









Symbol Glossary

The Symbol on this page are used in medical device labeling in accordance with ISO 15223. They are used to convey information on the IFUs and product packaging.

Symbol	Title/Description
	Manufacturer Indicates the medical device manufacturer.
	Date of Manufacturer Indicates the date when the medical device was manufactured.
	Batch Code Indicates the manufacturer's batch code so that the batch or lot can be identified.
	Catalogue Number Indicates the manufacturer's catalogue number so that the medical device can be identified.
	Medical device Indicates that this is a medical device.
	Quantity of devices Indicates quantity of devices.
	Importer Indicates the entity importing the medical device into the locale.
	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.

**CE Mark**

A marking by which the manufacturer indicates that a device is in conformity with the applicable requirements set out in the European legislation for medical devices and other applicable Union harmonisation legislation providing for its affixing.

**Prescription device**

Indicates the medical device manufacturer.

**Use-by date**

Indicates the date after which the medical is not to be used.

**Do not use if the package is damaged or opened**

Indicates a medical device that should not be used if the package has been damaged or opened.

**Do not re-use**

Indicates a medical device that is intended for one use, or for use on a single patient during a single procedure.

**No Latex**

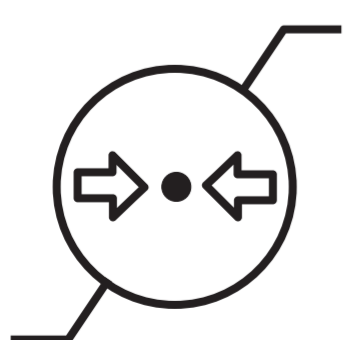
Indicates that the medical device does not contain presence of natural rubber latex.

**Contains or presence of natural rubber latex**

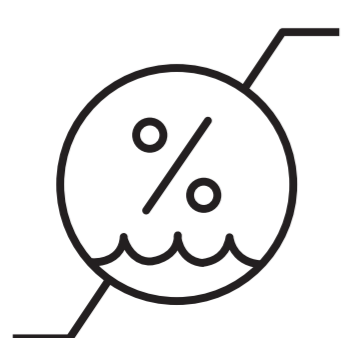
Indicates the presence of natural rubber or dry natural rubber latex as material of construction within the medical of the packaging of a medical device.

**Do not re-sterilize**

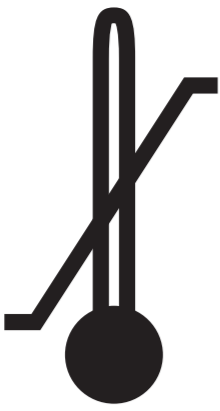

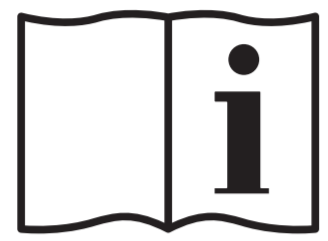


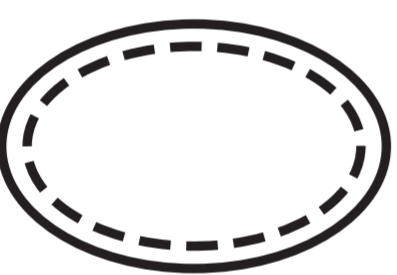
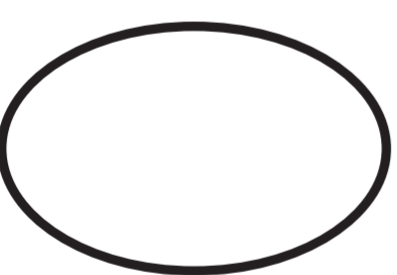


Indicates that the medical device is not to be re-sterilized.

**Atomic pressure limitation**

Indicates the range of atmospheric pressure to which the medical device can be safely exposed.

**Humidity limitation**

Indicates the range of humidity to which the medical device can be safely exposed.

	<p>Temperature limit</p> <p>Indicates the temperature limit to which the medical device can be safely exposed.</p>
	<p>Collect separately</p> <p>Separate collection for waste of electrical and electronic equipment. Do not dispose of battery in municipal waste. The symbol indicates separate collection for battery is required.</p>
 <p>xodusmedical.com/eifu</p>	<p>Consult instructions for use or consult electronic instructions for use</p> <p>Indicates the need for the user to consult the instructions for use.</p>
	<p>Non-Sterile</p> <p>Indicates a medical device that has not been subjected to a sterilization process.</p>
	<p>Sterilized using irradiation</p> <p>Indicates that the medical device that has been sterilized using irradiation.</p>
	<p>Single sterile barrier system with protective packaging inside</p> <p>Indicates a single sterile barrier system with protective packaging inside.</p>
	<p>Single sterile barrier system</p> <p>Indicates a single sterile barrier system.</p>
	<p>Authorized Representative in the European Community/ European Union</p> <p>Indicates the authorized representative in the European Community / European Union.</p>
	<p>Authorized Representative in the Switzerland Community</p> <p>Indicates the authorized representative in the Switzerland Community.</p>