BISCO, Inc.
1100 W. Irving Park Road
Schaumburg, IL 60193 U.S.A.

en - Symbols Glossary

Symbol	Title of Symbol	Standard	Description of Symbol
	Manufacturer	ISO 15223-1:2021 5.1.1	Indicates the medical device manufacturer
	Authorized representative in the European Community / European Union	ISO 15223-1:2021 5.1.2	Indicates the authorized representative in the European Community / European Union
	Date of manufacture	ISO 15223-1:2021 5.1.3	Indicates the date when the medical device was manufactured
	Use-by date	ISO 15223-1:2021 5.1.4	Indicates the date after which the medical device is not to be used
	Batch code	ISO 15223-1:2021 5.1.5	Indicates the manufacturer's batch code so that the batch or lot can be identified
	Catalogue number	ISO 15223-1:2021 5.1.6	Indicates the manufacturer's catalogue number so that the medical device can be identified
	Keep away from sunlight	ISO 15223-1:2021 5.3.2	Indicates a medical device that needs protection from light sources
	Temperature limit	ISO 15223-1:2021 5.3.7	Indicates the temperature limits to which the medical device can be safely exposed

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Symbol	Title of Symbol	Standard	Description of Symbol
	Do not re-use	ISO 15223-1:2021 5.4.2	Indicates a medical device that is intended for one single use only
	Consult instructions for use or consult electronic instructions for use	ISO 15223-1:2021 5.4.3	Indicates the need for the user to consult the instructions for use
	Caution	ISO 15223-1:2021 5.4.4	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
	Medical device	ISO 15223-1:2021 5.7.7	Indicates the item is a medical device
	CE mark / European Conformity / Conformité Européenne	European Medical Device Directive 93/42/EEC (as amended by Directive 2007/47/EC) and Regulation (EU) 2017/745	Indicates conformity of the product with applicable European legislations

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Symbol	Title of Symbol	Standard	Description of Symbol
CE 0459	CE mark / European Conformity / Conformité Européenne with notified body identification number	European Medical Device Directive 93/42/EEC (as amended by Directive 2007/47/EC) and Regulation (EU) 2017/745	Indicates conformity of the product with applicable European legislations where a notified body was responsible for the conformity assessment Notified Body Number 0459: GMED SAS, France
UK CA	UK conformity assessment mark	MHRA Guidance Medical devices: conformity assessment and the UKCA mark	Indicates conformity of the product with applicable UK legislations
UK REP	UK responsible person	ISO 20417:2021 6.1.2.d.1	Indicates the UK responsible person
	Skin or eye irritation or allergic skin reaction	Regulation (EC) No. 1272/2008	Indicates product may cause skin or eye irritation or an allergic skin reaction
	Flammable liquid and/or vapor	Regulation (EC) No. 1272/2008	Indicates product may contain substances that are flammable
	Corrosive	Regulation (EC) No. 1272/2008	Indicates product is corrosive

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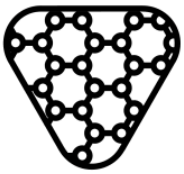
Symbol	Title of Symbol	Standard	Description of Symbol
	Acute toxicity	Regulation (EC) No. 1272/2008	Indicates product is fatal or toxic if inhaled, swallowed, or in contact with skin
	Health hazard	Regulation (EC) No. 1272/2008	Indicates product may be a health hazard such as, • product may cause allergy or asthma symptoms or breathing difficulties if inhaled • product may be carcinogenic, mutagenic or reprotoxic
	Contains nano materials	ISO 15223-1: 2021 5.4.11	Indicates a medical device that contains nano materials