














SYMBOL	TITLE	DESCRIPTION	STANDARD
	Medical device	Indicates the item is a medical device.	ISO/DIS 15223-1:2020(E) DRAFT 5.7.7
	Do not re-use	Indicates a medical device that is intended for one use, or for use on a single patient during a single procedure.	ISO 15223-1 5.4.2
	Non-sterile	Indicates a medical device that has not been subjected to a sterilization process.	ISO 15223-1 5.2.7
	Not made with natural rubber latex	Indicates a medical device that is not made with natural rubber latex.	ISO 15223-1:2012 Annex B
	Manufacturer	Indicates the medical device manufacturer.	ISO 15223-1 5.1.1
	Date of manufacture	Indicates the date when the medical device was manufactured.	ISO 15223-1 5.1.3
	Use-by date	Indicates the date after which the medical device is not to be used.	ISO 15223-1 5.1.4
	Temperature limit	Indicates the temperature limits to which the medical device can be safely exposed.	ISO 15223-1 5.3.7
	Do not use if packaging is damaged	Indicates a medical device should not be used if the package has been opened or damaged.	ISO 15223-1 5.2.8
	Catalog/model number	Indicates the manufacturer's catalog number so that the medical device can be identified.	ISO 15223-1 5.1.6
	Batch/lot code	Indicates the manufacturer's batch code so that the batch or lot can be identified.	ISO 15223-1 5.1.5
	Sterilized using ethylene oxide	Indicates a medical device that has been sterilized using ethylene oxide.	ISO 15223-1 5.2.3
	Consult instructions for use	Indicates the need for the user to consult the instructions for use.	ISO 15223-1 5.4.3