<table>
<thead>
<tr>
<th>Symbols</th>
<th>Description of symbol</th>
<th>Reference standards</th>
</tr>
</thead>
</table>
| ![Symbol](image1) | Indicates the medical device manufacturer | ISO 15223-1:2021 5.1.1  
ISO 15223-1:2016 5.1.1  
Manufacturer |
| ![Symbol](image2) | Indicates the authorized representative in the European Community/ European Union | ISO 15223-1:2016 5.1.2  
ISO 15223-1:2021 5.1.2  
Authorized representative in the European Community/ European Union |
| ![Symbol](image3) | Indicates the date when the medical device was manufactured | ISO 15223-1:2021 5.1.3  
ISO 15223-1:2021 5.1.3  
Date of manufacture |
| ![Symbol](image4) | Indicates the date after which the medical device is not to be used | ISO 15223-1:2016 5.1.4  
ISO 15223-1:2021 5.1.4  
Use-by date |
| ![Symbol](image5) | Indicates the manufacturer’s batch code so that the batch or lot can be identified | ISO 15223-1:2016 5.1.5  
ISO 15223-1:2021 5.1.5  
Batch code |
| ![Symbol](image6) | Indicates a medical device that has not been subjected to a sterilization process | ISO 15223-1:2016 5.2.7  
ISO 15223-1:2021 5.2.7  
Non-sterile |
<table>
<thead>
<tr>
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</table>
| ![No symbol](image) | 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 15223-1:2016 5.2.8  
ISO 15223-1:2021 5.2.8  
Do not use if package is damaged and consult *instructions for use* |
| ![Information symbol](image) | Indicates the need for the user to consult the *instructions for use* | ISO 15223-1:2016 5.4.3  
ISO 15223-1:2021 5.4.3  
Consult *instructions for use* or consult electronic *instructions for use* |
| ![UDI](image) | Indicates a carrier that contains unique device identifier information | ISO 15223-1:2021  
5.7.10  
Unique device identifier |
| ![Rx Only](image) | Prescription use only | 21 CFR 801.109 |
| ![CE](image) | European conformity | The requirements for accreditation and market surveillance relating to the marketing of products; Medical Device Directive & Medical Device Regulation. |
| ![CE0197](image) | European conformity | The requirements for accreditation and market surveillance relating to the marketing of products; Medical Device Directive & Medical Device Regulation. |