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CERTIFICATE OF REGISTRATION

This is to certify that the management system of:

Allied Healthcare Products, Inc.

Main Site: 1720 Sublette Avenue, St. Louis, Missouri 63110 United States (FIN F001542)

Additional Site: 46 New Street, Stuyvesant Falls, New York 12174 United States (FIN F002306)

has been registered by Intertek, an MDSAP recognized auditing organization, as conforming to the requirements of:

ISO 13485:2016

Australia: Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 1 (excluding Part 1.6)

Canada: Medical Devices Regulations – Part 1- SOR 98/282

United States: 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 (Subparts A to D)

Japan: MHLW Ministerial Ordinance 169, Article 4 to Article 68; PMD Act

The management system is applicable to:

Design, Manufacture and Final Testing for; Cannulas, Oxygen Masks, Tubing, Humidifiers, Nebulizers, Water Traps, Circumcision Clamps, CO2 Absorbents, Adapters, Cylinder Post Valves, Flowmeters, Medical Gas Hoses, Oxygen Regulators, Oxygen Timers, Selector Valves and Selector Valve Kits, Tubing and Accessory Dryers, Breast Pumps, Portable Suction Pumps and Aspirators, Vacuum Collection Bottles and Canisters, Vacuum Regulators, Backboards, Burn Sheets, Cinch Collars, Half Back Vests, Trauma Pants, Bag Mask Resuscitators, CPR-Timers, Demand Valves, Resuscitators, Ventilators, and Accessories of Emergency Medical Products and Respiratory Therapy Products. The servicing of all non-disposable devices.

Additional site: Additional Site: Manufacturing of CO2 absorbents.

Certificate Number:
0092750

Initial Certification Date:
2019-07-11

Certification Effective Date:
2019-07-11

Certification Expiry Date:
2022-07-10

ISO 13485:2016 CERTIFICATION ™

MDSAP MEDICAL DEVICE SINGLE AUDIT PROGRAM

Intertek

Calin Moldovean
President, Business Assurance

Intertek Testing Services NA, Inc.
900 Chelmsford Street
Lowell, MA, USA 01851

In the issuance of this certificate, Intertek assumes no liability to any party other than to the Client, and then only in accordance with the agreed upon Certification Agreement. This certificate's validity is subject to the organization maintaining their system in accordance with Intertek's requirements for systems certification. Validity may be confirmed via email at certificate.validation@intertek.com or by scanning the code to the right with a smartphone. The certificate remains the property of Intertek, to whom it must be returned upon request. Validity of this certificate may be verified at http://www.intertek.com/business-assurance/certificate-validation/