


























Explanation of Symbols

Symbol	Symbol title	Description	Standard reference
	Manufacturer	Indicates the medical device manufacturer	ISO 15223-1:2021 Ref. No. 5.1.1
	Use-by date	Indicates the date after which the medical device is not to be used	ISO 15223-1:2021 Ref. No. 5.1.4
	Batch code	Indicates the manufacturer's batch code so that the batch or lot can be identified	ISO 15223-1:2021 Ref. No. 5.1.5
	Catalogue number	Indicates the manufacturer's catalogue number so that the medical device can be identified	ISO 15223-1:2021 Ref. No. 5.1.6
	Keep away from sunlight	Indicates a medical device that needs protection from light sources	ISO 15223-1:2021 Ref. No. 5.3.2
	Keep dry	Indicates a medical device that needs to be protected from moisture	ISO 15223-1:2021 Ref. No. 5.3.4
	Temperature limit	Indicates the temperature limits to which the medical device can be safely exposed	ISO 15223-1:2021 Ref. No. 5.3.7
	Do not re-use	Indicates a medical device that is intended for one single use only	ISO 15223-1:2021 Ref. No. 5.4.2
	Consult instructions for use or consult electronic instructions for use	Indicates the need for the user to consult the instructions for use	ISO 15223-1:2021 Ref. No. 5.4.3
	Caution	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 15223-1:2021 Ref. No. 5.4.4
	In vitro diagnostic medical device	Indicates a medical device that is intended to be used as an in vitro diagnostic medical device	ISO 15223-1:2021 Ref. No. 5.5.1
	Contains sufficient for <n> tests	Indicates the total number of IVD tests that can be performed with the IVD medical device	ISO 15223-1:2021 Ref. No. 5.5.5
	Unique device identifier	Indicates a carrier that contains unique device identifier information	ISO 15223-1:2021 Ref. No. 5.7.10

Explanation of Symbols

Symbol	Symbol title	Description	Standard reference
	CE marking	CE marking of conformity' or 'CE marking' means a marking by which a manufacturer indicates that a device is in conformity with the applicable requirements set out in this Regulation and other applicable Union harmonisation legislation providing for its affixing	Regulation (EU) 2017/746, Ref. No. Annex V
	CE conformity mark Notified Body ID number	CE marking is followed by the registration number of the notified body involved in the conformity assessment	n/a
	Research Use Only	Indicates that the product is intended only for research purposes	n/a
	Contains <n> tubes for single use test	The symbol describes the total number of tubes in the package	n/a
	Concentrated solution (10x)	The symbol indicates, that the solution is 10x concentrated	n/a
	Concentrated solution (25x)	The symbol indicates, that the solution is 25x concentrated	n/a
	Contents	The symbol describes the contents of the package	n/a
	Indicates the authorized representative in Switzerland	The symbol indicates the authorized representative in Switzerland	n/a
	UKCA mark	Conformity marking used for products being placed on the market in Great Britain	n/a
	Flame (GHS02)	Pictogram that is intended to convey specific information on the hazard concerned	Regulation (EC) 1272/2008, Annex V
	Exclamation mark (GHS07)	Pictogram that is intended to convey specific information on the hazard concerned	Regulation (EC) 1272/2008, Annex V
	Health hazard (GHS08)	Pictogram that is intended to convey specific information on the hazard concerned	Regulation (EC) 1272/2008, Annex V