









Symbol Glossary

F-7.5.3-F32











PriMed Instruments Inc.





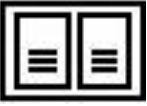
Ref. Number/ Symbol	Title	Description	Reference Document
5.2.7 	Non-sterile	Indicates a medical device that has not been subjected to a sterilization process.	ISO 15223-1
5.4.4 	Caution	Indicates the need for the user to consult the instructions for use for important cautionary information such as warnings and precautions that cannot, for a variety of reasons, be presented on the medical device itself.	ISO 15223-1
5.4.3 	Consult instructions for use	Indicates the need for the user to consult the instructions for use.	ISO 15223-1
5.3.4 	Keep dry	Indicates a medical device that needs to be protected from moisture.	ISO 15223-1
5.4.1 	Biological risks	Indicates that there are potential biological risks associated with the medical device.	ISO 15223-1
5.3.2 	Keep away from sunlight	Indicates a medical device that needs protection from light sources.	ISO 15223-1
5.2.8 	Do not use if package is damaged	Indicates a medical device that should not be used if the package has been damaged or opened.	ISO 15223-1
5.4.2 	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.	ISO 15223-1

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5.2.6 	Do not resterilize	Indicates a medical device that is not to be resterilized.	ISO 15223-1
5.1.1 	Manufacturer	Indicates the medical device manufacturer, as defined in EU Directives 90/385/EEC, 93/42/EEC and 98/79/EC.	ISO 15223-1
5.1.3 	Date of manufacture	Indicated the date when the medical device was manufactured.	ISO 15223-1
5.1.4 	Use-by date	Indicates the date after which the medical device is not to be used.	ISO 15223-1
5.2.3 	Sterilized using ethylene oxide	Indicates a medical device that has been sterilized using ethylene oxide.	ISO 15223-1
5.1.2 	Authorized representative in the European Community	Indicates the Authorized representative in the European Community	ISO 15223-1
5.1.5 	Batch code	Indicates the manufacturer's batch code so that the batch or lot can be identified.	ISO 15223-1
5.1.6 	Catalogue number	Indicates the manufacturer's catalogue number so that the medical device can be identified.	ISO 15223-1
801.109 (b)(1) 	Prescription only	Indicates the Federal Law (USA) restricts this device to sale by or on the order of a physician.	21 CFR 801 81 FR 38911
Article 17 Annex XII 	European Conformity	Indicates that the product may be legally placed on the market in the European Union.	MDD 93/42/EEC

	<p>Product number</p>	<p>Indicates the product number</p>	<p>—</p>
	<p>No Phthalate</p>	<p>Indicates the product is not made with Phthalate</p>	<p>—</p>
	<p>No Latex</p>	<p>Indicates the product is not made with Latex.</p>	<p>—</p>
	<p>Dispenser quantity</p>	<p>Indicates the quantity of units in dispenser.</p>	<p>—</p>
	<p>Case quantity</p>	<p>Indicates the quantity of units in a case.</p>	<p>—</p>

(ISO 15223-1) Medical devices – Symbols to be used with medical device labels, labelling and information to be supplied – Part 1: General requirements