

Symbols Glossary for DePuy







This document contains symbols that may appear on the product packaging or labeling. It provides information on the symbol including the standards title, reference, and description, where available. This document does not contain company created symbols, pictograms, or graphical images.







Please read the full Instructions For Use for detailed information on proper use, indications, contraindications, warnings, precautions and adverse events. This document does not replace the Instructions For Use.







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ISO 15223-1 Medical devices — Symbols to be used with medical device labels, labelling and information to be supplied — Part 1: General requirements

Image	Accompanying Text	Reference	Description
	Catalogue Number	5.1.6	Indicates the manufacturer's catalogue number so that the medical device can be identified.
	Batch code	5.1.5	Indicates the manufacturer's batch code so that the batch or lot can be identified.
	Serial Number	5.1.7	Indicates the manufacturer's serial number so that a specific medical device can be identified.
	Caution	5.4.4	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.
	Consult Instructions for use	5.4.3	Indicates the need for the user to consult the instructions for use.
	Manufacturer	5.1.1	Indicates the medical device manufacturer, as defined in EU Directives 90/385/EEC, 93/42/EEC and 98/79/EC. NOTE 5 The date of manufacture, as well as the name and address of the manufacturer, can be combined in one symbol.



	Date of Manufacture	5.1.3	Indicates the date when the medical device was manufactured.
	Use-by date	5.1.4	Indicates the date after which the medical device is not to be used.
	Do not re-sterilize	5.2.6	Indicates a medical device that is not to be re-sterilized.
	Do not re-use	5.4.2	Indicates a medical device that is intended for one use, or for use on a single patient during a single procedure.
	Authorized Representative in the European Community	5.1.2	Indicates the Authorized representative in the European Community.
	Fragile	5.3.1	Indicates a medical device that can be broken or damaged if not handled carefully.

	Humidity limitation	5.3.8	Indicates the range of humidity to which the medical device can be safely exposed.
	Keep away from sunlight	5.3.2	Indicates a medical device that needs protection from light sources.
	Keep dry	5.3.4	Indicates a medical device that needs to be protected from moisture.
	Sterile	5.2.1	Indicates a medical device that has been subjected to a sterilization process.
	Sterilized using aseptic processing techniques	5.2.2	Indicates a medical device that has been manufactured using using accepted aseptic techniques.
	Sterilized using ethylene oxide	5.2.3	Indicates a medical device that has been sterilized using ethylene oxide.





	Sterilized using irradiation	5.2.4	Indicates a medical device that has been sterilized using irradiation.
	Non-sterile	5.2.7	Indicates a medical device that has not been subjected to a sterilization process.
	Upper limit of temperature	5.3.6	Indicates the upper limit of temperature to which the medical device can be safely exposed.
	Temperature limit	5.3.7	Indicates the temperature limits to which the medical device can be safely exposed.
	Do not use if package is damaged	5.2.8	Indicates a medical device that should not be used if the package has been damaged or opened.
	Atmospheric pressure limitation	5.3.9	Indicates the range of atmospheric pressure to which the medical device can be safely exposed.

	Non-pyrogenic	5.6.3	Indicates a medical device that is non-pyrogenic.
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ISO 7000 Graphical symbols for use on equipment — Index and synopsis

Image	Accompanying Text	Reference
	Packaging unit	2794
	RFID tag, general	3010




IEC 60417 Graphical Symbols for Use on Equipment

Image	Accompanying Text	Reference
	Type CF applied part	5335
	Dangerous voltage	5036
	Defibrillation-proof type BF applied part	5334
	Type BF applied part	5333


ISO 7010 Graphical symbols – Safety colors and safety signs – Registered safety signs

Image	Accompanying Text	Reference
	Refer to instruction manual/booklet	M002


ASTM F2503-13 Standard Practice for Marking Medical Devices and Other Items for Safety in the Magnetic Resonance Environment

Image	Accompanying Text	Reference
	MR Conditional	Table 2, Figure 6
	MR Safe	Table 2, Figure 4
	MR Unsafe	Table 2, Figure 9


Title 21 Code of Federal Regulations Parts 801.109

Image	Accompanying Text
	Caution: Federal law restricts this device to sale by or on the order of a licensed healthcare practitioner.

DIRECTIVE 2012/19/EU OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 4 July 2012 on waste electrical and electronic equipment (WEEE)

Image	Accompanying Text
	Separate collection

EN 15986:2011 Symbols for use in the labelling of medical devices — Requirements for labelling of medical devices containing Phthalates

Image	Accompanying Text	Reference
	Contains or presence of phthalate	A.2, Figure A.2