



Product Service

EC Certificate

Production Quality Assurance System

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

No. G2 16 05 63105 038

Manufacturer:**CA-MI S.R.L.**

Via Ugo La Malfa, 13
Frazione Pilastro
43013 Langhirano (PR)
ITALY

**Facility(ies):**

CA-MI S.R.L.
Via Ugo La Malfa, 13, Frazione Pilastro, 43013 Langhirano (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 devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

Report No.:

ITA267696

Valid from:

2016-07-07

Valid until:

2019-12-01

**Date,** 2016-07-07

Stefan Preiß

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

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