

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

Corporation STERIS Canada
(also operating as STERIS Canada ULC)
490 boulevard Armand-Paris
Québec (Québec)
Québec
G1C 8A3
Canada

Facility ID Number: F000511

Holds Certificate No:

MDSAP 687114

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. 16/2013, RDC ANVISA n. 23/2012, RDC ANVISA n. 67/2009

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

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

USA: 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 - Subparts A to D

Design and manufacture of washing disinfection and drying systems and associated accessories and manufacture of sterile processing equipment for healthcare and associated accessories.

For and on behalf of BSI:

Graeme Tunbridge, Senior Vice President Medical Devices

Original Registration Date: 2018-10-23

Effective Date: 2022-06-29

Expiry Date: 2024-10-22



BSI Group America Inc. is an MDSAP recognised auditing organization

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