









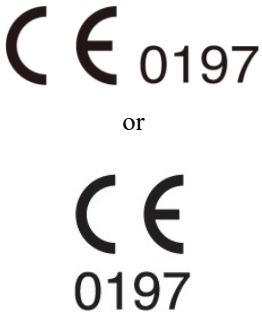


Description of symbol-001

Symbols	Description of symbol	Reference standards
	<p>Indicates the <i>medical device manufacturer</i></p>	<p>ISO 15223-1:2021 5.1.1 ISO 15223-1:2016 5.1.1 Manufacturer</p>
	<p>Indicates the authorized representative in the European Community/ European Union</p>	<p>ISO 15223-1:2016 5.1.2 ISO 15223-1:2021 5.1.2 Authorized representative in the European Community/ European Union</p>
	<p>Indicates the date when the medical device was manufactured</p>	<p>ISO 15223-1:2021 5.1.3 ISO 15223-1:2021 5.1.3 Date of manufacture</p>
	<p>Indicates the date after which the <i>medical device</i> is not to be used</p>	<p>ISO 15223-1:2016 5.1.4 ISO 15223-1:2021 5.1.4 Use-by date</p>
	<p>Indicates the <i>manufacturer's batch code</i> so that the batch or lot can be identified</p>	<p>ISO 15223-1:2016 5.1.5 ISO 15223-1:2021 5.1.5 <i>Batch code</i></p>
	<p>Indicates a <i>medical device</i> that has not been subjected to a sterilization process</p>	<p>ISO 15223-1:2016 5.2.7 ISO 15223-1:2021 5.2.7 <i>Non-sterile</i></p>

Description of symbol-001

Symbols	Description of symbol	Reference standards
	<p>Indicates that a <i>medical device</i> that should not be used if the package has been damaged or opened and that the user should consult the <i>instructions for use</i> for additional information</p>	<p>ISO 15223-1:2016 5.2.8            ISO 15223-1:2021 5.2.8            Do not use if package is damaged and consult <i>instructions for use</i></p>
	<p>Indicates the need for the user to consult the <i>instructions for use</i></p>	<p>ISO 15223-1:2016 5.4.3            ISO 15223-1:2021 5.4.3            Consult <i>instructions for use</i> or consult electronic <i>instructions for use</i></p>
	<p>Indicates a carrier that contains unique device identifier information</p>	<p>ISO 15223-1:2021            5.7.10            Unique device identifier</p>
<p>Rx Only</p>	<p>Prescription use only</p>	<p>21 CFR 801.109</p>
	<p>European conformity</p>	<p>The requirements for accreditation and market surveillance relating to the marketing of products; Medical Device Directive &amp; Medical Device Regulation.</p>
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