
















## Symbol Glossary

Symbol	Standard Reference	Standard Title	Symbol Title	Explanatory Text
	ISO 15223-1 Clause 5.1.5	Medical devices - Symbols to be used with medical device labels, labeling and information to be supplied.	Batch code	<i>Software version (ex: v1.7.0)</i>
	ISO 15223-1 Clause 5.1.6	Medical devices - Symbols to be used with medical device labels, labeling and information to be supplied.	Catalog or model number	<i>Software build ID (Date + Operating System + Supported computation mode)</i>
	ISO 15223-1 Clause 5.4.4	Medical devices - Symbols to be used with medical device labels, labeling and information to be supplied.	Caution	Indicates that caution is necessary when operating the device or control close to where the <i>symbol</i> is placed, or that the current situation needs operator awareness or operator action in order to avoid undesirable consequences.
	NA	NA	Warning	Indicates a limitation in use and steps that must be taken to prevent potential safety concerns.
	NA	NA	Prescription only	Caution: Federal Law (USA) restricts this device to sale by or on the order of a licensed healthcare practitioner. Requires prescription in the United States (made in compliance with 21 CFR Part 801.109).  Device is intended for use by trained medical professionals only.
  eIFU indicator	ISO 15223-1 Clause 5.4.3	Medical devices - Symbols to be used with medical device labels, labeling and information to be supplied.	Consult <i>instructions for use</i> or consult electronic <i>instructions for use</i>	Indicates that the user needs to consult the <i>instructions for use</i>

Symbol	Standard Reference	Standard Title	Symbol Title	Explanatory Text
	ISO 15223-1 Clause 5.1.1	Medical devices - Symbols to be used with medical device labels, labeling and information to be supplied.	<i>Manufacturer</i>	Indicates the medical device <i>manufacturer</i>
	ISO 15223-1 Clause 5.1.8	Medical devices - Symbols to be used with medical device labels, labeling and information to be supplied.	<i>Importer</i>	Indicates the entity importing the <i>medical device</i> into the local
	ISO 15223-1 Clause 5.1.9	Medical devices - Symbols to be used with medical device labels, labeling and information to be supplied.	<i>Distributor</i>	Indicates the entity distributing the <i>medical device</i> into the local
	ISO 15223-1 Clause 5.7.7	Medical devices - Symbols to be used with medical device labels, labeling and information to be supplied.	<i>Medical device</i>	Indicates the item is a <i>medical device</i>
	ISO 15223-1 Clause 5.7.10	Medical devices - Symbols to be used with medical device labels, labeling and information to be supplied.	<i>Unique Device Identifier</i>	Indicates a carrier that contains unique device identifier information
	N/A	N/A	CE Mark	<i>CE Mark followed by the identification number of the notified body responsible for the conformity assessment procedures (made in compliance with Regulation (EU) 2017/745, Article 20 and Annex V)</i>
	N/A	N/A	UKCA Mark	<i>UKCA mark (made in compliance with UK Medical Devices Regulations (2002))</i>
	ISO 15223-1 Clause 5.1.2	Medical devices - Symbols to be used with medical device labels, labeling and information to be supplied.	Authorized representative in the European Community/European Union	Indicates the authorized representative in the European Community/European Union

Symbol	Standard Reference	Standard Title	Symbol Title	Explanatory Text
<div>CH</div> <div>REP</div>	N/A	NA	Authorized representative for Switzerland	Indicates the authorized representative in Switzerland (Made in compliance with Swiss Medical Device Ordinance (MedDO; SR 812.213) and the associated Information sheet on Obligations of Economic Operators).

### Document History

<i>Revision</i>	<i>Date</i>	<i>Description</i>
01	2023-09-19	First Document Issue
02	2024-03-05	Addition of the identification number of the notified body responsible for the conformity assessment procedures under Regulation (EU) 2017/745; addition of the UKCA marking