

Symbol	Reference Number of Symbol	Title of Symbol	Description of Symbol
	5.1.1	Manufacturer	Indicates the medical device manufacturer.
	5.1.3	Date of manufacture	Indicates the date when the medical device was manufactured.
	5.1.11	Country of manufacture	To identify the country of manufacture of products.
	5.1.4	Use-by date	Indicates the date after which the medical device is not to be used.
	5.1.5	Batch Code	Indicates the manufacturer's batch code so that the batch or lot can be identified.
	5.1.6	Catalogue Number	Indicates the manufacturer's catalogue number so that the medical device can be identified.
	5.1.7	Serial number	Indicates the manufacturer's serial number so that a specific medical device can be identified.
	5.1.8	Importer	Indicates the entity importing the medical device into the locale.
	5.1.9	Distributor	Indicates the entity distributing the medical device into the locale.
	5.2.1	Sterile	Indicates a medical device that has been subjected to a sterilization process.
	5.2.3	Sterilized using ethylene oxide	Indicates a medical device that has been sterilized using ethylene oxide.
	5.2.4	Sterilized using irradiation	Indicates a medical device that has been sterilized using irradiation.
	5.2.6	Do not resterilize	Indicates a medical device that is not to be resterilized.
	5.2.7	Non-sterile	Indicates a medical device that has not been subjected to a sterilization process.
	5.2.8	Do not use if package is damaged and consult instructions for use	Indicates that 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.
	5.2.11	Single sterile barrier system	Indicates a single sterile barrier system.
	5.2.12	Double sterile barrier system	Indicates two sterile barrier systems.
	5.2.13	Single sterile barrier system with protective packaging inside	Indicates a single sterile barrier system with protective packaging inside.
	5.2.14	Single sterile barrier system with protective packaging outside	Indicates a single sterile barrier system with protective packaging outside.
	5.3.1	Fragile, handle with care	Indicates a medical device that can be broken or damaged if not handled carefully.
	5.3.2	Keep away from sunlight	Indicates a medical device that needs protection from light sources.

	5.3.4	Keep dry	Indicates a medical device that needs to be protected from moisture.
	5.3.7	Temperature limit	Indicates the temperature limits to which the medical device can be safely exposed.
	5.4.2	Do not re-use	Indicates a medical device that is intended for one single use only.
 eIFU Indicator	5.4.3	Consult instructions for use or consult electronic instructions for use	Indicates need for the user to consult the instructions for use.
	5.4.4	Caution	Indicates that caution is necessary when operating the device or control close to where the symbol is placed, or that the current situation needs operator awareness or operator action in order to avoid undesirable consequences.
	5.7.2	Patient Name	Indicates the name of the patient.
	5.7.3	Patient Identification	Indicates the identification data of the patient.
	5.7.4	Patient information website	Indicates a website where a patient can obtain additional information on the medical product.
	5.7.5	Health care centre or doctor	Indicates the address of the health care centre or doctor where medical information about the patient may be found.
	5.7.6	Date	Indicates the date that information was entered or a medical procedure took place.
	5.7.7	Medical device	Indicates the item is a medical device.
	5.7.10	Unique device identifier	Indicates a carrier that contains unique device identifier information.
	ASTM F2503	MR Conditional	Medical device that has been demonstrated to pose no known hazards in a specified MR environment with specified conditions of use.
	21 CFR 801.109 (b) (1)	Rx Only	Federal Law Restricts this Device to Sale by or on the order of a Physician.
	5.4.8	Contains biological material of animal origin	Indicates a medical device that contains biological tissue, cells, or their derivatives, of animal origin.
	ISO 7000/IEC 60417 2725	Contains or presence of	To indicate that the equipment contains the identified product or substance.
	ASTM F2503	MR Safe	An item that poses no known hazards resulting from exposure to any MR environment.