

APPROVAL
EC Directive 93/42/EEC Annex VI, Article 3
Quality Assurance System Product
Medical Devices

Registration No.: ED 60024942 0001

Report No.: 28300107 002

Manufacturer: UPPERMAXI S.L.
Zamora 95, at.

08018 Barcelona
Spain

Scope: Final inspection and testing of rotating instruments
and impression and restoration materials for
dental application


Replaces Approval, Registration No.: ED 60003294 0001

Date of Expiry: 11.05.2014

The Notified Body hereby authorizes the quality management system established and applied by the company mentioned above. The requirements of Annex VI, Article 3 of the directive have been met. This approval is subject to periodic surveillance, defined by Annex VI, Article 4 of the aforementioned EC Directive, and can be used by the company with the manufacturer's declaration of conformity.

Notified Body

Cologne, 20.05.2009


Dipl.-Ing. U. Frenkert



TÜV Rheinland Product Safety GmbH - Am Grauen Stein - D-51105 Köln
Accredited by Zentralstelle der Länder für Sicherheitstechnik (ZLS) and
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten (ZLG).

Notified under No. **0197** to the EC Commission.

CE The CE marking may be used if all relevant and effective EC Directives are complied with.

CE