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: N/A

Scope:

Contract Manufacturing of Components, Cannula, Tubes and Assemblies for use in 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)
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Valid Until:
20-MAY-2023

Bradley Chen
Vice President of Medical, Americas
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TUV USA, Inc.