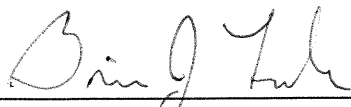


Declaration of Conformity

| PRODUCT IDENTIFICATION | | |
|--|--|---|
| Product Name | Model Number | |
| DiagnosticPRO Advantage | 16323 | |
| DosimetryPRO Advantage | 17500 | |
| CADPRO Advantage | 17032, 19094 | |
| DiagnosticPRO Edge | 19523 | |
| MANUFACTURER | | |
| Name of Company | Telephone | Representative |
| VIDAR Systems Corporation, a 3D Systems Company | 365 Herndon Parkway Herndon, VA 20170 USA | Carrie L. Brancart 703-471-7070 (phone) 703-471-1165 (fax) |
| AUTHORIZED REPRESENTATIVE | | |
| Name of Company | Address | Telephone/email |
| Emergo Europe | Prinsessegracht 20 2514 AP The Hague The Netherlands | +31.70.345.8570 - phone +31.70.346.7299 - fax europe@emergogroup.com |
| CONFORMITY ASSESSMENT | | |
| Device Classification | Route to Compliance | Standards Applied |
| Class 1, Rule 12 Non-sterile, non-measuring | Annex VII of MDD 93/42/EEC Council Directive | IEC 60601-1: (3 rd edition):2005 + CORR. 1(2006) + CORR. 2(2007) |

VIDAR Systems Corporation, a 3D Systems declares that the above-mentioned products meet the provision of the Council Directive 93/42/EEC for Medical Devices and Directive 2011/65/EU on the restriction of the use of certain hazardous substances in electrical and electronic equipment as transposed in the national laws of the Member States.

I am authorized as a company representative to sign as authority on this Declaration of Conformance:



Brian Loch, Director of R & D

12/06/2017
Date