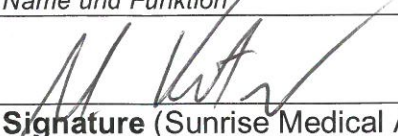




EC DECLARATION OF CONFORMITY

EU Konformitätserklärung

Company: <i>Die Firma</i>	Sunrise Medical GmbH Kahlbachring 2-4 D-69254 Malsch
Product: <i>Produkt</i> (May include accessories) <i>(Kann Zubehör beinhalten)</i>	Quickie Krypton F
<p>We, Sunrise Medical declare under our sole responsibility that the product(s) to which this declaration relates, is a class 1 device, and is in conformity with the requirements of EC Council Directive for Medical Devices 93/42/EEC.</p> <p>This was verified with conformity evaluation procedures according to Medical Device Directive Annex VII.</p>	

Michael Kutzer <i>Director R&D, PDM, Europe</i>	A	03.04.2017
Approval Name and Function <i>Name und Funktion</i>	Revision <i>Revision</i>	Approval Date <i>Genehmigungsdatum</i>
		
Signature (Sunrise Medical Approval representative) <i>Unterschrift</i>		

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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