

## Declaration of Conformity

<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>	INT 100 <i>*Available in power supply 50 Hz or 60 Hz</i>
<b>Software Version(s) (if applicable)</b>	4.2.11 5.1.1 5.2 6.0.9 6.1.2
<b>Global Medical Device Nomenclature (GMDN):</b>	58075
<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)
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 have been fulfilled. All supporting documentation is retained under the premises of the manufacturer or contract manufacturer.	
<b>Harmonized Standards Applied (in full or part):</b>	EN 1041, EN ISO 3826-1, EN ISO 14155-1, EN ISO 14971, EN ISO 15225, ISO 9001, ISO 7000, ISO 7064, ISO 15223-1, ISO/IEC 14882, 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, IEC 62304
<b>Certification Body (if applicable):</b>	Not applicable
<b>EC Certificate Number(s):</b>	Not applicable- Self declaration

## Declaration of Conformity

---

<b>Date of CE Mark:</b>	02 August 2018
This declaration is issued under the sole responsibility of Cerus Corporation.	

**Place and Date  
of Issue:**

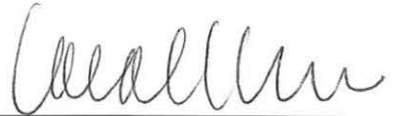
Concord, CA USA  
City

August 2, 2018  
Date

**Name and Title  
of the Issuer:**

Carol M. Moore  
(Name)

Sr. VP Regulatory  
Affairs and Quality  
(Title)

  
Signature