



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 15 09 32687 031**

**Manufacturer:** **C.G.M. s.p.a. Divisione  
Medicale META**

Via E. Villa, 7  
42124 Reggio Emilia  
ITALY



**Facility(ies):** C.G.M. s.p.a. Divisione Medica META  
Via E. Villa, 7, 42124 Reggio Emilia, ITALY

**Product  
Category(ies):** **Bone scraper and bone grafting collector,  
set for dental osteotome, surgical dental drills,  
device for infusion and hydration of  
granulated biomaterials, membrane fixation tacks,  
membrane tacks positioner**

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.:** ITA264004

**Valid from:** 2015-10-05

**Valid until:** 2018-05-14



**Date,** 2015-10-05

Hans-Heiner Junker

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

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