N/A	*In vitro* Diagnostic Medical Device	Indicates a medical device that is intended to be used as an in vitro diagnostic medical device.	ISO 15223-1:2016(E) Medical Devices – Symbols to be used with medical device labels, labeling and information to be supplied ISO 15223-1-5.5.1
2495	Negative Control	Indicates a control material that is intended to verify the results in the expected negative range.	ISO 15223-1:2016(E) Medical Devices – Symbols to be used with medical device labels, labeling and information to be supplied ISO 15223-1-5.5.3
2496	Positive Control	Indicates a control material that is intended to verify the results in the expected positive range.	ISO 15223-1:2016(E) Medical Devices – Symbols to be used with medical device labels, labeling and information to be supplied ISO 15223-1-5.5.4
0518	Contains Sufficient for <n> Tests	Indicates the total number of IVD tests that can be performed with the IVD.	ISO 15223-1:2016(E) Medical Devices – Symbols to be used with medical device labels, labeling and information to be supplied ISO 15223-1-5.5.5
3079	Open Here	Identifies the location where the package can be opened and to indicate the method of opening it.	ISO 7000 Graphical symbols for use on equipment.
2606	Do Not Use if Package is Damaged	Indicates a medical device that should not be	

GH508	Health Hazard	Indicates the possible presence following health hazards; carcinogen, respiratory sensitizer, reproductive toxicity, target organ toxicity, mutagenicity, aspiration toxicity.	OSHA's HCS, Appendix C to §1910.1200, Section C.4.11 (Classified in Accordance with Appendix A.8)
GH502	Flame	Indicates the possible presence of the following; flammables, self reactives, pyrophorics, self-heating, emits flammable gas, organic peroxides.	OSHA's HCS, Appendix C to §1910.1200, Section C.4.19 (Classified in Accordance with Appendix B.6)
GH503	Flame Over Cda	d in Accordance wit ac f. t i-ęes. đwa c nA's HCS, Appendix C to §1910.1200, Section C.4.19 Ï (Classified in Accordance with Appendix B.6)FlŸ ₃ c] ę 6fa¾ tl þ x	

European Conformity	Indicates the medical device conforms to European Medical Directive 93/42/EEC and meets applicable health, safety, and environmental requirements. If the mark is accompanied by a number, conformity is verified by the inf aata0

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2020-06