




















Symbol	Title of Symbol	Meaning of Symbol
	Batch code	Indicates the manufacturer's batch code so that the batch or lot can be identified.
	Catalogue number	Indicates the manufacturer's catalogue number so that the medical device can be identified.
	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.
	CE Certification mark	CE marking is the manufacturer's declaration that the product meets the requirements of the applicable EC legislation.
	Consult instructions for use	Indicates the need for the user to consult the instructions for use.
	Date of manufacture	To indicate the date on which a product was manufactured.
	Do not re-sterilize	Indicates a medical device that is not to be re-sterilized.
	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.
	Do not use if package is damaged	Indicates a medical device that should not be used if the package has been damaged or opened.
	European authorized representative	A European Authorised Representative (EAR.) serves as a legal entity designated by non European Union (EU) manufacturers, to represent them in the EU and ensure their compliance with the European Directives.
	Magnetic resonance conditional	A device or implant that may contain magnetic electrically conductive, or RF-reactive components that is safe for operations in proximity to the MRI, provided the conditions for safe operation are defined and observed.
	Magnetic resonance safe	The device or implant is completely non-magnetic, non-electrically conductive, and non-RF reactive, eliminating all of the primary potential threats during an MRI procedure.
	Manufacturer	Indicates the medical device manufacturer.
	Medical device	Indicates the item is a medical device.
	Non-sterile	Indicates a medical device that has not been subjected to a sterilization process.
	Rx only mark	U.S. federal law restricts this device to sale by or on the order of a dental professional.
	Sterilized using irradiation	Indicates a medical device that has been sterilized using irradiation.
Tooth#	Tooth number	Indicates tooth location.
	Unique device identifier	The UDI system is intended to provide a globally recognised identification system of medical devices for distribution and use.
	Use-by date	Indicates the date after which the medical device is not to be used.