can be obtained by contacting the TUV USA, Inc. Office.

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Bradley Chen
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