By Royal Charter

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

TIDI Products LLC
570 Enterprise Drive
Neenah
Wisconsin
54956
USA

Facility ID Number: F003249

Holds Certificate No: **MDSAP 703786**

The company listed on this certificate has been audited to and found to conform with ISO 13485:2016 including the following country specific requirements:

Australia: Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 1 (excluding Part 1.6) - Full Quality Assurance Procedure
Brazil: RDC ANVISA n. 67/2009, RDC ANVISA n. 665/2022 - Good Manufacturing Practices, RDC ANVISA n. 551/2021
Canada: Medical Devices Regulations - Part 1 - SOR 98/282
Japan: MHLW MO No 169 (2004), as amended by MHLW MO No 60 (2021), PMD Act
USA: 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 - Subparts A to D

Please see scope page.

For and on behalf of BSI:

Graeme Tunbridge, Senior Vice President Medical Devices

Original Registration Date: 2020-05-29 Effective Date: 2023-05-29 Expiry Date: 2026-05-28

MDSAP MEDICAL DEVICE SINGLE AUDIT PROGRAM
BSI Group America Inc. is an MDSAP recognised auditing organization

...making excellence a habit.™

Certificate No: **MDSAP 703786**

Registered Scope:

Design, manufacture, distribution, installation and servicing of sterile and non-sterile medical devices and personal protective equipment including scalpels, wound dressings and wound management, skin protection, patient monitors, protective covers and barriers, securement devices, urology bags, restraints and restraint alternatives, respiratory therapy, securement accessories for catheters, and fall prevention devices including hipsters, gait belts, cushions for use in hospitals, physical therapy and home.

Original Registration Date: 2020-05-29 Effective Date: 2023-05-29 Expiry Date: 2026-05-28