

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 ASKIR series (all except C30 series) Central Vacuum Plants
2000 ml	RE 210351/01	RE 210353	RE 210352/01	RE 210354	ASPIRET - ASKIR Series HOSPIVAC Series Central Vacuum Plants
4000 ml	RE 210006	RE 210007	RE 210008	RE 210306	HOSPIVAC Series Vuoto centralizzato
5000 ml	RE 210010	RE 210013	RE 210012	RE 210307	HOSPIVAC Series 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 ASKIR series (all except C30 series) Central Vacuum Plants
2000 ml	RE 210351/05	RE 210353/01	RE 210352/02	RE 210354	ASPIRET - ASKIR Series HOSPIVAC Series Central Vacuum Plants
5000 ml	RE 210010/01	RE 210013/01	RE 210012/01	RE 210307	HOSPIVAC Series 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			31843**	ASPIRET ASKIR series (all except C30 series)	
	1000 ml	31845			31854	31840	Central Vacuum Plants
	2000 ml	31846			31855	31841	ASPIRET - ASKIR Series HOSPIVAC Series Central Vacuum Plants
	3000 ml	31847			31856	31842	HOSPIVAC Series Central Vacuum Plants

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

Support ring for FLOVAC® container (all sizes) with 25x5 slide	970010210	1
Support ring for FLOVAC® container (all sizes) with 30x5 slide	970010220	
Support ring for FLOVAC® container (all sizes) with 41x4 slide (also suitable for BAXTER wall slide)	970010214	
Support ring for FLOVAC® container with 45x5 slide (also suitable for ABBOT wall slide)	970010215	
Support ring for autoclavable jars (1000 and 2000 ml) with 25x5 slide	920200422	2
ON-OFF tap for mounting on FLOVAC® container support ring	000010057	
ON-OFF tap with control vacuum gauge for mounting on FLOVAC® container support ring	000010056	3
Plastic wall slide 25x5	920200004	
Plastic wall slide 30x5	920200029	
Plastic wall slide 45x5	920200477	4
Metal clamp (chrome-plated) for 30x10 rail with slide 25x5	000230000	
Plastic clamp (technopolymer) for 30x10 rail with slide 25x5	000230500	5
4-place trolley for FLOVAC® container with support ring, ON-OFF tap and control vacuum gauge	000260950	
4-place trolley for FLOVAC® container with support ring, ON-OFF tap and control vacuum regulator	000260955	
4-place trolley for FLOVAC® with support ring	000260953	
2-place trolley for FLOVAC® container with support ring, ON-OFF tap and control vacuum gauge	000260951	
2-place trolley for FLOVAC® container with support ring, ON-OFF tap and control vacuum regulator	000260956	
2-place trolley for FLOVAC® with support ring	000260954	6
Tandem tube Ø 8x11 L=0.38 m for cascade connection	970010120	
Disposable vacuum breaker	000320000	7
Disposable vacuum breaker with 1.8m hose	000320010	8
Specimen container	000036100	9



	ASPIRET ASKIR 20 ASKIR 30 ASKIR 230-12V BR ASKIR 30 12V	ASKIR 36 BR ASKIR 36 LI-ION ASKIR 118 ASKIR 118 BASIC	EMIVAC	ASKIR C30 ASKIR C30 BR	HOSPIVAC 350 HOSPIVAC 400 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
	Tube Ø 8 x 14 mm Conical connector Antibacterial filter		SP 0036/02	SP 0036/02	SP 0032/01 (for 350 and 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
	Yankauer Handle Crown Tip with Hole	2044401	2044401	2044401	2044401
	Yankauer Tube L= 180	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



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



Certificate

No. Q5 063105 0045 Rev. 03

Holder of Certificate:



CA-MI S.R.L.

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

Certification Mark:



Scope of Certificate:

Design and development, production, service and sale of medical equipments for surgery (electrical and manual suction pumps), electric breast pumps, medical devices for breathing (aerosol therapy equipments and thermal water inhaler) and related accessories, medical devices for monitoring of vital physiological parameters (pulse oximeters, thermometers, blood pressure monitors, sphygmomanometer and stethoscopes), incentive spirometers and medical devices for phlebology (graduated compression medical stockings). 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. 03](http://www.tuvsud.com/ps-cert?q=cert:Q5_063105_0045_Rev_03)

Report No.: ITA1885389

Valid from: 2022-08-02

Valid until: 2025-08-01

Date, 2022-08-02



Christoph Dicks

Head of Certification/Notified Body

Certificate

No. Q5 063105 0045 Rev. 03

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, service and sale of medical equipments for surgery (electrical and manual suction pumps), electric breast pumps, medical devices for breathing (aerosol therapy equipments and thermal water inhaler) and related accessories, medical devices for monitoring of vital physiological parameters (pulse oximeters, thermometers, blood pressure monitors, sphygmomanometer and stethoscopes), incentive spirometers and medical devices for phlebology (graduated compression medical stockings). 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.

/



Italia

CERTIFICATO

Nr. 50 100 7022 Rev.008

Si attesta che / This is to certify that

IL SISTEMA QUALITÀ DI
THE QUALITY SYSTEM OF

CA-MI S.r.l.

SEDE LEGALE: / REGISTERED OFFICE:
**VIA UGO LA MALFA 13 - FRAZIONE PILASTRO
IT - 43013 LANGHIRANO (PR)**

SEDI OPERATIVE: **VEDI ALLEGATO 1**
OPERATIONAL SITES: **SEE ANNEX 1**

È CONFORME AI REQUISITI DELLA NORMA
HAS BEEN FOUND TO COMPLY WITH THE REQUIREMENTS OF

UNI EN ISO 9001:2015

QUESTO CERTIFICATO È VALIDO PER IL SEGUENTE CAMPO DI APPLICAZIONE
THIS CERTIFICATE IS VALID FOR THE FOLLOWING SCOPE

Progettazione e sviluppo, produzione, assistenza tecnica e commercializzazione di dispositivi medici per chirurgia (aspiratori elettrici e non), tiralatte elettrici, dispositivi medici per respirazione (aerosolterapia, inalatori termali) e relativi accessori, dispositivi medici per il monitoraggio di parametri fisiologici vitali (pulsossimetri, termometri, misuratori della pressione elettronica e non), dispositivi medici per la ginnastica respiratoria e dispositivi medici per flebologia (calze medicali a compressione graduata). Distribuzione di dispositivi medici attivi e non attivi non impiantabili (IAF 19, 12, 29)

Design and development, production, service and sale of medical equipment for surgery (electrical and manual suction pumps), electric breast pumps, medical devices for breathing (aerosol therapy equipment and thermal water inhaler) and related accessories, medical devices for monitoring of vital physiological parameters (pulse oximeters, thermometers, blood pressure monitors, sphygmomanometer and stethoscopes), incentive spirometers and medical devices for phlebology (graduated compression medical stockings). Distribution of active and non-active non implantable medical devices (IAF 19, 12, 29)



SGQ N° 049A

Membro degli Accordi di Mutuo Riconoscimento
EA, IAF e ILAC
Signatory of EA, IAF and ILAC Mutual
Recognition Agreements

Per l'Organismo di Certificazione
For the Certification Body
TÜV Italia S.r.l.

Validità / Validity

Dal / From: **2022-08-01**

Al / To: **2025-07-31**

Francesco Scarlata

Direttore Divisione Business Assurance
Business Assurance Division Manager

Data emissione /
Issuing Date

2022-07-31

PRIMA CERTIFICAZIONE / FIRST CERTIFICATION: 2007-09-03

"LA VALIDITÀ DEL PRESENTE CERTIFICATO È SUBORDINATA A SORVEGLIANZA PERIODICA A 12 MESI E AL RIESAME COMPLETO DEL SISTEMA DI GESTIONE AZIENDALE CON PERIODICITÀ TRIENNALE"
"THE VALIDITY OF THE PRESENT CERTIFICATE DEPENDS ON THE ANNUAL SURVEILLANCE EVERY 12 MONTHS AND ON THE COMPLETE REVIEW OF COMPANY'S MANAGEMENT SYSTEM AFTER THREE-YEARS"



Italia

ALLEGATO 1 AL CERTIFICATO NR 50 100 7022 Rev.008**ANNEX 1 TO CERTIFICATE NO 50 100 7022 Rev.008**

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IL CERTIFICATO NR 50 100 7022 (ultima revisione applicabile) COPRE ANCHE LE SEGUENTI SEDI OPERATIVE
THE CERTIFICATE N 50 100 7022 (last version) COVERS ALSO THE FOLLOWING OFFICES:

CA-MI S.r.l.

VIA UGO LA MALFA 13 - FRAZIONE PILASTRO
 IT - 43013 LANGHIRANO (PR)

Progettazione e sviluppo, produzione, assistenza tecnica e commercializzazione di dispositivi medici per chirurgia (aspiratori elettrici e non), tiralatte elettrici, dispositivi medici per respirazione (aerosolterapia, inalatori termali) e relativi accessori, dispositivi medici per il monitoraggio di parametri fisiologici vitali (pulsossimetri, termometri, misuratori della pressione elettronici e non), dispositivi medici per la ginnastica respiratoria e dispositivi medici per flebologia (calze medicali a compressione graduata). Distribuzione di dispositivi medici attivi e non attivi non impiantabili

Design and development, production, service and sale of medical equipment for surgery (electrical and manual suction pumps), electric breast pumps, medical devices for breathing (aerosol therapy equipment and thermal water inhaler) and related accessories, medical devices for monitoring of vital physiological parameters (pulse oximeters, thermometers, blood pressure monitors, sphygmomanometer and stethoscopes), incentive spirometers and medical devices for phlebology (graduated compression medical stockings). Distribution of active and non-active non implantable medical devices

VIA STRADA PER PARMA 34 - FRAZIONE PILASTRO
 IT - 43013 LANGHIRANO (PR)

Magazzino di dispositivi medici attivi e non attivi non impiantabili e componenti utilizzati in produzione

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

VIA UGO LA MALFA 27 - FRAZIONE PILASTRO
 IT - 43013 LANGHIRANO (PR)

Produzione di dispositivi medici per chirurgia (aspiratori elettrici e manuali), magazzino di dispositivi medici attivi e non attivi non impiantabili e componenti utilizzati in produzione

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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Dal / From: **2022-08-01**Al / To: **2025-07-31**
Francesco Scarlata

Direttore Divisione Business Assurance
 Business Assurance Division Manager

Data emissione /
Issuing Date**2022-07-31****PRIMA CERTIFICAZIONE / FIRST CERTIFICATION: 2007-09-03**

"LA VALIDITÀ DEL PRESENTE CERTIFICATO È SUBORDINATA A SORVEGLIANZA PERIODICA A 12 MESI E AL RIESAME COMPLETO DEL SISTEMA DI GESTIONE AZIENDALE CON PERIODICITÀ TRIENNALE"
 "THE VALIDITY OF THE PRESENT CERTIFICATE DEPENDS ON THE ANNUAL SURVEILLANCE EVERY 12 MONTHS AND ON THE COMPLETE REVIEW OF COMPANY'S MANAGEMENT SYSTEM AFTER THREE-YEARS"