SUNRISE MEDICAL

Declaration

Company:	Sunrise Medical GmbH Kahlbachring 2-4 69254 Malsch - Germany
Product: (May include accessories)	**F55**
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.	

Michael Kutzer, Director R&D and PDM Europe	01	20-JAN-2020
Approval Name and Function	**Revision**	**Approval Date**

Signature (Sunrise Medical Approval representative)