ZERTIFIKAT ◆ CERTIFICATE ◆ 認証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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 14 10 63105 030

Manufacturer:

CA-MI S.R.L.
Via Ugo La Malfa 13
43010 Pilastro (PR)
ITALY

Facility(ies):

CA-MI S.R.L.
Via Ugo La Malfa 13, 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 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.: ITA249277

Valid from: 2014-12-02
Valid until: 2019-12-01

603991

Date, 2014-11-20

Hans-Heiner Junker

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

TÜV SÜD Product Service GmbH · Zertifizierstelle · Ridlerstraße 65 · 80339 München · Germany

TÜV®