


















Graphical Symbols for Medical Device Labeling

Symbol	Title of Symbol	Description of Symbol	Standard/Source	Reference Number
	Manufacturer	Indicates the medical device manufacturer, as defined in EU Directives 90/385/EEC, 93/42/EEC and 98/79/EC.	ISO 15223-1:2021	5.1.1
	Authorized representative in the European Community	Indicates the authorized representative in the European Community.	ISO 15223-1:2021	5.1.2
	Date of manufacture	Indicates the date when the medical device was manufactured.	ISO 15223-1:2021	5.1.3
	Use-by date	Indicates the date after which the medical device is not to be used.	ISO 15223-1:2021	5.1.4
	Batch code	Indicates the manufacturer's batch code so that the batch or lot can be identified.	ISO 15223-1:2021	5.1.5
	Catalogue number	Indicates the manufacturer's catalogue number so that the medical device can be identified.	ISO 15223-1:2021	5.1.6
	Sterilized using irradiation	Indicates a medical device that has been sterilized using irradiation.	ISO 15223-1:2021	5.2.4
	Do not resterilize	Indicates a medical device that is not to be resterilized.	ISO 15223-1:2021	5.2.6
	Non-sterile	Indicates a medical device that has not been subjected to a sterilization process.	ISO 15223-1:2021	5.2.7
	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 15223-1:2021	5.2.8
	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 15223-1:2021	5.4.2
	Consult instructions for use	Indicates the need for the user to consult the instructions for use.	ISO 15223-1:2021	5.4.3
	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 15223-1:2021	5.4.4
	N/A	Caution: Federal (USA) law restricts this device to sale by or on the order of a physician.	21 CFR Part 801, Sec. 801.109	N/A
	Medical Device	Indicates the device is a medical device.	ISO 15223-1:2021	5.7.7

Symbol	Title of Symbol	Description of Symbol	Standard/Source	Reference Number
	Importer	Indicates the entity importing the medical device into the locale	ISO 15223-1:2021	5.1.8
	Authorized representative in Switzerland	Indicates the authorized representative in Switzerland.	N/A	N/A