



Product Service

EC-CERTIFICATE

Full Quality Assurance System

(Annex II, section 3 of the Directive 93/42/EEC on Medical Devices)

No. G1 08 10 68173 001

Manufacturer: **ZIL FOR S.R.L. Società Unipersonale**

Via Fossa, 37 31051

FOLLINA (TV) – Italy

EC-Representative: **DongBang Acuprime Ltd.**

Gater House, Gater Lane, Palace Gate

Exeter

EX1 1JL

UNITED KINGDOM

Product **Progettazione e produzione di mobile ed accessori**
Category(ies): **per studi dentistici e laboratory odontotecnici**

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 products / product categories according to Annex II section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system conforms to the provisions of this Directive and is subject to periodical surveillance. For marketing of class III products an additional Annex II.4 certificate is mandatory. See also notes overleaf.

Report No.: 74916921

Valid until: 2020-11-27



Date, 2008-11-24

Hans-Heiner Junker

TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.

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