

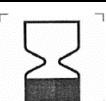



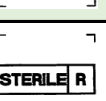
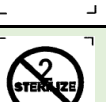
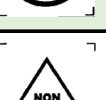








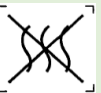



Symbole	Symbole Reference No	Title of Symbol	Description of Symbol
	5.1.1	Manufacturer	Indicates the medical device manufacturer, as defined in EU Directives 90/385/EEC, 93/42/EEC and 98/79/EC.
	5.1.3	Date of Manufacture	Indicates the date when the medical device was manufactured.
	5.1.4	Use by date	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	Catalog number	Indicates the manufacturer's catalog number so that the medical device can be identified.
	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 re-sterilize	Indicates a medical device that is not to be re-sterilized.
	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.	Indicates a medical device that should not be used if the package has been damaged or opened.
	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.
	5.4.2	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.
	5.2.3	Consult instructions for use	Consult instructions for use
	5.4.3	Consult instructions for use	Consult instructions for use
	5.4.4	Caution	Indicates that the instructions for use contain important cautionary information such as warnings and precautions that cannot, for a variety of reasons, be presented on the medical device itself.
	5.4.5	Contains or presence of natural rubber latex	Indicates the presence of natural rubber or dry natural rubber latex as a material of construction within the medical device or the packaging of a medical device which may cause allergic reactions.
	5.6.3	Non-pyrogenic	Indicates a medical device that is non-pyrogenic.
	---	Contains or presence of phthalate plasticizers DEHP	Indicates product that does contain the phthalate plasticizers DEHP.