Declaration of Conformity

Manufacturer Name and Address:	Cerus Corporation 2550 Stanwell Drive Concord, CA 94520 USA
EC Representative Name and Address:	Cerus Europe B.V Stationsstraat 79-D 3811 MH Amersfoort The Netherlands
Product Name:	INTERCEPT Illuminator
Product Code* or Model Number:	INT 100 **Available in power supply 50 Hz or 60 Hz*
Software Version(s): (if applicable)	4.2.11 5.1.1 5.2 6.0.9 6.1.2
Global Medical Device Nomenclature (GMDN):	58075
Directive:	Council Directive 93/42/EEC
Medical Device Classification:	Class I
Conformity Assessment Route:	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.	
Harmonized Standards Applied (in full or part):	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
Certification Body (if applicable):	Not applicable
EC Certificate Number(s):	Not applicable- Self declaration

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Date of CE Mark:	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] (Signature)