



Certificate

acc. to **ISO 13485:2016**

Medical Devices - Quality Management Systems - Requirements for Regulatory Purposes

Certificate Registration No.: 19-1618-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.

**Hanson Medical, Inc.
25960 Ohio Ave, Ste 200
Kingston, WA 98346, USA**

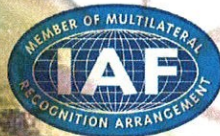
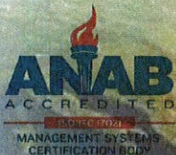
Additional sites covered by QM System: See Annex I

Scope:

Design, Development, Manufacture, and Distribution of Silicone Facial and Body Implants, Silicone Devices, Accessories, and Topical Scar Gel 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, 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.: **19-8029 SA2-UP**
Certificate Initial Issue Date: **09-JUL-2017**
Current Cycle Start Date: **09-JUL-2017**
Certificate Revised Date: **12-DEC-2019**

Effective Date:
12-DEC-2019 / ed. 3

Valid Until:
08-JUL-2020

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

Annex 1, page 1 of 1

(Annex 1 must be displayed with the main certificate)

Certificate Registration No.: 19-1618-Q / ed. 3

Company Name: Hanson Medical, Inc.

Central Office Address: 25960 Ohio Ave, Ste 200, Kingston, WA 98346, USA



Additional Site(s) covered by the QM System:

Location

Scope of Certification

Hanson Medical, Inc.
825 Riverside Ave, Ste 2
Paso Robles, CA 93446
USA

Design, Development, Production,
Quality Control, Document Control,
Training, Management, Warehouse,
and Shipping

---End of list---

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

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