

Certificate

acc. to ISO 13485:2016

Medical Devices - Quality Management Systems - Requirements for Regulatory Purposes

Certificate Registration No.: 18-1623-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.

ViaMed Holdings, LLC 40 Norfolk Ave. South Easton, MA 02375, USA

Additional sites covered by QM System: See Annex 1

Scope:

Manufacture of Precision Components and Assemblies per Customer Specification for Medical Industries

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.: 20-3906 RC Certificate Initial Issue Date: 2015-12-04 Current Cycle Start Date: 2021-11-03 Certificate Revised Date: 2021-11-03

Effective Date: 2021-12-04 / ed. 4

Valid Until: 2024-12-03

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)

Certificate Registration No.: 18-1623-Q / ed. 4

Certificate Registration Name:

ViaMed Holdings, LLC

Central Office Address:

40 Norfolk Ave., South Easton, MA 02375, USA



Additional Site(s) covered by the QM System:

Location

ViaMed Holdings, LLC 40 Norfolk Ave. South Easton, MA 02375, USA

ViaMed Holdings, LLC 8 Norfolk Ave. South Easton, MA 02375, USA

Scope of Certification

Manufacture of Precision Components and Assemblies per Customer Specification for Medical Industries

Manufacture of Precision Components and Assemblies per Customer Specification for Medical Industries

---End of list---

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