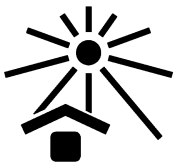
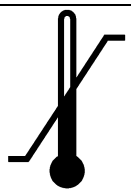
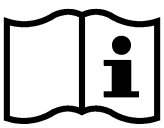
List of Applied Symbols

As of: 02/2020

Symbol	Reg.-Nr ISO 7000	Title	Description
	3082	Manufacturer	Indicates the medical device manufacturer.
	From ISO 15223-1	Authorized representative in the European Community	Indicates the Authorized representative in the European community.
	2497	Date of manufacture	Indicates the “date of manufacture “. The symbol shall be adjacent to the date that the product was manufactured, expressed as four digits for the year and two digits for the month and where appropriate, two digits for the day.
	2607	Use-by date	Indicates the date after which the medical device is not to be used.
	2492	Batch Code	Indicates the manufacturer´s batch code so that batch or lot can be identified.
	2493	Catalogue number	Indicates the manufacturer´s catalogue number so that the medical device can be identified.
	-	Medical Device	Indicates a Medical Device

Symbol	Reg.-Nr ISO 7000	Title	Description
	2501	Sterilized using ethylene oxide	Indicates the method of sterilization using ethylene oxide.
	2503	Sterilized using steam or dry heat	Indicates the method of sterilization using steam or dry heat
	2609	Non-sterile	Indicates that the device has not been sterilized.
	2606	Do not use if package is damaged	Indicates a medical device that should not be used if the package has been damaged or opened.
	0624	Keep away from sunlight	Indicates a medical device that needs protection from light sources.
	0626	Store dry	Indicates that the medical device should be kept dry.
	0632	Temperature limit	Indicates the temperature limits to which the medical device can be safely exposed.
	1051	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.
	1641	Consult instructions for use	Indicates the need for the user to consult the instructions for use.

Symbol	Reg.-Nr ISO 7000	Title	Description
	Derived from 2725	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
	From prEN 15986	Contains the presence of phthalates	Symbol for use in the labelling of medical devices - Requirements for labelling of medical devices containing phthalates
	From directive 93/42/EWG	CE Marking of Conformity	Indicates manufacturer declaration that the product complies with the essential requirements of the relevant European health, safety an environmental protection legislation.
	0434A	Caution	Indicates the need for the user to consult the instructions for use for important cautionary information such as warnings an precautions that cannot, for a variety of reasons, be presented on the medical device itself.