

Owlerton Green Sheffield, S6 2BJ

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## **Declaration of Conformity**

Manufacturer's Name:

Manufacturer's Address:

Single Registration Number:

**BUDI-DI** 

50339550NONSTSTDHANDLE42

Swann-Morton Limited

Sheffield, S6 2BJ, England GB-MF-000001890

Emergo Europe

Owlerton Green,

**European Authorised Representative Name: European Authorised Representative Address:** 

Westervoortsedijk 60 6827 AT Arnhem The Netherlands

Single Registration Number:

NL-AR-000000116

This Declaration of Conformity is issued under our sole responsibility as manufacturer of the devices covered by this declaration, Swann-Morton Limited, hereby ensure and declare that these devices meet the provisions of the medical devices regulations (EU) 2017/745.

The Notified Body used for our conformity assessment in accordance with Annex IV and Annex IX of the above Regulation is BSI NL (2797).

Certificates Issued:

## MDR 721051 R000 in respect of: Reusable Instruments

For Class 1r devices (class I re-usable surgical instruments), the Notified Body conformity assessment is limited to the aspects relating to the reuse of the device.

FM73368: Operates a Quality Management System which complies with the requirements of ISO 13485 for the following scope: The design, manufacture, packaging and distribution of surgical blades, disposable scalpels, handles and blade removers.

MDSAP 674417 – The company listed on this certificate has been audited to and found to conform with the following criteria: ISO 13485:2016, Australia – Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 1 (excluding Part 1.6) – Full Quality Assurance Procedure, Therapeutic Goods (Medical Devices) Regulations 2002, Schedule 3 Part 4 – Production Quality Assurance Procedure; Brazil – RDC ANVISA n.665/2022, RDC ANVISA n. 23/2012, RDC ANVISA n. 67/2009; Canada Medical Device Regulations – Part 1 – SOR 98/282; Japan – MHLW Ministerial Ordinance 169, Article 4 to 68, PMD Act AND USA – 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 – Subparts A to D. The design, manufacture, packaging and distribution of surgical blades, disposable scalpels, handles and blade removers.

## **Country Registrations:**

Canada Medical Device License: 72

U.S.A Establishment Registration & Device Listing (FDA) Registration No. 9611194 Owner/Operator No. 9003320.

Australian Register of Therapeutic Goods Certificate: 107772

Brazilian RDC number: 10302860262

Japan MHLW registration number: BG20500131

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Product Family:	NON-STERILE STAINLESS STANDARD METAL SURGICAL HANDLES		
Intended Use:	TO HOLD A No.3 or No.4 FITMENT SURGICAL BLADE		
Product Codes:	See Page 3		
Classification:	Class I (Reusable) (Annex VIII, Rule 6) (EU) Class I (MDR Schedule 1, Part 1, Rule 3 (Health Canada) Class I (FDA CFR 878.4800) (U.S.A – FDA) Class I (TG(MD)R 2002) Schedule 2 Part 3.2(4) (Australia) Class I (RDC Annex II, II, 1. Rule 1) (Brazil) Class I (JMDN: 12235000 Rule 6) (Japan)		
Standards Used:	See Table Below		
GMDN Code & Term	12235 Knife/Blade Handle A metal surgical instrument, e.g. stainless steel or brass, designed to mount a compatible blade used for cutting or dissecting tissue.		

## Standards applied in relation to this Declaration are:

STANDARD NUMBER	TITLE		
BS 2982	Materials and packaging of surgical scalpels with detachable blades		
BS EN 27740/ISO 7740	Instruments for surgery, scalpels with detachable blades, fitting dimensions		
BS EN ISO 15223-1	Medical devices – Symbols to be used with medical device labels, labelling & information to be supplied		
BS EN ISO 13485	Medical devices – Quality management systems – Requirements for regulatory purposes		
BS EN ISO 14971	Medical devices – Application of risk management to medical devices		
BS EN ISO 17664 – 1	Processing of healthcare products, information to be provided by the medical device manufacturer for the processing of medical devices		
BS EN ISO 20417	Medical Devices – Information to be supplied by the manufacturer		

PATTERN	PRODUCT CODE	UDI
No. 5B	0905	05033955009052
No. 6B	0906	05033955009069
No. 7	0907	05033955009076
No. 9	0909	05033955009090
No. 3L	0913	05033955009137
No. 4L	0914	05033955009144
No. 3 Graduated	0933	05033955009335
No. 4 Graduated	0934	05033955009342
В3	0923	05033955009236
	No. 5B  No. 6B  No. 7  No. 9  No. 3L  No. 4L  No. 3 Graduated  No. 4 Graduated	No. 5B       0905         No. 6B       0906         No. 7       0907         No. 9       0909         No. 3L       0913         No. 4L       0914         No. 3 Graduated       0933         No. 4 Graduated       0934

Signed for and on behalf of Swann-Morton Limited, Owlerton Green, Sheffield S6 2BJ

SIGNATURE	Rele
PRINT FULL NAME	Darren Hall
POSITION	QA/RA Systems Manager
PLACE & DATE	Swann-Morton Ltd, Sheffield S6 2BJ, England 11th May 2023