











Standard*/Source	Symbol	Reference Number	Title of Symbol	Description of Symbol Per Standard
ISO 15223-1:2021		5.1.6	Catalogue number	Indicates the manufacturer's catalogue number so that the medical device can be identified.
ISO 15223-1:2021		5.1.5	Batch Code	Indicates the manufacturer's batch code so that the batch or lot can be identified.
ISO 15223-1:2021		5.1.1	Manufacturer	Indicates the medical device manufacturer.
ISO 15223-1:2021		5.4.4	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 15223-1:2021		5.2.7	Non-sterile	Indicates a medical device that has not been subjected to a sterilization process.
21 CFR 801.109	Rx ONLY	N/A	Prescription use only	Caution: Federal (USA) law restricts this device to sale by or on the order of a licensed healthcare practitioner.
ISO 15223-1:2021		5.4.3	Consult instructions for use or consult electronic instructions for use	Indicates the need for the user to consult the instructions for use.
ISO 15223-1:2021		5.1.3	Date of manufacture	Indicates the date when the medical device was manufactured.
Pacific Instruments generated symbol		N/A	Quantity	Indicates the quantity.
ISO 15223-1:2021		5.7.7	Medical device	Indicates the item is a medical device.
ISO 15223-1:2021		5.7.10	Unique device identifier	Indicates a carrier that contains unique device identifier information.

*Standard title and reference number (FDA Recognition): ISO 15223-1:2021 Fourth Edition 2021-07, Medical devices - Symbols to be used with information to be supplied by the manufacturer - Part 1: General requirements.