SYMBOL	SYMBOL TITLE	EXPLANATORY TEXT	STANDARD REFERENCE	REFERENCE NUMBER
	Manufacturer	Indicates the medical device manufacturer	ISO 15223-1, Clause 5.1.1	5.1.1
	Date of manufacture	Indicates the date when the medical device was manufactured	ISO 15223-1, Clause 5.1.3	5.1.3
	Use-by date	Indicates the date after which the medical device is not to be used	ISO 15223-1, Clause 5.1.4	5.1.4
	Batch code	Indicates the manufacturer's batch code so that the batch or lot can be identified. Synonyms for “batch code” are “lot number”, “lot code” and “batch number”.	ISO 15223-1, Clause 5.1.5	5.1.5
	Catalogue number	Indicates the manufacturer's catalog number so that the medical device can be identified	ISO 15223-1, Clause 5.1.6	5.1.6
	Sterile	Indicates a medical device that has been subjected to a sterilization process	ISO 15223-1, Clause 5.2.1	5.2.1
	Sterilized using irradiation	Indicates a medical device that has been sterilized using irradiation.	ISO 15223-1, Clause 5.2.3	5.2.3
	Consult instructions for use or consult electronic instructions	Indicates the need for the user to consult the instructions for use	ISO 15223-1, Clause 5.4.3	5.4.3
	Do not re-use	Indicates a medical device that is intended for one single use only	ISO 15223-1, Clause 5.4.2	5.4.2
	Do not resterilize	Indicates a medical device that is not to be resterilized	ISO 15223-1, Clause 5.2.6	5.2.6
	Prescription Use Only	Caution: Federal law (USA) restricts this device to sale by or on the order of a licensed healthcare practitioner.	21 CFR 801.15(c)(1)(i)F	21 CFR 801.15(c)(1)(i)F
	Not made with natural rubber latex	Not made with natural rubber latex.	ISO 15223-1:2012 (Annex B, 15223-1:2012)	Annex B
	Unique device identifier	Indicates a carrier that contains unique device identifier information.	ISO 15223-1 Clause 5.7.10	5.7.10
	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.	ISO 15223-1, Clause 5.2.8	5.2.8
	Caution: Read all warnings and precautions in instructions for use	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, Clause 5.4.4	5.4.4
	Sterilized using ethylene oxide	Indicates a medical device that has been sterilized using ethylene oxide	ISO 15223-1, Clause 5.2.3	5.2.3
	Single sterile barrier system with protective packaging inside	Indicates a single sterile barrier system with protective packaging inside	ISO 15223-1, Clause 5.2.13	5.2.13
	Non-sterile	Indicates a medical device that has not been subjected to a sterilization process	ISO 15223-1, Clause 5.2.7	5.2.7

* Reference: ISO 15223-1:2021 Medical Devices - Symbols to be used with information to be supplied by the manufacturer - Part 1: General Requirements