By Royal Charter

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

Steris Corporation
St. Louis Operations
7501 Page Avenue
St. Louis
Missouri
63133
USA

Facility ID Number: F000423

Holds Certificate No: **MDSAP 687958**

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
Canada: Medical Devices Regulations - Part 1 - SOR 98/282
USA: 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 - Subparts A to D

The design, manufacture and distribution of healthcare personnel handwash/disinfection products, skin care products and instrument and hard surface cleaning/decontamination products.

For and on behalf of BSI:

Graeme Tunbridge, Senior Vice President Global Regulatory & Quality

Original Registration Date: 2018-11-21 Effective Date: 2024-06-04 Expiry Date: 2027-06-03

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

...making excellence a habit.™

Certificate No: **MDSAP 687958**

Location	Registered Activities
Steris Corporation 8525 Page Avenue St. Louis Missouri 63133 USA Facility ID Number: F000423	The storage and distribution of healthcare personnel handwash/disinfection products, skin care products, instrument and hard surface cleaning/decontamination products, and surgical light handles and camera covers.
Steris Corporation St. Louis Operations 7501 Page Avenue St. Louis Missouri 63133 USA Facility ID Number: F000423	The design, manufacture and distribution of healthcare personnel handwash/disinfection products, skin care products and instrument and hard surface cleaning/decontamination products.

Original Registration Date: 2018-11-21 Effective Date: 2024-06-04 Expiry Date: 2027-06-03