








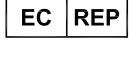










## Symbols Glossary

Symbol	Title and Meaning	Reference
 <small>www.acumed.net/ifu</small>	<b>Consult instructions for use</b> Indicates the need to consult the instructions for use (IFU). An electronic instructions for use (eIFU) indicator (website address) may accompany the symbol when used to indicate an instruction to consult an eIFU.	5.4.3**
	<b>Caution</b> 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.	5.4.4*
	<b>Use-by date</b> Indicates the date after which the medical device is not to be used.	5.1.4*
	<b>Do not use if package is damaged</b> Indicates a medical device that should not be used if the package has been damaged or opened.	5.2.8*
	<b>Upper limit of temperature</b> Indicates the upper limit of temperature to which the medical device can be safely exposed.	5.3.6*
	<b>Do not re-use</b> Indicates a medical device that is intended for one use, or for use on a single patient during a single procedure.	5.4.2*
	<b>Non-sterile</b> Indicates a medical device that has not been subjected to a sterilization process.	5.2.7*
	<b>Sterilized using ethylene oxide</b> Indicates a medical device that has been sterilized using ethylene oxide.	5.2.3*
	<b>Sterilized using irradiation</b> Indicates a medical device that has been sterilized using irradiation.	5.2.4*
	<b>Authorized representative in the European Community</b> Indicates the authorized representative in the European Community.	5.1.2*
	<b>Do not re-sterilize</b> Indicates a medical device that is not to be re-sterilized.	5.2.6*
	<b>Date of manufacture</b> Indicates the date when the medical device was manufactured.	5.1.3*
	<b>Catalog number</b> Indicates the manufacturer's catalogue number so that the medical device can be identified.	5.1.6*
	<b>Batch code</b> Indicates the manufacturer's batch code so that the batch or lot can be identified.	5.1.5*
	<b>Manufacturer</b> Indicates the medical device manufacturer, as defined by EU Directives 90/385/EEC, 93/42/EEC and 98/79/EC.	5.1.1*
	<b>MR Conditional</b> An item with demonstrated safety in the MR environment within defined conditions.	7.3.2***
	<b>MR Safe</b> An item that poses no known hazards resulting from exposure to any MR environment. MR Safe items are composed of materials that are electrically nonconductive, nonmetallic, and nonmagnetic.	7.3.1***
	<b>MR Unsafe</b> An item which poses unacceptable risks to the patient, medical staff or other persons within the MR environment.	7.3.3***

\* ISO 15223-1 2nd Ed. 2012-07-01, Medical Devices - Symbols to be used with medical devices labels, labeling, and information to be supplied - Part 1: General requirements

\*\* An eIFU indicator below symbol 5.4.3 indicates the website URL where instructions for use are made available in electronic format.

\*\*\* ASTM - F2503 International Standard Practice for Marking Medical Devices and Other Items for Safety in the Magnetic Resonance Environment