

SYMBOL	SYMBOL TITLE	EXPLANATORY TEXT	STANDARD REFERENCE	REFERENCE NUMBER
	Manufacturer	Indicates the medical device manufacturer	ISO 15223-1, Clause 5.1.1	5.1.1
	5		V50 45222 4 6	5.4.3
IП	Date of manufacture	Indicates the date when the medical device was manufactured	ISO 15223-1, Clause 5.1.3	5.1.3
(, , , ,		manufactureu		
	Use-by date	Indicates the date after which the medical device is not	ISO 15223-1, Clause 5.1.4	5.1.4
><	,	to be used		
	Batch code	Indicates the manufacturer's batch code so that the	ISO 15223-1, Clause 5.1.5	5.1.5
LOT		batch or lot can be identified. Synonyms for "batch		
		code" are "lot number", "lot code" and "batch number".		
	Catalogue number	Indicates the manufacturer's catalog number so that the	ISO 15223-1, Clause 5.1.6	5.1.6
REF		medical device can be identified		
	Charila	Indicates a madical device that has been subjected to a	ISO 15222 1 Clause 5 2 1	F 2.1
STERILE	Sterile	Indicates a medical device that has been subjected to a sterilization process	ISO 15223-1, Clause 5.2.1	5.2.1
STERILE		Stermzation process		
	Sterilized using	Indicates a medical device that has been sterilized using	ISO 15223-1, Clause 5.2.3	5.2.3
STERILE R	irradiation	irradiation.		
	Consult instructions for	Indicates the need for the user to consult the	ISO 15223-1, Clause 5.4.3	5.4.3
_ i	use or consult	instructions for use		
	electronic instructions			
\bigcirc	Do not re-use	Indicates a medical device that is intended for one	ISO 15223-1, Clause 5.4.2	5.4.2
$ \langle X \rangle $		single use only		
$\stackrel{\smile}{\sim}$	Do not resterilize	Indicates a medical device that is not to be resterilized	ISO 15223-1, Clause 5.2.6	5.2.6
STERNIZE	Do not resternize	indicates a medical device that is not to be resternized	130 13223 1, Clause 3.2.0	5.2.0
	Prescription Use Only	Caution: Federal law (USA) restricts this device to sale	21 CFR 801.15(c)(1)(i)F	21 CFR 801.15(c)(1)(i)F
R _x Only		by or on the order of a licensed healthcare practitioner.		
		Not made with natural rubber latex.	ISO 15223-1:2012 (Annex B, 15223-1:2012)	Annex B
(LARTEX)	rubber latex			
	Unique device	Indicates a covier that contains unique device identifies	ISO 15222 1 Clause 5 7 10	5.7.10
UDI	Unique device identifier	Indicates a carrier that contains unique device identifier information.	ISO 15223-1 Clause 5.7.10	5.7.10
ODI	lacitanci	iniornation.		
	Do not use if package is	Indicates a medical device that should not be used if the	ISO 15223-1, Clause 5.2.8	5.2.8
(652)	damaged and consult	package has been damaged or opened.		
	instructions for use.			
_		Indicates the need for the user to consult the		
/I \	Caution: Read all	instructions for use for important cautionary		
<u> </u>	warnings and precautions in	information such as warnings and precautions that cannot, for a variety of reasons, be presented on the		
	instructions for use	medical device itself.	ISO 15223-1, Clause 5.4.4	5.4.4
CTED!! EEC	Sterilized using	Indicates a medical device that has been sterilized using	ISO 15223-1, Clause 5.2.3	
STERILE	ethylene oxide	ethylene oxide		5.2.3
	Single sterile			
(3	barrier system	Indicator a cingle storile barrier system with most attra-		
\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\	with protective packaging inside	Indicates a single sterile barrier system with protective packaging inside	ISO 15223-1, Clause 5.2.13	5.2.13
_	packaging marac	Page 1910	TO TOTAL I, CHARGE O.E. I.J	5.2.13
		Indicates a medical device that has not been subjected		
NON STERILE	Non-sterile	to a sterilization process	ISO 15223-1, Clause 5.2.7	5.2.7
* D-f ICC		Provides - Symbols to be used with information to be suppli		

^{*} Reference: ISO 15223-1:2021 Medical Devices - Symbols to be used with information to be supplied by the manufacturer - Part 1: General Requirements