

## EC DECLARATION OF CONFORMITY

### *Declaración de Conformidad CE*

<b>Company:</b> <i>Campania:</i>	Sunrise Medical S.L. Polígono Bakiola, 41 48498 Arrankudiaga Vizcaya / ESPAÑA
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<b>Product:</b> <i>Producto:</i> <b>(May include accessories)</b> <i>(Incluye accesorios)</i>	<b>Steel manual wheelchair</b> <b>Silla manual de acero</b> <b>BREEZY 250</b>
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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 of 14 June 1993 amended by 2007/47/EEC of 21 March 2010.

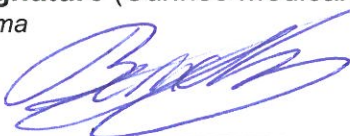
*Por la presente declaramos que los productos arriba mencionados, son clase I, y que son conformes a los requisitos de la Directiva de dispositivos Médicos 93/42/CEE, modificada por la Directiva 2007/47/CE de 21 de marzo de 2010*

This was verified with conformity evaluation procedures according to Medical Device Directive Annex VII.

*It ha sido verificado con conformidad a los procedimientos de evaluación acorde al anexo VII de la directiva de productos sanitarios..*

<b>Bernd Krebs - SVP PM, Marketing Services &amp; Engineering</b>	<b>A</b>	<b>23.09.2011</b>
<b>Approval Name and Function</b> <i>Nombre y Función</i>	<b>Revision</b> <i>Revision</i>	<b>Approval Date</b> <i>Fecha de aprobación</i>

**Signature (Sunrise Medical Approval representative)**  
*Firma*



GMS Form Number: GL 3.1.1.1.F35	Revision: <b>B</b>	Effective Date: <b>01.02.2010</b>
Form Owner: Heads of Engineering	Form Approver: Global Head of Engineering	GMS Change Number: 808