Certificate

acc. to ISO 13485:2016

Medical Devices - Quality Management Systems - Requirements for Regulatory Purposes

Certificate Registration No.: 18-1614-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.

Veridiam, Inc.
1717 N. Cuyamaca St.
El Cajon, CA 92020
USA

Additional sites covered by QM System: None

Scope:

Contract Manufacturing of Components, Cannula, Tubes, and Assemblies for Medical and Dental Devices

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
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SCC Accredited CB-MS OCSM Accrédité CCN

Audit Report Reference No.: 18-8018 SA1-UP
Current Cycle Start Date: 21-MAY-2017
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Effective Date:
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Valid Until:
20-MAY-2020

Bradley Chen
Director, Medical Products Division
TUV USA, Inc.

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