Symbols Glossary

This document contains symbols that may appear on the product packaging or labeling. It provides information on the symbol including the standard title and reference, symbol title, and description, where available.

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

Symbols from standards

Symbol	Standard reference	Standard title	Symbol title	Explanatory text
	ISO 15223-1, Clause 5.1.1	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	Manufacturer	Indicates the medical device manufacturer
	ISO 15223-1, Clause 5.1.2	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	Authorized representative in the European Community/European Union	Indicates the authorized representative in the European Community/European Union
	ISO 15223-1, Clause 5.1.3	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	Date of manufacture	Indicates the date when the medical device was manufactured
	ISO 15223-1, Clause 5.1.4	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	Use-by date	Indicates the date after which the medical device is not to be used
	ISO 15223-1, Clause 5.1.5	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	Batch code	Indicates the manufacturer’s batch code so that the batch or lot can be identified

Symbol	Standard reference	Standard title	Symbol title	Explanatory text
	ISO 15223-1, Clause 5.1.6	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	Catalogue number	Indicates the manufacturer's catalogue number so that the medical device can be identified
	ISO 15223-1, Clause 5.1.7	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	Serial number	Indicates the manufacturer's serial number so that a specific medical device can be identified
	ISO 15223-1, Clause 5.2.8	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	Do not use if package is damaged and consult instructions for use	Indicates that a medical device that should not be used if the package has been damaged or opened and that the user should consult the instructions for use for additional information
	ISO 15223-1, Clause 5.3.2	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	Keep away from sunlight	Indicates a medical device that needs protection from light sources
	ISO 15223-1, Clause 5.3.4	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	Keep dry	Indicates a medical device that needs to be protected from moisture
	ISO 15223-1, Clause 5.3.7	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	Temperature limit	Indicates the temperature limits to which the medical device can be safely exposed
	ISO 15223-1, Clause 5.3.8	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	Humidity limitation	Indicates the range of humidity to which the medical device can be safely exposed
	ISO 15223-1, Clause 5.4.2	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	Do not re-use	Indicates a medical device that is intended for one single use only

Symbol	Standard reference	Standard title	Symbol title	Explanatory text
	ISO 15223-1, Clause 5.4.3	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	Consult instructions for use or consult electronic instructions for use	Indicates the need for the user to consult the instructions for use
	ISO 15223-1, Clause 5.4.4	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	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
	ISO 15223-1, Clause 5.5.1	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	In vitro diagnostic medical device	Indicates a medical device that is intended to be used as an in vitro diagnostic medical device
	ISO 15223-1, Clause 5.7.10	Medical devices — Symbols to be used with information to be supplied by the manufacturer — Part 1: General requirements	Unique device identifier	Indicates a carrier that contains unique device identifier information
	EN 50419	Marking of electrical and electronic equipment (EEE) in respect to separate collection of waste EEE (WEEE)	Separate collection for electrical and electronic equipment	This electrical and electronic equipment is subject to separate waste collection with a view to minimizing the disposal of waste electrical and electronic equipment as unsorted municipal waste
	IEC 61010-1, Clause 5.1.3	Safety requirements for electrical equipment for measurement, control, and laboratory use - Part 1: General requirements	Direct current	Direct current

Symbols not from standards

Symbol	Reference	Title	Symbol title	Explanatory text
	Regulation (EC) No 1223/2009, Annex VII (2)	Regulation (EC) No 1223/2009 of the European Parliament and of the Council of 30 November 2009 on cosmetic products	Period-after-opening	The period of time after opening for which the product is safe and can be used without any harm to the consumer.
	Regulation (EU) 2017/746, Annex V (1)	Regulation (EU) 2017/746 of the European Parliament and of the Council of 5 April 2017 on in vitro diagnostic medical devices and repealing Directive 98/79/EC and Commission Decision 2010/227/EU	CE marking	A marking by which a manufacturer indicates that a device is in conformity with the applicable requirements set out in this Regulation and other applicable Union harmonisation legislation providing for its affixing.
	Trademarks of PRO EUROPE	N/A	Green dot	In order to make it clear that one has thus assumed responsibility for one's packaging, a contract can be concluded on a voluntary basis with the individual PRO Europe systems for the use of the trademark Green Dot on packaging.
	Trademarks of Bluetooth Special Interest Group (SIG)	N/A	Bluetooth	Bluetooth wireless technology.