

## **Declaration**

Company: Sunrise Medical GmbH Kahlbachring 2-4

D-69254 Malsch / Heidelberg

**Product:** 

(May include accessories)

Q300M Mini

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.

Mr. D.S.Davies Director Project Management & R&D Europe	A	17.11.2020	
Approval Name and Function	Revision	Approval Date	
Allander.			
Signature (Sunrise Medical Approval representative	re)		

GMS Form Numbe	r:	Revision:	В	Effective Date:	01.02.2010
Form Owner:	Heads of Engineering	Form Approver: Global Head of Engineering		GMS Change Number:	
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