

Declaration

	Sunrise Medical GmbH
Company:	Kahlbachring 2-4
	69254 Malsch - Germany

Product:	Quickie Q500 H
(May include accessories)	QUICKIE Q300 FI

We, Sunrise Medical declare under our sole responsibility that the product(s) to which this declaration relates, is a class 1 device, does not contain latex.

This was verified with conformity evaluation procedures according to Medical Device Directive Annex VII.

Approval Name and Function	Revision	Approval Date
Michael Kutzer, Director R&D and PDM Europe	Α	25.10.2019

Signature (Sunrise Medical Approval representative)

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