






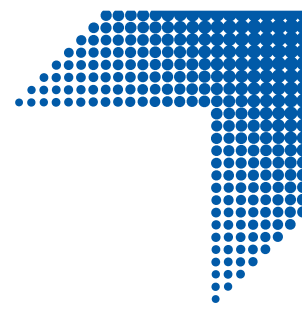








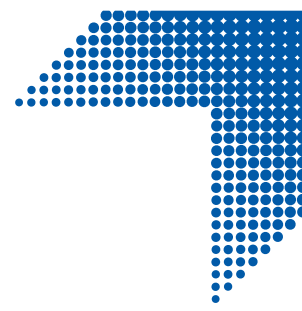



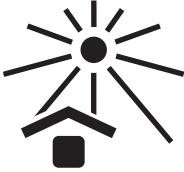





Referring to ISO 15223-1:2016 (E)

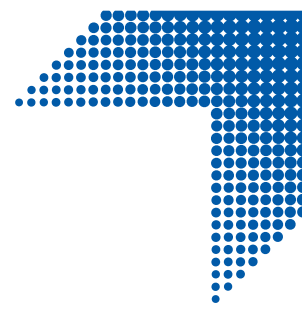
Symbol	Title of Symbol	Description of Symbol	Reg. No.	
MANUFACTURE		Manufacturer	Indicates the medical device manufacturer, as defined in EU Directives 90/385/EEC, 93/42/EEC and 98/79/EC.	3082
		Authorized representative in the European Community	Indicates the Authorized representative in the European Community.	
		Date of manufacture	Indicates the date when the medical device was manufactured.	2497
		Use-by date	Indicates the date after which the medical device is not to be used.	2607
		Batch code	Indicates the manufacturer's batch code so that the batch or lot can be identified.	2492
		Catalogue number	Indicates the manufacturer's catalogue number so that the medical device can be identified.	2493
		Serial number	Indicates the manufacturer's serial number so that a specific medical device can be identified.	2498







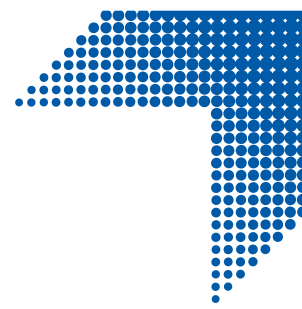
Symbol	Title of Symbol	Description of Symbol	Reg. No.	
STERILITY		Sterile	Indicates a medical device that has been subjected to a sterilization process.	2499
		Sterilized using aseptic processing techniques	Indicates a medical device that has been manufactured using accepted aseptic techniques.	2500
		Sterilized using ethylene oxide	Indicates a medical device that has been sterilized using ethylene oxide.	2501
		Sterilized using irradiation	Indicates a medical device that has been sterilized using irradiation.	2502
		Sterilized using steam or dry heat	Indicates a medical device that has been sterilized using steam or dry heat.	2503
		Do not resterilize	Indicates a medical device that is not to be resterilized.	2608
		Non-sterile	Indicates a medical device that has not been subjected to a sterilization process.	2609
		Do not use if package is damaged	Indicates a medical device that should not be used if the package has been damaged or opened.	2606






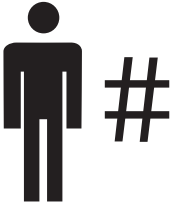

Symbol	Title of Symbol	Description of Symbol	Reg. No.	
STORAGE		Fragile, handle with care	Indicates a medical device that can be broken or damaged if not handled carefully.	0621
		Keep away from sunlight	Indicates a medical device that needs Pp protection from light sources.	0624
		Protect from heat and radioactive sources	Indicates a medical device that needs protection from heat and radioactive sources.	0615
		Keep dry	Indicates a medical device that needs to be protected from moisture.	0626
		Lower limit of temperature	Indicates the lower limit of temperature to which the medical device can be safely exposed.	0534
		Upper limit of temperature	Indicates the upper limit of temperature to which the medical device can be safely exposed.	0533
		Temperature limit	Indicates the temperature limits to which the medical device can be safely exposed.	0632



	Symbol	Title of Symbol	Description of Symbol	Reg. No.
SAFE USE		Biological risks	Indicates that there are potential biological risks associated with the medical device.	0659
		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.	1051
		Consult instructions for use	Indicates the need for the user to consult the instructions for use.	1641
		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.	0434



	Symbol	Title of Symbol	Description of Symbol	Reg. No.
IVD-SPECIFIC		In vitro diagnostic medical device	Indicates a medical device that is intended to be used as an in vitro diagnostic medical device.	
		Control	Indicates a control material that is intended to verify the performance characteristics of another medical device.	2494
		Negative control	Indicates a control material that is intended to verify the results in the expected negative range.	2495
		Positive control	Indicates a control material that is intended to verify the results in the expected positive range.	2496

OTHER		Patient number	Indicates a unique number associated with an individual patient.	2610
		Medical device	Indicates the item is a medical device.	