



Product Service

EC - CERTIFICATE

Production Quality Assurance System

(Annex V of the Directive 93/42/EEC on Medical Devices)

No. G2 10 05 63105 013

Manufacturer:
CA-MI S.R.L.

 Via Ugo La Malfa, 31
 43010 Pilastro (PR)
 ITALY

Facility(ies):
CA-MI S.R.L.

Via Ugo La Malfa, 31, 43010 Pilastro (PR), ITALY

**Product
Category(ies):**
**Aerosol therapy equipment,
kits for aerosol therapy and
thermal water inhaler**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective products / product categories according to Annex V, 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 IIb and III products an additional Annex III - certificate is mandatory. See also notes overleaf.

Report No.:

ITA 203959

Valid until:

2014-12-01

Date, 2010-07-15

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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