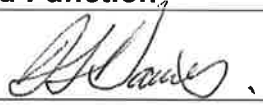
SUNRISE MEDICAL

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**
		
Signature (Sunrise Medical Approval representative)		