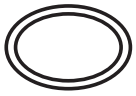
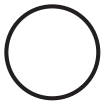
Symbol & Title	Definition	Title & Designation Number of Standard	Symbol Reference Number
	Medical Device	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.7.7
	CE mark	Directive 93/42/EEC MDR 2017/745	Annex V
	Catalogue number	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.1.6
	Batch code	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.1.5
	Sterilized using irradiation	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.2.4
	Sterilized using ethylene oxide	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.2.3
	Sterile	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.2.1
	Single sterile barrier system	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.2.11
	Single sterile barrier system with protective packaging inside	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.2.13
	Single sterile barrier system with protective packaging outside	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.2.14
	Double sterile barrier system	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.2.12
	Use by date	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.1.4

Symbol & Title	Definition	Title & Designation Number of Standard	Symbol Reference Number
SN	Serial number	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.1.7
℞ ONLY or ℞ ONLY	CAUTION: Federal (USA) law restricts this device to sale by or on the order of a physician.	21 CFR 801.109	B (1)
	Caution	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.4.4
		IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	Annex D, Table D.1, No. 10
	Do not reuse	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.4.2
	Do not resterilize	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.2.6
	Do not use if package is damaged and consult instructions for use	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.2.8
 www.applied medical.com/IFU	Consult instructions for use or consult electronic instructions for use	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.4.3
	Follow operating instructions	IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	Annex D, Table D.1, Item #11
	Consult instructions for use / This safety sign is blue in color per ISO 7010-M002.	ISO 20417;2021;Corr1;2021: Information to be supplied by the manufacturer. Symbol derived from ISO 7010 M002	6.1.5
	Follow instructions for use	IEC 60601-1;ed3.2;2020 Medical Electrical Equipment - Part 1: General requirements for basic safety and essential performance	7.2.3 7.2.5
	Non-pyrogenic	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.6.3

Symbol & Title	Definition	Title & Designation Number of Standard	Symbol Reference Number
	Contains or presence of phthalate	BS EN 15986:2011 Symbol for use in the labelling of medical devices – Requirements for labelling of medical devices containing phthalates	4.2 Figure 1
	Non-ionizing radiation	EN 60601-1-2;ED4.1;2020 - Medical electrical equipment - Part 1-2: General requirements for basic safety and essential performance - Collateral standard: Electromagnetic compatibility - Requirements and tests	N/A
	Temperature limitation	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	5.3.7
	Upper limit of temperature	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	5.3.6
	Lower limit of temperature	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	5.3.5
	Humidity limitation	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	5.3.8
	Atmospheric pressure limitation	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	5.3.9
	Separate collection for waste of electrical and electronic equipment.	Directive 2002/96/EC Directive 2012/19/EU BS EN 50419:2022	Annex IX 4.1.2(b)
IP2X	Protected against solid foreign objects of 12.5 mm diameter and greater	IEC 60529;ed2.0;2001;am2;2013 CSV/COR2:2015 Degrees of protection provided by enclosures (IP code) IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	2.1

Symbol & Title	Definition	Title & Designation Number of Standard	Symbol Reference Number
IPX4	Protected against splashing water	IEC 60529;ed2.0;2001;am2;2013 CSV/COR2:2015 Degrees of protection provided by enclosures (IP code)	4.1
		IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	Annex D, Table D.3, No. 2
	Stand-by	IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	Annex D, Table D.1, No. 29
	Alternating current	IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	Annex D, Table D.1, No. 1
	Type BF Applied Part	IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	Annex D, Table D.1, No. 20
	Type CF applied part	IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	Annex D, Table D.1, No. 21
	Defibrillation-proof type CF applied part	IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	Annex D, Table D.1, No. 27
	On	IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	Annex D, Table D.1, No. 12
	Off	IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	Annex D, Table D.1, No. 13
	Equipotentiality	IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	Annex D, Table D.1, No. 8
	Low priority alarm	IEC 60601-1;ed3.2;2020 Medical electrical equipment: Part 1-8: General requirements for basic safety and essential performance – Collateral standard: General requirements, tests, and guidance for alarm systems in medical electrical equipment and medical electrical systems	Annex C, Table C.1
	Dangerous Voltage	IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	Annex D, Table D.1, No. 24
	HF Isolated Patient Circuit	EN 60601-1-2;ED4.1;2020 – Medical electrical equipment – Part 1-2: General requirements for basic safety and essential performance – Collateral standard: Electromagnetic compatibility – Requirements and tests	5.2.2.6

Symbol & Title	Definition	Title & Designation Number of Standard	Symbol Reference Number
	Class II Equipment	IEC 60601-1;ed3.2;2020 Medical electrical equipment – Part 1: General requirements for basic safety and essential performance	Annex D, Table D.1, Item 9
	UL Recognized component mark	N/A	N/A
			
			
	UL Listed	N/A	N/A
	UL Classified	N/A	N/A
	Non-sterile	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.2.7
	Magnetic resonance conditional	ASTM F 2503:2020 Standard Practice for Marking Medical Devices and Other Items for Safety in the Magnetic Resonance Environment	Table 2
	Magnetic resonance safe	ASTM F 2503:2020 Standard Practice for Marking Medical Devices and Other Items for Safety in the Magnetic Resonance Environment	Table 2
	Magnetic resonance unsafe	ASTM F 2503:2020 Standard Practice for Marking Medical Devices and Other Items for Safety in the Magnetic Resonance Environment	Table 2
	Unique device identifier	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.7.10
	Authorized representation in the European Community/European Union	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.1.2

Symbol & Title	Definition	Title & Designation Number of Standard	Symbol Reference Number
CH REP	Swiss Authorised Representative	Swissmedic MedDO	Art. 51, para. 1
UK CA	UK Conformity Assessment	UK MDR 2002	5.1.2
	Importer	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.1.8
	Manufacturer	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.1.1
	Manufacturer and date of manufacture		
	Date of manufacture	ISO 15223-1:2021 Medical devices-Symbols to be used with information to be supplied by the manufacturer-Part 1: General requirements	5.1.3
 or 	Not made with natural rubber latex	FDA guidance document-Recommendations for Labeling Medical Products to Inform Users that the Product or Product Container is not Made with Natural Rubber Latex Guidance (2014)	4.0