



Product Service

# EC Certificate

## Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)  
(Devices in Class IIa, IIb or III)

No. G1 11 05 75731 015

**Manufacturer:****Limacorporate S.p.A.**

Via Nazionale, 52  
33038 Villanova San Daniele del Friuli (UDINE)  
ITALY

**Facility(ies):**

Limacorporate S.p.A.  
Via Nazionale, 52, 33038 Villanova San Daniele del Friuli  
(UDINE), ITALY

**Product****Category(ies):**

**ORTHOPEDIC IMPLANTS** (knee, hip, shoulder);  
**TRAUMA SYSTEMS** (screws, bone cerclage bands);  
Kits for bone cement;  
Instruments in class IIa for orthopaedic  
and trauma

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: ITA 213005

Valid from: 2011-05-27

Valid until: 2016-05-26

Date, 2011-05-27

Hans-Heiner Junker



TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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