

CERTIFICATE OF REGISTRATION

This is to certify that the management system of:

Parker Hannifin Corp. -Precision Fluidics Division

(FIN F000918)

Main Site: 245 Township Line Road

Hatfield, Pennsylvania, 19440, United States

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, distribution, installation and serving of nitrous oxide - oxygen conscious sedation flow meters, gas scavenging systems and gas distribution products for use by a physician, dentist or licensed healthcare professionals.

Certificate Number: 0093668

Revision Level: 03

Initial Certification Date: 2019-08-12

Certification Effective Date: 2025-08-11

Certification Expiry Date: 2028-08-11



intertek

Rathin Grover President, Business Assurance

Intertek Testing Services NA, Inc. 4700 Broadmoor SE, Suite 200 Kentwood, MI, USA, 49512



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/ CT-MDSAP-2016-NA-EN-LT-P-16Jan.25

