
















Symbol Glossary

Symbol	Symbol Ref. No.	Symbol Title	Symbol Description	Standard
	5.1.1	Manufacturer	Indicates the medical device manufacturer, as defined in EU Directives 90/385/EEC, 93/42/EEC and 98/79/EC.	ISO 15223-1:2016 Medical Devices – Symbols to Be Used with Medical Device Labels, Labeling, and Information to be Supplied
	5.1.2	Authorized Representative in the European Community	Indicates the Authorized Representative in the European Community.	ISO 15223-1:2016 Medical Devices – Symbols to Be Used with Medical Device Labels, Labeling, and Information to be Supplied
	5.1.4	Use-by Date	Indicates the date after which the medical device is not to be used.	ISO 15223-1:2016 Medical Devices – Symbols to Be Used with Medical Device Labels, Labeling, and Information to be Supplied
	5.1.5	Batch Code	Indicates the manufacturer's batch code so that the batch or lot can be identified NOTE: Synonyms for "batch code" are "lot number" and "batch number"	ISO 15223-1:2016 Medical Devices – Symbols to Be Used with Medical Device Labels, Labeling, and Information to be Supplied
	5.1.6	Catalogue Number	Indicates the manufacturer's catalog number so that the medical device can be identified NOTE: Synonyms for "catalogue number" are "reference number" and "reorder number"	ISO 15223-1:2016 Medical Devices – Symbols to Be Used with Medical Device Labels, Labeling, and Information to be Supplied
	5.2.3	Sterilized Using Ethylene Oxide	Indicates a medical device that has been sterilized using ethylene oxide.	ISO 15223-1:2016 Medical Devices – Symbols to Be Used with Medical Device Labels, Labeling, and Information to be Supplied
	5.2.4	Sterilized Using Irradiation	Indicates a medical device that has been sterilized using irradiation.	ISO 15223-1:2016 Medical Devices – Symbols to Be Used with Medical Device Labels, Labeling, and Information to be Supplied
	5.2.6	Do Not Resterilize	Indicates a medical device that is not to be resterilized.	ISO 15223-1:2016 Medical Devices – Symbols to Be Used with Medical Device Labels, Labeling, and Information to be Supplied
	5.2.7	Non-Sterile	Indicates a medical device that has not been subjected to a sterilization process.	ISO 15223-1:2016 Medical Devices – Symbols to Be Used with Medical Device Labels, Labeling, and Information to be Supplied
	5.2.8	Do Not Use if Package is Damaged	Indicates a medical device that should not be used if the package has been damaged or opened.	ISO 15223-1:2016 Medical Devices – Symbols to Be Used with Medical Device Labels, Labeling, and Information to be Supplied
	5.4.2	Do Not Reuse	Indicates a medical device that is intended for one use, or for use on a single patient during a single procedure.	ISO 15223-1:2016 Medical Devices – Symbols to Be Used with Medical Device Labels, Labeling, and Information to be Supplied

Symbol	Symbol Ref. No.	Symbol Title	Symbol Description	Standard
	5.4.3	Consult Instructions for Use	Indicates the need for the user to consult the instructions for use.	ISO 15223-1:2016 Medical Devices – Symbols to Be Used with Medical Device Labels, Labeling, and Information to be Supplied
	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 15223-1:2016 Medical Devices – Symbols to Be Used with Medical Device Labels, Labeling, and Information to be Supplied
	3.1.11	MR Conditional	An item with demonstrated safety in the MR environment within defined conditions.	ASTM F2503-13 Standard Practice for Marking Medical Devices and Other Items for Safety in the Magnetic Resonance Environment
	2725	Contains or presence of	On medical devices: to indicate that the equipment contains the identified product or substance.	ISO 7000:2014 Graphical symbols for use on equipment -- Registered symbols

Symbol	Symbol Title	Symbol Description	Additional Information
R_x Only	Rx only	Federal (USA) law restricts this device to sale by or on the order of a physician. An alternative to the text "Rx Only."	21 CFR 801.109 Prescription devices
CE	CE Marking	The product conforms to European Medical Directive 93/42/EEC and meets applicable health, safety and environmental requirements. If the mark is accompanied by a number, conformity is verified by the indicated notified body.	Medical Device Directive 93/42/EEC
QTY	Quantity	Indicates unit quantity	Symbol used by A&E Medical Corporation

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