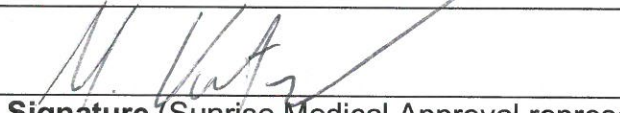




Declaration

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

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


Signature (Sunrise Medical Approval representative)

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