

Certificate

acc. to ISO 13485:2016

Medical Devices - Quality Management Systems - Requirements for Regulatory Purposes

Certificate Registration No.: 18-1617-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.

MAJnMAR Medical Products 203-1130 Austin Avenue Coquitlam, BC V3K 3P5, Canada

Additional sites covered by QM System: See Annex 1

Scope:

Design, Development and Manufacture of Disposable and Reusable Trocars, Laparoscopic Instruments, Surgical Devices, Surgical Staplers, Silicone Drainage System, Sponge Laparotomy, Gauze, EVA Parenteral Nutrition Infusion Sets

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

2021-10-30 / ed. 4



Audit Report Reference No.: 20-3894 RC-SA3
Certificate Initial Issue Date: 2018-10-30
Current Cycle Start Date: 2021-10-30

Certificate Revised Date: 2021-10-27
Effective Date:

Valid Until: 2024-10-29

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

Annex 1, page 1 of 1

(Annex 1 MUST be displayed with the main certificate)

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Company Name:

MAJnMAR Medical Products

Central Office Address:

203-1130 Austin Avenue, Coquitlam, BC V3K 3P5, Canada



Additional Site(s) covered by the QM System:

Location

Scope of Certification

Headquarters

MAJnMAR Medical Products 203-1130 Austin Avenue

Coquitlam, BC V3K 3P5, Canada

Design and development, purchasing,

storage

Site 01

MAJnMAR Medical Products EN193-163 Schoolhouse Street Coguitlam, BC V3K 4X8, Canada Storage

---End of list---

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