

EC Declaration of Conformity



Document No.: GE 0176 / 03

<i>Manufacturer:</i>	Sirona Dental Systems GmbH
<i>Address:</i>	Fabrikstraße 31 64625 Bensheim Germany
<i>Product Category:</i>	Dental Unit
<i>Medical device:</i>	INTEGO / INTEGO pro
<i>Product Identification (e.g. Ref.- / Type-Number):</i>	Type D3543
<i>Classification according to Annex IX (93/42/EEC):</i>	Class II a

We declare the compliance of the medical system concerned with the requirements of the Council Directive 93/42/EEC.

Any modification to the product, not authorized by us, will invalidate this declaration.

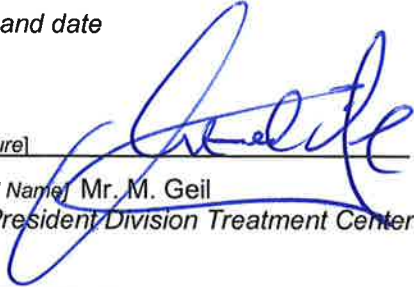

The conformity of the full quality assurance system (Directive 93/42/EEC, Annex II excluding 4) is certified by:

**TÜV SÜD Product Service GmbH
Ridlerstraße 65
80339 München
Germany**

The identification number of the notified body for implementation of the procedure is 0123.

Bensheim: September 11th, 2015

Place and date

	
<i>[Signature]</i>	<i>[Signature]</i>
<i>[Printed Name] Mr. M. Geil Vice President Division Treatment Centers</i>	<i>[Printed Name] Mr. K. Hennemann Approval Engineer</i>

*The declaration certifies the compliance according to Annex II of Directive 93/42/EEC.
Conditions of guarantee and liability are dealt within our General Conditions of Sale.*