# Declaration of Conformity

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| **Manufacturer Name and Address:** | Cerus Corporation<br>2550 Stanwell Drive<br>Concord, CA 94520 USA |
| **EC Representative Name and Address:** | Cerus Europe B.V.<br>Stationsstraat 79-D<br>3811 MH Amersfoort<br>The Netherlands |
| **Product Name:** | INTERCEPT Illuminator |
| **Product Code or Model Number:** | INT100-50<br>INT100-60 |
| **Global Medical Device Nomenclature (GMDN)** | 44185 |
| **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 and 2002/96/EC (WEEE) have been fulfilled. All supporting documentation is retained under the premises of the manufacturer. | |
| **Standards Applied:** | EN 1041, EN ISO 13485, 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 |
| **Date of CE Mark:** | 21 August 2006 |
| This declaration is issued under the sole responsibility of Cerus Corporation. | |

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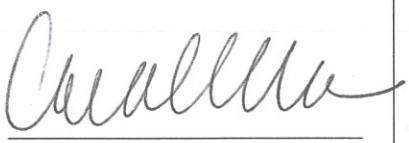
| **Place, Date of Issue** | Concord, CA USA<br>City | | August 26, 2014<br>Date |
| **Name and Title of the Issuer:** | Carol M. Moore<br>(Name) | Sr. VP, Regulatory Affairs, Quality and Clinical<br>(Title) |  <br>Signature |