REUSABLE JARS are equipped with screw cover with handle for easy grip, autoclavable silicone o-ring for tight seal, overflow valve system integrated in the cover, clear graduated scale in ml with 100ml or 200ml intervals, CA-MI branding. Suitable for central vacuum systems and CA-MI suction units.Available in two types of polycarbonate, Makrolon® and Apec®.

## REUSABLE JARS in MAKROLON® – FOR STEAM STERILIZATION UP TO 121 °C

| | Jar with: cover overflow valve o-ring | Jar | Cover with: overflow valve o-ring | O-ring | Suitable for: |
|---|---|---|---|---|---|
| 400 ml | RE 210301 | RE 210305 | RE 210302 | RE 210304 | EMIVAC |
| 1000 ml | RE 210001/02 | RE 210003 | RE 210352/01 | RE 210354 | ASPIRET<br>ASKIR series (all except C30 series)<br>Central Vacuum Plants |
| 2000 ml | RE 210351/01 | RE 210353 | RE 210352/01 | RE 210354 | ASPIRET - ASKIR Series<br>HOSPIVAC Series<br>Central Vacuum Plants |
| 4000 ml | RE 210006 | RE 210007 | RE 210008 | RE 210306 | HOSPIVAC Series<br>Vuoto centralizzato |
| 5000 ml | RE 210010 | RE 210013 | RE 210012 | RE 210307 | HOSPIVAC Series<br>Central Vacuum Plants |

## REUSABLE JARS in APEC® 1745 – FOR STEAM STERILIZATION UP TO 143 °C

| | Jar with: cover overflow valve o-ring | Jar | Cover with: overflow valve o-ring | O-ring | Suitable for: |
|---|---|---|---|---|---|
| 1000 ml | RE 210009 | RE 210002 | RE 210352/02 | RE 210354 | ASPIRET<br>ASKIR series (all except C30 series)<br>Central Vacuum Plants |
| 2000 ml | RE 210351/05 | RE 210353/01 | RE 210352/02 | RE 210354 | ASPIRET - ASKIR Series<br>HOSPIVAC Series<br>Central Vacuum Plants |
| 5000 ml | RE 210010/01 | RE 210013/01 | RE 210012/01 | RE 210307 | HOSPIVAC Series<br>Vuoto centralizzato |

FLOVAC® DISPOSABLE LINERS equipped with polyethylene disposable liner, hydrophobic filter, antibacterial filter, overflow system, reusable container with clear graduated scale in ml with 50ml intervals. The gelling kit is a powder inside liners with germicidal function turning the sucked liquid into a semisolid mass, preventing cross-contamination risks of staff in charge for hygiene and waste disposal.

## FLOVAC® DISPOSABLE LINERS & CONTAINERS

| | Liner with: cover filter | Liner with: cover filter gelling kit | Reusable Container | Suitable for |
|---|---|---|---|---|
| 1000 ml | 31848 | 31858 | 31843** | ASPIRET<br>ASKIR series (all except C30 series) |
| 1000 ml | 31845 | 31854 | 31840 | Central Vacuum Plants |
| 2000 ml | 31846 | 31855 | 31841 | ASPIRET - ASKIR Series<br>HOSPIVAC Series<br>Central Vacuum Plants |
| 3000 ml | 31847 | 31856 | 31842 | HOSPIVAC Series<br>Central Vacuum Plants |

** Required additional round spacer (SP.0220) when ordered for ASKIR Series

# SPARE ACCESSORIES

| | ASPIRET<br>ASKIR 20<br>ASKIR 30<br>ASKIR 230-12V BR<br>ASKIR 30 12V | ASKIR 36 BR<br>ASKIR 36 LI-ION<br>ASKIR 118<br>ASKIR 118 BASIC | EMIVAC | ASKIR C30<br>ASKIR C30 BR | HOSPIVAC 350<br>HOSPIVAC 400<br>HOSPIVAC BR |
|---|---|---|---|---|---|
| **SET of silicone TUBES, FILTERS and CONICAL CONNECTORS** | | | | | |
|  Tube Ø 6x10mm Conical connector | RE 210355 | | RE 210355/01 | | |
| Tube Ø 6x10mm Conical connector Antibacterial filter | SP 0036 | | SP 0043 | | |
|  Tube Ø 8 x 14 mm Conical connector | | RE 210355/03 | | RE 210355/03 | RE 210355/03 |
|  Tube Ø 8 x 14 mm Conical connector Antibacterial filter | | SP 0036/02 | | SP 0036/02 | SP 0032/01 (for 350 and BR)<br>SP 0032 (for Hospivac 400) |
|  FLOVAC® liners Tube Ø 6x10mm Conical connector | SP 0158/01 | | | | |
| FLOVAC® liners Tube Ø 8x14mm Conical connector | | SP 0160/01 | | SP 0160/01 | SP 0160/01 |
| Roll of silicone tube Ø 6x10 mm | Length 1m = SP 0045/02 - Length 10m = SP 0045/03 - Length 50m = SP 0045/04 | | | | |
| Roll of silicone tube Ø 8X14 mm | Length 1m = SP 0045/05 - Length 10m = SP 0045/06 - Length 50m = SP 0045/07 | | | | |
| **MALE CONNECTORS** | | | | | |
|  Ø 8-9-10 mm (pack of 5's) | SP 0223 | SP 0223 | | SP 0223 | SP 0223 |
| **CONICAL CONNECTORS** | | | | | |
|  Ø 8-9-10 mm | RE 210410 | | RE 210410 | | |
| Ø 10-11-12 mm | | RE 210420 | | RE 210420 | RE 210420 |
| **FILTERS (Antibacterial and Hydrophobic)** | | | | | |
|  Ø 64 with 8 mm connector | SP 0046 | | SP 0046 | | |
| Ø 64 with 11mm connector | | SP 0121 | | SP 0121 | SP 0121 (350 and BR only) |
| Ø 90 with 11mm connector | | | | | SP 0047 (for 400 only) |
| **ASPIRATION PROBES** | | | | | |
|  CH20 | RE 210400 (10 pcs) | RE 210400 (10 pcs) | | RE 210400 (10 pcs) | |
| **YANKAUER CANNULAS** | | | | | |
|  Yankauer Handle Flat Tip with Hole | 2044403 | 2044403 | 2044403 | 2044403 | 2044403 |
|  Yankauer Handle Crown Tip with Hole | 2044401 | 2044401 | 2044401 | 2044401 | 2044401 |
|  Yankauer Tube L= 180 | 204413018 | 204413018 | 204413018 | 204413018 | 204413018 |
| **CATHETER CONTAINER** | | | | | |
|  Tube of polycarbonate Ø 54 mm by 400 mm length. Fully autoclavable (121°C - 15 min) | | | | | 000032 |
| **SILICONE FETAL VACUUM CUPS** | | | | | |
|  Length 210 mm, Ø 50 mm, size XS | | | | VC-95100 | VC-95100 |
| Length 210 mm, Ø 60 mm, size S | | | | VC-95200 | VC-95200 |
| Length 210 mm, Ø 70 mm, size M | | | | VC-95300 | VC-95300 |

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

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 063105 0047 Rev. 01**

**Manufacturer:**

**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, Thermal Water Inhaler, Suction Unit, Surgical Suction Equipment, Breast Pump, Kit Accessory for Electric Breast Pump, Blood Pressure Monitor, Electronic Thermometer, Infrared Thermometer, Tens Device, Pulse Oximeter**

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. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:G2 063105 0047 Rev. 01

**Report No.:** ITA1626749

**Valid from:** 2021-02-09
**Valid until:** 2024-05-26

**Date,** 2021-02-09

Christoph Dicks
Head of Certification/Notified Body


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

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

DAkkS Deutsche Akkreditierungsstelle D-ZM-11321-01-00

TÜV SÜD Product Service

# Certificate

## No. Q5 063105 0045 Rev. 02

**Holder of Certificate:**

**CA-MI S.R.L.**
Via Ugo La Malfa, 13
Frazione Pilastro
43013 Langhirano (PR)
ITALY

**Certification Mark:**

TÜV SÜD EN ISO 13485 tuv-sud.com/ps-cert

**Scope of Certificate:**

**Design and development, production, sale and after-sales technical assistance of medical equipments for surgery (electrical and manual suction pumps), electric breast pumps, medical devices for breathing (aerosol therapy equipments and thermal water inhaler), medical devices for stimulation (tens) and related accessories. Placing on the market under its own name of devices for monitoring of vital physiological parameters (pulse oximeters, thermometers, blood pressure monitors, sphygmomanometer and stethoscopes), incentive spirometers, devices for phlebology (graduated compression medical stockings) and anti-decubitus mattress.**
**Distribution of active and non-active non implantable medical devices.**

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:Q5 063105 0045 Rev. 02

**Report No.:** ITA1620011

**Valid from:** 2021-03-25
**Valid until:** 2022-07-31

**Date,** 2021-03-25

Christoph Dicks
Head of Certification/Notified Body


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DAkkS Deutsche Akkreditierungsstelle D-ZM-11321-01-00

TÜV SÜD Product Service

# Certificate

## No. Q5 063105 0045 Rev. 02

**Applied Standard(s):**

EN ISO 13485:2016
Medical devices - Quality management systems - Requirements for regulatory purposes
(ISO 13485:2016)
DIN EN ISO 13485:2016

**Facility(ies):**

CA-MI S.R.L.
Via Ugo La Malfa, 13, Frazione Pilastro, 43013 Langhirano (PR), ITALY

**Design and development, production, sale and after-sales technical assistance of medical equipments for surgery (electrical and manual suction pumps), electric breast pumps, medical devices for breathing (aerosol therapy equipments and thermal water inhaler), medical devices for stimulation (tens) and related accessories. Placing on the market under its own name of devices for monitoring of vital physiological parameters (pulse oximeters, thermometers, blood pressure monitors, sphygmomanometer and stethoscopes), incentive spirometers, devices for phlebology (graduated compression medical stockings) and anti-decubitus mattress. Distribution of active and non-active non implantable medical devices.**

CA-MI S.r.l.
Via Strada per Parma 34, Frazione Pilastro, 43013 Langhirano (PR), ITALY

**Warehouse of active and non-active non implantable medical devices and components used in production.**

CA-MI S.r.l.
Via Ugo La Malfa 27, Frazione Pilastro, 43013 Langhirano (PR), ITALY

**Production of medical devices for surgery (electrical and manual suction pumps), warehouse of active and non-active non implantable medical devices and components used in production.**

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