



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 033

<b>Manufacturer:</b>	<b>CA-MI S.R.L.</b> Via Ugo La Malfa 13 43010 Pilastro (PR) ITALY
<b>Facility(ies):</b>	CA-MI S.R.L. Via Ugo La Malfa 13, 43010 Pilastro (PR), ITALY
<b>Product Category(ies):</b>	<b>Suction Unit, Surgical Suction equipment, Breast Pump and kit accessories for electric breast pumps</b>

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



**Date,** 2014-11-20

Hans-Heiner Junker

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

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