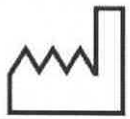
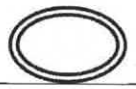
APTIS
Putting Ideas in Motion

Glossary of Symbols

Symbol	Standard / Symbol Reference	Symbol Title	Description of Symbol
	ISO 15223-1 / 5.1.1	Manufacturer	Indicates the medical device manufacturer.
	ISO 15223-1 / 5.1.2	Authorized representative in the European Community/European Union	Indicates the authorized representative in the European Community/ European Union.
	ISO 15223-1 / 5.1.3	Date of manufacture	Indicates 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 / 5.1.5	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 / 5.1.7	Serial Number	Indicates the *manufacturer's serial number* so that a specific *medical device* can be identified. Used for tracking instrument sets.
	ISO 15223-1 / 5.2.3	Sterilized using ethylene oxide	Indicates a medical device that has been sterilized using ethylene oxide.

Symbol	Standard / Symbol Reference	Symbol Title	Description of Symbol
	ISO 15223-1 / 5.2.6	Do not re-sterilize	Indicates a medical device that is not to be re-sterilized.
	ISO 15223-1 / 5.2.7	Non-sterile	Indicates a medical device that has not been subjected to a sterilization process.
	ISO 15223-1 / 5.2.8	Do not use if package is damaged and consult instructions for use	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 / 5..12	Double sterile barrier system	Indicates two sterile barrier systems.
	ISO 15223-1 / 5.4.2	Do not re-use	Indicates a medical device that is intended for one single use only.
	ISO 15223-1 / 5.4.3	Consult instructions for use or consult electronic instructions for use	Indicates the need for the user to consult the instructions for use.
	ISO 15223-1 / 5.4.4	Caution	Indicates the need for the user to consult the instructions for use 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.
	ISO 15223-1 / 5.4.10	Contains hazardous substances	Indicates a medical device that contains substances that can be carcinogenic, mutagenic, reprotoxic (CMR), or substances with endocrine disrupting properties.

Symbol	Standard / Symbol Reference	Symbol Title	Description of Symbol
	ISO 15223-1 / 5.7.7	Medical device	Indicates the item is a medical device.
	ISO 15223-1 / 5.7.10	Unique device identifier	Indicates a carrier that contains unique device identifier information.
	MDD 93/42/EEC / Article 17	CE Mark	Signifies European technical conformity.
	ASTM F2503 Figure 6	MR Conditional	Demonstrates safety in the MR environment within defined conditions including conditions for the static magnetic field, the time-varying gradient magnetic fields and the radiofrequency fields.
	ISO 15223-1 / 5.7.3	Patient identification	Indicates the identification data of the patient.
	ISO 15223-1 / 5.7.6	Date	Indicates the date that information was entered or a medical procedure took place
	ISO 15223-1 / 5.7.5	Health care center or doctor	Indicates the address of the health care center or doctor where medical information about the patient may be found.
	ISO 15223-1 / 5.7.4	Patient information website	Indicates a website where a patient can obtain additional information on the medical product.