

Certificate

acc. to **ISO 13485:2016**

Medical Devices - Quality Management Systems - Requirements for Regulatory Purposes

Certificate Registration No.: **21-1603-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.

Inovo, Inc.
401 Leonard Blvd. North
Lehigh Acres, FL 33971, USA

Additional sites covered by QM System: **N/A**

Scope:

Design and Development, Production, Distribution, Final Testing, and Servicing of Respiratory 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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Audit Report Reference No.: **21-3935 RC**

Certificate Initial Issue Date: **2021-03-26**

Current Cycle Start Date: **2021-06-22**

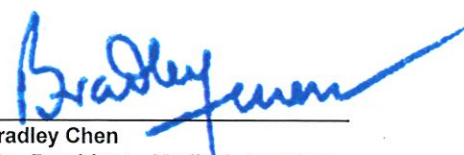
Certificate Revised Date: **2021-06-17**

Effective Date:

2021-06-22 / ed. 2

Valid Until:

2024-06-21



Bradley Chen
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Medical Products Division
TUV USA, Inc.