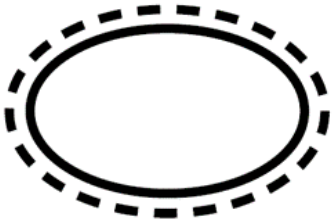
Reason for change:	MR1260 First issue.

Symbol	Standard	Ref #	Title	Description
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.1.1	Manufacturer	Indicates the medical device manufacturer
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.1.2	Authorized representative in the European Community/ European Union	Indicates the authorized representative in the European Community/ European Union
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.1.3	Date of manufacture	Indicates the date when the medical device was manufactured
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	Permitted by standard	Country of manufacture	Used in place of 5.1.3 and will have date of manufacture nearby. Indicates that the product was manufactured in Great Britain (also known as the UK or United Kingdom of Great Britain and Northern Ireland)

	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.1.4	Use-by date	Indicates the date after which the medical device is not to be used
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.1.5	Batch code	Indicates the manufacturer's batch code so that the batch or lot can be identified
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.1.6	Catalogue number	Indicates the manufacturer's catalogue number so that the medical device can be identified.
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.1.7	Serial number	Indicates the manufacturer's serial number so that a specific medical device can be identified

	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.1.8	Importer	Indicates the entity importing the medical device into the locale
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.1.9	Distributor	Indicates the entity distributing the medical device into the locale
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.2.3	Sterilized using ethylene oxide	Indicates a medical device that has been sterilized using ethylene oxide.
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.2.7	Non-sterile	Indicates a medical device that has not been subjected to a sterilization process

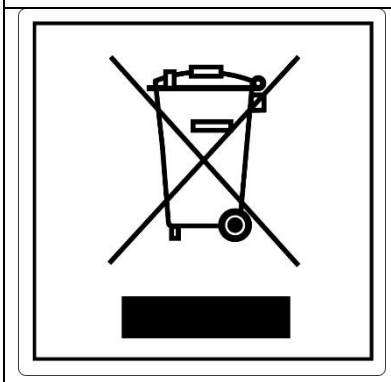
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.2.8	Do not use if package is damaged and consult instructions for use	Indicates a medical device that should not be used if the package has been damaged or opened and that the user should consult the instructions for use for additional information.
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.2.14	Single sterile barrier system with protective packaging outside	Indicates a single sterile barrier system with protective packaging outside
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	Permitted by standard		Combines 5.2.3 and 5.2.14 (As permitted by ISO 15223-1) to indicate that the device is sterilized by ethylene oxide and has a single sterile barrier with protective packaging outside

	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.3.4	Keep dry	Indicates a medical device that needs to be protected from moisture. **NOTE**: This symbol indicates that the packaging must be kept dry to maintain the integrity of the packaging material. The device itself is designed for use in wet or fluid-filled environments and is not affected by moisture.
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.3.7	Temperature limit	Indicates the temperature limits to which the medical device can be safely exposed.
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.3.8	Humidity limitation	Indicates the range of humidity to which the medical device can be safely exposed.

	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.4.2	Do not re-use/ Single use only	Indicates a medical device that is intended for one single use only.
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.4.3	Consult instructions for use	Indicates the need for the user to consult the instructions for use.
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.4.4	Caution	Indicates the need for the user to consult the instructions for use for important cautionary information such as warnings and precautions that cannot, for a variety of reasons, be presented on the medical device itself.

	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.7.7	Medical Device	Indicates the item is a medical device.
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	5.7.10	Unique Device Identifier	Indicates a carrier that contains Unique Device Identifier information.
	ISO 7010-M002, IEC 60601-1, Table D.2, Symbol 10		Follow instructions for use or follow electronic instructions for use.	Follow instructions for use.

CE 2797	MDD 93/42/EEC MDR 2017/745 Regulation (EC) 765/2008	Annex XII Article 20 Annex II	CE marking, may include Notified Body Reference no. 2797	Signifies European technical conformity.
UK CA 0086				Signifies UKCA technical conformity
Rx only	21 CFR 801.15 21 CFR 801.109	(c) (1) (i) (F) (b) (1)	Prescription only	Caution: Federal (US) law restricts this device to sale by or on the order of a physician.
	ISO 15223-1 Medical devices — Symbols to be used with information to be supplied by the manufacturer	A.16	Consult instructions for use or consult electronic instructions for use	

GTIN				Global Trade Item Number
	Directive 2002/96/ EC (repealed). Replaced by DIRECTIVE 2012/19/EU which does NOT contain this symbol.		Waste stream disposal status	Do not dispose of electronic products in the general waste stream
	DIRECTIVE 2012/19/ EU (WEEE)	Annex IX	Collect separately	indicating separate collection for EEE
SGS C US 800526				SGS North America Certification Mark