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

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