

## Declaration of Conformity

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<b>Manufacturer Name and Address:</b>	Cerus Corporation 2550 Stanwell Drive Concord, CA 94520 USA
<b>EC Representative Name and Address:</b>	Cerus Europe B.V. Stationsstraat 79-D 3811 MH Amersfoort The Netherlands
<b>Product Name:</b>	INTERCEPT Illuminator
<b>Product Code or Model Number:</b>	INT100-50 INT100-60
<b>Global Medical Device Nomenclature (GMDN)</b>	44185
<b>Directive:</b>	Council Directive 93/42/EEC
<b>Medical Device Classification:</b>	Class I
<b>Conformity Assessment Route:</b>	Annex VII and IX (Rules 1 and 12)
<p>We declare that the product(s) described above meets the applicable provisions of the above-mentioned Directive. In addition, the requirements of 2007/47/EC and 2002/96/EC (WEEE) have been fulfilled. All supporting documentation is retained under the premises of the manufacturer.</p>	
<b>Standards Applied:</b>	EN 980, EN 1041, EN ISO 13485, EN ISO 14155-1, EN ISO 14971, EN ISO 15225, ISO 9001, ISO 7000, ISO 7064, EN 61000-3-2, EN 61000-3-3, EN 61010-1, EN 55011, EN 61000-4-1, EN 61000-4-2, EN 61000-4-3, EN 61000-4-4, EN 61000-4-5, EN 61000-4-6, EN 61000-4-8, EN 61000-4-11, EN 50419, EN 61326
<b>Certification Body (if applicable):</b>	Not applicable
<b>EC Certificate Number(s):</b>	Not applicable- Self declaration
<b>Date of CE Mark:</b>	21 August 2006
<p>This declaration is issued under the sole responsibility of Cerus Corporation.</p>	

