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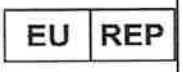
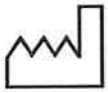
Ultimate Wireforms, Inc.

Label Symbols Glossary

Symbol	Associated Standard	Title of Symbol & Reference #	Meaning of Symbol
	ISO 15223-1: 2021(E)	Manufacturer 5.1.1	Indicates the medical device manufacturer
	ISO 15223-1:2021/Amd 1:2025(en)	Authorized representative in the European Community/European Union 5.1.2	Indicates the authorized representative in the European Community/European Union
	ISO 15223-1: 2021(E)	5.1.2	Indicates the date when the medical device was manufactured
	ISO 15223-1: 2021(E)	Batch Code (Lot Number) 5.1.5	Indicates the manufacturer's batch code so that the batch or lot can be identified
	ISO 15223-1: 2021(E)	Catalog number (Reference number) 5.1.6	Indicates the manufacturer's catalog number so that the medical device can be identified
	ISO 15223-1: 2021(E)	Non-sterile 5.2.7	Indicates a medical device that has not been subjected to a sterilization process.
	ISO 15223-1: 2021(E)	Do not re-use 5.4.2	Indicates a medical device that is intended for one single use only
	ISO 15223-1: 2021(E)	Caution 5.4.4	Indicates that caution is necessary in regard to use; operator awareness and/or action is needed in order to avoid undesirable consequences.
	ISO 20417, 6.1.3 c	ISO 7000-2725	Indicates Nickel within, related to allergen.
	ISO 15223-1: 2021(E)	Medical Device 5.7.7	Indicates the item is a medical device
	ASTM F2503-23	MR Unsafe 6.3.3	Indicate MR Unsafe; UWI products have not been evaluated for compatibility in a MRI environment.
	ISO 15223-1: 2021(E)	Unique Device Identifier 5.7.10	Indicates a carrier that contains unique device identifier information.
	MDD 93/42/EEC Annex II; MDR 2017/745 Annex V	CE Mark European Conformity	Indicates the product conforms to the requirements for accreditation and market surveillance relating to the marketing of products; Medical Device Directive and Medical Device Regulation.
	(US) 21 CFR 801.109 Labeling; Prescription devices	Prescription use only	Caution: Federal law (USA) restricts this device to sale by or on the order of a licensed healthcare practitioner.