



Regulation (EU) 2017/745, Annex IX Chapter I and III

## MDR 721051 R000

Manufacturer: Swann-Morton Limited

**Address:**Owlerton Green
Sheffield
S6 2BJ
United Kingdom

Single Registration Number: GB-MF-000001890

EU Authorised Representative: Emergo Europe

Address:

Prinsessegracht 20 2514 AP The Hague The Netherlands

#### Scope: See attached Device Schedule

On the basis of our examination of the quality system in accordance with Regulation (EU) 2017/745, Annex IX Chapter I and III, the quality system meets the requirements of the Regulation. For the placing on the market of Class III and Class IIb implantable devices an Annex IX Chapter II certificate is required.

For and on behalf of BSI, a Notified Body for the above Regulation (Notified Body Number 2797):

Gary E Slack, Senior Vice President Medical Devices

Gary C Stade

First Issued: 2021-01-20

Date: 2021-11-23

Expiry Date: 2026-01-19

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Validity of this certificate is conditional on the Manufacturer's quality system being maintained to the requirements of the Regulation as demonstrated through the required surveillance activities of the Notified Body.

This certificate was issued electronically and is bound by the conditions of the contract.

NB Contact: BSI Group The Netherlands B.V., Say Building, John M. Keynesplein 9, 1066 EP, Amsterdam, Netherlands. Tel: + 31 (0) 20 346 07 80 Corporate Contact: BSI Group Assurance Limited, registered in England under number 05435540 at 389 Chiswick High Road, London, W4 4AL, UK. A Member of the BSI Group of Companies.





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#### Device Schedule: Class IIa, Custom-made and other devices

Device(s)	Risk Classification
Single use surgical scalpels and blades	Class IIa
Sterile suture remover	Class Is
Reusable instruments 'Orthopaedic Instruments'	Class Ir

For Class Is devices, the Notified Body conformity assessment is limited to the aspects relating to establishing, securing and maintaining sterile conditions.

For class Ir devices (class I re-usable surgical instruments), the Notified Body conformity assessment is limited to the aspects relating to the reuse of the device

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### **Certificate History**

(References to applicable Common Specifications, Harmonized Standards complied with, and the relevant test and audit reports that support any of the below certificate changes may be requested from Certificate. Verification@bsigroup.com)

Date	Reference number	Action
2021-01-20	3103832	Issued
Current	3539989	Supplemented - Addition of Class Ir devices  Amended - Removal of subcontractor Woodland Works  Amended - Addition of SRN code: GB-MF-000001890  Amended - Administrative update on activity for "gamma irradiation" to "Radiation (Gamma Sterilization)" for Swann-Morton (Services) Limited Penn Works and on history section for "First issue" to "Issued"

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## List of Critical Subcontractors and Crucial Suppliers

Recognised as being involved in services related to the products covered by:

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Date: 2021-11-23

United Kingdom

Critical Subcontractor/Crucial Supplier	Service(s) supplied
Andersen Caledonia Limited Caledonian House Phoenix Crescent Strathclyde Business Park Lanarkshire ML4 3NJ United Kingdom	ETO Sterilization
Jewel Blade Ltd 442 Penistone Road Sheffield S6 2FU United Kingdom	Crucial Supplier
Swann-Morton (Microbiological Laboratory Services) Limited Owlerton Green Sheffield S6 2BJ United Kingdom	Microbiology Service
Swann-Morton (Services) Limited Penn Works Owlerton Green Sheffield S6 2BJ	Radiation (Gamma Sterilization)

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