SYMBOLS GLOSSARY
ENGLISH

SYMBOL	SYMBOL TITLE	SYMBOL DESCRIPTION
	CATALOG NUMBER	Indicates the manufacturer's catalog number so that the medical device can be identified
	LOT NUMBER	Indicates the manufacturer's lot number so that the batch or lot can be identified
	USE BY (YYYY-MM-DD)	Indicates the date after which the medical device is not to be used
	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
	SINGLE USE ONLY	Indicates a medical device that is intended for one single use only
	DO NOT RESTERILIZE	Indicates a medical device that is not to be resterilized
	DO NOT USE IF PACKAGE IS DAMAGED	Indicates that a medical device should not be used if the packaging has been damaged or opened and that the user should consult the instructions for use for additional information
	PRODUCT QUANTITY	Indicates the quantity of product
	MEDICAL DEVICE	Indicates the item is a medical device
	PRESCRIPTION ONLY	Caution: Federal (USA) law restricts this device to sale by or on the order of a physician. Indicates the product is indicated for professional use only
	MR CONDITIONAL	Indicates a medical device that has defined conditions to enter safely into the MR environment
	MR UNSAFE	Indicates a medical device that has unacceptable risks and should not enter the MR environment
	SERIAL NUMBER	Indicates the manufacturer's serial number so that a specific medical device can be identified
	TEMPERATURE RANGE	Indicates the temperature limits to which the medical device can be safely exposed
	UPPER TEMPERATURE LIMIT	Indicates the upper limit of temperature to which the medical device can be safely exposed
	LOWER TEMPERATURE LIMIT	Indicates the lower limit of temperature to which the medical device can be safely exposed

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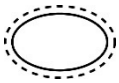
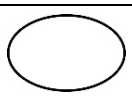
SYMBOL	SYMBOL TITLE	SYMBOL DESCRIPTION
	HUMIDITY RANGE	Indicates the range of humidity to which the medical device can be safely exposed
	ATMOSPHERIC PRESSURE RANGE	Indicates the range of atmospheric pressure to which the medical device can be safely exposed
	SINGLE PATIENT MULTIPLE USE	Indicates a medical device that may be used multiple times (in multiple procedures) on a single patient
	TYPE B APPLIED PART	Indicates an applied part complying with specific requirements for protection against electric shock
	WASTE ELECTRICAL AND ELECTRONIC EQUIPMENT (WEEE)	Indicates electronic equipment to be disposed of properly
	MANUFACTURER	Indicates the medical device manufacturer
	DATE OF MANUFACTURE (YYYY-MM-DD)	Indicates the date when the medical device was manufactured
	NON-STERILE	Indicates a medical device that has not been subjected to a sterilization process
	STERILIZED BY IRRADIATION	Indicates a medical device that has been sterilized using irradiation
	STERILIZED BY ETHYLENE OXIDE	Indicates a medical device that has been sterilized using ethylene oxide
	STERILIZED BY ASEPTIC PROCESSING	Indicates a medical device that has been manufactured using accepted aseptic techniques
	STERILIZED BY STEAM	Indicates a medical device that has been sterilized using steam or dry heat
	STERILIZED BY MULTIPLE METHODS	Indicates a medical device that has been sterilized using irradiation, ethylene oxide, aseptic techniques, and steam or dry heat
	STERILE BARRIER SYSTEM	Indicates a single sterile barrier system with protective packaging outside
	STERILE BARRIER SYSTEM	Indicates a single sterile barrier system
	AUTHORIZED REPRESENTATIVE IN THE EUROPEAN COMMUNITY	Indicates the authorized representative in the European Community/European Union
	AUTHORIZED REPRESENTATIVE IN SWITZERLAND	Indicates the authorized representative in Switzerland

SYMBOLS GLOSSARY
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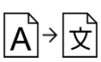
SYMBOL	SYMBOL TITLE	SYMBOL DESCRIPTION
	CONSULT INSTRUCTIONS FOR USE	Indicates the need for the user to consult the instructions for use/electronic IFU
	FOLLOW USER MANUAL	Indicates that the user manual must be read
	KEEP DRY	Indicates a medical device that needs to be protected from moisture
	KEEP AWAY FROM SUNLIGHT	Indicates a medical device that needs protection from light sources or heat
	FLAMMABLE	Indicates that the medical device contains materials that are highly flammable
	IONIC RADIATION	Indicates generally elevated, potentially hazardous, levels of non-ionizing radiation, or to indicate equipment or systems
	TYPE BF APPLIED PART	Indicates an applied part complying with a higher degree of protection against electric shock
	DISTRIBUTED BY	Indicates the entity distributing the medical device into the locale
	CONTAINS NO NRL	Indicates that no presence of dry natural rubber or natural rubber latex as a material of construction within the medical device or the packaging of a medical device
	LATEX-FREE	Indicates that there is no presence of natural rubber latex as a material of construction within the medical device or the packaging of a medical device
	CONTAINS BIO ANIMAL	Indicates a medical device that contains biological tissue, cells, or their derivatives, of animal origin
	CONTAINS BIO HUMAN	Indicates a medical device that contains biological tissue, cells, or their derivatives, of human origin
	CLASS II	Indicates Class II or double insulated equipment protection against electric shock
	DIRECT CURRENT	Indicates on the rating plate that the equipment is suitable for direct current only
	ALTERNATING CURRENT	Indicates on the rating plate that the equipment is suitable for alternating current only
	TRANSLATION	Indicates that the original medical device information has undergone a translation which supplements or replaces the original information. Translation: LOGOS - Via Curtatona 5/2 - 41126 Modena (Italy) | Taxpayer ID Number (TIN): IT02018930368 - REA Modena No. 259448