Symbol Glossary

SYMBOL	SYMBOL TITLE	STANDARD REFERENCE	STANDARD TITLE	EXPLANATORY TEXT
	Catalog number	ISO 15223-1 Reference no. 5.1.6. (ISO 7000-2493)	Medical devices — Symbols to be used with information to be supplied by the manufacturer – Part 1: General requirements.	Indicates the manufacturer's catalog number so the medical device can be identified.
	Lot or Batch code	ISO 15223-1 Reference no. 5.1.5. (ISO 7000-2492)	Medical devices — Symbols to be used with information to be supplied by the manufacturer – Part 1: General requirements.	Indicates the manufacturer's batch code so the batch or lot number can be identified.
	Use by date	ISO 15223-1 Reference no. 5.1.4. (ISO 7000-2607)	Medical devices — Symbols to be used with information to be supplied by the manufacturer – Part 1: General requirements.	Indicates the date after which the medical device is not to be used.
	Date of manufacture	ISO 15223-1 Reference no. 5.1.3. (ISO 7000-2497)	Medical devices — Symbols to be used with information to be supplied by the manufacturer – Part 1: General requirements.	Indicates the date when the medical device was manufactured.
	Manufacturer	ISO 15223-1 Reference no. 5.1.1. (ISO 7000-3082)	Medical devices — Symbols to be used with information to be supplied by the manufacturer – Part 1: General requirements.	Indicates the medical device manufacturer.
	Sterilized using irradiation	ISO 15223-1 Reference no. 5.2.4. (ISO 7000-2502)	Medical devices — Symbols to be used with information to be supplied by the manufacturer – Part 1: General requirements.	Indicates a medical device that has been sterilized using irradiation.
	Do not re-use	ISO 15223-1 Reference no. 5.4.2. (ISO 7000- 1051)	Medical devices — Symbols to be used with information to be supplied by the manufacturer – Part 1: General requirements.	Indicates a medical device that is intended for one single use only. NOTE: Synonyms for “Do not reuse” are “single use” and “use only once”.

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	Do not use if package is damaged and consult instructions for use	ISO 15223-1 Reference no. 5.2.8. (ISO 7000-2606)	Medical devices — Symbols to be used with information to be supplied by the manufacturer – Part 1: General requirements.	Indicates 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.
	Keep dry Keep away from rain	ISO 15223-1 Reference no. 5.3.4. (ISO 7000-0626)	Medical devices — Symbols to be used with information to be supplied by the manufacturer – Part 1: General requirements.	Indicates a medical device that needs protection from moisture.
	Temperature limit	ISO 15223-1 Reference no. 5.3.7. (ISO 7000-0632)	Medical devices — Symbols to be used with information to be supplied by the manufacturer – Part 1: General requirements.	Indicates the temperature limits to which the medical device can be safely exposed. PLEASE NOTE FOR USA ONLY: The limits 20°C to 25°C (68°F to 77°F) also mean that there are excursions permitted between 15°C to 30°C (59°F to 86°F) [See USP Controlled Room Temperature].
	Consult instructions for use	ISO 15223-1 Reference no. 5.4.3. (ISO 7000-1641)	Medical devices — Symbols to be used with information to be supplied by the manufacturer – Part 1: General requirements.	Indicates the need for the user to consult the instructions for use.
	Caution	ISO 15223-1 Reference no. 5.4.4. (ISO 7000-0434A)	Medical devices — Symbols to be used with information to be supplied by the manufacturer – Part 1: General requirements.	To indicate that caution is necessary when operating the device or control close to where the symbol is placed, or to indicate that the current situation needs operator awareness or operator action in order to avoid undesirable consequences
	Prescription Use Only	N/A	N/A	Caution: Federal law (USA) restricts this device to sale by or on the order of a licensed healthcare practitioner.

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QTY	Quantity	N/A	N/A	Indicates the package quantity or number of medical devices included in the package.
CE	CE marking	REGULATION (EU) 2017/745 Reference no. Annex V.[a]	REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 5 April 2017 on medical devices, amending Directive 2001/83/ EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/ EEC and 93/42/EEC	(43) 'CE marking of conformity' or 'CE marking' means 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.
EC REP	Authorized Representative in the European Community / European Union	ISO 15223-1:2021 Reference no. 5.1.2.	Medical devices — Symbols to be used with information to be supplied by the manufacturer – Part 1: General requirements.	Indicates the Authorized Representative in the European Community / European Union.

[a] The Luer lock applicator was issued CE marking under the Medical Devices Directive 93/42/EEC and is transitioning to Regulation (EU) 2017/745.