

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)
215 Main Street, Suite 1, Salem, NH 03079, USA

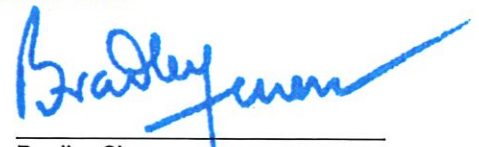
Tel: 001-603-870-8023, Fax: 001-603-870-8026, Email: medical-usa@tuv-nord.com



Audit Report Reference No.: **19-3712 RC**
Certificate Initial Issue Date: **21-MAY-2017**
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Valid Until:
20-MAY-2023



Bradley Chen
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TUV USA, Inc.